NTP TECHNICAL REPORT

ON THE

TOXICOLOGY AND CARCINOGENESIS

STUDIES OF ETHYLBENZENE

(CAS NO. 100-41-4)

IN F344/N RATS AND B6C3F₁ MICE

(INHALATION STUDIES)

NATIONAL TOXICOLOGY PROGRAM P.O. Box 12233 Research Triangle Park, NC 27709

January 1999

NTP TR 466

NIH Publication No. 99-3956

U.S. DEPARTMENT OF HEALTH AND HUMAN SERVICES
Public Health Service
National Institutes of Health

FOREWORD

The National Toxicology Program (NTP) is made up of four charter agencies of the U.S. Department of Health and Human Services (DHHS): the National Cancer Institute (NCI), National Institutes of Health; the National Institute of Environmental Health Sciences (NIEHS), National Institutes of Health; the National Center for Toxicological Research (NCTR), Food and Drug Administration; and the National Institute for Occupational Safety and Health (NIOSH), Centers for Disease Control. In July 1981, the Carcinogenesis Bioassay Testing Program, NCI, was transferred to the NIEHS. The NTP coordinates the relevant programs, staff, and resources from these Public Health Service agencies relating to basic and applied research and to biological assay development and validation.

The NTP develops, evaluates, and disseminates scientific information about potentially toxic and hazardous chemicals. This knowledge is used for protecting the health of the American people and for the primary prevention of disease.

The studies described in this Technical Report were performed under the direction of the NIEHS and were conducted in compliance with NTP laboratory health and safety requirements and must meet or exceed all applicable federal, state, and local health and safety regulations. Animal care and use were in accordance with the Public Health Service Policy on Humane Care and Use of Animals. The prechronic and chronic studies were conducted in compliance with Food and Drug Administration (FDA) Good Laboratory Practice Regulations, and all aspects of the chronic studies were subjected to retrospective quality assurance audits before being presented for public review.

These studies are designed and conducted to characterize and evaluate the toxicologic potential, including carcinogenic activity, of selected chemicals in laboratory animals (usually two species, rats and mice). Chemicals selected for NTP toxicology and carcinogenesis studies are chosen primarily on the bases of human exposure, level of production, and chemical structure. The interpretive conclusions presented in this Technical Report are based only on the results of these NTP studies. Extrapolation of these results to other species and quantitative risk analyses for humans require wider analyses beyond the purview of these studies. Selection *per se* is not an indicator of a chemical's carcinogenic potential.

Listings of all published NTP reports and ongoing studies are available from NTP Central Data Management, NIEHS, P.O. Box 12233, MD E1-02, Research Triangle Park, NC 27709 (919-541-3419). The Abstracts and other study information for 2-year studies are also available at the NTP's World Wide Web site: http://ntp-server.niehs.nih.gov.

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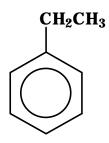
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CONTENTS

ABSTRACT		5
EXPLANATIO	ON OF LEVELS OF EVIDENCE OF CARCINOGENIC ACTIVITY	10
TECHNICAL	REPORTS REVIEW SUBCOMMITTEE	11
SUMMARY O	F TECHNICAL REPORTS REVIEW SUBCOMMITTEE COMMENTS	12
INTRODUCT	ION	15
MATERIALS	AND METHODS	21
RESULTS		29
DISCUSSION	AND CONCLUSIONS	47
REFERENCE	s	5 1
APPENDIX A	Summary of Lesions in Male Rats in the 2-Year Inhalation Study of Ethylbenzene	57
APPENDIX B	Summary of Lesions in Female Rats in the 2-Year Inhalation Study of Ethylbenzene	99
Appendix C	Summary of Lesions in Male Mice in the 2-Year Inhalation Study of Ethylbenzene	131
Appendix D	Summary of Lesions in Female Mice in the 2-Year Inhalation Study of Ethylbenzene	167
Appendix E	Genetic Toxicology	201
Appendix F	Chemical Characterization and Generation of Chamber Concentrations	21 1
Appendix G	Ingredients, Nutrient Composition, and Contaminant Levels in NIH-07 Rat and Mouse Ration	22 1
Appendix H	Sentinel Animal Program	225

ABSTRACT



ETHYLBENZENE

CAS No. 100-41-4

Chemical Formula: C₈H₁₀ Molecular Weight: 106.16

Synonyms: EB; ethylbenzol; phenylethane

Ethylbenzene is mainly used in the manufacture of styrene. Ethylbenzene is also a major component of mixed xylenes used as solvents in agricultural and home insecticide sprays, rubber and chemical manufacturing, and household degreasers, paints, adhesives, and rust preventives. Ethylbenzene is also used as an antiknock agent in aviation and motor fuels. Ethylbenzene was nominated for study by the National Institute for Occupational Safety and Health (NIOSH) and the Occupational Safety and Health Administration (OSHA) because of its potential for widespread human exposure and because of its structural similarity to benzene and toluene. Male and female F344/N rats and B6C3F₁ mice were exposed to ethylbenzene (greater than 99% pure) by inhalation for 2 years. Genetic toxicology studies were conducted in Salmonella typhimurium, mouse lymphoma cells, cultured Chinese hamster ovary cells, and mouse peripheral blood erythrocytes. In previously reported 13-week toxicity studies in which F344/N rats and B6C3F₁ mice were exposed to ethylbenzene by whole body inhalation exposure, no histopathologic changes were observed (NTP, 1992).

2-YEAR STUDY IN RATS

Groups of 50 male and 50 female F344/N rats were exposed to 0, 75, 250, or 750 ppm ethylbenzene by inhalation, 6 hours per day, 5 days per week, for 104 weeks.

Survival and Body Weights

Survival of male rats in the 750 ppm group was significantly less than that of the chamber controls. Mean body weights of 250 and 750 ppm males were generally less than those of the chamber controls beginning at week 20. Mean body weights of exposed groups of females were generally less than those of chamber controls during the second year of the study.

Pathology Findings

In male rats exposed to 750 ppm, the incidences of renal tubule adenoma and adenoma or carcinoma (combined) were significantly greater than the chamber control incidences. In addition, the incidence of renal tubule hyperplasia in 750 ppm males was significantly greater than that in the chamber controls.

The findings from an extended evaluation (step section) of the kidneys showed a significant increase in the incidences of renal tubule adenoma and hyperplasia in 750 ppm males and females; the incidence of renal tubule adenoma or carcinoma (combined) was significantly increased in 750 ppm males. The severities of nephropathy in 750 ppm male and all exposed female rats were significantly increased relative to the chamber controls.

The incidence of interstitial cell adenoma in the testis of 750 ppm males was significantly greater than that in the chamber control group and slightly exceeded the historical control range for inhalation studies.

2-YEAR STUDY IN MICE

Groups of 50 male and 50 female $B6C3F_1$ mice were exposed to 0, 75, 250, or 750 ppm ethylbenzene by inhalation, 6 hours per day, 5 days per week, for 103 weeks.

Survival and Body Weights

Survival of exposed groups of male and female mice was similar to that of the chamber controls. Mean body weights of female mice exposed to 75 ppm were greater than those of the chamber controls from week 72 until the end of the study.

Pathology Findings

In 750 ppm males, the incidences of alveolar/bronchiolar adenoma and alveolar/bronchiolar adenoma or carcinoma (combined) were significantly greater than those in the chamber control group but were within the NTP historical control ranges. The incidence of alveolar epithelial metaplasia in 750 ppm males was significantly greater than that in the chamber controls.

In 750 ppm females, the incidences of hepatocellular adenoma and hepatocellular adenoma or carcinoma (combined) were significantly greater than those in the chamber control group but were within the historical control ranges. The incidence of eosinophilic foci in 750 ppm females was significantly increased compared to that in the chamber controls. There was a spectrum of nonneoplastic liver changes related to ethylbenzene exposure in male mice, including syn-

cytial alteration of hepatocytes, hepatocellular hypertrophy, and hepatocyte necrosis.

The incidences of hyperplasia of the pituitary gland pars distalis in 250 and 750 ppm females and the incidences of thyroid gland follicular cell hyperplasia in 750 ppm males and females were significantly increased compared to those in the chamber control groups.

GENETIC TOXICOLOGY

Ethylbenzene gave little indication of mutagenicity, in vitro or in vivo. No induction of mutations was noted in Salmonella typhimurium strain TA97, TA98, TA100, or TA1535 with or without S9 metabolic activation, and no increases in sister chromatid exchanges or chromosomal aberrations were observed in cultured Chinese hamster ovary cells treated with ethylbenzene, with or without S9. In the mouse lymphoma assay, a significant mutagenic response was noted in the absence of S9, but only at the highest nonlethal dose tested and with accompanying cytotoxicity; the test was not performed with S9. increases in the frequency of micronucleated erythrocytes were observed in vivo in peripheral blood samples from male and female mice exposed to ethylbenzene for 13 weeks.

CONCLUSIONS

Under the conditions of these 2-year inhalation studies, there was clear evidence of carcinogenic activity* of ethylbenzene in male F344/N rats based on increased incidences of renal tubule neoplasms. The incidences of testicular adenoma were also increased. There was some evidence of carcinogenic activity of ethylbenzene in female F344/N rats based on increased incidences of renal tubule adenomas. There was some evidence of carcinogenic activity of ethylbenzene in male $B6C3F_1$ mice based on increased incidences of alveolar/bronchiolar neoplasms. There was some evidence of carcinogenic activity of ethylbenzene in female $B6C3F_1$ mice based on increased incidences of hepatocellular neoplasms.

Exposure of male and female rats to ethylbenzene resulted in increased incidences of renal tubule

hyperplasia and increased severities of nephropathy. Exposure of male mice to ethylbenzene resulted in increased incidences of alveolar epithelial metaplasia, syncytial alteration of hepatocytes, hepatocellular hypertrophy, hepatocyte necrosis, and thyroid gland

follicular cell hyperplasia. In female mice, ethylbenzene exposure resulted in increased incidences of eosinophilic foci of the liver, pituitary gland pars distalis hyperplasia, and thyroid gland follicular cell hyperplasia.

^{*} Explanation of Levels of Evidence of Carcinogenic Activity is on page 10. A summary of the Technical Reports Review Subcommittee comments and the public discussion on this Technical Report appears on page 12.

Summary of the 2-Year Carcinogenesis and Genetic Toxicology Studies of Ethylbenzene

	Male F344/N Rats	Female F344/N Rats	Male B6C3F ₁ Mice	Female B6C3F ₁ Mice
Concentrations in air	Chamber control, 75, 250, or 750 ppm	Chamber control, 75, 250, or 750 ppm	Chamber control, 75, 250, or 750 ppm	Chamber control, 75, 250, or 750 ppm
Body weights	250 and 750 ppm groups less than chamber controls	Exposed groups less than chamber controls	Exposed groups similar to chamber controls	75 ppm group greater than chamber controls
Survival rates	15/50, 14/50, 13/50, 2/50	31/50, 31/50, 34/50, 35/49	28/50, 36/50, 32/50, 30/50	35/50, 38/50, 40/50, 37/50
Nonneoplastic effects	Kidney: renal tubule hyperplasia (standard evaluation - 2/50, 2/50, 4/50, 12/50; standard and extended evaluations combined - 11/50, 9/50, 11/50, 23/50); severity of nephropathy (2.3, 2.4, 2.3, 3.5)	Kidney: renal tubule hyperplasia (standard evaluation - 0/50, 1/50, 3/50, 3/49; standard and extended evaluations combined - 1/50, 2/50, 4/50, 10/49); severity of nephropathy (1.3, 1.6, 1.7, 2.3)	Lung: alveolar epithelial metaplasia (0/50, 1/50, 2/50, 6/50) Liver: syncytial alteration (0/50, 5/50, 8/50, 23/50); hypertrophy (1/50, 0/50, 0/50, 17/50); necrosis (1/50, 1/50, 3/50, 10/50) Thyroid gland: follicular cell hyperplasia (21/50, 21/50, 29/50, 32/50)	Liver: eosinophilic focus (5/50, 7/50, 6/50, 22/50) Pituitary gland (pars distalis): hyperplasia (10/48, 12/49, 23/47, 22/49) Thyroid gland: follicular cell hyperplasia (18/50, 23/50, 25/50, 35/50)
Neoplastic effects	Kidney: renal tubule adenoma (standard evaluation - 0/50, 3/50, 2/50, 4/50; standard and extended evaluations combined - 3/50, 5/50, 7/50, 20/50); renal tubule adenoma or carcinoma (standard evaluation - 0/50, 3/50, 3/50, 7/50; standard and extended evaluations combined - 3/50, 5/50, 8/50, 21/50) Testes: adenoma (36/50, 33/50, 40/50, 44/50)	Kidney: renal tubule adenoma (standard evaluation - 0/50, 0/50, 0/50, 1/49; standard and extended evaluations combined - 0/50, 0/50, 1/50, 8/49)	Lung: alveolar/bronchiolar adenoma (5/50, 9/50, 10/50, 16/50); alveolar/bronchiolar adenoma or carcinoma (7/50, 10/50, 15/50, 19/50)	<u>Liver</u> : hepatocellular adenoma (6/50, 9/50, 12/50, 16/50); hepatocellular adenoma or carcinoma (13/50, 12/50, 15/50, 25/50)

Summary of the 2-Year Carcinogenesis and Genetic Toxicology Studies of Ethylbenzene (continued)

	Male F344/N Rats	Female F344/N Rats	Male B6C3F ₁ Mice	Female B6C3F ₁ Mice		
Level of evidence of carcinogenic activity	carcinogenic		Some evidence	Some evidence		
Genetic toxicology						
Salmonella typhimu	rium gene mutations:	Negative in strains TA97, TA98, TA100, and TA1535 with and without S9				
Mouse lymphoma g Sister chromatid ex		Positive	without S9			
	ese hamster ovary cells in vitro:	Negative				
Cultured Chin	ese hamster ovary cells in vitro:	Negative	e with and without S9			
Micronucleated ery Mouse peripho	throcytes eral blood <i>in vivo</i> :	Negative	Negative			

EXPLANATION OF LEVELS OF EVIDENCE OF CARCINOGENIC ACTIVITY

The National Toxicology Program describes the results of individual experiments on a chemical agent and notes the strength of the evidence for conclusions regarding each study. Negative results, in which the study animals do not have a greater incidence of neoplasia than control animals, do not necessarily mean that a chemical is not a carcinogen, inasmuch as the experiments are conducted under a limited set of conditions. Positive results demonstrate that a chemical is carcinogenic for laboratory animals under the conditions of the study and indicate that exposure to the chemical has the potential for hazard to humans. Other organizations, such as the International Agency for Research on Cancer, assign a strength of evidence for conclusions based on an examination of all available evidence, including animal studies such as those conducted by the NTP, epidemiologic studies, and estimates of exposure. Thus, the actual determination of risk to humans from chemicals found to be carcinogenic in laboratory animals requires a wider analysis that extends beyond the purview of these studies.

Five categories of evidence of carcinogenic activity are used in the Technical Report series to summarize the strength of the evidence observed in each experiment: two categories for positive results (**clear evidence** and **some evidence**); one category for uncertain findings (**equivocal evidence**); one category for no observable effects (**no evidence**); and one category for experiments that cannot be evaluated because of major flaws (**inadequate study**). These categories of interpretative conclusions were first adopted in June 1983 and then revised in March 1986 for use in the Technical Report series to incorporate more specifically the concept of actual weight of evidence of carcinogenic activity. For each separate experiment (male rats, female rats, male mice, female mice), one of the following five categories is selected to describe the findings. These categories refer to the strength of the experimental evidence and not to potency or mechanism.

- Clear evidence of carcinogenic activity is demonstrated by studies that are interpreted as showing a dose-related (i) increase of malignant neoplasms, (ii) increase of a combination of malignant and benign neoplasms, or (iii) marked increase of benign neoplasms if there is an indication from this or other studies of the ability of such tumors to progress to malignancy.
- Some evidence of carcinogenic activity is demonstrated by studies that are interpreted as showing a chemical-related increased
 incidence of neoplasms (malignant, benign, or combined) in which the strength of the response is less than that required for clear
 evidence.
- Equivocal evidence of carcinogenic activity is demonstrated by studies that are interpreted as showing a marginal increase of neoplasms that may be chemical related.
- No evidence of carcinogenic activity is demonstrated by studies that are interpreted as showing no chemical-related increases in malignant or benign neoplasms.
- **Inadequate study** of carcinogenic activity is demonstrated by studies that, because of major qualitative or quantitative limitations, cannot be interpreted as valid for showing either the presence or absence of carcinogenic activity.

When a conclusion statement for a particular experiment is selected, consideration must be given to key factors that would extend the actual boundary of an individual category of evidence. Such consideration should allow for incorporation of scientific experience and current understanding of long-term carcinogenesis studies in laboratory animals, especially for those evaluations that may be on the borderline between two adjacent levels. These considerations should include:

- adequacy of the experimental design and conduct;
- occurrence of common versus uncommon neoplasia;
- progression (or lack thereof) from benign to malignant neoplasia as well as from preneoplastic to neoplastic lesions;
- some benign neoplasms have the capacity to regress but others (of the same morphologic type) progress. At present, it is impossible to identify the difference. Therefore, where progression is known to be a possibility, the most prudent course is to assume that benign neoplasms of those types have the potential to become malignant;
- · combining benign and malignant tumor incidence known or thought to represent stages of progression in the same organ or tissue;
- latency in tumor induction;
- · multiplicity in site-specific neoplasia;
- metastases:
- supporting information from proliferative lesions (hyperplasia) in the same site of neoplasia or in other experiments (same lesion in another sex or species);
- presence or absence of dose relationships;
- statistical significance of the observed tumor increase;
- concurrent control tumor incidence as well as the historical control rate and variability for a specific neoplasm;
- survival-adjusted analyses and false positive or false negative concerns;
- structure-activity correlations; and
- in some cases, genetic toxicology.

NATIONAL TOXICOLOGY PROGRAM BOARD OF SCIENTIFIC COUNSELORS TECHNICAL REPORTS REVIEW SUBCOMMITTEE

The members of the Technical Reports Review Subcommittee who evaluated the draft NTP Technical Report on ethylbenzene on 11 and 12 December 1996 are listed below. Subcommittee members serve as independent scientists, not as representatives of any institution, company, or governmental agency. In this capacity, subcommittee members have five major responsibilities in reviewing the NTP studies:

- · to ascertain that all relevant literature data have been adequately cited and interpreted,
- · to determine if the design and conditions of the NTP studies were appropriate,
- to ensure that the Technical Report presents the experimental results and conclusions fully and clearly,
- to judge the significance of the experimental results by scientific criteria, and
- to assess the evaluation of the evidence of carcinogenic activity and other observed toxic responses.

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SUMMARY OF TECHNICAL REPORTS REVIEW SUBCOMMITTEE COMMENTS

On 11 and 12 December 1996, the draft Technical Report on the toxicology and carcinogenesis studies of ethylbenzene received public review by the National Toxicology Program's Board of Scientific Counselors' Technical Reports Review Subcommittee. The review meeting was held at the National Institute of Environmental Health Sciences, Research Triangle Park, NC.

Dr. P.C. Chan, NIEHS, introduced the toxicology and carcinogenesis studies of ethylbenzene by discussing the uses of the chemical and the rationale for study, describing the experimental design, reporting on survival and body weight effects, and commenting on compound-related neoplastic and nonneoplastic lesions in rats and mice. The proposed conclusions were *clear evidence of carcinogenic activity* in male F344/N rats and *some evidence of carcinogenic activity* in female F344/N rats and male and female B6C3F₁ mice.

Dr. Reddy, a principal reviewer, agreed with the proposed conclusions. He said that for the purpose of contrasting findings with those of Maltoni *et al.* (1985), the Technical Report should cite information on types, sites, and incidences of neoplasms from that study. Dr. Chan said that in that study, the total number of neoplasms was provided but not differentiated by target organ. Dr. Reddy noted that the methods, such as immunochemistry, used to rule out $\alpha 2\mu$ -globulin nephropathy in male rats should be described in the Technical Report. Dr. J. Mahler, NIEHS, responded that the hematoxylin-eosin stain, a good screen for hyaline droplet accumulation, was used.

Dr. Goldsworthy, the second principal reviewer, agreed with the proposed conclusions for rats and female mice. He agreed that the inhalation route was appropriate, but he noted that ethylbenzene has been detected in surface and ground water. Dr. Goldsworthy thought that the additional information obtained from renal step sections was helpful but asked for justification of the decision to step section kidneys but not other organs, such as thyroid and pituitary glands. Dr. J.R. Hailey, NIEHS, said that

the major reason to step section organs is to help interpret equivocal or uncertain effects, and that endocrine organs such as thyroid and pituitary glands are too small to step section. Dr. Goldsworthy suggested that *clear evidence of carcinogenic activity* may have been a better call in male mice, based on a positive exposure-response trend and the presence of metaplasia in the target tissue. Dr. Mahler said that metaplasia is an unusual lesion and is generally not recognized as a precursor to neoplasia.

Dr. Ryan, the third principal reviewer, agreed with the proposed conclusions for rats. She said that one of the reasons for studying the chemical was its structural similarity to benzene and toluene, and she questioned why the Technical Report did not include more discussion comparing the toxic effects of the three chemicals (see Table 12, page 49). expressed concern that the 750 ppm exposure in female rats and in male and female mice may have been too low because there were no survival or body weight effects in these groups. Dr. J.R. Bucher, NIEHS, commented that prechronic studies were performed with ethylbenzene and that an NTP study report was published in 1992. Because there were essentially no histopathologic findings in the 13-week studies, the exposure selection for the 2-year study was based on a body weight deficit in male rats. Dr. Ryan said that it could be argued that there was clear evidence of carcinogenic activity in male mice based on an exposure-related increase in combined benign and malignant lung neoplasms and in female mice based on an exposure-related increase of combined benign and malignant hepatic neoplasms. Dr. J.K. Haseman. NIEHS. said that there were three reasons for the level of evidence chosen: first, the neoplasm rates fell within the historical control range; second, the neoplasms were primarily benign; and third, the lung neoplasms were seen only in males and the liver neoplasms only in females.

Dr. LeBoeuf commented that survival in 750 ppm male rats was only 4% but the level of evidence of carcinogenic activity in male rats was based on increased incidences of renal tubule neoplasms in the 750 ppm group. He said that he was uncomfortable

basing the level of evidence of carcinogenic activity on findings accompanied by such poor survival. Dr. Bucher responded that the fact that increased renal neoplasms were seen in both males and females and were accompanied by severe nephropathy, which is rarely if ever seen in females, suggests an intrinsic carcinogenic activity of ethylbenzene.

Dr. Ryan moved that the Technical Report on ethylbenzene be accepted with the revisions discussed and the conclusions as written for male rats, *clear evidence of carcinogenic activity*, and for female rats and male and female mice, *some evidence of carcinogenic activity*. Dr. Reddy seconded the motion, which was accepted unanimously with nine votes.

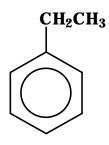
Later in the meeting, Dr. LeBoeuf made a motion to reopen the discussion on the neoplasm response in male rats. Dr. Taylor thought that the maker and seconder of the original motion should have to agree. Drs. Ryan and Reddy agreed to reopen the discussion. Dr. Goldsworthy seconded the motion to reopen the discussion, which was accepted by six yes votes to two no votes (Drs. Brown and Reddy). Dr. Ward was not present.

Dr. LeBoeuf stated that his primary concerns were the mortality in 750 ppm male rats and the interpretation of the data at that dose. He said that one of the original National Cancer Institute guidelines for the 2-year bioassays is that particular treatments should not affect survival, unless reduced survival is a result of neoplasia, and should not cause more than a 10% decrease in body weight gain. He said that in the ethylbenzene Technical Report, it was clear that

the majority of the neoplasms in male rats were considered to be incidental to the cause of death. For this reason, he recommended changing the conclusion in male rats to some evidence of carcinogenic activity. Dr. Haseman pointed out that at week 84, the survival in 750 ppm male rats was still 70%. Dr. Goldsworthy stated that one issue to consider is when the first neoplasms arose. Dr. Bucher commented that nephropathy was likely the primary contributor to mortality. Dr. Haseman suggested that the conclusion for male rats, as with the report on oxazepam, could indicate that there was clear evidence of carcinogenic activity only at concentrations resulting in enhanced nephropathy. Dr. Bucher noted that in many past studies, the conclusions for carcinogenic activity were confirmed even when the maximum tolerated doses were exceeded. He noted that in most studies in which renal tubule neoplasms are associated with nephropathy in male rats, carcinomas are generally not seen; he further noted that in female rats, the incidences of nephropathy are generally less than in male rats and that 21 neoplasms is exceptionally high. Dr. Goldsworthy reminded the reviewers of the stipulation "Under the conditions of these studies..." Dr. Ryan pointed out that in the standard evaluation, the renal tubule neoplasm incidences in male rats exceeded the historical control range even in the 75 ppm group.

Dr. LeBoeuf moved that the conclusion for male rats be changed to *some evidence of carcinogenic activity*. Dr. Ryan seconded the motion, which was defeated by six no votes to two yes votes (Drs. LeBoeuf and Russo). Dr. Ward was not present.

INTRODUCTION



ETHYLBENZENE

CAS No. 100-41-4

Chemical Formula: C₈H₁₀ Molecular Weight: 106.16

Synonyms: EB; ethylbenzol; phenylethane

CHEMICAL AND PHYSICAL PROPERTIES

Ethylbenzene is a colorless, flammable, aromatic liquid with a melting point of -95.0° C, a boiling point of 136.2° C at 760 mm Hg, and a density of 0.866 at 25° C. Its vapor pressure is 10 mm Hg at 25.9° C, and its vapor density is 3.66. It is practically insoluble in water (0.014 g/100 mL) at 15° C but is soluble in most organic solvents (Verschueren, 1983; Merck Index, 1989). Ethylbenzene has a flash point of 15° C and an autoignition temperature of 432° C (Lewis, 1992).

PRODUCTION, USE, AND HUMAN EXPOSURE

Ethylbenzene is produced by two primary processes: heating of benzene and ethylene in the presence of aluminum chloride and by fractionation directly from the mixed xylene stream during petroleum refining (Hawley's, 1987). The United States production of ethylbenzene was 7.56 billion pounds in 1984 (USITC, 1985), 8.5 billion pounds in 1986 (Heylin, 1987), 11.11 billion pounds in 1992, and 11.76 billion pounds in 1993 (Chem. Eng. News, 1994). Ethylbenzene was the eighteenth highest in production volume for chemicals produced in the United States in

1985 (*Hawley's*, 1987). Ethylbenzene is mainly used in the manufacture of styrene (*Fed. Regist.*, 1987) and cellulose acetate (ILO, 1983). It has also been used as an intermediate in the production of diethylbenzene, acetophenone, and ethyl anthraquinone. Ethylbenzene is a major component (15% to 20%) of mixed xylenes (Toftgard and Nilsen, 1982), which are used as solvents in agricultural and household insecticide sprays, rubber and chemical manufacturing industries, and household degreasing cleaners, paint, adhesives, and rust preventives (Fishbein, 1985). The United States produced 6.49 billion pounds of mixed xylenes in 1984 (USITC, 1985). Ethylbenzene has also been used in motor and aviation fuels as an antiknock agent (NIOSH, 1979; ILO, 1983).

Ethylbenzene is widely distributed in the environment due to its use as a solvent and fuel additive; it is also naturally present in crude petroleum. It has been detected in ambient air, surface water and groundwater, and in human milk (National Research Council, 1981). Ethylbenzene concentrations of 10 to 26 mg/L have been detected in the Missouri River (STORET, 1986) and concentrations up to 7 mg/L have been found in samples of potable water in Canada (Otson *et al.*, 1982). Ethylbenzene has also been found in wastewater effluents from pulpwood

mills (Nestmann *et al.*, 1980). Ethylbenzene was in a water sample from New Jersey, in eight air samples, and in 12 breath samples from workers exposed to ethylbenzene (Wallace *et al.*, 1984). Atmospheric air samples collected in the Los Angeles basin contained ethylbenzene, probably derived from vehicle exhaust (Lonneman *et al.*, 1968). No evidence of ethylbenzene bioaccumulation has been reported.

Based on irritant properties of ethylbenzene vapor, the American Conference of Governmental Industrial Hygienists (ACGIH, 1996) has set a threshold limit value of 100 ppm (435 mg/m³), with a short-term exposure limit of 125 ppm (545 mg/m³). The Occupational Safety and Health Administration (OSHA) set the permissible exposure limit at 100 ppm as an 8-hour time-weighted average and 125 ppm as a 15-minute short-term exposure limit (*Fed. Regist.*, 1989).

ABSORPTION, DISTRIBUTION, METABOLISM, AND EXCRETION

Structurally, ethylbenzene is related to other aliphatic derivatives of aromatic compounds. Many of the biological activities of these chemicals are similar. For example, benzene, ethylbenzene, and toluene are well absorbed after inhalation exposure and are distributed to adipose tissue, liver, kidney, bone marrow, and nervous tissue. These chemicals are metabolized mainly by the hepatic cytochrome P_{450} systems and are central nervous system depressants (Tegeris and Balster, 1994). The toxic effect on the central nervous system, at least in part, is exerted by inhibiting the membrane-bound ATPase activities in astrocytes (Naskali *et al.*, 1994; Vaalavirta and Tähti, 1995), thereby disturbing the ATPase-dependent astrocytic regulatory functions.

Toluene is metabolized by the liver cytochrome P_{450} enzyme system to benzyl alcohol, benzaldehyde, and benzoic acid via methyl hydroxylation and is excreted in the urine as hippuric acid. A minor pathway of metabolism is via ring hydroxylation and excretion as cresol sulphates and glucuronides (Dean, 1978). In NTP (1990a) inhalation studies, toluene was neither genotoxic nor carcinogenic. Ono *et al.* (1995) did not find toluene to be teratogenic in inhalation studies.

Benzene is metabolized primarily by the hepatic cytochrome P₄₅₀ system to benzene oxide and then rearranged to form phenol, catechol, and benzoquinones (hydroquinones) and excreted in the urine or exhaled (NTP, 1986). Alternatively, oxidation and ring opening of catechol give rise to trans, transmuconaldehyde and muconic acid. Hydration of benzene oxide to dihydrodiol and ring oxidation to diolepoxide have also been postulated (Busby et al., 1990). Inhalation exposure to benzene in BDF₁ mice caused DNA damage in peripheral blood cells, bone marrow, and liver (Plappert et al., 1994). hematotoxicity of benzene observed in rats and mice is mainly due to the metabolites hydroquinone and benzoquinone (Zhu et al., 1995). Xylene undergoes oxidation of the methyl group to give rise to methyl benzyl alcohols or aromatic hydroxylation to xylenols before excretion in the urine (Dean, 1978).

Percutaneous absorption rates of benzene, toluene, ethylbenzene, and aniline in male HRS/J hairless mice following an application of 5 mL of $^{14}\mathrm{C}$ -labeled test solution were 56, 49, 37, and 2.3 µg/cm² per minute, respectively (Susten et al., 1990). The excretion of benzene and aniline in expired air was greater during the first 15 minutes of exposure, whereas that of toluene and ethylbenzene was greatest during the second 15 minutes of exposure. These data suggested a two-compartment model might better describe the kinetics of the appearance of toluene and ethylbenzene in expired breath.

Differences in the metabolism of ethylbenzene in rats, rabbits, and humans are minor (Chin *et al.*, 1980; Climie *et al.*, 1983). Ethylbenzene metabolism appears to involve side-chain hydroxylation by liver microsomal enzymes (Pyykko *et al.*, 1987). Ring oxidation may also occur (Engström, 1984).

Experimental Animals

Ethylbenzene is readily absorbed from the atmosphere in Harlan-Wistar rats. In rats exposed to radiolabeled ethylbenzene for 6 hours by inhalation, radioactivity was found in the liver, gastrointestinal tract, and adipose tissue 42 hours after exposure (Chin *et al.*, 1980). One day following oral administration of radioactive ethylbenzene, radioactivity was found in the intestine, liver, kidney, and fat of rats (Climie

et al., 1983). Freundt et al. (1989) reported that the blood concentration of ethylbenzene was dose-dependent after a 2-hour inhalation of 120, 240, 350, or 650 ppm in rats.

In rats, ethylbenzene is metabolized to mandelic acid and phenylglyoxylic acid by side-chain oxidation and then excreted in the urine (Bardodej and Bardodejova, 1970; Engström, 1984; Gromiec and Piotrowski, 1984). Other minor metabolites found in urine included 1-phenylethanol, omega-hydroxyacetophenone, hippuric acid, benzoic acid, phenylacetic acid, and phenaceturic acid (Engström, 1984; Engström et al., 1985). Engström (1984) showed that in male Wistar rats exposed to ethylbenzene by inhalation for 6 hours per day, 5 days per week, for 3, 5, and 9 weeks at 50, 300, or 600 ppm, the total urinary elimination of ethylbenzene metabolites in 24 hours was dose dependent. Excretion of metabolites into urine increased in a dose-related manner but less than linearly. The total amount of metabolites excreted at each time point at each dose was constant. These data implied induction of a metabolic enzyme.

Humans

Human exposure to ethylbenzene is mainly via inhalation of vapor and/or mist. To a smaller extent, absorption also occurs through dermal contact or by ingestion (Dutkiewicz and Tyras, 1967). Ethylbenzene is readily absorbed from the atmosphere through the lungs in humans (Bardodej and Bardodejova, 1970; Gromiec and Piotrowski, 1984), and orally administered ethylbenzene is quickly and effectively absorbed as well (Climie et al., 1983). Absorption of liquid ethylbenzene through the skin is rapid when compared to similar hydrocarbon compounds such as benzene or styrene (Dutkiewicz and Tyras, 1967). Trace amounts of ethylbenzene were found in the subcutaneous fat (Wolf et al., 1977) and body fat (Engström and Bjurstrom, 1978) of humans exposed to the chemical either by the dermal or inhalation route.

In humans, as in rats, most of the absorbed ethylbenzene is metabolized by liver microsomal enzymes to mandelic acid and phenylglyoxylic acid by side-chain oxidation and then excreted in the urine (Bardodej and Bardodejova, 1970; Engström, 1984; Gromiec and Piotrowski, 1984). However, a small amount of phenolic derivatives (2- and 4-ethylphenol)

is also found in the urine (Angerer and Lehnert, 1979; Engström, 1984), indicating the occurrence of ring oxidation. The presence of phenaceturic acid in urine implies oxidation of the ω -methyl group of the side chain (Figure 1; Engström, 1984).

TOXICITY

Experimental Animals

The oral LD_{50} for ethylbenzene in male and female Wistar rats was estimated to be 3.5 g/kg (Wolf *et al.*, 1956), and the intraperitoneal LD_{50} for mice was 2.27 g/kg (DFG, 1985; Lewis, 1992). The 4-hour LC_{50} in female rats was 4,000 ppm, and the 1-hour LC_{50} was 8,000 ppm (Smyth *et al.*, 1962).

Ethylbenzene is a mucous membrane irritant; guinea pigs exposed for 1 minute to 0.2% ethylbenzene vapor experienced moderate eye and nasal irritation. Exposure to 0.1% ethylbenzene produced slight nasal irritation that ceased after 30 minutes. At 1%, ethylbenzene caused ataxia, loss of consciousness, tremor (Lewis, 1992), central nervous system depression, and death (ACGIH, 1986).

In Wistar rats, oral administration of ethylbenzene at 408 or 680 mg/kg per day or inhalation exposure at 1,250 or 2,200 ppm, 7 to 8 hours per day, 5 days per week for 6 months induced slight increases in kidney and liver weights and cloudy swelling of the tubular epithelium of the kidney and parenchymal cells of the liver (Wolf et al., 1956). Male Wistar rats exposed to ethylbenzene by inhalation at 300 or 600 ppm for 16 weeks exhibited increased activities of liver enzymes, including NADPH cytochrome c reductase, 7-ethoxycoumarin-O-deethylase, UDP-glucuronosyltransferase, and D-glucuronolactone dehydrogenase. Kidney 7-ethoxycoumarin-O-deethylase and UDPglucuronosyl-transferase activities were also increased (Elovaara et al., 1985). Electron microscopy showed that the cloudy swelling of the renal tubule epithelium was due to an increase in endoplasmic reticulum as a result of an adaptive response of increased microsomal enzyme activity (Elovaara et al., 1985). Rats exposed to ethylbenzene at 2,000 ppm for 3 days had increased hepatic cytochrome P_{450} and NADPH cytochrome c reductase activities (Toftgard and Nilsen, 1982). F344/N rats and B6C3F₁ mice

FIGURE 1 Metabolism of ethylbenzene as reconstructed from urinary metabolites found in rat and human urine. The thickness of the arrows represents the extent of the respective route; the broken arrows indicate that only trace amounts were found. Unclear pathways are depicted by open arrows (Reproduced from Engström, 1984).

exposed to ethylbenzene by inhalation at 382 or 782 ppm 5 days per week for 4 weeks had significantly increased absolute and relative liver weights (Cragg *et al.*, 1989). The authors concluded that the no-observed-adverse-effect-level for rats and mice was 382 ppm.

In 13-week toxicity studies performed by the NTP (1992), F344/N rats and B6C3F₁ mice were exposed to ethylbenzene by inhalation at 0, 100, 250, 500, 750, or 1,000 ppm. Signs of toxicity included increased liver, lung, and kidney weights in exposed male and female rats and increased liver weights in exposed male and female mice. No evidence of histopathologic injury was noted in these studies. No animals died, and the mean body weight gains of the exposed rats and mice did not differ from those of the respective controls. Sperm or vaginal cytology evaluations of the exposed rats and mice revealed no changes from normal. Based on the changes in organ weights, the high dose selected for the 2-year studies was 750 ppm.

Humans

Ethylbenzene is a skin, eye, and respiratory irritant and a central nervous system depressant at an atmospheric concentration of 0.2%. Human volunteers breathing 0.1% ethylbenzene vapor reported initial eye irritation which gradually decreased, while exposure to a 0.2% atmospheric concentration was accompanied by extreme irritation of the eyes, nose, and throat (Yant et al., 1930) and central nervous system depression. Symptoms of central nervous system depression included headache; nausea; weakness; dizziness; sleepiness; loss of coordination, judgment, and consciousness; and coma or death (Lewis, 1992). Erythema and inflammation of the skin developed after dermal contact (Lewis, 1992). Prolonged exposure to ethylbenzene vapor may result in leukopenia and lymphocytosis, neurofunctional disorder, and hepatitis (ILO, 1983).

REPRODUCTIVE

AND DEVELOPMENTAL TOXICITY

Ethylbenzene is embryotoxic and teratogenic. The offspring of rats exposed to ethylbenzene at 1,000 ppm, 7 hours per day, 5 days per week for 3 weeks before mating, then exposed daily through day 19 of gestation had a higher incidence of super-

numerary ribs (Hardin *et al.*, 1981). In the offspring of CFY rats exposed to ethylbenzene at 552 ppm, 24 hours per day from days 7 to 15 of gestation, retardation of skeletal development, increased incidence of supernumerary ribs, and anomalies of the uropoietic apparatus were observed (Ungvary and Tatrai, 1985). An increased rate of malformation was also found in CFLP mice exposed to ethylbenzene. Maternal toxicity reported by these investigators included increased liver, kidney, and spleen weights. Increased postimplantation loss of fetuses in dams was also observed.

CARCINOGENICITY

Maltoni *et al.* (1985) reported a study in which Sprague-Dawley rats were administered 500 mg ethylbenzene/kg per day in olive oil by gavage, 4 or 5 days per week for 104 weeks. Incidences of malignant neoplasms were 35.0% (versus 26.7% in controls) in dosed males and 45.9% (versus 22.4% in controls) in dosed females. The results of this study were considered inconclusive. No other information on the carcinogenicity of ethylbenzene in experimental animals or humans was found in the literature. Benzene, a homologue of ethylbenzene, is carcinogenic in rats and mice (NTP, 1986; Farris *et al.*, 1993) and is a human carcinogen inducing acute myelogenous leukemia and aplastic anemia.

GENETIC TOXICITY

Ethylbenzene was not mutagenic in Salmonella typhimurium strain TA97, TA98, TA100, or TA1535 when tested up to toxic doses (1,000 µg/plate) in the presence or absence of exogenous metabolic activation (S9) (Zeiger et al., 1992). It was also reported to be negative, with and without S9, in S. typhimurium strains TA1537 and TA1538 (Nestmann et al., 1980), in Escherichia coli WP2 and WP2uvrA, and in Saccharomyces cerevisiae JD1 (Dean et al., 1985). A weakly positive response was reported in a sister chromatid exchange test with human lymphocytes cultured in the presence of S9 (Norppa and Vainio, 1983), and an increase in mutant L5178Y mouse lymphoma cell colonies was observed at the highest nonlethal dose (80 µg/mL) of ethylbenzene tested in the absence of S9 (McGregor et al., 1988). Micronucleus assays in mouse peripheral blood were negative (NTP, 1992; Appendix E).

STUDY RATIONALE

Ethylbenzene was nominated for toxicity study by OSHA and NIOSH and was selected for study by the NTP because of its potential for widespread consumer exposure and its structural similarity to benzene and toluene. The present studies were undertaken following the designation of ethylbenzene as a priority chemical for toxicologic testing by the Interagency Agreement (Superfund) between the NTP and the United States Environmental Protection Agency

(EPA). The studies were designed to determine the toxicologic and carcinogenic effects of ethylbenzene in F344/N rats and B6C3F $_1$ mice after a 2-year inhalation exposure. Data were needed for the EPA to make regulatory decisions mandated by the Clean Air Act (42 U.S.C. § 7412). The inhalation route of exposure was selected because human exposure to ethylbenzene is mainly by inhalation.

MATERIALS AND METHODS

PROCUREMENT AND CHARACTERIZATION OF ETHYLBENZENE

Ethylbenzene was obtained from ARCO Chemical Company (Newtown Square, PA) in two lots (A060989 and A051890). Identity, purity, and stability analyses were conducted by the analytical chemistry laboratory, Midwest Research Institute (Kansas City, MO). Reports on analyses performed in support of the ethylbenzene studies are on file at the National Institute of Environmental Health Sciences (NIEHS).

The chemical, a clear, colorless, pungent smelling, volatile liquid, was identified as ethylbenzene by infrared, ultraviolet/visible (lot A060989 only), and nuclear magnetic resonance spectroscopy. All spectra were consistent with the literature spectra of ethylbenzene. The boiling point and density of the chemical were also consistent with literature references.

The purity of lot A060989 was determined by elemental analyses, Karl Fischer water analysis, iodometric titration for peroxide determination, and gas chromatography. Elemental analyses for carbon and hydrogen were in agreement with the theoretical values for ethylbenzene. Karl Fischer water analysis indicated less than 0.05% water. Iodometric titration revealed no peroxide. Gas chromatography by two systems revealed a major peak and no impurities with areas greater than 0.1% relative to the major peak. Major peak comparisons of lot A060989 with a previously analyzed lot of ethylbenzene (lot K061786) not used in the current studies indicated a purity of $101.0\% \pm 0.5\%$ for lot A060989 relative to lot K061786. The overall purity of lot A060989 was determined to be greater than 99%.

Additional analyses of lot A060989 were performed with gas chromatography/mass spectrometry to identify and quantify cumene in the bulk ethylbenzene. In these analyses, 62 ± 3.1 ppm cumene was detected.

The purity of lot A051890 was determined by iodometric titration for peroxide and by gas chromatography. Less than 2 ppm peroxide was detected. Gas chromatography indicated one impurity with an area of 0.1% relative to the major peak. The overall purity of lot A051890 was determined to be greater than 99%.

Accelerated stability studies of the bulk chemical were performed by the analytical chemistry laboratory. These studies indicated that ethylbenzene is stable as a bulk chemical for at least 2 weeks when stored protected from light at temperatures up to 60° C. To ensure stability, the bulk chemical was stored at room temperature in the original steel containers until just prior to use, when it was transferred to amber glass bottles with Teflon®-lined caps and a nitrogen head-space. The rapid use and small shipment sizes of ethylbenzene made stability monitoring unnecessary during the studies; however, the peroxide content of the bulk chemical was tested monthly with iodometric titration. The concentration of peroxide ranged from 1.12 to 10.7 ppm.

VAPOR GENERATION AND EXPOSURE SYSTEM

Ethylbenzene vapor was produced by flash evaporator units. Nitrogen gas carried ethylbenzene vapor from the condensing column into heated stainless-steel transfer lines that led to exposure chambers. Each exposure chamber was supplied by a separate flash evaporator unit. Exposure concentrations for individual exposure chambers were created by varying the ethylbenzene flow rate to the individual flash evaporation units. At the chamber inlets, the ethylbenzene vapor was mixed with HEPA- and charcoal-filtered Stainless-steel chambers (Hazleton H-2000®) manufactured by Lab Products, Inc. (Maywood, NJ) were used throughout the studies. The 750 ppm chambers were sampled once during the first full week of exposure for the presence of aerosol by a Quartz Crystal Microbalance Cascade Impactor

(California Measurements, Sierra Madre, CA). Results indicated that aerosol formation due to test atmosphere generation was not significant.

VAPOR CONCENTRATION MONITORING

The chamber concentrations of ethylbenzene were monitored by an on-line gas chromatograph using a flame ionization detector. Samples were drawn from supply lines leading to exposure chambers and the control chamber at least once every hour. Summaries of chamber concentrations are presented in Table F1.

CHAMBER ATMOSPHERE CHARACTERIZATION

The times for the exposure concentration to build up to 90% of the final exposure concentration (T_{90}) and to decay to 10% of the exposure concentration (T_{10}) were measured in the 750 ppm exposure chambers with animals present. At a chamber airflow rate of 15 air changes per hour, the theoretical value for both T_{90} and T_{10} is 10 minutes; analysis of chamber concentrations during the first 2 weeks of the studies indicated T_{90} and T_{10} values of 15 minutes; therefore, 15 minutes was used for the T_{90} throughout the studies.

Inhalation chambers were sampled to determine the uniformity of ethylbenzene concentrations; samples from 12 shelf positions within the exposure chambers were analyzed by gas chromatography. Chamber concentration uniformity was maintained throughout the studies.

The persistence of ethylbenzene following exposure was monitored by gas chromatography in the 750 ppm chambers with and without animals present. No ethylbenzene was detectable in the chambers 2 hours after exposure (detection limit 0.44 ppm).

The stability of ethylbenzene was monitored in the generator reservoirs of the 75 and 750 ppm chambers. No significant contaminants or degradation products were found in any of the generator reservoir samples.

Samples from occupied and unoccupied 75 and 750 ppm chambers were analyzed for degradation products before studies began, during the first week of the studies, and every 90 days thereafter. One small

impurity was detected in samples taken from the 750 ppm chambers.

2-YEAR STUDIES Study Design

Groups of 50 male and 50 female F344/N rats and $B6C3F_1$ mice were exposed by inhalation to 0, 75, 250, and 750 ppm ethylbenzene for 6 hours plus T_{90} (15 minutes) per day, 5 days per week, for 103 (mice) or 104 (rats) weeks. The high exposure concentration selected for these studies was 750 ppm ethylbenzene, roughly 19% of the 4-hour LC_{50} for rats reported by Smyth $et\ al.\ (1962)$. Following the last day of exposure, rats and mice were observed for 9 to 12 days prior to necropsy.

Source and Specification of Animals

Male and female F344/N rats and $B6C3F_1$ mice were obtained from Simonsen Laboratories, Inc. (Gilroy, CA) for use in the 2-year studies. Five male and five female rats and mice were randomly selected for parasite evaluation and gross observation of disease. Rats and mice were approximately 6 weeks old at the beginning of the studies. The health of the animals was monitored during the studies according to the protocols of the NTP Sentinel Animal Program (Appendix H).

Animal Maintenance

Rats and mice were housed individually. Feed and water were available *ad libitum*. Cages were rotated once weekly. Further details of animal maintenance are given in Table 1. Information on feed composition and contaminants is provided in Appendix G.

Clinical Examinations and Pathology

Animals were observed twice daily. Clinical findings were recorded approximately monthly. Body weights were recorded initially, weekly for the first 13 weeks, at week 16, monthly through the end of exposure, and prior to terminal necropsy. A complete necropsy and microscopic examination were performed on all rats and mice. At necropsy, all organs and tissues were examined for grossly visible lesions, and all major tissues were fixed and preserved in 10% neutral buffered formalin, processed and trimmed, embedded in paraffin, sectioned to a thickness of 5 to 6 μ m, and stained with hematoxylin and eosin for microscopic examination. For all paired organs (i.e., adrenal

gland, kidney, and ovary), samples from each organ were examined. Tissues examined microscopically are listed in Table 1.

Microscopic evaluations were completed by the study laboratory pathologist, and the pathology data were entered into the Toxicology Data Management System. The slides, paraffin blocks, and residual wet tissues were sent to the NTP Archives for inventory, slide/block match, and wet tissue audit. The slides, individual animal data records, and pathology tables were evaluated by an independent quality assessment laboratory. The individual animal records and tables were compared for accuracy, the slide and tissue counts were verified, and the histotechnique was evaluated. A quality assessment pathologist evaluated slides from all tumors and all potential target organs. which included the kidney, liver, lung, and nose of male and female rats; bone marrow, parathyroid gland, prostate gland, and testis of male rats; pituitary gland of female rats; heart, kidney, liver, lung, nose, and thyroid gland of male and female mice; and pituitary gland of female mice.

The quality assessment report and the reviewed slides were submitted to the NTP Pathology Working Group (PWG) chairperson, who reviewed the selected tissues and addressed any inconsistencies in the diagnoses made by the laboratory and quality assessment pathol-Representative histopathology slides containing examples of lesions related to chemical administration, examples of disagreements in diagnoses between the laboratory and quality assessment pathologists, or lesions of general interest were presented by the chairperson to the PWG for review. The PWG consisted of the quality assessment pathologist and other pathologists experienced in rodent toxicologic pathology. This group examined the tissues without any knowledge of dose groups or previously rendered diagnoses. When the PWG consensus differed from the opinion of the laboratory pathologist, the diagnosis was changed. Final diagnoses for reviewed lesions represent a consensus between the laboratory pathologist, reviewing pathologist(s), and the PWG. Details of these review procedures have been described, in part, by Maronpot and Boorman (1982) and Boorman et al. (1985). For subsequent analyses of the pathology data, the decision of whether to evaluate the diagnosed lesions for each tissue type separately or combined was generally based on the guidelines of McConnell et al. (1986).

STATISTICAL METHODS Survival Analyses

The probability of survival was estimated by the product-limit procedure of Kaplan and Meier (1958) and is presented in the form of graphs. Animals found dead of other than natural causes or missing were censored from the survival analyses; animals dying from natural causes were not censored. Statistical analyses for possible dose-related effects on survival used Cox's (1972) method for testing two groups for equality and Tarone's (1975) life table test to identify dose-related trends. All reported P values for the survival analyses are two sided.

Calculation of Incidence

The incidences of neoplasms or nonneoplastic lesions are presented in Tables A1, A5, B1, B4, C1, C5, D1, and D5 as the numbers of animals bearing such lesions at a specific anatomic site and the numbers of animals with that site examined microscopically. For calculation of statistical significance, the incidences of most neoplasms (Tables A3, B3, C3, and D3) and all nonneoplastic lesions are given as the numbers of animals affected at each site examined microscopically. However, when macroscopic examination was required to detect neoplasms in certain tissues (e.g., harderian gland, intestine, mammary gland, and skin) before microscopic evaluation, or when neoplasms had multiple potential sites of occurrence (e.g., leukemia or lymphoma), the denominators consist of the number of animals on which a necropsy was performed. Tables A3, B3, C3, and D3 also give the survival-adjusted neoplasm rate for each group and each site-specific neoplasm, i.e., the Kaplan-Meier estimate of the neoplasm incidence that would have been observed at the end of the study in the absence of mortality from all other competing risks (Kaplan and Meier, 1958).

Analysis of Neoplasm Incidences

The majority of neoplasms in these studies were considered to be incidental to the cause of death or not rapidly lethal. Thus, the primary statistical method used was logistic regression analysis, which assumed that the diagnosed neoplasms were discovered as the result of death from an unrelated cause and thus did not affect the risk of death. In this approach, neoplasm prevalence was modeled as a logistic function of chemical exposure and time. Both linear and quadratic terms in time were incorporated initially,

and the quadratic term was eliminated if the fit of the model was not significantly enhanced. The neoplasm incidences of exposed and control groups were compared on the basis of the likelihood score test for the regression coefficient of dose. This method of adjusting for intercurrent mortality is the prevalence analysis of Dinse and Lagakos (1983), further described and illustrated by Dinse and Haseman (1986). When neoplasms are incidental, this comparison of the time-specific neoplasm prevalences also provides a comparison of the time-specific neoplasm incidences (McKnight and Crowley, 1984).

In addition to logistic regression, other methods of statistical analysis were used, and the results of these tests are summarized in the appendixes. These methods include the life table test (Cox, 1972; Tarone, 1975), appropriate for rapidly lethal neoplasms, and the Fisher exact test and the Cochran-Armitage trend test (Armitage, 1971; Gart *et al.*, 1979), procedures based on the overall proportion of neoplasm-bearing animals.

Tests of significance included pairwise comparisons of each exposed group with controls and a test for an overall dose-related trend. Continuity-corrected tests were used in the analysis of neoplasm incidence, and reported P values are one sided. The procedures described in the preceding paragraphs were also used to evaluate selected nonneoplastic lesions. For further discussion of these statistical methods, refer to Haseman (1984).

Analysis of Nonneoplastic Lesion Incidences

Because all nonneoplastic lesions in this study were considered to be incidental to the cause of death or not rapidly lethal, the primary statistical analysis used was a logistic regression analysis in which nonneoplastic lesion prevalence was modeled as a logistic function of chemical exposure and time. For lesions detected at the interim evaluation, the Fisher exact test, a procedure based on the overall proportion of affected animals, was used.

Analysis of Continuous Variables

Average severity values were analyzed for significance with the Mann-Whitney U test (Hollander and Wolfe, 1973).

Historical Control Data

Although the concurrent control group is always the first and most appropriate control group used for evaluation, historical control data can be helpful in the overall assessment of neoplasm incidence in certain instances. Consequently, neoplasm incidences from the NTP historical control database, which is updated yearly, are included in the NTP reports for neoplasms appearing to show compound-related effects.

QUALITY ASSURANCE METHODS

The studies were conducted in compliance with Food and Drug Administration Good Laboratory Practice Regulations (21 CFR, Part 58). In addition, as records from the studies were submitted to the NTP Archives, these studies were audited retrospectively by an independent quality assurance contractor. Separate audits covered completeness and accuracy of the pathology data, pathology specimens, final pathology tables, and a draft of this NTP Technical Report. Audit procedures and findings are presented in the reports and are on file at NIEHS. The audit findings were reviewed and assessed by NTP staff, so all comments had been resolved or were otherwise addressed during the preparation of this Technical Report.

GENETIC TOXICOLOGY

The genetic toxicity of ethylbenzene was assessed by testing the ability of the chemical to induce mutations in various strains of *Salmonella typhimurium*, mutations in L5178Y mouse lymphoma cells, sister chromatid exchanges and chromosomal aberrations in cultured Chinese hamster ovary cells, and increases in the frequency of micronucleated erythrocytes in mouse peripheral blood. The protocols for these studies and the results are given in Appendix E.

The genetic toxicity studies of ethylbenzene are part of a larger effort by the NTP to develop a database that would permit the evaluation of carcinogenicity in experimental animals from the molecular structure and the effects of the chemical in short-term *in vitro* and *in vivo* genetic toxicity tests. These genetic toxicity tests were originally developed to study mechanisms

of chemical-induced DNA damage and to predict carcinogenicity in animals, based on the electrophilicity theory of chemical mutagenesis and the somatic mutation theory of cancer (Miller and Miller, 1977; Straus, 1981; Crawford, 1985).

There is a strong correlation between a chemical's potential electrophilicity (structural alert to DNA reactivity), mutagenicity in *Salmonella*, and carcinogenicity in rodents. The combination of electrophilicity and *Salmonella* mutagenicity is highly correlated with the induction of carcinogenicity in rats and mice and/or at multiple tissue sites (Ashby and Tennant, 1991). Other *in vitro* genetic toxicity tests correlate less well with rodent carcinogenicity (Tennant *et al.*, 1987; Zeiger *et al.*, 1990), although these other tests can provide information on the types of DNA and chromosome effects that can be induced by the chemical being investigated. Data from NTP studies show that a positive response in *Salmonella* is

the most predictive *in vitro* test for rodent carcinogenicity (89% of the *Salmonella* mutagens are rodent carcinogens) and that there is no complementarity among the *in vitro* genetic toxicity tests. That is, no battery of tests that included the *Salmonella* test improved the predictivity of the *Salmonella* test alone.

The predictivity for carcinogenicity of a positive response in bone marrow chromosome aberration or micronucleus tests appears to be less than the *Salmonella* test (Shelby *et al.*, 1993; Shelby and Witt, 1995). Positive responses in long-term peripheral blood micronucleus tests have not been formally evaluated for their predictivity for rodent carcinogenicity. However, because of the theoretical and observed associations between induced genetic damage and adverse effects in somatic and germ cells, the determination of *in vivo* genetic effects is important to the overall understanding of the risks associated with exposure to a particular chemical.

TABLE 1

Experimental Design and Materials and Methods in the 2-Year Inhalation Studies of Ethylbenzene

Study Laboratory

IIT Research Institute (Chicago, IL)

Strain and Species

Rats: F344/N Mice: B6C3F₁

Animal Source

Simonsen Laboratories, Inc. (Gilroy, CA)

Time Held Before Studies

Rats: 13 days Mice: 11 days

Average Age When Studies Began

Rats: 6 weeks Mice: 6 weeks

Date of First Exposure

Rats: 7 March 1990 Mice: 5 March 1990

Duration of Exposure

Rats: 5 days per week for 104 weeks Mice: 5 days per week for 103 weeks

Date of Last Exposure

Rats: 28 February 1992 Mice: 21 February 1992

Necropsy Dates

Rats: 9-11 March 1992 Mice: 2-5 March 1992

Average Age at Necropsy

Rats: 111 weeks Mice: 110 weeks

Size of Study Groups

50 males and 50 females

Method of Distribution

Animals were distributed randomly into groups of approximately equal initial mean body weights.

Animals per Cage

1

Method of Animal Identification

Tail tattoo

Diet

NIH-07 open formula pelleted diet (Zeigler Brothers Inc., Gardners, PA), available ad libitum

Water Distribution

Untreated coarse-filtered City of Chicago drinking water provided via automatic watering system (Edstrom Industries, Waterford, WI), available ad libitum

TABLE 1

Experimental Design and Materials and Methods in the 2-Year Inhalation Studies of Ethylbenzene (continued)

Cages

Rats: Models R-16 and R-20 (males) and models R-20 and R-24 (females) stainless steel inhalation cages

(Lab Products Inc., Maywood, NJ), rotated weekly

Mice: Model M-40 stainless steel inhalation cages (Lab Products Inc., Maywood, NJ), rotated weekly

Cage Board

Techsorb® (Shepherd Specialty Papers Inc., Kalamazoo, MI)

Chamber Air Supply Filters

Coarse prefilter, activated carbon absorber, and HEPA filter (R & R Equipment Sales, Rosemont, IL)

Inhalation Chambers

Model H-2000® 2 m³ stainless steel (Lab Products Inc., Maywood, NJ)

Racks

Stainless steel (Lab Products Inc., Maywood, NJ)

Chamber Environment

Temperature: $21\,^\circ$ to $28\,^\circ$ C (rats)

 21° to 27° C (mice) Relative humidity: 37% to 76% (rats)

32% to 72% (mice)

Fluorescent light: 12 hours/day Chamber air flow: 500 ± 66 L/minute

Exposure Concentrations

0, 75, 250, or 750 ppm

Type and Frequency of Observation

Observed twice daily; clinical findings recorded approximately monthly; body weights recorded initially, weekly for the first 13 weeks, at week 16, monthly through the end of exposure, and at study termination.

Method of Sacrifice

CO₂ asphyxiation

Necropsy

Necropsy performed on all animals.

Histopathology

Complete histopathologic examinations were performed on all chamber control and exposed rats and mice surviving to the end of the study as well as on animals that died early. In addition to gross lesions and tissue masses, the tissues examined included: adrenal gland, blood vessel (aorta), bone and marrow, brain, clitoral gland, esophagus, gallbladder (mice), heart, large intestine (cecum, colon, and rectum), small intestine (duodenum, jejunum, and ileum), kidney, larynx, liver, lung, lymph nodes (bronchial, mandibular, mesenteric, and mediastinal), mammary gland, nose, ovary, pancreas, parathyroid gland, pituitary gland, preputial gland, prostate gland, salivary gland, spleen, stomach (forestomach and glandular), testis with epididymis and seminal vesicle, thymus, thyroid gland, trachea, urinary bladder, and uterus.

RESULTS

RATS

Survival

Estimates of 2-year survival probabilities for male and female rats are shown in Table 2 and in the Kaplan-Meier survival curves (Figure 2). Survival of male rats in the 750 ppm group was significantly less than that of the chamber controls. The survival of male rats followed a negative trend, decreasing with increasing dose.

Body Weights and Clinical Findings

Mean body weights of 250 and 750 ppm males were generally less than those of the chamber controls from week 20 until the end of the study (Figure 3; Tables 3 and 4). The mean body weights of exposed groups of females were generally less than those of the chamber controls during the second year of the study. No clinical findings were attributed to ethylbenzene exposure.

TABLE 2
Survival of Rats in the 2-Year Inhalation Study of Ethylbenzene

	Chamber Control	75 ppm	250 ppm	750 ppm
Male				
Animals initially in study	50	50	50	50
Moribund	28	20	26	26
Natural deaths	7	16	11	22
Animals surviving to study termination	15	14	13	2
Percent probability of survival at the end of the study	a 30	28	26	4
Mean survival (days) ^b	651	639	651	604
Survival analysis ^c	P< 0.001	P= 0.888	P = 0.953	P< 0.001
Female				
Animals initially in study	50	50	50	50
Missing ^d	0	0	0	1
Moribund	7	14	8	6
Natural deaths	12	5	8	8
Animals surviving to study termination	31 ^e	31	34	35
Percent probability of survival at the end of the study		62	68	72
Mean survival (days)	661	690	696	706
Survival analysis	P = 0.248N	P= 1.000N	P = 0.620N	P = 0.326N

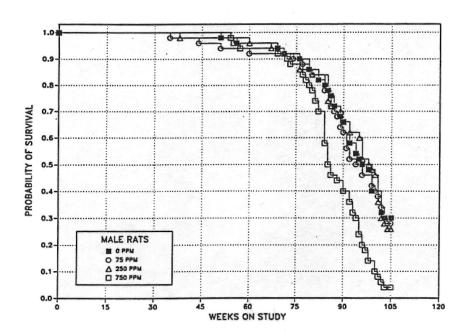
^a Kaplan-Meier determinations

b Mean of all deaths (uncensored, censored, and terminal sacrifice)

^c The result of the life table trend test (Tarone, 1975) is in the chamber control column, and the results of the life table pairwise comparisons (Cox, 1972) with the chamber controls are in the exposed group columns. A negative trend or lower mortality in an exposure group is indicated by **N**.

d Censored from survival analyses

e Includes one animal that died during the last week of the study



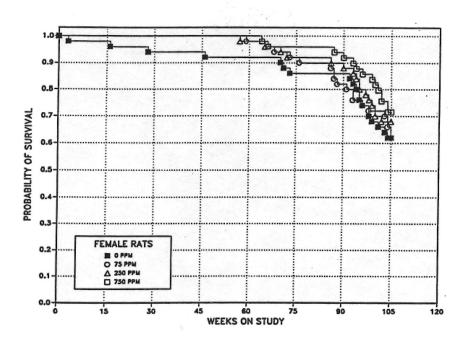
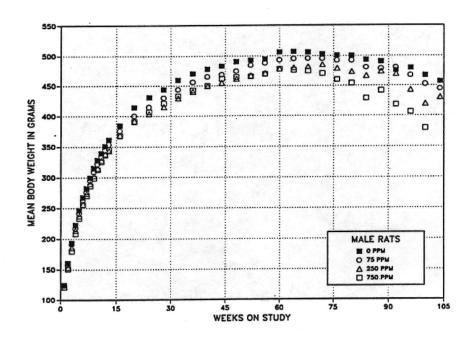


FIGURE 2 Kaplan-Meier Survival Curves for Male and Female Rats Exposed to Ethylbenzene by Inhalation for 2 Years



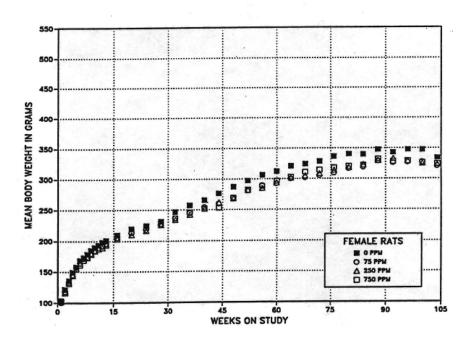


FIGURE 3
Growth Curves for Male and Female Rats Exposed to Ethylbenzene by Inhalation for 2 Years

TABLE 3
Mean Body Weights and Survival of Male Rats in the 2-Year Inhalation Study of Ethylbenzene

Weeks	Chamber Control		75 ppm			250 ppm			750 ppm		
on	Av. Wt.	No. of	Av. Wt	. Wt. (% of	No. of	Av. Wt.	Wt. (% of		Av. Wt.	Wt. (% of	
Study	(g)	Survivors	(g)		Survivors	(g)		Survivors	(g)		Survivors
1	124	50	124	100	50	122	98	50	121	98	50
2	161	50	158	99	50	154	96	50	151	94	50
3	193	50	192	99	50	186	96	50	180	93	50
4	223	50	218	98	50	214	96	50	209	94	50
5	247	50	242	98	50	238	96	50	233	95	50
6	267	50	263	98	50	257	96	50	256	96	50
7	282	50	276	98	50	274	97	50	270	96	50
8	300	50	295	99	50	290	97	50	286	96	50
9	315	50	309	98	50	303	96	50	299	95	50
10	328	50	322	98	50	315	96	50	313	95	50
11	339	50	333	98	50	328	97	50	326	96	50
12	351	50	344	98	50	338	97	50	336	96	50
13	361	50	354	98	50	344	95	50	345	96	50
16	384	50	376	98	50	369	96	50	368	96	50
20	414	50	400	97	50	392	95	50	391	94	50
24	431	50	415	96	50	404	94	50	407	95	50
28	444	50	430	97	50	415	94	50	421	95	50
32	460	50	444	97	50	430	94	50	433	94	50
36	470	50	456	97	49	441	94	50	443	94	50
40	478	50	465	97	49	450	94	49	450	94	50
44	483	50	469	97	49	455	94	49	462	96	50
48	490	50	474	97	48	461	94	49	464	95	50
52	493	49	484	98	47	467	95	49	465	94	50
56	495	49	488	99	47	471	95	49	469	95	48
60	506	48	493	98	47	479	95	49	478	94	47
64	507	48	495	98	46	480	95	48	476	94	47
68	506	48	496	98	46	482	95	47	475	94	47
72	503	46	496	99	46	485	97	46	471	94	46
76	501	46	493	98	45	479	96	44	460	92	44
80	500	43	492	99	43	473	95	43	454	91	39
84	493	41	480	97	41	466	95	42	429	87	35
88	490	36	478	98	35	474	97	36	442	90	23
92	475	33	480	101	27	469	99	33	418	88	20
96	480	26	467	98	25	443	92	30	406	85	12
100	467	20	452	97	20	420	90	24	380	81	7
104	457	15	445	97	14	431	94	14	417	91	2
Mean for	weeks										
1-13	269		264	98		259	96		256	95	
14-52	455		441	97		428	94		430	95	
53-104	491		481	98		466	95		444	90	

TABLE 4
Mean Body Weights and Survival of Female Rats in the 2-Year Inhalation Study of Ethylbenzene

Weeks	Chamber Control		75 ppm			250 ppm			750 ppm		
on	Av. Wt.	No. of	Av. Wt	. Wt. (% of	No. of	Av. Wt.	Wt. (% of		Av. Wt.	Wt. (% of	
Study	(g)	Survivors	(g)		Survivors	(g)		Survivors	(g)		Survivors
1	104	50	100	97	50	104	100	50	102	98	50
2	122	50	119	98	50	119	98	50	116	96	50
3	135	50	134	99	50	133	99	50	131	97	50
4	149	49	146	98	50	145	97	50	144	96	50
5	158	49	157	99	50	156	99	50	154	98	50
6	168	49	166	98	50	164	97	50	161	96	50
7	173	49	172	99	50	169	98	50	170	98	50
8	179	49	176	98	50	173	97	50	174	98	50
9	185	49	180	98	50	179	97	50	180	98	50
10	190	49	185	98	50	184	97	50	185	98	50
11	194	49	190	98	50	188	97	50	188	97	50
12	198	49	193	98	50	189	96	50	192	97	50
13	201	49	196	98	50	194	97	50	196	97	50
16	210	49	206	98	50	204	97	50	206	98	50
20	220	48	213	97	50	210	96	50	215	98	50
24	225	48	219	97	50	217	97	50	222	99	50
28	231	48	227	98	50	226	98	50	229	99	50
32	247	47	237	96	50	234	95	50	237	96	50
36	257	47	245	95	50	243	94	50	246	96	50
40	266	47	255	96	50	252	95	50	252	95	50
44	276	47	260	94	50	262	95	50	253	92	50
48	287	46	270	94	50	269	94	50	269	94	50
52	297	46	281	95	50	282	95	50	283	95	50
56	306	46	290	95	50	285	93	50	288	94	50
60	312	46	293	94	49	294	94	49	296	95	50
64	321	46	300	94	49	301	94	49	303	95	50
68	324	46	302	93	48	305	94	48	311	96	48
72	328	44	305	93	47	308	94	47	314	96	48
76	336	43	309	92	46	314	94	46	317	94	47
80	340	43	317	93	45	318	94	46	320	94	47
84	340	43	319	94	45	323	95	46	322	95	47
88	347	43	329	95	41	330	95	45	332	96	46
92	343	42	326	95	40	333	97	44	328	96	45
96	347	37	327	94	37	329	95	40	329	95	42
100	347	34	327	94	36	325	94	36	326	94	41
104	334	32	320	96	34	324	97	34	325	97	35
Mean for	weeks										
1-13	166		162	98		161	97		161	97	
14-52	252		241	96		240	95		241	96	
53-104	333		313	94		315	95		316	95	

Pathology and Statistical Analyses

This section describes the statistically significant or biologically noteworthy changes in the incidences of mononuclear cell leukemia as well as neoplasms and/or nonneoplastic lesions of the kidney, testis, and other organs. Summaries of the incidences of neoplasms and nonneoplastic lesions, individual animal tumor diagnoses, statistical analysis of primary neoplasms that occurred with an incidence of at least 5% in at least one animal group, and historical incidences for the neoplasms mentioned in this section are presented in Appendix A for male rats and Appendix B for female rats.

Kidney: In male rats exposed to 750 ppm, the incidences of renal tubule proliferative lesions were significantly increased relative to those in the chamber control group (Tables 5 and A3). The incidences of renal tubule adenoma and adenoma or carcinoma (combined) in this group were significantly greater than the chamber control group incidences. Renal tubule carcinomas were found in four exposed male rats, one in the 250 ppm group and three in the 750 ppm group. The incidences of renal tubule adenoma in 75 and 750 ppm males, renal tubule carcinoma in 250 and 750 ppm males, and renal tubule adenoma or carcinoma (combined) in all exposed groups of males exceeded the historical control ranges (Tables 5 and A4a). In addition, the incidence of renal tubule hyperplasia in 750 ppm males was significantly greater than that in the chamber control group (Tables 5 and A5).

Renal tubule hyperplasia, adenoma, and carcinoma constitute a morphologic and biologic continuum. Hyperplasia was a focal lesion consisting of tubules which were enlarged up to two to three times the diameter of a normal tubule and which were lined by increased numbers of epithelial cells that partially or totally filled the tubule lumen (Plate 1). Hyperplasia was considered a preneoplastic lesion and was distinguished from regenerative epithelial changes commonly seen as a component of chronic nephropathy. Renal tubule adenomas were discrete proliferative lesions, which were larger than focal hyperplasia and which tended to form more complex, usually multilobulated structures (Plate 2). Most adenomas ranged in size from 0.4 to 1 mm in size. Carcinomas were macroscopic tumors, 0.5 to 1.5 cm in size, which projected beyond the capsular surface (Plate 3). Microscopically, carcinomas were characterized by more pleomorphic cells, more prominent vascular supply, and large central areas of necrosis (Plate 4).

Initially, a single section of each kidney was examined microscopically. Because of the increased incidences of proliferative lesions in exposed males and a suggestion of a similar effect in females, additional step sections of kidney were prepared from remaining formalin-fixed tissues. Four additional sections per kidney from each male and female rat were prepared and examined. Numerous additional incidences of focal hyperplasia and adenoma were identified in the kidneys of both males and females. The incidences of these proliferative lesions observed in the extended evaluation and the combined incidences of standard and step sections are presented in Table 5. In males, there were significant increases in the incidences of renal tubule adenoma and hyperplasia in the step sections of the 750 ppm group compared to those of the chamber controls. Incidences of multiple adenomas were found in both 250 and 750 ppm males. No additional renal tubule carcinomas were identified. In the extended evaluation of females, additional incidences of renal tubule adenoma were found only in the 250 and 750 ppm groups, and the adenoma incidence in the 750 ppm group was significantly increased over chamber controls in which no adenomas were identified in either the standard or step sections. The incidence of renal tubule hyperplasia in the extended evaluation was also significantly increased in 750 ppm females.

The severities of nephropathy in 750 ppm male and all exposed female rats were significantly increased relative to chamber controls (Table 5). Nephropathy was characterized by a spectrum of changes, including dilation of renal tubules with hyaline or cellular casts. interstitial fibrosis and mononuclear inflammatory cell infiltration, foci of tubular regeneration, and transitional epithelial hyperplasia of the renal papilla. The enhanced nephropathy was more severe in males than in females, generally moderate to marked in severity, and involved most of the renal parenchyma. Several nonrenal changes which were considered secondary to the exacerbated nephropathy in 750 ppm males were significantly increased in severity relative to controls, including parathyroid gland hyperplasia, mineralization of blood vessel walls and the stomach, and fibrous osteodystrophy of bone.

TABLE 5 Incidences of Neoplasms and Nonneoplastic Lesions of the Kidney in Rats in the 2-Year Inhalation Study of Ethylbenzene

	Chambe	er Control	75 լ	ppm	250) ppm	750	ppm
Male								
Number Examined Microscopically	50		50		50		50	
Single Sections (Standard Evaluation) Nephropathy ^a Renal Tubule Hyperplasia	47 2	(2.3) ^b (3.0)	43	(2.4) (2.0)	47 4	(2.3) (1.3)	48 12**	(3.5)** (1.8)
Renal Tubule Adenoma ^c Renal Tubule Carcinoma ^d Renal Tubule Adenoma or Carcinoma ^c	0 0 0		3 0 3		2 1 3		4* 3 7**	
Step Sections (Extended Evaluation) Renal Tubule Hyperplasia	10		7		9		17*	
Renal Tubule Adenoma, Multiple Renal Tubule Adenoma (includes multiple) Renal Tubule Carcinoma Renal Tubule Adenoma or Carcinoma	0 3 0 3		0 2 0 2		2 7 1 8		4 17** 3 18**	
Single Sections and Step Sections (Combined) Renal Tubule Hyperplasia Renal Tubule Hyperplasia, Oncocytic	11 2	(2.0) (3.0)	9	(2.3) (2.3)	11 0	(2.1)	23** 1	(2.5) (2.0)
Renal Tubule Adenoma, Multiple Renal Tubule Adenoma (includes multiple) Renal Tubule Carcinoma Renal Tubule Adenoma or Carcinoma Oncocytoma	0 3 0 3 0		0 5 0 5 1		2 7 1 8 1		4 20** 3 21** 2	
Female								
Number Examined Microscopically	50		50		50		49	
Single Sections (Standard Evaluation) Nephropathy Renal Tubule Hyperplasia	38 0	(1.3)	42 1	(1.6)* (1.0)	43	(1.7)** (2.3)	46 3	(2.3)** (1.3)
Renal Tubule Adenoma	0		0		0		1	
Step Sections (Extended Evaluation) Renal Tubule Hyperplasia	1		1		1		8*	
Renal Tubule Adenoma	0		0		1		7*	
Single Sections and Step Sections (Combined) Renal Tubule Hyperplasia	1	(1.0)	2	(1.0)	4	(2.2)	10**	(1.8)
Renal Tubule Adenoma	0		0		1		8**	

Significantly different $(P \le 0.05)$ from the chamber control group by the logistic regression test (incidence) or by the Mann-Whitney U test (severity)
** P≤0.01

Number of animals with lesion

Average severity grade of lesions in affected animals: 1= minimal; 2= mild; 3= moderate; 4= marked

Historical incidence for 2-year inhalation studies with chamber control groups (mean \pm standard deviation): 6/652 (0.9% \pm 1.3%); range, 0%-4%

Historical incidence: 0/652

Testis: The incidence of interstitial cell adenoma in 750 ppm males was significantly greater than that in the chamber control group and slightly exceeded the historical control range; the incidence of bilateral testicular adenoma was also significantly increased in 750 ppm males (Tables 6, A3, and A4b). This common neoplasm in male F344/N rats is composed of nodular aggregates of large polyhedral cells with

foamy or eosinophilic cytoplasm that extend between and cause compression of the surrounding seminiferous tubules. This neoplasm will develop in nearly all male rats if they are allowed to complete their natural life span; ethylbenzene appeared to enhance its development. The incidence of interstitial cell hyperplasia in 750 ppm males was significantly decreased.

TABLE 6
Incidences of Neoplasms and Nonneoplastic Lesions of the Testis in Male Rats in the 2-Year Inhalation Study of Ethylbenzene

	Chamber Control	75 ppm	250 ppm	750 ppm
Number Examined Microscopically Interstitial Cell Hyperplasia ^a	50 14 (1.5) ^b	50 19 (1.2)	50 12 (1.3)	50 8* (1.1)
Bilateral Adenoma				
Overall rate ^c	27/50 (54%)	23/50 (46%)	32/50 (64%)	40/50 (80%)
Adjusted rate ^d	96.0%	91.0%	96.5%	100.0%
Terminal rate e	14/15 (93%)	12/14 (86%)	12/13 (92%)	2/2 (100%)
First incidence (days)	608	538	590	500
Logistic regression test ^f	P< 0.001	P = 0.313N	P = 0.177	P< 0.001
Adenomag				
Overall rate	36/50 (72%)	33/50 (66%)	40/50 (80%)	44/50 (88%)
Adjusted rate	100.0%	100.0%	100.0%	100.0%
Terminal rate	15/15 (100%)	14/14 (100%)	13/13 (100%)	2/2 (100%)
First incidence (days)	497	538	420	483
Logistic regression test	P< 0.001	P = 0.404N	P = 0.194	P = 0.001

^{*} Significantly different ($P \le 0.05$) from the chamber control group by the logistic regression test

a Number of animals with lesion

b Average severity grade of lesions in affected animals: 1= minimal; 2= mild; 3= moderate; 4= marked

^c Number of animals with neoplasm per number of animals with testis examined microscopically

d Kaplan-Meier estimated neoplasm incidence at the end of the study after adjustment for intercurrent mortality

e Observed incidence in animals surviving until the end of the study

In the chamber control column are the P values associated with the trend test. In the exposed group columns are the P values corresponding to the pairwise comparisons between the chamber controls and that exposed group. The logistic regression test regards neoplasms in animals dying prior to terminal kill as nonfatal. A lower incidence in an exposure group is indicated by N.

Historical incidence for 2-year inhalation studies with chamber control groups (mean ± standard deviation): 450/655 (68.7% ± 8.7%); range, 54%-83%

Other organs: The incidences of several nonneoplastic lesions were significantly greater in the 750 ppm males than in chamber controls (Table A5). Incidences of edema (chamber control, 1/50; 75 ppm, 0/50; 250 ppm, 0/50; 750 ppm, 6/50), congestion (1/50, 2/50, 0/50, 6/50), and hemorrhage (0/50, 2/50, 1/50, 8/50) in the lungs as well as hemorrhage in mesenteric (3/49, 5/50, 4/50, 8/50) and renal (0/9, 0/8, 1/9, 8/14) lymph nodes were slightly increased. These circulatory lesions were considered to be agonal changes in moribund animals and not directly related to chemical toxicity. The incidence of cystic degeneration of the liver was also increased in 750 ppm males (15/50, 12/50, 19/50, 30/49); the biologic significance of this increase in the absence of other hepatotoxic changes is unclear.

Compared to the chamber control group, the incidences of prostate gland inflammation in all exposed groups of males were significantly increased (11/50, 29/50, 22/50, 25/50; Table A5). This inflammatory change consisted of infiltration by predominantly mononuclear inflammatory cells into glandular acini

and interstitium, increased interstitial fibrosis, and loss of secretory material in affected areas. Relative to chamber controls, males exposed to 75 or 750 ppm exhibited increased incidences of hyperplasia of the bone marrow characterized by hypercellularity due to increased numbers of erythroid and myeloid precursor cells (7/49, 16/49, 9/50, 19/50). The relationship of these changes to ethylbenzene exposure is uncertain due to the lack of clear concentration-dependent responses.

Mononuclear cell leukemia: The incidence of mononuclear cell leukemia was decreased in 750 ppm males (27/50, 26/50, 32/50, 9/50; Table A3). While this decrease was statistically significant by logistic regression, it was not significant by life table analysis, the more appropriate test for this generally fatal neoplasm. This decrease was due in large part to the reduced survival in the 750 ppm group as a result of nephropathy and, therefore, was not considered to be related to ethylbenzene exposure.

MICE

Survival

Estimates of 2-year survival probabilities for male and female mice are shown in Table 7 and in the Kaplan-Meier survival curves (Figure 4). Survival of exposed groups of male and female mice was similar to that of the chamber controls.

Body Weights and Clinical Findings

Mean body weights of female mice exposed to 75 ppm were greater than those of the chamber controls from week 72 until the end of the study; mean body weights of 750 ppm females were generally less than those of the chamber controls from week 24 through week 68 but were similar to those of the chamber controls from week 72 until the end of the study (Tables 8 and 9; Figure 5). No clinical findings were attributed to ethylbenzene exposure.

TABLE 7
Survival of Mice in the 2-Year Inhalation Study of Ethylbenzene

	Chamber Control	75 ppm	250 ppm	750 ppm
Лаle				
nimals initially in study	50	50	50	50
accidental deaths ^a	1	0	0	1
Moribund	6	2	5	6
Jatural deaths	15	12	13,	13
nimals surviving to study termination	28	36	32^{d}	30
ercent probability of survival at the end of the study b	57	72	64	61
Mean survival (days) ^C	636	684	692	665
urvival analysis ^e	P = 0.975	P= 0.177N	P = 0.459N	P = 0.673N
'emale				
nimals initially in study	50	50	50	50
accidental deaths ^a	1	0	1	0
Toribund	5	6	1	4
Jatural deaths	9,	6	8	9
'erminal sacrifice	35^{d}	38	40	37
ercent probability of survival at the end of the study	71	76	82	74
Mean survival (days)	689	700	701	692
urvival analysis	P = 0.995N	P= 0.762N	P = 0.304N	P = 0.886N

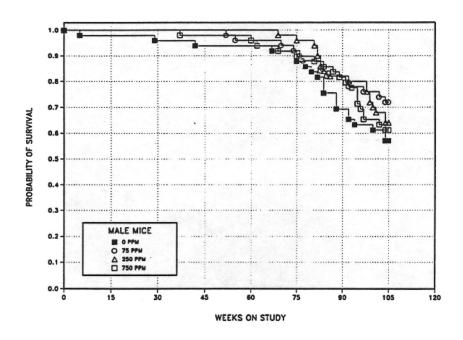
a Censored from survival analyses

Kaplan-Meier determinations

Mean of all deaths (uncensored, censored, and terminal sacrifice)

Includes one animal that died during the last week of the study

^e The result of the life table trend test (Tarone, 1975) is in the chamber control column, and the results of the life table pairwise comparisons (Cox, 1972) with the chamber controls are in the exposed group columns. A negative trend or lower mortality in an exposure group is indicated by **N**.



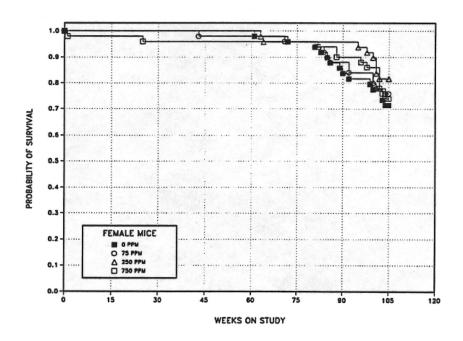


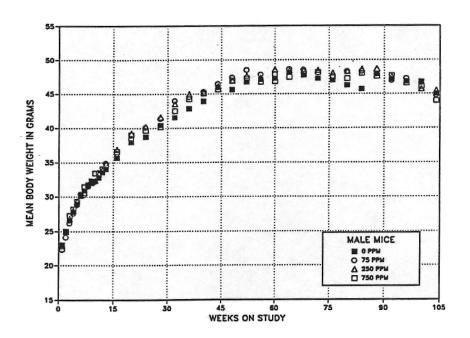
FIGURE 4
Kaplan-Meier survival Curves for Male and Female Mice Exposed to Ethylbenzene by Inhalation for 2 Years

TABLE 8
Mean Body Weights and Survival of Male Mice in the 2-Year Inhalation Study of Ethylbenzene

on Av. Wt. No. of Av. Wt. Wt. (% of No. of No. of Av. Wt. Wt. (% of No. of N		750 ppm			250 ppm			75 ppm		r Control	Chambe	Weeks
1			Av Wt	No of		Av Wt	No of		Δv Wt			
1 23.0 50 22.3 97 50 22.9 100 50 22.4 97 2 24.9 50 24.2 97 50 25.0 100 50 25.0 100 3 26.6 50 26.2 99 50 28.1 101 50 28.2 101 5 29.0 48 28.8 99 50 29.4 101 50 29.3 101 6 30.2 48 30.4 101 50 29.3 101 7 30.9 48 30.3 98 50 30.7 99 50 31.5 102 8 31.6 48 31.5 100 50 31.8 101 50 31.7 100 9 32.0 48 32.2 101 50 32.4 100 50 33.7 101 50 33.5 104 11 32.3 101	s) Survivors											
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	31	98	44.0	33	101	45.4	37	100	44.8	29	45.0	104
Mean for weeks											wooks	Mean for
1-13 29.9 29.8 100 30.1 101 30.3 101		101	30.3		101	30.1		100	29 R			
14-52 41.9 43.2 103 43.2 103 42.7 102												
53-104 46.9 47.6 101 47.7 102 47.0 100												

TABLE 9
Mean Body Weights and Survival of Female Mice in the 2-Year Inhalation Study of Ethylbenzene

Weeks	Chambe	er Control		75 ppm			250 ppm			750 ppm	
on	Av. Wt.	No. of	Av Wt	Wt. (% of	No. of	Av Wt	Wt. (% of		Av Wt	Wt. (% of	No. of
Study	(g)	Survivors	(g)		Survivors	(g)		Survivors	(g)		Survivors
1	18.5	50	18.6	101	50	18.6	101	50	18.0	97	50
2	20.1	50	19.9	99	50	20.0	100	50	19.5	97	49
3	21.4	50	20.7	97	50	21.1	99	50	21.4	100	49
4	22.4	50	21.6	96	50	22.3	100	50	22.5	100	49
5	23.3	50	22.8	98	50	23.2	100	50	23.5	101	49
6	24.3	50	23.4	96	50	24.2	100	50	23.8	98	49
7	24.4	50	24.0	98	50	24.4	100	50	24.5	100	49
8	25.1	50	24.6	98	50	24.9	99	50	25.1	100	49
9	26.0	50	25.7	99	50	25.6	99	50	26.1	100	49
10	26.0	50	25.4	98	50	25.9	100	50	26.1	100	49
11	26.7	50	26.2	98	50	26.0	97	50	27.0	101	49
12	26.6	50	26.5	100	50	26.7	100	50	26.6	100	49
13	27.3	50	27.2	100	50	26.7	98	49	27.0	99	49
16	28.4	50	28.0	99	50	27.3	96	49	28.8	101	49
20	31.0	50	29.9	97	50	28.8	93	49	30.5	98	49
24	32.0	50	31.2	98	50	29.9	93	49	29.8	93	49
28	33.1	50	32.9	99	50	30.9	93	49	30.5	92	48
32	34.1	49	33.8	99	50	32.5	95	49	31.7	93	48
36	35.7	49	35.5	99	50	35.3	99	49	33.1	93	48
40	36.3	49	36.7	101	50	35.8	99	49	33.4	92	48
44	39.1	49	38.4	98	49	36.8	94	49	35.4	90	48
48	39.8	49	40.0	101	49	40.0	101	49	37.2	94	48
52	40.9	49	41.8	101	49	41.2	101	49	39.4	96	48
56	40.5	49	43.6	102	49	43.0	101	49	39.4	93	48
60	42.0	49	44.4	102	49	43.7	101	49	40.7	94	48
64	43.1	49	44.4	103	49	43.7	101	49	40.7	91	48
68		48	47.3	101			100				48
	46.3				49	46.5		47	43.2	93	
72	45.6	48	48.6	107	48	47.3	104	47	44.3	97	48
76	46.0	47	48.1	105	48	47.8	104	47	44.4	97	48
80	46.0	47	49.0	107	48	49.5	108	47	44.9	98	48
84	45.8	45	49.8	109	46	49.5	108	47	45.0	98	47
88	45.8	43	50.8	111	44	49.0	107	47	45.2	99	46
92	47.0	40	51.8	110	43	49.9	106	47	45.6	97	45
96	47.0	40	50.3	107	42	49.0	104	46	45.7	97	44
100	47.0	38	49.9	106	42	47.6	101	44	46.4	99	43
104	45.9	36	49.9	109	38	45.2	99	40	45.5	99	37
Mean for	weeks										
1-13	24.0		23.6	98		23.8	99		23.9	100	
14-52	35.0		34.8	99		33.9	97		33.0	94	
53-104	45.6		48.4	106		47.1	103		43.9	96	



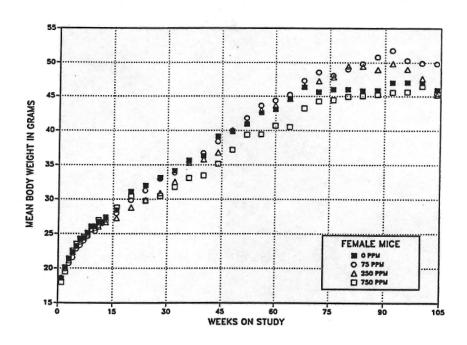


FIGURE 5 Growth Curves for Male and Female Mice Exposed to Ethylbenzene by Inhalation for 2 Years

Pathology and Statistical Analysis

This section describes the statistically significant or biologically noteworthy changes in the incidences of neoplasms and/or nonneoplastic lesions of the lung, liver, and other organs. Summaries of the incidences of neoplasms and nonneoplastic lesions, individual animal tumor diagnoses, statistical analysis of primary neoplasms that occurred with an incidence of at least 5% in at least one animal group, and historical incidences for the neoplasms mentioned in this section are presented in Appendix C for male mice and Appendix D for female mice.

Lung: Incidences of alveolar/bronchiolar adenoma and alveolar/ bronchiolar adenoma or carcinoma (combined) in males increased with a positive trend (Tables 10 and C3). In 750 ppm males, the incidences of alveolar/bronchiolar adenoma and

alveolar/bronchiolar adenoma or carcinoma (combined) were significantly greater than those in the chamber control group but were within the historical control ranges (Tables 10, C3, and C4). In 750 ppm females, the incidence of alveolar/ bronchiolar adenoma was greater than that in the chamber control group. This difference was not significant, but the incidence exceeded the historical control range (Tables 10, D3, and D4a). Alveolar/bronchiolar neoplasms were nodular proliferations within the lung parenchyma which caused variable compression depending on size (Plate 5). Adenomas were typically well circumscribed nodules composed of monomorphic cuboidal cells arranged in solid or papillary patterns. In carcinomas, the borders were less distinct and the neoplastic cells were cuboidal to columnar in shape and exhibited greater cytologic atypia.

TABLE 10 Incidences of Neoplasms and Nonneoplastic Lesions of the Lung in Mice in the 2-Year Inhalation Study of Ethylbenzene

	Chambe	er Control	75 լ	ppm	250) ppm	750	ppm
Male								
Number Examined Microscopically	50		50		50		50	
Alveolar Epithelium, Hyperplasia ^a	1	$(1.0)^{b}$	5	(2.6)	2	(1.5)	4	(2.0)
Alveolar Epithelium, Metaplasia	0		1	(1.0)	2	(1.0)	6*	(1.2)
Alveolar/bronchiolar Adenoma ^c	5		9		10		16**	k
Alveolar/bronchiolar Carcinoma	2		1		5		3	
Alveolar/bronchiolar Adenoma or Carcinoma ^d	7		10		15		19**	k
Female								
Number Examined Microscopically	50		50		49		50	
Alveolar Epithelium, Hyperplasia	0		1	(2.0)	3	(2.0)	1	(3.0)
Alveolar Epithelium, Metaplasia	0		0		0		1	(2.0)
Alveolar/bronchiolar Adenoma ^e	4		4		5		8	
Alveolar/bronchiolar Adenoma or Carcinoma ^t	4		6		5		8	

^{*} Significantly different ($P \le 0.05$) from the chamber control group by the logistic regression test

^{**} $(P \le 0.01)$

a Number of animals with lesion

b Average severity grade of lesions in affected animals: 1= minimal; 2= mild; 3= moderate; 4= marked

^c Historical incidence for 2-year inhalation studies with chamber control groups (mean ± standard deviation): 141/947 (14.9% ± 7.0%); range, 6%-36%

d Historical incidence: 205/947 (21.7% ± 8.0%); range, 10%-42%

Historical incidence: $61/939 (6.5\% \pm 3.2\%)$; range, 0%-14%

f Historical incidence: $97/939 (10.3\% \pm 3.7\%)$; range, 0%-16%

Another proliferative change in the lung was observed only in exposed mice and was diagnosed as alveolar epithelial metaplasia. In males, the incidence of this lesion increased with increasing exposure concentration and was significantly increased in the 750 ppm group (Tables 10 and C5). Alveolar epithelial metaplasia was also observed in one 750 ppm female. Metaplasia was characterized by the presence of cells morphologically similar to bronchiolar epithelial cells lining the alveolar spaces adjacent to terminal bronchioles (Plate 6).

The incidences of hepatocellular adenoma Liver: and adenoma or carcinoma (combined) in females occurred with a positive trend (Table D3). These incidences in 750 ppm females were significantly greater than those in the chamber controls but did not exceed the historical control ranges (Tables 11, D3, and D4b). Although hepatocellular carcinomas also occurred with a positive trend, incidences in exposed groups were not significantly greater than in chamber controls and did not exceed historical control ranges. Multiple adenomas were found in all exposed groups of females, and multiple carcinomas were found in two 750 ppm females, but multiple liver neoplasms were not found in chamber control females. Hepatocellular adenomas consisted of nodules of hepatocytes which compressed adjacent liver parenchyma and lacked the normal lobular and sinusoidal pattern. Hepatocellular carcinomas were large masses composed of anaplastic hepatocytes forming solid sheets or trabecular patterns.

In addition to liver neoplasms, the incidence of eosinophilic foci in the liver was significantly greater in 750 ppm females than in chamber controls (Tables 11 and D5). This lesion, composed of focal collections of cells, which have altered staining characteristics and which blend into surrounding hepatic cords with little or no compression, is considered to be a precursor to hepatocellular neoplasia.

A spectrum of nonneoplastic liver changes related to ethylbenzene exposure in male mice included syncytial alteration of hepatocytes, hepatocellular hypertrophy, and hepatocyte necrosis (Tables 11 and C5). These changes were minimal to mild in severity. Syncytial alteration was seen in all groups of exposed males, with concentration-dependent increases in incidence. This change consisted of the presence of greatly enlarged hepatocytes containing multiple nuclei, generally five or more, either randomly scattered throughout the liver lobule or with a tendency to cluster in centrilobular areas (Plate 7). Hypertrophy of hepatocytes occurred in the centrilobular zones of 750 ppm males and was characterized by cells with increased amounts of cytoplasm and enlarged nuclei. Syncytial alteration and hypertrophy frequently occurred in the same animal. Hepatocellular necrosis was evident as random single cell necrosis, generally of hypertrophied cells.

Other organs: Significantly increased incidences of hyperplasia of the pituitary gland pars distalis were limited to 250 and 750 ppm females (chamber control, 10/48; 75 ppm, 12/49; 250 ppm, 23/47; 750 ppm, 22/49; Table D5). This hyperplasia was seen as focal, poorly delineated, monomorphic increases of cells which had no compressive features or altered arrangement. Positive trends in the incidences of thyroid follicular cell hyperplasia occurred in both males (21/50, 21/50, 29/50, 32/50; Table C5) and females (18/50, 23/50, 25/50, 35/50; Table D5), with significant increases in incidences relative to chamber controls in 750 ppm males and females. Thyroid hyperplasia was typically a focal noncompressive proliferation with simple papillary infoldings of follicular epithelial cells. There were no corresponding increases in the incidences of adenomas of either the pituitary gland or thyroid gland (Tables C1 and D1).

TABLE 11
Incidences of Neoplasms and Nonneoplastic Lesions of the Liver in Mice in the 2-Year Inhalation Study of Ethylbenzene

	Chambe	er Control	75 <u>j</u>	opm	25() ppm	750 ppm
Male							
Number Examined Microscopically Hepatocyte, Hypertrophy ^a Hepatocyte, Necrosis Hepatocyte, Syncytial Alteration	50 1 1 0	(1.0) ^b (1.0)	50 0 1 5	(2.0) (1.0)	50 0 3 8*	(1.3) * (1.4)	50 17** (1.1) 10** (1.8) 23** (1.1)
Female							
Number Examined Microscopically Eosinophilic Focus	50 5	(1.8)	50 7	(1.4)	50 6	(1.5)	50 22** (2.0)
Hepatocellular Adenoma, Multiple Hepatocellular Adenoma (includes multiple) ^C Hepatocellular Carcinoma, Multiple Hepatocellular Carcinoma (includes multiple) Hepatocellular Adenoma or Carcinoma ^d	0 6 0 7 13		1 9 0 4 12		3 12 0 3 15		4 16* 2 12 25*

^{*} Significantly different (P≤0.05) from the chamber control group by the logistic regression test

GENETIC TOXICOLOGY

Ethylbenzene was not mutagenic in *Salmonella typhimurium* strain TA97, TA98, TA100, or TA1535 with or without Aroclor-induced rat or hamster liver S9 (Table E1; Zeiger *et al.*, 1988). A positive response was observed with ethylbenzene in the L5178Y mouse lymphoma cell assay in the absence of S9 at the highest nonlethal dose tested (80 μ g/mL); the assay was not performed with S9 (Table E2; McGregor *et al.*, 1988). A significant amount of

cytotoxicity was noted at this dose level (relative total growth was reduced to 34% and 13% of the control level in each of two trials). No increases in sister chromatid exchanges (Table E3) or chromosomal aberrations (Table E4) were induced by ethylbenzene in cultured Chinese hamster ovary cells, with or without S9. *In vivo*, no increases in micronucleated erythrocytes were observed in peripheral blood samples from male and female mice exposed to ethylbenzene for 13 weeks by inhalation (Table E5).

^{**} $(P \le 0.01)$

^a Number of animals with lesion

Average severity grade of lesions in affected animals: 1= minimal; 2= mild; 3= moderate; 4= marked

Historical incidence for 2-year inhalation studies with chamber control groups (mean ± standard deviation): 114/937 (12.2% ± 9.7%); range. 0%-40%

d Historical incidence: 200/937 (21.3% \pm 11.9%); range, 3%-54%

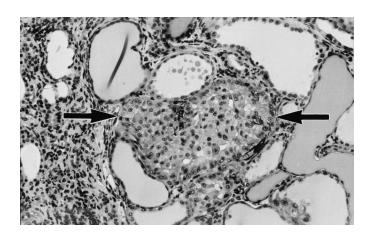


Plate 1

Renal tubule hyperplasia in the kidney of a male F344/N rat exposed to 750 ppm ethylbenzene by inhalation for 2 years. The hyperplastic tubule (between arrows) consists of epithelial cells which fill the lumen. Note the changes of chronic nephropathy in the surrounding parenchyma including dilated tubules and thickened interstitium. H&E; $100\times$

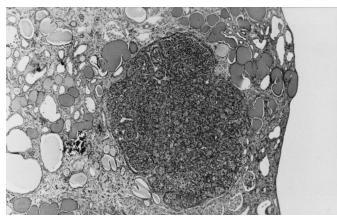


Plate 2

Renal tubule adenoma in the kidney of a male F344/N rat exposed to 750 ppm ethylbenzene by inhalation for 2 years. The adenoma is just under the capsular surface and is well circumscribed and multilobulated. Note the changes of chronic nephropathy in the surrounding parenchyma including dilated tubules filled with protein casts, thickened interstitium, and focal mineralization. H&E; $35\times$

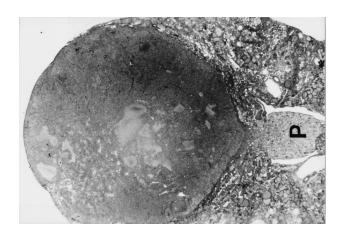


Plate 3

Renal tubule carcinoma in the kidney of a male F344/N rat exposed to 750 ppm ethylbenzene by inhalation for 2 years. The 1 cm diameter mass protrudes coutside the capsular surface and extends deep into the parenchyma near the papilla (P). Note the cystic necrosis of the center. H&E; $6\times$

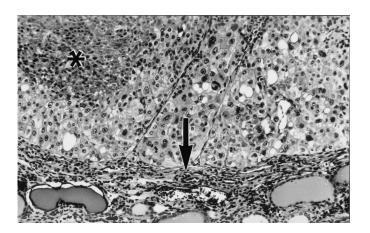


Plate 4

Higher magnification of Plate 3. Irregular lobules of pleomorphic tumor cells are separated by fine septae and compress the surrounding parenchyma (arrow). Note the central necrosis of one lobule (*). H&E; $100\times$

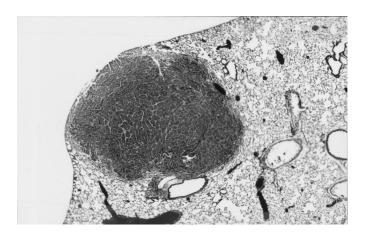


Plate 5 Alveolar/bronchiolar adenoma in the lung of a male B6C3F $_1$ mouse exposed to 750 ppm ethylbenzene by inhalation for 2 years. The adenoma is well demarcated from the adjacent compressed lung parenchyma, and there is bulging of the pleural surface. H&E; $20\times$

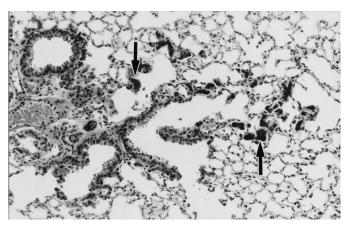


Plate 6 Alveolar epithelial metaplasia in the lung of a male $B6C3F_1$ mouse exposed to 750 ppm ethylbenzene by inhalation for 2 years. Multiple foci of dark-staining epithelial cells (arrows) are in the alveolar spaces adjacent to one branch of a terminal bronchiole bifurcation. H&E; $85\times$

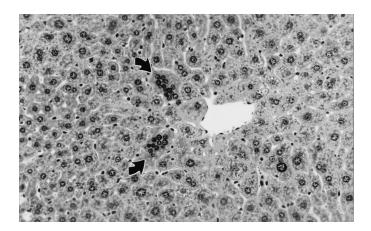


Plate 7 Syncytial alteration of hepatocytes in the liver of a male $B6C3F_1$ mouse exposed to 750 ppm ethylbenzene by inhalation for 2 years. Two large syncytial cells (arrows), each containing approximately 10 nuclei, are adjacent to a central vein. H&E; $140\times$

DISCUSSION AND CONCLUSIONS

Ethylbenzene is mainly used in the manufacture of styrene. Ethylbenzene is also a major component of mixed xylenes used as solvents in agricultural and home insecticide sprays, rubber and chemical manufacturing, and household degreasers, paints, adhesives, and rust preventives (Fishbein, 1985). Ethylbenzene has been used as an antiknock agent in aviation and motor fuels (NIOSH, 1979).

The National Institute for Occupational Safety and Health and the Occupational Safety and Health Administration nominated ethylbenzene for study because of its widespread human exposure and because of its structural similarity to benzene and toluene.

In previous studies, male and female F344/N rats and B6C3F $_1$ mice were exposed to ethylbenzene by inhalation for 13 weeks at concentrations of 0, 100, 250, 500, 750, or 1,000 ppm (NTP, 1992). In the current studies, male and female F344/N rats and B6C3F $_1$ mice were exposed to ethylbenzene by inhalation for 2 years at concentrations of 0, 75, 250, or 750 ppm.

In the 2-year study, the survival rate and the mean body weights of the 750 ppm male rats were less than those of the chamber control group after week 75 of the study. Female rats generally had higher survival rates than males, and this was probably related to the typical occurrence of nephropathy in male F344/N rats, which was enhanced by ethylbenzene exposure. The mean body weights of exposed groups of female rats were less than those of the chamber controls during the second year of the study. Survival rates of exposed male and female mice were similar to those of the respective chamber controls. Female mice exposed to 75 ppm had greater mean body weights than those of the chamber controls.

In the 13-week studies, increased absolute and relative kidney weights were observed in male rats exposed to 750 or 1,000 ppm ethylbenzene, although no accompanying histopathologic changes were seen (NTP, 1992). In the standard histopathologic evaluation of the kidney in the 2-year study, the incidence of renal tubule adenoma in the 750 ppm male rats was signifi-

cantly greater than that in the chamber control group. The incidence in 750 ppm males exceeded the NTP historical control range. An extended evaluation of the kidneys in the 2-year study identified many more adenomas. In addition, multiple renal tubule adenomas were found in the 250 and 750 ppm males. The standard evaluation and extended evaluation (combined) showed significantly increased incidences of renal tubule adenoma, renal tubule adenoma or carcinoma (combined), and renal tubule hyperplasia in 750 ppm male rats, as well as positive trends across exposure groups. No renal lesions were observed in females in the NTP 13-week study. In the standard evaluation in the 2-year study, no significant increases in incidences of renal lesions were observed in female rats. In the extended evaluation of the kidneys, the incidences of renal tubule hyperplasia and renal tubule adenoma were significantly increased in the 750 ppm female rats compared to those in the chamber controls.

Kurokawa et al. (1983) first reported that a greater incidence of rat kidney lesions was found when multiple kidney sections were examined compared with single sections. This was expected, considering the very small size of many of the tubule cell adenomas typically seen in the kidney. The NTP has compared lesions from single and multiple kidney sections and found increased incidences of renal tubule hyperplasia and renal tubule adenoma in multiple sections from male rats (Eustis et al., 1994), agreeing with the findings of Kurokawa et al. (1983). However, few additional neoplasms were identified in female rats or in male or female mice (Eustis et al., 1994). In the present studies, additional incidences of renal tubule hyperplasia and renal tubule adenoma were found in step sections from male and female rats.

Nephropathy is commonly found in aging male and, to a lesser degree, female rats; in the current study, the severities of nephropathy were increased in 750 ppm male rats and in all exposed female rat groups. In the extended evaluation of the kidneys in the 2-year study of ethylbenzene, the incidences of renal tubule hyperplasia in 750 ppm males and females were increased. Ethylbenzene may have

exacerbated the age-related nephropathy development in rats or exerted toxic injury to the renal cells and induced compensatory cell replication of the renal tubule epithelium. Whether they were a direct effect of ethylbenzene or an indirect result of ethylbenzene-induced cytotoxicity, the renal tubule lesions in male and female rats were considered exposure related. Males appeared to be more sensitive to the renal toxic effect of ethylbenzene than females, and that may account for the early deaths in 750 ppm males.

Following exposure to certain hydrocarbons, male rats develop renal tubule hyaline droplets, attributed to accumulation of $\alpha 2\mu$ -globulin in the kidney. The accumulation of $\alpha 2\mu$ -globulin is known to lead to nephropathy and renal tubule neoplasm development in male rats. This spectrum of nonneoplastic changes differs from the chronic progressive nephropathy commonly found in aging male rats (USEPA, 1991). No clear evidence of hyaline droplets was seen in the kidneys of male F344/N rats exposed to ethylbenzene for 13-weeks (NTP, 1992) or 2 years, and, thus, this proposed mechanism did not appear to contribute to the proliferative renal tubule lesions in male or female F344/N rats in the studies reported here.

After a 6-hour inhalation exposure to ethylbenzene in male Wistar rats, the major metabolites identified in the urine were 1-phenylethanol (α-methylbenzyl alcohol), mandelic acid, phenylglyoxylic acid, phenylacetic acid, and benzoic acid. Minor metabolites included omega-hydroxyacetophenone, 1-phenyl-1,2ethanediol, acetophenone, p-hydroxyacetophenone, and phenylglyoxal. Blood metabolites were difficult to identify and measure (Engström, 1984). None of the urinary metabolites, except 1-phenylethanol, are considered ultimate carcinogens likely reactive with cellular macromolecules. Ethylbenzene is neither mutagenic nor clastogenic. Both 1- and 2-phenylethanol were negative for mutagenicity and did not induce sister chromatid exchanges in cultured human lymphocytes (NTP, 1990b; Norppa and Vainio, 1983). It is possible that in the process of metabolizing ethylbenzene to 1- and 2-phenylethanol, an epoxide intermediate is formed. A gender difference in epoxide formation may account for the differential sensitivity for neoplasia between the male and female mouse lungs.

α-Methylbenzyl alcohol (1-phenylethanol), a metabolite of ethylbenzene (Engström, 1984), has been shown to enhance nephropathy and induce renal tubule adenoma or adenocarcinoma in male F344/N rats (NTP, 1990b) but had no effect on nephropathy or renal tubule lesions in female F344/N rats. Since kidney toxicity and carcinogenicity were observed in both male and female rats in the present studies, the data suggested that the renal effect of ethylbenzene is more potent than that of α -methylbenzyl alcohol. Other metabolites, such as an epoxide or diolepoxide after ring oxidation (Engström, 1984), phenylglyoxal bearing a reactive aldehyde group, or those metabolites postulated in benzene metabolism, such as hydroquinone, benzoquinone, or benzene diolepoxide (NTP, 1986; Busby et al., 1990), may contribute to the renal toxicity and carcinogenicity of ethylbenzene. However, no reactive metabolite has been identified. Further studies to identify the active species are needed. It should be noted that neither ethylbenzene nor α-methylbenzyl alcohol is mutagenic or clastogenic.

Structurally, ethylbenzene is related to benzene and toluene (Table 12). Toluene is negative for carcinogenic activity (NTP, 1990a). Benzene is a multipotential carcinogen suppressing bone marrow cellularity and inducing leukopenia and leukemia and neoplasms in the Zymbal's gland, oral cavity, and skin in rats and Zymbal's gland, lymph gland, lung, harderian gland, preputial gland, mammary gland, ovary, forestomach, and liver in mice after gavage dosing (NTP, 1986). Benzene is metabolized to benzene oxide, benzene oxepin, benzene dihydrodiol, phenol, hydroquinone, trihydroxybenzene, catechol, benzoquinone, and trans, trans-muconaldehyde (Snyder and Hedli, 1996; Sabourin et al., 1989, 1992). The metabolites proposed for the hematotoxicity in rats are hydroquinone, benzoquinone, and trans, trans-muconaldehyde (Zhu et al., 1995), and for lung tumors in mice, the metabolite is benzene diolepoxide-2 (Busby et al., 1990). Although benzene and ethylbenzene are structurally related, the metabolites, target organs, and mechanism of action of benzene appear quite different from those of ethylbenzene.

TABLE 12
Results of Carcinogenicity and Mutagenicity Tests of Benzene, Toluene, and Ethylbenzene in Male and Female F344/N Rats and Male and Female B6C3F Mice in 2-Year Studies 1 ^a

		Carcino	genicity		Salmonella
Chemical and Route	Male Rat	Female Rat	Male Mouse	Female Mouse	Test Result
Benzene (gavage) (NTP, 1986)	Zymbal's gland, oral cavity, skin	+ Zymbal's gland, oral cavity		Zymbal's gland, lymph gland, lung, harderian gland, mammary gland, ovary, forestomach, liver	_
Toluene (inhalation) (NTP, 1990a)	_	_	_	_	_
Ethylbenzene (inhalation) CH ₂ CH ₃	+ kidney, testis	+ kidney	+ lung	+ liver	_

aCarcinogenic response: + = some or clear evidence of carcinogenic activity; — = no evidence of carcinogenic activity

The increased incidence of testicular adenoma observed in male rats in the 750 ppm group was considered related to ethylbenzene exposure. This is evidenced by the finding that 92% (22/24) of the 750 ppm male rats that died between days 400 and 600 had testicular adenoma, whereas only 33% (3/9) of the chamber controls that died early had testicular adenoma. The incidence of bilateral adenoma was also increased in 750 ppm males. Testicular adenoma develops in nearly all rats in the latter part of their lives, but in inhalation studies, the incidence is low compared with those in feed and gavage studies (Haseman et al., 1997); ethylbenzene appeared to

hasten the development of testicular adenoma. How ethylbenzene accomplishes this effect is not clear. There were no testicular effects detected in the 13-week studies (NTP, 1992). α -Methylbenzyl alcohol, a metabolite of ethylbenzene, may not be involved because it inhibits testicular adenoma (NTP, 1990b).

In addition to inducing renal tubule neoplasms in rats, ethylbenzene exposure may have induced bone marrow hyperplasia characterized by hypercellularity of erythroid and myeloid precursor cells. Cragg et al. (1989) also reported that Fischer 344/N rats exposed

to ethylbenzene at 782 ppm for 4 weeks had a small increase in leukocyte counts. On the other hand, ethylbenzene depressed mononuclear cell leukemia in 750 ppm males, but this was considered to be due largely to reduced survival in this group.

In the 2-year mouse studies, the incidences of alveolar epithelial metaplasia and alveolar/bronchiolar adenoma or carcinoma (combined) were significantly increased in 750 ppm male mice but not in 750 ppm female mice. Estimated differences in inhaled air volume alone (male mice have a greater ventilation volume per body weight than do female mice) could not explain the difference between males and females in their responses to ethylbenzene inhalation.

The incidence of hepatocellular adenoma or carcinoma (combined) was significantly greater in the 750 ppm group of female mice compared to that in the chamber controls. The incidence of liver eosinophilic foci was also significantly greater in the 750 ppm group of female mice. A significant increase in absolute liver weight was observed in male and female mice exposed to ethylbenzene at 750 ppm and higher in the 13-week studies (NTP, 1992). Increased absolute and relative liver weights were also reported in female B6C3F₁ mice exposed to ethylbenzene by inhalation (Cragg *et al.*, 1989). The female mouse liver appears to be more sensitive to the effects of ethylbenzene.

It is also not clear why male and female $B6C3F_1$ mice had different neoplasm responses to ethylbenzene exposure. There are little data available on which to judge why ethylbenzene affected the male and female endocrine systems differently, although an exposure-related increase in the incidence of pituitary gland (pars distalis) hyperplasia was seen in female mice. Ethylbenzene induced an exposure-related increase in the incidences of hyperplasia in the thyroid gland of male and female mice, but there was no difference between males and females in incidence observed.

Phenylglyoxylic and mandelic acids were effective in causing brain dopamine depletion *in vitro* (Mutti and Franchini, 1987) and *in vivo* (Mutti *et al.*, 1988). On the other hand, Andersson *et al.* (1981) reported that

male Sprague-Dawley rats exposed to ethylbenzene by inhalation at 2,000 ppm, 6 hours per day for 3 days had increases in dopamine and noradrenaline levels in the hypothalamus and the median eminence. Such neurotoxic effects would disturb brain function and cause neurobehavioral and neuroendocrine changes and may be related to the gender difference in response to ethylbenzene exposure.

The gender and species differences and organ specificity in the carcinogenic effects of ethylbenzene are unexpected findings. The mechanisms of action of ethylbenzene carcinogenesis in rats and mice remain to be defined.

CONCLUSIONS

Under the conditions of these 2-year inhalation studies, there was clear evidence of carcinogenic activity* of ethylbenzene in male F344/N rats based on increased incidences of renal tubule neoplasms. The incidences of testicular adenoma were also increased. There was some evidence of carcinogenic activity of ethylbenzene in female F344/N rats based on increased incidences of renal tubule adenomas. There was some evidence of carcinogenic activity of ethylbenzene in male $B6C3F_1$ mice based on increased incidences of alveolar/bronchiolar neoplasms. There was some evidence of carcinogenic activity of ethylbenzene in female $B6C3F_1$ mice based on increased incidences of hepatocellular neoplasms.

Exposure of male and female rats to ethylbenzene resulted in increased incidences of renal tubule hyperplasia and increased severities of nephropathy. Exposure of male mice to ethylbenzene resulted in increased incidences of alveolar epithelial metaplasia, syncytial alteration of hepatocytes, hepatocellular hypertrophy, hepatocyte necrosis, and thyroid gland follicular cell hyperplasia. In female mice, ethylbenzene exposure resulted in increased incidences of eosinophilic foci of the liver, pituitary gland pars distalis hyperplasia, and thyroid gland follicular cell hyperplasia.

^{*} Explanation of Levels of Evidence of Carcinogenic Activity is on page 10. A summary of the Technical Reports Review Subcommittee comments and the public discussion on this Technical Report appears on page 12.

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APPENDIX A SUMMARY OF LESIONS IN MALE RATS IN THE 2-YEAR INHALATION STUDY OF ETHYLBENZENE

TABLE A1	Summary of the Incidence of Neoplasms in Male Rats	
	in the 2-Year Inhalation Study of Ethylbenzene	57
TABLE A2	Individual Animal Tumor Pathology of Male Rats	
	in the 2-Year Inhalation Study of Ethylbenzene	62
TABLE A3	Statistical Analysis of Primary Neoplasms in Male Rats	
	in the 2-Year Inhalation Study of Ethylbenzene	84
TABLE A4a	Historical Incidence of Renal Tubule Neoplasms	
	in Chamber Control Male F344/N Rats	91
TABLE A4b	Historical Incidence of Testicular Adenoma	
	in Chamber Control Male F344/N Rats	91
TABLE A5	Summary of the Incidence of Nonneoplastic Lesions in Male Rats	
	in the 2-Year Inhalation Study of Ethylbenzene	92

TABLE A1
Summary of the Incidence of Neoplasms in Male Rats in the 2-Year Inhalation Study of Ethylbenzene^a

	Chamber Control	75 ppm	250 ppm	750 ppm
Disposition Summary	70	50	50	70
Animals initially in study	50	50	50	50
Early deaths	0.0	00	0.0	0.0
Moribund	28	20	26	26
Natural deaths	7	16	11	22
Survivors Terminal sacrifice	15	1.4	13	2
	15	14		
Animals examined microscopically	50	50	50	50
Alimentary System				
Esophagus	(50)	(50)	(50)	(50)
Intestine large, colon	(50)	(48)	(48)	(48)
Sarcoma	1 (2%)			
Intestine large, rectum	(48)	(49)	(48)	(48)
Intestine large, cecum	(46)	(44)	(46)	(39)
Intestine small, duodenum	(48)	(48)	(50)	(50)
Intestine small, jejunum	(42)	(39)	(44)	(34)
Intestine small, ileum	(45)	(44)	(45)	(37)
Liver	(50)	(50)	(50)	(49)
Hepatocellular adenoma		3 (6%)		
Histiocytic sarcoma		1 (2%)		
Osteosarcoma, metastatic, spleen	1 (2%)			
Mesentery	(4)	(5)	(3)	(3)
Lipoma			1 (33%)	
Sarcoma	1 (25%)			
Oral mucosa	(2)		(1)	(1)
Pharyngeal, squamous cell papilloma	2 (100%)		1 (100%)	1 (100%)
Pancreas	(50)	(49)	(50)	(50)
Duct, carcinoma		1 (2%)		
Salivary glands	(50)	(49)	(50)	(50)
Stomach, forestomach	(50)	(50)	(50)	(50)
Stomach, glandular	(50)	(49)	(50)	(50)
Tongue	(1)			
Squamous cell papilloma	1 (100%)			
Cardiovascular System				
Heart	(50)	(50)	(50)	(50)
Endocrine System				
Adrenal cortex	(50)	(50)	(50)	(50)
Osteosarcoma, metastatic, spleen	1 (2%)	• /	` '	. ,
Adrenal medulla	(50)	(50)	(49)	(48)
Osteosarcoma, metastatic, spleen	1 (2%)	, ,	, ,	• •
Pheochromocytoma malignant	,	1 (2%)		2 (4%)
Pheochromocytoma benign	6 (12%)	10 (20%)	6 (12%)	9 (19%)
Bilateral, pheochromocytoma benign	7 (14%)	3 (6%)	3 (6%)	3 (6%)
Islets, pancreatic	(50)	(50)	(50)	(50)
Adenoma	3 (6%)	4 (8%)	4 (8%)	4 (8%)
Carcinoma	2 (4%)	1 (2%)	(- · · ·)	· · · · · · · · · · · · · · · · · · ·
Parathyroid gland	(45)	(46)	(46)	(46)
Adenoma	. ,	1 (2%)	` '	. ,
		• •		

TABLE A1
Summary of the Incidence of Neoplasms in Male Rats in the 2-Year Inhalation Study of Ethylbenzene (continued)

	Chamber Control	75 ppm	250 ppm	750 ppm
Endocrine System (continued)				
Pituitary gland	(49)	(50)	(50)	(45)
Pars distalis, adenoma	23 (47%)	18 (36%)	18 (36%)	18 (40%)
Pars distalis, adenoma, multiple	2 (4%)	1 (2%)	1 (2%)	10 (1070)
Thyroid gland	(50)	(49)	(50)	(50)
Bilateral, C-cell, adenoma	2 (4%)	(10)	(00)	(00)
C-cell, adenoma	1 (2%)	6 (12%)	3 (6%)	2 (4%)
C-cell, carcinoma	2 (4%)	0 (1270)	0 (070)	2 (170)
Follicular cell, carcinoma	1 (2%)	1 (2%)	1 (2%)	1 (2%)
General Body System				
Peritoneum		(1)		
Genital System				
Epididymis	(50)	(50)	(50)	(50)
Preputial gland	(49)	(50)	(49)	(50)
Adenoma	3 (6%)	1 (2%)	1 (2%)	2 (4%)
Bilateral, adenoma	(- · · ·)	,	(/	2 (4%)
Prostate	(50)	(50)	(50)	(50)
Seminal vesicle	(49)	(49)	(50)	(50)
Testes	(50)	(50)	(50)	(50)
Bilateral, interstitial cell, adenoma	27 (54%)	23 (46%)	32 (64%)	40 (80%)
Interstitial cell, adenoma	9 (18%)	10 (20%)	8 (16%)	4 (8%)
Hematopoietic System Bone marrow Histiocytic sarcoma Lymph node Lymph node, bronchial Histiocytic sarcoma Lymph node, mandibular Lymph node, mesenteric	(49) (9) (44) (47) (49)	(49) 1 (2%) (8) (34) 1 (3%) (48) (50)	(50) (9) (39) (49) (50)	(50) (14) (28) (50) (50)
Histiocytic sarcoma	(40)	1 (2%)	(30)	(30)
Lymph node, mediastinal Histiocytic sarcoma	(48)	(48) 1 (2%)	(50)	(47)
Spleen	(50)	(49)	(50)	(50)
Histiocytic sarcoma	(00)	1 (2%)	(00)	(00)
Osteosarcoma	1 (2%)	- (~/0)		
Γhymus	(46)	(44)	(46)	(44)
Histiocytic sarcoma	(/	1 (2%)	(==/	\ = =/
Thymoma benign			1 (2%)	
Integumentary System Mammary gland	(46)	(47)	(46)	(49)
Adenoma			1 (2%)	
Fibroadenoma	2 (4%)	2 (4%)	2 (4%)	
Fibroadenoma, multiple		1 (2%)		
Fibroma		2 (4%)		

TABLE A1
Summary of the Incidence of Neoplasms in Male Rats in the 2-Year Inhalation Study of Ethylbenzene (continued)

•	Chamber Control	75 ppm	250 ppm	750 ppm
Integumentary System (continued)				
Skin	(50)	(50)	(50)	(50)
Basal cell adenoma			1 (2%)	1 (00/)
Basal cell carcinoma Keratoacanthoma	3 (6%)	2 (4%)	2 (4%)	1 (2%) 2 (4%)
Squamous cell papilloma	2 (4%)	1 (2%)	1 (2%)	2 (170)
Pinna, schwannoma benign		1 (2%)		
Pinna, schwannoma malignant			1 (2%)	
Sebaceous gland, adenoma Subcutaneous tissue, fibroma	1 (2%)	1 (2%)	1 (2%) 3 (6%)	
Subcutaneous tissue, fibrosarcoma	1 (270)	1 (2%)	3 (070)	
Subcutaneous tissue, lipoma		- ()		1 (2%)
Subcutaneous tissue, myxoma			1 (2%)	
Subcutaneous tissue, sarcoma		2 (4%)		
Musculoskeletal System				
Bone	(49)	(50)	(50)	(50)
Histiocytic sarcoma	, ,	1 (2%)		, ,
Turbinate, chondroma	400	4.5	1 (2%)	443
Skeletal muscle	(1)	(1)		(1)
Histiocytic sarcoma Osteosarcoma, metastatic, spleen	1 (100%)	1 (100%)		
Sarcoma	1 (10070)			1 (100%)
Navyana Svotom				
Nervous System Brain	(50)	(50)	(50)	(50)
Glioma malignant	(00)	(00)	1 (2%)	1 (2%)
Respiratory System				
Larynx	(40)	(44)	(41)	(35)
Lung	(50)	(50)	(50)	(50)
Alveolar/bronchiolar adenoma	2 (4%)	1 (2%)		1 (2%)
Alveolar/bronchiolar carcinoma	1 (2%)			
Carcinoma, metastatic, thyroid gland Histiocytic sarcoma	2 (4%)	1 (2%)		
Osteosarcoma, metastatic, spleen	1 (2%)	1 (2/0)		
Mediastinum, osteosarcoma, metastatic, sple				
Nose	(49)	(49)	(50)	(50)
Гrachea	(50)	(50)	(50)	(50)
Leiomyosarcoma			1 (2%)	
Special Senses System				
Harderian gland			(1)	
Carcinoma			1 (100%)	
Zymbal's gland	(1)		(1)	(1)
Carcinoma	1 (100%)		1 (100%)	

TABLE A1 Summary of the Incidence of Neoplasms in Male Rats in the 2-Year Inhalation Study of Ethylbenzene (continued)

	Chamber Control	75 ppm	250 ppm	750 ppm
Urinary System				
Kidney	(50)	(50)	(50)	(50)
Histiocytic sarcoma		1 (2%)		
Lipoma	1 (2%)			
Renal tubule, adenoma		3 (6%)	2 (4%)	4 (8%)
Renal tubule, carcinoma			1 (2%)	3 (6%)
Urinary bladder	(49)	(49)	(50)	(49)
Transitional epithelium, papilloma		1 (2%)		
Systemic Lesions Multiple organs ^b Histiocytic sarcoma Leukemia mononuclear Mesothelioma malignant	(50) 27 (54%)	(50) 1 (2%) 26 (52%) 2 (4%)	(50) 32 (64%) 1 (2%)	(50) 9 (18%)
Neoplasm Summary				
Total animals with primary neoplasms ^C	49	45	50	50
Total primary neoplasms	134	131	134	111
Total animals with benign neoplasms	48	44	48	48
Total benign neoplasms	97	95	94	93
Total animals with malignant neoplasms	33	32	37	17
Total malignant neoplasms	37	36	40	18
Total animals with metastatic neoplasms	3			
Total metastatic neoplasms	8			

Number of animals examined microscopically at the site and the number of animals with neoplasm Number of animals with any tissue examined microscopically Primary neoplasms: all neoplasms except metastatic neoplasms

X: Lesion present

Blank: Not examined

TABLE A2 Individual Animal Tumor Pathology of Male Rats in the 2-Year Inhalation Study of Ethylbenzene: Chamber Control

Number of Days on Study	3 5 4	3 9 1	4 7 9	4 9 7	5 2 8	5 4 7	5	7	5 5 7 8 4 4		0	6 0 8		6 1 8	6 2 3	6 3 0	6 3 9	6 4 0	6 4 0	6 4 4	6 5 2	6 5 3	6 6 1	6 6 8
Carcass ID Number	0 4 8	0 3 8	0 0 9	0 3 9	0 2 5	0 0 8	2	4	0 0 3 5 6 0	1	0	0 2 3	0	2	0 3 2	2	0 4 1	0	0 2 8	0 1 1	0 1 2	4	0 1 4	2
Alimentary System																								
Esophagus	+	+	+	+	+	+	+	+	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Intestine large, colon	+	+	+	+	+	+	+	+	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Sarcoma																								
Intestine large, rectum	+	+	+	+	+	+			+ +				+	+	+	+	+	+	+	+	+	+	+	+
Intestine large, cecum	+	+	+	+	+	+	+ ,	A	+ +	. <i>P</i>	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Intestine small, duodenum	+	+	+	+	+	+			+ +			+	+	+	+	+	+	+	+	+	+	+	+	+
Intestine small, jejunum	A	Α	+	Α	+	+	+ ,	A	+ +	. <i>P</i>	+	+	+	+	+	+	+	+	+	Α	+	+	+	+
Intestine small, ileum	A	+	+	+	+	+	+ .	-	+ +			+		+	+	+	+	+	+	Α	+	+	+	+
Liver	+	+	+	+	+	+	+	+	+ +			+	+	+	+	+	+	+	+	+	+	+	+	+
Osteosarcoma, metastatic, spleen										Σ	(
Mesentery				+																				
Sarcoma				X																				
Oral mucosa																								
Pharyngeal, squamous cell papilloma																								
Pancreas	+	+	+	+	+	+	+ .	+	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Salivary glands	+	+	+	+	+	+	+ -	+	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Stomach, forestomach	+	+	+	+	+	+	+ -	+	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Stomach, glandular	+	+	+	+	+	+	+	+	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Tongue								+																
Squamous cell papilloma								X																
Cardiovascular System																								
Blood vessel	+	+	+	+	+	+	+	+	+ +	. 4	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Heart	+	+	+	+	+	+	+	+	+ +	. 4	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Endocrine System Adrenal cortex																								
Osteosarcoma, metastatic, spleen	+	+	+	+	+	+	+	+	+ +	· -}	· +	+	+	+	+	+	+	+	+	+	+	+	+	+
Adrenal medulla						,			, .														+	
	+	+	+	+	+	+	+	+	+ +	· >		+	+	+	+	+	+	+	+	+	+	+	+	+
Osteosarcoma, metastatic, spleen									Σ		`				X								Х	
Pheochromocytoma benign Bilateral, pheochromocytoma benign										_		X			Λ								Λ	X
Islets, pancreatic						,						Λ.												Λ +
Adenoma	+	+	+	+	+	+	+	_	+ +	. +	+	+	+	+	+	+	+	+	+	+	+	+	+	+
										Σ	,													
Carcinoma													1.1						ъ.	1.1				М
Parathyroid gland	+	+	+	+	+	+	+ ·	+	+ +	+		+		+			+	+		M				M
Pituitary gland	+	+ V	+ v	+	+ v	+ V		+	+ +	+		+		+ V		+	+	+	+	+				+
Pars distalis, adenoma		Λ	X		Λ	X	Λ				X		Λ	X	Λ						Λ	X		
Pars distalis, adenoma, multiple																								
Thyroid gland	+	+	+	+	+	+	+	+	+ +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Bilateral, C-cell, adenoma																								
C-cell, adenoma																							٠,	
C-cell, carcinoma																							X	
Follicular cell, carcinoma																								

M: Missing tissue

I: Insufficient tissue

^{+:} Tissue examined microscopically A: Autolysis precludes examination

TABLE A2
Individual Animal Tumor Pathology of Male Rats in the 2-Year Inhalation Study of Ethylbenzene: Chamber Control (continued)

(continued)																										
Number of Days on Study	6		6	6	_	7	7	7	7	7								7		7		7	7	7	7	
Number of Days on Study	8 5		8 9	8 9	9	1 0	0	3	4	5	3 4	3 4	3 4	3 4		3 4										
	0		0	0	0	0			0	0	0	0	0	0				0	0	0	0	0	0	0	0	Total
Carcass ID Number	4		1 7	3	1 3	0 2		0 6	4 6	4 7	0 7	1 0	1 5	1 6	1 8	2 0		2 6	3 0	3 1	3 5	3 7	4 0	4 2	4 9	Tissues/ Tumors
Alimentary System																										
Esophagus	+	· N	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	49
ntestine large, colon Sarcoma	+	. +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+ X	+	+	+	+	50 1
ntestine large, rectum	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	48
ntestine large, cecum	+	. A	+	+	+	+	Α	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	46
ntestine small, duodenum	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	48
ntestine small, jejunum	+	A	+	+	+	+	Α	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	42
ntestine small, ileum	+	A	+	+	+	+	Α	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	45
Liver	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Osteosarcoma, metastatic, spleen																										1
Mesentery			+																+						+	4
Sarcoma																										1
Oral mucosa			+																						+	2
Pharyngeal, squamous cell papilloma			X																						X	2
Pancreas	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Salivary glands	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Stomach, forestomach	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Stomach, glandular	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Tongue Tongue																										1
Squamous cell papilloma																										1
Cardiovascular System																										
Blood vessel	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Heart	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Endocrine System																										
Adrenal cortex	+	. +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Osteosarcoma, metastatic, spleen																										1
Adrenal medulla	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Osteosarcoma, metastatic, spleen																										1
Pheochromocytoma benign										X									X	X						6
Bilateral, pheochromocytoma benign			X			X					X					X										7
slets, pancreatic	+	+	+	+		+		+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+		+	50
Adenoma					X			X																X		3
Carcinoma										X																2
Parathyroid gland	+	+	+	+	+	+	+	+							+			+	+	+	+	+	+	+	+	45
Pituitary gland	+	+	+	+	+	+					+				+			M	+		+	+	+		+	49
Pars distalis, adenoma							X	X	X	X	X	X	X		X		X			X	X			X		23
Pars distalis, adenoma, multiple		X												X												2
Thyroid gland	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Bilateral, C-cell, adenoma																			X	X						2
C-cell, adenoma													X													1
C-cell, carcinoma									X																	2
Follicular cell, carcinoma																									X	1

TABLE A2 Individual Animal Tumor Pathology (continued)	y of Male Rats in the 2-Year Inhalation Study of Ethylbenzene: Chamber Control
Number of Days on Study	3 3 4 4 5 5 5 5 5 5 5 6 6 6 6 6 6 6 6 6 6 6
Carcass ID Number	0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0
General Body System Tissue NOS	+
Genital System Epididymis Preputial gland Adenoma	+++++++++++++++++++++++++++++++++++++++
Prostate Seminal vesicle Testes Bilateral, interstitial cell, adenoma Interstitial cell, adenoma	X + + + + + + + + + + + + + + + + + + +
Hematopoietic System Bone marrow Lymph node Lymph node, bronchial	+ + + + M + + + + + + + + + + + + + + +
Lymph node, mandibular Lymph node, mesenteric Lymph node, mediastinal Spleen	+ + + M + + + + + + + + + + M + + + + +
Osteosarcoma Thymus	X + M + + M + M + + + + + + + + + + + + +
Integumentary System Mammary gland Fibroadenoma Skin	+ + + + + + + + + + + + + + + + + + +
Keratoacanthoma Squamous cell papilloma Subcutaneous tissue, fibroma	X X X
Musculoskeletal System Bone Skeletal muscle Osteosarcoma, metastatic, spleen	+ + + + M + + + + + + + + + + + + + + +
Nervous System Brain	+ + + + + + + + + + + + + + + + + + + +
Respiratory System Larynx Lung Alveolar/bronchiolar adenoma	M M M M M + + + M M + + M M + + + H + + + +
Alveolar/bronchiolar carcinoma Carcinoma, metastatic, thyroid gland Osteosarcoma, metastatic, spleen Mediastinum, osteosarcoma, metastatic, spleen	X X
Nose Trachea	+ + + + M + + + + + + + + + + + + + + +

TABLE A2 Individual Animal Tumor Pathology (continued)	of Male	e R	Rat	s iı	n tl	he 2	2-Y	ear	· In	ha	lat	ion	St	tud	ly o	of I	Eth	ylh	en	zei	ne:	C	ha	ml	ber	Control
Number of Days on Study	6 8 5	6 8 9	8	8	3 9		7 1 0	7 1 3	7 1 4	7 1 5	7 3 4	7 3 4	7 3 4	7 3 4	7 3 4	7 3 4	7 3 4	7 3 4	7 3 4	7 3 4	7 3 4	7 3 4	7 3 4	7 3 4	7 3 4	
Carcass ID Number	0 4 3	0	1	. 3	3 1	. 0	3	0	0 4 6	0 4 7	0 0 7	0 1 0	1		0 1 8	0 2 0	0 2 1	0 2 6	0 3 0	3	0 3 5	0 3 7	0 4 0		4	Total Tissues/ Tumors
General Body System Tissue NOS																										1
Genital System Epididymis Preputial gland Adenoma Prostate Seminal vesicle Testes Bilateral, interstitial cell, adenoma Interstitial cell, adenoma	+ + + + X	+ + + +	- + - + - +	+ + + + + + + + >	+ + + + + + + + X X	+ + + + + + X X	+ + + A + X X	+ + + + X	+ + + + X	+ + + + +	+ + + + X	+ + + + X	+ + + + X	+ + + + X	+ + + + X	+ + X + + X	+ + + + +	+ + + + X	+ + + + X	+ + + + X	+ M + + X	+++++	+ + + + X	+ + + + X	+ + + + X	50 49 3 50 49 50 27
Hematopoietic System Bone marrow Lymph node Lymph node, bronchial Lymph node, mandibular Lymph node, mesenteric Lymph node, mediastinal Spleen Osteosarcoma Thymus	+ + + + + +	+ N + + +		- + + + + + + + + + + + + + + + + + + +	+ + + + + + + + + + + + + + + + + + +	+ + + + + + + +	+ + + +	+ + + +	+ + + + + +	+ + + + + + +	+ + + + + + + +	+ + + + + + +	+ + + + + + + +	+ + + + + + +	+ + + + + + +	+ + + + + + +	+ + + + + + + +	+ + + + + + + +	+ + + + + + + +	+ + + + + + +	+ M + + +	+ + + + + + +	+ + + + + + +	+ + + + + + +	+ + + + + + +	49 9 44 47 49 48 50 1
Integumentary System Mammary gland Fibroadenoma Skin Keratoacanthoma Squamous cell papilloma Subcutaneous tissue, fibroma	+	+	- +	⊦ + ⊦ +	+ + + +	+ +	+	+	+	M +	+				+		+ X +		+	+	+ + X	+	M +	+	+ X +	46 2 50 3 2 1
Musculoskeletal System Bone Skeletal muscle Osteosarcoma, metastatic, spleen	+	+	- 4	⊦ +	⊦ ⊣	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	49 1 1
Nervous System Brain	+	+	- +	+ +	+ +	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Respiratory System Larynx Lung Alveolar/bronchiolar adenoma Alveolar/bronchiolar carcinoma Carcinoma, metastatic, thyroid gland Osteosarcoma, metastatic, spleen	+	+	- +	⊦ + ⊦ +	+ + + +	+ +	+	+ +	+ + X X	+	+ +	+ + X	+ +	+ +	+ +	+ +	+ +	+++	+++	+ +	+ + X	+ +	+	+	+ +	40 50 2 1 2
Mediastinum, osteosarcoma, metastatic, spleen Nose Trachea	+	+	- + - +	⊦ + ⊦ +	⊦	+ + +	+	+	++	+	+	+	++	+	++	++	++	++	++	++	+	+	+	++	++	1 49 50

Individual Animal Tumor Pathol (continued)	logy of Mal	e R	ats	in	th	e 2	- Y (ear	In	ha	lat	ior	ı St	tud	ly o	of 1	Eth	yll	en	ıze	ne	• (Cha	m	ber	Control
	3	3	4	4	5	5	5	5	5	5	5	6	6	6	6	6	6	6	6	6	6	6	6	6	6	
Number of Days on Study	5	9	7	9	2	4	5	7	7	8	9	0	0	0	1	2	3	3	4	4	4	5	5	6	6	
•	4	1	9	7	8	7	3	0	4	4	0	2	8	9	8	3	0	9	0	0	4	2	3	1	8	
	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	
Carcass ID Number	4	3	0	3	2	0	2	4	3	5	1	0	2	0	2	3	2	4	0	2	1	1	4	1	2	
	8	8	9	9	5	8	4	5	6	0	9	3	3	5	9	2	7	1	4	8	1	2	4	4	2	
Special Senses System Eye										+																
Zymbal's gland Carcinoma																									+ X	
Urinary System																										
Kidney Lipoma	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Urinary bladder	+	+	+	+	+	+	+	A	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Systemic Lesions																										
Multiple organs	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Leukemia mononuclear								X	X	X			X			X	X	X	X	X	X	X	X		X	

Individual Animal Tumor Patho (continued)	ology of Male	R	ats	s in	th	e 2	- Y (ear	· Ir	ıha	lat	ior	s St	tud	y o	f I	Eth	yll	en	ze	ne:	C	h	ım	bei	Control
Number of Days on Study	6 8 5	6 8 9	6 8 9	6 8 9	6 9 2	7 1 0	7 1 0	7 1 3	7 1 4	7 1 5	7 3 4															
Carcass ID Number	0 4 3	0 0 1	0 1 7	0 3 3	0 1 3	0 0 2	0 3 4	0 0 6	0 4 6	0 4 7	0 0 7	0 1 0	0 1 5	0 1 6	0 1 8	0 2 0	0 2 1	0 2 6	0 3 0	0 3 1	0 3 5	0 3 7	0 4 0	0 4 2	0 4 9	Total Tissues/ Tumors
Special Senses System Eye Zymbal's gland Carcinoma																										1 1 1
Urinary System Kidney Lipoma Urinary bladder	+	+	+	+	+	+	+	+	+	+	+	+	+	+ X +	+	+	+	+	+	+	+	+	+	+	+	50 1 49
Systemic Lesions Multiple organs Leukemia mononuclear	+ X	+ X	+ X	+	+ X	+ X	+ X	+	+	+ X	+	+ X	+	+	+	+	+ X	+ X	+ X	+ X	+	+ X	+ X		+	50 27

	9	3	3	4	5	5	5	5 5	5 1	5 5	5	6	6	6	6	6	6	6	6	6	6	6	6	6
Number of Days on Study	4	0	5	1	1			6 8		88	9	0	0		1		1	2	3	3	3	3	4	5
Tumber of Days on Study	4	8	2					0 2						7		7		6	1	4	7	9	0	
	1	1	1	1	1	1	1	1 1	1	1 1	1	1	1	1	1	1	1	1	1	1	1	1	1	1
Carcass ID Number	2	0	5	0	2) 2			0		4	4				1	1	4		0
	2	3						7 2													9			
Alimentary System																								
Esophagus	+	+	+	+	+	+	+	+ -	+ -	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Intestine large, colon	+	Α	+	Α	+	+	+	+ -	+ -	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Intestine large, rectum	+	Α	+	+	+	+	+	+ -	+ -	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Intestine large, cecum	+	Α	Α	Α	+	+	+	+ -	+ -	+ +	+	Α	+	+	+	+	+	+	Α	+	+	+	+	+
Intestine small, duodenum	+	A	+	+	+	+	+	+ -	+ -	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Intestine small, jejunum	+	A	A	A	+			+ -		+ +		Α		A	+	+	+	+	A	A	+	+	+	+
Intestine small, ileum	+	Α	A	+	+					+ +			+	+	+	+	+	+	A	+	+	+	+	
Liver	+	+	+	+	+	+	+		+ -	+ +	+	+	+	+	+	+	+	+	+	+	+	+		
Hepatocellular adenoma								X					17										X	
Histiocytic sarcoma													X											
Mesentery	+									+					+		,							
Pancreas Duct carcinoma	+	A	+	+	+	+	+	+ -	+ -	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Duct, carcinoma Salivary glands		.1	J.	_	_			_	_	μ,	1	ر	_	_	+	M	_	+	۔	ر	J	.1		_
Sanvary grands Stomach, forestomach	+		+	+	+	T +	+	+ ·	 -	r + L J		+	+	+ _	+	141	T +	+	+	+	+		+	⊤
Stomach, glandular	⊤	Δ	+	+	+	+	+	+ -	 + .	, T + +	+	+	+	+	+	+	+	+	+	+	+	+	+	+
-																								
Cardiovascular System																								
Blood vessel	+	+	+	+	+	+	+	+ -	+ -	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Heart	+	+	+	+	+	+	+	+ -	+ -	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Endocrine System																								
Adrenal cortex	+	+	+	+	+	+	+	+ -	+ -	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Adrenal medulla	+	+	+	+	+	+	+	+ -	+ -	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Pheochromocytoma malignant																								
Pheochromocytoma benign										X								X						
Bilateral, pheochromocytoma benign																								
Islets, pancreatic	+	+	+	+	+	+		+ -	+ -	+ +	+	+	+	+	+	+	+	+	+	+	+	+		+
Adenoma								X															X	
Carcinoma																								
Parathyroid gland	M	[+	+	+	+	+		+ -	+ -	+ +	+	+	+	M	+	+	+	+	+	+	+	+	+	+
Adenoma							X																	
Pituitary gland	+	+	+	+	+ V	+	+ V			+ + v	+		+	+	+	+ V	+	+	+	+	+	+		
Pars distalis, adenoma					X		X	Χ	2	X		X				X	X						X	X
Pars distalis, adenoma, multiple		٨															,	,						
Thyroid gland C-cell, adenoma	+	А	+	+	+	+	+	+ -	+ -	+ +	+	+	+	+	+	+ X	+	+	+	+	+	+	+	+
C-ceii, adenoma Follicular cell, carcinoma																Λ								
i omequal cen, caremonia																								
General Body System																								
Peritoneum																								
Genital System																								
Epididymis	+	+	+	+	+	+	+	+ -	+ -	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Preputial gland	+	+	+	+	+	+	+	+ -	+ -	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Adenoma								X																
Prostate	+	+	+	+	+	+	+	+ -	+ -	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Seminal vesicle	+	A	+	+	+	+	+	+ -	+ -	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Testes	+	+	+	+	+	+	+	+ -	+ -	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Bilateral, interstitial cell, adenoma						X					X		X			X				X				
Interstitial cell, adenoma								7	X	X				X								X	X	

	-	,	,	,	,	-	~	~	-	~	~	~	~	~	~	~	~		~	~					~	
	6	6		6	6	7	7	7	7	7		7	7	7	7	7	7	7	7	7	7	7	7	7	7	
Number of Days on Study	6 8	7	9	9	9 5	0 7	1	1 4	1	2 0	2	3 4														
	1	1	1	1	1	1	1	1		1	1	1	1	1	1		1	1	1	1	1	1				Total
Carcass ID Number	2	3	4	2	1 2	1	1 4	1	1 2	1	1 4	0	1		1	1	1 1	1	1 2		3	1	3		1 4	Tissues/
Carcas ID Tuniber	3	5			7																					Tumors
Alimentary System																										
Esophagus	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Intestine large, colon	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	48
Intestine large, rectum	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	49
Intestine large, cecum	+	+	+	+	A	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	44
Intestine small, duodenum	+	+	+	+	+	+	Α	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	48
Intestine small, jejunum	+	A	. A	+	+	+	A	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	39
Intestine small, ileum	+	+	A	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	44
Liver	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Hepatocellular adenoma Histiocytic sarcoma	X																									3 1
Mesentery						+														+						5
Pancreas	+	+	+	+	+	+	+	+	+	+	+		+	+	+	+	+	+	+	+	+	+	+	+	+	49
Duct, carcinoma												X														1
Salivary glands	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	49
Stomach, forestomach	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Stomach, glandular	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	49
Cardiovascular System																										
Blood vessel	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Heart	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Endocrine System																										
Adrenal cortex	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Adrenal medulla	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Pheochromocytoma malignant																	X									1
Pheochromocytoma benign					X			X	X			X					X		X		X			X		10
Bilateral, pheochromocytoma benign						X									X							X				3
Islets, pancreatic	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Adenoma								X												X						4
Carcinoma																X						٠.				1
Parathyroid gland	+	+	+	+	+	+	+	+	+	+	+	+	M	+	+	+	+	+	+	+	+	M	+	+	+	46
Adenoma																										1
Pituitary gland	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+			+	+	+	+	+	+	+	+	50
Pars distalis, adenoma	Х	X								X				Х	X		Х	X	Х		X					18
Pars distalis, adenoma, multiple																X										1
Thyroid gland C-cell, adenoma	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+		+			+	+	+	49
C-cen, adenoma Follicular cell, carcinoma									Λ	X		X	X						X		X					6 1
,													Λ										_	_		1
General Body System Peritoneum											+															1
											•												—	—		
Genital System				,																	,					F0.
Epididymis	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Preputial gland	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Adenoma																										1
Prostate	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Seminal vesicle	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	49
Pastas										- 1			+	+	+	+	+	+	+	+	+				+	50
Testes Bilateral, interstitial cell, adenoma	+	+	+	+ X	17	_	X	-	X	т	37	X	37		v	1/2	37	37	X	37		37	X	17		23

Individual Animal Tumor Patholog	<i>y</i> 01 11 141 1.	_													_			_						_	
	2	;	3	, 4	4 5				5					6								6	6	6	-
Number of Days on Study	4) 5		1 1					8	8	9		0			1			3	3	3	3	4	
	4	8	3 2	. (8 6	8	9	0	2	4	4	6	0	1	7	5	7	9	6	1	4	7	9	0	4
	1		l 1		1 1	. 1	. 1	1	1			1	1	1	1	1	1	1	1	1	1	1	1	1	1
Carcass ID Number	2) 5				2 4					0	3		3		4	0			1			3	
	2	;	3 () {	8 9) 4	9	7	2	4	1	7	2	1	8	4	6	5	3	5	1	9	0	7	9
Hematopoietic System																									
Bone marrow	+	. 1	4 -		+ +		+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Histiocytic sarcoma														X											
Lymph node						4	+				+											+			
Lymph node, bronchial	+	- 1	M -		+ +	- N	1 +	+	+	+	+	M	+	+	+	+	+	M	+	+	+	+	M	+	M
Histiocytic sarcoma														X											
Lymph node, mandibular	+	1	A -		+ +	+ +	+ +	+	- +	+	+		+							+	+	+	+	+	+
Lymph node, mesenteric	+	-	+ +		+ +		+ +	+	+	+	+	+	+		+	+	+	+	+	+	+	+	+	+	+
Histiocytic sarcoma					, 1	π.				у. л		,	,	X		,		,	,						
Lymph node, mediastinal Histiocytic sarcoma	+		- +	-	- I	v1 ∃	- +	. +	+	11/1	+	+	+	+ X	+	+	+	+	+	+	+	+	+	+	T
Spleen	_		Δ _						+	_	_	_	+		+	+	+	+	+	_	_	_	_	_	+
Histiocytic sarcoma		1	. 7		. 7	, ,	Т.	-T	7	т	г		-	X	-	-	-	-	-	г	т	т'	т	Τ'	•
Thymus	+		+ N	Λ.	+ +	- 1	√ 1 →	. 4	- M	М	+	+	+		+	+	+	+	+	+	+	+	+	+	M
Histiocytic sarcoma			•	_		-	- '	Ċ	.,,			•	•	X			•				•	•	•	•	· -
International Creaters																									
Integumentary System		,												3.7											
Mammary gland	+	1	M -		+ +		+ +	. +	- +	+	+	+	+	M	+	+	+	+	+	+	+	+	+	+	+
Fibroadenoma																									
Fibroadenoma, multiple Fibroma							τ.	,																	
Skin							Χ	٠.	+																
Keratoacanthoma	+				+ +		- +	. +	- +	+	+	+	+	+	+	+	+	+	+	+	+	X	+	+	+
Squamous cell papilloma																						71			
Pinna, schwannoma benign																									
Subcutaneous tissue, fibroma																									
Subcutaneous tissue, fibrosarcoma								Χ	(
Subcutaneous tissue, sarcoma																									
Museuleskalatal System																									
Musculoskeletal System Bone	_		- -							_	_	_	+	+	+	+	+	+	+	_	_	_	_	_	+
Histiocytic sarcoma			. 7		. 7	, ,	Т.	-T	7	т	г		-	X	-	-	-	-	-	г	т	т'	т	Τ'	•
Skeletal muscle														+											
Histiocytic sarcoma														X											
Nervous System																									
Brain	+		+ +	<u>.</u>	+ +		+ +	. +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
	·								-	_	_		•				-	•		_	-	_	_	_	
Respiratory System				_			-																		
Larynx	+	1	A N	Λ.					- M											+	+	+	+	+	+
Lung	+	-	+ +		+ +		+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Alveolar/bronchiolar adenoma														17											
Histiocytic sarcoma			۸ .											X				,							
Nose Trachea	+	1	A +		+ +		+ +	+	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
I I a Clied	+	_	-		+ +	- +	- +	. +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Special Senses System																									
Eye																									

	_			_	^	~	~	~	~	~	~	~	~	~	~	~	~	~	~	~	~	~	~	~	~	
Name have of Davis on Charles	6	(/	1		7	/	_	7	7		7				7	1	7	7		7		
Number of Days on Study	6 8	1	7 9 l 2			7	3	1 4	8	0	2 3	3 4														
	1	1	1 1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	Total
Carcass ID Number	2	9				3	4	3	2	1	4	0	0	1		1	1	1	2	2	3	3	3	4		Tissues
	3				7										2				0					7		Tumors
Hematopoietic System																										
Bone marrow	+	-	+ +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	49
Histiocytic sarcoma																										1
Lymph node				+ -	-			+							+						+		+			8
Lymph node, bronchial	+	-	+ N	Λ +	+	M	M	+	M	M	+	+	M	+	M	+	+	M	M	+	+	+	M	+	+	34
Histiocytic sarcoma																										1
Lymph node, mandibular	+	-	+ +	- +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	48
Lymph node, mesenteric	+	-	+ +	- +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50 1
Histiocytic sarcoma Lymph node, mediastinal	_	_				_	_	_	_		_	_	_		_		_	_	_	_		_	_		_	48
Histiocytic sarcoma	7			_		т	_	_	т	_	т	_	т	_	т	_	_	т	т	_	_	_	т	_	_	1
Spleen	+	_	+ 4	- 4	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	49
Histiocytic sarcoma			. '	ľ						•	Ċ	•	•	•		•		•	•						•	1
Thymus	+		+ +	- 4	- +	+	+	M	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	44
Histiocytic sarcoma																										1
Integumentary System																										
Mammary gland	+		+ +	- 4	- +	+	M	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	47
Fibroadenoma																			X					X		2
Fibroadenoma, multiple														X												1
Fibroma						X																				2
Skin	+	-	+ +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Keratoacanthoma									X																	2
Squamous cell papilloma														X												1
Pinna, schwannoma benign																						X				1
Subcutaneous tissue, fibroma							X																			1
Subcutaneous tissue, fibrosarcoma	3.					3.7																				1
Subcutaneous tissue, sarcoma	Х					X																				2
Musculoskeletal System																										
Bone	+	-	+ +	- +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Histiocytic sarcoma																										1
Skeletal muscle																										1
Histiocytic sarcoma																										1
Nervous System																										5.0
Brain	+	_	+ +	- +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Respiratory System																										
Larynx	+	-	+ +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	44
Lung	+	-	+ +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Alveolar/bronchiolar adenoma							X																			1
Histiocytic sarcoma																										1
Nose	+	-	+ +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	49
Trachea	+	_	+ +	- +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Special Senses System																										
- r																										

TABLE A2 Individual Animal Tumor Patholog	y of Mal	e R	ats	in	th	e 2	- Y	ear	· In	ıha	ılat	ioi	ı St	tud	ly (of l	Eth	yll	ben	ze	ne	: 7	′5 ֈ	pr	n (c	ontinued)
	2	3	3	4	5	5	5	5	5	5	5	5	6	6	6	6	6	6	6	6	6	6	6	6	6	
Number of Days on Study	4	0	5	1	1	3	4	6	8	8	8	9	0	0	0	1	1	1	2	3	3	3	3	4	5	
· ·	4	8	2	6	8	8	9	0	2	4	4	6	0	1	7	5	7	9	6	1	4	7	9	0	4	
	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	
Carcass ID Number	2	0	5	0	2	2	4	1	4	0	2	0	3	0	3	4	4	0	3	1	1	1	4	3	0	
	2	3	0	8	9	4	9	7	2	4	1	7	2	1	8	4	6	5	3	5	1	9	0	7	9	
Urinary System																										
Kidney	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Histiocytic sarcoma														X												
Renal tubule, adenoma																	X		X							
Urinary bladder	+	Α	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Transitional epithelium, papilloma																										
Systemic Lesions																										
Multiple organs	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Histiocytic sarcoma														X												
Leukemia mononuclear									X		X	X			X	X	X		X		X	X	X		X	
Mesothelioma malignant																										

Individual Animal Tumor Patholog	y of Mal	e R	ats	in	th	e 2	-Ye	ar	In	ha	lat	ion	St	tud	ly o	of l	Eth	ıyl	bei	ıze	ne	: 7	′5 լ	pn	n (c	ontinued)
	6	6	6	6	6	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	
Number of Days on Study	6	7	9	9	9	0	1	1	1	2	2	3	3	3	3	3	3	3	3	3	3	3	3	3	3	
	8	1	2	3	5	7	3	4	8	0	3	4	4	4	4	4	4	4	4	4	4	4	4	4	4	
	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	Total
Carcass ID Number	2	3	4	2	2	3	4	3	2	1	4	0	0	1	1	1	1	1	2	2	3	3	3	4	4	Tissues/
	3	5	1	5	7	0	3	9	8	6	5	2	6	0	2	3	4	8	0	6	1	4	6	7	8	Tumors
Urinary System																										
Kidney	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Histiocytic sarcoma																										1
Renal tubule, adenoma								X																		3
Urinary bladder	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	49
Transitional epithelium, papilloma																X										1
Systemic Lesions																										
Multiple organs	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Histiocytic sarcoma																										1
Leukemia mononuclear		X		X	X			X	X	X	X	X	X			X		X			X		X	X	X	26
Mesothelioma malignant			X								X															2

			4		r	r	r	E	E	E	-	E	E			0 4	,		,	c	e	e	c	c	C
Normhan of Davis on Ctude.	2		_	4	5	5				5				5 (3 (6	
Number of Days on Study	6 6	2 0	6 7	9 6	0 9	2 7										2 2 4 9		4 4					7 1	7 1	8 6
	2	2	2	2	2	2	2	2	2	2	2	2	2	2 :	2 :	2 2	2 :	2 2	2	2	2	2	2	2	2
Carcass ID Number	3 8	3 7	1 9	0 4	0 9	4		0 3						2 3 5 5		3 : 0 :		4 2 5 (0 1	3 5	5 0
Alimentary System																							_	_	
Esophagus	+	+	+	+	+	+	+	+	+	+	+	+	+	+ -	+ -	+ -	+ -	+ -	+	+	+	+	+	+	+
Intestine large, colon	+	+	+	+	Α	+	+	+	+	+	+	+	+	+ -	+ -	+ -	+ ,	Α -	+	+	+	+	+	+	+
Intestine large, rectum	+	+	+	+	Α	+	+	+	+	+	+	+	+	+ -	+ -	+ -	+ .	Α -	+	+	+	+	+	+	+
Intestine large, cecum	A	+	+	+	Α	+	+	+	+	+	+	+	+	+ 1	Α .	+ -	+ .	Α -	+	+	+	+	+	+	+
Intestine small, duodenum	+	+	+	+	+	+	+	+	+	+	+	+	+	+ -	+ -	+ -	+ -	+ -	+	+	+	+	+	+	+
Intestine small, jejunum	A	+	+	+	Α	+	+	+	+	+	+	+	+	+ 1	Α.	+ -	+ -	+ -	+	+	+	+	Α	+	+
Intestine small, ileum	A	+	+	+	Α	+	+	+	+	+	+	+	+	+ -	+ -	+ -	+ .	Α -	+	+	+	Α	+	+	+
Liver	+	+	+	+	+	+	+	+	+	+	+	+	+	+ -	+ -	+ -	+ -	+ -	+	+	+	+	+	+	+
Mesentery																-	+								+
Lipoma																									X
Oral mucosa																									
Pharyngeal, squamous cell papilloma																									
Pancreas	+	+	+	+	+	+	+	+	+	+	+	+	+	+ .	+ -	+ -	+ -	+ -	+	+	+	+	+	+	+
Salivary glands	+	+	+	+	+	+	+	+	+	+	+	+	+	+ -	+ -	+ -	+ -	+ -	+	+	+	+	+	+	+
Stomach, forestomach	+	+	+	+	+	+	+	+	+	+	+	+	+	+ -	+ -	+ -	+ -	+ -	+	+	+	+	+	+	+
Stomach, glandular	+	+	+	+	+	+	+	+	+	+	+	+	+	+ ·	+ ·	+ -	+ ·	+ -	+	+	+	+	+	+	+
Cardiovascular System																									
Blood vessel	+	+	+	+	+	+	+	+	+	+	+	+	+	+ -	+ .	+ -	+ -	+ -	+	+	+	+	+	+	+
Heart	+	+	+	+	+	+	+	+	+	+	+	+	+	+ -	+	+ -	+ .	+ -	+	+	+	+	+	+	+
				_	_																		—		
Endocrine System																									
Adrenal cortex	+	+	+	+	+	+	+	+	+	+	+	+	+	+ -	+ -	+ -	+ -	+ -	+	+	+	+	+	+	+
Adrenal medulla	+	+	+	+	+	+	+	+		+	M	+	+	+ -	+ -	+ -	+ -	+ -	+	+	+	+		+	+
Pheochromocytoma benign										X													X		
Bilateral, pheochromocytoma benign																				X					
Islets, pancreatic	+	+	+	+	+	+	+	+	+	+	+	+	+	+ .	+ -	+ -	+ -	+ -	+	+	+	+		+	+
Adenoma																							X		
Parathyroid gland	+	+	+	+	+	+	+	+	+	+	+	+		Μ -		+ -	+ -	+ -	+	+	+	+	+	+	+
Pituitary gland	+	+	+	+	+	+			+	+	+	+		+ ·			+ -			+	+	+	+	+	+
Pars distalis, adenoma		Χ	X			X	X	X						X		X		2	X			X		X	
Pars distalis, adenoma, multiple																									
Thyroid gland	+	+	+	+	+	+	+	+	+	+	+			+ -	+ .	+ -	+ -	+ -	+	+ V	+	+	+	+	+
C-cell, adenoma Follicular cell, carcinoma													X							X					
2 omediai con, caremonia																							_	_	
General Body System																									
None																									
Genital System																									
Epididymis	+	+	+	+	+	+	+	+	+	+	+	+	+	+ -	+ -	+ -	+ -	+ -	+	+	+	+	+	+	+
Preputial gland	+	+	+	+	+	+	+	+	+	+	+	+	+	+ -	+ -	+ -	+ -	+ -	+	M	+	+	+	+	+
Adenoma																									
Prostate	+	+	+	+	+	+	+	+	+	+	+	+	+	+ -	+ -	+ -	+ -	+ -	+	+	+	+	+	+	+
Seminal vesicle	+	+	+	+	+	+	+	+	+	+	+	+	+	+ -	+ -	+ -	+ -	+ -	+	+	+	+	+	+	+
Testes	+	+	+	+	+	+	+	+	+	+	+	+	+	+ -	+ -	+ -	+ -	+ -	+	+	+	+	+	+	+
Bilateral, interstitial cell, adenoma										X				X	X			X		X	X	X	X	X	X
Interstitial cell, adenoma		χ			X				X						,	X	V								

6	6	7	- /	- /	- /	/	/	/	- /	- /	- /	- /	7	7	7	7	7	7	/	-/-	- /	- /	- /	7	
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2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	Total
1	0	3	4	4	0	2	4	3	1	3	3	0	0	0	1	1	1	1	1	2	2	4	4	4	Tissues/
8	6	4	6	8	7	0	4	9	5	1	6	2	5	8	0	1	2	3			7	0	1	7	Tumors
+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
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+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
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	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
-T	+	→	T	+	+		+	⊤	+	_	_	+	+	+	+	+	_	_	+	+	+	_	_	+	50
+					т.	т	т	-T	-	-	-		7	77	т,	7"	7	-	-	-T	т	-T	-T	-	
Y	×	¥		X	X	Y		X		X		X	X	X	X	X	X	X	X		V	Y	Y	X	32
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Individual Animal Tumor Patholo	gy of Mal	e .	Kai	.5 1	U	ne <i>i</i>	Z- Y	ear	r In	ha	lat	ion	St	ud	y o	t E	ith;	yIb	en	zei	1e:	Z.	5U	p p	m (continued)
	2		4 4	1	4 5	5 5	5	5	5	5	5	5	5	5	6	6	6	6	6	6	6	6	6	6	6
Number of Days on Study	6		2 (9 (2	2	5	8	9	9	9	9	9		2	2	4	4	6	6	6	7	7	
JJ	6		0		6 9		8	6	9				1	6					4	5		9	1		
	2		2 2	2	2 2	2 2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2
Carcass ID Number	3		3	l	0 () 4	4	0	3	2	2	2	4	2	3	3	1	4	2	1	2	2	0	3	5
	8		7 9)	4 9	3	2	3	3	2	3	8	9	5	2	0	7	5	6	6	9	4	1	5	0
Hematopoietic System																									
Bone marrow	+	_	+ -	+	+ -	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Lymph node											+			+					+						
Lymph node, bronchial	+	-	Μ -	+	+ -	- N	4 +	M	+	M	+	+	M	+	+	M	+	+	+	+	M	+	+	+	+
Lymph node, mandibular	+	-	+ -	+	+ -	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Lymph node, mesenteric	+	-	+ -	+	+ -	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Lymph node, mediastinal	+	-	+ -	+	+ -	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Spleen	+	-	+ -	+	+ -	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Thymus	+	-	Μ -	+	М -	+ +	+	M	+	+	+	+	M	+	+	+	+	+	+	+	+	+	+	+	+
Thymoma benign																									
Integumentary System																									
Mammary gland	+	-	+ -	+	+ -	+ +	+	+	+	+	+	M	+	+	+	+	M	+	+	+	+	M	+	+	+
Adenoma																		X							
Fibroadenoma																									
Skin	+	-	+ -	+	+ -	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Basal cell adenoma																									
Keratoacanthoma																									
Squamous cell papilloma																									
Pinna, schwannoma malignant																			X						
Sebaceous gland, adenoma																								X	
Subcutaneous tissue, fibroma																									
Subcutaneous tissue, myxoma	X	ζ.																							
Musculoskeletal System																									
Bone	4	_	+ -	+	+ -	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Turbinate, chondroma	'				-				,		•	•			•	•	•		•	•	•				
Nervous System																									
Brain											,	,	,							,		,			
	+		+ - X	г	+ -	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	т
Glioma malignant			Λ																						
Peripheral nerve Spinal cord													,												+
													+												т
Respiratory System		ī		ır	۱.						3.5		1	3.4								1 •			
Larynx	N	/1	+ 1	VI	IVI -	+ +	+	M	+	+	M	+	M	M	+	+	+	+	+	+	+	M	+	+	+
Lung	+	-	+ -	+	+ -	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Nose	+	-	+ -	+	+ -	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Trachea	+	-	+ -	+	+ -	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Leiomyosarcoma																									
Special Senses System																									
Harderian gland																									+
Carcinoma																									X
Zymbal's gland																									
Carcinoma																									
Urinary System																									
Kidney	+	-	+ -	+	+ -	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Renal tubule, adenoma																								X	
Renal tubule, carcinoma																								-	
Urinary bladder	4	-	+ -	+	+ -	- +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+

TABLE A2 Individual Animal Tumor Patholo	ogy of Male	e I	Ra	ts	in	the	e 2	- Y	ear	· In	ha	lat	ioı	ı S	tud	ly (of :	Etł	ıyl	ber	ıze	ne	: 2	50	pp	m	(continued)
Number of Days on Study	6 8 8	9	9	7 0 3	7 0 3	7 0 3	7 0 4	7 0 7	7 0 8	7 1 2	7 1 3	7 2 0	7 2 3	7 3 4	7 3 4	7 3 4	7 3 4	7 3 4	7 3 4	7 3 4	7 3 4	7 3 4	7 3 4	7 3 4	7 3 4	7 3 4	
Carcass ID Number	2 1 8	(0	2 3 4	2 4 6	2 4 8	2 0 7	2 2 0	2 4 4	2 3 9	2 1 5	2 3 1	2 3 6	2 0 2	2 0 5	2 0 8	2 1 0	2 1 1	2 1 2	2 1 3	2 1 4	2 2 1	2 2 7	2 4 0	2 4 1	2 4 7	Total Tissues/ Tumors
Hematopoietic System Bone marrow Lymph node Lymph node, bronchial Lymph node, mandibular Lymph node, mesenteric Lymph node, mediastinal Spleen Thymus Thymoma benign	+ + + + +	-	+ + + + + + +	+ + + + + + + + +	+ + + + + + +	+ + + + + + +	+ + M + + +	+ + + + + +	+ + + + + + +	+ + + + + + + +	+ + + + + + + +	+ M + + + +	+ + + + + +	+ + + + + +	+ + + + + + +	+ + + + + X	+ + + + + + +	+ + + + + + +	+ + + + + + +	+ + + + + + +	+ M + + + +	+ + + 1 M + + + + +	+ + + + +	+ + + + + + +	+ M + + + +	+ [+ + + +	50 9 39 49 50 50 50 46
Integumentary System Mammary gland Adenoma Fibroadenoma Skin Basal cell adenoma Keratoacanthoma Squamous cell papilloma Pinna, schwannoma malignant Sebaceous gland, adenoma Subcutaneous tissue, fibroma Subcutaneous tissue, myxoma	+ + X		+	M +	+	+	+	+	+	+	+	+ X +	+	+	+	+ X + X	+	+ + X	+	+ + X	+	+	+	+ + X	+	+	46 1 2 50 1 2 1 1 1 3
Musculoskeletal System Bone Turbinate, chondroma	+		+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+ X	+	+	+	+	50 1
Nervous System Brain Glioma malignant Peripheral nerve Spinal cord	+	-	+	+	+	+ + + +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50 1 2 3
Respiratory System Larynx Lung Nose Trachea Leiomyosarcoma	+ + + +	-	+ + +	+ + +	+ + + +	+ + + +	+ + + +	+ + + +	M + +	+ + + +	+ + + +	+ + + +	+ + + +	+ + + X	+ + + +	+ + + +	+ + + +	+ + + +	+ + + +	+ + + +	+ + + +	+ + + +	+ + + +	+ + + +	+ + + +	+ + + +	41 50 50 50 1
Special Senses System Harderian gland Carcinoma Zymbal's gland Carcinoma			-														-				+ X						1 1 1 1
Urinary System Kidney Renal tubule, adenoma Renal tubule, carcinoma Urinary bladder	+	-	+	+	+	+	+	+	+	+	+	+	+	+	+	+ X +	+	+	+	+	+	+	+	+	+ X	+	50 2 1 50

TABLE A2 Individual Animal Tumor Pathologore	gy of Male	R	ats	in	th	e 2	- Y (ear	In	ha	lat	ior	s St	tud	ly (of 1	Eth	ıyll	ber	ıze	ne	2	50	pp	m	(continued)
	2	4	4	4	5	5	5	5	5	5	5	5	5	5	6	6	6	6	6	6	6	6	6	6	6	
Number of Days on Study	6	2	6	9	0	2	2	5	8	9	9	9	9	9	1	2	2	4	4	6	6	6	7	7	8	
•	6	0	7	6	9	7	8	6	9	0	0	1	1	6	9	4	9	3	4	5	8	9	1	1	6	
	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	
Carcass ID Number	3	3	1	0	0	4	4	0	3	2	2	2	4	2	3	3	1	4	2	1	2	2	0	3	5	
	8	7	9	4	9	3	2	3	3	2	3	8	9	5	2	0	7	5	6	6	9	4	1	5	0	
Systemic Lesions																										
Multiple organs	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Leukemia mononuclear				X	X			X	X	X	X	X	X	X				X	X	X	X					
Mesothelioma malignant						X																				

Individual Animal Tumor Patho	logy of Male	R	ats	in	th	e 2	-Y	ear	In	ha	lat	ior	ı S	tud	ly (of l	Eth	yll	en	ıze	ne:	2	50	pp	m	(continued)
	6	6	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	
Number of Days on Study	8	9 6	0 3	U	0 3	0 4	0 7	0 8	1 2	1 3	2 0	2 3	3 4	3 4	3 4	3 4	-	3 4								
Carcass ID Number	-	2 0	•	2 4	2 4	2 0	2 2	2 4	2	2 1	2	2	2 0	0	2 0	2 1	2 1	2 1	2 1	2 1	2 2	2	4	2 4	2 4	Total Tissues/
Systemic Lesions	8	6	4	6	8	7	0	4	9	5	1	6	2	5	8	0	1	2	3	4	1	7	0	1	7	Tumors
Multiple organs Leukemia mononuclear Mesothelioma malignant	+ X	+ X	+ X	+ X	+ X	+ X	+ X	+ X	+ X	+ X	+ X	+ X	+	+	+ X	+ X	+	+ X	+ X	+	+	+ X	+ X	+ X	+	50 32 1

Individual Animal Tumor Pathology	oi mai	_													٠	<i>J</i>			J						1.1		
	3		3 3	3	4	5	5	5	5	5	5	5	5	5	5	5	5	5	5	5	5	5	5	5	5	5	
Number of Days on Study	7		3 9	9	8	0	0	3	3	4	4	5	6	6	6	7	8	8	8	8	8	8	9	9	9	9	
, ,	7	:	3 3	3	3	0	7	8	9	6	7	5	3	7	9	2	4	6	7	7	7	8	0	0	1	4	
	3	:	3 :	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	
Carcass ID Number	4		1 2	2	2	4	3	1	5	1	0	0	2	0	2	1	2	1	1	1	2	0	1	3	4	4	
	4	•	7 9	9	7	3	6	5	0	6	9	8	1	4	6	7	4	3	1	9	5	6	4	7	5	9	
Alimentary System																											
Esophagus	+		+ -	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Intestine large, colon	+		+ 1	4	+	+	+	+	+	+	+	+	+	+	Α	+	+	+	+	+	+	+	+	+	+	+	
Intestine large, rectum	+		+ -	+	+	+	+	+	+	+	+	+	+	+	+	+	Α	+	+	+	+	+	+	+	+	+	
Intestine large, cecum	+		+ 1	4	Α	+	Α	+	+	+	Α	Α	+	+	Α	Α	Α	+	+	+	+	+	+	+	+	+	
Intestine small, duodenum	+		+ -	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Intestine small, jejunum	+	. ,	Α .	4	Α	Α	Α	+	+	+	Α	Α	+	+	Α	Α	+	+	+	+	+	+	+	+	+	+	
Intestine small, ileum	+	. ,	Α .	4	Α	Α	Α	+	+	+	+	Α	+	+	Α	Α	Α	+	+	+	+	+	+	+	+	+	
Liver	+		+ -	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Mesentery														+					+					+			
Oral mucosa																											
Pharyngeal, squamous cell papilloma																											
Pancreas	+		+ -	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Salivary glands	+		+ -	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Stomach, forestomach	+		+ -	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Stomach, glandular	+		+ -	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Cardiovascular System																											
Blood vessel																											
	+		+ -	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Heart	+		+ -	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Endocrine System																											
Adrenal cortex	+		+ -	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Adrenal medulla	+		+ -	+	+	+	+	+	+	+	+	+	+	+	+	M	+	+	+	+	+	+	+	M	+]	+	
Pheochromocytoma malignant							X												X								
Pheochromocytoma benign											X									X		X					
Bilateral, pheochromocytoma benign										X																	
Islets, pancreatic	+		+ -	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Adenoma																											
Parathyroid gland	+		+ -	+	+	M	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Pituitary gland	+		+ -												M								· [+	+	+	+	
Pars distalis, adenoma	X							X	•			-	X						X		•				X		
Thyroid gland	+		+ -	+	+	+	+		+	+	+	+			+	+	+	+			+	+			+		
C-cell, adenoma						•					•	•	•	•			•		•							•	
Follicular cell, carcinoma																											
General Body System																											
None																											
Genital System																											
Epididymis	+		+ -	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Preputial gland			· ·	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	, +	· +	+	4	4	+	4	+	
Adenoma	+			_	Τ'	т.	7-	7	7	-	7	-	7	7	7-	7	7	7		т	_	Т	_	_	_	т	
Bilateral, adenoma						X				X																	
Prostate			_	_	_		_	_	_	Λ +	_	ر	_	_	_	_	_	_	,							_	
Prostate Seminal vesicle	+			_	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Testes	+			_	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
	+		+ -	+	+	+ V	+ V	+	+ V	+	+ V	+	+ v	+ v	+ v	+ v	+ v	+ v	+ V								
Bilateral, interstitial cell, adenoma					X	X	Λ		Λ	٨	Λ	٨	Λ	Λ	X	Λ	X	X		X	X	X	X	X	X	Λ	
Interstitial cell, adenoma					х																						

	5	5	6	6	6	6	6	6	6	6	6	6	6	6	6	6	6	6	6	6	7	7	7	7	7	
Number of Days on Study	9	9				4	4	4	5	5	6	6	6	6	7	7	8	8	9	9	0	0	1	3	3	
Number of Days on Study	6	7				1	3	5	1	2	1	3	5	8	2	9	0	1	6	6	2	9	9	4		
	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	Total
Carcass ID Number	3	4	1	3	3	4	0	4	2	2	2	3	3	0	3	3	2	1	4	4	0	3	0	0	1	Tissues/
	3	1	. 8	4	9	0	5	2	0				5		1		3		6	8	1		7	3	0	Tumors
Alimentary System																										
Esophagus	+	4	+ +	- +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Intestine large, colon	+	Н	+ +	- +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	48
Intestine large, rectum	+	Н	- 4	- +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	Α	+	+	48
Intestine large, cecum	+	Н	- 4	- 1	A A	+	+	+	+	+	+	+	+	Α	+	+	+	+	+	+	+	+	+	+	+	39
Intestine small, duodenum	+	Н	- 4	- 4	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Intestine small, jejunum	+	4	+ +	- 1	٠ +	Α	+	+	+	+	Α	Α	+	Α	+	+	+	Α	+	+	+	+	Α	+	+	34
Intestine small, ileum	+	4	+ +	- +	- A	+	+	+	+	+	+	+	+	Α	+	+	+	Α	+	+	+	+	Α	+	+	37
Liver	+	4		- +	- +	+	+	+	+		+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	49
Mesentery																										3
Oral mucosa	+																									1
Pharyngeal, squamous cell papilloma	X																									1
Pancreas	+	4		- +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Salivary glands	+	4	+ +	- +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Stomach, forestomach	+	4		- +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Stomach, glandular	+	+		- +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Cardiovascular System																										
Blood vessel	+	4	+ +	- 4	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Heart	+	+	- 4	- +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Endocrine System																										
Adrenal cortex	+	4	+ +	- 4	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Adrenal medulla	+	4		- 4	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	48
Pheochromocytoma malignant																										2
Pheochromocytoma benign			}	(X											X	X				X	X	9
Bilateral, pheochromocytoma benign																X							X			3
Islets, pancreatic	+	4		- 4	- +	+	+	+	+	+	+	+	+	+	+		+	+	+	+	+	+	+		+	50
Adenoma									X								X								X	4
Parathyroid gland	M	[+		- 4	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	Μ	+	M	46
Pituitary gland	+	4		- 4	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	45
Pars distalis, adenoma		>	ζ.	Σ	ζ.	X				X			X				X					X		X		18
Thyroid gland	+						+	+	+	+	+		+	+	+	+		+	+	+	+		+	+	+	50
C-cell, adenoma													X	•		·						X				2
Follicular cell, carcinoma											X															1
General Body System																										
None																										
Genital System																										
Epididymis	+	4		- +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Preputial gland	+	Н		- +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Adenoma																X							X			2
Bilateral, adenoma																										2
Prostate	+	4		- +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Seminal vesicle	+	Н		- +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Testes	+	4	+ +	- +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Bilateral, interstitial cell, adenoma	X	>	()	()	X	X	X	X			X	X		X	X	X	X	X	X	X	X	X	X	X	X	40
Interstitial cell, adenoma									v	X																4

TABLE A2 Individual Animal Tumor Patholo	gy of Male Rats in the 2-Year Inhalation Study of Ethylbenzene: 750 ppm (continued)
Number of Days on Study	3 3 3 4 5 5 5 5 5 5 5 5 5 5 5 5 5 5 5 5
Carcass ID Number	3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3
Hematopoietic System Bone marrow Lymph node Lymph node, bronchial Lymph node, mandibular Lymph node, mesenteric Lymph node, mediastinal Spleen Thymus	+ + + + + + + + + + + + + + + + + + +
Integumentary System Mammary gland Skin Basal cell carcinoma Keratoacanthoma Subcutaneous tissue, lipoma	+ + + + + + + + + + + + + + + + + + + +
Musculoskeletal System Bone Skeletal muscle Sarcoma	+ + + + + + + + + + + + + + + + + + + +
Nervous System Brain Glioma malignant	+ + + + + + + + + + + + + + + + + + +
Respiratory System Larynx Lung Alveolar/bronchiolar adenoma Nose	M + + + M + M + + + + + M M M + + M + + M M + M M M + + + + + + + + + + + + + + + + + + + +
Trachea Special Senses System Zymbal's gland	++++++++++++++++++++
Urinary System Kidney Renal tubule, adenoma Renal tubule, carcinoma Urinary bladder	+ + + + + + + + + + + + + + + + + + +
Systemic Lesions Multiple organs Leukemia mononuclear	+ + + + + + + + + + + + + + + + + + +

Number of Days on Study	5 9	9						6 4	6 5	6 5	6	6	6 6	6 6	6 7	6 7	6 8	6 8	6 9	6 9	7 0	7 0	7	7	7 3	
Number of Days on Study	6							5	1	2	1	3	5	8	2	9	0	1	6	6	2	9	9	4	4	
	3	3	3 3	; ;	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	Total
Carcass ID Number	3		1 1		3 1 9			4 2	2 0		2 8	3 2	3 5	0 2		3 0		1 2	4 6		0 1	3 8	0 7	0 3	1 0	Tissues/ Tumors
Hematopoietic System																										
Bone marrow	+	-	+ +		+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Lymph node	+		+	٠,		+			+						+				+					+		14
Lymph node, bronchial	IV.	1 -	+ +	- I	M +	- 1\	1 +	M	IVI	M	M	M	+	+	M	+	M	M	+	+	+	+	+	M	+	28 50
Lymph node, mandibular Lymph node, mesenteric	+		+ + 		+ +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50 50
Lymph node, mediastinal					+ +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	47
Spleen	+		+ +		+ +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Г̂hymus	+	-	+ +		+ +	- +	+	+	+	+	+	+	M	+	M	+	+	+	+	+	+	+	+	+	+	44
Integumentary System																										
Mammary gland	+	-	+ +	- 1	M +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	49
Skin	+	-	+ +		+ +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Basal cell carcinoma															37								X		37	1
Keratoacanthoma Subcutaneous tissue, lipoma															X									X	X	2 1
•																								Λ		1
Musculoskeletal System																										50
Bone Shalatal musala	+	-	+ +		+ +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Skeletal muscle Sarcoma																						+ X				1 1
Nervous System																										
Brain	+		⊢ ⊣		+ +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Glioma malignant							·	·	·		·	·	·	·					·		•		·	·		1
Respiratory System																										
Larynx	N	1 -	+ +		+ +	- +	+	+	+	M	+	+	+	+	+	+	+	+	+	+	+	+	+	+	M	35
Lung	+	-	+ +		+ +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Alveolar/bronchiolar adenoma																X										1
Nose Frachea	+		+ + + +		+ +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50 50
Special Senses System Zymbal's gland																						+				1
U rinary System Kidney			_	_	L .1			_	_		_	_	_	_	_	_	_	_	_	_	_	_		_	_	50
Renal tubule, adenoma	+	_	+ + }		r +	- +	+	+	+	+	+	+ X	+	+	_	т	_	т	+	+	_	+	+	+	+ X	4
Renal tubule, carcinoma		2		-					X			. 1													. 1	3
Urinary bladder	+				+ +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	49
Systemic Lesions																										
Multiple organs	+	-	+ +		+ +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Leukemia mononuclear															X										X	9

TABLE A3
Statistical Analysis of Primary Neoplasms in Male Rats in the 2-Year Inhalation Study of Ethylbenzene

	Chamber Control	75 ppm	250 ppm	750 ppm
Adrenal Medulla: Benign Pheochromocytoma				
Overall rate ^a	13/50 (26%)	13/50 (26%)	9/49 (18%)	12/48 (25%)
Adjusted rate ^b	48.8%	62.0%	42.6%	100.0%
Terminal rate ^c	4/15 (27%)	7/14 (50%)	4/13 (31%)	2/2 (100%)
First incidence (days)	584	584	590	546
Life table test ^d	P = 0.003	P = 0.545	P = 0.279N	P = 0.014
Logistic regression test ^d	P = 0.211	P = 0.552	P = 0.233N	P = 0.307
Cochran-Armitage test ⁰	P = 0.516N			
Fisher exact test ^d		P = 0.590N	P = 0.251N	P = 0.547N
Adrenal Medulla: Benign or Malignant Pheochron	ocytoma			
Overall rate	13/50 (26%)	13/50 (26%)	9/49 (18%)	14/48 (29%)
Adjusted rate	48.8%	62.0%	42.6%	100.0%
Terminal rate	4/15 (27%)	7/14 (50%)	4/13 (31%)	2/2 (100%)
First incidence (days)	584	584	590	507
Life table test	P< 0.001	P = 0.545	P = 0.279N	P = 0.005
Logistic regression test	P = 0.106	P = 0.552	P = 0.233N	P = 0.214
Cochran-Armitage test	P = 0.379			
Fisher exact test		P = 0.590N	P = 0.251N	P = 0.450
Kidney (Renal Tubule): Adenoma (Single Sections)				
Overall rate	0/50 (0%)	3/50 (6%)	2/50 (4%)	4/50 (8%)
Adjusted rate	0.0%	11.2%	11.0%	56.9%
Terminal rate	0/15 (0%)	0/14 (0%)	1/13 (8%)	1/2 (50%)
First incidence (days)	e	617	671	587
Life table test	P = 0.006	P = 0.120	P = 0.236	P = 0.008
Logistic regression test	P = 0.064	P = 0.119	P = 0.240	P = 0.037
Cochran-Armitage test	P = 0.109	D 0.404	D 0.047	D 0.050
Fisher exact test		P = 0.121	P = 0.247	P = 0.059
Kidney (Renal Tubule): Adenoma (Step Sections)				
Overall rate	3/50 (6%)	2/50 (4%)	7/50 (14%)	17/50 (34%)
Adjusted rate	13.4%	14.3%	39.7%	88.8%
Terminal rate	0/15 (0%)	2/14 (14%)	4/13 (31%)	1/2 (50%)
First incidence (days)	685	734 (T)	671	572
Life table test	P< 0.001	P= 0.519N	P = 0.144	P< 0.001
Logistic regression test	P< 0.001	P = 0.516N	P = 0.159	P< 0.001
Cochran-Armitage test Fisher exact test	P< 0.001	P = 0.500N	P= 0.159	P< 0.001
Tighti Cauci test		1 - 0.0001	1 – 0.100	1 < 0.001
Kidney (Renal Tubule): Adenoma (Single and Step	•	E/EO (100/)	7/50 (140/)	90/50 (400/)
Overall rate	3/50 (6%)	5/50 (10%)	7/50 (14%)	20/50 (40%)
Adjusted rate	13.4%	23.9%	39.7%	100.0%
Terminal rate First incidence (days)	0/15 (0%) 685	2/14 (14%) 617	4/13 (31%) 671	2/2 (100%) 572
Life table test	080 P< 0.001	P = 0.343	P = 0.144	P< 0.001
Logistic regression test	P< 0.001 P< 0.001	P = 0.343 P = 0.337	P = 0.144 P = 0.159	P< 0.001 P< 0.001
Cochran-Armitage test	P< 0.001	1 - 0.337	1 - 0.133	1 < 0.001
Fisher exact test	1 < 0.001	P = 0.357	P = 0.159	P< 0.001
and different tool		1 0.001	1 0.100	1 0.001

TABLE A3
Statistical Analysis of Primary Neoplasms in Male Rats in the 2-Year Inhalation Study of Ethylbenzene (continued)

	Chamber Contr	ol 75 ppm	250 ppm	750 ppm
Kidney (Renal Tubule): C	arcinoma (Single Sections)			
Overall rate	0/50 (0%)	0/50 (0%)	1/50 (2%)	3/50 (6%)
Adjusted rate	0.0%	0.0%	7.7%	12.5%
Terminal rate	0/15 (0%)	0/14 (0%)	1/13 (8%)	0/2 (0%)
First incidence (days)	_		734 (T)	587
Life table test	P = 0.002	_f	P = 0.471	P = 0.063
Logistic regression test	P = 0.018	_	P = 0.471	P = 0.129
Cochran-Armitage test	P = 0.021			
Fisher exact test		_	P = 0.500	P = 0.121
Kidney (Renal Tubule): C	` • •			
Overall rate	0/50 (0%)	0/50 (0%)	1/50 (2%)	3/50 (6%)
Adjusted rate	0.0%	0.0%	7.7%	12.5%
Cerminal rate	0/15 (0%)	0/14 (0%)	1/13 (8%)	0/2 (0%)
First incidence (days)	— B. 0.000	_	734 (T)	587
Life table test	P = 0.002	_	P = 0.471	P= 0.063
Logistic regression test	P = 0.018	_	P = 0.471	P = 0.129
Cochran-Armitage test Fisher exact test	P = 0.021		P = 0.500	P= 0.121
isher exact test		_	r = 0.300	r=0.121
	arcinoma (Single and Step Sections)	0 (70 (00))	. (50 (00 ()	0.470 (004)
Overall rate	0/50 (0%)	0/50 (0%)	1/50 (2%)	3/50 (6%)
Adjusted rate	0.0%	0.0%	7.7%	12.5%
Terminal rate First incidence (days)	0/15 (0%)	0/14 (0%)	1/13 (8%) 734 (T)	0/2 (0%) 587
Life table test	P= 0.002	_	P= 0.471	P= 0.063
ogistic regression test	P = 0.002		P = 0.471	P = 0.129
Cochran-Armitage test	P = 0.021		1 - 0.171	1 - 0.120
Fisher exact test	1 01021	_	P = 0.500	P= 0.121
(idnev (Renal Tuhule) · A	denoma or Carcinoma (Single Sections)			
Overall rate	0/50 (0%)	3/50 (6%)	3/50 (6%)	7/50 (14%)
Adjusted rate	0.0%	11.2%	18.4%	62.4%
Terminal rate	0/15 (0%)	0/14 (0%)	2/13 (15%)	1/2 (50%)
First incidence (days)	-	617	671	587
Life table test	P< 0.001	P = 0.120	P = 0.111	P< 0.001
ogistic regression test	P = 0.003	P = 0.119	P = 0.121	P = 0.006
Cochran-Armitage test	P = 0.007			
isher exact test		P = 0.121	P = 0.121	P = 0.006
Kidney (Renal Tubule): A	denoma or Carcinoma (Step Sections)			
Overall rate	3/50 (6%)	2/50 (4%)	8/50 (16%)	18/50 (36%)
Adjusted rate	13.4%	14.3%	46.4%	89.1%
Terminal rate	0/15 (0%)	2/14 (14%)	5/13 (38%)	1/2 (50%)
First incidence (days)	685	734 (T)	671	572
ife table test	P< 0.001	P = 0.519N	P = 0.087	P< 0.001
ogistic regression test	P< 0.001	P = 0.516N	P = 0.098	P< 0.001
Cochran-Armitage test	P< 0.001			
Fisher exact test		P = 0.500N	P = 0.100	P< 0.001

TABLE A3
Statistical Analysis of Primary Neoplasms in Male Rats in the 2-Year Inhalation Study of Ethylbenzene (continued)

	Chamber Control	75 ppm	250 ррт	750 ppm
Kidney (Renal Tubule): Adenoma or Carcinon	na (Single and Sten Section	s)		
Overall rate	3/50 (6%)	5/50 (10%)	8/50 (16%)	21/50 (42%)
Adjusted rate	13.4%	23.9%	46.4%	100.0%
Ferminal rate	0/15 (0%)	2/14 (14%)	5/13 (38%)	2/2 (100%)
First incidence (days)	685	617	671	572
Life table test	P< 0.001	P = 0.343	P = 0.087	P< 0.001
Logistic regression test	P< 0.001	P = 0.337	P = 0.098	P< 0.001
Cochran-Armitage test	P< 0.001			
Fisher exact test		P = 0.357	P = 0.100	P< 0.001
Liver: Hepatocellular Adenoma				
Overall rate	0/50 (0%)	3/50 (6%)	0/50 (0%)	0/49 (0%)
Adjusted rate	0.0%	9.7%	0.0%	0.0%
Ferminal rate	0/15 (0%)	0/14 (0%)	0/13 (0%)	0/2 (0%)
First incidence (days)	—	560	—	— —
Life table test	P = 0.326N	P = 0.112	_	_
Logistic regression test	P = 0.246N	P = 0.125	_	_
Cochran-Armitage test	P = 0.259N			
Fisher exact test		P = 0.121	_	_
Lung: Alveolar/bronchiolar Adenoma or Carc	inoma			
Overall rate	3/50 (6%)	1/50 (2%)	0/50 (0%)	1/50 (2%)
Adjusted rate	18.4%	5.3%	0.0%	10.0%
Ferminal rate	2/15 (13%)	0/14 (0%)	0/13 (0%)	0/2 (0%)
First incidence (days)	714	713	_	679
Life table test	P= 0.643	P = 0.309N	P = 0.146N	P= 0.593
Logistic regression test	P = 0.635N	P = 0.310N	P = 0.119N	P = 0.740N
Cochran-Armitage test	P = 0.339N			
Fisher exact test		P = 0.309N	P = 0.121N	P = 0.309N
Mammary Gland: Fibroadenoma				
Overall rate	2/50 (4%)	3/50 (6%)	2/50 (4%)	0/50 (0%)
Adjusted rate	13.3%	21.4%	13.8%	0.0%
Terminal rate	2/15 (13%)	3/14 (21%)	1/13 (8%)	0/2 (0%)
First incidence (days)	734 (T)	734 (T)	720	_ ` ´
Life table test	P = 0.547N	P = 0.467	P = 0.657	P = 0.726N
Logistic regression test	P = 0.503N	P = 0.467	P = 0.681	P = 0.726N
Cochran-Armitage test	P = 0.116N			
isher exact test		P = 0.500	P = 0.691N	P = 0.247N
Mammary Gland: Fibroma, Fibroadenoma, o	r Adenoma			
Overall rate	2/50 (4%)	5/50 (10%)	3/50 (6%)	0/50 (0%)
Adjusted rate	13.3%	27.1%	16.5%	0.0%
Ferminal rate	2/15 (13%)	3/14 (21%)	1/13 (8%)	0/2 (0%)
First incidence (days)	734 (T)	549	643	_
Life table test	P = 0.382N	P = 0.193	P = 0.470	P = 0.726N
Logistic regression test	P = 0.180N	P = 0.199	P = 0.507	P = 0.726N
Cochran-Armitage test	P = 0.073N			
Fisher exact test		P = 0.218	P = 0.500	P = 0.247N

TABLE A3
Statistical Analysis of Primary Neoplasms in Male Rats in the 2-Year Inhalation Study of Ethylbenzene (continued)

	Chamber Control	75 ppm	250 ppm	750 ppm
Oral Cavity (Oral Mucosa and Tongue): Squ	ramous Cell Panilloma			
Overall rate	3/50 (6%)	0/50 (0%)	1/50 (2%)	1/50 (2%)
Adjusted rate	12.6%	0.0%	7.7%	4.0%
Ferminal rate	1/15 (7%)	0/14 (0%)	1/13 (8%)	0/2 (0%)
First incidence (days)	570	- -	734 (T)	596
Life table test	P= 0.618	P = 0.132N	P = 0.335N	P= 0.639N
ogistic regression test	P = 0.505N	P = 0.132N	P = 0.303N	P = 0.348N
Cochran-Armitage test	P = 0.442N	1 - 0.12411	1 - 0.00011	1 – 0.04014
Fisher exact test	1 0.11211	P = 0.121N	P = 0.309N	P = 0.309N
ancreatic Islets: Adenoma				
Overall rate	3/50 (6%)	4/50 (8%)	4/50 (8%)	4/50 (8%)
Adjusted rate	16.0%	17.5%	22.5%	60.5%
Cerminal rate	1/15 (7%)	1/14 (7%)	2/13 (15%)	1/2 (50%)
First incidence (days)	692	560	671	645
ife table test	P = 0.033	P = 0.492	P = 0.477	P = 0.043
Logistic regression test	P = 0.220	P = 0.483	P = 0.509	P = 0.163
Cochran-Armitage test	P = 0.493			
Fisher exact test		P = 0.500	P = 0.500	P = 0.500
Pancreatic Islets: Adenoma or Carcinoma				
Overall rate	5/50 (10%)	5/50 (10%)	4/50 (8%)	4/50 (8%)
Adjusted rate	23.3%	23.9%	22.5%	60.5%
Cerminal rate	1/15 (7%)	2/14 (14%)	2/13 (15%)	1/2 (50%)
First incidence (days)	590	560	671	645
ife table test	P = 0.116	P = 0.617	P = 0.523N	P = 0.152
ogistic regression test	P = 0.464	P = 0.612	P = 0.492N	P = 0.462
Cochran-Armitage test	P = 0.429N			
isher exact test		P = 0.630N	P = 0.500N	P = 0.500N
Pituitary Gland (Pars Distalis): Adenoma				
Overall rate	25/49 (51%)	19/50 (38%)	19/50 (38%)	18/45 (40%)
Adjusted rate	79.7%	66.7%	65.8%	82.7%
Cerminal rate	9/14 (64%)	7/14 (50%)	6/13 (46%)	1/2 (50%)
First incidence (days)	391	518	420	377
Life table test	P = 0.034	P = 0.222N	P = 0.237N	P = 0.068
ogistic regression test	P = 0.355N	P = 0.147N	P = 0.135N	P = 0.234N
Cochran-Armitage test	P = 0.314N			
isher exact test		P = 0.135N	P = 0.135N	P = 0.194N
reputial Gland: Adenoma				
Overall rate	3/49 (6%)	1/50 (2%)	1/49 (2%)	4/50 (8%)
Adjusted rate	12.5%	2.3%	7.7%	42.7%
Terminal rate	1/14 (7%)	0/14 (0%)	1/13 (8%)	0/2 (0%)
'irst incidence (days)	574	560	734 (T)	500
Life table test	P = 0.048	P = 0.318N	P = 0.314N	P = 0.172
ogistic regression test	P = 0.228	P = 0.291N	P = 0.302N	P = 0.502
Cochran-Armitage test	P = 0.227			
Fisher exact test		P = 0.301N	P = 0.309N	P = 0.511

TABLE A3
Statistical Analysis of Primary Neoplasms in Male Rats in the 2-Year Inhalation Study of Ethylbenzene (continued)

	Chamber Control	75 ppm	250 ppm	750 ppm
Skin: Keratoacanthoma				
Overall rate	3/50 (6%)	2/50 (4%)	2/50 (4%)	2/50 (4%)
Adjusted rate	11.2%	9.1%	15.4%	54.5%
Terminal rate	1/15 (7%)	0/14 (0%)	2/13 (15%)	1/2 (50%)
First incidence (days)	528	637	734 (T)	672
Life table test	P= 0.306	P = 0.516N	P = 0.543N	P = 0.462
ogistic regression test	P = 0.608	P = 0.501N	P = 0.500N	P = 0.606N
Cochran-Armitage test	P = 0.491N	1 0.00111	1 0100011	1 0100011
isher exact test	1 0/1011	P = 0.500N	P = 0.500N	P = 0.500N
skin: Squamous Cell Papilloma or Keratoac	anthoma			
overall rate	5/50 (10%)	3/50 (6%)	3/50 (6%)	2/50 (4%)
Adjusted rate	20.1%	15.6%	23.1%	54.5%
Cerminal rate	2/15 (13%)	1/14 (7%)	3/13 (23%)	1/2 (50%)
First incidence (days)	528	637	734 (T)	672
ife table test	P = 0.483	P = 0.393N	P = 0.407N	P= 0.646
ogistic regression test	P = 0.483 P = 0.422N	P = 0.368N	P = 0.407N P = 0.353N	P = 0.040 P = 0.342N
Cochran-Armitage test	P = 0.422N P = 0.225N	1 - 0.30011	1 - 0.33311	1 - 0.0421N
isher exact test	1 — 0.22014	P = 0.357N	P = 0.357N	P = 0.218N
skin: Squamous Cell Papilloma, Keratoacan	thoma Rasal Call Adanoma	or Rasal Call C	arcinoma	
overall rate	5/50 (10%)	3/50 (6%)	4/50 (8%)	3/50 (6%)
Adjusted	20.1%	15.6%	26.2%	69.7%
erminal	2/15 (13%)	1/14 (7%)	3/13 (23%)	1/2 (50%)
irst incidence (days)	528	637	688	672
ife table	P = 0.183	P = 0.393N	P= 0.548N	P = 0.358
ogistic regression	P = 0.535	P = 0.368N	P = 0.497N	P = 0.575N
cochran-Armitage	P = 0.389N	1 – 0.3001	1 – 0.43711	1 – 0.37311
isher exact	1 - 0.0001	P = 0.357N	P = 0.500N	P= 0.357N
skin (Subcutaneous Tissue): Fibroma				
Overall rate	1/50 (2%)	1/50 (2%)	3/50 (6%)	0/50 (0%)
Adjusted rate	6.7%	5.3%	18.8%	0.0%
Cerminal rate	1/15 (7%)	0/14 (0%)	2/13 (15%)	0.0%
irst incidence (days)	734 (T)	713	688	U/2 (U/0) —
ifst incluence (days) ife table test	P = 0.624	P = 0.759	P = 0.269	P= 0.882N
ogistic regression test	P = 0.624 P = 0.690N	P = 0.759 P = 0.759	P = 0.209 P = 0.302	P = 0.882N
Cochran-Armitage test	P = 0.030N P = 0.339N	1 – 0.733	1 – 0.302	1 - U.0021V
isher exact test	1 – 0.33311	P = 0.753N	P = 0.309	P = 0.500N
W. (G.)				
Skin (Subcutaneous Tissue): Fibrosarcoma o		0.480 (00.4)	0.450 (00.1)	0 (#0 (00))
Overall rate	0/50 (0%)	3/50 (6%)	0/50 (0%)	0/50 (0%)
Adjusted rate	0.0%	10.9%	0.0%	0.0%
Germinal rate	0/15 (0%)	0/14 (0%)	0/13 (0%)	0/2 (0%)
irst incidence (days)	— D. 0.00017	560	_	_
ife table test	P = 0.380N	P = 0.122	_	_
ogistic regression test	P = 0.259N	P = 0.120	_	_
	_			
Cochran-Armitage test Fisher exact test	P = 0.255N	P= 0.121		

TABLE A3
Statistical Analysis of Primary Neoplasms in Male Rats in the 2-Year Inhalation Study of Ethylbenzene (continued)

	Chamber Control	75 ppm	250 ppm	750 ppm
Skin (Subcutaneous Tissue): Fibroma, Myxoma	ı. Fibrosarcoma, or Sarco	ma		
Overall rate	1/50 (2%)	4/50 (8%)	4/50 (8%)	0/50 (0%)
Adjusted rate	6.7%	15.6%	20.4%	0.0%
Terminal rate	1/15 (7%)	0/14 (0%)	2/13 (15%)	0/2 (0%)
First incidence (days)	734 (T)	560	266	_
Life table test	P = 0.487N	P = 0.182	P = 0.157	P = 0.882N
Logistic regression test	P = 0.323N	P = 0.171	P = 0.181	P= 0.882N
Cochran-Armitage test	P = 0.155N			
Fisher exact test		P = 0.181	P = 0.181	P = 0.500N
Testes: Bilateral Adenoma				
Overall rate	27/50 (54%)	23/50 (46%)	32/50 (64%)	40/50 (80%)
Adjusted rate	96.0%	91.0%	96.5%	100.0%
Terminal rate	14/15 (93%)	12/14 (86%)	12/13 (92%)	2/2 (100%)
First incidence (days)	608	538	590	500
Life table test	P< 0.001	P = 0.364N	P = 0.185	P< 0.001
Logistic regression test	P< 0.001	P = 0.313N	P = 0.177	P< 0.001
Cochran-Armitage test	P< 0.001			
Fisher exact test		P = 0.274N	P = 0.208	P = 0.005
Testes: Adenoma				
Overall rate	36/50 (72%)	33/50 (66%)	40/50 (80%)	44/50 (88%)
Adjusted rate	100.0%	100.0%	100.0%	100.0%
Terminal rate	15/15 (100%)	14/14 (100%)	13/13 (100%)	2/2 (100%)
First incidence (days)	497	538	420	483
Life table test	P< 0.001	P = 0.480N	P = 0.259	P< 0.001
Logistic regression test	P< 0.001	P = 0.404N	P = 0.194	P = 0.001
Cochran-Armitage test	P = 0.010			
Fisher exact test		P = 0.333N	P = 0.241	P = 0.039
Thyroid Gland (C-cell): Adenoma				
Overall rate	3/50 (6%)	6/49 (12%)	3/50 (6%)	2/50 (4%)
Adjusted rate	20.0%	32.7%	13.0%	30.8%
Terminal rate	3/15 (20%)	3/14 (21%)	1/13 (8%)	0/2 (0%)
First incidence (days)	734 (T)	617	591	665
Life table test	P = 0.366	P = 0.223	P = 0.633	P = 0.208
Logistic regression test	P = 0.539N	P = 0.217	P = 0.659N	P = 0.390
Cochran-Armitage test	P = 0.217N			
Fisher exact test		P = 0.233	P = 0.661N	P = 0.500N
Thyroid Gland (C-cell): Adenoma or Carcinom	a			
Overall rate	5/50 (10%)	6/49 (12%)	3/50 (6%)	2/50 (4%)
Adjusted rate	27.5%	32.7%	13.0%	30.8%
Terminal rate	3/15 (20%)	3/14 (21%)	1/13 (8%)	0/2 (0%)
First incidence (days)	661	617	591	665
Life table test	P = 0.553	P = 0.481	P = 0.384N	P = 0.444
Logistic regression test	P = 0.342N	P = 0.474	P = 0.349N	P = 0.676N
Cochran-Armitage test	P = 0.112N			
Fisher exact test		P = 0.486	P = 0.357N	P = 0.218N

TABLE A3
Statistical Analysis of Primary Neoplasms in Male Rats in the 2-Year Inhalation Study of Ethylbenzene (continued)

	Chamber Control	75 ppm	250 ppm	750 ppm
All Organs: Mononuclear Cell Leukemia				
Overall rate	27/50 (54%)	26/50 (52%)	32/50 (64%)	9/50 (18%)
Adjusted rate	74.7%	79.3%	83.1%	62.2%
Terminal rate	7/15 (47%)	8/14 (57%)	7/13 (54%)	1/2 (50%)
First incidence (days)	570	582	496	383
ife table test	P = 0.412N	P = 0.543	P = 0.264	P = 0.287N
ogistic regression test	P< 0.001N	P = 0.555N	P = 0.166	P< 0.001N
Cochran-Armitage test	P< 0.001N			
isher exact test		P = 0.500N	P = 0.208	P< 0.001N
All Organs: Benign Neoplasms				
Overall rate	48/50 (96%)	44/50 (88%)	48/50 (96%)	48/50 (96%)
Adjusted rate	100.0%	100.0%	100.0%	100.0%
Terminal rate	15/15 (100%)	14/14 (100%)	13/13 (100%)	2/2 (100%)
First incidence (days)	391	518	266	377
ife table test	P< 0.001	P = 0.452N	P = 0.475	P< 0.001
ogistic regression test	P = 0.184	P = 0.205N	P = 0.667	P = 0.586
Cochran-Armitage test	P = 0.296			
isher exact test		P = 0.134N	P = 0.691N	P = 0.691N
All Organs: Malignant Neoplasms				
Overall rate	33/50 (66%)	32/50 (64%)	37/50 (74%)	17/50 (34%)
Adjusted rate	83.6%	85.7%	89.6%	85.3%
'erminal rate	9/15 (60%)	9/14 (64%)	9/13 (69%)	1/2 (50%)
'irst incidence (days)	497	560	420	383
ife table test	P = 0.225	P = 0.532	P = 0.310	P = 0.368
ogistic regression test	P = 0.001N	P = 0.551N	P = 0.247	P = 0.003N
Cochran-Armitage test	P< 0.001N			
isher exact test		P = 0.500N	P = 0.257	P = 0.001N
All Organs: Benign or Malignant Neoplasms				
Overall rate	49/50 (98%)	45/50 (90%)	50/50 (100%)	50/50 (100%)
adjusted rate	100.0%	100.0%	100.0%	100.0%
'erminal rate	15/15 (100%)	14/14 (100%)	13/13 (100%)	2/2 (100%)
irst incidence (days)	391	518	266	377
ife table test	P< 0.001	P = 0.454N	P = 0.430	P< 0.001
ogistic regression test	P = 0.059	P = 0.151N	P = 0.349	_
Cochran-Armitage test	P = 0.081			
isher exact test		P = 0.102N	P = 0.500	P = 0.500

(T)Terminal sacrifice

^a Number of neoplasm-bearing animals/number of animals examined. Denominator is number of animals examined microscopically for adrenal gland, kidney, liver, lung, pancreatic islets, pituitary gland, preputial gland, testes, and thyroid gland; for other tissues, denominator is number of animals necropsied.

b Kaplan-Meier estimated neoplasm incidence at the end of the study after adjustment for intercurrent mortality

Observed incidence at terminal kill

Beneath the chamber control incidence are the P values associated with the trend test. Beneath the exposure group incidence are the P values corresponding to pairwise comparisons between the chamber controls and that exposure group. The life table test regards neoplasms in animals dying prior to terminal kill as being (directly or indirectly) the cause of death. The logistic regression test regards these lesions as nonfatal. The Cochran-Armitage and Fisher exact tests compare directly the overall incidence rates. For all tests, a negative trend or a lower incidence in an exposure group is indicated by **N**.

Not applicable; no neoplasms in animal group

Value of statistic cannot be computed.

TABLE A4a Historical Incidence of Renal Tubule Neoplasms in Chamber Control Male F344/N Rats^a

		Incidence in Controls							
Study	Adenoma	Carcinoma	Adenoma or Carcinoma						
Historical Incidence at IIT Research	ch Institute								
Isobutyl Nitrite	0/45	0/45	0/45						
Overall Historical Incidence									
Total Standard deviation Range	6/652 (0.9%) 1.3% 0%-4%	0/652 (0%)	6/652 (0.9%) 1.3% 0%-4%						

^a Data as of 12 May 1995

TABLE A4b Historical Incidence of Testicular Adenoma in Chamber Control Male F344/N Rats^a

Study	Incidence in Controls
Historical Incidence at IIT Research Institute Isobutyl Nitrite	31/46
Overall Historical Incidence	
Total Standard deviation Range	450/655 (68.7%) 8.7% 54%-83%

^a Data as of 12 May 1995

TABLE A5
Summary of the Incidence of Nonneoplastic Lesions in Male Rats in the 2-Year Inhalation Study of Ethylbenzene^a

	Chambe	r Control	75	5 ppm	250) ppm	75	0 ppm
Disposition Summary								
Animals initially in study	5	50		50		50		50
Early deaths								
Moribund	2	28		20		26		26
Natural deaths		7		16		11		22
urvivors				4.4		10		
Terminal sacrifice]	15		14		13		2
Animals examined microscopically	5	50		50		50		50
Alimentary System								
ntestine large, colon	(50)		(48)		(48)		(48)	
Hemorrhage			1					
Inflammation		4		(2%)				
Mineralization		(2%)		(2%)	(10)			(2%)
ntestine large, rectum	(48)		(49)		(48)	(00/)	(48)	
Thrombosis	(40)		(4.4)			(2%)	(00)	
ntestine large, cecum	(46)	(00/)	(44)	(50/)	(46)		(39)	
Inflammation	1	(2%)	2	(5%)				(3%)
Mineralization			1	(90/)			1	(3%)
Necrosis Ulcer	1	(2%)	1	(2%)				
ntestine small, duodenum	(48)	(2/0)	(48)		(50)		(50)	
Mineralization		(2%)	(40)		(30)		(30)	
Necrosis	1	(270)			1	(2%)		
ntestine small, jejunum	(42)		(39)		(44)	(270)	(34)	
Inflammation	(12)		(00)		(11)			(3%)
ntestine small, ileum	(45)		(44)		(45)		(37)	
Inflammation	()		()		()			(3%)
iver	(50)		(50)		(50)		(49)	
Angiectasis	` ′		` ,			(4%)		(2%)
Basophilic focus	6	(12%)	5	(10%)	2	(4%)		(8%)
Clear cell focus	2	(4%)	3	(6%)	3	(6%)		
Cyst			1	(2%)				
Degeneration		(2%)						
Degeneration, cystic		(30%)		(24%)		(38%)		(61%)
Eosinophilic focus	5	(10%)		(22%)	4	(8%)	9	(18%)
Fibrosis				(6%)				
Hemorrhage				(4%)				(00.1)
Hepatodiaphragmatic nodule	-	(00/)		(2%)	-	(00/)	1	(2%)
Inflammation, chronic	1	(2%)		(2%)	1	(2%)		
Inflammation, chronic active			1	(2%)				(00/)
Mineralization	4	(90/)		(40/)			1	(2%)
Mixed cell focus		(2%)		(4%)			O	(169/)
Necrosis Pigmentation		(4%) (2%)	4	(8%)			8	(16%)
Pigmentation Thrombosis	1	(£ /0)			1	(2%)		
Vacuolization cytoplasmic	Q	(16%)	10	(20%)		(14%)	1	(8%)
Bile duct, hyperplasia	0	(10/0)	10	(20/0)	,	(17/0)		(2%)
Bile duct, inflammation, suppurative			1	(2%)			1	(~ /U)
Kupffer cell, hyperplasia			1	(~ /0)	1	(2%)		
raprici cen, nyperpiasia					1	(~ /0)		

^a Number of animals examined microscopically at the site and the number of animals with lesion

TABLE A5
Summary of the Incidence of Nonneoplastic Lesions in Male Rats in the 2-Year Inhalation Study of Ethylbenzene (continued)

	Chamber Control	75 ppm	250 ppm	750 ppm
Alimentary System (continued)				
Mesentery	(4)	(5)	(3)	(3)
Inflammation	(-)	1 (20%)	(5)	(-)
Fat, necrosis	3 (75%)	4 (80%)	2 (67%)	3 (100%)
Pancreas	(50)	(49)	(50)	(50)
Inflammation	(00)	2 (4%)	1 (2%)	1 (2%)
Acinus, atrophy	24 (48%)	21 (43%)	20 (40%)	18 (36%)
Acinus, hyperplasia	4 (8%)	21 (4070)	1 (2%)	10 (3070)
Artery, degeneration	1 (0/0)	1 (2%)	1 (2%)	
Artery, inflammation		1 (2%)	1 (270)	
Artery, mineralization		1 (2%)		
Stomach, forestomach	(50)	(50)	(50)	(50)
Hyperkeratosis	1 (2%)	(30)	1 (2%)	(30)
Hyperplasia	8 (16%)	5 (10%)	8 (16%)	8 (16%)
Inflammation	1 (2%)	1 (2%)	3 (6%)	3 (6%)
Mineralization	2 (4%)	1 (2%)	1 (2%)	5 (10%)
Ulcer	9 (18%)	9 (18%)	9 (18%)	10 (20%)
Stomach, glandular	(50)	(49)	(50)	(50)
	(30)	(40)		(30)
Degeneration			1 (2%)	1 (2%)
Degeneration, cystic Inflammation	2 (4%)	1 (2%)	3 (6%)	1 (2%) 1 (2%)
Inflammation Mineralization	2 (4%) 4 (8%)	1 (2%) 4 (8%)	3 (6%)	1 (2%) 18 (36%)
				18 (36%)
Necrosis Ulcer	5 (10%)	2 (4%) 2 (4%)	2 (4%)	2 (4%)
Oicei		2 (4/0)		2 (470)
Condinues only Contant				
Cardiovascular System	(50)	(50)	(50)	(50)
Blood vessel	(50)	(50)	(50)	(50)
Mineralization			4 (00/)	1 (2%)
Aorta, inflammation	0 (40()	0 (10/)	1 (2%)	1 (2%)
Aorta, mineralization	2 (4%)	2 (4%)	4 (8%)	14 (28%)
Heart	(50)	(50)	(50)	(50)
Cardiomyopathy	26 (52%)	21 (42%)	15 (30%)	30 (60%)
Inflammation			1 (2%)	
Mineralization	2 (4%)	1 (2%)		7 (14%)
Atrium, thrombosis	5 (10%)	7 (14%)	7 (14%)	
Valve, fibrosis	1 (2%)			
Endonino Sustan				
Endocrine System	(50)	(50)	(50)	(50)
Adrenal cortex	(50)	(50)	(50)	(50)
Cytoplasmic alteration		1 (2%)	1 (00/)	
Degeneration	1 (00/)	0 (40/)	1 (2%)	0 (00/)
Degeneration, cystic	1 (2%)	2 (4%)	1 (2%)	3 (6%)
Hyperplasia	1 (2%)		1 (2%)	2 (4%)
Hypertrophy			4 (00.0)	1 (2%)
Necrosis			1 (2%)	2 (4%)
Pigmentation	1 (2%)	40 (5)	40 (
Vacuolization cytoplasmic	13 (26%)	18 (36%)	16 (32%)	11 (22%)
Bilateral, atrophy	1 (2%)			
Capsule, inflammation				1 (2%)
Adrenal medulla	(50)	(50)	(49)	(48)
Hyperplasia	10 (20%)	7 (14%)	13 (27%)	8 (17%)
Necrosis				1 (2%)
Bilateral, hyperplasia	2 (4%)	2 (4%)	1 (2%)	4 (8%)

Inflammation

Inflammation

Mineralization

Atrophy Degeneration

Hemorrhage Mineralization

Arteriole, inflammation Bilateral, atrophy

Bilateral, necrosis Interstitial cell, hyperplasia

Seminal vesicle

Testes

Infiltration cellular, lymphocyte

TABLE A5 Summary of the Incidence of Nonneenlastic Legions in Male Pats in the 2-Vear Inhalation Study of Ethylhenzene

	Chambe	r Control	75	5 ppm	250	ppm	750) ppm
Endocrine System (continued)								
Islets, pancreatic	(50)		(50)		(50)		(50)	
Hyperplasia	2	(4%)	5	(10%)	4	(8%)		
Parathyroid gland	(45)		(46)		(46)		(46)	
Fibrosis					1	(2%)		
Hyperplasia	12	(27%)	6	(13%)	16	(35%)	35	(76%)
Pituitary gland	(49)		(50)		(50)		(45)	
Pars distalis, angiectasis	5	(10%)	11	(22%)	5	(10%)	4	(9%)
Pars distalis, cyst	1	(2%)	6	(12%)	5	(10%)	4	(9%)
Pars distalis, degeneration					1	(2%)		
Pars distalis, hemorrhage				(2%)		(2%)		
Pars distalis, hyperplasia	12	(24%)	11	(22%)		(24%)	12	(27%)
Pars distalis, necrosis						(2%)		
Pars distalis, pigmentation		(2%)			2	(4%)		
Pars intermedia, angiectasis		(2%)						
Thyroid gland	(50)		(49)		(50)		(50)	
Cyst				(2%)				
C-cell, hyperplasia		(12%)		(10%)	5	(10%)		
Follicle, cyst	1	(2%)	1	(2%)			2	(4%)
General Body System None								
Genital System Epididymis	(50)		(50)		(50)		(50)	
Granuloma sperm	(30)			(2%)		(2%)	(30)	
Inflammation				(2%)		(2%)	1	(2%)
Mineralization			1	(~ /U)	1	(2/0)		(2%)
Preputial gland	(49)		(50)		(49)		(50)	(2/0)
Atrophy	(49)		(30)			(2%)	(30)	
Hyperplasia	9	(4%)	9	(4%)	1	(2/0)	1	(2%)
Inflammation		(39%)		(14%)	Q	(16%)		(20%)
Prostate	(50)	(0070)	(50)	(11/0)	(50)	(10/0)	(50)	(2070)
Hyperplasia		(2%)		(2%)		(2%)	(30)	
Infiltration collular lymphocyte	1	(2/0)		(20/)	1	(~ / U)		

1 (2%)

29 (58%)

1 (2%)

(50) 7 (14%)

1 (2%)

1 (2%) 7 (14%)

19 (38%)

(49)

22 (44%)

1 (2%)

(50) 10 (20%)

1 (2%) 5 (10%)

1 (2%) 12 (24%)

(50)

25 (50%)

6 (12%)

1 (2%)

2 (4%)

4 (8%)

8 (16%)

(50)

(50)

11 (22%)

1 (2%)

10 (20%)

1 (2%)

9 (18%)

14 (28%)

(49)

(50)

TABLE A5
Summary of the Incidence of Nonneoplastic Lesions in Male Rats in the 2-Year Inhalation Study of Ethylbenzene (continued)

	Chamber Contro	ol 7:	5 ppm	250) ppm	750) ppm
Hematopoietic System							
Bone marrow	(49)	(49)		(50)		(50)	
Atrophy	()	(/			(2%)		(2%)
Hemorrhage	1 (2%)				(4%)		(8%)
Hyperplasia	7 (14%)	16	(33%)		(18%)		(38%)
Inflammation	. ()		(2%)		()		(00.0)
Myelofibrosis	3 (6%)	_	()			5	(10%)
Myeloid cell, atrophy	1 (2%)					_	(==, =)
Lymph node	(9)	(8)		(9)		(14)	
Hemorrhage	(-7	(-)			(11%)	()	
Lumbar, hemorrhage		1	(13%)	_	()		
Lumbar, hyperplasia, plasma cell			(13%)				
Pancreatic, fibrosis		-	(1070)	1	(11%)		
Pancreatic, pigmentation					(11%)		
Renal, ectasia					(22%)	1	(7%)
Renal, hemorrhage					(11%)		(57%)
Renal, hyperplasia, lymphoid				•	(11/0)		(7%)
Renal, hyperplasia, plasma cell				1	(11%)		(7%)
Renal, infiltration cellular, histocyte		1	(13%)	1	(1170)	1	(170)
Renal, pigmentation		•	(1070)	1	(11%)	2	(14%)
Lymph node, bronchial	(44)	(34)		(39)	(1170)	(28)	(1170)
Ectasia	(11)	(01)			(3%)	(20)	
Hemorrhage	7 (16%)	4	(12%)		(10%)	7	(25%)
Infiltration cellular, histiocyte	7 (1070)	-	(12/0)		(5%)	,	(2370)
Pigmentation	3 (7%)	3	(9%)	٤	(370)	5	(18%)
Lymph node, mandibular	(47)	(48)	(370)	(49)		(50)	(1070)
Atrophy	(41)		(2%)	(40)		(30)	
Hemorrhage	1 (2%)	1	(270)			1	(2%)
Hyperplasia, plasma cell	4 (9%)	1	(2%)	1	(2%)		(8%)
Inflammation	1 (070)	1	(270)		(2%)	1	(070)
Pigmentation	1 (2%)				(270)		
Lymph node, mesenteric	(49)	(50)		(50)		(50)	
Atrophy	(43)		(2%)	(30)		(30)	
Ectasia		1	(2/0)	1	(2%)		
Hemorrhage	3 (6%)	5	(10%)	4	1 1	Q	(16%)
Inflammation	3 (070)	J	(1070)		(2%)		(2%)
Lymph node, mediastinal	(48)	(48)		(50)	(2/0)	(47)	(2/0)
Edema	1 (2%)		(2%)	(30)			(6%)
Hemorrhage	12 (25%)			10	(20%)		, ,
	12 (25%)		(21%) (2%)	10	(20%)	17	(36%)
Hyperplasia, plasma cell Infiltration cellular, histiocyte	2 (4%)		(2%)			1	(2%)
	2 (4%)	1	(2%)	1	(90/)	1	(2%)
Inflammation	0 (100/)	0	(170/)		(2%)	7	(150/)
Pigmentation	9 (19%)		(17%)		(14%)		(15%)
Spleen	(50)	(49)		(50)		(50)	(40/)
Atrophy	1 (00/)	1	(00/)			Z	(4%)
Congestion	1 (2%)		(2%)	0	(00/)		(00/)
Depletion cellular	0 (00/)		(4%)		(6%)		(8%)
Fibrosis	3 (6%)		(2%)		(8%)		(2%)
Hematopoietic cell proliferation	3 (6%)	4	(8%)		(2%)	2	(4%)
Inflammation, chronic	2 (12)			1	(2%)	=	(40/)
Necrosis	2 (4%)		(00.1)			2	(4%)
Pigmentation		1	(2%)	-	(00/)		
Red pulp, depletion cellular				1	(2%)		

TABLE A5
Summary of the Incidence of Nonneoplastic Lesions in Male Rats in the 2-Year Inhalation Study of Ethylbenzene (continued)

	Chamber Control	75 ppm	250 ppm	750 ppm
Hematopoietic System (continued)				
Thymus	(46)	(44)	(46)	(44)
Čyst		1 (00/)		1 (2%)
Hemorrhage		1 (2%)		1 (2%)
Integumentary System				
Mammary gland	(46)	(47)	(46)	(49)
Fibrosis			1 (2%)	- 4
Galactocele	11 (24%)	11 (23%)	11 (24%)	9 (18%)
Hyperplasia	3 (7%)	3 (6%)	4 (9%)	3 (6%)
Inflammation	1 (2%)	2 (4%)	1 (2%)	1 (90/)
Mineralization	1 (90/)	4 (9%)	2 (4%)	1 (2%)
Pigmentation	1 (2%)	` '	` ,	6 (12%)
Skin Cyst epithelial inclusion	(50) 1 (2%)	(50) 4 (8%)	(50) 2 (4%)	(50)
Hyperkeratosis	1 (2%)	4 (6/0)	2 (4/0)	1 (2%)
Inflammation	1 (270)			1 (2%)
Inflammation, granulomatous		1 (2%)		1 (270)
Epidermis, hyperplasia		1 (2/0)	1 (2%)	
Subcutaneous tissue, inflammation		2 (4%)	- ()	
Musculoskeletal System Bone Fibrous osteodystrophy Hyperostosis Turbinate, hyperostosis	(49) 1 (2%) 1 (2%)	(50) 1 (2%) 1 (2%)	(50) 5 (10%) 2 (4%)	(50) 9 (18%) 1 (2%)
Nervous System				
Brain	(50)	(50)	(50)	(50)
Hemorrhage		1 (2%)		
Hydrocephalus	1 (2%)	2 (4%)	1 (2%)	
Mineralization	0 (40/)	1 (00/)	1 (2%)	1 (00/)
Necrosis	2 (4%)	1 (2%)	3 (6%)	1 (2%)
Respiratory System				
Larynx	(40)	(44)	(41)	(35)
Foreign body			1 (2%)	
Infiltration cellular, lymphocyte	1 (3%)	1 (2%)	1 (2%)	1 (3%)
Inflammation	1 (3%)	3 (7%)	3 (7%)	1 (3%)
Necrosis		1 (2%)		
Respiratory epithelium, hyperplasia	1 (3%)	4 (9%)	1 (2%)	1 (3%)
Respiratory epithelium, metaplasia, squamo	us 1 (3%)	1 (2%)	1 (2%)	2 (6%)

TABLE A5
Summary of the Incidence of Nonneoplastic Lesions in Male Rats in the 2-Year Inhalation Study of Ethylbenzene (continued)

(Chamber Control	75 ppm	250 ppm	750 ppm
Respiratory System (continued)				
Lung	(50)	(50)	(50)	(50)
Congestion	1 (2%)	2 (4%)		6 (12%)
Edema	1 (2%)			6 (12%)
Fibrosis			1 (2%)	1 (2%)
Foreign body			1 (2%)	
Hemorrhage		2 (4%)	1 (2%)	8 (16%)
Infiltration cellular, histiocyte	2 (4%)	1 (2%)		
Inflammation, acute		1 (2%)	1 (2%)	1 (2%)
Inflammation, chronic	1 (2%)		1 (2%)	2 (4%)
Inflammation, chronic active	2 (4%)	3 (6%)	2 (4%)	7 (14%)
Inflammation, granulomatous	1 (2%)	1 (2%)		
Mineralization	1 (2%)	1 (2%)		3 (6%)
Alveolar epithelium, hyperplasia	2 (4%)	2 (4%)	1 (2%)	2 (4%)
Artery, mineralization				1 (2%)
Goblet cell, hyperplasia			1 (2%)	
Interstitium, fibrosis		1 (2%)		4 (00.1)
Interstitium, inflammation	(40)	(40)	(50)	1 (2%)
Nose	(49)	(49)	(50)	(50)
Angiectasis			1 (2%)	4 (00()
Congestion	0 (40()	0 (40()	0 (40()	1 (2%)
Foreign body	2 (4%)	2 (4%)	2 (4%)	1 (2%)
Infiltration cellular, lymphocyte	0 (100/)	1 (2%)	0 (100/)	0 (100/)
Inflammation	8 (16%)	8 (16%)	9 (18%)	9 (18%)
Necrosis		1 (00/)		1 (2%)
Glands, cyst	0 (40/)	1 (2%)	1 (90/)	
Goblet cell, hyperplasia	2 (4%)	1 (90/)	1 (2%)	
Nasolacrimal duct, inflammation	1 (2%)	1 (2%)	1 (2%)	1 (90/)
Olfactory epithelium, inflammation Olfactory epithelium, metaplasia				1 (2%)
Respiratory epithelium, hyperplasia	0 (100/)	7 (140/)	0 (190/)	1 (2%)
Respiratory epithelium, nyperplasia Respiratory epithelium, inflammation	9 (18%)	7 (14%)	9 (18%)	6 (12%)
Respiratory epithelium, metaplasia, squamou	1 (2%) us 1 (2%)			3 (6%) 1 (2%)
Respiratory epithelium, ulcer	1 (2%) 1 (2%)		1 (2%)	1 (470)
Frachea	(50)	(50)	(50)	(50)
Mineralization	(30)	(30)	(30)	1 (2%)
Willeralization				1 (2/0)
Special Senses System	(4)	(4)		
Eye	(1)	(1)		
Lens, cataract	1 (100%)			
Retina, degeneration	1 (100%)		(4)	745
Zymbal's gland	(1)		(1)	(1)
Cyst				1 (100%)
Hyperplasia				1 (100%)

TABLE A5
Summary of the Incidence of Nonneoplastic Lesions in Male Rats in the 2-Year Inhalation Study of Ethylbenzene (continued)

	Chamber Control		75	5 ppm	250 ррт		750 ppm	
Urinary System								
Kidney	(50)		(50)		(50)		(50)	
Cyst			4	(8%)	1	(2%)	10	(20%)
Hemorrhage			1	(2%)				
Infarct	2	(4%)					1	(2%)
Inflammation			1	(2%)				
Mineralization	1	(2%)	1	(2%)	1	(2%)	9	(18%)
Necrosis	1	(2%)					1	(2%)
Nephropathy	47	(94%)	43	(86%)	47	(94%)	48	(96%)
Pigmentation	9	(18%)	6	(12%)	9	(18%)	2	(4%)
Renal tubule, hyperplasia	2	(4%)	2	(4%)	4	(8%)	12	(24%)
Transitional epithelium, hyperplasia	12	(24%)	14	(28%)	15	(30%)	34	(68%)
Urinary bladder	(49)		(49)		(50)		(49)	
Hemorrhage	1	(2%)	2	(4%)	, ,		1	(2%)
Inflammation	1	(2%)	3	(6%)	1	(2%)		
Necrosis				(2%)		, ,		
Transitional epithelium, hyperplasia				(4%)	1	(2%)		

APPENDIX B SUMMARY OF LESIONS IN FEMALE RATS IN THE 2-YEAR INHALATION STUDY OF ETHYLBENZENE

TABLE B1	Summary of the Incidence of Neoplasms in Female Rats	
	in the 2-Year Inhalation Study of Ethylbenzene	100
TABLE B2	Individual Animal Tumor Pathology of Female Rats	
	in the 2-Year Inhalation Study of Ethylbenzene	104
TABLE B3	Statistical Analysis of Primary Neoplasms in Female Rats	
	in the 2-Year Inhalation Study of Ethylbenzene	120
TABLE B4	Summary of the Incidence of Nonneoplastic Lesions in Female Rats	
	in the 2-Year Inhalation Study of Ethylbenzene	125

TABLE B1 Summary of the Incidence of Neoplasms in Female Rats in the 2-Year Inhalation Study of Ethylbenzene^a

	Chamber Control	75 ppm	250 ppm	750 ppm
Disposition Summary				
Animals initially in study Early deaths	50	50	50	50
Moribund Natural deaths	7 12	14 5	8 8	6 8
Survivors				
Died last week of study	1	0.4		0.7
Terminal sacrifice Missing	30	31	34	35 1
Animals examined microscopically	50	50	50	49
Alimentary System				
Intestine large, colon	(47)	(49)	(48)	(49)
Intestine large, rectum	(49)	(50)	(47)	(49)
Polyp adenomatous			1 (2%)	
Intestine large, cecum	(44)	(50)	(47)	(49)
Intestine small, duodenum	(47)	(48)	(47)	(48)
Intestine small, jejunum	(41)	(49)	(47)	(45)
ntestine small, ileum	(41)	(48)	(47)	(46)
Liver	(50)	(50)	(50)	(49)
Histiocytic sarcoma	1 (2%)	(4)	(0)	(7)
Mesentery	(7)	(4)	(6)	(7)
Oral mucosa				(1)
Pharyngeal, squamous cell papilloma	(40)	(70)	(50)	1 (100%)
Pancreas	(49)	(50)	(50)	(49)
Histiocytic sarcoma	1 (2%)	(50)	(50)	(40)
Salivary glands	(50)	(50)	(50)	(49)
Stomach, forestomach	(49)	(50)	(50)	(49)
Stomach, glandular	(49)	(49)	(49)	(49)
Гоngue Schwannoma malignant			(1)	(1) 1 (100%)
Squamous cell papilloma			1 (100%)	1 (100%)
Squamous cen papmonia			1 (100%)	
Cardiovascular System	(70)	(50)	(50)	(40)
Heart	(50)	(50)	(50)	(49)
Endocrine System				
Adrenal cortex	(50)	(50)	(50)	(49)
Adenoma		1 (2%)	1 (2%)	
Carcinoma	(50)	(50)	1 (2%)	(40)
Adrenal medulla	(50)	(50)	(50)	(49)
Pheochromocytoma malignant	0 (10)	1 (2%)	2 (4%)	
Pheochromocytoma benign	2 (4%)	0 (40()		
Bilateral, pheochromocytoma benign	(50)	2 (4%)	(50)	(40)
Islets, pancreatic	(50)	(50)	(50)	(49)
Adenoma	1 (2%)		1 (2%)	1 (2%)

TABLE B1
Summary of the Incidence of Neoplasms in Female Rats in the 2-Year Inhalation Study of Ethylbenzene (continued)

	Chamber Control	75 ppm	250 ppm	750 ppm
Endocrine System (continued)				
Pituitary gland	(49)	(49)	(50)	(49)
Pars distalis, adenoma	27 (55%)	17 (35%)	24 (48%)	24 (49%)
Pars distalis, adenoma, multiple	3 (6%)	6 (12%)	1 (2%)	3 (6%)
Pars distalis, carcinoma		1 (2%)	1 (2%)	
Гhyroid gland	(48)	(50)	(50)	(49)
Bilateral, C-cell, adenoma		1 (2%)		
C-cell, adenoma	2 (4%)	3 (6%)	2 (4%)	3 (6%)
C-cell, carcinoma				1 (2%)
General Body System None				
Genital System				
Genital System Clitoral gland	(47)	(49)	(48)	(47)
Adenoma	2 (4%)	(40)	(40)	(41)
Carcinoma	1 (2%)		1 (2%)	
Ovary	(50)	(50)	(50)	(49)
Histiocytic sarcoma	(30)	(30)	(30)	1 (2%)
Uterus	(50)	(50)	(50)	(49)
Polyp stromal	2 (4%)	3 (6%)	4 (8%)	3 (6%)
Bilateral, polyp stromal	~ (1/0)	J (370)	1 (2%)	J (0/0)
Endometrium, sarcoma stromal		1 (2%)	- (2/0)	
Hematopoietic System	(40)	(50)	(50)	(40)
Bone marrow	(49)	(50)	(50)	(49)
Lymph node	(3)	(3)	(4)	(4)
Lumbar, histiocytic sarcoma	1 (33%)	(0.4)	(41)	(00)
Lymph node, bronchial	(37)	(34)	(41)	(38)
Lymph node, mandibular	(49)	(50)	(50)	(49)
Lymph node, mesenteric	(49)	(50)	(50)	(49)
Lymph node, mediastinal	(49)	(49)	(50)	(49)
Rhabdomyosarcoma, metastatic,			1 (90/)	
uncertain primary site	(40)	(50)	1 (2%)	(40)
Spleen	(49)	(50) (47)	(49)	(49)
Thymus Phobdomyosarcoma motostotic	(48)	(47)	(47)	(47)
Rhabdomyosarcoma, metastatic,			1 (90/)	
uncertain primary site			1 (2%)	
Integumentary System				
Mammary gland	(48)	(50)	(49)	(49)
Adenoma	1 (2%)	2 (4%)		1 (2%)
Carcinoma	2 (4%)	1 (2%)	2 (4%)	
Carcinoma, multiple	1 (2%)			1 (2%)
Fibroadenoma	13 (27%)	18 (36%)	18 (37%)	15 (31%)
Fibroadenoma, multiple	6 (13%)	1 (2%)	3 (6%)	6 (12%)

TABLE B1
Summary of the Incidence of Neoplasms in Female Rats in the 2-Year Inhalation Study of Ethylbenzene (continued)

	Chamber Control	75 ppm	250 ppm	750 ppm
Integumentary System (continued) Skin	(50)	(50)	(50)	(49)
Squamous cell carcinoma	(30)	(30)	1 (2%)	(49)
Squamous cell papilloma Sebaceous gland, carcinoma			2 (4%)	1 (2%)
Subcutaneous tissue, fibroma Subcutaneous tissue, fibrosarcoma	1 (2%)	1 (2%)		
Subcutaneous tissue, lipoma		1 (2%)		
Subcutaneous tissue, sarcoma		1 (2%)		
Musculoskeletal System Skeletal muscle		(1)		
okeietai iliusete		(1)		
Nervous System	(50)	(50)	(5.0)	(40)
Brain Astrocytoma malignant	(50) 1 (2%)	(50)	(50)	(49)
Carcinoma, metastatic, pituitary gland	- ()	1 (2%)	1 (2%)	
Respiratory System				
Larynx	(45)	(43)	(44)	(45)
Lung Alveolar/bronchiolar adenoma	(50) 1 (2%)	(50)	(50) 1 (2%)	(49)
Alveolar/bronchiolar adenoma, multiple	- ()	1 (2%)		
Carcinoma, metastatic, mammary gland Histiocytic sarcoma	1 (2%)		1 (2%)	
Sarcoma, metastatic, uncertain primary sit		1 (2%)		
Mediastinum, sarcoma, metastatic, uncertain primary site		1 (2%)		
Nose	(50)	(50)	(50)	(49)
Glands, adenoma	(50)	1 (2%)	(70)	(40)
Гrachea	(50)	(50)	(50)	(49)
Special Senses System				
Ear Entermal con concerns	(3)			
External ear, sarcoma Zymbal's gland	1 (33%) (1)	(1)		
Adenoma	1 (100%)			
Carcinoma		1 (100%)		
Urinary System				
Kidney Panal tuhula adanoma	(50)	(50)	(50)	(49)
Renal tubule, adenoma Jrinary bladder	(48)	(49)	(49)	1 (2%) (48)

TABLE B1 Summary of the Incidence of Neoplasms in Female Rats in the 2-Year Inhalation Study of Ethylbenzene (continued)

	Chamber Control	75 ppm	250 ppm	750 ppm
Systemic Lesions				
Multiple organs ^b	(50)	(50)	(50)	(49)
Histiocytic sarcoma	2 (4%)	, ,	• •	1 (2%)
Leukemia granulocytic		1 (2%)		
Leukemia mononuclear	13 (26%)	18 (36%)	16 (32%)	11 (22%)
Lymphoma malignant	1 (2%)	1 (2%)		
Neoplasm Summary Total animals with primary neoplasms Total primary neoplasms Total animals with benign neoplasms Total benign neoplasms Total animals with malignant neoplasms Total malignant neoplasms	42 84 37 62 20 22	45 84 39 57 24 27	43 84 37 60 21 24	46 74 39 58 14 16
Total animals with metastatic neoplasms		2	3	
Total metastatic neoplasms Total animals with malignant neoplasms		3	4	
			1	

Number of animals examined microscopically at the site and the number of animals with neoplasm Number of animals with any tissue examined microscopically Primary neoplasms: all neoplasms except metastatic neoplasms

TABLE B2 Individual Animal Tumor Pathology of Female Rats in the 2-Year Inhalation Study of Ethylbenzene: Chamber Control

Number of Days on Study	0 1 1 3 4 4 5 6 6 6 6 6 6 6 6 6 6 7 7 7 7 7 7 7 7 7
Carcass ID Number	$ \begin{array}{cccccccccccccccccccccccccccccccccccc$
Alimentary System	
Esophagus	+ + + + + + + + + + + + + + + + + + + +
Intestine large, colon	A A A + + + + + + + + + + + + + + + + +
Intestine large, rectum	+ + A + + + + + + + + + + + + + + + + +
Intestine large, cecum	A A A + + + + + + + + A A + A + + + + +
Intestine small, duodenum	A A A + + + + + + + + + + + + + + + + +
Intestine small, jejunum	A A A A A + + + + + + + A + + A + + A + + + + + + + + + + + + + + + + + + + +
Intestine small, ileum	A A A A + + + + + + A A A + A + A + + + + + + + + + + + + + + + + + + + +
Liver	+ + + + + + + + + + + + + + + + + + + +
Histiocytic sarcoma	X
Mesentery	+ + + + + +
Pancreas	+ A + + + + + + + + + + + + + + + + + +
Histiocytic sarcoma	X
Salivary glands	+ + + + + + + + + + + + + + + + + + + +
Stomach, forestomach	+ A + + + + + + + + + + + + + + + + + +
Stomach, glandular	+ A + + + + + + + + + + + + + + + + + +
Tooth	+
Cardiovascular System	
Blood vessel	+ + + + + M + + + + + + + + + + + + + +
Heart	+ + + + + + + + + + + + + + + + + + + +
Endocrine System	
Adrenal cortex	
Adrenal medulla	+ + + + + + + + + + + + + + + + + + + +
	X
Pheochromocytoma benign Islets, pancreatic	
Adenoma	+ + + + + + + + + + + + + + + + + + +
Parathyroid gland	
Pituitary gland	
Pars distalis, adenoma	
Pars distalis, adenoma, multiple	X X
Thyroid gland C-cell, adenoma	+ A M + + + + + + + + + + + + + + + + +
C cen, auchoma	ΛΛ
General Body System None	
Genital System	
	+ + + + + + + + + + + + + + M + + + + M + + + +
Clitoral gland	
Clitoral gland Adenoma	
Adenoma Carcinoma	+ + + + + + + + + + + + + + + + + + + +
Adenoma	+ + + + + + + + + + + + + + + + + + + +

+: Tissue examined microscopically A: Autolysis precludes examination

M: Missing tissue I: Insufficient tissue

X: Lesion present Blank: Not examined

TABLE B2

Individual Animal Tumor Patholog (continued)	gy of Female Rats in the 2-Year Inhalation Study of Ethylbenzene: Chamber												er Contro														
	7		7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	
Number of Days on Study	3			3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	
in the contract of	5		5	5	5	5	5	5	5	5	5	5		5	6	6	6		6	6	6	6	6	6	6	6	
	0	()	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	Total
Carcass ID Number	5		5	5	6	6	6	7	7	7	7	8	9	9	5	6	6	7	7	7	8	8	8	9	9	9	Tissues/
	4		5	9	1	3	4	3	4	7	9	5	2	5	1	6	8	2	5	8	0	2	8	4	8	9	Tumors
Alimentary System																											
Esophagus	+		+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Intestine large, colon	+		+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	47
Intestine large, rectum	+		+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	49
Intestine large, cecum	+		+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	44
Intestine small, duodenum	+		+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	47
Intestine small, jejunum	+		+	+	+	+	M	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	41
Intestine small, ileum	+		+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	41
Liver	+		+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Histiocytic sarcoma																											1
Mesentery																									+		7
Pancreas	+		+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	49
Histiocytic sarcoma																											1
Salivary glands	+		+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Stomach, forestomach	+		+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	49
Stomach, glandular	+		+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	49
Γooth																											1
Cardiovascular System																											
Blood vessel	+		+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	49
Heart	+		+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Endocrine System																											
Adrenal cortex	+		+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Adrenal medulla	+		+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Pheochromocytoma benign									X								•			·							2
Islets, pancreatic	+		+	+	+	+	+	+			+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Adenoma							·	·		Ċ	Ċ		·	Ċ			Ċ	·		·		•		·			1
Parathyroid gland	+		+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	48
Pituitary gland			+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	49
Pars distalis, adenoma		,	X					X	Ċ		X		•		X		X									X	27
Pars distalis, adenoma, multiple				21	71			71			21	X		71	71		21		21	71	21	71	21	71	21	71	3
Thyroid gland	_		_	_	_	_	_	_	_	_	_	Λ _	_	_	_	_	_	_	_	_	_	_	_	_	_		48
C-cell, adenoma			i.	٢	т		т	~	т	7'	т.	т	т	т		-	Т	-	-	Т		т	т.	7	7	т	2
General Body System None																											
Genital System																											
Clitoral gland			_	_	_	_	_	_				M			5	_	_	ر	_	ر	5	_					47
Adenoma	+		+ X	т	_	+	+	+ X	+	+	+	1V1	+	+	+	+	+	+	+	+	+	+	+	+	+	+	2
Carcinoma		•	^			X		Λ																			
						^										,		,		,							1
Ovary Uterus	+		+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
	+	,	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50 2
Polyp stromal	X																										Z.

TABLE B2 Individual Animal Tumor Patholo (continued)	gy of Female Rats in the 2-Year Inhalation Study of Ethylbenzene: Chamber Control
Number of Days on Study	0 1 1 3 4 4 5 6 6 6 6 6 6 6 6 6 6 7 7 7 7 7 7 7 7 7
Carcass ID Number	0 0 0 1 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0
Hematopoietic System Bone marrow Lymph node Lumbar, histiocytic sarcoma	+ + A + + + + + + + + + + + + + + + + +
Lymph node, bronchial Lymph node, mandibular Lymph node, mesenteric Lymph node, mediastinal	+ + M + + + + + + + + + + + + + + + + +
Spleen Thymus	+ A + + + + + + + + + + + + + + + + + +
Integumentary System Mammary gland Adenoma Carcinoma Carcinoma, multiple	+ + M + + + + + + + + M + + + + + + + +
Fibroadenoma Fibroadenoma, multiple Skin Subcutaneous tissue, fibroma	X X X X X X X X X X X X X X X X X X X
Musculoskeletal System Bone	+ + + + + + + + + + + + + + + + + + + +
Nervous System Brain Astrocytoma malignant	+ + + + + + + + + + + + + + + + + + +
Respiratory System Larynx Lung Alveolar/bronchiolar adenoma	+ + + M + + M + + + + + + + + + + + + +
Histiocytic sarcoma Nose Trachea	X + + + + + + + + + + + + + + + + + + +
Special Senses System Ear External ear, sarcoma	+ X
Eye Zymbal's gland Adenoma	⁺ X
Urinary System Kidney Urinary bladder	+ + + + + + + + + + + + + + + + + + +
Systemic Lesions Multiple organs Histocytic sarcoma	+ + + + + + + + + + + + + + + + + + +
Leukemia mononuclear Lymphoma malignant	X X X X X X X X X X X X X X X X X X X

gy of Fem	alo	e R	at	s in	th	e 2	-Ye	ar	In	hal	ati	on	Stı	ıdy	o of	f E	thy	lb	enz	zen	e:	C	ha	mb	er Contro
7 3 5	3	3	3	3 3	3	3	7 3 5	7 3 5	7 3 5	7 3 5	7 3 5	7 3 5	7 3 6	7 3 6	7 3 6	7 3 6	7 3 6	3	7 3 6	7 3 6	7 3 6	7 3 6	7 3 6	7 3 6	
5	5	5 5	5 (6 6	6	7	0 7 4	0 7 7	7	8	9	9	5	6	6	7	7	7	0 8 0	0 8 2	0 8 8	0 9 4	0 9 8	0 9 9	Total Tissues/ Tumors
+			+	+ +	- +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	49 3 1
+ + +	· +	- - - -	+ -	+ N + + + +	И N - + - +	M M	1 + + + +	+ + + + +	+ + + +	+ + + + +	+ + +	M + +	M + +	+ + + +	+ + + + +	M + +	+ + + + +	+ + + + +	+ + + +	M + +	+ + + +	M + +	+ + + + +	+ + + + +	37 49 49 49
+		- 	+ -	+ +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	49 48
+	. +	+ - 2	+ ·	+ +	- +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	48 1 2
+	. 4	- -						X +	+	X +	+	+	+	+	X +	+				X	X +	X +	X +	+	1 13 6 50
+	. 4	+ -	+ -	+ +	- +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
+	. 4	-	+	+ +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50 1
+	- +	⊦ -	+ -	+ +			+	+	+++	+	+++	+	+	++	+++	+	+++	+++	M +	+++	+++	+	+	++	45 50 1
+	. 4	- -	+ -	+ +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	++	+	++	+	+	1 50 50
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+		- -	+ -	+ +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	++	+	+	+	+	50 48
+	. 4	-	+ -	+ +	- +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50 2
	7 3 5 0 5 4	7 7 3 3 5 5 5 5 5 5 5 5 5 5 5 5 5 5 5 5	7 7 7 3 3 3 5 5 5 5 5 5 5 5 5 5 5 5 5 5	7 7 7 7 3 3 3 3 5 5 5 5 5 6 0 0 0 0 0 5 5 5 5 4 5 9 9 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	7 7 7 7 7 7 7 3 3 3 3 3 3 3 5 5 5 5 5 5	7 7 7 7 7 7 7 7 3 3 3 3 3 3 3 3 5 5 5 5	7 7 7 7 7 7 7 7 7 7 7 3 3 3 3 3 3 3 3 3	7 7 7 7 7 7 7 7 7 7 7 7 7 3 3 3 3 3 3 3	7 7 7 7 7 7 7 7 7 7 7 7 7 7 3 3 3 3 3 3	7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3	7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7	7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7	7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7	7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7	7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7	7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7	7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7	7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7	7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7	7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7	7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7	7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7	7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7	7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7	3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3

ndividual Animal Tumor Pathology	or rem	aic					~ -	Ca	. 11	ша	utt	UII	ы	uy	U,			,					' Р.	եա
	4	4	4	5	5	6	6	6 6	6	6	6	6	6	7	7	7	7	7	7	7	7	7	7	7
Jumber of Days on Study	1	6	7	0	3	0	0 (0 1	. 3	4	4	6	8	1	2	2	3	3	3	3	3	3	3	3
<u> </u>	2	2	5	6	1	1	9 9	9 0	6	6	7	8	6	9	4	4	1	1	4	4	4	4	5	5
	1	1	1	1	1	1	1	1 1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1
rcass ID Number	7	5	5	7	8	6	8 9	9 7	8	5	8	7	6	7	9	9	5	9	6	6	8	9	5	6
	4	2	9	5	6	8	8 9	9 3	5	8	2	7	6	9	2	5	5	1	3	4	3	8	6	2
mentary System																								
phagus	+	+	+	+	+	+	+ -	+ +	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
estine large, colon	+	· A	+	+	+	+	+ -	+ +	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
estine large, rectum	+	+	+	+	+	+	+ -	+ +	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
estine large, cecum	+	+	+	+	+	+	+ -	+ +	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
estine small, duodenum	+	· A	+	+	+	Α	+ -	+ +	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
estine small, jejunum	+	+	+	+	+	+	+ -	+ 1	١ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
stine small, ileum	+	+	+	Α	+	+	+ -	+ <i>A</i>	4 +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
er	+	+	+	+	+	+	+ -	+ +	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
entery	+																							
creas	+	+	+	+	+	+	+ -	+ +	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
vary glands	+	+	+	+	+	+	+ -	+ +	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
nach, forestomach	+	+	+	+	+	+	+ -	+ +	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
ach, glandular	+	· A	+	+	+	+	+ -	+ +	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
liovascular System																								
od vessel	4		+	+	+	+	+ -	+ +	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Vesser	+	+	+	+	+	+	+ -	+ +	· ·	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
• 6 •																								
ocrine System						,																		
nal cortex	+	+	+	+	+	+	+ -		+ .,	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
denoma									ζ															
nal medulla	+	+	+	+	+	+	+ -	+ +	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
heochromocytoma malignant														v		v								
ilateral, pheochromocytoma benign														X		X	,							
s, pancreatic	+	+	+	+	+	+	+ .	+ +	+ +	+	+	+					+	+	+	+	+	+	+	+
thyroid gland	+	+	+	+ T	+	M	+ ·	+ +	+ +	+	+	+	+			+	+	+	+	+		+		
tary gland	+	+	+ V	1	+ v	+			+ +	+	+	+ V	+		+ v	+	+	+		+	+		+ V	+
Pars distalis, adenoma			X		X		4	X				X			X			X	X			X	X	
Pars distalis, adenoma, multiple														X				Λ						
ars distalis, carcinoma oid gland				,	.1	ر	_		ι,		+	+	+		+			+	,				,	. 1
roid giand Bilateral, C-cell, adenoma	+	+	+	+	+	+	+ .	- +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
-cell, adenoma																		X						
eral Body System																								
enital System																								
																	,							
oral gland ry	+	+	+	+	+	+	+ .	+ +	- + 	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
ry us	+	+	+	+	+	+	+ .	+ +	- + 	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
is olyp stromal	+	+	+	+	+	+ X	+ .	+ +	- +	+	+ X	+	+	+	+	+ X	+	+	+	+	+	+	+	+
nyp stromai idometrium, sarcoma stromal		Х				Λ					Λ					Λ								
		Λ			,																			
na e e e e e e e e e e e e e e e e e e e					+	+										+								
atopoietic System																								
marrow	+	+	+	+	+	+	+ .	+ +	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
oh node											+		+				+							
ph node, bronchial	+	+	+	M	M	+	Μ .	+ +	+ +	+	+	+	+	+	+	M	+	M	+	+	+	M	M	M
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0 0 1 1 3 4 4 6 8 1 2 2 3 3 3 3 2 2 5 6 1 1 1 9 9 0 6 6 7 8 6 9 4 4 1 1 1 4 1 1 1 1 1 1 1 1 1 1 1 1 1</td><td>4 4 4 5 5 6 6 6 6 6 6 6 6 6 6 7 7 7 7 7 7 7 7</td><td>4 4 4 5 5 6 6 6 6 6 6 6 6 6 6 7 7 7 7 7 7 7 7</td><td>4 4 4 5 5 6 6 6 6 6 6 6 6 6 6 7 7 7 7 7 7 7 7</td><td>4 4 4 5 5 6 6 6 6 6 6 6 6 6 6 7 7 7 7 7 7 7 7</td><td>1 6 7 0 3 0 0 0 1 3 4 4 6 8 1 2 2 3 3 3 3 3 3 3 3 3 3 3 3 3 3 2 2 2 5 6 1 1 9 9 0 6 6 7 8 6 9 4 4 1 1 1 4 4 4 4 4 5 5 5 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1</td></td<></td></td<>	4 4 4 5 5 6 6 6 6 6 6 6 6 6 6 7 7 7 7 7 1 1 6 7 0 3 0 0 0 0 1 1 3 4 4 6 8 1 1 2 2 3 2 2 5 6 1 1 1 9 9 0 6 6 7 8 6 9 4 4 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	4 4 4 5 5 6 6 6 6 6 6 6 6 6 6 6 6 6 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 <td< td=""><td>4 4 4 5 5 6 6 6 6 6 6 6 6 6 7 7 7 7 7 7 7 1 6 7 7 0 3 0 0 0 1 1 3 4 4 6 8 1 2 2 3 3 3 3 2 2 5 6 1 1 1 9 9 0 6 6 7 8 6 9 4 4 1 1 1 4 1 1 1 1 1 1 1 1 1 1 1 1 1</td><td>4 4 4 5 5 6 6 6 6 6 6 6 6 6 6 7 7 7 7 7 7 7 7</td><td>4 4 4 5 5 6 6 6 6 6 6 6 6 6 6 7 7 7 7 7 7 7 7</td><td>4 4 4 5 5 6 6 6 6 6 6 6 6 6 6 7 7 7 7 7 7 7 7</td><td>4 4 4 5 5 6 6 6 6 6 6 6 6 6 6 7 7 7 7 7 7 7 7</td><td>1 6 7 0 3 0 0 0 1 3 4 4 6 8 1 2 2 3 3 3 3 3 3 3 3 3 3 3 3 3 3 2 2 2 5 6 1 1 9 9 0 6 6 7 8 6 9 4 4 1 1 1 4 4 4 4 4 5 5 5 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1</td></td<>	4 4 4 5 5 6 6 6 6 6 6 6 6 6 7 7 7 7 7 7 7 1 6 7 7 0 3 0 0 0 1 1 3 4 4 6 8 1 2 2 3 3 3 3 2 2 5 6 1 1 1 9 9 0 6 6 7 8 6 9 4 4 1 1 1 4 1 1 1 1 1 1 1 1 1 1 1 1 1	4 4 4 5 5 6 6 6 6 6 6 6 6 6 6 7 7 7 7 7 7 7 7	4 4 4 5 5 6 6 6 6 6 6 6 6 6 6 7 7 7 7 7 7 7 7	4 4 4 5 5 6 6 6 6 6 6 6 6 6 6 7 7 7 7 7 7 7 7	4 4 4 5 5 6 6 6 6 6 6 6 6 6 6 7 7 7 7 7 7 7 7	1 6 7 0 3 0 0 0 1 3 4 4 6 8 1 2 2 3 3 3 3 3 3 3 3 3 3 3 3 3 3 2 2 2 5 6 1 1 9 9 0 6 6 7 8 6 9 4 4 1 1 1 4 4 4 4 4 5 5 5 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1

Individual Animal Tumor Patholog	y of Fem	lal						_	10	аі	111	uai	au	UII	Su	uu	U	L	ııı	yıD	CII	ZCII	ю.	•	P	hш	(continued
	7	,	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	
Number of Days on Study	3	}	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	
	5	i	5	5	5	5	5	5	5	5	5	6	6	6	6	6	6	6	6	6	6	6	6	6	6	6	
	1		1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	2	Total
Carcass ID Number	6			7	7	8	8	8	9	9	9	5	5	5	5		6	6	6	7		8	8	9	9	0	Tissues/
	9)	0	1	8	0	4	7	3	6	7	1	3	4	7	0	1	5	7	2	6	1	9	0	4	0	Tumors
Hematopoietic System (continued)																											
Lymph node, mediastinal	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	49
Spleen	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Thymus	+	+	+	+	+	+	+	+	+	+	+	+	M	+	+	+	+	+	+	+	+	+	+	+	+	+	47
Integumentary System																											
Mammary gland	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Adenoma	y	<																	X								2
Carcinoma Fibroadenoma	Σ	,						X		X				X			X		X				v	X	v	v	1 18
Fibroadenoma Fibroadenoma, multiple		1						Λ		Λ				Λ			Λ						Λ	Λ	Λ	Λ	18
Skin	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Subcutaneous tissue, fibrosarcoma																											1
Subcutaneous tissue, lipoma								3.7													X						1
Subcutaneous tissue, sarcoma								X																			1
Musculoskeletal System																											
Bone	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Skeletal muscle																											1
Nervous System																											
Brain	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Carcinoma, metastatic,																											
pituitary gland																											1
Respiratory System																											
Larynx	+	+	+	+	+	M	+	+	+	+	M	+	+	+	+	+	+	M	M	+	+	+	+	M	+	+	43
Lung Alveolar/bronchiolar adenoma,	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
multiple						X																					1
Sarcoma, metastatic,						••																					
uncertain primary site																											1
Mediastinum, sarcoma, metastatic,																											-
uncertain primary site							,																,				1
Nose Glands, adenoma	+	٢	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50 1
Trachea	4	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Snacial Conses System																											
Special Senses System Zymbal's gland																											1
Carcinoma																											1
Urinary System					,											,		,								,	F0
Kidney Urinary bladder	+	⊢ ⊢	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50 49
·			•	-	- 1"	-	Г	-	-	٢	-	-	-	- 1	- 1	- 11	1"	- 1"	-	1-	-	-	٢		-	-	10
Systemic Lesions																											F0
Multiple organs	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Leukemia granulocytic Leukemia mononuclear						X		X		X			X	X					X	X					X	X	1 18
Lymphoma malignant						/1		/1		/1			1	1					2 1	41					/1	21	1

Individual Animal Tumor Patholog	ω ·															, -			,						
	3		4 4	Į :	5 6	3 (6	6 6	6	6	6	6	6	6	6	7	7	7	7	7	7	7	7	7	7
Number of Days on Study	9		4 8	3 (0 () 2	2	5 5	5	6	7	8	8	9	9	0	3	3	3	3	3	3	3	3	3
- v	3	. !	9 6	3	2 2	2 4	4	1 8	3 9	2	4	5	9	4	9	9	4	4	5	5	5	5	5	5	5
	2		2 3	3	2 2	2 2	2	2 2	2 2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2
Carcass ID Number	5		7 (6 9					5	6	7	8	6	7	5	5	5	6	6	6	6
Curcus 12 Tumber	6		2 (9 5															3		
Alimentary System																									
Esophagus	+		+ -	+ -	+ -	+ -	+	+ -	+ +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Intestine large, colon	+		· + -		+ /	Δ.		+ -		٠ +				+	+	+	+	+	+	+	+	+	+	+	+
Intestine large, rectum	-		Д -	⊢ .	+ /			· + -						+	+	+	+	+	+	+	+	+	+	+	+
Polyp adenomatous	'				' '	•			' '														X		'
Intestine large, cecum						٨		+ -	. ,	٠ +	+	+	Α	+									23		+
	-				+ 1										+	+	+	+	+	+	+	+	+	+	+
Intestine small, duodenum	+			Α.				+ -						+	+	+	+	+	+	+	+	+	+	+	+
Intestine small, jejunum	+	- 4	Α -					+ -			+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Intestine small, ileum	+	- ,	Α -		+ 1	Α.	+		+ <i>A</i>		+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Liver	+		+ -	+ -	+ -	+ -	+	+ -	+ +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Mesentery					+					+	+														
Pancreas	+		+ -	+ -	+ -	+ -	+	+ -	+ +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Salivary glands	+		+ -	+ -	+ -	+ -	+	+ -	+ +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Stomach, forestomach	_		+ -	<u>.</u>	+ -	+ -	+	+ -			. +	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Stomach, glandular			· 	L .	· 	· 	· -	· + -	- A	٠	. +			+	Ţ	·	·	·	į.	·	·	·	·	·	·
			_			_	-	_	<i>-</i>	1 1				_	_			-				_			т
Tongue Squamous cell papilloma																									
Cardiovascular System Blood vessel Heart	+		+ -	+ -	+ -	+ -	+ -	+ -	+ + + +	- +	· +	+	++	++	+ +	++	++	++	++	++	+	+	+	+	++
																				_		_			
Endocrine System																									
Adrenal cortex	+		+ -	+ -	+ -	+ -	+	+ -	+ +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Adenoma																									
Carcinoma																									
Adrenal medulla	+		+ -	+ -	+ -	+ -	+	+ -	+ +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Pheochromocytoma malignant									Σ	(X											
Islets, pancreatic	+		+ -	+ .	+ -	+ -	+	+ -			+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Adenoma										·	,				X		•						•		
Parathyroid gland			_	_	_ 1	Λſ	_	+ 1	ν Γ .,			1./	ſ⊥			М		М		_	_	_	_	+	_
Pituitary gland					. 1								+										+		
	+		v -		т ,		+		- +					+	+				+	+	+				
Pars distalis, adenoma			X		2	X				X		X				Х	X		T 7			X	X	X	Λ
Pars distalis, adenoma, multiple																			X						
Pars distalis, carcinoma																									
Thyroid gland	+		+ -	+ -	+ -	+ -	+	+ -	+ +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
C-cell, adenoma																	X								
General Body System																									
None																									
Genital System																									
Clitoral gland	4		+ -	+ -	+ -	+ -	+	+ -	⊢ →	- 4	. +	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Carcinoma									. 1	T	-			'	'				'	-		X			
																						Λ.			
Ovary	+		+ -		+ -	+ -	+	+ -	+ +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Uterus	+		+ -	+ -	+ -	+ -	+	+ -	+ +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Polyp stromal Bilateral, polyp stromal																		X							

Individual Animal Tumor Patholog	50															_											(
	7	-	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	
Number of Days on Study	3	;	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	
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	2	4	2 2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	Total
Carcass ID Number	7	,	, ;	7	7	8	8	8	9	9	9	9	5	5	5	6	6	7	7	8	8	8	9	9	9	9	Tissues/
	3		3						0	2		4			9				5	0				6	8		Tumors
Alimentary System																											
Esophagus	+		+ -	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Intestine large, colon	+			+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	48
Intestine large, rectum			L .	<u>.</u>	i	·	Ţ	Ţ	Ţ	·	·	Ţ	·	·	i	<u>.</u>	·	·	·	į.	·	·	·	·	·	·	47
Polyp adenomatous				-	_	т	_	_	_		_	_	_		_	-	_	-	-		_		_			-	1
Intestine large, cecum	+		+ -	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	47
Intestine small, duodenum	+		+ -	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	47
Intestine small, jejunum	+		+ -	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	47
Intestine small, ileum	+		+ -	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	47
Liver	+		+ -	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Mesentery														+			+	+									6
Pancreas	+		+ -	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Salivary glands			∟ .	· -	·	· +	· -	·	_		·	·	· +		·	<u>.</u>	· +		_	·	· +	·	·	· +			50
Stomach, forestomach									Ċ			- 1	Ċ	Ċ		Ċ	Ċ	Ċ	Ċ		Ċ	•	Ċ		÷	Ċ	50
	Ţ				Τ.	Τ.	Τ.	Τ.	Τ.						Τ.	Τ.	Τ.	+	Τ.								49
Stomach, glandular	+			+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Tongue													+														1
Squamous cell papilloma													X														1
Cardiovascular System																											
Blood vessel	+		+ -	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Heart	+		+ -	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Endocrine System																											
Adrenal cortex	+		. -	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Adenoma																X									•		1
Carcinoma																Λ										X	1
Adrenal medulla	+		+ -	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Pheochromocytoma malignant																											2
Íslets, pancreatic	+		+ -	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Adenoma																											1
Parathyroid gland	+		+ -	+	+	+	+	+	+	M	+	+	+	+	+	+	+	+	+	M	M	+	+	+	+	+	42
Pituitary gland	+		+ -	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Pars distalis, adenoma		-	X			X	X	X	X			X				X		X				X	X	X	X		24
Pars distalis, adenoma, multiple		•	-																				• •				1
Pars distalis, carcinoma																			X								1
																,											
Thyroid gland	+			+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+		50
C-cell, adenoma																										X	2
General Body System																											
None																											
Genital System																											
Clitoral gland	_		- -	+	+	M	+	+	+	+	+	+	+	+	+	+	+	+	M	+	+	+	+	+	+	+	48
Carcinoma	,					171			•	'		'	'	'		'		'	141	'			-	-			1
																,											
Ovary	+				+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Uterus	+		+ -	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Polyp stromal								X		X								X		X							4
Bilateral, polyp stromal																											1

TABLE B2 Individual Animal Tumor Pathology o	of Fem	al	e I	Rai	ts i	in t	the	2-	Ye	ar	In	ha	ati	on	St	ud	y o	f E	th	ylb	en	zer	ıe:	2	50	ppm (cont	inued
Number of Days on Study	3 9	4	1		5 0	6 0	6 2	6 5	6 5	6 5	6	6 7	6 8	6 8	6 9	6 9	7 0	7	7 3	7 3	7 3	7 3	7	7	7 3	7 3	
Number of Days on Study	3	(2					9			5	9			9	4	4	5	5	5	5	5	5	5	
Carcass ID Number	2 5				2 6	2	2 9	2 6	2 9	2 5	2 7	2	2	2 5	2 6	2 7	2	2 6	2 7	2 5	2 5	2 5	2 6	2 6	2	2 6	
cureus 12 Number	6						7		5									2									
Hematopoietic System																											
Bone marrow Lymph node	+		+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Lymph node, bronchial	+	1	VI	+	M	+		M	+	+	+	+	+	+	+	+	+	+	+	+	+	М	M	М	+	+	
Lymph node, mandibular	+		+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Lymph node, mesenteric	+		+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Lymph node, mediastinal	+	-	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Rhabdomyosarcoma, metastatic,																											
uncertain primary site	X																										
Spleen	+	-	+	+	+	+	+	+	+	A	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Гһутиѕ	+	•	+	+	+	M	+	M	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Rhabdomyosarcoma, metastatic, uncertain primary site	X																										
ntegumentary System																											
Mammary gland	+	-	+			+	+	+	+	M			+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Carcinoma					X			.,			X				.,	٠,				•	٠,						
Fibroadenoma								X					X		Х	X				Х	X			X			
Fibroadenoma, multiple Skin																											
Squamous cell carcinoma Squamous cell papilloma	+	•	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+ X	+	+	+	+	+	+	+	+	
Musculoskeletal System Sone	+		+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
	•			_	_		_		_	•					_	_	•									•	
Nervous System																											
Brain	+	•	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Carcinoma, metastatic,																											
pituitary gland																											
Respiratory System		•	. т	ъſ		N									λſ												
arynx	10					M	+	+	+	+	+	+	+	+	M		+	+	+	+	+	+	+	+	+	+	
Lung Alveolar/bronchiolar adenoma	+		+		+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Carcinoma, metastatic, mammary gland					X																						
Nose Frachea	+	•	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
.1 actiea	+	_	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Special Senses System Eye			+																			+					
Jrinary System																											
Kidney Urinary bladder	+		+	+	+	+ A	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Systemic Lesions																											
Multiple organs	+			+	+	+	+	+	+	+	+		+				+	+	+	+	+	+	+	+		+	
Leukemia mononuclear				X			X				X	X		X	X	X					X				X		

Individual Animal Tumor Pathology o																			_							
Number of Days on Study	7 3	7 3		7 3	7 3	7 3	7 3	7 3	7 3	7 3	7 3	7 3	7 3	7 3	7 3	7 3	7 3	7 3	7 3	7 3	7 3	7 3	7 3	7 3	7 3	
	5	5	5	5	5	5	5	5	5	5	5	6	6	6	6	6	6	6	6	6	6	6	6	6	6	
	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	Total
Carcass ID Number	7 3	7 6	7 7	7 9	8 1	8 2	8 9	9	9 2	9	9 4	5 2	5 3	5 9	6 0	6 8	7 0	7 5	8 0	8 5	8	9 1	9 6	9 8	9 9	Tissues/ Tumors
Hematopoietic System																										
Bone marrow	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Lymph node Lymph node, bronchial	_	4		_	_	_	_	М	+	+	+	М	+	+	+	+	_	_	_	_	_	_	М	+	+	4 41
Lymph node, mandibular	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Lymph node, mesenteric	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Lymph node, mediastinal Rhabdomyosarcoma, metastatic,	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
uncertain primary site																										1
Spleen Thymus	+	+	· +	+ 1 +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	49 47
Rhabdomyosarcoma, metastatic, uncertain primary site	т	_	. 10	1 —	_	_	_	_	_	_	_	_	_	_	_	_	_	_	_	_	_		_	_	Т	1
Integumentary System																										
Mammary gland	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	49
Carcinoma		_	, .																							2
Fibroadenoma	37		X		X	X	X	X	X						37	X		17				X	X		X	18
Fibroadenoma, multiple Skin	X +				_	_		_	_	_	_	_		_	X	+	_	X +	_	_	_	_	_	_	_	3 50
Squamous cell carcinoma	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	_	1
Squamous cell papilloma														X							X					2
Musculoskeletal System Bone																										50
						Т	Т	Т			+	Т	Т			Т			Т		+	+	Т		+	
Nervous System																										50
Brain Carcinoma, metastatic,	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
pituitary gland																		X								1
Respiratory System																										
Larynx	+	N	1 +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	44
Lung	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Alveolar/bronchiolar adenoma Carcinoma, metastatic, mammary gland														X												1 1
Nose	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Trachea	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Special Senses System Eye																				+						3
Urinary System																				-						
Kidney	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Urinary bladder	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	49
Systemic Lesions																										
Multiple organs	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Leukemia mononuclear		Х	X				X				X						X						X		X	16

Individual Animal Tumor Pathology	oi reili	aı,											<u> </u>	<u> </u>					,	CII				ус ррш
	4	4	1 6	6	6	6	6	6	6	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7
Number of Days on Study	4	6	0	2	4	5	6	9	9	0	0	1	2	2	3	3	3	3	3	3	3	3	3	3
3	8	2	2 4	9	6	5	7	0	9	3	9	1	2	3	4	4	4	4	4	5	5	5	5	5
	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3
Carcass ID Number	5	5	5	5	7	9	7	7	8	9	6	6	7	5	6	6	8	8	9	5	6	6	7	7
	3	6	8	1	7	6	3	8	8	5	7	3	5	2		9	2	7	2	7	1	5	4	6
Alimentary System																								
Esophagus	+	_	+ +	- 4	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Intestine large, colon	+	_	+ +	- +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Intestine large, rectum	+	_	+ +	- 4	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Intestine large, cecum	+	_		- 4	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Intestine small, duodenum	A			. 4	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Intestine small, jejunum	+		⊢ +	- A	ÀÀ	À	+	À	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Intestine small, ileum	À		` .	- 4					+	+	+	·	· -	<u>.</u>	<u>.</u>	<u>.</u>	·		·	·	· -		·	· -
Liver			ж Т L .I		- A				+	+	+	_	_	+	+	+	+						+	±
Mesentery	+	-	- 1	- 7		+		_	_	_		_	_	_	_	т	_	_		_	_	_	_	T
Mesentery Oral mucosa							+				+								+					
Pharyngeal, squamous cell papilloma																								
Pancreas	+	-	+ +	- +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Salivary glands	+	-	+ +	- +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Stomach, forestomach	+	-	+ +	- +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Stomach, glandular	+	-	+ +	- +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Tongue														+										
Schwannoma malignant														X										
Cardiovascular System																								
Blood vessel	+	-	+ +	- +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Heart	+	-	+ +	- +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Endocrine System																								
Adrenal cortex	+		- →	. 4	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Adrenal medulla							·	<u>.</u>		<u>.</u>	·	<u>.</u>	<u>.</u>	·	<u>.</u>	· -								
Islets, pancreatic					- +	+	+	+	+	<u>.</u>	·	·	· -	<u>.</u>	<u>.</u>	<u>.</u>	·		·	·	· -		·	· -
Adenoma	7		' 7	7	т	X		Τ'	-	Т	-	-	Т	Т		Т	-		-	7'	Τ'	Τ'	7-	'
Parathyroid gland					,	+		+		,	,	Ŋſ	,	,			,							
	+	-	r +	- +	- +							M		+	+	+	+	+	+	+	+	+	+	-
Pituitary gland	+	-	+ +						+		+	+ v	+	+			+	+ v	+		+ v	+		+
Pars distalis, adenoma				Σ		X	X	X		X		X			X	X		X		X	X		X	V
Pars distalis, adenoma, multiple																								X
Thyroid gland	+	-	+ +	- +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
C-cell, adenoma																					X			
C-cell, carcinoma																								
General Body System																								
None																								
Genital System																								
Clitoral gland			۱. ا				_	_	+	_	M	_	_	_	+	+	_	_	_				_	_
Ovary	+	-	. 7					T	-T		+	+		+	+	+	+	-T		-T	-T	T		+
Histiocytic sarcoma	+	-	- +	- 1	- +	+	+	+	+	+	+	+	+	+	т	_	+	+	+	+	+	+	+	т
Uterus																								
	+	-	- +	- +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Polyp stromal																								

	~	. ~	~	~	~	~	~	~	~	~	~	~	~	~	~	~	~	~	~	~	~	~	~	~	~	
N I CD CLI	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	
Number of Days on Study	3 5				3 5	3	3 5	3 5	3 5	3 5	3	3	3 6	3	3	3	3	3 6	3	3	3	3	3 6	3	3	
	0) j	Э	Э	Э	5	Э	Э	Э	Э	6	6	b	6	6	6	6	b	6	6	6	6	0	6	6	
	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	4	Total
Carcass ID Number	7	8	8	8	8	9	9	9	9	9	5	5	5	6	6	6	6	7	7	8	8	8	9	9	0	Tissues/
	9	1	4	5	6	0	3	4	7	9	4	5	9	0	2	4	6	0	2	0	3	9	1	8	0	Tumors
Alimentary System																							_			
Esophagus	4	- +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	49
Intestine large, colon	-	- +	- +		+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	49
Intestine large, rectum		- 4			·		·	·	·	·	·	·	<u>.</u>	·	<u>.</u>	<u>.</u>	<u>.</u>	<u>.</u>	<u>.</u>	<u>.</u>	·	·	·	·	+	49
Intestine large, rectum		- 4			+	_	_	·	· +	_	·	+	<u>.</u>	<u>.</u>	<u>.</u>	<u>.</u>	<u>.</u>	<u>.</u>	<u>'</u>	· +	· +	· +	· +	· +	<u>.</u>	49
Intestine small, duodenum		' ــــــــــــــــــــــــــــــــــــ			·			i	<u>.</u>	i	i	·	<u>.</u>	i	_	·	·	·		·			<u>.</u>		+	48
Intestine small, jejunum	7																								T.	45
Intestine small, jejunum Intestine small, ileum	7		- +	. +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
	+	- +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	46
Liver	+	- +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	49
Mesentery														+			+		+						+	7
Oral mucosa															+											1
Pharyngeal, squamous cell papilloma															X											1
Pancreas	+	- +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	49
Salivary glands	+	- +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	49
Stomach, forestomach	+	- +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	49
Stomach, glandular	+	- +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	49
Tongue																										1
Schwannoma malignant																										1
Cardiovascular System																										
Blood vessel	4	- 4	- +	. +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	49
Heart		- 4	- +	. +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	49
													<u>'</u>	_	_								<u> </u>			- 10
Endocrine System																										
Adrenal cortex	+	- +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	49
Adrenal medulla	+	- +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	49
Islets, pancreatic	+	- +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	49
Adenoma																										1
Parathyroid gland	+	- +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	M	+	+	+	47
Pituitary gland	+	- +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	49
Pars distalis, adenoma		Σ			X						X	X	X		X	X	X		X	X		X			X	24
Pars distalis, adenoma, multiple			X																					X		3
Гhyroid gland	+	- +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	49
C-cell, adenoma		Σ							X																	3
C-cell, carcinoma																						X				1
General Body System																							_			
None																										
Genital System																										
Clitoral gland								J	5	5	5	+	M	_	_	_	ر	_	ر				.1		_	47
	-	- +	- +	+	+	+	+	+	+	+	+	+	. 1VI	+	+	+	+	+	+	+	+	+	+	+	+	47
Ovary Histografia saraama	+	- + ,	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Histiocytic sarcoma	>																									1
Uterus	+	- +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+ X	+	+	+ X	+	+	+	+	+	+	49
Polyp stromal				X																						3

Individual Animal Tumor Pathol	y of Female Rats in the 2-Y	ear Inhalation Study of Ethylbenzene:	750 ppm (continued
Number of Days on Study	4 4 6 6 6 6 6 6 6 6 4 6 0 2 4 5 6 9		7 7 3 3
Tumber of Dujo on Study	8 2 4 9 6 5 7 0		5 5
a	3 3 3 3 3 3 3 3		3 3
Carcass ID Number	5 5 5 5 7 9 7 7 3 6 8 1 7 6 3 8	8 9 6 6 7 5 6 6 8 8 9 5 6 6 8 5 7 3 5 2 8 9 2 7 2 7 1 5	7 7 4 6
Hematopoietic System			
Bone marrow Lymph node	+ + + + + + + + + +	+ + + + + + + + + + + + + + + + + + + +	+ + +
Lymph node Lymph node, bronchial	+ + M + M + + +	- + + + M + + + M M M M + + +	+ +
Lymph node, mandibular	+ + + + + + +	- + + + + + + + + + + + + + +	+ +
Lymph node, mesenteric	+ + + + + + +	- + + + + + + + + + + + + + +	+ +
Lymph node, mediastinal	+ + + + + + + +	+ + + + + + + + + + + + + + +	+ +
Spleen	+ + + + + + + +	. + + + + + + + + + + + + + +	+ +
Thymus	+ + + + + + +	. + + + + + + + + + + + + + +	+ +
Integumentary System			
Mammary gland Adenoma	+ + + + + + + +	- + + + + + + + + + + + + + + + + X	+ +
Carcinoma, multiple			
Fibroadenoma		X X X X	X
Fibroadenoma, multiple		$X \qquad X \qquad X$	
Skin Sebaceous gland, carcinoma	+ + + + + + +	- + + + + + + + + + + + + + + +	+ +
Musculoskeletal System			
Bone	+ + + + + + + +	+ + + + + + + + + + + + + + + +	+ +
Nervous System			
Brain	+ + + + + + +	+ + + + + + + + + + + + + + + + + + + +	+ +
Respiratory System			
Larynx	+ + M + + + +	- M + + M + + + + + + + + + +	+ +
Lung	+ + + + + + + +	. + + + + + + + + + + + + + +	+ +
Nose	+ + + + + + + +	. + + + + + + + + + + + + + + +	+ +
Trachea	+ + + + + + +		+ +
Special Senses System Eye			
Harderian gland	+		
Urinary System			
Kidney	+ + + + + + + +	- + + + + + + + + + + + + + +	+ +
Renal tubule, adenoma			
Urinary bladder	+ + + + A + + +	+ + + + + + + + + + + + + + +	+ +
Systemic Lesions			
Multiple organs	+ + + + + + + +	. + + + + + + + + + + + + + +	+ +
Histiocytic sarcoma	V V V V V	V	
Leukemia mononuclear	X X X X X X	X	

Individual Animal Tumor Pathol	ogy of Fem	ale	·K	aus _		<u> </u>	e 2-	1 6	ar	III	nai	au	OII	วแ	iuy	-01		шу	/ IU	en.	LCII			_	եհո	(continue
	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	
Number of Days on Study	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	
, ,	5	5	5	5	5	5	5	5	5	5	6	6	6	6	6	6	6	6	6	6	6	6	6	6	6	
	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	4	Total
Carcass ID Number	7	8	8	8	8	9	9	9	9	9	5	5	5	6	6	6	6	7	7	8	8	8	9	9	0	Tissues/
	9					0			7		4				2						3		1		0	Tumors
Hematopoietic System																										
Bone marrow												,														49
Lymph node				٠ ٦		т		т	т	_	т	т	_	т	т	Τ	_	_	т	_	т	_	т	_	_	43
						1./	1 1/1		Nπ								+	_							M	38
Lymph node, bronchial	+	+	- +	. 1	- +	10	I M	. +	M	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+		
Lymph node, mandibular	+	+	. +	. +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	49
Lymph node, mesenteric	+	+	+	. +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	49
Lymph node, mediastinal	+	+	+	+	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	49
Spleen	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	49
Thymus	+	+	+	+	- M	+	+	+	+	+	+	+	+	M	+	+	+	+	+	+	+	+	+	+	+	47
ntegumentary System																										
Mammary gland	+	+	+	. 4	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	49
Adenoma																										1
Carcinoma, multiple																			X							1
		τ.		,				v	v			v				v			Λ	v	v		v			
Fibroadenoma		2	X					Λ	X			X	37	3.7		X				Λ	X		X			15
Fibroadenoma, multiple														X												6
Skin	+	+	+	. +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	49
Sebaceous gland, carcinoma										X																1
Musculoskeletal System																										
Bone	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	49
Nervous System																										
Brain	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	49
Respiratory System																							_			
arynx												,				M			,							45
	Τ.												Τ.	Τ.	Τ.		Τ.	Τ.	Τ.	Τ.	Τ.	Τ.			Τ.	
Lung	+	+	- +	. 4	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	49
Nose	+	+	+	. +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	49
<u> </u>	+	+	+	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	49
Special Senses System																										
Eye										+																1
Harderian gland																										1
Jrinary System																										
Kidney	+	+	+	. 4	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	49
Renal tubule, adenoma																	X									1
Jrinary bladder	+	+	+	- 4	- +	+	+	+	+	+	+	+	+	+	+	+		+	+	+	+	+	+	+	+	48
Systemic Lesions																										
Multiple organs	+	+	. 4	. 4	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	49
Histiocytic sarcoma	X			1	-				'			'							'		,		'			1
Leukemia mononuclear	Λ								v	X						X		X	v							11
Leukenna mononuciear									Λ	Λ						^		^	^							11

TABLE B3
Statistical Analysis of Primary Neoplasms in Female Rats in the 2-Year Inhalation Study of Ethylbenzene

	Chamber Control	75 ppm	250 ppm	750 ppm			
Adrenal Medulla: Benign or Malignant Pheoch	romocytoma						
Overall rate ^a	2/50 (4%)	3/50 (6%)	2/50 (4%)	0/49 (0%)			
Adjusted rate ^b	6.5%	8.6%	5.0%	0.0%			
Terminal rate ^c	2/31 (6%)	1/31 (3%)	0/34 (0%)	0/35 (0%)			
First incidencę (days)	734 (T)	719	659	e			
Life table test ^d	P = 0.101N	P = 0.516	P = 0.664N	P = 0.212N			
Logistic regression test ^d	P = 0.105N	P = 0.514	P = 0.678N	P = 0.212N			
Cochran-Armitage test ^d	P = 0.120N						
Fisher exact test ^d		P = 0.500	P = 0.691N	P = 0.253N			
Clitoral Gland: Adenoma or Carcinoma							
Overall rate	3/47 (6%)	0/49 (0%)	1/48 (2%)	0/47 (0%)			
Adjusted rate	10.3%	0.0%	3.1%	0.0%			
Γerminal rate	3/29 (10%)	0/30 (0%)	1/32 (3%)	0/34 (0%)			
First incidence (days)	734 (T)	_	734 (T)	_			
Life table test	P = 0.137N	P = 0.114N	P = 0.269N	P = 0.094N			
Logistic regression test	P = 0.137N	P = 0.114N	P = 0.268N	P = 0.094N			
Cochran-Armitage test	P = 0.162N						
Fisher exact test		P = 0.113N	P = 0.301N	P = 0.121N			
Kidney (Renal Tubule): Adenoma (Step Section							
Overall rate	0/50 (0%)	0/50 (0%)	1/50 (2%)	7/49 (14%)			
Adjusted rate	0.0%	0.0%	2.9%	19.4%			
Ferminal rate	0/31 (0%)	0/31 (0%)	1/34 (3%)	6/35 (17%)			
First incidence (days)	— D. 0.004	f	734 (T)	722			
Life table test	P< 0.001		P = 0.518	P= 0.015			
Logistic regression test	P< 0.001	_	P = 0.518	P = 0.014			
Cochran-Armitage test Fisher exact test	P< 0.001	_	P = 0.500	P = 0.006			
(C) 1 (C) 1 (C) 1 1 (C	G. G						
Kidney (Renal Tubule): Adenoma (Single and S	-	0/50 (00/)	1/50 (00/)	0/40 (100/)			
Overall rate	0/50 (0%)	0/50 (0%)	1/50 (2%)	8/49 (16%)			
Adjusted rate	0.0%	0.0%	2.9%	22.2%			
Cerminal rate	0/31 (0%)	0/31 (0%)	1/34 (3%)	7/35 (20%)			
First incidence (days)	P< 0.001	_	734 (T) P= 0.518	722 P= 0.008			
_ife table test _ogistic regression test	P< 0.001 P< 0.001	_	P = 0.518 P = 0.518	P = 0.008 P = 0.007			
Cochran-Armitage test	P< 0.001	_	r=0.316	r = 0.007			
Fisher exact test	r< 0.001	_	P = 0.500	P = 0.003			
Mammany Clands Ethnoodanama							
Mammary Gland: Fibroadenoma Overall rate	19/50 (38%)	19/50 (38%)	21/50 (42%)	21/49 (43%)			
Adjusted rate	55.8%	49.7%	55.0%	53.8%			
Terminal rate	16/31 (52%)	12/31 (39%)	17/34 (50%)	17/35 (49%)			
First incidence (days)	687	609	651	699			
Life table test	P = 0.512N	P = 0.564N	P = 0.561	P= 0.541N			
Logistic regression test	P = 0.549N	P = 0.554N	P = 0.501	P = 0.543N			
Cochran-Armitage test	P = 0.333	2 0.00111	1 0.000	2 0.01011			
Fisher exact test	1 0.000	P = 0.582N	P = 0.419	P = 0.387			
			*				

TABLE B3
Statistical Analysis of Primary Neoplasms in Female Rats in the 2-Year Inhalation Study of Ethylbenzene (continued)

	Chamber Control	75 ppm	250 ppm	750 ppm
Mammary Gland: Fibroadenoma or Adenoma				
Overall rate	20/50 (40%)	20/50 (40%)	21/50 (42%)	21/49 (43%)
Adjusted rate	56.9%	52.3%	55.0%	53.8%
Terminal rate	16/31 (52%)	13/31 (42%)	17/34 (50%)	17/35 (49%)
First incidence (days)	651	609	651	699
Life table test	P = 0.421N	P = 0.567N	P = 0.519N	P = 0.458N
Logistic regression test	P = 0.454N	P = 0.550N	P = 0.575N	P = 0.466N
Cochran-Armitage test	P = 0.419			
Fisher exact test		P = 0.581N	P = 0.500	P = 0.466
Mammary Gland: Carcinoma				
Overall rate	3/50 (6%)	1/50 (2%)	2/50 (4%)	1/49 (2%)
Adjusted rate	9.2%	3.2%	4.5%	2.9%
Terminal rate	2/31 (6%)	1/31 (3%)	0/34 (0%)	1/35 (3%)
First incidence (days)	704	734 (T)	502	734 (T)
Life table test	P = 0.306N	P = 0.301N	P = 0.471N	P = 0.263N
Logistic regression test	P = 0.353N	P = 0.295N	P = 0.511N	P = 0.266N
Cochran-Armitage test	P = 0.351N			
Fisher exact test		P = 0.309N	P = 0.500N	P = 0.316N
Mammary Gland: Adenoma or Carcinoma				
Overall rate	4/50 (8%)	2/50 (4%)	2/50 (4%)	2/49 (4%)
Adjusted rate	11.4%	6.5%	4.5%	5.7%
Terminal rate	2/31 (6%)	2/31 (6%)	0/34 (0%)	2/35 (6%)
First incidence (days)	651	734 (T)	502	734 (T)
Life table test	P = 0.327N	P = 0.342N	P = 0.313N	P = 0.290N
Logistic regression test	P = 0.378N	P = 0.322N	P = 0.349N	P = 0.303N
Cochran-Armitage test	P = 0.385N			
Fisher exact test		P = 0.339N	P = 0.339N	P = 0.349N
Mammary Gland: Fibroadenoma, Adenoma, o		00/70 (400/)	00/50 (400/)	00/10/(170/)
Overall rate	22/50 (44%)	20/50 (40%)	23/50 (46%)	22/49 (45%)
Adjusted rate	61.0%	52.3%	57.0%	56.3%
Terminal rate	17/31 (55%)	13/31 (42%)	17/34 (50%)	18/35 (51%)
First incidence (days)	651	609	502	699
Life table test	P = 0.396N	P = 0.412N	P = 0.509N	P= 0.367N
Logistic regression test	P = 0.449N	P = 0.377N	P = 0.561N	P = 0.368N
Cochran-Armitage test Fisher exact test	P = 0.436	P = 0.420N	P = 0.500	P= 0.545
Pituitary Gland (Pars Distalis): Adenoma Overall rate	30/49 (61%)	23/49 (47%)	25/50 (50%)	27/49 (55%)
Adjusted rate	78.5%	61.2%	63.6%	65.4%
Terminal rate	23/31 (74%)	17/31 (55%)	20/34 (59%)	21/35 (60%)
First incidence (days)	496	475	449	629
Life table test	P = 0.298N	P = 0.122N	P = 0.112N	P = 0.155N
Logistic regression test	P = 0.377N	P = 0.061N	P = 0.096N	P = 0.151N
Cochran-Armitage test	P = 0.546			
Fisher exact test		P = 0.112N	P = 0.178N	P = 0.341N

TABLE B3
Statistical Analysis of Primary Neoplasms in Female Rats in the 2-Year Inhalation Study of Ethylbenzene (continued)

	Chamber Control	75 ppm	250 ppm	750 ppm			
Pituitary Gland (Pars Distalis): Adeno	ma or Carcinoma						
Overall rate	30/49 (61%)	24/49 (49%)	26/50 (52%)	27/49 (55%)			
Adjusted rate	78.5%	62.3%	66.2%	65.4%			
Terminal rate	23/31 (74%)	17/31 (55%)	21/34 (62%)	21/35 (60%)			
First incidence (days)	496	475	449	629			
Life table test	P = 0.264N	P = 0.167N	P = 0.149N	P = 0.155N			
Logistic regression test	P = 0.332N	P = 0.089N	P = 0.134N	P = 0.151N			
Cochran-Armitage test	P = 0.511N						
Fisher exact test		P = 0.155N	P = 0.235N	P = 0.341N			
Skin: Squamous Cell Papilloma or Squ	uamous Cell Carcinoma						
Overall rate	0/50 (0%)	0/50 (0%)	3/50 (6%)	0/49 (0%)			
Adjusted rate	0.0%	0.0%	8.8%	0.0%			
Terminal rate	0/31 (0%)	0/31 (0%)	3/34 (9%)	0/35 (0%)			
First incidence (days)	_ ` ´	_ ` ′	734 (T)	_ ` ´			
Life table test	P = 0.617N	_	P = 0.137	_			
Logistic regression test	P = 0.617N	_	P = 0.137	_			
Cochran-Armitage test	P = 0.656N						
Fisher exact test		_	P = 0.121	_			
Thyroid Gland (C-cell): Adenoma							
Overall rate	2/48 (4%)	4/50 (8%)	2/50 (4%)	3/49 (6%)			
Adjusted rate	6.2%	12.4%	5.9%	8.6%			
Terminal rate	1/31 (3%)	3/31 (10%)	2/34 (6%)	3/35 (9%)			
First incidence (days)	721	731	734 (T)	734 (T)			
Life table test	P = 0.580N	P = 0.348	P = 0.666N	P= 0.554			
Logistic regression test	P = 0.585N	P = 0.354	P = 0.675N	P= 0.558			
Cochran-Armitage test	P= 0.567	1 0.001	1 0101011	1 0.000			
Fisher exact test	1 0.00	P = 0.359	P = 0.676N	P = 0.510			
Thyroid Gland (C-cell): Adenoma or (Carcinoma						
Overall rate	2/48 (4%)	4/50 (8%)	2/50 (4%)	4/49 (8%)			
Adjusted rate	6.2%	12.4%	5.9%	11.4%			
Terminal rate	1/31 (3%)	3/31 (10%)	2/34 (6%)	4/35 (11%)			
First incidence (days)	721	731	734 (T)	734 (T)			
Life table test	P = 0.436	P = 0.348	P = 0.666N	P= 0.394			
Logistic regression test	P= 0.431	P = 0.354	P = 0.675N	P= 0.396			
Cochran-Armitage test	P= 0.371	1 0.001	1 0.07014	1 0.000			
Fisher exact test	1 – 0.371	P = 0.359	P = 0.676N	P = 0.349			
Uterus: Stromal Polyp							
Overall rate	2/50 (4%)	3/50 (6%)	5/50 (10%)	3/49 (6%)			
Adjusted rate	6.2%	7.5%	14.7%	8.6%			
Adjusted rate Terminal rate	1/31 (3%)	0/31 (0%)	5/34 (15%)	3/35 (9%)			
First incidence (days)	721	601	734 (T)	734 (T)			
Life table test	P = 0.569	P = 0.511	P = 0.252	734 (1) P= 0.554			
	P = 0.369 P = 0.546	P = 0.511 P = 0.508	P = 0.232 P = 0.238	P= 0.554 P= 0.558			
Logistic regression test	P = 0.346 P = 0.495	r = 0.300	r = 0.230	1 = 0.336			
Cochran-Armitage test Fisher exact test	r=0.490	P = 0.500	P = 0.218	P = 0.490			
FISHEL CARCULESU		1 – 0.300	1 - 0.210	1 — 0.400			

TABLE B3
Statistical Analysis of Primary Neoplasms in Female Rats in the 2-Year Inhalation Study of Ethylbenzene (continued)

	Chamber Control	75 ppm	250 ppm	750 ppm
Uterus: Stromal Polyp or Stromal Sarcoma				
Overall rate	2/50 (4%)	4/50 (8%)	5/50 (10%)	3/49 (6%)
Adjusted rate	6.2%	9.3%	14.7%	8.6%
Terminal rate	1/31 (3%)	0/31 (0%)	5/34 (15%)	3/35 (9%)
First incidence (days)	721	462	734 (T)	734 (T)
Life table test	P = 0.545N	P = 0.355	P = 0.252	P = 0.554
Logistic regression test	P = 0.585	P = 0.309	P = 0.238	P = 0.558
Cochran-Armitage test	P = 0.568			
Fisher exact test		P = 0.339	P = 0.218	P = 0.490
All Organs: Mononuclear Cell Leukemia				
Overall rate	13/50 (26%)	18/50 (36%)	16/50 (32%)	11/49 (22%)
Adjusted rate	34.0%	46.2%	38.3%	25.0%
Terminal rate	7/31 (23%)	11/31 (35%)	9/34 (26%)	5/35 (14%)
First incidence (days)	507	412	486	448
Life table test	P = 0.118N	P = 0.217	P = 0.429	P = 0.315N
Logistic regression test	P = 0.195N	P = 0.222	P = 0.370	P = 0.458N
Cochran-Armitage test	P = 0.196N			
Fisher exact test		P = 0.194	P = 0.330	P = 0.430N
All Organs: Benign Neoplasms				
Overall rate	37/50 (74%)	39/50 (78%)	37/50 (74%)	39/49 (80%)
Adjusted rate	90.1%	88.5%	88.0%	86.7%
Terminal rate	27/31 (87%)	26/31 (84%)	29/34 (85%)	29/35 (83%)
First incidence (days)	496	475	449	629
Life table test	P = 0.293N	P = 0.441	P = 0.341N	P = 0.386N
Logistic regression test	P = 0.444N	P = 0.565	P = 0.381N	P = 0.500N
Cochran-Armitage test	P = 0.346			
Fisher exact test		P = 0.408	P = 0.590N	P = 0.337
All Organs: Malignant Neoplasms				
Overall rate	20/50 (40%)	25/50 (50%)	22/50 (44%)	14/49 (29%)
Adjusted rate	47.8%	56.7%	49.2%	31.9%
Terminal rate	10/31 (32%)	13/31 (42%)	12/34 (35%)	7/35 (20%)
First incidence (days)	318	412	393	448
Life table test	P = 0.029N	P = 0.264	P = 0.538	P = 0.107N
Logistic regression test	P = 0.055N	P = 0.335	P = 0.527	P = 0.165N
Cochran-Armitage test	P = 0.044N			
Fisher exact test		P = 0.211	P = 0.420	P = 0.162N

TABLE B3
Statistical Analysis of Primary Neoplasms in Female Rats in the 2-Year Inhalation Study of Ethylbenzene (continued)

	Chamber Control	75 ppm	250 ppm	750 ppm
All Organs: Benign or Malignant Neoplasms				
Overall rate	42/50 (84%)	45/50 (90%)	44/50 (88%)	46/49 (94%)
Adjusted rate	93.3%	90.0%	89.8%	93.9%
Terminal rate	28/31 (90%)	26/31 (84%)	29/34 (85%)	32/35 (91%)
First incidence (days)	318	412	393	448
Life table test	P = 0.389N	P = 0.395	P = 0.470N	P = 0.494N
Logistic regression test	P = 0.236	P = 0.442	P = 0.595	P = 0.318
Cochran-Armitage test	P = 0.124			
Fisher exact test		P = 0.277	P = 0.387	P = 0.106

(T)Terminal sacrifice

^b Kaplan-Meier estimated neoplasm incidence at the end of the study after adjustment for intercurrent mortality

Value of statistic cannot be computed.

Number of neoplasm-bearing animals/number of animals examined. Denominator is number of animals examined microscopically for adrenal gland, clitoral gland, kidney, pituitary gland, thyroid gland, and uterus; for other tissues, denominator is number of animals necropsied.

Observed incidence at terminal kill

Beneath the chamber control incidence are the P values associated with the trend test. Beneath the exposure group incidence are the P values corresponding to pairwise comparisons between the chamber controls and that exposure group. The life table test regards neoplasms in animals dying prior to terminal kill as being (directly or indirectly) the cause of death. The logistic regression test regards these lesions as nonfatal. The Cochran-Armitage and Fisher exact tests compare directly the overall incidence rates. For all tests, a negative trend or a lower incidence in an exposure group is indicated by N.

Not applicable; no neoplasms in animal group

TABLE B4
Summary of the Incidence of Nonneoplastic Lesions in Female Rats in the 2-Year Inhalation Study of Ethylbenzene^a

	Chamber Control	75	ppm	250	ppm	750) ppm
Disposition Summary							
Animals initially in study	50		50		50		50
Early deaths							
Moribund	7	1	14		8		6
Natural deaths	12		5		8		8
Survivors Died last week of study	1						
Terminal sacrifice	30		31		34		35
Missing	00	`	01		01		1
							_
Animals examined microscopically	50		50		50		49
Alimentary System							
Intestine large, colon	(47)	(49)		(48)		(49)	
Inflammation	1 (2%)						
Intestine large, rectum	(49)	(50)		(47)		(49)	
Arteriole, inflammation	1 (2%)	(50)		(47)		(40)	
Intestine large, cecum	(44)	(50)		(47)		(49)	
Inflammation Intestine small, ileum	1 (2%) (41)	(48)		(47)		(46)	
Hyperplasia	(41)	(40)			(2%)	(40)	
Inflammation					, ,		
Liver	(50)	(50)		(50)	(270)	(49)	
Angiectasis	3 (6%)	(00)		(00)			(12%)
Basophilic focus	23 (46%)	29 ((58%)	33	(66%)		(59%)
Clear cell focus	3 (6%)		(6%)		(2%)		(8%)
Congestion	1 (2%)				(2%)		(2%)
Degeneration		1 ((2%)	2	(4%)		
Eosinophilic focus	2 (4%)		(6%)	8	(16%)	5	(10%)
Fibrosis		1 ((2%)				
Hematopoietic cell proliferation	1 (2%)						
Hemorrhage	1 (2%)		(00/)		(00.4)	_	(4.007)
Hepatodiaphragmatic nodule	4 (8%)	4 ((8%)		(8%)	5	(10%)
Infiltration cellular, lymphocyte Inflammation, acute				1	(2%)	1	(20/)
Inflammation, acute Inflammation, chronic	3 (6%)	9 ((4%)	Q	(6%)		(2%) (4%)
Mixed cell focus	5 (10%)	۷ ((1/0)		(2%)	۷	(1/0)
Necrosis	1 (2%)	1 ((2%)		(4%)	2	(4%)
Vacuolization cytoplasmic	11 (22%)		(24%)		(28%)		(29%)
Centrilobular, degeneration	1 (2%)	(/		· · - · - /	11	,
Portal vein, thrombosis	, ,	1 ((2%)				
Mesentery	(7)	(4)	•	(6)		(7)	
Artery, degeneration				1	(17%)		
Artery, inflammation	1 (14%)				(17%)		
Fat, necrosis	6 (86%)		(100%)		(83%)		(100%)
Pancreas	(49)	(50)		(50)	(00/)	(49)	
Cyst	0 (40/)			1	(2%)		
Inflammation	2 (4%)	10 /	(260/)	10	(260/)	10	(200/)
Acinus, atrophy Arteriole, inflammation	18 (37%)	18 ((36%)		(36%) (2%)		(39%) (2%)
Artery, inflammation	1 (2%)				(2%)	1	(270)
Artery, Illianimadoli	1 (470)			1	(2 /0)		

 $^{^{\}mathrm{a}}$ Number of animals examined microscopically at the site and the number of animals with lesion

TABLE B4
Summary of the Incidence of Nonneoplastic Lesions in Female Rats in the 2-Year Inhalation Study of Ethylbenzene (continued)

	Chamber Control	75 ppm	250 ppm	750 ppm
Alimentary System (continued)				
Stomach, forestomach	(49)	(50)	(50)	(49)
Hemorrhage	()	()	(==)	1 (2%)
Hyperkeratosis	2 (4%)		1 (2%)	- ()
Hyperplasia	3 (6%)	1 (2%)	3 (6%)	8 (16%)
Inflammation	1 (2%)	1 (2%)	0 (070)	6 (12%)
Ulcer	3 (6%)	4 (8%)	1 (2%)	0 (12/0)
Stomach, glandular	(49)	(49)	(49)	(49)
Cyst	(43)	(43)	(43)	1 (2%)
Hyperplasia	1 (2%)			1 (270)
Inflammation	1 (2/0)			2 (4%)
	F (100/)		1 (90/)	, ,
Necrosis	5 (10%)		1 (2%)	1 (2%)
Pigmentation	1 (2%)	4 (00()		4 (00()
Ulcer		4 (8%)		1 (2%)
Glands, cyst				1 (2%)
Cardiovascular System				
Blood vessel	(49)	(50)	(50)	(49)
Degeneration Degeneration	(20)	(00)	1 (2%)	(20)
Inflammation			1 (2%)	
Aorta, inflammation	1 (2%)		1 (270)	
	(50)	(50)	(50)	(49)
Heart			(50)	
Cardiomyopathy	6 (12%)	1 (2%)		6 (12%)
Fibrosis			1 (00/)	1 (2%)
Inflammation			1 (2%)	
Atrium, inflammation			1 (2%)	1 (00/)
Atrium, thrombosis		4 (00()	4 (00/)	1 (2%)
Endocardium, hyperplasia		1 (2%)	1 (2%)	
Myocardium, hypertrophy	4 (00/)	1 (2%)		
Valve, degeneration	1 (2%)			
Endocrine System				
Adrenal cortex	(50)	(50)	(50)	(49)
Angiectasis	3 (6%)	3 (6%)	4 (8%)	3 (6%)
Cytoplasmic alteration	1 (2%)	3 (6%)	1 (2%)	- ()
Degeneration	- (~,0)	- (0.0)	- (2/0)	1 (2%)
Degeneration, cystic	5 (10%)	3 (6%)	2 (4%)	4 (8%)
Hemorrhage	4 (8%)	5 (10%)	2 (4%)	1 (2%)
Hyperplasia	3 (6%)	4 (8%)	2 (4%)	3 (6%)
	3 (0/0)	4 (0/0)	3 (6%)	3 (0/0)
Hypertrophy	1 (90/)			1 (90/)
Necrosis	1 (2%)		1 (2%)	1 (2%)
Pigmentation	10 (040/)	r (400/)	1 (2%)	1 (2%)
Vacuolization cytoplasmic	12 (24%)	5 (10%)	12 (24%)	6 (12%)
Adrenal medulla	(50)	(50)	(50)	(49)
Hemorrhage		1 (2%)		
Hyperplasia	4 (8%)		2 (4%)	2 (4%)
Infiltration cellular, lymphocyte			1 (2%)	
Necrosis		1 (2%)		
Parathyroid gland	(48)	(46)	(42)	(47)
Atrophy	1 (2%)			
Hyperplasia	5 (10%)	2 (4%)	4 (10%)	5 (11%)

TABLE B4
Summary of the Incidence of Nonneoplastic Lesions in Female Rats in the 2-Year Inhalation Study of Ethylbenzene (continued)

	Chamber Control	75 ppm	250 ppm	75 0 ppm
Endocrine System (continued)				
Pituitary gland	(49)	(49)	(50)	(49)
Cyst	1 (2%)			
Infiltration cellular, mixed cell	1 (2%)			
Necrosis	1 (2%)			
Pars distalis, angiectasis	2 (4%)	16 (33%)	6 (12%)	2 (4%)
Pars distalis, cyst	5 (10%)	1 (2%)	3 (6%)	3 (6%)
Pars distalis, degeneration		1 (2%)		
Pars distalis, hemorrhage	44 (000)	2 (4%)	44 (000)	45 (050)
Pars distalis, hyperplasia	11 (22%)	9 (18%)	14 (28%)	17 (35%)
Pars distalis, pigmentation		2 (4%)		
Pars intermedia, angiectasis	(40)	2 (4%)	(50)	(40)
Thyroid gland	(48)	(50)	(50)	(49)
Hyperplasia	1 (90/)		1 (2%)	
Inflammation	1 (2%)			1 (2%)
Bilateral, C-cell, hyperplasia C-cell, hyperplasia	5 (10%)	5 (10%)	5 (10%)	1 (2%) 5 (10%)
C-cell, inflammation	5 (10%) 1 (2%)	3 (10%)	J (10%)	5 (10%)
Follicle, cyst	1 (2/0)		1 (2%)	
Tomele, cyst			1 (270)	
Genital System	(47)	(40)	(40)	(47)
Clitoral gland	(47)	(49)	(48)	(47)
Cyst Hyperplasia	4 (9%)	3 (6%)	1 (2%)	1 (2%) 3 (6%)
Inflammation	6 (13%)	5 (10%)	4 (8%)	4 (9%)
Bilateral, hyperplasia	0 (1370)	J (1070)	4 (670)	1 (2%)
Ovary	(50)	(50)	(50)	(49)
Cyst	5 (10%)	6 (12%)	5 (10%)	5 (10%)
		0 (12/0)	0 (1070)	0 (1070)
Corpus luteum, hyperplasia	1 (2%)			
Corpus luteum, hyperplasia Uterus	1 (2%) (50)	(50)	(50)	(49)
Uterus		(50)	(50)	(49)
	(50)	(50) 3 (6%)	1 (2%)	(49) 4 (8%)
Uterus Angiectasis	(50) 1 (2%)	, ,		4 (8%) 1 (2%)
Uterus Angiectasis Hydrometra	(50) 1 (2%)	3 (6%)	1 (2%)	4 (8%)
Uterus Angiectasis Hydrometra Inflammation Endometrium, cyst Vagina	(50) 1 (2%)	3 (6%)	1 (2%)	4 (8%) 1 (2%)
Uterus Angiectasis Hydrometra Inflammation Endometrium, cyst Vagina Fibrosis	(50) 1 (2%)	3 (6%) (3) 1 (33%)	1 (2%)	4 (8%) 1 (2%)
Uterus Angiectasis Hydrometra Inflammation Endometrium, cyst Vagina	(50) 1 (2%)	3 (6%)	1 (2%)	4 (8%) 1 (2%)
Uterus Angiectasis Hydrometra Inflammation Endometrium, cyst Vagina Fibrosis Arteriole, degeneration	(50) 1 (2%)	3 (6%) (3) 1 (33%)	1 (2%)	4 (8%) 1 (2%)
Uterus Angiectasis Hydrometra Inflammation Endometrium, cyst Vagina Fibrosis Arteriole, degeneration Hematopoietic System	(50) 1 (2%) 1 (2%)	3 (6%) (3) 1 (33%) 1 (33%)	1 (2%) 1 (2%)	4 (8%) 1 (2%) 1 (2%)
Uterus Angiectasis Hydrometra Inflammation Endometrium, cyst Vagina Fibrosis Arteriole, degeneration Hematopoietic System Bone marrow	(50) 1 (2%) 1 (2%) (49)	3 (6%) (3) 1 (33%)	1 (2%)	4 (8%) 1 (2%) 1 (2%)
Uterus Angiectasis Hydrometra Inflammation Endometrium, cyst Vagina Fibrosis Arteriole, degeneration Hematopoietic System Bone marrow Atrophy	(50) 1 (2%) 1 (2%) (49) 1 (2%)	3 (6%) (3) 1 (33%) 1 (33%) (50)	1 (2%) 1 (2%)	4 (8%) 1 (2%) 1 (2%) (49) 1 (2%)
Uterus Angiectasis Hydrometra Inflammation Endometrium, cyst Vagina Fibrosis Arteriole, degeneration Hematopoietic System Bone marrow Atrophy Hemorrhage	(49) 1 (2%) 1 (2%) (49) 1 (2%) 3 (6%)	3 (6%) (3) 1 (33%) 1 (33%) (50) 2 (4%)	1 (2%) 1 (2%)	(49) 1 (2%) 1 (2%) 1 (2%)
Uterus Angiectasis Hydrometra Inflammation Endometrium, cyst Vagina Fibrosis Arteriole, degeneration Hematopoietic System Bone marrow Atrophy Hemorrhage Hyperplasia	(50) 1 (2%) 1 (2%) (49) 1 (2%)	3 (6%) (3) 1 (33%) 1 (33%) (50) 2 (4%) 8 (16%)	1 (2%) 1 (2%)	4 (8%) 1 (2%) 1 (2%) (49) 1 (2%)
Uterus Angiectasis Hydrometra Inflammation Endometrium, cyst Vagina Fibrosis Arteriole, degeneration Hematopoietic System Bone marrow Atrophy Hemorrhage Hyperplasia Hyperplasia, mast cell	(49) 1 (2%) 1 (2%) (49) 1 (2%) 3 (6%) 7 (14%)	3 (6%) (3) 1 (33%) 1 (33%) (50) 2 (4%)	1 (2%) 1 (2%)	(49) 1 (2%) 1 (2%) 1 (2%)
Uterus Angiectasis Hydrometra Inflammation Endometrium, cyst Vagina Fibrosis Arteriole, degeneration Hematopoietic System Bone marrow Atrophy Hemorrhage Hyperplasia	(49) 1 (2%) 1 (2%) (49) 1 (2%) 3 (6%)	3 (6%) (3) 1 (33%) 1 (33%) (50) 2 (4%) 8 (16%) 1 (2%)	1 (2%) 1 (2%)	(49) 1 (2%) 1 (2%) 1 (2%)
Uterus Angiectasis Hydrometra Inflammation Endometrium, cyst Vagina Fibrosis Arteriole, degeneration Hematopoietic System Bone marrow Atrophy Hemorrhage Hyperplasia Hyperplasia, mast cell Myelofibrosis	(49) 1 (2%) 1 (2%) (49) 1 (2%) 3 (6%) 7 (14%)	3 (6%) (3) 1 (33%) 1 (33%) (50) 2 (4%) 8 (16%)	1 (2%) 1 (2%) (50) 7 (14%)	(49) 1 (2%) 1 (2%) 1 (2%)

TABLE B4
Summary of the Incidence of Nonneoplastic Lesions in Female Rats in the 2-Year Inhalation Study of Ethylbenzene (continued)

	Chamber Control	75 ppm	250 ppm	750 ppm
Hematopoietic System (continued)				
Lymph node	(3)	(3)	(4)	(4)
Pancreatic, hemorrhage		• •	1 (25%)	1 (25%)
Pancreatic, infiltration cellular, histiocyte				1 (25%)
Renal, hemorrhage				2 (50%)
Renal, pigmentation				1 (25%)
Lymph node, bronchial	(37)	(34)	(41)	(38)
Atrophy	1 (3%)			
Ectasia			1 (2%)	
Hemorrhage	4 (11%)	6 (18%)	5 (12%)	3 (8%)
Hyperplasia, lymphoid		1 (3%)		
Infiltration cellular, histiocyte			1 (2%)	
Necrosis			1 (2%)	
Pigmentation	6 (16%)	7 (21%)	7 (17%)	5 (13%)
Lymph node, mandibular	(49)	(50)	(50)	(49)
Ectasia		1 (2%)	1 (2%)	
Hemorrhage	3 (6%)	2 (4%)	1 (2%)	3 (6%)
Hyperplasia, plasma cell	2 (4%)	9 (18%)	4 (8%)	3 (6%)
Infiltration cellular, histiocyte	1 (2%)			
Pigmentation	()	41	()	1 (2%)
Lymph node, mesenteric	(49)	(50)	(50)	(49)
Amyloid deposition	1 (2%)			
Atrophy	1 (2%)			
Ectasia			3 (6%)	
Hemorrhage	8 (16%)	6 (12%)	8 (16%)	7 (14%)
Hyperplasia				1 (2%)
Hyperplasia, plasma cell			1 (2%)	
Infiltration cellular, histiocyte	1 (2%)			
Inflammation	1 (2%)		1 (2%)	
Lymph node, mediastinal	(49)	(49)	(50)	(49)
Edema		1 (2%)		
Hemorrhage	21 (43%)	15 (31%)	13 (26%)	21 (43%)
Hyperplasia, plasma cell	1 (2%)			
Infiltration cellular, histiocyte	1 (2%)			
Necrosis	1 (2%)			
Pigmentation	22 (45%)	22 (45%)	25 (50%)	27 (55%)
Spleen	(49)	(50)	(49)	(49)
Hematopoietic cell proliferation	6 (12%)	3 (6%)	3 (6%)	3 (6%)
Hemorrhage		2 (4%)		
Hyperplasia, lymphoid		1 (2%)		
Necrosis				1 (2%)
Pigmentation	1 (2%)	1 (2%)		
Red pulp, atrophy	1 (2%)			
Chymus	(48)	(47)	(47)	(47)
Atrophy	1 (2%)			
Cyst				2 (4%)
Integumentary System				
Mammary gland	(48)	(50)	(49)	(49)
Galactocele	10 (21%)	10 (20%)	10 (20%)	11 (22%)
Hyperplasia	13 (27%)	19 (38%)	21 (43%)	18 (37%)
Infiltration cellular, lymphocyte	13 (21/0)	1 (2%)	£1 (43/0)	10 (37/0)
minu ation central, tymphocyte		1 (2/0)		

TABLE B4
Summary of the Incidence of Nonneoplastic Lesions in Female Rats in the 2-Year Inhalation Study of Ethylbenzene (continued)

	Chamber Control	75 ppm	250 ppm	750 ppm
Integumentary System (continued)				
Skin	(50)	(50)	(50)	(49)
Cyst epithelial inclusion Infiltration cellular, lymphocyte		1 (2%)	1 (2%)	
Inflammation, chronic	1 (2%)	1 (270)		
Ulcer	1 (2%)		1 (2%)	2 (4%)
Epidermis, hyperplasia			2 (4%)	1 (2%)
Subcutaneous tissue, fibrosis		1 (2%)	1 (00/)	
Subcutaneous tissue, inflammation			1 (2%)	
Musculoskeletal System				
Bone	(50)	(50)	(50)	(49)
Fibrous osteodystrophy	- ()	1 (2%)	- 4	. (==)
Hyperostosis	2 (4%)	5 (10%)	5 (10%)	1 (2%)
Turbinate, hyperostosis		2 (4%)	2 (4%)	1 (2%)
Nervous System				
Brain	(50)	(50)	(50)	(49)
Hemorrhage	1 (2%)	4 (00/)		1 (2%)
Hydrocephalus Necrosis	1 (2%)	1 (2%)		1 (2%)
Respiratory System				
Larynx	(45)	(43)	(44)	(45)
Infiltration cellular, lymphocyte	, ,	3 (7%)	1 (2%)	, ,
Inflammation		2 (5%)	1 (2%)	2 (4%)
Metaplasia, squamous	1 (2%)		4 (00/)	0 (40()
Respiratory epithelium, hyperplasia Respiratory epithelium, metaplasia, squamo	us 1 (2%)	1 (2%)	1 (2%)	2 (4%) 2 (4%)
Lung	(50)	(50)	(50)	(49)
Congestion	5 (10%)	(30)	1 (2%)	(10)
Edema	0 (10/0)	1 (2%)	1 (2%)	
Fibrosis	1 (2%)	1 (2%)	1 (2%)	
Hemorrhage	1 (2%)	2 (4%)	2 (4%)	
Infiltration cellular, histiocyte	2 (4%)	3 (6%)	1 (2%)	3 (6%)
Inflammation, chronic	2 (4%)		1 (00/)	3 (6%)
Inflammation, chronic active	1 (90/)	1 (90/)	1 (2%)	9 (40/)
Inflammation, granulomatous Alveolar epithelium, hyperplasia	1 (2%) 1 (2%)	1 (2%) 5 (10%)	1 (2%) 2 (4%)	2 (4%) 5 (10%)
A A vectar epithenum, hyperpiasia	1 (2/0)	J (1070)	ω (4/0)	J (1070)

TABLE B4
Summary of the Incidence of Nonneoplastic Lesions in Female Rats in the 2-Year Inhalation Study of Ethylbenzene (continued)

	Chamber Control	75 ppm	250 ppm	750 ppm
Respiratory System (continued)				
Nose	(50)	(50)	(50)	(49)
Angiectasis			1 (2%)	
Congestion	1 (2%)			
Foreign body	4 (8%)	1 (2%)		1 (2%)
Infiltration cellular, lymphocyte				1 (2%)
Inflammation	10 (20%)	10 (20%)	5 (10%)	1 (2%)
Necrosis			1 (2%)	
Thrombosis	1 (2%)			
Glands, cyst			1 (2%)	
Glands, hyperplasia		1 (2%)	2 (4%)	1 (2%)
Goblet cell, hyperplasia				1 (2%)
Nasolacrimal duct, inflammation	2 (4%)	1 (2%)		
Nasolacrimal duct, metaplasia, squamous	1 (2%)	~ (440/)	~ (400()	
Respiratory epithelium, hyperplasia	6 (12%)	7 (14%)	5 (10%)	
Respiratory epithelium, metaplasia, squamo	us	1 (2%)		
Respiratory epithelium, ulcer Trachea	(50)	2 (4%) (50)	(50)	(49)
Tractica	(30)	(30)	(30)	(43)
Special Senses System	(4)		(0)	(4)
Eye	(1)		(3)	(1)
Lens, cataract	1 (100%)		2 (67%)	1 (100%)
Retina, degeneration	1 (100%)			1 (100%)
Harderian gland Inflammation				(1) 1 (100%)
miammation				1 (100%)
Urinary System				
Kidney	(50)	(50)	(50)	(49)
Cyst	1 (2%)	1 (2%)	(30)	4 (8%)
Infarct	1 (2%)	1 (2/0)	1 (2%)	T (0/0)
Mineralization	8 (16%)	11 (22%)	7 (14%)	
Necrosis	0 (2070)	11 (~~,0)	1 (2%)	
Nephropathy	38 (76%)	42 (84%)	43 (86%)	46 (94%)
Pigmentation	4 (8%)	10 (20%)	8 (16%)	3 (6%)
Arteriole, inflammation	1 (2%)	• ,	` ,	` '
Renal tubule, degeneration	, ,		1 (2%)	
Renal tubule, hyperplasia		1 (2%)	3 (6%)	3 (6%)
Transitional epithelium, hyperplasia			, ,	2 (4%)

APPENDIX C SUMMARY OF LESIONS IN MALE MICE IN THE 2-YEAR INHALATION STUDY OF ETHYLBENZENE

TABLE C1	Summary of the Incidence of Neoplasms in Male Mice	
	in the 2-Year Inhalation Study of Ethylbenzene	132
TABLE C2	Individual Animal Tumor Pathology of Male Mice	
	in the 2-Year Inhalation Study of Ethylbenzene	136
TABLE C3	Statistical Analysis of Primary Neoplasms in Male Mice	
	in the 2-Year Inhalation Study of Ethylbenzene	158
TABLE C4	Historical Incidence of Alveolar/bronchiolar Neoplasms	
	in Chamber Control Male B6C3F ₁ Mice	161
TABLE C5	Summary of the Incidence of Nonneoplastic Lesions in Male Mice	
	in the 2-Year Inhalation Study of Ethylbenzene	162

TABLE C1 Summary of the Incidence of Neoplasms in Male Mice in the 2-Year Inhalation Study of Ethylbenzene^a

•	Chamber Control	75 ppm	250 ppm	750 ppm
Disposition Summary				
Animals initially in study	50	50	50	50
Early deaths				
Accidental deaths	1			1
Moribund	6	2	5	6
Natural deaths	15	12	13	13
Survivors				
Died last week of study			1	
Terminal sacrifice	28	36	31	30
Animals examined microscopically	50	50	50	50
Alimentary System				
Intestine large, cecum	(42)	(46)	(44)	(43)
Intestine small, jejunum	(44)	(46)	(44)	(43)
Epithelium, carcinoma	(==/	1 (2%)	\==/	(/
Intestine small, ileum	(42)	(46)	(44)	(41)
Liver	(50)	(50)	(50)	(50)
Alveolar/bronchiolar carcinoma, metastatic,	()	()	()	()
lung				1 (2%)
Cholangiocarcinoma		1 (2%)		()
Fibrosarcoma, metastatic, stomach, glandula	r 1 (2%)	(/		
Hemangioma	1 (2%)			
Hemangiosarcoma	, ,	1 (2%)		
Hepatoblastoma		1 (2%)		
Hepatocellular carcinoma	17 (34%)	8 (16%)	11 (22%)	10 (20%)
Hepatocellular carcinoma, multiple		1 (2%)	2 (4%)	
Hepatocellular adenoma	11 (22%)	12 (24%)	17 (34%)	17 (34%)
Hepatocellular adenoma, multiple	1 (2%)	4 (8%)		1 (2%)
Hepatocholangiocarcinoma	1 (2%)		1 (2%)	1 (2%)
Mesentery	(1)		(1)	
Hepatocellular carcinoma, metastatic, liver	1 (100%)			
Hepatocholangiocarcinoma, metastatic, liver			1 (100%)	
Pancreas	(49)	(50)	(48)	(48)
Acinus, hepatocholangiocarcinoma, metastat	ic,			
liver			1 (2%)	
Stomach, forestomach	(48)	(50)	(50)	(47)
Fibrosarcoma, metastatic, stomach, glandula	r 1 (2%)			
Squamous cell papilloma	()	1 (2%)	45.51	()
Stomach, glandular	(48)	(50)	(50)	(47)
Fibrosarcoma	1 (2%)		(1)	443
Tooth			(1)	(1)
Odontoma				1 (100%)
Cardiovascular System				
Blood vessel	(48)	(48)	(49)	(47)
Aorta, fibrosarcoma, metastatic, stomach,				
glandular	1 (2%)			
Aorta, hepatocellular carcinoma, metastatic,				
liver	1 (2%)			
Aorta, sarcoma				1 (2%)

TABLE C1
Summary of the Incidence of Neoplasms in Male Mice in the 2-Year Inhalation Study of Ethylbenzene (continued)

(Chamber Control	75 ppm	250 ppm	750 ppm
Cardiovascular System (continued)	(50)	(50)	(50)	(50)
Alveolar/bronchiolar carcinoma, metastatic, lung				1 (2%)
Fibrosarcoma, metastatic, stomach, glandular Hepatocholangiocarcinoma, metastatic, liver	1 (2%)		1 (2%)	, ,
Pericardium, hepatocholangiocarcinoma, metastatic, liver	1 (2%)			
Endocrine System	(45)	(45)	(40)	(10)
Adrenal cortex Adenoma Carcinoma	(47) 1 (2%)	(47)	(48) 1 (2%) 1 (2%)	(48)
Hepatocellular carcinoma, metastatic, liver	1 (2%)			
Islets, pancreatic Adenoma	(49)	(50)	(48)	(48) 1 (2%)
Carcinoma	(44)	(45)	(45)	1 (2%)
Pituitary gland Pars distalis, carcinoma	(44)	(45)	(45) 1 (2%)	(47)
Thyroid gland Follicular cell, adenoma	(50) 3 (6%)	(50) 2 (4%)	(50) 1 (2%)	(50) 5 (10%)
Follicular cell, adenoma, multiple	0 (0/0)	2 (170)	1 (270)	1 (2%)
General Body System				
Tissue NOS	(2)	(3)	(1)	(2)
Fibrosarcoma Fat, hepatocholangiocarcinoma, metastatic, liver	1 (50%)		1 (100%)	
Thoracic, hepatocholangiocarcinoma,			1 (100%)	
metastatic, liver Thoracic, sarcoma	1 (50%)			1 (50%)
Genital System Epididymis	(49)	(50)	(50)	(50)
Leiomyoma Seminal vesicle	(49)	1 (2%) (50)	(50)	(50)
Hepatocholangiocarcinoma, metastatic, liver	, ,	, ,	1 (2%)	, ,
Testes Interstitial cell, adenoma	(49) 1 (2%)	(50)	(50) 1 (2%)	(50) 1 (2%)
Hematopoietic System Bone marrow	(50)	(50)	(50)	(50)
Lymph node	(4)	(7)	(11)	(3)
Fibrosarcoma, metastatic, stomach, glandular Pancreatic, carcinoma	1 (25%)			1 (33%)
Popliteal, hemangioma		4 (440)	1 (9%)	- (0070)
Renal, cholangiocarcinoma, metastatic, liver Renal, fibrosarcoma, metastatic, stomach,		1 (14%)		
glandular	1 (25%)			
Renal, hepatocholangiocarcinoma, metastatic liver	,		1 (9%)	

TABLE C1
Summary of the Incidence of Neoplasms in Male Mice in the 2-Year Inhalation Study of Ethylbenzene (continued)

•	Chamber Control	75 ppm	250 ppm	750 ppm
Hematopoietic System (continued)				
Lymph node, bronchial Alveolar/bronchiolar carcinoma, metastatic, lung	(14)	(24)	(27)	(27) 1 (4%)
Fibrosarcoma, metastatic, stomach, glandula Hepatocholangiocarcinoma, metastatic, liver	r 1 (7%)		1 (4%)	
Sarcoma Lymph node, mandibular Sarcoma, metastatic, nose	(43) 1 (2%)	(45)	(46)	1 (4%) (44)
Lymph node, mesenteric Hepatocholangiocarcinoma, metastatic, liver Lymph node, mediastinal	(45) (24)	(46) (25)	(47) 1 (2%) (27)	(48) (25)
Fibrosarcoma, metastatic, stomach, glandula Hepatocholangiocarcinoma, metastatic, liver Sarcoma		(20)	(21)	1 (4%)
Spleen Thymus Alveolar/bronchiolar carcinoma, metastatic,	(50) (37)	(50) (37)	(49) (39)	(49) (34)
lung Hepatocholangiocarcinoma, metastatic, liver Sarcoma	1 (3%)			1 (3%) 1 (3%)
Integumentary System	(50)	(50)	(70)	(70)
Skin Fibrosarcoma Hemangioma	(50) 1 (2%)	(50)	(50) 1 (2%)	(50)
Musculoskeletal System				
Bone Sternum, fibrosarcoma, metastatic, stomach, glandular	(50) 1 (2%)	(49)	(50)	(50)
Skeletal muscle Alveolar/bronchiolar carcinoma, metastatic,	(2)		(2)	
lung Fibrosarcoma, metastatic, stomach, glandula Hepatocellular carcinoma, metastatic, liver Hepatocholangiocarcinoma, metastatic, liver	r 1 (50%) 1 (50%)		1 (50%) 1 (50%)	
Nervous System Brain	(50)	(50)	(50)	(50)

TABLE C1 Summary of the Incidence of Neoplasms in Male Mice in the 2-Year Inhalation Study of Ethylbenzene (continued)

	Chamber Control	75 ppm	250 ppm	750 ppm
Respiratory System				
Larynx	(48)	(49)	(46)	(49)
Lung	(50)	(50)	(50)	(50)
Alveolar/bronchiolar adenoma	5 (10%)	8 (16%)	9 (18%)	15 (30%)
Alveolar/bronchiolar adenoma, multiple		1 (2%)	1 (2%)	1 (2%)
Alveolar/bronchiolar carcinoma	2 (4%)	1 (2%)	5 (10%)	3 (6%)
Cholangiocarcinoma, metastatic, liver		1 (2%)		
Fibrosarcoma, metastatic, stomach, glandula				
Hepatocellular carcinoma, metastatic, liver	5 (10%)	3 (6%)	5 (10%)	3 (6%)
Hepatocholangiocarcinoma, metastatic, liver	1 (2%)		1 (2%)	1 (2%)
Bronchiole, polyp adenomatous		1 (2%)		
Mediastinum, sarcoma				1 (2%)
Nose	(50)	(50)	(50)	(50)
Sarcoma	1 (2%)			(4)
Pleura				(1)
Alveolar/bronchiolar carcinoma, metastatic,				4 (4000()
lung	(50)	(50)	(50)	1 (100%)
Trachea	(50)	(50)	(50)	(50)
Special Senses System				
Harderian gland	(2)	(3)	(2)	
Adenoma	1 (50%)	3 (100%)	2 (100%)	
Urinary System Kidney Alveolar/bronchiolar carcinoma, metastatic, lung Cholangiocarcinoma, metastatic, liver Fibrosarcoma, metastatic, stomach, glandula Hepatocellular carcinoma, metastatic, liver Renal tubule, adenoma	(50) ur 1 (2%) 1 (2%)	(50) 1 (2%) 1 (2%)	(50) 1 (2%) 1 (2%)	(50) 1 (2%)
Systemic Lesions				
Multiple organs ^b	(50)	(50)	(50)	(50)
Leukemia granulocytic	\/	N= =7	1 (2%)	1 (2%)
Lymphoma malignant	2 (4%)	2 (4%)	3 (6%)	2 (4%)
Neoplasm Summary				
Total animals with primary neoplasms ^c	35	34	40	41
Total primary neoplasms	50	49	60	68
Total animals with benign neoplasms	20	25	26	30
Total benign neoplasms	24	33	35	43
Total suimals with malignant nasalasma	21	15	21	16
rotat animais with matignant neodiasms				
Total malignant neoplasms Total malignant neoplasms	26	16	25	۷.)
Total animals with malignant neoplasms Total malignant neoplasms Total animals with metastatic neoplasms	26 9	$\frac{16}{4}$	25 6	25 5

Number of animals examined microscopically at the site and the number of animals with neoplasm Number of animals with any tissue examined microscopically Primary neoplasms: all neoplasms except metastatic neoplasms

X: Lesion present Blank: Not examined

TABLE C2 Individual Animal Tumor Pathology of Male Mice in the 2-Year Inhalation Study of Ethylbenzene: Chamber Control

Number of Days on Study	0 2	0 2	2	2 8	4 6	5 1	5 2	5 4		5 6	5 8	5 8				6 1	3	6 4	6 5	6 9	7 2	7 2	7 2	7 2	7 2	
	4	9	2	9	7	9	2	4	5	8	4	5	7	0	0	6	9	2	2	8	3	5	9	9	9	
	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	
Carcass ID Number	2	4	3	1	1	4	1	4	4	4	3	2	2	1	4	1	4	3	3	0	0	0	0	0		
Curcuss ID Ivaliable	9	9	1	8	2	7	1	8							1	5		9	6	2	5		3	6		
Alimontony System																										
Alimentary System Esophagus						A								+	+	+										
Gallbladder	Δ	Δ	Δ												M		+	+	+	+	Α	Δ	+	+	T	
Intestine large, colon	+					A		+			+				+			+	+	+	+	+	+	+	+	
Intestine large, rectum	Δ	+	+		+	+	+	+	+	+	+	+			À	+	+	+	+	+	· +	·	+	_	<u>'</u>	
Intestine large, rectum	A	À	À		+	À	+	+	+	+	+	+		À	+	+	+	+	À	+	À	+	+	+	+	
Intestine small, duodenum			A		+	A	+				+	+		+	+	+	+		A		Δ		+	_	<u>'</u>	
Intestine small, jejunum	Δ		+	+	+	A	+	+	+	+	+	+	+		À		+	+	A	+			+	_	<u>'</u>	
Intestine small, ileum	A	+	À		+	A	+	+	+	+	+	+		À			M		A	+	À	+	+	+	+	
Liver	+	+	+	+	+	+	+	+	+	+		+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Fibrosarcoma, metastatic, stomach, glandular													X					Ċ		Ċ						
Hemangioma													11													
Hepatocellular carcinoma				X		X			X	X			X	X	X	X	X	X		X		X			X	
Hepatocellular adenoma							X								X						X				••	
Hepatocellular adenoma, multiple																							X			
Hepatocholangiocarcinoma					X																					
Mesentery																		+								
Hepatocellular carcinoma, metastatic, liver																		X								
Pancreas	+	+	+	+	+	Α	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Salivary glands	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Stomach, forestomach	+	+	Α	+	+	Α	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Fibrosarcoma, metastatic, stomach, glandular													X													
Stomach, glandular	+	+	Α	+	+	Α	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Fibrosarcoma													X													
Cardiovascular System																										
Blood vessel	+	M	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	M	+	+	+	+	+	+	+	
Aorta, fibrosarcoma, metastatic,																										
stomach, glandular													X													
Aorta, hepatocellular carcinoma,																										
metastatic, liver				X																						
Heart	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Fibrosarcoma, metastatic, stomach, glandular													X													
Pericardium, hepatocholangiocarcinoma,																										
metastatic, liver					X																					
Endocrine System																										
Adrenal cortex	+	+	+	+	+	M	М	+	+	М	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Adenoma	,-					141	141			141		•											'			
Hepatocellular carcinoma, metastatic, liver																		X								
Adrenal medulla	+	+	+	+	+	M	М	+	+	М	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Islets, pancreatic	+	+	+	+	+	A		+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Parathyroid gland		М	+	+	+							+		+	+	М	М			М	М	М	+	+	M	
Pituitary gland	+	+	+	+	+				M							+		+	+	+	+	+	+	+	+	
Thyroid gland	+	+	+	+	+				+									+	+	+	+	+	+	+	+	
		'			,							- 1											'	,		
Follicular cell, adenoma																										

^{+:} Tissue examined microscopically A: Autolysis precludes examination

M: Missing tissue I: Insufficient tissue

TABLE C2
Individual Animal Tumor Pathology of Male Mice in the 2-Year Inhalation Study of Ethylbenzene: Chamber Control

(continued) **Number of Days on Study** 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 Total **Carcass ID Number** 1 2 3 4 4 0 0 Tissues/ Tumors **Alimentary System** Esophagus M +Gallbladder Intestine large, colon Intestine large, rectum Intestine large, cecum Intestine small, duodenum Intestine small, jejunum Intestine small, ileum Liver Fibrosarcoma, metastatic, stomach, glandular Hemangioma Hepatocellular carcinoma X X XX X Hepatocellular adenoma X X X Hepatocellular adenoma, multiple Hepatocholangiocarcinoma Mesentery Hepatocellular carcinoma, metastatic, liver Pancreas Salivary glands Stomach, forestomach Fibrosarcoma, metastatic, stomach, glandular Stomach, glandular Fibrosarcoma Cardiovascular System Blood vessel Aorta, fibrosarcoma, metastatic, stomach, glandular Aorta, hepatocellular carcinoma, metastatic, liver Fibrosarcoma, metastatic, stomach, glandular Pericardium, hepatocholangiocarcinoma, metastatic, liver **Endocrine System** Adrenal cortex Adenoma Hepatocellular carcinoma, metastatic, liver Adrenal medulla Islets, pancreatic Parathyroid gland MM +M M M M M M MMMMMMM Pituitary gland M + Thyroid gland XX Follicular cell, adenoma X

Individual Animal Tumor Pathology of (continued)	Male	M	ice	in	th	e 2	-Ye	ear	· In	ıha	lat	ioı	n S	tud	ly (of 1	Eth	ıyl	bei	ıze	ne	: (Cha	am	ber C	ontrol
	0			2	4		5	5					5					6	6	6	7	7				
Number of Days on Study	2 4	2 9	0 2	8 9	6 7	1 9	2 2	4	5 5	6 8		8 5	8 7	1 0	1 0	1 6	3 9	4 2	5 2	9 8	2 3	2 5	2 9	2 9	2 9	
	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	
Carcass ID Number	2 9	4 9	3 1	1 8	1 2	4 7	1 1	4 8	4 6	4 4	3 0	2 4	2 8	1 4	4 1	1 5	4 3	3 9	3 6	0 2	0 5	0 8	0 3	0 6		
General Body System																										
Гissue NOS Fibrosarcoma					+								+ X													
Thoracic, hepatocholangiocarcinoma, metastatic, liver					X								21													
Genital System																										
Epididymis Ponia	+	M	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Penis Preputial gland	+	М	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	Α	+	+	+	+	+	+	
Prostate	+			+	+		+	+	+	+	+	+	+	+	+	M		M		+	+	+	+	+	+	
Seminal vesicle	+		+		+		+		+	+	+	+	+	+		+		+		+	+	+	+	+	+	
Гestes	+	M	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Interstitial cell, adenoma																			X							
Hematopoietic System																										
Bone marrow	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Lymph node			+				+					+	+													
Fibrosarcoma, metastatic, stomach, glandular Renal, fibrosarcoma, metastatic,													X													
stomach, glandular	1.4			M	N			ъ.	N	N	ъſ	ъ.	X +	1.1	ъſ	N/I		ъ.	1.1	1.1			N		M	
Lymph node, bronchial Fibrosarcoma, metastatic, stomach, glandular	IVI	+	+	IVI	IVI	+	+	IVI	IVI	IVI	IVI	IVI	+ V	IVI	IVI	IVI	+	IVI	IVI	IVI	+	+	IVI	+	IVI	
Lymph node, mandibular	_	_	_	_	_	_	_	_	_	_	М	_	+	_	М	_	_	М	_	М	_	_	_	_	_	
Sarcoma, metastatic, nose	'	'						x			141		'		141			111	'	171	'					
Lymph node, mesenteric	+	+	+	+	+	M	+	+	+	+	+	+	+	+	+	+	+	+	+	M	+	+	+	M	+	
Lymph node, mediastinal	+												+													
Fibrosarcoma, metastatic, stomach, glandular													X													
Hepatocholangiocarcinoma, metastatic, liver					X																					
Spleen	+	+	+										+													
Thymus Hepatocholangiocarcinoma, metastatic, liver	+	+	+	+	+ X	M	M	+	+	M	+	+	M	+	M	M	+	M	+	M	M	ı M	+	+	+	
Integumentary System Mammary gland	M	1\1	М	М	М	_	М	М	М	М	М	M	M	М	М	М	М	M	1/I	М	1/1	I M	Ŋſ	Ŋſ	М	
Skin													+													
Fibrosarcoma				·		•	•	•	Ċ				·			•			Ċ					·		
Musculoskeletal System																										
Bone	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Sternum, fibrosarcoma, metastatic,						•													·				·			
stomach, glandular													X													
Skeletal muscle													+					+								
Fibrosarcoma, metastatic, stomach, glandular													X					* *								
Hepatocellular carcinoma, metastatic, liver																		X								

V 1 6D 6.1	7	7	7	7	7	7			7			7		7		7			7		7		7	7		
Number of Days on Study	9	9	2 9	9	2 9	9	2 9	2 9	2 9	2 9	2 9	2 9	2 9	2 9	2 9	3 0	3 0	3 0	3 0	3 0	3 0	3 0	3 0	3 0	3 0	
Carcass ID Number	0 1 3	1	0 1 7	2	0 2 2	0 2 3	2	2	0 3 4	0 3 5	3	0 3 8	4	0 4 5	0 5 0	0 0 1	0	0 0 7	0 0 9	1	0 2 0		0 3 2	0 3 3	0 4 2	Total Tissues/ Tumors
General Body System Tissue NOS Fibrosarcoma Thoracic, hepatocholangiocarcinoma, metastatic, liver																										2 1
Genital System																										40
Epididymis Penis	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	49 1
Preputial gland	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	48
Prostate	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	M	+	+	+	+	+	46
Seminal vesicle	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	49
Testes Interstitial cell, adenoma	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	49 1
Hematopoietic System																										
Bone marrow	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Lymph node Fibrosarcoma, metastatic, stomach, glandular																										4
Renal, fibrosarcoma, metastatic,																										1
stomach, glandular																										1
Lymph node, bronchial	+	+	M	M	M	+	M	M	M	M	M	M	M	M	M	M	M	M	M	+	M	+	M	M	M	14
Fibrosarcoma, metastatic, stomach, glandular																										1
Lymph node, mandibular	M	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	M	+	+	M	+	+	+	+	43
Sarcoma, metastatic, nose			1.4											N.f												1
Lymph node, mesenteric Lymph node, mediastinal					+																				+	45 24
Fibrosarcoma, metastatic, stomach, glandular	171	. 141		141	'	141	141		141			141			141		141	141		141		141				1
Hepatocholangiocarcinoma, metastatic, liver																										1
Spleen	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Thymus Hepatocholangiocarcinoma, metastatic, liver	+	+	+	+	+	M	+	+	+	+	+	+	+	M	+	+	+	+	M	+	+	+	+	+	+	37 1
Integumentary System																										
Mammary gland	M	M	M		M	M	M	M	M	M	+	M	M	+	M	M	M	M	M	M	M	M	M	M	M	3
Skin Fibrosarcoma	+	+	+	+ X	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50 1
Musculoskeletal System																										_
Bone	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Sternum, fibrosarcoma, metastatic, stomach, glandular																										1
Skeletal muscle																										1 2
Fibrosarcoma, metastatic, stomach, glandular																										1
Hepatocellular carcinoma, metastatic, liver																										1

Individual Animal Tumor Pathology of (continued)	Male	M	ice	in	th	e 2	2-Y	eai	r Iı	nha	ala	tio	n S	tuo	ły (of 1	Etł	ıyl	bei	ıze	ne	: (Cha	am	ber	Control
Number of Days on Study	0 2	0 2	2	2	4	5 1	5 2	5 4	5 5	5 6	5 8	5 8	5 8	6	6	6	6	6	6 5	6	7	7	7	7	7	
compet of Buys on Study	4	9	2	9	7	9	2		5	8		5	7	0	0	6	9	2	2	8	3	5	9	9	9	
	0	0	0	0	0	0	0	0	0	0	0	0		0	0	0	0	0	0	0	0	0	0	0	0	
Carcass ID Number	2 9	4 9	3 1	1 8	1 2	4 7	1	4 8	4 6	4	3 0	2 4	2 8	1 4	4 1	1 5	4 3	3 9	3 6	0 2	0 5	0 8		0 6		
Nervous System Brain	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Respiratory System																										
Larynx	+	+	+	+	+	+		+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Lung Alveolar/bronchiolar adenoma	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+		+ X			+	+	+	+	+	+	
Alveolar/bronchiolar carcinoma														X		Λ	Λ	Λ								
Fibrosarcoma, metastatic, stomach, glandular													Х													
Hepatocellular carcinoma, metastatic, liver				X		X			X									Х		Х						
Hepatocholangiocarcinoma, metastatic, liver					X																					
Nose	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Sarcoma								X																		
Trachea	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Special Senses System Harderian gland Adenoma																										
Urinary System																										
Kidney Fibrosarcoma, metastatic, stomach, glandular Hepatocellular carcinoma, metastatic, liver	+	+	+	+	+	+	+	+	+	+	+	+	+ X	+	+	+ X	+	+	+	+	+	+	+	+	+	
Urinary bladder	+	M	+	+	+	A	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Systemic Lesions																										
Multiple organs Lymphoma malignant	+	+	+	+	+	+	+ X	+	+	+	+	+ X	+	+	+	+	+	+	+	+	+	+	+	+	+	

I ABLE CA					
Individual	Animal Tumor Pathology	of Male Mice in the 2-Year	Inhalation Study of	f Ethylbenzene:	Chamber Control
(continued)					

(continued)																										
Number of Days on Study	7 2 9	7 3 0																								
Carcass ID Number	0 1 3	0 1 6	0 1 7	0 2 1	0 2 2	0 2 3	0 2 6	0 2 7	0 3 4	0 3 5	0 3 7	0 3 8	0 4 0	0 4 5	0 5 0	0 0 1	0 0 4	0 0 7	0 0 9	0 1 9	0 2 0	0 2 5	0 3 2	0 3 3	0 4 2	Total Tissues/ Tumors
Nervous System Brain	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Respiratory System Larynx Lung Alveolar/bronchiolar adenoma Alveolar/bronchiolar carcinoma Fibrosarcoma, metastatic, stomach, glandular Hepatocellular carcinoma, metastatic, liver Hepatocholangiocarcinoma, metastatic, liver Nose	+ +	+ +	+ +	+ +	+ + X	+ +	+ +	+ +	+ +	+ + X	+ + X	+ +	+ +	M +	+ +	+ +	+ +	+ +	+ +	+ +	+ +	+ +	+ +	+ +	+ +	48 50 5 2 1 5 1
Sarcoma Trachea	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	1 50
Special Senses System Harderian gland Adenoma	+ X																+									2 1
Urinary System Kidney Fibrosarcoma, metastatic, stomach, glandular Hepatocellular carcinoma, metastatic, liver Urinary bladder	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50 1 1 48
Systemic Lesions Multiple organs Lymphoma malignant	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50 2

Individual Animal Tumor Pathology															_			_								_
	3	3	3 4	5	5		5	5	5	6	6	6	7	7	7	7	7	7	7	7	7	7	7	7	7	
Number of Days on Study	6	8			3		7		9		4	7	0	2	2		2	2	2	2	2	2	2	2	2	
	0	() 5	4	1		8	0	4	0	1	7	9	5	9	9	9	9	9	9	9	9	9	9	9	
	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	
Carcass ID Number	1	3	3 2	1	. 0	2	4	3	1	1	3	0	2	1	0	0	0	0	0	0	1	2	2	2	2	
	3	() 9	2	9	0	8	4	9	0	6	1	4	1	2	3	4	6	7	8	5	1	5	7	8	
Alimentary System																										
Esophagus	+	+	+ +		+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Gallbladder	N.	1	+ +		V I +				M						M	+	+	+	+	+	+	+	+	+	+	
Intestine large, colon	+	+	+ +		+ +											+	+	+	+	+	+	+	+	+	+	
Intestine large, rectum	+	+	+ +		A +				+					+			+	+	+	+	+	+	+	+	+	
Intestine large, cecum	+	+	+ +		⊦ A												+	+	+	+	+	+	+	+	+	
Intestine small, duodenum	+	+	+ +				M										+	+	+	+	+	+	+	+	+	
Intestine small, jejunum	+	+	+ +	- <i>F</i>	A A	. +	A	+	+	+	A	+	+		+	+	+	+	+	+	+	+	+	+	+	
Epithelium, carcinoma					,									X												
Intestine small, ileum	A		+ +		+ A		Α	+	+					+			+	+	+	+	+	+	+	+	+	
Liver	+	+	+ +		+ +	+	+	+	+	+	+		+	+	+	+	+	+	+	+	+	+	+	+	+	
Cholangiocarcinoma												X														
Hemangiosarcoma																										
Hepatoblastoma				,	7	1,		v			v															
Hepatocellular carcinoma				2	1	X		X			X															
Hepatocellular carcinoma, multiple	v		χ	7					X						v	v					v	v			X	
Hepatocellular adenoma	X		Χ						X		X				X	Λ		X			X	X			Λ	
Hepatocellular adenoma, multiple Pancreas					,						X +	,	,	,				X +	,							
rancreas Salivary glands	+	-	+ + ∟ '	_	r +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Sanvary gianus Stomach, forestomach	+	7	r + L J		- + - J		+	+	+	+	+	+	+	+	T +	+	+	T +	+	+	+	+	+	+	+	
Squamous cell papilloma	т		гт		г т		_	т	т	_	т	т	т	т	т	_	_	т	_	_	_	X		_	т	
Stomach, glandular	+	+	+ +		+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+		+	+	
Cardiavasaular System																										_
Cardiovascular System	1.	r .																								
Blood vessel	N.	1 -	+ +		+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Heart	+	-	+ +	_	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Endocrine System																										
Adrenal cortex	+	+			M N						+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Adrenal medulla	+	+	+ +		M N												+	+	+	+	+	+	+	+	+	
Islets, pancreatic	+	4	+ +				+						+		+		+	+	+	+	+	+	+	+	+	
Parathyroid gland	+	7	_ `	Λ -		+			M											M	+	+	+	+	+	
Pituitary gland	+	ľ	M +						M								+	+	+	+	+	+	+	+	+	
Thyroid gland	+	+	+ +		+ +	+	+	+	+		+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Follicular cell, adenoma										X																
General Body System Tissue NOS																										
Genital System																										
Epididymis	+	4	+ +		+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Leiomyoma						•	•	•	•	•	•		•	•		•		X		•	•		•	•	•	
Penis					+																					
Preputial gland	+	4	+ +		· + +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	M	+	+	+	+	+	
Prostate	+	4	+ +		+ +	+	+	+	+	+	+	+	+	+	+	+	+	M	+	+	+	+	+	+	+	
Seminal vesicle	+	4	+ +		+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Testes																										

FABLE C2 Individual Animal Tumor Patholog	y of Male	e N	1ic	e in	th	e 2	Y - Y -	ear	· In	ha	lat	ion	ı St	ud	y o	f E	cth;	yll	en	ıze	ne:	7	′5 բ	pr	n (c	ontinued)
	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	
Number of Days on Study	2	2		2	2	2	2	2	3	3	3	3	3	3	-	-	3	3	3	3	3	3	3	3	3	
value of Days on Stady	9	9		9	9	9	9	9	0	0	0	-							0	0	0	0	0	0		
	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	Total
Carcass ID Number	3	3	3	3	4	4	4	4	0	1	1	1	1	2	2	2	3	3	3	4	4	4	4	4	5	Tissues/
	2			9	3	4	6	9	5	4	6		8						5	0	1		5	7	0	Tumors
Alimentary System																										
Esophagus	+	+	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Gallbladder	+	+	- +	+	M	+	+	+	+	+	+	+	+	+	+	M	M	+	+	+	+	+	+	+	+	39
ntestine large, colon	+	+	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	48
Intestine large, rectum	+	+	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	49
ntestine large, cecum	+	+	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	46
ntestine small, duodenum		+	. +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	48
Intestine small, jejunum		+	. +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	46
Epithelium, carcinoma		,			'				,														'			1
Intestine small, ileum	_	_1		4	_	4	_	+	+	+	+	+	+	+	+	+	+	+	+	_	+	_	_	4	+	46
Liver		-T						_	_	_	_	_	_	+	· +	· +	_	+	_						+	50
Cholangiocarcinoma	+	+	Т	_	т	-	-	-	7	7	-	7	7	т′	т	Т	т	т,	-	-	т	-	т	-	7	1
Hemangiosarcoma	Х																									1
Hepatoblastoma	Λ																			X						1
					X							X								Λ	X			X		
Hepatocellular carcinoma					Λ							Λ			37						Λ			Λ		8
Hepatocellular carcinoma, multiple							3.7						3.7		X			37		37						1
Hepatocellular adenoma							X						X					X		X						12
Hepatocellular adenoma, multiple									X																X	4
Pancreas	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Salivary glands	+	+	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Stomach, forestomach	+	+	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Squamous cell papilloma																										1
Stomach, glandular	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Cardiovascular System																										
Blood vessel	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	M	+	+	+	+	+	+	48
Heart	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Endocrine System																										
Adrenal cortex	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	47
Adrenal medulla	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	46
slets, pancreatic	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Parathyroid gland	+	+	- N	I M	+	+	+	+	M	M	+	+	M	+	M	M	+	+	+	+	+	+	+	+	M	34
Pituitary gland	+	+	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	M	+	+	+	+	+	+	+	+	45
Гhyroid gland	+	+	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Follicular cell, adenoma					·	•		•		-	•		•					•		•	•		•	•	X	2
General Body System Fissue NOS		+	-			+																			+	3
Conital System																										
Genital System																										F0
Epididymis	+	+	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Leiomyoma																										1
Penis																			+							2
Preputial gland	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	49
Prostate	+	+	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	49
Seminal vesicle	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Testes																										50

																										ed)
	3	3	4	5	5	5	5	5	5	6	6	6	7	7	7	7	7	7	7	7	7	7	7	7	7	
Number of Days on Study	6	8					7	8	9	4	4	7	0	2	2	2	2	2	2	2	2	2	2	2	2	
	0	0	5	4	1	6	8	0	4	0	1	7	9	5	9	9	9	9	9	9	9	9	9	9	9	
	1	1	1	1	. 1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	
Carcass ID Number	1	3	2	1	. 0	2	4	3	1	1	3	0	2	1	0	0	0	0	0	0	1	2	2	2	2	
	3	0	9	2	9	0	8	4	9	0	6	1	4	1	2	3	4	6	7	8	5	1	5	7	8	
Hematopoietic System																										
Bone marrow	+	4	- +		+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Lymph node					+		+					+		+		+					+				+	
Renal, cholangiocarcinoma,																										
metastatic, liver												X														
Lymph node, bronchial	M	[-	- +		+ +	+	+	+	+	M	M		+	+	M	M	M	M	+	+	M	+	+	M	M	
Lymph node, mandibular	+				+ +																			+	+	
Lymph node, mesenteric					- N																				+	
Lymph node, mediastinal		[-			- N											+										
Spleen	+				+ +													+			+		+	+		
Гhymus	M				M N																+		+	+	+	
Integumentary System																										_
	1. /	Γ λ	<i>1</i> .	1	Л +	λ.	[], /	1.1	Ŋ./f	Ŋ.Æ	Ŋſ	Ŋ./	Ŋſ	Ŋſ	ŊΛ	ŊЛ	Ŋ.f	Ŋ.f	ŊЛ	Ŋ.	1, 1	1,1	1.1	1,1	М	
Mammary gland																										
Skin	+		- +	_	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	_
Musculoskeletal System																										
Bone	+	+	- +		+ +	+	+	+	+	+	+	+	M	+	+	+	+	+	+	+	+	+	+	+	+	
Nervous System																										
Brain	+	4	- +		+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Respiratory System																										
		1	1 +																							
Larynx	+		/1 + - +		- + - +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Lung Alveolar/bronchiolar adenoma	+	4	- +	_	X		+	+	+	+	+	+ V	+ X	+	+	+	+	+	+	+	+	+ X	+	+	+	
Alveolar/bronchiolar adenoma, multiple					Λ							Λ	Λ							v		Λ				
															v					X						
Alveolar/bronchiolar carcinoma												X			X											
Cholangiocarcinoma, metastatic, liver								X			X	Λ														
Hepatocellular carcinoma, metastatic, liver Bronchiole, polyp adenomatous								Λ			Λ														X	
Nose					,					,			,						,						Λ +	
Nose Trachea	+	٦.	- +	_	- + _ ,	+	+	+	+	+	+	+	+	+	+	+	_	_	+	+	+	+	+	+	+	
HULICHU		_						-	-	-	-	-	7	-	т*	7'	7"	7"	7	-	т_	т_	т	т	Г	
Special Senses System																										
Harderian gland													+													
Adenoma													X													
Urinary System																										
Kidney	J				ر ا		_			_	_		_	_	_	_	_	_	_	_		_	_	_	_	
Cholangiocarcinoma, metastatic, liver	+	٦	+	_			Т	-T	-	-	Т	X	7	7	т.	т.	7-	7-	7	-	т	т	-	т	Г	
Hepatocellular carcinoma, metastatic, liver											X	Λ														
Urinary bladder	+	4	- +		+ +	+	+	+	+	+		+	+	+	+	+	+	+	+	+	+	+	+	+	+	
•																										_
Systemic Lesions										,			,													
Multiple organs Lymphoma malignant	+	+	- +	 }		+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+ X	+	+	+	+	
LVIIIDIOMA MAIIONANI				,																	×					

7	7	7	7	7	7	7	7	7	7	7	7	7	7	•		7	7	7	7	7	7	7	7	7	
2 9								-	-					-					3 0						
																									1
																									Total
2																							-		Tissues/ Tumors
+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
																									7
																									1
M	M	+	M																M		M	+	+	+	24
+	+	+	+																+	-	+	+	+	+	45 46
+	+	+	+																			+			25
+	+	+	+	+	+	+															+	+	+	+	50
+	+	+	+	+	+	+															+	+	+	+	37
M	M	M	M	M	M	M	M	M	M	M	M	M	M	M	M	M	M	M	M	M	M	M	M	M	2
+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	49
+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	49
+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
				X			X											X						X	8
																									1
																									1
				v																					1 3
				Λ																					1
+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
		+					+																		3
		X					X																		3
+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
																									1
																									1
+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
+	_	- 1	- 1	- 1																					50
	1 3 2 + M + + + + + + M M	M M M + + + + + + + + + + + + + + + + +	M M + + + + + + + + + + + + + + + + + +	2 2 2 2 9 9 9 9 9 1 1 1 1 1 3 3 3 3 3 2 7 8 9	2 2 2 2 2 2 9 9 9 9 9 9 1 1 1 1 1 1 1 3 3 3 3 3 4 2 7 8 9 3	2 2 2 2 2 2 2 9 9 9 9 9 9 9 9 9 9 9 9 9	2 2 2 2 2 2 2 2 9 9 9 9 9 9 9 9 9 9 9 9	2 2 2 2 2 2 2 2 2 2 9 9 9 9 9 9 9 9 9 9	2 2 2 2 2 2 2 2 2 2 3 9 9 9 9 9 9 9 9 9	2 2 2 2 2 2 2 2 3 3 3 9 9 9 9 9 9 9 0 0 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 3 3 3 3	2 2 2 2 2 2 2 2 3 3 3 3 9 9 9 9 9 9 9 9	2 2 2 2 2 2 2 2 2 3 3 3 3 3 3 9 9 9 9 9	2 2 2 2 2 2 2 2 2 2 3 3 3 3 3 3 3 9 9 9 9	2 2 2 2 2 2 2 2 2 2 3 3 3 3 3 3 3 3 3 3	2 2 2 2 2 2 2 2 2 2 2 3 3 3 3 3 3 3 3 3	2 2 2 2 2 2 2 2 2 2 2 2 3 3 3 3 3 3 3 3	2 2 2 2 2 2 2 2 2 2 2 2 3 3 3 3 3 3 3 3	2 2 2 2 2 2 2 2 2 2 2 3 3 3 3 3 3 3 3 3	2 2 2 2 2 2 2 2 2 2 2 3 3 3 3 3 3 3 3 3	2 2 2 2 2 2 2 2 2 2 2 3 3 3 3 3 3 3 3 3	2 2 2 2 2 2 2 2 2 2 3 3 3 3 3 3 3 3 3 3	2 2 2 2 2 2 2 2 2 2 2 3 3 3 3 3 3 3 3 3	2 2 2 2 2 2 2 2 2 2 2 3 3 3 3 3 3 3 3 3	2 2 2 2 2 2 2 2 2 2 2 3 3 3 3 3 3 3 3 3	2 2 2 2 2 2 2 2 2 2 2 3 3 3 3 3 3 3 3 3

	4	E	r	E	E	E	E	E	c	e	e	e	e	e	e	7	7	7	~	~	~	~	~	7	7	
N		5		5	5	5	5						6					7	7	1	7	1	7		7	
Number of Days on Study	8	2 4	6 3	7 0	7 4	7 9									-		2	2 4	2 9	2 9	2 9	9	9	2 9	2 9	
		0	9		0	9																9	9	0	0	_
Carcass ID Number	2 2	2	2	2	2	2	2 4	2		2			2					2	2	2	2	2	2	2	2	
ourous 12 Munior	2	0		4				5								0								6		
Alimentary System																										_
Esophagus	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Gallbladder	A	Α	M	+	A	A	M	M	+	Α	Α	+	M	+	M	+	+	+	M	+	+	+	+	+	M	
Intestine large, colon	+	+	+	+	+	+	+	A	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Intestine large, rectum	+	+	+	+	+	A	+	A	+	A	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Intestine large, cecum	+	+	+	+	+			A									+	A	+	+	+	+	+	+	+	
Intestine small, duodenum	+	Α	+	+	+			A								+	+	A	+	+	+	+	+	+	+	
Intestine small, jejunum	+	A	A	+	+			+										A	+	+	+	+	+	+	+	
Intestine small, ileum	+	+	+	+				+											+	+	+	+	+	+	+	
Liver	+	+	+	+	+		+	+	+					+			+	+	+	+	+	+	+	+	+	
Hepatocellular carcinoma	X		X	X	X	X		v			X	X	X			X										
Hepatocellular carcinoma, multiple								X						v	v		v		v	v				v		
Hepatocellular adenoma							X							X	Λ		X		X	Λ				X		
Hepatocholangiocarcinoma Mesentery																										
Hepatocholangiocarcinoma, metastatic, liver																										
Pancreas	_	+	+	+	+	А	+	+	+	М	+	+	+	+	+	+	+	+	+	+	+	+	_	+	+	
Acinus, hepatocholangiocarcinoma,	7	-	-			. 1	1	1		171	1		•					1	1.	r	-	٢		٢	•	
metastatic, liver																										
Salivary glands	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Stomach, forestomach	+	+	+	+	+	+	+	+					+	+	+	+	+	+	+	+	+	+	+	+	+	
Stomach, glandular	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Tooth								_															+			
Cardiovascular System																										
Blood vessel	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Heart	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Hepatocholangiocarcinoma, metastatic, liver																										
Endocrine System																										
Adrenal cortex	+	+	+	Μ	+	+	+	+	+	+	+	+	М	+	+	+	+	+	+	+	+	+	+	+	+	
Adenoma		•	•			•		•		•	-	•					•		-		•	·	•	•	•	
Carcinoma															X											
Adrenal medulla	+	+	+	M	+	+	+	+	+	+	+	+	M			+	+	+	+	+	+	+	+	+	+	
Islets, pancreatic	+							+											+	+	+	+	+	+	+	
Parathyroid gland	M	M	M	+	M	M	+	M	M	M	+	M	+	M	M	+	M	M	+	M	+	+	M	+	+	
Pituitary gland	+			+					+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Pars distalis, carcinoma										X																
Thyroid gland	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Follicular cell, adenoma																										
General Body System																										
Tissue NOS																										
Fat, hepatocholangiocarcinoma,																										
metastatic, liver																										
Genital System																										_
Epididymis	+	_	_	_	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	_	_	_	_	_	+	
Preputial gland	M	+	+	+	М	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Prostate	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Seminal vesicle	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Hepatocholangiocarcinoma, metastatic, liver			•																							
Testes	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Interstitial cell, adenoma																										

	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	
Number of Days on Study	2	2	2	2	2	2	2	2	2	2	2	2	2	2	3	3	3	3	3	3	3	3	3	3	3	
	9	9	9	9	9	9	9	9	9	9	9	9	9	9			0	0	0	0	0	0		0		
	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	Total
Carcass ID Number	2	2	2	2	3	3	3	3	3	3	3	3	4	4	0	0	1	1	4	4	4	4	4	4	5	Tissues/
	3	6	7	9	1	2	3	4	6	7	8	9	3	7	6	8	3	7	0	2	4	6	8	9	0	Tumors
Alimentary System																										
Esophagus	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Gallbladder	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	37
Intestine large, colon	A	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	48
Intestine large, rectum	M	+	+	+	+	+	M	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	45
Intestine large, cecum	Α	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	44
ntestine small, duodenum	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	43
ntestine small, jejunum	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	44
ntestine small, ileum	Α	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	44
Liver	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Hepatocellular carcinoma																			X			X				11
Hepatocellular carcinoma, multiple											X															2
Hepatocellular adenoma			X	X			X			X				X	X	X	X						X			17
Hepatocholangiocarcinoma																		X								1
Mesentery																		+								1
Hepatocholangiocarcinoma, metastatic, liver																		X								1
Pancreas	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	48
Acinus, hepatocholangiocarcinoma,																		37								
metastatic, liver																		X								1
Salivary glands	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50 50
Stomach, forestomach Stomach, glandular	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50 50
Footh	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	1
10001																										1
Cardiovascular System																										
Blood vessel	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	M	+	+	+	+	+	+	49
Heart	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Hepatocholangiocarcinoma, metastatic, liver																		X								1
Endocrine System																										
Adrenal cortex	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	48
Adenoma															X											1
Carcinoma																										1
Adrenal medulla	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	48
slets, pancreatic	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+				48
Parathyroid gland	+	+	+	M	+	+	+	+	M	+	+	M	+	+	+	+	+	M	+	+	M	+	M	ĺ +	M	28
Pituitary gland	+	M	+	+	+	+	+	+	+	+	+	+	M	+	+	+	+	+	+	+	+	+	+	+	+	45
Pars distalis, carcinoma																										1
Гhyroid gland	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Follicular cell, adenoma												X														1
General Body System																										
Γissue NOS																		+								1
Fat, hepatocholangiocarcinoma,																										
metastatic, liver																		X								1
Genital System																							_	_		
					,	,	,	,	,	,	,	,						,	,							50
Epididymis Preputial gland	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	30 48
reputiai giand Prostate	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	48 50
Seminal vesicle	+	+	+	+	ر +		+		ر +				_	_	T	_	_	ر +		+	+	+	+	+	+	50
Hepatocholangiocarcinoma, metastatic, liver	+	+	+	+	+	+	+	+	+	+	+	+	_	_	_	_	_	X	+	+	+	+	+	+	+	1
rrepatoenotangiocaremonia, metastatie, mver																		Λ								
Γestes					.1	. 1	. 1		.1	. 1		. 1	-1	-1		J	_1	. 1	. 1	. 1					. 1	50

Individual Animal Tumor Pathology of	Male	M	lice	e in	th	e 2	2-Y	eai	r Iı	ıha	ılat	ioı	n S	tud	ly (of 1	Eth	yll	oen	ze	ne:	2	50	рĮ	m	(continued)
	4	5	5	5	5	5	5	5	6	6	6	6	6	6	6	7	7	7	7	7	7	7	7	7	7	
Number of Days on Study	8	2	6	7	7	7	7	8	0	4	8	8	8	9	9	0	2	2	2	2	2	2	2	2	2	
	0	4	3	0	4	9	9	8	2	2	0	3	8	1	6	5	2	4	9	9	9	9	9	9	9	
	2	2	2	2	2	2	2		2			2			2	2	2	2	2	2	2	2	2		2	
Carcass ID Number	2 2	3		2			4 1		1				1		0			0 9	0			1				
		U	0	_	-	0	_	0	_	-			0	-		0	J	0	~	_	<u> </u>		~	0	0	
Hematopoietic System Bone marrow	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Lymph node		+	Ċ	+		•		Ċ	+	•	+	+		+		+	+			Ċ	•		Ċ		Ċ	
Popliteal, hemangioma														X												
Renal, hepatocholangiocarcinoma, metastatic, liver																										
Lymph node, bronchial	M	N	[+	Μ	+	+	M	M	M	M	M	+	+	+	+	+	M	+	M	+	+	M	M	M	+	
Hepatocholangiocarcinoma, metastatic, liver																										
Lymph node, mandibular	+	+	+				+															M				
Lymph node, mesenteric	+	+	+	+	+	M	+	+	+	M	+	+	+	+	+	+	+	+	+	M	+	+	+	+	+	
Hepatocholangiocarcinoma, metastatic, liver	1.4		1.	r .	3.4	1.4	1.7	1.7			1.			1.4			1.4	,	,						1.1	
Lymph node, mediastinal Spleen							M +															+ M			M	
Thymus							+																			
Integumentary System																										
Mammary gland	М	+	M	[+	М	М	M	М	м	М	М	М	М	М	М	М	М	М	М	М	М	М	М	М	М	
Skin							+																			
Hemangioma		·	•	•		•	•	•	•	•	•	•	•		•	•		X			•	•	•	•		
Musculoskeletal System																										
Bone	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Skeletal muscle								+																		
Alveolar/bronchiolar carcinoma,																										
metastatic, lung								X																		
Hepatocholangiocarcinoma, metastatic, liver																										
Nervous System																										
Brain	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Respiratory System								_				_														
Larynx	+	+	+	+	+	+			+										+	+	+	+	+	+	+	
Lung Alveolar/bronchiolar adenoma	+	+	+	+	+	+	+	+	+	+	+	+	+	+ X	+	+	+ X		+	+	+	+	+	+	+	
Alveolar/bronchiolar adenoma, multiple									X					Λ			Λ	Λ								
Alveolar/bronchiolar carcinoma								X		X					X					X						
Hepatocellular carcinoma, metastatic, liver					X			X			X		X													
Hepatocholangiocarcinoma, metastatic, liver																										
Nose	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Trachea	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Special Senses System																										
Harderian gland																										
Adenoma																										
Urinary System																										
Kidney	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Alveolar/bronchiolar carcinoma,								* *																		
metastatic, lung								X																		
Renal tubule, adenoma Ureter																		,								
Oreter Urinary bladder	_	_	_	_	_	_	_	_		_		_	+	_	+	+	+	+ Δ	_	_	_	_	_	_	_	
Ormary bladuct		τ	$\overline{}$			- T		-T	- T		-T			-	-10	7		Γ								

TABLE C2 Individual Animal Tumor Pathology of	Male	M	[ic	e iı	ı tl	ıe 2	2-Y	eai	r Iı	ıha	ılat	tioi	n S	tuc	ly c	of 1	Eth	ıyll	ber	ıze	ne	: 2	250	рŗ	m	(continued)
Number of Days on Study	7 2 9	7 2 9	7 2 9	7 2 9	7 2 9	7 2 9	7 2 9	7 2 9	7 2 9	7 2 9	7 2 9	7 2 9	7 2 9	7 2 9	7 3 0	7 3 0	7 3 0	7 3 0	7 3 0	7 3 0	7 3 0	7 3 0	7 3 0	7 3 0	7 3 0	
Carcass ID Number	2 2 3	2 2 6	2 2	2 2	2 3	2 3	2 3	2 3	2 3 6	2 3 7	2 3	2 3	2 4	2 4 7	2 0	2 0	2 1 3	2	2 4	2 4	2 4 4	2 4 6	2 4 8	2 4 9	2 5 0	Total Tissues/ Tumors
Hematopoietic System Bone marrow Lymph node Popliteal, hemangioma	+	+	+	+	+	++	+	+	+	+	+	+	+	+	+	+	+	+++	+	+	+	+	+	+	+	50 11 1
Renal, hepatocholangiocarcinoma, metastatic, liver Lymph node, bronchial Hepatocholangiocarcinoma, metastatic, liver Lymph node, mandibular	M +	[M	Ι Ν +	1 N	1 +	+	+	M +	+	M +	M M		M +	+	+	M +	+	X + X +	+	+	M +	+	+	+	+	1 27 1 46
Lymph node, mesenteric Hepatocholangiocarcinoma, metastatic, liver Lymph node, mediastinal Spleen	+ +	+ + +	+ + +	+ N +	1 +	+ M +	+	+	+	+	+	+	+ + + +	+	+	+	+	+	+	+	+	+	+	+	+	47 1 27 49 39
Thymus Integumentary System Mammary gland Skin Hemangioma	M +	+ [M +	1 M	1 M	1 M	+ [+ +							+ M +													39 3 50 1
Musculoskeletal System Bone Skeletal muscle Alveolar/bronchiolar carcinoma,	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+++	+	+	+	+	+	+	+	50 2
metastatic, lung Hepatocholangiocarcinoma, metastatic, liver																		X								1 1
Nervous System Brain	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Respiratory System Larynx Lung Alveolar/bronchiolar adenoma Alveolar/bronchiolar adenoma, multiple	M +	[+ +	+		+ + X X	+ +	+	+ + X	+	+ +	+ + X	+	+	+	+++	+	+ + X	+	+	+	+	+	+	+	+ + X	46 50 9 1
Alveolar/bronchiolar carcinoma Hepatocellular carcinoma, metastatic, liver Hepatocholangiocarcinoma, metastatic, liver Nose Trachea	+	++++	+	+	+	++	++	++	+++	X + +	++	++	+++	+++	+++	++	++	X + +	X + +	+++	++	+ +	++	++	++	5 5 1 50 50
Special Senses System Harderian gland Adenoma														+ X					+ X							2 2
Urinary System Kidney	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Alveolar/bronchiolar carcinoma, metastatic, lung Renal tubule, adenoma Ureter																		X								1 1 2
Urinary bladder	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	49

TABLE C2 Individual Animal Tumor Patho	ology of Mal	e N	Лi	ce	in	th	e 2	2- Y	eai	r Iı	nha	ılat	tioı	ı S	tud	ly (of I	Etl	ıyl	bei	ıze	ne	: 2	250	рĮ	pm	(continued)
Number of Days on Study	4 8 (•	5 6 3	7	5 7 4	5 7 9	5 7 9		6 0 2	4	•	6 8 3	6 8 8	6 9 1	6 9 6					7 2 9	7 2 9	7 2 9	7 2 9	7 2 9	7 2 9	
Carcass ID Number	2 2 2	2 3	? })	2 4 5	2 2 4	2 3 5	2 0 3	4	2 2 5	1	2 1 8	2 2 8	2 2 1	2 1 5	2 1 9	2 0 1	1	0	2 0 9	2 0 2	2 0 4	2 0 7	2 1 1	2 1 2	2 1 6	2 2 0	
Systemic Lesions Multiple organs Leukemia granulocytic Lymphoma malignant	-		+ K	+	+	+	+	+	+	+	+	+ X	+	+	+	+	+	+	+ X	+	+	+	+	+	+	+	

Individual Animal Tumor Patho	logy of Mal	e N	Iic	e in	th	e 2	-Y	ear	· Iı	ıha	lat	ioi	ı S	tud	ly (of 1	Etł	ıyl	ber	ıze	ne	2	250	pp	m	(continued)
Number of Days on Study	7 2 9	7 3 0																								
Carcass ID Number	2 2 3	2 2 6	2 2 7	2 2 9	2 3 1	2 3 2	2 3 3	2 3 4	2 3 6	2 3 7	2 3 8	2 3 9	2 4 3	2 4 7	2 0 6	2 0 8	2 1 3	2 1 7	2 4 0	2 4 2	2 4 4	2 4 6	2 4 8	2 4 9	2 5 0	Total Tissues/ Tumors
Systemic Lesions Multiple organs Leukemia granulocytic Lymphoma malignant	+	+ X	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50 1 3

		2		4		5 5			6	6	6	6	6	6		6	6	7	7	7	7	7	7	7
Number of Days on Study	7	5	1	3		2 6			1	3	4	5	5	6	6	7	7	0	2	2	2	2	2	2
	9	9	8	0	0	0 5	2	7	8	2	7	9	9	0	9	9	9	8	5	9	9	9	9	9
	3	3	3	3	3	3 3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3
Carcass ID Number	2	1	3	2	3	2 (0	3	0	2	4	0	4	3	1	1	3	1	2	0	0	1	1	1
	6	5	8	4	9	0 6	7	4	5	8	6	1	7	7	4	9	3	0	9	2	4	2	3	6
llimentary System																								
Csophagus	+	+	+	+	+	+ -	- +	+	+	+	M	+	+	+	+	+	+	+	+	+	+	+	+	+
Gallbladder	+	Α	. A	Α	Α	Α -	- A	+	+	+	Α	+	+	M	+	+	Α	+	+	+	+	+	+	+
ntestine large, colon	+	+	+	+	Α	Α -	- A	+	+	+	Α	+	+	+	+	+	+	M	+	+	+	+	+	+
ntestine large, rectum	+	Α	+	+	+	Α -	- A	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
ntestine large, cecum	+	Α	M	Α	Α	Α -	- A	+	+	+	Α	+	+	+	+	+	+	+	+	+	+	+	+	+
ntestine small, duodenum	+	+				Α -					Α	Α	+	+	+	+	+	+	+	+	+	+	+	+
ntestine small, jejunum	+	+	M	Α	A	Α -	- A	+	Α	+	+	+	+	+	+	+	Α	+	+	+	+	+	+	+
Intestine small, ileum	+	A	M	Α	A	Α -	- A	+	Α	+	Α	+	+	+	+	+	M	+	+	+	+	+	+	+
Liver	+	+	+	+	+	+ -	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Alveolar/bronchiolar carcinoma,																								
metastatic, lung																		X						
Hepatocellular carcinoma				X	X	X		X				X	X			X		X	X				X	
Hepatocellular adenoma									X							X								X
Hepatocellular adenoma, multiple			X																					
Hepatocholangiocarcinoma Pancreas			Λ.			М -	۸																	
alicreas Salivary glands	+	+	+	+	+	IVI -	- A	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
tomach, forestomach	+	+	+	+	Ā	+ - A -	- +	+	+	+	+	+	+	+	+	_			+	+	+	+	+	+
tomach, glandular		+	+	+		A -			+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	T
Congue		Ċ					1	• '			'									•	•		Ċ	
ooth										+														
Odontoma										X														
N																								
Cardiovascular System																								
Blood vessel	+	+		+	+	+ -	- +	+	+	+	+	+	+	+	+	M	+	+	+	+	+	+	+	+
Aorta, sarcoma			X																					
Heart Alveolar/bronchiolar carcinoma,	+	+	+	+	+	+ -	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
metastatic, lung																		X						
Indooring System																								
Endocrine System Adrenal cortex	1.	1 +	_	_	_	_	_ ^		_			_	_	_	_	_	_	_	_	_	_	_	_	_
Adrenal medulla		1 +	+	+	+	+ -	- Δ	. +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	4
slets, pancreatic	+	+	+	+	+	M -	- A	. +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Adenoma					•		•		•		•	•	•		•				•	•	•	·	•	
Carcinoma															X									
Parathyroid gland	N	1 +	+	+	+	М -	- N	1 +	+	+	+	M	M	+		M	+	+	+	M	M	M	M	+
ituitary gland	+	+	+	M		М -		+					+			+			+	+	+			+
hyroid gland	+	+	+	+	+	+ -	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Follicular cell, adenoma											X		X											X
Follicular cell, adenoma, multiple																								
eneral Body System																								
Cissue NOS											+													
Thoracic, sarcoma			+ X								+													
THOTACIC, SALCOINA			Λ																					
enital System																								
pididymis	+	+	+	+	+	+ -	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
reputial gland	+	+	+	+	+	+ -	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Prostate	+	+	+	+	+	+ -	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Seminal vesicle	+	+	+	+	+	+ -	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Testes	+	+	+	+	+	+ -	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Interstitial cell, adenoma																								

TARLE C2

	7				7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	
Number of Days on Study	2 9				2 9	2 9	2 9	3 0																		
	3	_			3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	Total
Carcass ID Number	1				3	4	4	0	0	0	3 1	1	2	2	2	3	3	3	4	4	4	4	4	4	5	Tissues
Carcago ID Transpor	8					2	8	3	8		1	7	1	3	5	0	1	6	0	1	3	4	5	9	0	Tumors
Alimentary System																										
Esophagus	+	- 4	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	49
Gallbladder	+	- +	- N	1 +	+	+	+	+	M	+	+	+	+	+	+	+	+	+	+	+	+	M	+	+	+	38
Intestine large, colon	+	- +	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	45
Intestine large, rectum	+	- +	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	47
Intestine large, cecum	+	- +	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	43
Intestine small, duodenum	+	- +	+ +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	45
Intestine small, jejunum	+	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	43
Intestine small, ileum	+	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	41
Liver Alvoolar/bronchiolar carcinoma	+	- +	- +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Alveolar/bronchiolar carcinoma,																										1
metastatic, lung Hepatocellular carcinoma																										10
Hepatocellular adenoma		3	ΧX	· v		Y	X	Y	Y			v	Х						Y	Y	X			Y	X	17
Hepatocellular adenoma, multiple			1 /		X	Λ	Λ	Λ	Λ			Λ	Λ						Λ	Λ	Λ			Λ	Λ	17
Hepatocholangiocarcinoma					71																					1
Pancreas	4		⊢ +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	48
Salivary glands	-		' ' ⊦ +	. +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Stomach, forestomach	+	. 4	· ·		+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	47
Stomach, glandular	+		- +	. +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	47
Tongue											+															1
Tooth																										1
Odontoma																										1
Cardiovascular System																										
Blood vessel	+	- 4	+ +	- +	+	+	+	+	+	+	+	+	+	+	M	+	+	+	+	+	+	+	+	+	M	47
Aorta, sarcoma																										1
Heart	+	- 4	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Alveolar/bronchiolar carcinoma,																										
metastatic, lung																										1
Endocrine System																										
Adrenal cortex	+	- +	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	48
Adrenal medulla	+	- +	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	48
Islets, pancreatic	+	- +	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	48
Adenoma			χ	(1
Carcinoma						٠.,		٠.					٠,											٠.		1
Parathyroid gland	N	1 N	VI +	- IV	I M	M														+	M	M	+	M	l +	25
Pituitary gland	+	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	47
Thyroid gland	+	- +	+ +	- +	+	+ v	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Follicular cell, adenoma Follicular cell, adenoma, multiple				Х		X							X													5 1
																										1
General Body System																										0
Tissue NOS																										2
Thoracic, sarcoma																										1
Genital System																										
Epididymis	+	- +	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Preputial gland	+	- +	+ +	- +	+	+	+	+	+	+	+	+	M	+	+	+	+	+	+	+	+	+	+	+	+	49
Prostate	+	- +	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Seminal vesicle	+	- +		+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50 50
Testes Interstitial cell, adenoma	+	- +	- +	+	+	+	+	+	+	+ X	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50 1
micronnai cen, auchonia										Λ																1

	1	2	4	4	4	5	5	5	6	6	6	6	6	6	6	6	6	6	7	7	7	7	7	7	7
Number of Days on Study	7	5	1	3	8	2	6		0	1	3	4	5	5			7	7	0	2	2	2	2	9	2
Number of Days on Study	9	9	8	0		0	5	2	7	8	2		9	9		9	9	9	8	5	9	9	9	9	9
	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3
Carcass ID Number	2	1													3										
Carcass 12 Ivaniber	6														7										
Hematopoietic System																									
Bone marrow	_	+	_	_	_	_	_	_	_	_	_	_	_	_	_	+	_	_	_	_	_	_	_	_	_
Lymph node						'	+		'	+	'	'	'			+	'				'		'	'	'
Pancreatic, carcinoma							'			'						X									
Lymph node, bronchial	М	М	+	М	М	М	М	+	М	М	+	М	+	М	M		+	М	+	М	М	+	М	М	M
Alveolar/bronchiolar carcinoma,	111			141	171	141	171		171	171	ľ	171		171	.,,			111		171	141		171	111	111
metastatic, lung																			X						
Sarcoma			X																						
Lymph node, mandibular	М	+		+	+	М	+	М	+	+	+	+	+	+	+	+	+	M	+	+	+	+	+	+	+
Lymph node, mesenteric															+										
Lymph node, mediastinal															+										
Sarcoma	141	243	X	Ċ		.,,	•	.,,	.,,	.,,	. 7 2	.,,	.,,		•				.,,	.,,	.,,	. 71			
Spleen	+	+		+	+	М	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Thymus															M					+	+	+	+	+	M
Alveolar/bronchiolar carcinoma,	·			·						•							·	•	·		·			·	
metastatic, lung																			X						
Sarcoma			X																						
Internation Creton																							_	_	
Integumentary System				١.			١,		1 (1 (1 (1 (3.7	1 (1 (١.	1 (3.4	1 (M
Mammary gland	+														M										
Skin	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+			+
Musculoskeletal System																									
Bone	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Nervous System																									
Brain	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
D																									
Respiratory System	3.4					,	,		,	,		,	,							,	,				
Larynx		+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Lung	+	+	+ V	+	+ V	+	+	+	+	+	+	+	+	+	+	+	+	+ V	+	+	+	+		+	+
Alveolar/bronchiolar adenoma			X		X													X			X		X	X	
Alveolar/bronchiolar adenoma, multiple																			37			3.7			
Alveolar/bronchiolar carcinoma					37												v		X			X		37	
Hepatocellular carcinoma, metastatic, liver			37		X												X							X	
Hepatocholangiocarcinoma, metastatic, liver			X																						
Mediastinum, sarcoma			X																						
Nose	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Pleura																			+						
Alveolar/bronchiolar carcinoma,																			37						
metastatic, lung						,	,		,	,									X		,				
Trachea												+	+	-	-	+	+	+	+	+	-	+	+	+	+

	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	
Number of Days on Study	2	2	2	2	2	2	2	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	
	9	9	9	9	9	9	9	0	0		0	0	0	0		0	0	0	0	0	0	0	0	0		
	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	Total
Carcass ID Number	1	2	2	3	3	4	4	0	0	0	1	1	2	2	2	3	3	3	4	4	4	4	4	4	5	Tissues/
	8	2	7																				5			Tumors
Hematopoietic System																										
Bone marrow	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Lymph node																										3
Pancreatic, carcinoma																										1
Lymph node, bronchial	+	+	+	+	M	+	M	+	+	+	M	+	+	+	+	+	+	+	+	+	+	Μ	M	+	+	27
Alveolar/bronchiolar carcinoma,																										
metastatic, lung																										1
Sarcoma																										1
Lymph node, mandibular	+	+	+	+	+	+	+	+	+	M	+	+	+	+	+	+	+	+	+	+	+	M	+	+	+	44
Lymph node, mesenteric	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	48
Lymph node, mediastinal	M	M	[M	+	M	+	+	+	+	+	+	+	M	M	+	+						M	+	+	+	25
Sarcoma																										1
Spleen	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	49
Γhymus	+	+	+	+	+	M	M	M	+	+	M	+	+	+	+	+	+	M	+	+	+	+	+	+	+	34
Alveolar/bronchiolar carcinoma,																										
metastatic, lung																										1
Sarcoma																										1
Integumentary System																										
Mammary gland	M	M	М	М	М	М	М	М	М	+	М	М	М	М	М	М	М	М	М	М	М	М	M	М	М	3
Skin	+	+		+		+		+	+		+				+				+				+			50
	•	_		_	_	_	_	_		_	_	_			_	_	_	_	_		_	<u> </u>		<u> </u>		
Musculoskeletal System																										
Bone	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Nervous System																										
Brain	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Respiratory System																										
Larynx	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	49
Lung	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Alveolar/bronchiolar adenoma		X		X	X				X				X				X		X		X	X				15
Alveolar/bronchiolar adenoma, multiple																									X	1
Alveolar/bronchiolar carcinoma																X										3
Hepatocellular carcinoma, metastatic, liver																										3
Hepatocholangiocarcinoma, metastatic, liver																										1
Mediastinum, sarcoma																										1
Nose	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Pleura																										1
Alveolar/bronchiolar carcinoma,																										
metastatic, lung																										1
																										50

Individual Animal Tumor Patholog	y of Male	· IVI	ice	ın	tn	e z	- Y (ear	. III	na	uat	10I	1 51	tua	ly o	1	Etn	ıyı	bei	ıze	ene	: '	/50	p])m	(continued)
	1	2	4	4	4	5	5	5	6	6	6	6	6	6	6	6	6	6	7	7	7	7	7	7	7	
Number of Days on Study	7	5	1	3	8	2	6	8	0	1	3	4	5	5	6	6	7	7	0	2	2	2	2	2	2	
, ,	9	9	8	0	0	0	5	2	7	8	2	7	9	9	0	9	9	9	8	5	9	9	9	9	9	
	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	
Carcass ID Number	2	1	3	2	3	2	0	0	3	0	2	4	0	4	3	1	1	3	1	2	0	0	1	1	1	
artass 1D Number	6	5	8	4	9	0	6	7	4	5	8	6	1	7	7	4	9	3	0	9	2	4	2	3	6	
Urinary System																										
Kidney	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Alveolar/bronchiolar carcinoma,																										
metastatic, lung																			X							
Ureter										+																
Urinary bladder	+	+	+	+	+	Α	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Systemic Lesions																										
Multiple organs	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	
Leukemia granulocytic		X	'		,			'	'		'	'	'			•				'			'		'	
Lymphoma malignant		71																		X						

Individual Animal Tumor Patholog	gy of Mal	e N	Лiс	ce i	in t	he	2 -Y	/ea	r Iı	nha	alat	tio	n S	tud	ly o	of I	Etl	ıyl	bei	nze	ne	: 7	750	p j	рn	1 (continued)
	7	7	7	7 1	7 7	7 7	7 7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	1
Number of Days on Study	2 9	9	2 2	2 2	2 2	2 2	2 2	3 0	3 0	(3															
_	3	3	3	3 :	3 3	3 3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	9	B Total
Carcass ID Number	1 8	2	2 7	2 : 7 :	3 3 2 5	3 4 5 2	1 4 2 8	0 3	0 8	0 9	1 1	1 7	2 1	2	2 5	3 0	3 1	3 6	4 0	4 1	4 3	4	4 5	4 9	(
Urinary System Kidney Alveolar/bronchiolar carcinoma,	+	- 4	- -	+ -	+ -	+ -	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+		→ 50
metastatic, lung Ureter																										1
Urinary bladder	+	- 4		+ -	+ -	+ -	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+		+ 49
Systemic Lesions	_																									_
Multiple organs Leukemia granulocytic	+	- +	⊦ -	+ -	+ -	+ -	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+		+ 50 1
Lymphoma malignant															X											2

TABLE C3
Statistical Analysis of Primary Neoplasms in Male Mice in the 2-Year Inhalation Study of Ethylbenzene

	Chamber Control	75 ppm	250 ppm	750 ppm
Harderian Gland: Adenoma				
Overall rate ^a	1/50 (2%)	3/50 (6%)	2/50 (4%)	0/50 (0%)
Adjusted rate ^b	3.6%	8.0%	6.3%	0.0%
Геrminal rate ^с	1/28 (4%)	2/36 (6%)	2/32 (6%)	0/30 (0%)
First incidencę (days)	729 (T)	709	729 (T)	e
Life table test ^d	P = 0.192N	P = 0.398	P = 0.547	P = 0.486N
Logistic regression test ^d	P = 0.182N	P = 0.375	P = 0.547	P = 0.486N
Cochran-Armitage test ^d	P = 0.183N			
Fisher exact test ^a		P = 0.309	P = 0.500	P = 0.500N
Liver: Hepatocellular Adenoma				
Overall rate	12/50 (24%)	16/50 (32%)	17/50 (34%)	18/50 (36%)
Adjusted rate	37.6%	39.1%	46.7%	55.8%
Terminal rate	9/28 (32%)	12/36 (33%)	13/32 (41%)	16/30 (53%)
First incidence (days)	522	360	579	618
Life table test	P = 0.142	P = 0.503	P = 0.324	P = 0.186
Logistic regression test	P = 0.182	P = 0.322	P = 0.329	P = 0.189
Cochran-Armitage test	P = 0.178		D 0.400	D 0.400
Fisher exact test		P = 0.252	P = 0.189	P = 0.138
Liver: Hepatocellular Carcinoma				
Overall rate	17/50 (34%)	9/50 (18%)	13/50 (26%)	10/50 (20%)
Adjusted rate	41.1%	21.5%	28.5%	23.8%
Γerminal rate	5/28 (18%)	5/36 (14%)	3/32 (9%)	1/30 (3%)
First incidence (days)	289	514	480	430
Life table test	P = 0.228N	P = 0.032N	P = 0.170N	P = 0.083N
Logistic regression test	P = 0.196N	P = 0.064N	P = 0.321N	P = 0.091N
Cochran-Armitage test	P = 0.200N	D 0.055N	D 0.957N	D O OOON
Fisher exact test		P = 0.055N	P = 0.257N	P = 0.088N
Liver: Hepatocellular Adenoma or Carcinoma				
Overall rate	27/50 (54%)	24/50 (48%)	30/50 (60%)	27/50 (54%)
Adjusted rate	63.7%	54.9%	64.6%	66.7%
Ferminal rate	13/28 (46%)	17/36 (47%)	16/32 (50%)	17/30 (57%)
First incidence (days)	289	360	480	430
Life table test	P = 0.413	P = 0.127N	P = 0.505N	P= 0.433N
Logistic regression test Cochran-Armitage test	P= 0.465 P= 0.447	P = 0.321N	P = 0.389	P = 0.521N
Fisher exact test	r=0.447	P = 0.345N	P = 0.343	P = 0.579N
Liver: Hepatocellular Carcinoma or Hepatoblas Overall rate	toma 17/50 (34%)	10/50 (20%)	13/50 (26%)	10/50 (20%)
Adjusted rate	41.1%	24.1%	28.5%	23.8%
Terminal rate	5/28 (18%)	6/36 (17%)	3/32 (9%)	1/30 (3%)
First incidence (days)	289	514	480	430
Life table test	P = 0.202N	P = 0.049N	P = 0.170N	P = 0.083N
Logistic regression test	P = 0.167N	P = 0.101N	P = 0.321N	P = 0.0001V
Cochran-Armitage test	P = 0.173N			****=:
Fisher exact test		P = 0.088N	P = 0.257N	P = 0.088N

TABLE C3
Statistical Analysis of Primary Neoplasms in Male Mice in the 2-Year Inhalation Study of Ethylbenzene (continued)

	Chamber Control	75 ppm	250 ppm	75 0 ppm
Liver: Hepatocellular Adenoma, Hepatocellular Ca	rcinoma, or Hepatobl	astoma		
Overall rate	27/50 (54%)	24/50 (48%)	30/50 (60%)	27/50 (54%)
Adjusted rate	63.7%	54.9%	64.6%	66.7%
Terminal rate	13/28 (46%)	17/36 (47%)	16/32 (50%)	17/30 (57%)
First incidence (days)	289	360	480	430
Life table test	P = 0.413	P = 0.127N	P = 0.505N	P = 0.433N
Logistic regression test	P = 0.465	P = 0.321N	P = 0.389	P = 0.521N
Cochran-Armitage test	P = 0.447			
Fisher exact test		P = 0.345N	P = 0.343	P = 0.579N
Lung: Alveolar/bronchiolar Adenoma				
Overall rate	5/50 (10%)	9/50 (18%)	10/50 (20%)	16/50 (32%)
Adjusted rate	15.1%	22.7%	27.4%	47.3%
Terminal rate	2/28 (7%)	6/36 (17%)	6/32 (19%)	13/30 (43%)
First incidence (days)	616	531	602	418
Life table test	P = 0.005	P = 0.341	P = 0.218	P = 0.014
Logistic regression test	P = 0.006	P = 0.234	P = 0.193	P = 0.009
Cochran-Armitage test	P = 0.006	D 0.104	D 0 101	D 0.000
Fisher exact test		P = 0.194	P = 0.131	P = 0.006
Lung: Alveolar/bronchiolar Carcinoma				
Overall rate	2/50 (4%)	1/50 (2%)	5/50 (10%)	3/50 (6%)
Adjusted rate	6.2%	2.8%	13.1%	9.6%
Terminal rate	1/28 (4%)	1/36 (3%)	2/32 (6%)	2/30 (7%)
First incidence (days)	610 P= 0.341	729 (T)	588 P= 0.285	708 P= 0.534
Life table test Logistic regression test	P = 0.341 P = 0.351	P = 0.430N P = 0.474N	P = 0.285 P = 0.227	P= 0.534 P= 0.529
Cochran-Armitage test	P = 0.331 P = 0.348	$\Gamma = 0.4741$	$\Gamma = 0.227$	$\Gamma = 0.329$
Fisher exact test	1 – 0.340	P = 0.500N	P = 0.218	P = 0.500
I islici cauci test		1 - 0.0001	1 – 0.210	1 – 0.000
Lung: Alveolar/bronchiolar Adenoma or Carcinom		10/70 (000)	4 T (T 0 (0 0 0 ())	40 (70 (000))
Overall rate	7/50 (14%)	10/50 (20%)	15/50 (30%)	19/50 (38%)
Adjusted rate	20.6%	25.2%	37.9%	55.0%
Terminal rate	3/28 (11%)	7/36 (19%)	8/32 (25%)	15/30 (50%)
First incidence (days) Life table test	610 P= 0.004	531 P= 0.482	588 P= 0.114	418 P= 0.014
Logistic regression test	P = 0.004 P = 0.004	P = 0.462 P = 0.355	P = 0.114 P = 0.064	P= 0.014 P= 0.008
Cochran-Armitage test	P = 0.004 P = 0.004	$\Gamma = 0.333$	r = 0.004	I = 0.008
Fisher exact test	1 – 0.004	P = 0.298	P = 0.045	P = 0.006
Thyroid Gland (Follicular Cell): Adenoma Overall rate	3/50 (6%)	2/50 (4%)	1/50 (2%)	6/50 (12%)
Adjusted rate	10.7%	5.1%	3.1%	17.8%
Terminal rate	3/28 (11%)	1/36 (3%)	1/32 (3%)	4/30 (13%)
First incidence (days)	729 (T)	640	729 (T)	647
Life table test	P = 0.069	P = 0.390N	P = 0.257N	P= 0.294
Logistic regression test	P = 0.072	P = 0.437N	P = 0.257N	P = 0.278
Cochran-Armitage test	P = 0.073			
Fisher exact test		P = 0.500N	P = 0.309N	P = 0.243

TABLE C3
Statistical Analysis of Primary Neoplasms in Male Mice in the 2-Year Inhalation Study of Ethylbenzene (continued)

	Chamber Control	75 ppm	250 ppm	750 ppm
All Organs: Malignant Lymphoma				
Overall rate	2/50 (4%)	2/50 (4%)	3/50 (6%)	2/50 (4%)
Adjusted rate	4.8%	4.8%	8.4%	6.5%
Terminal rate	0/28 (0%)	1/36 (3%)	1/32 (3%)	1/30 (3%)
First incidence (days)	522	514	680	725
Life table test	P = 0.608	P = 0.655N	P = 0.564	P = 0.673N
Logistic regression test	P = 0.609N	P = 0.629	P = 0.474	P = 0.691
Cochran-Armitage test	P = 0.606N			
Fisher exact test		P = 0.691N	P = 0.500	P = 0.691N
All Organs: Benign Neoplasms				
Overall rate	20/50 (40%)	25/50 (50%)	26/50 (52%)	30/50 (60%)
Adjusted rate	56.4%	56.4%	68.1%	78.3%
Terminal rate	13/28 (46%)	17/36 (47%)	20/32 (63%)	22/30 (73%)
First incidence (days)	522	360	579	418
Life table test	P = 0.045	P = 0.564	P = 0.364	P = 0.100
Logistic regression test	P = 0.046	P = 0.304	P = 0.341	P = 0.053
Cochran-Armitage test	P = 0.046			
Fisher exact test		P = 0.211	P = 0.158	P = 0.036
All Organs: Malignant Neoplasms				
Overall rate	21/50 (42%)	15/50 (30%)	21/50 (42%)	16/50 (32%)
Adjusted rate	46.4%	35.2%	44.5%	36.4%
Terminal rate	5/28 (18%)	9/36 (25%)	7/32 (22%)	4/30 (13%)
First incidence (days)	289	514	480	259
Life table test	P = 0.349N	P = 0.082N	P = 0.381N	P = 0.184N
Logistic regression test	P = 0.287N	P = 0.179N	P = 0.448	P = 0.212N
Cochran-Armitage test	P = 0.309N			
Fisher exact test		P = 0.149N	P = 0.580N	P = 0.204N
All Organs: Benign or Malignant Neoplasms				
Overall rate	35/50 (70%)	34/50 (68%)	40/50 (80%)	41/50 (82%)
Adjusted rate	76.0%	70.8%	80.0%	89.0%
Terminal rate	17/28 (61%)	22/36 (61%)	22/32 (69%)	25/30 (83%)
First incidence (days)	289	360	480	259
Life table test	P = 0.131	P = 0.159N	P = 0.549N	P = 0.379
Logistic regression test	P = 0.069	P = 0.423N	P = 0.248	P = 0.162
Cochran-Armitage test	P = 0.063			
Fisher exact test		P = 0.500N	P = 0.178	P = 0.121

(T)Terminal sacrifice

Number of neoplasm-bearing animals/number of animals examined. Denominator is number of animals examined microscopically for liver, lung, and thyroid gland; for other tissues, denominator is number of animals necropsied.

b Kaplan-Meier estimated neoplasm incidence at the end of the study after adjustment for intercurrent mortality

Observed incidence at terminal kill

Beneath the chamber control incidence are the P values associated with the trend test. Beneath the exposure group incidence are the P values corresponding to pairwise comparisons between the chamber controls and that exposure group. The life table test regards neoplasms in animals dying prior to terminal kill as being (directly or indirectly) the cause of death. The logistic regression test regards these lesions as nonfatal. The Cochran-Armitage and Fisher exact tests compare directly the overall incidence rates. For all tests, a negative trend or a lower incidence in an exposure group is indicated by N.

e Not applicable; no neoplasms in animal group

 $\begin{tabular}{ll} TABLE~C4\\ Historical~Incidence~of~Alveolar/bronchiolar~Neoplasms~in~Chamber~Control~Male~B6C3F_1~Mice^a\\ \end{tabular}$

		Incidence in Controls		
Study	Adenoma	Carcinoma	Adenoma or Carcinoma	
Historical Incidence at IIT Research	Institute			
Isobutyl Nitrite	7/50	1/50	8/50	
Overall Historical Incidence				
Total Standard deviation Range	141/947 (14.9%) 7.0% 6%-36%	75/947 (7.9%) 5.7% 0%-16%	205/947 (21.7%) 8.0% 10%-42%	

^a Data as of 12 May 1995

TABLE C5
Summary of the Incidence of Nonneoplastic Lesions in Male Mice in the 2-Year Inhalation Study of Ethylbenzene^a

	Chamber Control	75 ppm	250 ppm	750 ppm
Disposition Summary				
Animals initially in study	50	50	50	50
Early deaths		•		
Accidental deaths	1			1
Moribund	6	2	5	6
Natural deaths	15	12	13	13
Survivors				
Died last week of study			1	
Terminal sacrifice	28	36	31	30
Animals examined microscopically	50	50	50	50
Alimentary System				
Gallbladder	(33)	(39)	(37)	(38)
Epithelium, hyperplasia	(00)	(00)	1 (3%)	(00)
Intestine small, duodenum	(45)	(48)	(43)	(45)
Parasite metazoan	(10)	(10)	1 (2%)	(10)
Epithelium, hyperplasia			1 (2%)	1 (2%)
Intestine small, jejunum	(44)	(46)	(44)	(43)
Cyst	(41)	1 (2%)	(11)	(40)
Epithelium, dysplasia		1 (2%)		
Peyer's patch, hyperplasia	1 (2%)	1 (2%)		
Intestine small, ileum	(42)	(46)	(44)	(41)
Peyer's patch, hyperplasia	1 (2%)	(10)	(11)	(41)
Liver	(50)	(50)	(50)	(50)
Angiectasis	(00)	(30)	(30)	1 (2%)
Basophilic focus	3 (6%)	3 (6%)	5 (10%)	4 (8%)
Clear cell focus	5 (10%)	4 (8%)	7 (14%)	3 (6%)
Cyst	1 (2%)	1 (2%)	7 (1470)	3 (070)
Eosinophilic focus	6 (12%)	8 (16%)	8 (16%)	12 (24%)
Eosinophilic focus, multiple	1 (2%)	0 (1070)	0 (1070)	12 (2470)
Fibrosis	1 (270)	1 (2%)		1 (2%)
Hemorrhage		1 (270)		2 (4%)
Hepatodiaphragmatic nodule			1 (2%)	2 (470)
Inflammation, chronic		2 (4%)	1 (270)	
Mineralization		~ (T/U)	1 (2%)	
Mixed cell focus	3 (6%)	2 (4%)	1 (2/0)	1 (2%)
Necrosis	7 (14%)	8 (16%)	10 (20%)	10 (20%)
Thrombosis	(11/0)	0 (10/0)	10 (20/0)	1 (2%)
Hepatocyte, hyperplasia				1 (2%)
Hepatocyte, hypertrophy	1 (2%)			17 (34%)
Hepatocyte, necrosis	1 (2%)	1 (2%)	3 (6%)	10 (20%)
Hepatocyte, syncytial alteration	1 (2/0)	5 (10%)	8 (16%)	23 (46%)
Hepatocyte, vacuolization cytoplasmic	4 (8%)	2 (4%)	4 (8%)	3 (6%)
Vein, thrombosis	1 (2%)	~ (T/U)	2 (4%)	3 (0/0)
Pancreas	(49)	(50)	(48)	(48)
Inflammation	1 (2%)	(00)	(10)	(10)
Acinus, hyperplasia	1 (2/0)			1 (2%)
Duct, cyst	1 (2%)	1 (2%)		1 (2/0)
Duct, cyst Duct, degeneration	1 (2/0)	1 (2/0)		1 (2%)
Duct, degeneration Duct, fibrosis	1 (2%)			1 (2/0)
Salivary glands	(50)	(50)	(50)	(50)
Infiltration cellular	21 (42%)	26 (52%)	17 (34%)	20 (40%)
Initia actori certarat	ω1 (Tω/U)	20 (02/0)	11 (01/0)	20 (10/0)

^a Number of animals examined microscopically at the site and the number of animals with lesion

TABLE C5
Summary of the Incidence of Nonneoplastic Lesions in Male Mice in the 2-Year Inhalation Study of Ethylbenzene (continued)

	Chamber Control	75 ppm	250 ppm	750 ppm
Alimentary System (continued)				
Stomach, forestomach	(48)	(50)	(50)	(47)
Cyst	4 (00)			1 (2%)
Ulcer	1 (2%)			1 (00/)
Epithelium, hyperplasia Serosa, inflammation				1 (2%) 1 (2%)
Stomach, glandular	(48)	(50)	(50)	(47)
Infiltration cellular	1 (2%)	(30)	(30)	(11)
Inflammation	1 (270)		1 (2%)	
Metaplasia	1 (2%)			
Mineralization	, ,		2 (4%)	
Tongue				(1)
Inflammation, granulomatous				1 (100%)
Tooth			(1)	(1)
Developmental malformation			1 (100%)	
Cardiovascular System				
Heart	(50)	(50)	(50)	(50)
Cardiomyopathy	18 (36%)	36 (72%)	29 (58%)	25 (50%)
Inflammation	2 (4%)			
Myocardium, mineralization	1 (2%)			
Pericardium, hyperplasia	1 (2%)			
Endocrine System				
Adrenal cortex	(47)	(47)	(48)	(48)
Accessory adrenal cortical nodule	1 (2%)			
Degeneration	1 (2%)	2 (4%)	1 (2%)	
Hemorrhage	1 (2%)			1 (2%)
Hyperplasia	13 (28%)	8 (17%)	9 (19%)	4 (8%)
Vacuolization cytoplasmic	1 (2%)	00 (470/)	00 (400/)	17 (050/)
Capsule, hyperplasia	19 (40%)	22 (47%)	20 (42%)	17 (35%)
Adrenal medulla	(47)	(46)	(48) 1 (2%)	(48)
Degeneration Hyperplasia			2 (4%)	
Mineralization			1 (2%)	
Islets, pancreatic	(49)	(50)	(48)	(48)
Degeneration	(/	()	1 (2%)	(/
Hyperplasia	5 (10%)	5 (10%)	8 (17%)	1 (2%)
Pituitary gland	(44)	(45)	(45)	(47)
Pars distalis, cyst		1 (2%)	2 (4%)	1 (2%)
Pars distalis, hyperplasia	1 (2%)	1 (2%)	1 (2%)	
Thyroid gland	(50)	(50)	(50)	(50)
Follicle, cyst	04 (122.1)	04 (1001)	1 (2%)	00 (0:00)
Follicular cell, hyperplasia	21 (42%)	21 (42%)	29 (58%)	32 (64%)
General Body System				
Tissue NOS	(2)	(3)	(1)	(2)
Cyst	• •	1 (33%)	, ,	• •
Fat, necrosis		2 (67%)		1 (50%)

TABLE C5
Summary of the Incidence of Nonneoplastic Lesions in Male Mice in the 2-Year Inhalation Study of Ethylbenzene (continued)

	Chamber Control	75 ppm	250 ppm	750 ppm
Genital System				
Epididymis	(49)	(50)	(50)	(50)
Atypia cellular	1 (2%)	1 (2%)		
Cyst	1 (2%)			0 (40()
Degeneration		4 (00/)		2 (4%)
Fibrosis	9 (40/)	1 (2%)	1 (90/)	
Granuloma sperm Infiltration cellular	2 (4%)	1 (2%)	1 (2%)	
Influration centuar Inflammation	1 (2%)	1 (2%)		
Mineralization	1 (2%)			
Bilateral, fibrosis	1 (2/0)		1 (2%)	
Penis	(1)	(2)	1 (270)	
Concretion	(1)	1 (50%)		
Inflammation	1 (100%)	1 (50%)		
Preputial gland	(48)	(49)	(48)	(49)
Cyst	(10)	(10)	1 (2%)	(10)
Degeneration Degeneration	3 (6%)	3 (6%)	9 (19%)	6 (12%)
Degeneration, cystic	3 (0/0)	1 (2%)	0 (10/0)	0 (12/0)
Fibrosis		I (w/U)	1 (2%)	
Hyperplasia	1 (2%)		1 (2%)	
Infiltration cellular	1 (2%)	5 (10%)	6 (13%)	4 (8%)
Inflammation	6 (13%)	11 (22%)	11 (23%)	13 (27%)
Mineralization	2 (22.3)	1 (2%)	(,	()
Necrosis		- ()	1 (2%)	
Prostate	(46)	(49)	(50)	(50)
Atrophy	1 (2%)	()	()	()
Infiltration cellular	1 (2%)	2 (4%)	1 (2%)	2 (4%)
Inflammation	6 (13%)	5 (10%)	5 (10%)	6 (12%)
Seminal vesicle	(49)	(50)	(50)	(50)
Atrophy	2 (4%)	1 (2%)	3 (6%)	1 (2%)
Degeneration	19 (39%)	26 (52%)	16 (32%)	22 (44%)
Inflammation	1 (2%)	1 (2%)	2 (4%)	
Testes	(49)	(50)	(50)	(50)
Atrophy		1 (2%)		1 (2%)
Mineralization				1 (2%)
Germinal epithelium, atrophy		1 (2%)		
Germinal epithelium, degeneration				1 (2%)
Interstitial cell, hyperplasia		1 (2%)		
Tunic, fibrosis			1 (2%)	
Hematopoietic System				
Bone marrow	(50)	(50)	(50)	(50)
Hematopoietic cell proliferation	•	1 (2%)	•	
Hyperplasia	1 (2%)	•		
Pigmentation, hemosiderin	2 (4%)	2 (4%)		1 (2%)
Myeloid cell, hyperplasia	2 (4%)	4 (8%)	9 (18%)	4 (8%)
Lymph node	(4)	(7)	(11)	(3)
Inguinal, hyperplasia	1 (25%)	1 (14%)		
Inguinal, pigmentation		1 (14%)		
Lumbar, congestion			1 (9%)	
Lumbar, hyperplasia	1 (25%)	4 (57%)	4 (36%)	2 (67%)
Lumbar, inflammation			2 (18%)	
Lumbar, pigmentation		1 (14%)		1 (33%)
Renal, congestion			2 (18%)	
Renal, hyperplasia		3 (43%)	3 (27%)	

TABLE C5
Summary of the Incidence of Nonneoplastic Lesions in Male Mice in the 2-Year Inhalation Study of Ethylbenzene (continued)

	Chamber Control	75 ppm	250 ppm	750 ppm
Hematopoietic System (continued)				
Lymph node, mandibular	(43)	(45)	(46)	(44)
Pigmentation, hemosiderin		1 (2%)		1 (2%)
Lymph node, mesenteric	(45)	(46)	(47)	(48)
Atrophy				1 (2%)
Congestion	1 (2%)	3 (7%)	1 (2%)	2 (4%)
Hematopoietic cell proliferation		3 (7%)	- 4	
Hyperplasia	1 (2%)	2 (4%)	3 (6%)	4 (00.1)
Inflammation	(0.4)	2 (4%)	(0.7)	1 (2%)
Lymph node, mediastinal	(24)	(25)	(27)	(25)
Hyperplasia	1 (4%)	(50)	(40)	(40)
Spleen	(50)	(50)	(49)	(49)
Atrophy	1 (2%)	4 (00/)	0 (40/)	
Hematopoietic cell proliferation	3 (6%)	4 (8%)	2 (4%)	1 (00/)
Lymphoid follicle, atrophy	2 (4%)		1 (2%)	1 (2%)
Lymphoid follicle, hyperplasia	(27)	(97)	2 (4%)	(24)
Thymus	(37)	(37)	(39)	(34)
Atrophy	18 (49%)	11 (30%)	20 (51%)	11 (32%)
Cyst	1 (3%)			
Integumentary System				
Mammary gland	(3)	(2)	(3)	(3)
Atrophy	1 (33%)	1 (50%)	2 (67%)	1 (33%)
Skin	(50)	(50)	(50)	(50)
Cyst		1 (2%)		
Infiltration cellular, melanocyte		1 (2%)		
Inflammation	3 (6%)	6 (12%)	4 (8%)	3 (6%)
Necrosis	1 (2%)	- 4	- 4	- 4
Ulcer	1 (2%)	5 (10%)	2 (4%)	3 (6%)
Hair follicle, atrophy		1 (2%)		
Prepuce, degeneration			1 (2%)	
Prepuce, hyperplasia, lymphoid			1 (2%)	
Prepuce, inflammation			2 (4%)	
Prepuce, ulcer		4 (00/)	2 (4%)	
Sebaceous gland, cyst		1 (2%)		
Musculoskeletal System				
Bone	(50)	(49)	(50)	(50)
Vertebra, degeneration	1 (2%)			1 (2%)
Nervous System				
Brain	(50)	(50)	(50)	(50)
Mineralization	21 (42%)	18 (36%)	19 (38%)	19 (38%)
Respiratory System				
Larynx	(48)	(49)	(46)	(49)
Foreign body	(10)	(10)	1 (2%)	(10)
Hemorrhage			± (₩/O)	1 (2%)
Infiltration, cellular	1 (2%)	4 (8%)	4 (9%)	- (~/~/
	- (~,0)	1 (2%)	3 (7%)	2 (4%)
Glands, degeneration		1 (2%)	3 (7%)	۵ (470)

3 (6%)

1 (100%)

(49) 2 (4%)

1 (2%)

8 (16%)

(1)

1 (2%)

1 (2%)

1 (2%)

1 (50%)

1 (50%)

1 (2%)

6 (12%)

1 (2%)

1 (2%) 1 (2%)

(49) 2 (4%)

(2)

Papilla, necrosis

Pelvis, dilatation

Degeneration Inflammation

Urinary bladder

Inflammation

Ulcer

Infiltration cellular

Muscularis, inflammation

Muscularis, necrosis

Serosa, fibrosis

Ureter

Renal tubule, vacuolization cytoplasmic

Calculus, microscopic observation only

TABLE C5

	Chamber Control	75 ppm	250 ppm	750 ppm
Respiratory System (continued)				
Lung	(50)	(50)	(50)	(50)
Congestion	` ,	` '	1 (2%)	` '
Hemorrhage			` ,	1 (2%)
Infiltration cellular, histiocyte	2 (4%)		1 (2%)	2 (4%)
Inflammation	1 (2%)		` ,	` ′
Pigmentation, hemosiderin	(1 (2%)	
Thrombosis			1 (2%)	
Alveolar epithelium, hyperplasia	1 (2%)	5 (10%)	2 (4%)	4 (8%)
Alveolar epithelium, metaplasia	- (2.3)	1 (2%)	2 (4%)	6 (12%)
Nose	(50)	(50)	(50)	(50)
Edema	(66)	(00)	(00)	1 (2%)
Hemorrhage	1 (2%)			1 (2%)
Inflammation	7 (14%)	3 (6%)	4 (8%)	1 (2%)
Polyp, inflammatory	2 (4%)	1 (2%)	2 (4%)	1 (270)
Nasolacrimal duct, inflammation	3 (6%)	1 (270)	1 (2%)	1 (2%)
Respiratory epithelium, inflammation	0 (070)		1 (270)	1 (2%)
Respiratory epithelium, metaplasia, squa	imous	1 (2%)		1 (270)
Pleura	inous	1 (270)		(1)
Trachea	(50)	(50)	(50)	(50)
Glands, cyst	(00)	(00)	(00)	1 (2%)
Glands, hemorrhage				1 (2%)
Special Senses System None				
Urinary System				
Kidney	(50)	(50)	(50)	(50)
Degeneration	1 (2%)			
Infarct	1 (2%)	# (400A)	T (100)	a (5 - 1)
Inflammation	3 (6%)	5 (10%)	5 (10%)	3 (6%)
Metaplasia, osseous		1 (2%)		
Mineralization		1 (2%)		
Nephropathy	34 (68%)	38 (76%)	40 (80%)	36 (72%)
Pigmentation, bile	1 (2%)			
Cortex, cyst	1 (2%)	8 (16%)	5 (10%)	4 (8%)
Papilla, inflammation	3 (6%)	4 (8%)	3 (6%)	2 (4%)
Papilla, necrosis			1 (2%)	

2 (4%)

(50) 1 (2%)

12 (24%)

1 (2%)

1 (2%)

7 (15%)

(48) 1 (2%)

APPENDIX D SUMMARY OF LESIONS IN FEMALE MICE IN THE 2-YEAR INHALATION STUDY OF ETHYLBENZENE

TABLE D1	Summary of the Incidence of Neoplasms in Female Mice	
	in the 2-Year Inhalation Study of Ethylbenzene	168
TABLE D2	Individual Animal Tumor Pathology of Female Mice	
	in the 2-Year Inhalation Study of Ethylbenzene	172
TABLE D3	Statistical Analysis of Primary Neoplasms in Female Mice	
	in the 2-Year Inhalation Study of Ethylbenzene	190
TABLE D4a	Historical Incidence of Alveolar/bronchiolar Neoplasms	
	in Chamber Control Female B6C3F ₁ Mice	194
TABLE D4b	Historical Incidence of Hepatocellular Neoplasms	
	in Chamber Control Female B6C3F ₁ Mice	194
TABLE D5	Summary of the Incidence of Nonneoplastic Lesions in Female Mice	
	in the 2-Year Inhalation Study of Ethylbenzene	195

TABLE D1
Summary of the Incidence of Neoplasms in Female Mice in the 2-Year Inhalation Study of Ethylbenzene^a

	Chamber Control	75 ppm	250 ppm	750 ppm
Disposition Summary				
Animals initially in study	50	50	50	50
Early deaths				
Accidental deaths	1		1	
Moribund	5	6	1	4
Natural deaths	9	6	8	9
Survivors				
Died last week of study	1			
Terminal sacrifice	34	38	40	37
Animals examined microscopically	50	50	50	50
Alimentary System	(40)	(40)	(50)	(50)
Esophagus	(48)	(48)	(50)	(50)
Gallbladder	(44)	(44)	(44)	(46)
Intestine large, rectum	(49)	(48)	(49)	(47)
Intestine large, cecum	(49)	(47)	(48)	(44)
Intestine small, duodenum	(45)	(48)	(47)	(46)
Polyp adenomatous	(46)	(46)	1 (2%)	(45)
Intestine small, jejunum Intestine small, ileum	(46) (47)	(46) (47)	(46) (47)	(45) (46)
Liver	(50)	(50)	(50)	(50)
Cholangiocarcinoma	(30)	1 (2%)	(30)	(30)
Fibrosarcoma, metastatic, pancreas	1 (2%)	1 (~/0)		
Hemangioma	1 (2/0)	1 (2%)		
Hepatocellular carcinoma	7 (14%)	4 (8%)	3 (6%)	10 (20%)
Hepatocellular carcinoma, multiple	. (==.0)	- (0.0)	5 (512)	2 (4%)
Hepatocellular adenoma	6 (12%)	8 (16%)	9 (18%)	12 (24%)
Hepatocellular adenoma, multiple		1 (2%)	3 (6%)	4 (8%)
Pancreas	(50)	(50)	(50)	(49)
Fibrosarcoma	1 (2%)			
Salivary glands	(50)	(50)	(50)	(50)
Stomach, forestomach	(50)	(49)	(48)	(50)
Squamous cell papilloma	1 (2%)	2 (4%)	1 (2%)	1 (2%)
Stomach, glandular	(50)	(49)	(48)	(50)
Serosa, sarcoma, metastatic, uterus		1 (2%)		
Cardiovascular System				
Blood vessel	(46)	(48)	(48)	(50)
Adventitia, hepatocellular carcinoma,				
metastatic, liver	1 (2%)			
Heart	(50)	(49)	(50)	(50)
Fibrosarcoma, metastatic, pancreas	1 (2%)			
Endocrine System				
Adrenal cortex	(47)	(50)	(50)	(49)
Adenoma			1 (2%)	
Adrenal medulla	(47)	(50)	(50)	(49)
Pheochromocytoma malignant			1 (2%)	
Pheochromocytoma benign		1 (2%)	1 (2%)	
Islets, pancreatic	(50)	(50)	(50)	(49)
Adenoma			1 (2%)	

TABLE D1
Summary of the Incidence of Neoplasms in Female Mice in the 2-Year Inhalation Study of Ethylbenzene (continued)

	Chamber Control	75 ppm	250 ppm	750 ppm
Endocrine System (continued)				
Pituitary gland	(48)	(49)	(47)	(49)
Pars distalis, adenoma	4 (8%)	8 (16%)	7 (15%)	5 (10%)
Pars intermedia, adenoma Thyroid gland	(50)	(50)	1 (2%) (50)	(50)
Follicular cell, adenoma	5 (10%)	4 (8%)	3 (6%)	4 (8%)
Follicular cell, adenoma, multiple	0 (1076)	2 (4%)	0 (070)	1 (0/0)
General Body System				
Tissue NOS	(1)	(6)	(4)	(1)
Hemangiosarcoma		1 (17%)		
Leiomyosarcoma	1 (100%)	1 (170/)		
Abdominal, osteosarcoma Pelvic, sarcoma		1 (17%)	1 (25%)	
Genital System				
Clitoral gland	(41)	(47)	(48)	(48)
Fibrosarcoma	(11)	(11)	(10)	1 (2%)
Ovary	(49)	(50)	(49)	(49)
Cystadenoma	2 (4%)			2 (4%)
Fibrosarcoma, metastatic, pancreas	1 (2%)			
Granulosa cell tumor benign	1 (2%)	(50)	(50)	(50)
Uterus	(50) 1 (2%)	(50)	(50)	(50)
Leiomyosarcoma Polyp stromal	2 (4%)	1 (2%)	1 (2%)	
Sarcoma	£ (470)	1 (2%)	1 (2/0)	
Endometrium, adenoma	1 (2%)	1 (270)		
Myometrium, hemangioma	` ,		1 (2%)	
Vagina	(1)			
Leiomyosarcoma	1 (100%)			
Hematopoietic System				
Bone marrow	(48)	(50)	(50)	(50)
Lymph node	(3)	(7)	(2)	(5)
Iliac, hemangioma	JOC	1 (14%) 1 (14%)		
Lumbar, osteosarcoma, metastatic, tissue N Renal, hemangiosarcoma	103	1 (14%)		
Lymph node, bronchial	(32)	(40)	(29)	(38)
Fibrosarcoma, metastatic, pancreas	1 (3%)	(/	\/	\/
Hepatocellular carcinoma, metastatic, liver				
Lymph node, mandibular	(47)	(48)	(47)	(44)
Lymph node, mesenteric	(48)	(48)	(46)	(44)
Fibrosarcoma, metastatic, pancreas	1 (2%)	(40)	(44)	(0.1)
Lymph node, mediastinal	(34)	(42)	(41)	(31)
Fibrosarcoma, metastatic, pancreas	1 (3%)			
Hepatocellular carcinoma, metastatic, liver Spleen	1 (3%) (50)	(50)	(50)	(49)
Capsule, fibrosarcoma, metastatic, pancrea		(00)	(00)	(10)
Thymus	(42)	(44)	(45)	(46)
Hepatocellular carcinoma, metastatic, liver		` '	\ -/	\ -/

TABLE D1
Summary of the Incidence of Neoplasms in Female Mice in the 2-Year Inhalation Study of Ethylbenzene (continued)

	Chamber Control	75 ppm	250 ppm	750 ppm
Integumentary System				
Mammary gland	(49)	(50)	(48)	(49)
Carcinoma	1 (2%)	3 (6%)		
Skin	(50)	(50)	(49)	(50)
Fibroma				1 (2%)
Fibrosarcoma		2 (4%)		4 (00.1)
Fibrous histiocytoma		1 (00/)	0 (40()	1 (2%)
Hemangioma		1 (2%)	2 (4%)	
Squamous cell carcinoma Sebaceous gland, adenoma		1 (2%)	1 (2%)	
Musculoskeletal System				
Bone	(49)	(50)	(50)	(50)
Rib, sarcoma, metastatic, tissue NOS	(10)	(00)	1 (2%)	(00)
Vertebra, osteosarcoma			1 (2%)	
Skeletal muscle		(2)	- (~,0)	
Carcinoma, metastatic, mammary gland		1 (50%)		
Rhabdomyosarcoma		1 (50%)		
Vervous System				
Brain	(50)	(50)	(50)	(50)
Cerebrum, oligodendroglioma benign	1 (2%)	, ,	· ,	` ,
D				
Respiratory System	(40)	(40)	(47)	(40)
Larynx	(49)	(49)	(47)	(48)
Lung Alveolar/bronchiolar adenoma	(50) 3 (6%)	(50) 4 (8%)	(49) 4 (8%)	(50) 8 (16%)
Alveolar/bronchiolar adenoma, multiple	1 (2%)	4 (0/0)	1 (2%)	8 (1076)
Alveolar/bronchiolar carcinoma	1 (270)	2 (4%)	1 (2%)	
Carcinoma, metastatic, harderian gland	1 (2%)	L (470)		
Carcinoma, metastatic, mammary gland	1 (270)	1 (2%)		
Hepatocellular carcinoma, metastatic, liver	3 (6%)	1 (2%)	2 (4%)	1 (2%)
Osteosarcoma, metastatic, tissue NOS	0 (070)	1 (2%)	2 (170)	1 (273)
Sarcoma, metastatic, tissue NOS		- ()	1 (2%)	
Sarcoma, metastatic, uterus		1 (2%)	, ,	
Squamous cell carcinoma, metastatic, lacrimal gland				1 (2%)
Vose	(49)	(50)	(50)	(50)
Carcinoma, metastatic, harderian gland	1 (2%)			
Pleura	(1)			
Hepatocellular carcinoma, metastatic, liver	1 (100%)	(50)	(50)	(50)
Trachea	(50)	(50)	(50)	(50)
Special Senses System				
Tarderian gland	(1)		(1)	(3)
Adenoma			1 (100%)	3 (100%)
Carcinoma	1 (100%)			
Lacrimal gland				(1)
Squamous cell carcinoma				1 (100%)

TABLE D1 Summary of the Incidence of Neoplasms in Female Mice in the 2-Year Inhalation Study of Ethylbenzene (continued)

	Chamber Control	75 ppm	250 ppm	750 ppm
Urinary System				
Kidney	(50)	(50)	(50)	(50)
Cholangiocarcinoma, metastatic, liver		1 (2%)		
Cortex, fibrosarcoma, metastatic, pancreas	1 (2%)			
Ureter		(1)		(1)
Urinary bladder	(47)	(48)	(47)	(49)
Serosa, sarcoma, metastatic, uterus		1 (2%)		
Systemic Lesions Multiple organs ^b Leukemia granulocytic	(50) 1 (2%)	(50)	(50)	(50)
Lymphoma malignant	3 (6%)	6 (12%)	5 (10%)	5 (10%)
Neoplasm Summary				
Total animals with primary neoplasms ^c	29	38	31	38
Total primary neoplasms	44	58	50	60
Total animals with benign neoplasms	20	26	27	28
Total benign neoplasms	27	34	39	40
Total animals with malignant neoplasms	13	20	9	18
Total malignant neoplasms	17	24	11	20
Total animals with metastatic neoplasms	5	5	3	2
Total metastatic neoplasms	18	9	4	2

Number of animals examined microscopically at the site and the number of animals with neoplasm Number of animals with any tissue examined microscopically Primary neoplasms: all neoplasms except metastatic neoplasms

TABLE D2 Individual Animal Tumor Pathology of Female Mice in the 2-Year Inhalation Study of Ethylbenzene: Chamber Control

Number of Days on Study	2 4 5 5 5 5 5 6 6 6 6 6 7 7 7 7 7 7 7 7 7 7
Jan San San San San San San San San San S	5 3 1 5 8 2 7 8 9 9 8 4 5 8 5 0 0 0 0 0 1 1 1 1 1
	0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0
Carcass ID Number	8 9 9 6 6 9 7 9 7 6 6 9 6 7 5 5 7 7 7 9 5 5 5 6
	0 7 0 4 7 2 5 4 3 5 0 9 6 7 8 6 1 6 9 5 1 2 5 9 1
Alimentary System	
Esophagus	M + + + + + + + + + + + + + + + + + + +
Gallbladder	A + + + + A + + + M A + + + + + + + + +
Intestine large, colon	+ + + + + + + + M + + A + + + + + + + +
Intestine large, rectum	A + + + + + + + + + + + + + + + + + + +
Intestine large, cecum	A + + + + + + + + + + + + + + + + + + +
Intestine small, duodenum	M + + + + + + + + + + + + + + + + + + +
Intestine small, jejunum	A + + + + + + + + A + A + A + + + + + +
Intestine small, ileum	A + + + + + + + + A A + + + + + + + + +
Liver	+ + + + + + + + + + + + + + + + + + + +
Fibrosarcoma, metastatic, pancreas	X
Hepatocellular carcinoma	X X X X X X
Hepatocellular adenoma	X X
Mesentery	+
Pancreas	+ + + + + + + + + + + + + + + + + + + +
Fibrosarcoma	X
Salivary glands	+ + + + + + + + + + + + + + + + + + + +
Stomach, forestomach	+ + + + + + + + + + + + + + + + + + +
Squamous cell papilloma	X
Stomach, glandular	+ + + + + + + + + + + + + + + + + + + +
Cardiovascular System	
Blood vessel	+ + + + + + M + + M + + + + + + + + + +
Adventitia, hepatocellular carcinoma,	
metastatic, liver	X
Heart	+ + + + + + + + + + + + + + + + + + + +
Fibrosarcoma, metastatic, pancreas	X
Endocrine System	
Adrenal cortex	M + + + + + + + M + + + M + + + + + + +
Adrenal medulla	M + + + + + + + M + + + M + + + + + + +
Islets, pancreatic	+ + + + + + + + + + + + + + + + + + + +
Parathyroid gland	+ + M + M + + + M + + + + + M M + + + +
Pituitary gland	+ + + + + + + + + M M + + + + + + + + +
Pars distalis, adenoma	X X
Thyroid gland	+ + + + + + + + + + + + + + + + + + + +
Follicular cell, adenoma	X X X
General Body System	
Tissue NOS	+
Leiomyosarcoma	X
•	
Genital System	M . M MMM W
Clitoral gland	+ + M + M + + + + M M M + + + M + + + +
Ovary	+ + + + + + + + + + + + + + + + + + +
Cystadenoma	X X
Fibrosarcoma, metastatic, pancreas	X
Granulosa cell tumor benign	

^{+:} Tissue examined microscopically A: Autolysis precludes examination

M: Missing tissue I: Insufficient tissue

X: Lesion present Blank: Not examined

TABLE D2 Individual Animal Tumor Pathology (continued)	of Fema	ale	M	ice	in	th	e 2	-Ye	ear	In	ha	lat	ion	St	ud	y o	f E	Eth	ylb	en	zei	ıe:	C	ha	mb	er Control
(**************************************	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	
Number of Days on Study	3 1	3 1	3 1	3 1	3 1	3 1	3 1	3 1	3 1	3 1	3 1	3 2														
	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	1	Total
Carcass ID Number	6 2	6 8	6 9	7 2	7 8	8 2	8 4	8 5	8 6	8 9	9	5 3	5 4	5 7	6 3	7 0	7 4	8 1	8 3	8 7	8	9 1	9 6	9 8	0 0	Tissues/ Tumors
Alimentary System																										
Esophagus	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	M		+	+	+	+	+	+	+	48
Gallbladder	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	M	+	+	+	+	+	+	+	44
Intestine large, colon	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	48
Intestine large, rectum	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	49
Intestine large, cecum	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	49
Intestine small, duodenum	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	45
Intestine small, jejunum	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	46
Intestine small, ileum	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	47
Liver	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Fibrosarcoma, metastatic, pancreas				3.7																						1
Hepatocellular carcinoma				X						v					v	v							v			7
Hepatocellular adenoma										X					A	X							X			6
Mesentery Pancreas															+											2
Fibrosarcoma	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
																										1 50
Salivary glands	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50 50
Stomach, forestomach	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50 1
Squamous cell papilloma Stomach, glandular																										50
Stomach, glandular		_	_	_		_	_	_		_	_	_	_	_	_	_	_	_	_	_	_	_	_	_		30
Cardiovascular System																										
Blood vessel	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	46
Adventitia, hepatocellular carcinoma,																										
metastatic, liver																										1
Heart	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Fibrosarcoma, metastatic, pancreas																										1
Endocrine System																										
Adrenal cortex	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	47
Adrenal medulla	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	47
Islets, pancreatic	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Parathyroid gland	N.	1 +	+	N	1 M	[+	M	M	M	M	M	M	+	+	M	M	M	M	+	+	+	M	+	+	+	26
Pituitary gland	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	48
Pars distalis, adenoma																							X		X	4
Thyroid gland	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Follicular cell, adenoma								X														X				5
General Body System																										
Tissue NOS																										1
Leiomyosarcoma																										1
Genital System																										
Clitoral gland		1.4			1.4	[+			J.	_	5	J.	_1	M	5		_							. 1		41
	+	171	. +	+	101	. +	+	+	+	+	+	+ M	+	171	+	+	+	+	+	+	+	+	+	+	+	41
Ovary Cystadenoma	+	+	+	+	+	+	+	+	+	+	+	ıVI	+	+	+	+	+	+	+	+	+	+	+	+	+	49 2
Fibrosarcoma, metastatic, pancreas															X											1
Granulosa cell tumor benign															Λ											1

TABLE D2 Individual Animal Tumor Pathology of (continued)	Fema	ıle	M	ice	in	the	e 2-	Ye	ear	In	hal	lati	ion	St	ud	y o	of I	Eth	ylł	en	zei	ıe:	C	ha	mber Control
Number of Days on Study	2 1 5	4 2 3	5 0 1	5 6 5	5 7 8	5 9 2	5 9 7	6 1 8	6 2 9	3	6 8 8	6 9 4	7 1 5	7 1 8	7 2 5	7 3 0	7 3 0	7 3 0	7 3 0	7 3 0	7 3 1	7 3 1	7 3 1	7 3 1	7 3 1
Carcass ID Number	0 8 0	0 9 7	0 9 0	0 6 4	0 6 7	0 9 2	7	0 9 4	0 7 3	0 6 5	6	0 9 9	0 6 6	0 7 7	5	0 5 6	0 7 1	0 7 6	0 7 9	0 9 5	0 5 1	0 5 2	0 5 5	0 5 9	6
Genital System (continued) Uterus Leiomyosarcoma Polyp stromal Endometrium, adenoma Vagina Leiomyosarcoma	+	+	+	+ X X		+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+ X	+	+
Hematopoietic System Bone marrow Lymph node Lymph node, bronchial Fibrosarcoma, metastatic, pancreas	+ M	+ + +	+ M	+	+ + M	+	+ + +	+	+ M	+	+		+ + X	+						+ M	+	+	+	+	+ M
Hepatocellular carcinoma, metastatic, liver Lymph node, mandibular Lymph node, mesenteric Fibrosarcoma, metastatic, pancreas		+				+		+	+	+	M	+	+ X	+	+		+	+	+	+	+	+	+	+	+++
Lymph node, mediastinal Fibrosarcoma, metastatic, pancreas Hepatocellular carcinoma, metastatic, liver Spleen	M +	+	+	M +	+]	M	M +						+ X +	X						M +	+	+	+	M +	+
Capsule, fibrosarcoma, metastatic, pancreas Thymus Hepatocellular carcinoma, metastatic, liver	M	+	+	+	M	. +	M						X							+	+	+	M	+	+
Integumentary System Mammary gland Carcinoma	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	M	+	+	+	+	+	+	+	+
Skin Musculoskeletal System	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Nervous System Brain Cerebrum, oligodendroglioma benign Spinal cord	+	+	+	+	+ X	+	+ + +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Respiratory System Larynx Lung Alveolar/bronchiolar adenoma Alveolar/bronchiolar adenoma, multiple Carcinoma, metastatic, harderian, gland	+	+	+	+	+	+	+++	+++	+	M +	+++	+ + X	+++	+++	+	+++	+	+	+	+	+	+	+	+	+++
Hepatocellular carcinoma, metastatic, liver Nose Carcinoma, metastatic, harderian gland Pleura Hepatocellular carcinoma, metastatic, liver	М	+	+	+	+	+	+	+	+		X +		+	X + + X	+	+	+	+	+	+	+	+	X +	+	+
Trachea	+	+	+	+	+	+	+	+	+	+	+	+	+		+	+	+	+	+	+	+	+	+	+	+

							,- I	eai	111	na	iau	1011	St	ua	y u	IE	un	УШ	en	ZEI	ıe:	C	пä	mD(er Control
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0 6 2	6	6 (3	7 7	' 8	8	8	0 8 6	0 8 9	0 9 3	0 5 3	0 5 4	0 5 7			0 7 4	0 8 1	0 8 3	0 8 7	0 8 8	0 9 1	0 9 6	9	0	Total Tissues/ Tumors
+	• +	+ -	+	+ -	⊦ -	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+ X	+	+	+	+	+	50 1 2 1 1
+	· +	+ - √1 -	+ -	+ - M -	⊦ - ∟ N	⊦ +	+	+	+	+ M	+	+	+ M	+	+ M	+	+ M	+	+	+	+	+	+ M	+ M	48 3 32
+	. 4	•• + -	+ -	+ -		v1 · ⊢ + ⊢ +	+ +	++	++	+ +					+ +	++	+ +	++	+++	+++	+++	+ +	+ +	+ +	1 1 47 48
M	1 ⊣	+ l	M I	М -	⊦ -	+ +	+	M	M	M	+	+	M	+	+	+	+	+	+	+	M	+	+	+	1 34 1 1
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+	. 4	+ - + -	+ -	+ -	⊦ - ⊦ -	⊦ + ⊦ +			+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	49 1 50
+	. 4	+ -	+ -	+ -	-	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	49
+	. 4	+ -	+	+ -	-	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50 1 1
+	. 4	+ -	+ -	+ -	⊦ -	+ + + +	+	+	+	+	+ + X	+	+++	+++	+ + X	+ +	+++	+++	+++	+++	+++	+++	+ + X	+	49 50 3 1
+	. 4	+ -	+ -	+ -	⊦ -	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	3 49 1 1 1 50
	++++++	1 1 1 0 0 0 6 6 6 2 8 + + + + + + + + + + + + + + + + +	1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	1 1 1 0 0 0 0 6 6 6 2 8 9 + + + + M + + + + + + +	1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	1 1 1 1 1 1 1 1 1 1 1 1 1 1 2 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 6 6 6 6	1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 2 2 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0	1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 2 2 2 2 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0	1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 2 2 2 2 2 2 2 2 3 3 3 4 7 3 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 3 4 7 3 4 7 3 4 7 3 4 7 3 4 7 3 4 7 4 7	1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 2 2 2 2	1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 2 2 2 2	1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 2 2 2 2	1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2	1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2	1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2	1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2

TABLE D2 Individual Animal Tumor Pathology of (continued)	f Fema	ale	Mi	ice	in	the	e 2	-Ye	ear	In	ha	lat	ion	St	ud	y o	of I	Eth	yll	en	ze	ne:	C	ha	mb	er Con	tro
	2	4	5	5	5	5	5	6	6	6	6	6	7	7	7	7	7	7	7	7	7	7	7	7	7		_
Number of Days on Study	1	2	0	6	7	9	9	1	2	3	8	9	1	1	2	3	3	3	3	3	3	3	3	3	3		
•	5	3	1	5	8	2	7	8	9	9	8	4	5	8	5	0	0	0	0	0	1	1	1	1	1		
	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0		
Carcass ID Number	8	9	9	6	6	9	7	9	7	6	6	9	6	7	5	5	7	7	7	9	5	5	5	5	6		
	0	7	0	4	7	2	5	4	3	5	0	9	6	7	8	6	1	6	9	5	1	2	5	9	1		
Special Senses System Harderian gland Carcinoma										+ X																	
Urinary System																											
Kidney	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+		
Cortex, fibrosarcoma, metastatic, pancreas												٠.	X														
Urinary bladder	+	+	+	+	+	+	+	+	М	+	+	M	+	+	+	+	+	+	+	+	+	+	+	+	+		
Systemic Lesions																											
Multiple organs	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+		
Leukemia granulocytic						X																					
Lymphoma malignant		X							X										X								

TABLE D2 Individual Animal Tumor Pathology of (continued)	f Fema	ale	M	ice	in	the	2 -	Ye	ear	In	ha	lat	ion	St	ud	y o	f I	Eth	yll	en	zei	ne:	C	ha	mb	er Control
Number of Days on Study	7 3 1	7 3 2																								
Carcass ID Number	0 6 2	0 6 8	0 6 9	0 7 2	0 7 8	0 8 2	0 8 4	0 8 5	0 8 6	0 8 9	0 9 3	0 5 3	0 5 4	0 5 7	0 6 3	0 7 0	0 7 4	0 8 1	0 8 3	0 8 7	0 8 8	0 9 1	0 9 6	0 9 8	1 0 0	Total Tissues/ Tumors
Special Senses System Harderian gland Carcinoma																										1 1
Urinary System Kidney Cortex, fibrosarcoma, metastatic, pancreas Urinary bladder	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+ M	+	50 1 47
Systemic Lesions Multiple organs Leukemia granulocytic Lymphoma malignant	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50 1 3

Individual Animal Tumor Pathology	Temale Mice in the 2-1		11
·	3 4 5 5 5 6 6	6 6 6 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7	7 7 7
Number of Days on Study	0 9 6 8 9 0 3	4 9 9 0 2 3 3 3 3 3 3 3 3 3 3 3	3 3 3
v	0 5 2 4 0 2 9	2 7 7 3 3 0 0 0 1 1 1 1 1 1 1	1 1 1
	1 1 1 1 1 1 1	1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	1 1 1
Carcass ID Number			7 7 8
Cui cuis 12 Tumboi	2 3 7 8 5 1 4		4 6 5
Alimentary System			
Esophagus	+ + + + + + +	+ + + + + + + + + + + + + + + +	+ + +
Gallbladder	A + + + A + + 1	M + + + M + + + M + + + + + + + + + + +	+ + +
Intestine large, colon	+ + + + + A + + + + + + + + + + + + + +	+ + + + + + + + + + + + + + + +	+ + +
Intestine large, rectum	$A + + + + A + \cdots$	+ + + + + + + + + + + + + + + +	+ + +
Intestine large, cecum	A + + + A A + +	+ + + + + + + + + + + + + + + +	+ + +
Intestine small, duodenum	$A + + + A + + + \cdots$	+ + + + + + + + + + + + + + + +	+ + +
Intestine small, jejunum	+ $+$ $+$ $+$ $+$ $+$ $+$ $+$ $+$ $+$	+ + + + + + + + + + + + + + + +	+ + +
Intestine small, ileum	A + + + A A + +	+ + + + + + + + + + + + + + + +	+ + +
Liver		+ + + + + + + + + + + + + + + +	+ + +
Cholangiocarcinoma	X		
Hemangioma		X	
Hepatocellular carcinoma	X	. X	
Hepatocellular adenoma	X	X X X	X
Hepatocellular adenoma, multiple			
Pancreas	+ + + + + + +	+ + + + + + + + + + + + + + + + + + + +	+ + +
Salivary glands	+ + + + + + + +	+ + + + + + + + + + + + + + + + + + + +	+ + +
Stomach, forestomach	+ + + + A + + +	+ + + + + + + + + + + + + + + + + + + +	+ + +
Squamous cell papilloma	X	X	
Stomach, glandular	+ + + + A + + +	+ + + + + + + + + + + + + + + + + + + +	+ + +
Serosa, sarcoma, metastatic, uterus	X		
Cardiovascular System			
Blood vessel	A + + + + + + +	+ + + + + + + + + + + + + + + + + + +	+ + +
Heart	+ + + + + + +	+ + + + + + + + + + + + + + + +	+ + +
Endocrine System			
Adrenal cortex	+ + + + + + +	+ + + + + + + + + + + + + + + +	+ + +
Adrenal medulla	+ + + + + + +	+ + + + + + + + + + + + + + +	+ + +
Pheochromocytoma benign			
Islets, pancreatic	+ + + + + + +	+ + + + + + + + + + + + + + + +	+ + +
Parathyroid gland		$M \ M \ + \ + \ M \ M \ + \ M \ M \ M \ $	
Pituitary gland	$+ + + + + M + \cdots$	+ + + + + + + + + + + + + + + +	
Pars distalis, adenoma		X X X	
Thyroid gland	+ + + + + + +	+ + + + + + + + + + + + + + + +	+ + +
Follicular cell, adenoma		X	
Follicular cell, adenoma, multiple		X X	
General Body System			
Гissue NOS	+ +	+	
Hemangiosarcoma			
Abdominal, osteosarcoma	X		
Genital System			
Clitoral gland	+ M + + M M + +	+ + + + + + + + + + + + + + + +	+ + +
Ovary	+ + + + + + +	+++++++++++++	+ + +
Uterus	+ + + + + + +	+++++++++++++	+ + +
Polyp stromal	•		
Sarcoma	X		

Individual Animal Tumor Patholog	,															`	_			•						_	(00110111010
	7	,	7 7	7	7 1	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	
Number of Days on Study	3	:	3	3	3 3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	
3	1		l 1	l	1 :	1	1	1	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	
	1		l 1		1 :	1	1	2	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	Total
Carcass ID Number	8	8	3 9)	9 9	9	9	0	5	5	5	6	6	6	6	6	7	7	7	8	8	8	9	9	9	9	Tissues/
	6	9	9 3	3	5 8	8	9	0	6	8	9	2		4		8	3	5	9	0	1	2	0	1	2	7	Tumors
Alimentary System																											
Esophagus	+		+ -	+	+]	M	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	M	+	48
Gallbladder	+		+ -	+ :	Μ -	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	44
Intestine large, colon	+	-	+ -	+	+ -	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	49
Intestine large, rectum	+	-	+ -	+	+ -	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	48
Intestine large, cecum	+	-	+ -	+	+ -	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	47
Intestine small, duodenum	+	-	+ -	+	+ -	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	48
Intestine small, jejunum	+	-	+ -	+	+ -	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	46
Intestine small, ileum	+	-	+ -	+	+ -	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	47
Liver	+	-	+ -	+	+ -	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Cholangiocarcinoma																											1
Hemangioma																											1
Hepatocellular carcinoma																		X					X				4
Hepatocellular adenoma				Κ.	X								X					X									8
Hepatocellular adenoma, multiple		2	X																								1
Pancreas	+	-	+ -	+	+ -	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Salivary glands	+	-	+ -	+	+ -	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Stomach, forestomach	+	-	+ -	+	+ -	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	49
Squamous cell papilloma																											2
Stomach, glandular	+		+ -	+	+ -	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	49
Serosa, sarcoma, metastatic, uterus																											1
Cardiovascular System																											
Blood vessel	+		+ -	+	+ -	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	48
Heart	+		+ -	+	+ -	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	49
Endocrine System																											
Adrenal cortex	+		+ -	+	+ -	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Adrenal medulla	+		+ -	+	+ -	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Pheochromocytoma benign																										X	1
Islets, pancreatic	+		+ -	+	+ -	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+		+		50
Parathyroid gland	N	1 -	+ -	+	+]	M	M	M	+	+	M	+	+	M	M	M	+	+	M	M	M	+	M	M	+	M	24
Pituitary gland	+		+ -	+	+ -	+	+	+	+		+		+	+	+	+	+	+	+	+	+	+	+	+	+	+	49
Pars distalis, adenoma										X		X					X				X						8
Thyroid gland	+		+ -	+		+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Follicular cell, adenoma					2	X									X	X											4
Follicular cell, adenoma, multiple																											2
General Body System																											
Tissue NOS			-	+								+						+									6
Hemangiosarcoma																		X									1
Abdominal, osteosarcoma																											1
Genital System																											
Clitoral gland	+	-	+ -	+	+ -	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	47
Ovary	+	-	+ -	+	+ -	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Uterus	+	-	+ -	+	+ -	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Polyp stromal								X																			1
Sarcoma																											1

Individual Animal Tumor Pathology of	1 Cine												.	-	uu,	, ,			,	~				· P	pm (continued,
	3	4	5	5	5	6	6	6	6	6	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7
Number of Days on Study	0	9	6	8	9	0	3	4	9	9	0	2	3	3	3	3	3	3	3	3	3	3	3	3	3
	0	5	2	4	0	2	9	2	7	7	3	3	0	0	0	1	1	1	1	1	1	1	1	1	1
_	1				1													1				1	1	1	
Carcass ID Number	5	5	8	8	5	6	9	9	7	8	6	8	6	7	7	5	5	5	6	6	7	7	7	7	8
	2	3	/	8	5	1	4	b	8	3	7	4	0	1	1	1	4	1	b	9	0	2	4	6	5
Hematopoietic System																									
Bone marrow	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Lymph node Iliac, hemangioma			+	+			+				+			+		+									
Lumbar, osteosarcoma, metastatic,																									
tissue NOS			X																						
Renal, hemangiosarcoma																									
Lymph node, bronchial	+	+	+	+	+				M						M		+	+	+	+	M	+	+	+	+
Lymph node, mandibular	+	+	+	+	+	+	+		M			+	+		M	+	+	+	+	+	+	+	+	+	+
Lymph node, mesenteric Lymph node, mediastinal	M +		+ +]	+			+		+	+	+	+ M	+ M	+	+	+	+	+	+	+	+	+	+	+	+ M
Spleen	+	1V.	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Thymus	M	+	M	+	+			M	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Integrimentary System																									
Integumentary System Mammary gland	_	_	_	_	_	_	_	_	_	_	_	_	_	_	_	_	_	_	_	_	_	_	_	_	+
Carcinoma	+	+	+	+	+	+	+	+ X	+	+	+	+ X	+	+ X	т	+	_	+	_	+	+	+	+	+	т'
Skin	+	+	+	+	+	+	+	+	+	+	+		+		+	+	+	+	+	+	+	+	+	+	+
Fibrosarcoma															X										
Hemangioma																					X				
Squamous cell carcinoma																									
Musculoskeletal System																									
Bone	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Skeletal muscle										+		+													
Carcinoma, metastatic, mammary gland										37		X													
Rhabdomyosarcoma										X															
Nervous System																									
Brain	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Respiratory System																									
Larynx	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Lung	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Alveolar/bronchiolar adenoma																					X		X		
Alveolar/bronchiolar carcinoma					X			v																	
Carcinoma, metastatic, mammary gland Hepatocellular carcinoma, metastatic, liver								X				X													
Osteosarcoma, metastatic, tissue NOS			X									21													
Sarcoma, metastatic, uterus		X																							
Nose	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Trachea	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Special Senses System None																									
Urinary System																									
Kidney	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Cholangiocarcinoma, metastatic, liver					X																				
Ureter							+																		
Urinary bladder	+	+	+	+	+	+	+	+	+	+	+	+	+	M	+	+	+	+	+	+	+	+	M	+	+
Serosa, sarcoma, metastatic, uterus		X																							
Systemic Lesions																									
Multiple organs	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Lymphoma malignant	X			X			X				X														

Individual Animal Tumor Pathology of	rema	ue	IVI	ıce	ın	th	e Z	- Y (ear	ın	na	ıat	ion	St	ud	y o	1 1	th	yib	en	zeı	1e:	7	o p	pm	(continue
Number of Days on Study	7 3 1	7 3 1	7 3 1	7 3 1	7 3 1	7 3 1	7 3 1	7 3 2	7 3 2	7 3 2	7 3 2	7 3 2	7 3 2	7 3 2	7 3 2	7 3 2	7 3 2	7 3 2	7 3 2	7 3 2	7 3 2	7 3 2	7 3 2	7 3 2	7 3 2	
Carcass ID Number	1 8 6	1 8 9	1 9 3	1 9 5	1 9 8	1 9 9	2 0 0	1 5 6	1 5 8	1 5 9	1 6 2	1 6 3	1 6 4	1 6 5	1 6 8	1 7 3	1 7 5	1 7 9	1 8 0	1 8 1	1 8 2	1 9 0	1 9 1	1 9 2	1 9 7	Total Tissues/ Tumors
Hematopoietic System Bone marrow Lymph node Iliac, hemangioma Lumbar, osteosarcoma, metastatic,	+	+	+	+	+	+	+	+	+	+	+ + X	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50 7 1
tissue NOS Renal, hemangiosarcoma Lymph node, bronchial Lymph node, mandibular Lymph node, mesenteric Lymph node, mediastinal Spleen Thymus	M + + + +	+ + + + + +	+ + + + N	+ + + + +	+ + + + +	+ + + + + +	+ + + + + +	+ + + + + +	+ + + + + +	+ + + + + +	X + + + +	+ + + + + +	M + + +	+ + + + + +	+ + + + + +	+ + + + + +	M + + +	+ + + + + +	+ + M + +	+	+ + + + + +	+ + + + + +	+ + + + + +	M + + + +	+ + + M +	1 40 48 48 42 50 44
ntegumentary System Mammary gland Carcinoma ikin Fibrosarcoma Hemangioma Squamous cell carcinoma	+	+	+ X		+	+	+ + X	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50 3 50 2 1 1
Musculoskeletal System Bone Skeletal muscle Carcinoma, metastatic, mammary gland Rhabdomyosarcoma	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50 2 1 1
Nervous System Brain	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Respiratory System Larynx Lung Alveolar/bronchiolar adenoma Alveolar/bronchiolar carcinoma Carcinoma, metastatic, mammary gland Hepatocellular carcinoma, metastatic, liver Osteosarcoma, metastatic, tissue NOS Sarcoma, metastatic, uterus Nose Trachea	+ + + +	+ + +	+++++	++++	+ + + + + + + + + + + + + + + + + + + +	+ + + + + + + + + + + + + + + + + + + +	+ + + +	+ + + +	+ + + +	+ + + +	+ + + +	+ + X	+ + + + + + + + + + + + + + + + + + + +	+ + + +	+ + + +	+ + + +	+ + + + +	M + X X + + + +	+ + + +	+ + + +	+ + + +	+ + + +	+ + X	+ + + +	+ + + +	49 50 4 2 1 1 1 1 50
Special Senses System None																										
Urinary System Kidney Cholangiocarcinoma, metastatic, liver Jreter Jrinary bladder Serosa, sarcoma, metastatic, uterus	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50 1 1 48 1
Systemic Lesions Multiple organs Lymphoma malignant	+	+ X	+	+	+	+	+ X	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50 6

	0	4	4	6	6	6	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7
Number of Days on Study	8	3	4	5	8	9	0	0	0	0	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3
	6	7	6	9	2	4	1	5	7	8	0	0	0	0	1	1	1	1	1	1	1	1	1	1	1
	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2
Carcass ID Number	6	7	6	9	7	9	8	8	8	7	7	7	8	9	5	5	5	5	5	5	6	6	6	7	7
	8	9		4			5												8						
limentary System																									
Esophagus	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Gallbladder							A						+	+	+	+	+	+	+	+	+	+	+	+	+
ntestine large, colon			+				+					+	+	+		+	+	+	+	+	+	+	+	+	+
ntestine large, rectum			+							+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
ntestine large, cecum							+					+	+	+		+	+	+	+	+	+	+	+	+	+
ntestine small, duodenum	A	+	+	+	A	+	A	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Polyp adenomatous																					X				
ntestine small, jejunum	A	+	+	+			M					+	+	+	+	+	+		+		+	+	+	+	+
ntestine small, ileum	A	+	+	+	+		A				+	+	+	+			+			+	+	+	+	+	+
Liver Hanatacallular carcinoma	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+ X	+	+	+	+	+	+	+	+	+	+
Hepatocellular carcinoma Hepatocellular adenoma				X				X							Λ							X			Х
Hepatocellular adenoma, multiple				Λ	X			Λ											X			Λ			Λ
Pancreas	_	_		+	+	+	+	+	+	+	+	+	_	+	+	+	+	+	+	_	_	_	_	+	+
Salivary glands		4	+	+	+	+	+		+	+		+	+	+	+	+	+	+	+	+	+	+	+	+	
Stomach, forestomach	+	+	. +	+	+	+			À					+		+	+	+	+	+	+	+	+	+	+
Squamous cell papilloma						Ċ	11															•	Ċ		
Stomach, glandular	+	+	+	+	+	+	Α	+	Α	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
-																									
Cardiovascular System Blood vessel	M	[+			м	+	+																		
Heart	101	۱ +	+	+				+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
icui t	'				'											_		_			_				
Endocrine System																									
Adrenal cortex	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Adenoma																			X						
Adrenal medulla	+	+	+			+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Pheochromocytoma malignant				X																					
Pheochromocytoma benign																					X				
slets, pancreatic	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Adenoma		3	r .		1.4					ъ.	1 . #					1 . 1	1.1		1 1		1. /			1.	
Parathyroid gland	+						+																		
Pituitary gland Pars distalis, adenoma	+	+	+	+ X		+	M	+ X		+	+ X	+	+	+	+	+	+ X	+	+	+	+	+	+	IV	+
Pars distalls, adenoma Pars intermedia, adenoma				Λ				Λ			Λ			X			Λ								
Fars intermedia, adenoma Fhyroid gland		J		_	_		+	_	_	_	_	_	_		_	_	_	_	_	_		_	_	_	_
Follicular cell, adenoma	+	+	+	+	+ X		+	+	+	+	+	+	+	_	_	_	_	т	_	+	+	+	+	+	_
					/1																				
General Body System																									
Cissue NOS			+								+													+	
Pelvic, sarcoma			X																						
enital System																									
Clitoral gland	+	+	+	+	+	+	+	+	+	M	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Ovary	+	+	+	+	+	M	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Uterus	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Polyp stromal																									
Myometrium, hemangioma				X																					

Individual Animal Tumor Patholog	5 01 1 0111														`	_			_							
	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	
Number of Days on Study	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	
•	1	1	. 1	1	1	1	1	1	1	1	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	
	2	2	2	2	2	2	2	2	2	3	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	Total
Carcass ID Number	7	8	8	8	8	9	9	9	9	0	5	5	5	6	6	6	6	6	7	7	8	8	9	9	9	Tissues/
	8	1	. 3	8	9	0	1	2															5	6		Tumors
Alimentary System																										
Esophagus	+	- 4	+ +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Gallbladder	+	+	- +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	44
Intestine large, colon	+	+	+ +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	48
Intestine large, rectum	+	+	+ +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	49
Intestine large, cecum	+	+	- +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	48
Intestine small, duodenum	+	+	- +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	47
Polyp adenomatous																										1
Intestine small, jejunum	+	+	- +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	46
Intestine small, ileum	+	+	+ +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	47
Liver	+	+	+ +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Hepatocellular carcinoma														X								X				3
Hepatocellular adenoma																	X	X		X			X	X		9
Hepatocellular adenoma, multiple												X														3
Pancreas	+	+	- +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Salivary glands	+	+	+ +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Stomach, forestomach	+	+	+ +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	48
Squamous cell papilloma																						X				1
Stomach, glandular	+	_	- +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	48
Cardiovascular System																										
Blood vessel	+	+	- +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	48
Heart	+	+	- +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Endocrine System																										
Adrenal cortex	+	+	+ +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Adenoma																										1
Adrenal medulla	+	+	+ +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Pheochromocytoma malignant																										1
Pheochromocytoma benign																										1
Islets, pancreatic	+	+	- +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Adenoma	1	•		1	ıπ	1	r .		N.f	1 A				1.1							X					1
Parathyroid gland	N	1 -	+ +				+ 1					+	+	M		+	+	+	+		M		+	+	+	34
Pituitary gland	+	-	- +			+ X		+	+	+	IVI	+	+	+	+	+	+	+	+	+ X	+	+	+	+	+	47
Pars distalis, adenoma Pars intermedia, adenoma				Σ	L	Λ														Λ						7 1
Γhyroid gland Follicular cell, adenoma	+	+	- +	- +	· +	. +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50 3
																										ა
General Body System																										
Fissue NOS																			+							4
Pelvic, sarcoma																										1
Genital System																										
Clitoral gland	+	+	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	M	+	+	+	+	+	+	+	48
Ovary	+	+	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	49
Uterus	+	+	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Polyp stromal																									X	1
Myometrium, hemangioma																										1

TABLE D2 Individual Animal Tumor Pathology of	emale Mice in th	e 2-Year Inhalation Study of Ethylbenzene: 2	250 ppm (continued)
Number of Days on Study	0 4 4 6 6 6 8 3 4 5 8 9 6 7 6 9 2 4	0 0 0 0 3 3 3 3 3 3 3 3 3 3 3 3 3	7 7 3 3 1 1
Carcass ID Number	2 2 2 2 2 2 6 7 6 9 7 9 8 9 0 4 4 3		7 7
Hematopoietic System Bone marrow Lymph node Lymph node, bronchial	+ + + + + + + + + + M + + + M	1 + + + + + + + + + + + + + + + + + + +	· + + 1 + +
Lymph node, mandibular Lymph node, mesenteric Lymph node, mediastinal Spleen Fhymus	M + + + + + + + M + + + + + M + + + + +	1 + + + + + + + + + + M + + M + M + + + + + + + + + + + + + + + + + + + +	+ + + + + + + + M
Integumentary System Mammary gland Skin Hemangioma Sebaceous gland, adenoma	M + + M + + M + + + + +		+ + +
Musculoskeletal System Bone Rib, sarcoma, metastatic, tissue NOS Vertebra, osteosarcoma	+ + + + + + + + X	+++++++++++++++	+ +
Nervous System Brain	+ + + + + +	. + + + + + + + + + + + + + + + +	+ +
Respiratory System Larynx Lung Alveolar/bronchiolar adenoma Alveolar/bronchiolar adenoma, multiple	+ M A + + + M + + + + + X	X X X	+ + +
Hepatocellular carcinoma, metastatic, liver Sarcoma, metastatic, tissue NOS Nose Frachea	X + + + + + + + + + + + + +	X + + + + + + + + + + + + + + + + + + +	· + + · + +
Special Senses System Harderian gland Adenoma			
Urinary System Kidney Urinary bladder	+ + + + + + + + + + + + + + + + + + + +	+ + + + + + + + + + + + + + + + + + +	- + + - M +
Systemic Lesions Multiple organs Lymphoma malignant	+ + + + + + + + + X X X	+ + + + + + + + + + + + + + + + + + +	+ +

Individual Animal Tumor Pathology o	f Fem	ale	M	ice	in	the	2-	Ye	ar	In	hal	lati	ion	St	ud	y o	f F	Eth	ylb	en	zei	ıe:	2	50	ppn	1 (continue
Number of Days on Study	7 3 1	7 3 1	7 3 1	7 3 1	7 3 1	7 3 1	7 3 1	7 3 1	7 3 1	7 3 1	7 3 2	7 3 2	7 3 2	7 3 2	7 3 2	7 3 2	7 3 2	7 3 2	7 3 2	7 3 2	7 3 2	7 3 2	7 3 2	7 3 2	7 3 2	
Carcass ID Number	2 7 8	2 8 1	2 8 3	2 8 8	2 8 9	2 9 0	2 9 1	2 9 2	2 9 7	3 0 0	2 5 3	2 5 4	2 5 7	2 6 1	2 6 2	2 6 3	2 6 4	2 6 6	2 7 0	2 7 5	2 8 2	2 8 7	2 9 5	2 9 6	2 9 8	Total Tissues/ Tumors
Hematopoietic System Bone marrow Lymph node Lymph node, bronchial Lymph node, mandibular Lymph node, mesenteric Lymph node, mediastinal Spleen Thymus	+ + + + + + + +	+ M + + +	+ I M + + + +	+ + + M + +	+	+ + + + + M	+ + +	+ + + +	M + +		+ + +	+ + M + +	+	+		+ + + + + + +	+		M +	+	+ M + + +	+ + + M + +	+	+ M + + +	+ + + + + + + +	50 2 29 47 46 41 50 45
Integumentary System Mammary gland Skin Hemangioma Sebaceous gland, adenoma	+	+	+	+	+	+	+	+	+ + X	+	+	+	+	+ + X	+	+	+	+	+	+	+	+	+	+ + X	+++	48 49 2 1
Musculoskeletal System Bone Rib, sarcoma, metastatic, tissue NOS Vertebra, osteosarcoma	+	+	+	+	+	+	+	+	+	+	+	+	+	+ X	+	+	+	+	+	+	+	+	+	+	+	50 1 1
Nervous System Brain	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Respiratory System Larynx Lung Alveolar/bronchiolar adenoma Alveolar/bronchiolar adenoma, multiple Hepatocellular carcinoma, metastatic, liver Sarcoma, metastatic, tissue NOS Nose Frachea	++++	+ + + +	+ + + +	+ + + + +	+ + + +	+ + + +	+ + + + +	+ + + +	+ + + +	+ + + +	+ + + +	+ + + +	+ + + +	+ + + +	+ + + + +	+ + + +	+ + + +	+ + + +	+ + X + +	+ + + + +	+ + + +	+ + X + +	+ + + +	+ + + +	+ + + +	47 49 4 1 2 1 50
Special Senses System Harderian gland Adenoma																				+ X						1 1
Urinary System Kidney Urinary bladder	+	+	+	+	+	++	+	++	++	++	+	++	++	++	++	++	++	+	+	+	++	+	++	+	+	50 47
Systemic Lesions Multiple organs Lymphoma malignant	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+ X	+	+	+	+	+	50 5

							_		_										-		_	_		_
		1		6	6	6	-	7 7	7	7	7	7		7		7			7		7		7	
Number of Days on Study	0	6	6	1	1	6	-	1 1			1	2	3	3	3	3	3	3	3	3	3	3	3	3
	- /	9	8	2	4	6	1	1 2	2 2	3	8	3	0	0	0	1	1	1	1	1	1	1	1	1
	3	3	3	3	3	4	3	3 3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3
Carcass ID Number	7	8	8	5	6	0	5	6 5	6	7	6	9	5	9	9	5	5	6	7	7	7	7	8	8
	2	0	4	9	3	0	6	1 8	3 9	0	0	6	1	3	5	4	5	2	5	7	8	9	2	3
Alimentary System																								
Esophagus	+	+	+	+	+	+	+	+ -	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Gallbladder	+	Α	+	+	+	Α	M	+ -	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Intestine large, colon	+	+	+	+	+	+	+	+ -	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Intestine large, rectum	+	Α	+	Α	+	+	Α	+ -	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Intestine large, cecum	N.	[A	M	Α	+	Α	Α	+ -	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Intestine small, duodenum	+	Α	+	Α	+	Α	M	+ -	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Intestine small, jejunum	A	N	1 +	Α	+	Α	+	+ -	+ +	+	+	Α	+	+	+	+	+	+	+	+	+	+	+	+
Intestine small, ileum	+	Α	. +	Α	+	+	A	+ -	+ +	+	Α	+	+	+	+	+	+	+	+	+	+	+	+	+
Liver	+	+	+	+	+	+	+	+ -	+ +		+	+	+	+	+	+	+	+	+	+	+	+	+	+
Hepatocellular carcinoma											X	X					X	X						X
Hepatocellular carcinoma, multiple				X					Σ															
Hepatocellular adenoma					X									X			X			X				
Hepatocellular adenoma, multiple															X									
Pancreas	+	+	+	+	+	+	A	+ -	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Salivary glands	+	+	+	+	+	+	+	+ -	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Stomach, forestomach	+	+	+	+	+	+	+	+ -	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Squamous cell papilloma																								
Stomach, glandular	+	+	+	+	+	+	+	+ -	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Cardiovascular System																								
Blood vessel	+	+	+	+	+	+	+	+ -	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Heart	+	+	+	+	+	+	+	+ -	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Endocrine System																								
Adrenal cortex		د			_	_	_	_	ا. ي		_	_	_	_	_	_	_	_	_	_	_	_	_	_
Adrenal cortex Adrenal medulla	+ +			⊤	+	+	+	+ -	. T	. ⊥	⊤	±	+	+	+	+	+	+	+	+	+	_	+	+
Islets, pancreatic	+ +			⊤	+	+	Ā	+ -	+ +	+	+	+	+	+	+	+	+	+	+	+	+	_	+	+
Parathyroid gland	+	+		+	M	+		т - М -				M						+ M	M	М	+	+	+	+
Pituitary gland	+	+	+	M		+			 		+	+	+	+		+		+	+	+	+	+	+	+
Pars distalis, adenoma	'			.,1	Ċ		•		. '		Ċ				•	•	•	•	X	•		X		-
Thyroid gland	+	+	+	+	+	+	+	+ -	+ +	+	+	+	+	+	+	+	+	+		+		+	+	+
Follicular cell, adenoma			•		·	-		•			·	•	X		X		•	-		-			•	
General Body System Fissue NOS																								
Genital System																								
Clitoral gland	+	+	+	+	+	+	+	М -	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Fibrosarcoma	'		X		Ċ		•		. '		Ċ				•	•	•	•		•	•			-
Ovary	+	+	+	+	+	+	+	+ -	+ +	+	+	M	+	+	+	+	+	+	+	+	+	+	+	+
Cystadenoma					·																			
Uterus	+	+	+	+	+	+	+	+ -		+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
T																								
Hematopoietic System																					,			
Bone marrow	+	+	+	+	+	+	+	+ -	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Lymph node	3.7	+	r .	1.0	3.4	+	+				1 A	1.7						+ 1.4			1 4	,		
Lymph node, bronchial					M			+ -	+ +			M	+	+	+	+	+	M	+	+	M	+	+	+
Lymph node, mandibular			1 + M	+	+	+		+ -	- + 	· M		+	+	+	+	+	+	+	+	+	+	+	+	+
Lymph node, mesenteric	_		M				M		+ + √1 N			+	+ 1\1	+	+	+	+	+	+	+	+	+	+	
Lymph node, mediastinal Spleen	IV.	ιIV	1 M	. +	M	+	+	M I		1 +	ıVI		M +	+	+	+	+	+	+	+	+	+	+	+
	+	+	+	+	+	+	+				1.1	+ M		+	+	+	+	+	+	+	+	+	+	+
Thymus	+	_	-	+	+	_	_	-T -	r (\	1 +	171	101	_	_										

TARLE D2

Individual Animal Tumor Pathology	of Fem	ale	M	ice	in	the	2-	Ye	ar	In	hal	lati	ion	St	uď	y o	f E	ith	ylb	en	zei	1e:	7	50	ppr	n (continue
	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	
Number of Days on Study	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	
Ç Ç	1	1	1	1	1	1	1	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	
	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	Total
Carcass ID Number	8	8	8	8	9	9	9	5	5	5	6	6	6	6	6	7	7	7	7	8	8	9	9	9	9	Tissues/
	5	6	7	8	1	4	9	2	3	7	4	5	6	7	8	1	3	4	6	1	9	0	2	7	8	Tumors
Alimentary System																										
Esophagus	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Gallbladder	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	M	+	+	+	+	+	+	+	+	46
Intestine large, colon	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Intestine large, rectum	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	47
Intestine large, cecum	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	44
ntestine small, duodenum	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	46
ntestine small, jejunum	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	45
ntestine small, ileum	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	46
Liver	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Hepatocellular carcinoma			X								X						X							X		10 2
Hepatocellular carcinoma, multiple	v			v	v								v				v				v	v		v		
Hepatocellular adenoma Hepatocellular adenoma, multiple	X			Λ	X					X			X			X	X				Λ	X	X	X		12 4
Pancreas					_	ر	_		+	+	+	+	_	_	+	Λ +	+	_	_			_	Λ.	.,	_	49
Salivary glands		+	+	+	+	+	T +	⊤	+	+	+	+	+	T _	⊤	+	+	+	+	+	+	+	+		⊤	50
Stomach, forestomach						_		_	_			_		т _	_	_	_	_	_							50
Squamous cell papilloma							_	_		-		_		-	_	_		-	X		-				-	1
Stomach, glandular	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+		+	+	+	+	+	+	50
S1																							_			
Cardiovascular System																										70
Blood vessel Heart	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50 50
reart	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+		+	+	30
Endocrine System																										
Adrenal cortex	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	M	49
Adrenal medulla	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	M	49
slets, pancreatic	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	49
Parathyroid gland	+	N	1 M	M	M	+	M	+	+	+	+		M	M	+	+	+	+	M	M	M	M	M		M	27
Pituitary gland	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	49
Pars distalis, adenoma										X															X	5
Гhyroid gland Follicular cell, adenoma	+	+	+	+	+	+	+	+	+	+	+ X	+	+	+	+	+ X	+	+	+	+	+	+	+	+	+	50
roniculai cen, adenonia											Λ					Λ										4
General Body System Cissue NOS				+																						1
Genital System																							_	_		
Clitoral gland	+	+	+	+	+	+	+	+	+	M	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	48
Fibrosarcoma	·	•	•	•	·	•						•		-	-		•				•	•	•			1
Ovary	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	49
Cystadenoma														X												2
Jterus	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Iematopoietic System																							_			
Bone marrow	+	_			_	_	_	_	_	4	_	_	_	_	_	_	_	_	_	_	_	4	_		_	50
Lymph node	+	_	_	т	-		7	7	-	-	-	-	7	7-	7	7		-r	7	-	-	-	-		7	5
Lymph node Lymph node, bronchial	+	+	+	+	+	+	+	+	+	+	М	М	M	+	+	+	+	M	+	+	+	+	+	+	+	38
Lymph node, mandibular	∓	+	M	+	+	+	M	+	M		+	+	+	+	+	+	+	M		+	+	+	+	+	+	44
Lymph node, mesenteric		+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	M		+	+	+	+	+	+	44
Lymph node, mediastinal	+	N	1 M	+	+	+	+	М	M	+	+		M				M		M	+	M	+	+	+	+	31
Spleen	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	49

Individual Animal Tumor Pathology of																		_						
Number of Days on Study	-	1	5		6	6						7		-	•	-	7	•		7	7	7		7 3
Number of Days on Study	0 7	6 9	6 8		1 4	6 6			1 2			2	3 0	3 0	3 0	3 1	3 1	3 1	3 1	3 1	3 1	1	3 1	1
	3	3	3	3	3	4			3 3	3 3	3		3	3	3	3	3	3	3	3	3	3	3	3
Carcass ID Number	7 2	8	8 4	5 9	6 3	0	5 6		5 (8 (5 1	9	9 5	5 4	5 5	6 2	7 5	7 7	7 8	7 9	8 2	8
Integumentary System																								
Mammary gland Skin	+	A +	. +	+	+	+	+	+ +	+ -	+ +	- + - +	+	+	+	+	+	+	+	+	+	+	+	+	+
Fibroma	'								'			'	X			'			'					,
Fibrous histiocytoma																				X				
Musculoskeletal System Bone																								
	+	+	+	+	+	+	+	+	+ -	+ +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+
Nervous System Brain	+	+	+	+	+	+	+	+	+ -	+ +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+
Respiratory System																								
Larynx	+	M	1 +	+	+	+	+	+	+ -	+ +	+	+	+	+	+	M	+	+	+	+	+	+	+	+
Lung	+	+	+	+	+	+	+	+	+ -	+ +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+
Alveolar/bronchiolar adenoma Hepatocellular carcinoma, metastatic, liver												X	X					X					X	
Squamous cell carcinoma, metastatic,												71												
lacrimal gland											X													
Nose	+	+	+	+	+	+	+	+	+ -	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Trachea	+	+	+	+	+	+	+	+	+ -	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Special Senses System																								
Harderian gland Adenoma																						+ X		
Lacrimal gland											+													
Squamous cell carcinoma											X													
Urinary System																								
Kidney	+	+	+	+	+	+	+	+	+ -	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Ureter Urinary bladder	+	A	. +	+	+	+	+	+	+ -	+ +	- +	+	+	+	+	+	+	+	+	+	+	+	+	+
Systemic Lesions																								
Multiple organs	+	+	+	+	+	+	+	+	+ -	+ +	+	+	+	+	+	+	+	+	+	+	+	+	+	+
Lymphoma malignant						X	X											X	X					

Individual Animal Tumor Pathology o	f Fem	al	e N	1ic	e i	n t	he	2-`	Ye	ar	In	ha	lati	ion	St	ud	y (f I	Eth	yll	en	zei	ne:	7	50	ppr	n (continued
Number of Days on Study	7 3 1		3 :	3	7 3 1	7 3 1			7 3 2																		
Carcass ID Number	3 8 5	8	3 8		8		9	9	5	3 5 3	3 5 7	3 6 4	3 6 5	3 6 6	3 6 7	3 6 8	3 7 1	3 7 3	3 7 4	3 7 6	3 8 1	3 8 9	3 9 0	3 9 2	3 9 7	3 9 8	Total Tissues/ Tumors
Integumentary System Mammary gland Skin Fibroma Fibrous histiocytoma	+		+ -	+ +	+ +	+ +	+ +	+ +	+	+ +	+	+++	+	+	++	+++	+	+	+	+	+	+	+	+	+	+	49 50 1 1
Musculoskeletal System Bone	+		+ -	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Nervous System Brain	+		+ -	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50
Respiratory System Larynx Lung Alveolar/bronchiolar adenoma Hepatocellular carcinoma, metastatic, liver	+		+ -		+ + X	+ +	+ +	+	+	+ +	+	+ + X	+	+++	+	+	+	+ + X	+	+	+	+ + X	+	+ + X	+	+	48 50 8 1
Squamous cell carcinoma, metastatic, lacrimal gland Nose Frachea	+		+ -	+	+	+	+	++	+	++	+	++	+	++	++	++	+	++	++	++	++	+	+	+	++	++	1 50 50
Special Senses System Harderian gland Adenoma Lacrimal gland Squamous cell carcinoma																		+ X			+ X						3 3 1 1
U rinary System Kidney Jreter Jrinary bladder	+		+ -	+	+		+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50 1 49
Systemic Lesions Vultiple organs Lymphoma malignant	+ X		- -	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	50 5

TABLE D3
Statistical Analysis of Primary Neoplasms in Female Mice in the 2-Year Inhalation Study of Ethylbenzene

farderian Gland: Adenoma verall rate ^a djusted rate ^b erminal rate ^c irst incidence (days) ife table test ^d ogistic regression test ^d ochran-Armitage test ^d isher exact test ^d farderian Gland: Adenoma or Carcinoma verall rate djusted rate	0/50 (0%) 0.0% 0/35 (0%)e P= 0.021 P= 0.021 P= 0.021	0/50 (0%) 0.0% 0/38 (0%) 	1/50 (2%) 2.5% 1/40 (3%) 730 (T) P= 0.527 P= 0.527 P= 0.500	3/50 (6%) 8.1% 3/37 (8%) 730 (T) P= 0.131 P= 0.121
verall rate ^a djusted rate ^b erminal rate ^c first incidence (days) fife table test ^d ochran-Armitage test ^d fisher exact test ^d fisher exact test ^d	0.0% 0/35 (0%) —e P= 0.021 P= 0.021 P= 0.021	0.0% 0/38 (0%)	2.5% 1/40 (3%) 730 (T) P= 0.527 P= 0.527	8.1% 3/37 (8%) 730 (T) P= 0.131 P= 0.131
djusted rate ^b erminal rate ^c irst incidence (days) ife table test ^d ogistic regression test ^d ochran-Armitage test ^d isher exact test ^d irst incidence (days) ife table test ^d ochran-Armitage test ^d isher exact test ^d isher exact test ^d isher according to the company of the compan	0.0% 0/35 (0%) —e P= 0.021 P= 0.021 P= 0.021	0.0% 0/38 (0%)	2.5% 1/40 (3%) 730 (T) P= 0.527 P= 0.527	8.1% 3/37 (8%) 730 (T) P= 0.131 P= 0.131
erminal rate ^c irst incidence (days) ife table test ^d ogistic regression test ^d ochran-Armitage test ^d isher exact test ^d iarderian Gland: Adenoma or Carcinoma verall rate		_ ` '	1/40 (3%) 730 (T) P= 0.527 P= 0.527	730 (T) P= 0.131 P= 0.131
irst incidence (days) ife table test ^d ogistic regression test ^d ochran-Armitage test ^d isher exact test ^d iarderian Gland: Adenoma or Carcinoma verall rate		_ ` '	730 (T) P= 0.527 P= 0.527	730 (T) P= 0.131 P= 0.131
ife table test ^d ogistic regression test ^d ochran-Armitage test ^d isher exact test ^d iarderian Gland: Adenoma or Carcinoma verall rate	P = 0.021 P = 0.021	_f _ _	P= 0.527 P= 0.527	P= 0.131 P= 0.131
ogistic regression test ^d ochran-Armitage test ^d isher exact test ^d f arderian Gland: Adenoma or Carcinoma verall rate	P = 0.021 P = 0.021	_	P= 0.527	P = 0.131
ochran-Armitage test ^d isher exact test ^d f arderian Gland: Adenoma or Carcinoma verall rate		_		P= 0.121
isher exact test ^d (arderian Gland: Adenoma or Carcinoma verall rate		_	P = 0.500	P = 0.121
verall rate	1/50 (2%)			
verall rate	1/50 (2%)			
	· - \	0/50 (0%)	1/50 (2%)	3/50 (6%)
	2.4%	0.0%	2.5%	8.1%
erminal rate	0/35 (0%)	0/38 (0%)	1/40 (3%)	3/37 (8%)
irst incidence (days)	639		730 (T)	730 (T)
ife table test	P = 0.083	P = 0.486N	P = 0.730N	P = 0.330
ogistic regression test	P = 0.081	P = 0.516N	P = 0.761	P = 0.310
ochran-Armitage test	P = 0.080			
isher exact test		P = 0.500N	P = 0.753N	P = 0.309
iver: Hepatocellular Adenoma				
verall rate	6/50 (12%)	9/50 (18%)	12/50 (24%)	16/50 (32%)
djusted rate	17.1%	22.1%	27.5%	41.8%
erminal rate	6/35 (17%)	7/38 (18%)	9/40 (23%)	15/37 (41%)
irst incidence (days)	730 (T)	562	659	614
ife table test	P = 0.013	P = 0.345	P = 0.165	P = 0.018
ogistic regression test	P = 0.014	P = 0.311	P = 0.128	P = 0.022
ochran-Armitage test	P = 0.011			
sher exact test		P = 0.288	P = 0.096	P = 0.014
iver: Hepatocellular Carcinoma				
verall rate	7/50 (14%)	4/50 (8%)	3/50 (6%)	12/50 (24%)
djusted rate	17.3%	9.7%	7.5%	28.3%
erminal rate	3/35 (9%)	2/38 (5%)	3/40 (8%)	7/37 (19%)
irst incidence (days)	565	602	730 (T)	612
ife table test	P = 0.029	P = 0.238N	P = 0.127N	P = 0.205
ogistic regression test	P = 0.022	P = 0.259N	P = 0.150N	P = 0.162
ochran-Armitage test	P = 0.022			
isher exact test		P = 0.262N	P = 0.159N	P = 0.154
iver: Hepatocellular Adenoma or Carcinoma				
verall rate	13/50 (26%)	12/50 (24%)	15/50 (30%)	25/50 (50%)
djusted rate	32.8%	28.2%	34.5%	57.9%
erminal rate	9/35 (26%)	8/38 (21%)	12/40 (30%)	19/37 (51%)
irst incidence (days)	565	562	659	612
ife table test	P = 0.004	P = 0.426N	P = 0.562	P = 0.029
ogistic regression test	P = 0.002	P = 0.420N	P = 0.302 P = 0.471	P = 0.025
ochran-Armitage test	P = 0.002	1 - 0.17011	1-0.111	1 - 0.010
isher exact test	1 – 0.002	P = 0.500N	P = 0.412	P = 0.011

TABLE D3
Statistical Analysis of Primary Neoplasms in Female Mice in the 2-Year Inhalation Study of Ethylbenzene (continued)

	Chamber Control	75 ppm	250 ppm	750 ppm
Lung: Alveolar/bronchiolar Adenoma				
Overall rate	4/50 (8%)	4/50 (8%)	5/49 (10%)	8/50 (16%)
Adjusted rate	10.9%	10.5%	11.4%	21.6%
Terminal rate	3/35 (9%)	4/38 (11%)	2/40 (5%)	8/37 (22%)
First incidence (days)	694	730 (T)	682	730 (T)
Life table test	P = 0.106	P = 0.598N	P = 0.579	P = 0.206
Logistic regression test	P = 0.111	P = 0.618N	P = 0.525	P = 0.218
Cochran-Armitage test	P = 0.096			
Fisher exact test		P = 0.643N	P = 0.487	P = 0.178
Lung: Alveolar/bronchiolar Adenoma or Carci	noma			
Overall rate	4/50 (8%)	6/50 (12%)	5/49 (10%)	8/50 (16%)
Adjusted rate	10.9%	15.0%	11.4%	21.6%
Terminal rate	3/35 (9%)	5/38 (13%)	2/40 (5%)	8/37 (22%)
First incidence (days)	694	590	682	730 (T)
Life table test	P = 0.184	P = 0.419	P = 0.579	P = 0.206
Logistic regression test	P = 0.181	P = 0.386	P = 0.525	P = 0.218
Cochran-Armitage test	P = 0.169			
Fisher exact test		P = 0.370	P = 0.487	P = 0.178
Mammary Gland: Carcinoma				
Overall rate	1/50 (2%)	3/50 (6%)	0/50 (0%)	0/50 (0%)
Adjusted rate	2.9%	7.3%	0.0%	0.0%
Terminal rate	1/35 (3%)	1/38 (3%)	0/40 (0%)	0/37 (0%)
First incidence (days)	730 (T)	642	_ ` `	_ ` '
Life table test	P = 0.144N	P = 0.336	P = 0.473N	P = 0.489N
Logistic regression test	P = 0.149N	P = 0.311	P = 0.473N	P = 0.489N
Cochran-Armitage test	P = 0.150N			
Fisher exact test		P = 0.309	P = 0.500N	P = 0.500N
Pituitary Gland (Pars Distalis): Adenoma				
Overall rate	4/48 (8%)	8/49 (16%)	7/47 (15%)	5/49 (10%)
Adjusted rate	11.1%	19.8%	17.0%	13.5%
Terminal rate	3/35 (9%)	6/38 (16%)	5/38 (13%)	5/37 (14%)
First incidence (days)	725	697	659	730 (T)
Life table test	P = 0.445N	P = 0.223	P = 0.322	P = 0.533
Logistic regression test	P = 0.425N	P = 0.203	P = 0.276	P = 0.542
Cochran-Armitage test	P = 0.459N			
Fisher exact test		P = 0.188	P = 0.249	P = 0.513
Thyroid Gland (Follicular Cell): Adenoma				
Overall rate	5/50 (10%)	6/50 (12%)	3/50 (6%)	4/50 (8%)
Adjusted rate	13.7%	15.2%	7.1%	10.8%
Terminal rate	4/35 (11%)	5/38 (13%)	2/40 (5%)	4/37 (11%)
First incidence (days)	694	697	682	730 (T)
Life table test	P = 0.355N	P = 0.557	P = 0.288N	P = 0.462N
Logistic regression test	P = 0.339N	P = 0.532	P = 0.320N	P = 0.449N
Cochran-Armitage test	P = 0.372N			
Fisher exact test		P = 0.500	P = 0.357N	P = 0.500N

TABLE D3
Statistical Analysis of Primary Neoplasms in Female Mice in the 2-Year Inhalation Study of Ethylbenzene (continued)

	Chamber Control	75 ppm	250 ppm	750 ppm
All Organs: Hemangioma				
Overall rate	0/50 (0%)	3/50 (6%)	3/50 (6%)	0/50 (0%)
Adjusted rate	0.0%	7.9%	7.0%	0.0%
Terminal rate	0/35 (0%)	3/38 (8%)	2/40 (5%)	0/37 (0%)
First incidence (days)	_ ` `	730 (T)	659	_ ` `
Life table test	P = 0.280N	P = 0.136	P = 0.149	_
Logistic regression test	P = 0.281N	P = 0.136	P = 0.123	_
Cochran-Armitage test	P = 0.291N			
Fisher exact test		P = 0.121	P = 0.121	_
All Organs: Hemangioma or Hemangiosarcoma				
Overall rate	0/50 (0%)	4/50 (8%)	3/50 (6%)	0/50 (0%)
Adjusted rate	0.0%	10.5%	7.0%	0.0%
Terminal rate	0/35 (0%)	4/38 (11%)	2/40 (5%)	0/37 (0%)
First incidence (days)	_	730 (T)	659	_
Life table test	P = 0.212N	P = 0.074	P = 0.149	_
Logistic regression test	P = 0.211N	P = 0.074	P = 0.123	_
Cochran-Armitage test	P = 0.222N	D 0.070	D 0 101	
Fisher exact test		P = 0.059	P = 0.121	_
All Organs: Malignant Lymphoma				
Overall rate	3/50 (6%)	6/50 (12%)	5/50 (10%)	5/50 (10%)
Adjusted rate	7.1%	13.4%	10.7%	12.2
Terminal rate	1/35 (3%)	2/38 (5%)	1/40 (3%)	3/37 (8%
First incidence (days)	423	300	437	666
Life table test	P = 0.496	P = 0.278 P = 0.175	P = 0.416	P= 0.393
Logistic regression test Cochran-Armitage test	P = 0.237 P = 0.469	r = 0.175	P = 0.340	P = 0.358
Fisher exact test	r = 0.405	P = 0.243	P = 0.357	P= 0.357
		1 0.210	1 0.00.	1 0,007
All Organs: Benign Neoplasms				
Overall rate	20/50 (40%)	26/50 (52%)	27/50 (54%)	28/50 (56%)
Adjusted rate	50.8%	63.2%	59.9%	71.6%
Terminal rate	16/35 (46%)	23/38 (61%)	22/40 (55%)	26/37 (70%)
First incidence (days)	565	562	659	614
Life table test	P= 0.163	P= 0.260	P= 0.284	P= 0.122
Logistic regression test	P= 0.176	P = 0.190	P = 0.163	P = 0.116
Cochran-Armitage test Fisher exact test	P = 0.132	P = 0.158	P = 0.115	P = 0.080
All Organs: Malignant Neoplasms Overall rate	13/50 (26%)	20/50 (40%)	9/50 (18%)	18/50 (36%)
Adjusted rate	28.5%	40.5%	19.3%	39.7%
Terminal rate	4/35 (11%)	9/38 (24%)	4/40 (10%)	10/37 (27%)
First incidence (days)	423	300	437	568
Life table test	P = 0.380	P = 0.181	P = 0.191N	P = 0.279
Logistic regression test	P = 0.075	P = 0.045	P = 0.265N	P = 0.192
Cochran-Armitage test	P = 0.322			
Fisher exact test		P = 0.101	P = 0.235N	P = 0.194

TABLE D3
Statistical Analysis of Primary Neoplasms in Female Mice in the 2-Year Inhalation Study of Ethylbenzene (continued)

	Chamber Control	75 ppm	250 ppm	75 0 ppm
All Organs: Benign or Malignant Neoplasms				
Overall rate	29/50 (58%)	38/50 (76%)	31/50 (62%)	38/50 (76%)
Adjusted rate	62.9%	76.0%	64.5%	80.8%
Terminal rate	18/35 (51%)	26/38 (68%)	23/40 (58%)	28/37 (76%)
First incidence (days)	423	300	437	568
Life table test	P = 0.229	P = 0.187	P = 0.466N	P = 0.161
Logistic regression test	P = 0.042	P = 0.041	P = 0.431	P = 0.045
Cochran-Armitage test	P = 0.116			
Fisher exact test		P = 0.044	P = 0.419	P = 0.044

(T)Terminal sacrifice

- Number of neoplasm-bearing animals/number of animals examined. Denominator is number of animals examined microscopically for liver, lung, pituitary gland, and thyroid gland; for other tissues, denominator is number of animals necropsied.
- b Kaplan-Meier estimated neoplasm incidence at the end of the study after adjustment for intercurrent mortality
- C Observed incidence at terminal kill
- Beneath the chamber control incidence are the P values associated with the trend test. Beneath the exposure group incidence are the P values corresponding to pairwise comparisons between the chamber controls and that exposure group. The life table test regards neoplasms in animals dying prior to terminal kill as being (directly or indirectly) the cause of death. The logistic regression test regards these lesions as nonfatal. The Cochran-Armitage and Fisher exact tests compare directly the overall incidence rates. For all tests, a negative trend or a lower incidence in an exposure group is indicated by N.
- Not applicable; no neoplasms in animal group
- Value of statistic cannot be computed.

TABLE D4a Historical Incidence of Alveolar/bronchiolar Neoplasms in Chamber Control Female B6C3F₁ Mice^a

		Incidence in Controls					
Study	Adenoma	Carcinoma	Adenoma or Carcinoma				
Historical Incidence at IIT Researc	h Institute						
Isobutyl Nitrite	4/51	2/51	6/51				
Overall Historical Incidence							
Total Standard deviation	61/939 (6.5%) 3.2%	38/939 (4.1%) 3.2%	97/939 (10.3%) 3.7%				
Range	0%-14%	0%-12%	0%-16%				

^a Data as of 12 May 1995

TABLE D4b Historical Incidence of Hepatocellular Neoplasms in Chamber Control Female B6C3F₁ Mice^a

	Incidence in Controls					
Study	Adenoma	Carcinoma	Adenoma or Carcinoma			
Historical Incidence at IIT Research	Institute					
Isobutyl Nitrite	6/51	4/51	10/51			
Overall Historical Incidence						
Total Standard deviation Range	114/937 (12.2%) 9.7% 0%-40%	103/937 (11.0%) 6.7% 0%-30%	200/937 (21.3%) 11.9% 3%-54%			

^a Data as of 12 May 1995

TABLE D5
Summary of the Incidence of Nonneoplastic Lesions in Female Mice in the 2-Year Inhalation Study of Ethylbenzene^a

	Chambe	r Control	75	5 ppm	250	ppm	75	0 ppm
Disposition Summary								
Animals initially in study	5	0		50		50		50
Carly deaths								
Accidental deaths		1				1		
Moribund		5		6		1		4
Natural deaths		9		6		8		9
urvivors								
Died last week of study		1		0.0		40		07
Terminal sacrifice	3	4		38		40		37
animals examined microscopically	5	0		50		50		50
Alimentary System								
Gallbladder	(44)		(44)		(44)		(46)	
Infiltration cellular		(2%)	,		\ -/		(•)	
ntestine small, duodenum	(45)		(48)		(47)		(46)	
Ulcer								(2%)
ntestine small, jejunum	(46)		(46)		(46)		(45)	
Peyer's patch, hyperplasia								(2%)
ntestine small, ileum	(47)		(47)	(00/)	(47)		(46)	
Peyer's patch, hyperplasia	(50)			(2%)	(50)		(50)	
iver Angiectasis	(50)		(50)	(2%)	(50)		(50)	
Basophilic focus	9	(6%)	1	(270)	1	(8%)	વ	(6%)
Clear cell focus		(2%)				(2%)	3	(0 /0)
Eosinophilic focus		(10%)	7	(14%)		(12%)	22	(44%)
Hemorrhage	Ū	(1070)	•	(11/0)		(2%)		(2%)
Hepatodiaphragmatic nodule						(4%)		())
Infiltration cellular	3	(6%)						
Inflammation	1	(2%)	3	(6%)	1	(2%)		(2%)
Mineralization								(2%)
Mixed cell focus						(2%)		(2%)
Necrosis		(2%)		(8%)	3	(6%)	4	(8%)
Pigmentation, hemosiderin	1	(2%)	1	(2%)				(00/)
Bile duct, cyst								(2%)
Hepatocyte, hypertrophy			1	(9%)			1	(2%)
Hepatocyte, necrosis Hepatocyte, syncytial alteration			1	(2%)	1	(2%)		
Hepatocyte, syncytial alteration Hepatocyte, vacuolization cytoplasmic	9	(4%)				(4%)	1	(2%)
Serosa, inflammation	۵	(1/0)			۵	(1/0)		(2%)
Mesentery	(2)						1	(~ /0)
Fat, necrosis	2	(100%)						
ancreas	(50)	,	(50)		(50)		(49)	
Angiectasis	, ,			(2%)	, ,		, ,	
Atrophy			2	(4%)				
Cyst				(2%)				
Degeneration				(2%)				
Fibrosis				(2%)				
Infiltration cellular	7	(14%)		(24%)	12	(24%)	10	(20%)
Necrosis		(00/)	1	(2%)				
Acinus, hyperplasia	1	(2%)						

^a Number of animals examined microscopically at the site and the number of animals with lesion

TABLE D5
Summary of the Incidence of Nonneoplastic Lesions in Female Mice in the 2-Year Inhalation Study of Ethylbenzene (continued)

	Chamber Control	75 ppm	250 ppm	750 ppm
Alimentary System (continued)				
Salivary glands	(50)	(50)	(50)	(50)
Atrophy		1 (2%)		
Infiltration cellular	30 (60%)	32 (64%)	33 (66%)	30 (60%)
Stomach, forestomach	(50)	(49)	(48)	(50)
Hyperplasia			1 (90/)	2 (4%)
Ulcer			1 (2%)	9 (40/)
Epithelium, cyst Epithelium, hyperplasia		2 (4%)		2 (4%) 1 (2%)
Stomach, glandular	(50)	(49)	(48)	(50)
Infiltration cellular	1 (2%)	(43)	(46)	(30)
Glands, cyst	1 (2%)			
Glands, hyperplasia	1 (270)			1 (2%)
Serosa, infiltration cellular				1 (2%)
Cardiovascular System				
Heart	(50)	(49)	(50)	(50)
Cardiomyopathy	10 (20%)	23 (47%)	23 (46%)	15 (30%)
Endocrine System				
Adrenal cortex	(47)	(50)	(50)	(49)
Accessory adrenal cortical nodule	2 (4%)	1 (2%)	(33)	()
Degeneration	12 (26%)	4 (8%)	4 (8%)	5 (10%)
Hemorrhage	2 (4%)	1 (2%)	2 (4%)	3 (6%)
Hyperplasia	3 (6%)	5 (10%)	3 (6%)	3 (6%)
Infiltration cellular	, ,	2 (4%)	, ,	`
Inflammation		3 (6%)		
Necrosis	1 (2%)			
Vacuolization cytoplasmic	1 (2%)			
Capsule, hyperplasia	46 (98%)	49 (98%)	48 (96%)	46 (94%)
Adrenal medulla	(47)	(50)	(50)	(49)
Hemorrhage	1 (2%)			
Hyperplasia	2 (4%)	3 (6%)	()	1 (2%)
Islets, pancreatic	(50)	(50)	(50)	(49)
Hyperplasia	4 (22.1)	2 (4%)	2 (4%)	
Infiltration cellular	1 (2%)	(0.4)	2 (4%)	(07)
Parathyroid gland	(26)	(24)	(34)	(27)
Infiltration cellular	(49)	1 (4%)	(47)	(40)
Pituitary gland	(48)	(49)	(47)	(49)
Pars distalis, angiectasis		1 (2%)	4 (9%)	2 (4%)
Pars distalis, cyst Pars distalis, hemorrhage	3 (6%)	1 (270)	1 (2%)	1 (2%)
Pars distalis, hyperplasia	10 (21%)	12 (24%)	23 (49%)	22 (45%)
Pars distalis, necrosis	10 (21/0)	12 (24/0)	£J (4J/0)	1 (2%)
Pars intermedia, hyperplasia	2 (4%)	1 (2%)		1 (2%)
Thyroid gland	(50)	(50)	(50)	(50)
Infiltration cellular	(00)	(00)	1 (2%)	(00)
Follicle, degeneration	1 (2%)		- (w/O)	
Follicular cell, hyperplasia	18 (36%)	23 (46%)	25 (50%)	35 (70%)

TABLE D5
Summary of the Incidence of Nonneoplastic Lesions in Female Mice in the 2-Year Inhalation Study of Ethylbenzene (continued)

	Chambe	r Control	75	ó ppm	250) ppm	750) ppm
General Body System								
Tissue NOS	(1)		(6)		(4)		(1)	
Fat, necrosis			4	(67%)	3	(75%)	1	(100%)
Genital System								
Clitoral gland	(41)		(47)		(48)		(48)	
Atrophy					1	(2%)		
Degeneration								(2%)
Ovary	(49)		(50)	4	(49)		(49)	
Angiectasis	2	(4%)		(2%)		4		
Atrophy		(4.007)		(2%)		(2%)	4.0	(000/)
Cyst		(16%)	8	(16%)		(20%)	10	(20%)
Hemorrhage	1	(2%)				(2%)		
Infiltration cellular Mineralization					2	(4%)	1	(90/)
Mineralization Uterus	(50)		(50)		(50)			(2%)
∪terus Angiectasis	(30)		(50)		(50)	(2%)	(50)	
Degeneration	9	(4%)	1	(2%)	1	(~ /O)	1	(2%)
Hemorrhage	۵	(470)	1	(270)	1	(2%)		(2%)
Infiltration cellular						(270)		(2%)
Inflammation	3	(6%)						(2%)
Thrombosis	_	(0.0)			1	(2%)	_	()
Endometrium, hyperplasia	44	(88%)	46	(92%)		(94%)	46	(92%)
Hematopoietic System Bone marrow Hematopoietic cell proliferation Infiltration cellular, histiocyte	(48)		(50) 1	(2%)	(50)		(50) 1	(2%)
Inflammation	1	(2%)						
Myelofibrosis								(4%)
Pigmentation, hemosiderin		(6%)	1	(2%)	1	(2%)		(2%)
Myeloid cell, hyperplasia		(2%)						(2%)
Lymph node	(3)		(7)	(4.40/)	(2)		(5)	
Iliac, hyperplasia			1	(14%)				(000/)
Inguinal, hyperplasia	4	(220/)					1	(20%)
Inguinal, pigmentation, hemosiderin Lumbar, hyperplasia		(33%) (33%)	0	(29%)				
Pancreatic, hyperplasia	1	(33/0)	۷	(20/0)			1	(20%)
Renal, hyperplasia	1	(33%)					1	(20/0)
Renal, necrosis	1	(00/0)	1	(14%)				
Lymph node, bronchial	(32)		(40)	(11/0)	(29)		(38)	
Hyperplasia	(52)			(5%)		(3%)		(5%)
Lymph node, mandibular	(47)		(48)		(47)	/	(44)	\$ 1 · · · · · · · · · · · · · · · · · ·
Hyperplasia		(2%)		(4%)	(' /			(5%)
Lymph node, mesenteric	(48)		(48)		(46)		(44)	
Hematopoietic cell proliferation		(2%)						
Hyperplasia		(8%)		(6%)	3	(7%)	2	(5%)
			2	(4%)				
Inflammation			~	()				
	1	(2%)	1	(2%) (2%)				

TABLE D5
Summary of the Incidence of Nonneoplastic Lesions in Female Mice in the 2-Year Inhalation Study of Ethylbenzene (continued)

	Chamber Control	75 ppm	250 ppm	750 ppm	
Hematopoietic System (continued)					
Lymph node, mediastinal	(34)	(42)	(41)	(31)	
Hyperplasia	3 (9%)	4 (10%)	` '	2 (6%)	
Hyperplasia, histiocytic	1 (3%)	(,		(3.7.3)	
Spleen	(50)	(50)	(50)	(49)	
Hematopoietic cell proliferation	4 (8%)	7 (14%)	2 (4%)	1 (2%)	
Hyperplasia	1 (2%)	, ,	, ,	, ,	
Necrosis		1 (2%)			
Pigmentation, hemosiderin	4 (8%)	1 (2%)	1 (2%)	4 (8%)	
Lymphoid follicle, hyperplasia	9 (18%)	5 (10%)	1 (2%)	3 (6%)	
Γhymus	(42)	(44)	(45)	(46)	
Åtrophy	5 (12%)	5 (11%)	5 (11%)	6 (13%)	
Thymocyte, hyperplasia	1 (2%)	2 (5%)	, ,		
Integumentary System					
Mammary gland	(49)	(50)	(48)	(49)	
Galactocele	(40)	1 (2%)	1 (2%)	(10)	
Hyperplasia		1 (2%)	1 (270)		
Skin	(50)	(50)	(49)	(50)	
Fibrosis	(30)	(30)	1 (2%)	(30)	
Inflammation	1 (90/)				
	1 (2%)		2 (4%)		
Necrosis	1 (2%)		9 (40/)		
Ulcer	1 (2%)		2 (4%)		
Musculoskeletal System					
Bone	(49)	(50)	(50)	(50)	
Arthrosis			1 (2%)		
Fracture		2 (4%)			
Periosteum, femur, inflammation	1 (2%)				
Nervous System					
Brain	(50)	(50)	(50)	(50)	
Hemorrhage	2 (4%)	(/	()	(/	
Mineralization	19 (38%)	17 (34%)	26 (52%)	25 (50%)	
Cerebellum, atrophy	1 (2%)	(/	,	()	
Cerebrum, atrophy	2 (4%)	1 (2%)	2 (4%)		
Cerebrum, gliosis	_ (2/0)	- (3/0)	- (2/0)	1 (2%)	
Cerebrum, hemorrhage				1 (2%)	
Medulla, atrophy	1 (2%)			- (~/~/	
Medulla, hemorrhage	1 (2%)				
Meninges, infiltration cellular	1 (270)	2 (4%)	1 (2%)		
Spinal cord	(1)	~ (1/0)	2 (2/0)		
Hemorrhage	1 (100%)				
	1 (100/0)				

TABLE D5
Summary of the Incidence of Nonneoplastic Lesions in Female Mice in the 2-Year Inhalation Study of Ethylbenzene (continued)

	Chamber Control	75 ppm	250 ppm	750 ppm
Respiratory System				
Larynx	(49)	(49)	(47)	(48)
Degeneration			1 (2%)	
Infiltration cellular	1 (2%)		1 (2%)	1 (2%)
Glands, degeneration		2 (4%)	2 (4%)	2 (4%)
Glands, inflammation			1 (2%)	
Lung	(50)	(50)	(49)	(50)
Hemorrhage	1 (2%)		1 (2%)	
Hyperplasia, lymphoid		1 (2%)		
Infiltration cellular, histiocyte	1 (2%)			
Alveolar epithelium, hyperplasia		1 (2%)	3 (6%)	1 (2%)
Alveolar epithelium, metaplasia				1 (2%)
Vein, thrombosis	1 (2%)			• •
Nose	(49)	(50)	(50)	(50)
Hemorrhage	1 (2%)			
Inflammation	3 (6%)	2 (4%)	3 (6%)	
Nasolacrimal duct, inflammation	1 (2%)			
Respiratory epithelium, metaplasia, squamo	us 2 (4%)		1 (2%)	
Pleura	(1)			
Гrachea	(50)	(50)	(50)	(50)
Special Senses System None				
U rinary System Kidney	(50)	(50)	(50)	(50)
Casts protein	1 (2%)	` '	• ,	1 (2%)
Infiltration cellular	` '	1 (2%)		` '
Mineralization		• •		1 (2%)
Nephropathy	13 (26%)	7 (14%)	9 (18%)	21 (42%)
Cortex, cyst			1 (2%)	1 (2%)
Cortex, metaplasia, osseous		1 (2%)		
Urinary bladder	(47)	(48)	(47)	(49)
Hemorrhage	1 (2%)			
Infiltration cellular	4 (9%)	4 (8%)	4 (9%)	5 (10%)
Illilia dioli celiulai				
Inflammation	1 (2%) 1 (2%)			

APPENDIX E GENETIC TOXICOLOGY

SALMONELLA	4 MUTAGENICITY TEST PROTOCOL	202
Mouse Lym	IPHOMA MUTAGENICITY TEST PROTOCOL	202
CHINESE HA	MISTER OVARY CELL CYTOGENETICS PROTOCOLS	203
Mouse Per	IPHERAL BLOOD MICRONUCLEUS TEST PROTOCOL	204
RESULTS		204
TABLE E1	Mutagenicity of Ethylbenzene in Salmonella typhimurium	205
TABLE E2	Induction of Trifluorothymidine Resistance in L5178Y Mouse Lymphoma Cells	
	by Ethylbenzene	206
TABLE E3	Induction of Sister Chromatid Exchanges in Chinese Hamster Ovary Cells	
	by Ethylbenzene	207
TABLE E4	Induction of Chromosomal Aberrations in Chinese Hamster Ovary Cells	
	by Ethylbenzene	208
TABLE E5	Frequency of Micronuclei in Peripheral Blood Erythrocytes of Mice	
	Following Treatment with Ethylbenzene by Inhalation for 13 Weeks	209

GENETIC TOXICOLOGY

SALMONELLA MUTAGENICITY TEST PROTOCOL

Testing was performed as reported by Zeiger *et al.* (1988). Ethylbenzene was sent to the laboratory as a coded aliquot from Radian Corporation (Austin, TX). It was incubated with the *Salmonella typhimurium* tester strains (TA97, TA98, TA100, and TA1535) either in buffer or S9 mix (metabolic activation enzymes and cofactors from Aroclor 1254-induced male Sprague-Dawley rat or Syrian hamster liver) for 20 minutes at 37° C. Top agar supplemented with L-histidine and d-biotin was added, and the contents of the tubes were mixed and poured onto the surfaces of minimal glucose agar plates. Histidine-independent mutant colonies arising on these plates were counted following incubation for 2 days at 37° C.

Each trial consisted of triplicate plates of concurrent positive and negative controls and five doses of ethylbenzene. The high dose was limited by toxicity. Trials performed in the absence of S9 were repeated. Trials initially performed with 10% S9 were repeated with 30% S9.

In this assay, a positive response is defined as a reproducible, dose-related increase in histidine-independent (revertant) colonies in any one strain/activation combination. An equivocal response is defined as an increase in revertants that is not dose related, not reproducible, or is of insufficient magnitude to support a determination of mutagenicity. A negative response is obtained when no increase in revertant colonies is observed following chemical treatment. There was no minimum percentage or fold increase required for a chemical to be judged positive or weakly positive.

MOUSE LYMPHOMA MUTAGENICITY TEST PROTOCOL

The experimental protocol is presented in detail by McGregor $\it et al.$ (1988). Ethylbenzene was supplied as a coded aliquot by Radian Corporation. The high dose of 160 µg/mL was determined by toxicity. L5178Y mouse lymphoma cells were maintained at 37° C as suspension cultures in supplemented Fischer's medium; normal cycling time was approximately 10 hours. To reduce the number of spontaneously occurring trifluorothymidine-resistant cells, subcultures were exposed to medium containing THMG (thymidine, hypoxanthine, methotrexate, and glycine) for 1 day, to medium containing THG (thymidine, hypoxanthine, and glycine) for 1 day, and to normal medium for 3 to 5 days. For cloning, the horse serum content was increased and Noble agar was added.

All treatment levels within an experiment, including concurrent positive and solvent controls, were replicated. Treated cultures contained 6×10^6 cells in 10 mL medium. This volume included the S9 fraction in those experiments performed with metabolic activation. Incubation with ethylbenzene continued for 4 hours, at which time the medium plus ethylbenzene was removed and the cells were resuspended in fresh medium and incubated for an additional 2 days to express the mutant phenotype. Cell density was monitored so that log phase growth was maintained. After the 48-hour expression period, cells were plated in medium and soft agar supplemented with trifluorothymidine (TFT) for selection of TFT-resistant (TK $^{-/-}$) cells, and cells were plated in nonselective medium and soft agar to determine cloning efficiency. Plates were incubated at 37 ° C in 5% CO $_2$ for 10 to 12 days. The test was initially performed without S9. Because a clearly positive response was obtained, the test was not performed with S9.

Minimum criteria for accepting an experiment as valid and a detailed description of the statistical analysis and data evaluation are presented in Caspary *et al.* (1988). All data were evaluated statistically for trend and peak responses. Both responses had to be significant ($P \le 0.05$) for ethylbenzene to be considered

positive, i.e., capable of inducing TFT resistance. A single significant response led to a "questionable" conclusion, and the absence of both a trend and peak response resulted in a "negative" call.

CHINESE HAMSTER OVARY CELL CYTOGENETICS PROTOCOLS

Testing was performed as reported by Galloway *et al.* (1987). Ethylbenzene was sent to the laboratory as a coded aliquot by Radian Corporation. It was tested in cultured Chinese hamster ovary (CHO) cells for induction of sister chromatid exchanges (SCEs) and chromosomal aberrations (Abs), both in the presence and absence of Aroclor 1254-induced male Sprague-Dawley rat liver S9 and cofactor mix. Cultures were handled under gold lights to prevent photolysis of bromodeoxyuridine-substituted DNA. Each test consisted of concurrent solvent and positive controls and of four doses of ethylbenzene; the high dose was limited by toxicity. A single flask per dose was used.

Sister Chromatid Exchange Test: In the SCE test without S9, CHO cells were incubated for 26 hours with ethylbenzene in supplemented McCoy's 5A medium. Bromodeoxyuridine (BrdU) was added 2 hours after culture initiation. After 26 hours, the medium containing ethylbenzene was removed and replaced with fresh medium plus BrdU and Colcemid, and incubation was continued for 1.5 hours. Cells were then harvested by mitotic shake-off, fixed, and stained with Hoechst 33258 and Giemsa. In the SCE test with S9, cells were incubated with ethylbenzene, serum-free medium, and S9 for 2 hours. The medium was then removed and replaced with medium containing serum and BrdU and no ethylbenzene, and incubation proceeded for an additional 25.8 hours, with Colcemid present for the final 2 hours. Harvesting and staining were the same as for cells treated without S9. All slides were scored blind and those from a single test were read by the same person. Fifty second-division metaphase cells were scored for frequency of SCEs/cell from each dose level.

Statistical analyses were conducted on the slopes of the dose-response curves and the individual dose points (Galloway et~al., 1987). An SCE frequency 20% above the concurrent solvent control value was chosen as a statistically conservative positive response. The probability of this level of difference occurring by chance at one dose point is less than 0.01; the probability for such a chance occurrence at two dose points is less than 0.001. An increase of 20% or greater at any single dose was considered weak evidence of activity; increases at two or more doses resulted in a determination that the trial was positive. A statistically significant trend (P< 0.005) in the absence of any responses reaching 20% above background led to a call of "equivocal."

Chromosomal Aberrations Test: In the Abs test without S9, cells were incubated in McCoy's 5A medium with ethylbenzene for 8.5 hours; Colcemid was added and incubation continued for 2 hours. The cells were then harvested by mitotic shake-off, fixed, and stained with Giemsa. For the Abs test with S9, cells were treated with ethylbenzene and S9 for 2 hours, after which the treatment medium was removed and the cells were incubated for 8.5 hours in fresh medium, with Colcemid present for the final 2 hours. Cells were harvested in the same manner as for the treatment without S9.

Cells were selected for scoring on the basis of good morphology and completeness of karyotype $(21 \pm 2 \text{ chromosomes})$. All slides were scored blind and those from a single test were read by the same person. One hundred first-division metaphase cells were scored at each dose level. Classes of aberrations included simple (breaks and terminal deletions), complex (rearrangements and translocations), and other (pulverized cells, despiralized chromosomes, and cells containing 10 or more aberrations).

Chromosomal aberration data are presented as percentage of cells with aberrations. To arrive at a statistical call for a trial, analyses were conducted on both the dose response curve and individual dose points. For a

single trial, a statistically significant ($P \le 0.05$) difference for one dose point and a significant trend ($P \le 0.015$) were considered weak evidence for a positive response; significant differences for two or more doses indicated the trial was positive. A positive trend test in the absence of a statistically significant increase at any one dose resulted in an equivocal call (Galloway *et al.*, 1987). Ultimately, the trial calls were based on a consideration of the statistical analyses as well as the biological information available to the reviewers.

MOUSE PERIPHERAL BLOOD MICRONUCLEUS TEST PROTOCOL

A detailed discussion of this assay can be found in MacGregor et~al.~(1990). Peripheral blood samples were obtained from male and female B6C3F1 mice at the end of a 13-week toxicity study (NTP, 1992). Smears were immediately prepared and fixed in absolute methanol, stained with a chromatin-specific fluorescent dye mixture of Hoechst 33258/pyronin Y (MacGregor et~al.~,1983), and coded. Slides were scanned at $630\times$ or $1,000\times$ to determine the frequency of micronuclei in 2,000 polychromatic erythrocytes (PCEs) and 10,000 normochromatic erythrocytes (NCEs) in each animal of each dose group. The criteria of Schmid (1976) were used to define micronuclei, with the additional requirement that the micronuclei exhibit the characteristic fluorescent emissions of DNA (blue with 360 nm and orange with 510 nm ultraviolet illumination); the minimum size limit was approximately one-twentieth the diameter of the NCE cell. In addition, the percentage of PCEs among the total erythrocyte population was determined.

Log transformation of the NCE data, testing for normality by the Shapiro-Wilk test, and testing for heterogeneity of variance by Cochran's test were performed before statistical analyses. The frequency of micronucleated cells among NCEs was analyzed by analysis of variance using the SAS GLM procedure. The NCE data for each dose group were compared with the concurrent solvent control using Student's *t*-test. The frequency of micronucleated cells among PCEs was analyzed by the Cochran-Armitage trend test, and individual dose groups were compared to the concurrent solvent control by Kastenbaum-Bowman's binomial test. The percentage of PCEs among total erythrocytes was analyzed by an analysis of variance on ranks (classed by sex), and individual dose groups were compared with the concurrent solvent control using a *t*-test on ranks.

RESULTS

Ethylbenzene was not mutagenic in *S. typhimurium* strain TA97, TA98, TA100, or TA1535 with or without Aroclor-induced rat or hamster liver S9 (Table E1; Zeiger *et al.*, 1988). A positive response was observed with ethylbenzene in the L5178Y mouse lymphoma cell assay in the absence of S9 at the highest nonlethal dose tested (80 μ g/mL); the assay was not performed with S9 (Table E2; McGregor *et al.*, 1988). A significant amount of cytotoxicity was noted at this dose level (relative total growth was reduced to 34% and 13% of the control level in each of two trials). No increases in SCEs (Table E3) or Abs (Table E4) were induced by ethylbenzene in cultured CHO cells, with or without S9. *In vivo*, no increases in frequencies of micronucleated erythrocytes were observed in peripheral blood samples from male and female mice treated for 13 weeks with ethylbenzene (Table E5).

TABLE E1 Mutagenicity of Ethylbenzene in Salmonella typhimurium^a

	Revertants/plate ^b								
Strain Dose	-S!)	+ hams	ster S9	+ rat S9				
(μg/plate)	Trial 1	Trial 2	10%	30%	10%	30%			
TA100									
0	112 ± 9.3	147 ± 4.0	114 ± 8.2	136 ± 3.3	111 ± 2.1	154 ± 7.8			
10	104 ± 0.9	161 ± 5.8	120 ± 11.5	138 ± 9.5	100 ± 5.0	155 ± 9.0			
33	100 ± 4.4	147 ± 4.1	137 ± 22.7	140 ± 138	110 ± 8.1	155 ± 9.3			
100	97 ± 4.8	157 ± 3.2	109 ± 7.1	138 ± 12.2	105 ± 2.3	161 ± 14.5			
333	97 ± 6.9	118 ± 11.5	97 ± 7.1	137 ± 1.2	111 ± 4.7	127 ± 13.2			
666	76 ± 6.2	$74 \pm 4.0^{\circ}$	0. =	101 = 112		12. = 10.2			
1,000	.0 = 0.2		$98~\pm~1.7$	112 ± 6.1	$77~\pm~8.2$	$109~\pm~8.8$			
Trial summary	Negative	Negative	Negative	Negative	Negative	Negative			
Positive control ^d	375 ± 12.3	394 ± 32.5	873 ± 46.0	740 ± 18.0	$1,304 \pm 306.0$	352 ± 19.8			
ТА1535									
0	14 ± 3.7	29 ± 3.8	7 ± 1.5	11 ± 2.3	9 ± 2.0	12 ± 1.2			
10	19 ± 1.3	26 ± 3.2	9 ± 1.3	14 ± 1.5	8 ± 0.7	13 ± 2.5			
33	21 ± 4.6	19 ± 2.5	6 ± 0.7	11 ± 1.5	9 ± 3.0	8 ± 0.6			
100	16 ± 1.5	25 ± 2.5	8 ± 1.5	10 ± 2.4	5 ± 0.6	10 ± 1.5			
333	16 ± 2.1	14 ± 0.3	9 ± 1.2	9 ± 2.7	8 ± 2.4	6 ± 0.9			
666	0 ± 0.0^{e}	0 ± 0.0							
1,000			5 ± 1.8	11 ± 1.9	5 ± 1.5	9 ± 1.5			
Trial summary	Negative	Negative	Negative	Negative	Negative	Negative			
Positive control	418 ± 23.1	$520~\pm~20.0$	703 ± 16.5	431 ± 36.9	$393~\pm~72.0$	101 ± 11.4			
TA97									
0	182 ± 1.5	111 ± 9.5	195 ± 12.3	184 ± 18.2	200 ± 10.0	218 ± 6.5			
10	203 ± 1.8	$120~\pm~16.3$	194 ± 10.3	210 ± 22.5	190 ± 15.1	$249~\pm~20.2$			
33	198 ± 6.9	$144~\pm~2.4$	195 ± 3.5	$186~\pm~22.4$	193 ± 5.3	227 ± 16.5			
100	195 ± 9.9	$124~\pm~5.2$	191 ± 7.1	227 ± 1.8	179 ± 7.8	12 ± 13.0			
333	188 ± 5.7	108 ± 9.1	173 ± 3.5	$202~\pm~8.3$	211 ± 3.3	211 ± 6.4			
666	103 ± 1.5	$6 \pm 5.7^{\circ}$							
1,000			$124~\pm~9.6$	$180~\pm~15.9$	189 ± 23.4	$195~\pm~15.3$			
Trial summary	Negative	Negative	Negative	Negative	Negative	Negative			
Positive control	856 ± 20.8	954 ± 47.1	$1,587 \pm 146.1$		647 ± 154.3	540 ± 12.7			
ТА98									
0	26 ± 1.8	29 ± 5.5	24 ± 3.2	35 ± 3.8	34 ± 3.3	34 ± 7.2			
10	16 ± 2.3	27 ± 4.4	29 ± 1.8	34 ± 4.7	26 ± 1.8	32 ± 4.1			
33	22 ± 4.8	35 ± 7.8	26 ± 0.6	34 ± 4.5	34 ± 3.5	32 ± 2.3			
100	21 ± 2.4	16 ± 2.1	28 ± 4.7	26 ± 1.2	32 ± 2.3	30 ± 4.2			
333	18 ± 1.5	20 ± 8.4	23 ± 3.0	30 ± 0.7	30 ± 2.3	28 ± 5.6			
666	13 ± 1.3 13 ± 1.2	$27 \pm 14.5^{\circ}$	20 ± 0.0	00 = 0.7	υυ ± ω.υ	20 - 0.0			
1,000	10 - 1.2	ωι ± 11.0	$21~\pm~2.3$	$30~\pm~0.9$	$26~\pm~1.5$	$30~\pm~3.5$			
Trial summary	Negative	Negative	Negative	Negative	Negative	Negative			
Positive control	845 ± 69.2	566 ± 45.0	$1,082 \pm 174.8$	285 ± 32.9	784 ± 214.8	149 ± 10.7			

Study was performed at SRI International. The detailed protocol and these data are presented in Zeiger et al. (1988). 0 µg/plate was the solvent control.

Revertants are presented as mean ± standard error from three plates.

Slight toxicity

The positive controls in the absence of metabolic activation were sodium azide (TA100 and TA1535), 9-aminoacridine (TA97), and 4-nitro-o-phenylenediamine (TA98). The positive control for metabolic activation with all strains was 2-aminoanthracene.

Precipitate on plate, toxic

TABLE E2 Induction of Trifluorothymidine Resistance in L5178Y Mouse Lymphoma Cells by Ethylbenzene^a

	centration μg/mL)	Cloning Efficiency (%)	Relative Total Growth (%)	Mutant Count	Mutant Fraction ^b	Average Mutant Fraction	
-S9							
Trial 1							
$Dimethyl sulfoxide^{\tt C}$		84	94	159	63		
		89	106	150	56		
		78 87	108 92	155 138	66 53	60	
		01	92	136	33	00	
Ethylmethane sulfonate ^d	250	81	85	357	147		
Eury iniculaire surronate	200	83	95	374	150	149*	
,							
Methylmethane sulfonated	¹ 15	61	40	251	138		
		52	39	238	152	145*	
ral II	10	0.1	100	100	F-1		
Ethylbenzene	10	81	103	123	51	50	
	20	86	106	157	61	56	
	20	81 81	90 93	130 127	54 52	53	
	40	87	82	175	67	JJ	
	40	73	72	144	66	67	
	80	74	36	1,235	559	07	
	00	71	32	1,335	619	589*	
	160	Lethal					
T.:-1 0							
Trial 2 Dimethylsulfoxide		85	106	87	34		
Difficulty is unoxide		69	95	63	30		
		82	98	75	30		
		100	101	91	30	31	
Ethylmethane sulfonate	250	50	67	302	201		
		51	67	381	250	225*	
36 1 1 1 10 .	4.5	40	2.4	100	0.4		
Methylmethane sulfonate	15	43	34	122	94	107*	
		42	32	152	120	107*	
Ethylbenzene	20	83	83	109	44		
Larymenzene	20	82	83	103	41	42	
	40	78	61	75	32	- M	
	-	73	54	58	27	29	
	60	64	37	91	48		
		68	60	79	39	43	
	80	48	10	228	159		
	100	55	15	233	142	150*	
	100	Lethal					

Significant positive response ($P \le 0.05$) versus the solvent control Study was performed at Inveresk Research International. The detailed protocol and these data are presented in McGregor *et al.* (1988). Mutant fraction (MF) (frequency) is a ratio of the mutant count to the cloning efficiency, divided by 3 [to arrive at MF/10⁶ cells treated]).
Solvent control
Positive control

TABLE E3 Induction of Sister Chromatid Exchanges in Chinese Hamster Ovary Cells by Ethylbenzene^a

Compound	Dose (µg/mL)	Total Cells Scored	No. of Chromo- somes	No. of SCEs	SCEs/ Chromo- some	SCEs/ Cell	Hrs in BrdU	Relative Change of SCEs/ Chromosome ^b (%)
- S9 Summary: Negative								
Dimethylsulfoxide ^c		50	1,045	555	0.53	11.1	25.5	
Mitomycin-C ^d	0.001 0.010	50 5	1,041 103	773 220	0.74 2.13	15.5 44.0	25.5 25.5	39.81 302.17
Ethylbenzene	75.5 99.5 125 ^e 151 ^e	50 50 50 0	1,046 1,049 1,033	551 522 590	0.52 0.49 0.57	11.0 10.4 11.8	25.5 25.5 25.5 25.5	-0.82 -6.31 7.54
					$P = 0.207^{f}$			
+ S9 Summary: Negative								
Dimethylsulfoxide		50	1,047	531	0.50	10.6	25.8	
Cyclophosphamide ^d	$0.35 \\ 2$	50 5	1,048 108	723 159	0.68 1.47	14.5 31.8	25.8 25.8	36.03 190.29
Ethylbenzene	125 137.5 150 ^e 175 ^e	50 50 50 50	1,044 1,041 1,037	561 531 516	0.53 0.51 0.49	11.2 10.6 10.3	25.8 25.8 25.8 25.8	5.95 0.58 -1.89
					P= 0.713			

Study was performed at Litton Bionetics, Inc. A detailed description of the protocol is presented in Galloway et al. (1987). SCE= sister chromatid exchange; BrdU= bromodeoxyuridine SCEs/chromosome in treated cells versus SCEs/chromosome in solvent control cells

Solvent control

Positive control

Precipitate on plate
Significance of SCEs/chromosome tested by the linear regression trend test versus log of the dose

TABLE E4 Induction of Chromosomal Aberrations in Chinese Hamster Ovary Cells by Ethylbenzene^a

		- S9			+ S9				
Dose	Dose Total Cells No. of Abs/ Cells with				Dose Total Cells No. of Abs/ Cells wi				
(μg/mL) Scored	Abs	Cell	Abs (%)	(μg/mL) Scored Abs Cell Abs (%				
Harvest time: 10.5 Summary: Negati					Harvest time: 10.5 hours Summary: Negative				
Dimethylsulfoxide ^l	100	3	0.03	3	Dimethylsulfoxide 100 3 0.03 3				
Mitomycin-C ^c					Cyclophosphamide ^C				
1	50	16	0.32	22	50 50 23 0.46 36				
Ethylbenzene					Ethylbenzene				
75	100	1	0.01	1	75 100 4 0.04 4				
100	100	3	0.03	3	100 100 1 0.01 1				
125	100	5	0.05	5	125 100 1 0.01 1				
150	0				150 0				
				$P = 0.150^{d}$	P= 0.917				

Study was performed at Litton Bionetics, Inc. The detailed protocol is presented in Galloway et al. (1987). Abs= aberrations

Solvent control Positive control

Significance of percent cells with aberrations tested by the linear regression trend test versus log of the dose

TABLE E5 Frequency of Micronuclei in Peripheral Blood Erythrocytes of Mice Following Treatment with Ethylbenzene by Inhalation for 13 Weeks^a

Compound	Dose	Number of Mice with Erythrocytes	Micronucleated	PCEs ^b	
o o mpo u mu	(ppm)	Scored	PCEs	NCEs	(%)
Male					
	0	8	2.18 ± 0.56	1.54 ± 0.16	$2.22~\pm~0.10$
	500	10	2.04 ± 0.31	1.68 ± 0.13	3.13 ± 0.94
	750	9	1.90 ± 0.53	1.90 ± 0.13	1.97 ± 0.09
	1,000	10	1.21 ± 0.20	1.59 ± 0.16	2.02 ± 0.14
rend test ^c			P = 0.928	P= 0.816	
NOVA ^d					P = 0.278
emale					
	0	10	1.54 ± 0.56	0.92 ± 0.11	1.74 ± 0.14
	500	10	2.64 ± 0.53	1.01 ± 0.12	1.83 ± 0.18
	750	10	1.87 ± 0.38	1.32 ± 0.22	1.85 ± 0.15
	1,000	10	1.01 ± 0.26	1.12 ± 0.12	1.80 ± 0.15
rend test			P = 0.817	P = 0.077	
NOVA					P = 0.886
verall trend			P = 0.951	P = 0.149	
verall ANOVA					P = 0.684

Study was performed at the USDA Western Regional Center. The protocol is presented in MacGregor *et al.* (1990). PCE= polychromatic erythrocyte; NCE= normochromatic erythrocyte. At least 2,000 PCEs and 10,000 NCEs were scored from each animal.

Mean ± standard error

Cochran-Armitage linear regression of proportions for PCEs or linear contrasts from analysis of variance for NCEs Analysis of variance on ranks

APPENDIX F CHEMICAL CHARACTERIZATION AND GENERATION OF CHAMBER CONCENTRATIONS

PROCUREME	NT AND CHARACTERIZATION OF ETHYLBENZENE	212
VAPOR GENE	ERATION AND EXPOSURE SYSTEM	213
VAPOR CONC	CENTRATION MONITORING	214
CHAMBER AT	TMOSPHERE CHARACTERIZATION	214
FIGURE F1	Infrared Absorption Spectrum of Ethylbenzene	216
FIGURE F2	Nuclear Magnetic Resonance Spectrum of Ethylbenzene	217
FIGURE F3	Schematic of Generation and Delivery System	218
FIGURE F4	Inhalation Suite	219
TABLE F1	Summary of Chamber Concentrations in the 2-Year Inhalation Studies	
	of Ethylbenzene	220

CHEMICAL CHARACTERIZATION AND GENERATION OF CHAMBER CONCENTRATIONS

PROCUREMENT AND CHARACTERIZATION OF ETHYLBENZENE

Ethylbenzene was obtained from ARCO Chemical Company (Newtown Square, PA) in two lots (A060989 and A051890) that were used during the 2-year studies. Identity, purity, and stability analyses were conducted by the analytical chemistry laboratory, Midwest Research Institute (Kansas City, MO). Reports on analyses performed in support of the ethylbenzene studies are on file at the National Institute of Environmental Health Sciences.

The chemical, a clear, colorless, pungent smelling, volatile liquid, was identified as ethylbenzene by infrared, ultraviolet/visible (lot A060989 only), and nuclear magnetic resonance spectroscopy. All spectra were consistent with the literature spectra (*Sadtler Standard Spectra*) of ethylbenzene. The infrared and nuclear magnetic resonance spectra are presented in Figures F1 and F2. The boiling point and density of the chemical were also consistent with literature references (*Merck Index*, 1983).

The purity of lot A060989 was determined by elemental analyses, Karl Fischer water analysis, peroxide determination, and gas chromatography. To determine peroxide concentrations, a sample of ethylbenzene was refluxed with isopropyl alcohol, glacial acetic acid, and sodium iodide; liberated iodine was titrated with 0.1 N sodium thiosulfate to the starch endpoint. Gas chromatography was performed using a flame ionization detector. Two systems were used:

- A) 1% SP-1000 on 80/100 Supelcoport glass column, with a nitrogen carrier gas at a flow rate of 70 mL/minute, and an oven temperature program of 50° C for 5 minutes, then 50° to 250° C at 10° C per minute.
- B) DB-5 Megabore capillary fused-silica column with a helium carrier gas at a flow rate of 10 mL/minute, a makeup gas of nitrogen at a flow rate of 20 mL/minute, and an oven temperature program of 50° C for 5 minutes, then 50° to 250° C at 10° C per minute.

Elemental analyses for carbon and hydrogen were in agreement with the theoretical values for ethylbenzene. Karl Fischer water analysis indicated less than 0.05% water. Iodometric titration revealed no peroxide. Gas chromatography by each system revealed a major peak and no impurities with areas greater than 0.1% relative to the major peak. Major peak comparisons of lot A060989 with a previously analyzed lot of ethylbenzene (lot K061786) not used in the current studies indicated a purity of $101.0\% \pm 0.5\%$ for lot A060989 relative to lot K061786. The overall purity of lot A060989 was determined to be greater than 99%.

Additional analyses of lot A060989 were performed with gas chromatography/mass spectrometry to identify and quantify cumene in the bulk ethylbenzene. The gas chromatograph system included a DB-5 fused-silica capillary column with a helium carrier gas at a linear flow rate of $30~\rm cm^3/second$ and an oven temperature program of 60° C for 5 minutes, then 60° to 200° C at 10° C per minute; injection was performed with a 30-second splitless delay. Tridecane was added as an internal standard to the cumene standard solution. Cumene was identified by comparison of retention times and specific ion ratios to the cumene standard. Cumene in lot A060989 had a retention time of 5.8 minutes and a specific ion ratio of 19:100:23, compared to a retention time of 5.6 minutes and an ion ratio of 20:100:24 for the standard. System B described for purity analyses, with cumene added as a standard, was used to quantify cumene; 62 ± 3.1 ppm was detected.

The purity of lot A051890 was determined by iodometric titration for peroxide and by gas chromatography with system A, but with a 10% SP-1000 on 80/100 Supelcoport glass column. Less than 2 ppm peroxide was detected. Gas chromatography indicated one impurity with an area of 0.1% relative to the major peak. The overall purity of lot A051890 was determined to be greater than 99%.

Accelerated stability studies of lot K061786 were performed by the analytical chemistry laboratory. Gas chromatography was performed with system A but with a 10% SP-1000 column and 87° C isothermal temperature. These studies indicated that ethylbenzene is stable as a bulk chemical for at least 2 weeks when stored protected from light at temperatures up to 60° C. To ensure stability, the bulk chemical was stored at room temperature in the original steel containers until just prior to use, when it was transferred to amber glass bottles with Teflon®-lined caps and a nitrogen headspace. The rapid use and small shipment sizes of ethylbenzene made stability monitoring unnecessary during the studies; however, the peroxide content of the bulk chemical was tested monthly with iodometric titration. The concentration of peroxide ranged from 1.12 to 10.7 ppm.

VAPOR GENERATION AND EXPOSURE SYSTEM

A diagram of the ethylbenzene generation and delivery system is shown in Figure F3. Ethylbenzene vapor was produced by flash evaporator units. Liquid ethylbenzene was pumped by fine metering pumps from a reservoir into the top of a 30-cm-long, 20-mm internal diameter, Hempel distillation column packed with 3-mm diameter glass beads. At its lower end, the column was fitted into a two-armed 500-mL glass flask. One arm of the flask allowed access by a thermocouple that, in conjunction with a thermostated heating tape, maintained the column temperature at $150^{\circ} \pm 15^{\circ}$ C. Nitrogen carrier gas at 95 psi was bled from a high-pressure liquid nitrogen tank monitored by a weight scale, was passed through a manifold, and entered the flask through the second arm; it was heated to $200^{\circ} \pm 50^{\circ}$ C by a mantle surrounding the flask. The nitrogen gas carried ethylbenzene vapor into stainless steel transfer lines heated to 75° C by a heating tape. Magnehelic gauges were installed in the carrier gas lines immediately before the flash evaporators to monitor for blockages; pressure alarms ensured that the nitrogen gas pressure remained within the appropriate range. Transfer lines led to exposure chambers. Each exposure chamber was supplied by a separate flash evaporator unit.

Exposure concentrations for individual exposure chambers were created by varying the ethylbenzene flow rate to the individual flash evaporation units. Ethylbenzene vapor concentrations of 75, 250, and 750 ppm were created by ethylbenzene flow rates of 0.19, 0.63, and 1.9 mL/minute. To prevent saturation of the vapor streams, nitrogen flow rates were maintained at 5 L/minute for 75 and 250 ppm chambers and 10 L/minute for 750 ppm chambers. Each carrier gas line was fitted with a pressure release valve to shield the glass flash evaporator from pressure buildup due to blockage. At the chamber inlets, the ethylbenzene vapor passed through venturi-type plenums to enhance complete mixing with HEPA- and charcoal-filtered air

Stainless-steel chambers (Hazleton $H-2000^{\circ}$) manufactured by Lab Products, Inc. (Maywood, NJ) were used throughout the studies. A diagram of the inhalation suite is shown in Figure F4. The total volume of each chamber was 2.3 m^3 ; the active mixing volume of each chamber was 1.7 m^3 . The chamber was designed so that uniform vapor concentrations could be maintained throughout the chamber when catch pans were in place.

The 750 ppm chambers were sampled once during the first full week of exposure for the presence of aerosol by a Quartz Crystal Microbalance Cascade Impactor (California Measurements, Sierra Madre, CA). Aerosol concentrations prior to and during exposure were $0.1492 \pm 0.0121 \text{ mg/m}^3$ and $0.0904 \pm 0.0270 \text{ mg/m}^3$, respectively, for rats and $0.1772 \pm 0.0633 \text{ mg/m}^3$ and $0.2522 \pm 0.0605 \text{ mg/m}^3$,

respectively, for mice. These results indicate that aerosol formation due to test atmosphere generation was not significant.

VAPOR CONCENTRATION MONITORING

The chamber concentrations of ethylbenzene were monitored automatically by an on-line gas chromatograph (Hewlett Packard Model 5880A; Hewlett Packard, Palo Alto, CA) with a flame ionization detector and a 10% SP-1000 on 80/100 Supelcoport glass column. Samples were drawn from supply lines leading to exposure chambers and the control chamber at least once every hour by a six-port gas sample valve in conjunction with a 10-port stream selector valve. A similarly equipped gas chromatograph was used as a backup and for the analysis of grab samples.

The on-line monitoring system was calibrated using certified gas standards prepared by Scott Specialty Gases (Troy, MI) and Air Products Specialty Gases (Chicago, IL). Calibration was then verified by analyzing liquid standards prepared gravimetrically with bulk ethylbenzene. Calibrations were performed prior to the beginning of the studies, weekly for the first 2 weeks of the studies, and monthly thereafter using the certified gas standards. Daily calibration checks were performed by analyzing a randomly selected standard gas sample; if the concentration deviated by more than 10% from the current calibration curve, a full-range recalibration was performed at the earliest convenient time.

Monthly calibrations of the backup gas chromatograph used in these studies were performed by collecting samples of the gas standards in gas-tight syringes and injecting them into the gas chromatograph. Daily calibration checks were performed by analyzing randomly selected standard gas samples; if the concentration deviated by more than 10% from the current calibration curve, a full range recalibration was performed. Summaries of the chamber concentrations are presented in Table F1.

CHAMBER ATMOSPHERE CHARACTERIZATION

The times for the exposure concentration to build up to 90% of the final exposure concentration (T_{90}) and to decay to 10% of the exposure concentration (T_{10}) were measured in the 750 ppm exposure chambers with animals present during the first 2 weeks of the studies. At a chamber airflow rate of 15 air changes per hour, the theoretical value for both T_{90} and T_{10} is 10 minutes. Plots of time-concentration histories during the first 2 weeks of the studies indicated T_{90} and T_{10} values of 15 minutes; therefore, 15 minutes was used for the T_{90} value throughout the studies. Actual T_{90} values ranged from 11.4 to 15 minutes for rats and from 8.9 to 12.3 minutes for mice. Actual T_{10} values ranged from 10.5 to 11.7 minutes for rats and from 9.9 to 10.7 minutes for mice.

Inhalation chambers were sampled to determine the uniformity of ethylbenzene concentrations; grab samples from 12 shelf positions within the exposure chamber were analyzed by an off-line gas chromatograph. Grab samples were collected in gas-tight syringes from sampling ports in the exposure chambers without animals present before exposures began and with animals present approximately every 90 days during the studies. Chamber concentration uniformity was maintained throughout the studies.

The persistence of ethylbenzene following exposure was monitored by gas chromatography in the 750 ppm chambers without animals present, at 4-minute intervals for at least 2 hours, and with animals present once per hour for at least 2 hours during the first week of the studies and at 90-day intervals afterward. No ethylbenzene was detectable after 2 hours (detection limit 0.44 ppm).

The stability of ethylbenzene was monitored in the generator reservoirs of the 75 and 750 ppm chambers. Samples were collected without animals in the chambers, before the studies began, over a 3-day simulated

exposure period; samples were collected at the beginning of the first day and after 6 hours of ethylbenzene generation on the third day. Samples were also collected on day 1 of the studies, during the first hour of exposure, and on day 5, during the sixth hour of exposure. Grab samples (1 mL) were diluted to 100 mL with methylene chloride and analyzed by gas chromatography. No contaminants or degradation products with peak areas of 0.1% or greater relative to the major peak were found in any of the generator reservoir samples.

Grab samples from occupied and unoccupied 75 and 750 ppm chambers were analyzed for degradation products. Grab samples (10 mL) of chamber atmospheres were collected in gas-tight syringes and analyzed by gas chromatography. Samples were collected without animals in the chambers, before the studies began, over a 3-day simulated exposure period; samples were collected at the beginning of the first day and after 6 hours of ethylbenzene generation on the third day. Sampling was also performed every 90 days throughout the study; samples were collected during the first hour of the first day of the exposure week and during the sixth hour of day 5 of the exposure week. One small impurity with an area less than 0.04% of the ethylbenzene peak area was detected in samples taken from the 750 ppm chambers.

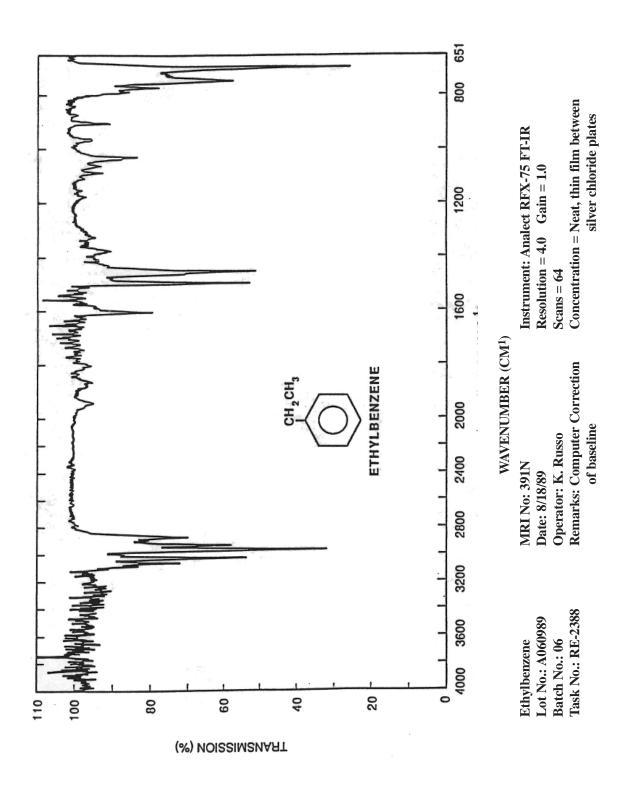


FIGURE F1
Infrared Absorption Spectrum of Ethylbenzene

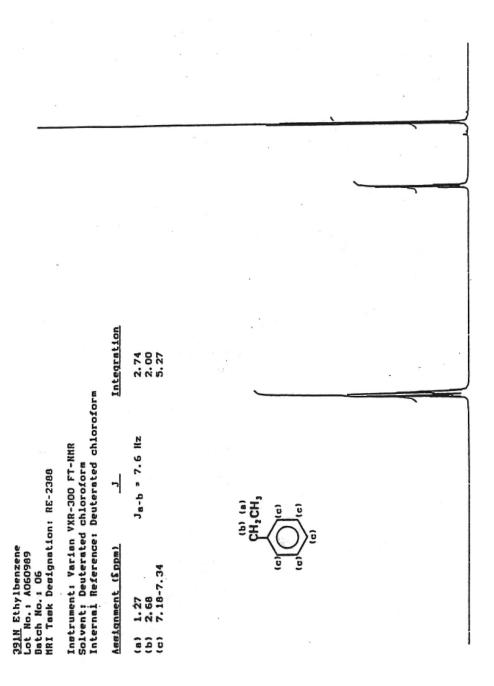


FIGURE F2 Nuclear Magnetic Resonance Spectrum of Ethylbenzene

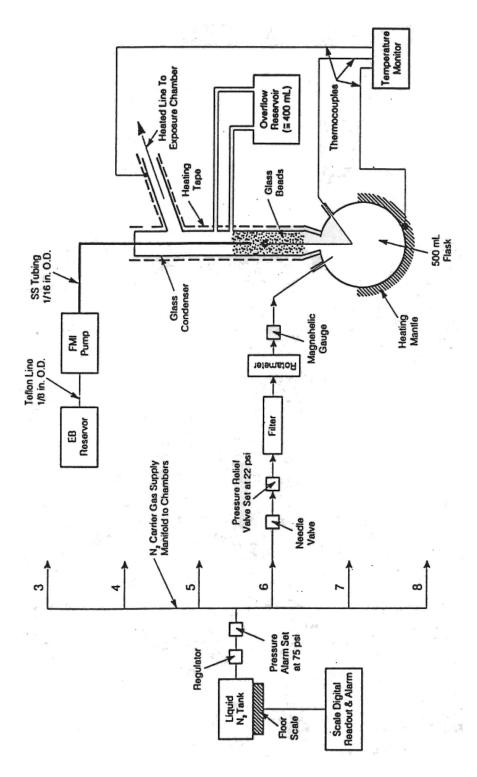


FIGURE F3
Schematic of Generation and Delivery System

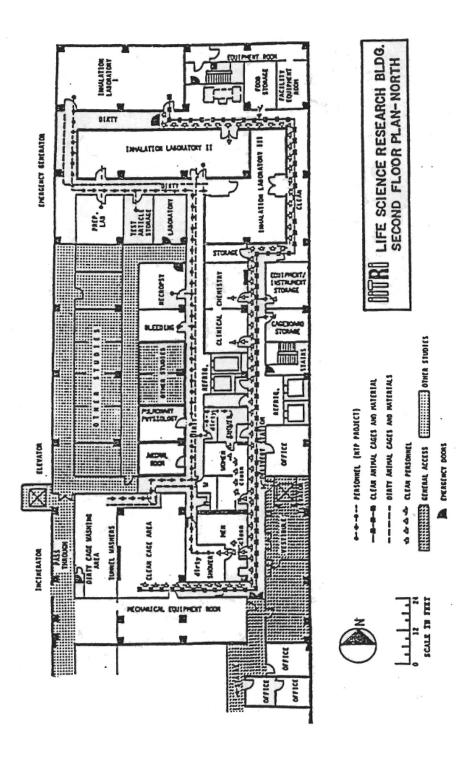


FIGURE F4
Inhalation Suite

TABLE F1 Summary of Chamber Concentrations in the 2-Year Inhalation Studies of Ethylbenzene

Target Concentration (ppm)	Total Number of Readings ^a	Average Concentration ^b (ppm)
Rat Chambers		
75	104	74.8 ± 1.7
250	104	250 ± 4
750	104	749 ± 7
Mouse Chambers		
75	103	75.2 ± 1.5
250	103	248 ± 5
750	103	748 ± 9

 $[\]begin{array}{ll} a & \text{Number of weekly means} \\ b & \text{Mean} \pm \text{ standard deviation; average of weekly means} \end{array}$

APPENDIX G INGREDIENTS, NUTRIENT COMPOSITION, AND CONTAMINANT LEVELS IN NIH-07 RAT AND MOUSE RATION

TABLE G1	Ingredients of NIH-07 Rat and Mouse Ration	222
TABLE G2	Vitamins and Minerals in NIH-07 Rat and Mouse Ration	222
TABLE G3	Nutrient Composition of NIH-07 Rat and Mouse Ration	223
TARLE G4	Contaminant Levels in NIH-07 Rat and Mouse Ration	224

TABLE G1 Ingredients of NIH-07 Rat and Mouse Ration^a

Ingredients ^b	Percent by Weight	
Ground #2 yellow shelled corn	24.50	
Ground hard winter wheat	23.00	
Soybean meal (49% protein)	12.00	
Fish meal (60% protein)	10.00	
Wheat middlings	10.00	
Dried skim milk	5.00	
Alfalfa meal (dehydrated, 17% protein)	4.00	
Corn gluten meal (60% protein)	3.00	
Soy oil	2.50	
Dried brewer's yeast	2.00	
Dry molasses	1.50	
Dicalcium phosphate	1.25	
Ground limestone	0.50	
Salt	0.50	
Premixes (vitamin and mineral)	0.25	
,		

TABLE G2 Vitamins and Minerals in NIH-07 Rat and Mouse Ration^a

	Amount	Source	
Vitamins			
A	5,500,000 IU	Stabilized vitamin A palmitate or acetate	
D_2	4,600,000 IU	D-activated animal sterol	
D ₃ K ₃	2.8 g	Menadione	
d - α -Tocopheryl acetate	20,000 IŬ		
Choline	560.0 g	Choline chloride	
Folic acid	2.2 g		
Niacin	30.0 g		
d-Pantothenic acid	18.0 g	d-Calcium pantothenate	
Riboflavin	3.4 g		
Thiamine	10.0 g	Thiamine mononitrate	
B ₁₂	4,000 µg		
Pyridoxine	1.7 g	Pyridoxine hydrochloride	
Biotin	140.0 mg	d-Biotin	
Minerals			
Iron	120.0 g	Iron sulfate	
Manganese	60.0 g	Manganous oxide	
Zinc	16.0 g	Zinc oxide	
Copper	4.0 g	Copper sulfate	
Iodine	1.4 g	Calcium iodate	
Cobalt	0.4 g	Cobalt carbonate	

 $^{^{\}rm a}$ Per ton (2,000 lb) of finished product

a NCI, 1976; NIH, 1978
 b Ingredients were ground to pass through a U.S. Standard Screen No. 16 before being mixed.

TABLE G3 Nutrient Composition of NIH-07 Rat and Mouse Ration

Mean ± Standard			
Nutrient	Deviation	Range	Number of Samples
Protein (% by weight)	23.42 ± 0.56	22.2 - 24.3	25
Crude fat (% by weight)	5.30 ± 0.16	5.00 - 5.60	25
Crude fiber (% by weight)	3.49 ± 0.41	2.60 - 4.30	25
Ash (% by weight)	6.37 ± 0.18	6.11 - 6.81	25
Amino Acids (% total diet)			
Arginine	1.280 ± 0.083	1.110 - 1.390	11
Cystine	0.308 ± 0.071	0.181 - 0.400	11
Glycine	1.158 ± 0.048	1.060 - 1.220	11
Histidine	0.584 ± 0.027	0.531 - 0.630	11
Isoleucine	0.917 ± 0.033	0.867 - 0.965	11
Leucine	1.975 ± 0.051	1.850 - 2.040	11
Lysine	1.274 ± 0.049	1.200 - 1.370	11
Methionine	0.437 ± 0.109	0.306 - 0.699	11
Phenylalanine	0.999 ± 0.120	0.665 - 1.110	11
Threonine	0.904 ± 0.058	0.824 - 0.985	11
Tryptophan	0.218 ± 0.153	0.107 - 0.671	11
Tyrosine	0.685 ± 0.094	0.564 - 0.794	11
Valine	1.086 ± 0.055	0.962 - 1.170	11
Essential Fatty Acids			
Linoleic	2.407 ± 0.227	1.830 - 2.570	10
Linolenic	0.259 ± 0.065	0.100 - 0.320	10
Vitamins			
Vitamin A (IU/kg)	$6,595 \pm 1,548$	4,180 - 11,450	25
Vitamin D (IU/kg)	$4,450 \pm 1,382$	3,000 - 6,300	4
α-Tocopherol (ppm)	35.43 ± 8.98	22.5 - 48.9	11
Thiamine (ppm)	18.16 ± 1.54	15.0 - 21.0	25
Riboflavin (ppm)	7.83 ± 0.923	6.10 - 9.00	11
Niacin (ppm)	99.22 ± 24.27	65.0 - 150.0	11
Pantothenic acid (ppm)	30.55 ± 3.52	23.0 - 34.6	11
Pyridoxine (ppm)	9.11 ± 2.53	5.60 - 14.0	11
Folic acid (ppm)	2.46 ± 0.63	1.80 - 3.70	11
Biotin (ppm)	0.268 ± 0.047	0.190 - 0.354	11
Vitamin B ₁₂ (ppb)	40.5 ± 19.1	10.6 - 65.0	11
Choline (ppm)	$2,991 \pm 382$	2,300 - 3,430	10
Minerals			
Calcium (%)	1.17 ± 0.10	1.00 - 1.49	25
Phosphorus (%)	0.93 ± 0.03	0.850 - 1.00	25
Potassium (%)	0.886 ± 0.063	0.772 - 0.971	9
Chloride(%)	0.529 ± 0.087	0.380 - 0.635	9
Sodium (%)	0.316 ± 0.033	0.258 - 0.371	11
Magnesium (%)	0.166 ± 0.010	0.148 - 0.181	11
Sulfur (%)	0.272 ± 0.059	0.208 - 0.420	10
Iron (ppm)	350.5 ± 87.3	255.0 - 523.0	11
Manganese (ppm)	92.48 ± 5.14	81.7 — 99.4	11
Zinc (ppm)	59.33 ± 10.2	46.1 - 81.6	11
Copper (ppm)	11.81 ± 2.50	8.09 - 15.4	11
Iodine (ppm) Chromium (ppm)	3.54 ± 1.19	1.52 - 5.83	10
Cobalt (ppm)	$\begin{array}{c} 1.66 \pm 0.46 \\ 0.76 \pm 0.23 \end{array}$	0.85 - 2.09 $0.49 - 1.15$	11 7
Copair (ppin)	0.70 ± 0.23	0.43 — 1.13	,

TABLE G4 Contaminant Levels in NIH-07 Rat and Mouse Ration^a

	$\begin{array}{cc} \textbf{Mean} \pm \textbf{Standard} \\ \textbf{Deviation}^{\text{b}} \end{array}$	Range	Number of Samples
Contaminants			
Arsenic (ppm)	0.37 ± 0.18	0.10 - 0.70	25
Cadmium (ppm)	0.10 ± 0.07	0.05 - 0.20	25
Lead (ppm)	0.30 ± 0.23	0.10 - 1.00	25
Mercury (ppm) ^c	0.02	0.02 - 0.03	25
Selenium (ppm)	0.33 ± 0.12	0.05 - 0.60	25
Aflatoxins (ppm)	< 5.0		25
Nitrate nitrogen (ppm) _d	11.72 ± 5.20	2.90 - 21.0	25
Nitrite nitrogen (ppm) ^d	0.23 ± 0.18	0.10 - 0.70	25
BHA (ppm) ^e	1.88 ± 1.94	1.00 - 10.0	25
BHT (ppm) ^e	1.56 ± 1.58	1.0 - 8.00	25
Aerobic plate count (CFU/g)	78,748 ± 143,028	4,100 - 710,000	25
Coliform (MPN/g)	3 ± 0.2	3 - 4	25
Escherichia coli (MPN/g)	< 3	0 1	25
Salmonella (MPN/g)	Negative		25
Total nitrosoamines (ppb) ^f	7.25 ± 1.71	4.80 - 11.40	25
N-Nitrosodimethylamine (ppb) ^f	5.50 ± 1.30	3.80 - 9.10	25
<i>N</i> -Nitrosopyrrolidine (ppb) ^f	1.75 ± 1.00	1.00 - 4.30	25
Pesticides (ppm)			
α-ВНС	< 0.01		25
β-ВНС	< 0.02		25
γ-BHC	< 0.01		25
δ-BHC	< 0.01		25
Heptachlor	< 0.01		25
Aldrin	< 0.01		25
Heptachlor epoxide	< 0.01		25
DDE	< 0.01		25
DDD	< 0.01		25
DDT	< 0.01		25
НСВ	< 0.01		25
Mirex	< 0.01		25
Methoxychlor	< 0.05		25
Dieldrin	< 0.01		25
Endrin	< 0.01		25
Telodrin	< 0.01		25
Chlordane	< 0.05		25
Toxaphene	< 0.10		25
Estimated PCBs	< 0.20		25
Ronnel	< 0.01		25
Ethion	< 0.02		25
Trithion	< 0.05		25
Diazinon	< 0.10		25
Methyl parathion	< 0.02		25
Ethyl parathion	< 0.02		25
Malathion	0.24 ± 0.21	0.05 - 0.97	25
Endosulfan I	< 0.01		25
Endosulfan II	< 0.01		25
Endosulfan sulfate	< 0.03		25

CFU= colony-forming units; MPN= most probable number; BHC= hexachlorocyclohexane or benzene hexachloride For values less than the limit of detection, the detection limit is given as the mean.

All but three values were less than the detection limit; the detection limit was used for the low end of the range. Sources of contamination: alfalfa, grains, and fish meal Sources of contamination: soy oil and fish meal All values were corrected for percent recovery.

APPENDIX H SENTINEL ANIMAL PROGRAM

METHODS		226
TABLE H1	Murine Virus Antibody Determinations for Rats and Mice	
	in the 2-Year Inhalation Studies of Ethylbenzene	227

SENTINEL ANIMAL PROGRAM

METHODS

Rodents used in the Carcinogenesis Program of the National Toxicology Program are produced in optimally clean facilities to eliminate potential pathogens that may affect study results. The Sentinel Animal Program is part of the periodic monitoring of animal health that occurs during the toxicologic evaluation of chemical compounds. Under this program, the disease state of the rodents is monitored via serology on sera from extra (sentinel) animals in the study rooms. These animals and the study animals are all subject to identical environmental conditions. The sentinel animals come from the same production source and weanling groups as the animals used for the studies of chemical compounds.

Serum samples were collected from randomly selected rats and mice during the 2-year studies. Blood from each animal was collected and allowed to clot, and the serum was separated. The samples were processed appropriately and sent to Microbiological Associates, Inc. (Bethesda, MD), for determination of antibody titers. The laboratory serology methods and viral agents for which testing was performed are tabulated below; the times at which the blood was collected during the studies are also listed.

Method and Test

Time of Analysis

RATS

ELISA

Mycoplasma arthritidis Mycoplasma pulmonis PVM (pneumonia virus of mice)

RCV/SDA

(rat coronavirus/sialodacryoadenitis virus)

Sendai

Hemagglutination Inhibition H-1 (Toolan's H-1 virus) KRV (Kilham rat virus)

Study termination Study termination

6, 12, and 18 months, study termination

6, 12, and 18 months, study termination 6, 12, and 18 months, study termination

6, 12, and 18 months, study termination 6, 12, and 18 months, study termination

MICE

ELISA

Ectromelia virus EDIM (epizootic diarrhea of infant mice) GDVII (mouse encephalomyelitis virus) LCM (lymphocytic choriomeningitis virus)

Mouse adenoma virus-FL MHV (mouse hepatitis virus)

M. arthritidis M. pulmonis PVM Reovirus 3

Sendai

6, 12, and 18 months, study termination 6 and 18 months, study termination

6, 12, and 18 months, study termination

6, 12, and 18 months, study termination

6, 12, and 18 months, study termination 6, 12, and 18 months, study termination

Study termination Study termination

6, 12, and 18 months, study termination 6, 12, and 18 months, study termination

6, 12, and 18 months, study termination

Method and Test

Time of Analysis

MICE (continued)

Immunofluorescence Assay

EDIM 6 and 12 months, study termination

GDVII 12 months
MHV 6 and 12 months
Mouse adenoma virus-FL Study termination
Reovirus 3 6 and 12 months
Sendai 6 months

Hemagglutination Inhibition

K (papovavirus) 6, 12, and 18 months, study termination

MVM (minute virus of mice) 6, 12, and 18 months, study termination

Polyoma virus 6, 12, and 18 months, study termination

Results of serology tests are presented in Table H1.

TABLE H1
Murine Virus Antibody Determinations for Rats and Mice in the 2-Year Inhalation Studies of Ethylbenzene

Interval	Incidence of Antibody in Sentinel Animals	Positive Serologic Reaction for
Rats		
6 Months	0/24	None positive
12 Months	0/24	None positive
18 Months	0/22	None positive
Study termination	1/10	Mycoplasma arthritidis ^a
Mice		
6 Months	0/10	None positive
12 Months	1/9	Reovirus 3
18 Months	0/8	None positive
Study termination	0/10	None positive

Further evaluation of the sample positive for *M. arthritidis* by immunoblot and Western blot procedures indicated that the positive titer may have been due to cross reaction with antibodies of nonpathogenic *Mycoplasma* or other agents. Only one sample was positive, and there were no clinical findings or histopathologic changes of *M. athritidis* infection in the rat with the positive titer. Accordingly, the *M. arthritidis* positive titer was considered to be a false positive.