Certificate for the approval in accordance with Council Directive 90/429/EEC of a semen collection centre as regards domestic animals of the porcine species

Name and address of the semen collection Centre	
Owner	
Person in charge	
Name and address of the responsible centre Veterinarian	
Name and address of the competent official Veterinarian	
I, the undersigned, certify that the semen collection centre detailed above has been inspected on the basis of the attached checklist and found in compliance with the requirements of Council Directive 90/429/EEC.	
Name and address of the central competent Authorities	
I, the undersigned, certify that the semen collection centre detailed above complies with the animal health requirements laid down in Council Directive 90/429/EEC for imports into the Community of porcine semen.	
Approval Date: [dd.mm.yyyy]	
Approval Number assigned to the centre	

Number	Reference	Question	Y= Yes N= No
1. General	criteria		1
1.1	90/429/EEC Council Directive Article 7 Paragraph 1	Is the country listed in Annex I to Commission Decision 2002/613/EC	
1.2	90/429/EEC Council Directive Article 14	Are the checks at origin carried out equivalent to the requirements laid down in Article 3 of Directive 90/425/EEC	
1.3	90/429/EEC Council Directive Article 10	Is a model animal health certificate available for imports into the EU of porcine semen from the country of dispatch	
1.4	90/429/EEC Council Directive Article 14	Are arrangements in place to pre-notify the arrival of a consignment at an approved border inspection post for the checks required in accordance with Council Directive 97/78/EC	
1.5	Article 15	Are the rules on disease notification established in Directives 90/425/EEC applied	
1.6	90/429/EEC Council Directive Annex A Chapter II Paragraph 1	Is the centre approved for dual purposes (national and EU –approved)	
2. Technic	al Conditions to be applied for se	emen collection centers	
2.1	90/429/EEC Council Directive Annex A Chapter I Paragraph 1	Is the centre placed under the permanent supervision of a centre veterinarian	
		Has the centre animal housing including facilities for the isolation of animals which have failed tests described in Annex B, Chapter II, or which show clinical signs of disease	
2.2	90/429/EEC Council Directive Annex A Chapter I Paragraph 2a,b,c,d	Has the centre semen collection facilities including a separate room for the cleaning and disinfection or sterilization of equipment	
		Has the centre a semen processing room which need not necessarily be on the same site Has the centre a semen storage room which	
2.3	90/429/EEC Council Directive Annex A Chapter I Paragraph 3	need not necessarily be on the same site Has the centre be so constructed or isolated that contact with livestock outside is prevented	
2.4	90/429/EEC Council Directive Annex A Chapter I Paragraph 4	Has the centre be so constructed that the animal housing and semen collection, processing and storage facilities can be readily cleaned and disinfected	
2.5	90/429/EEC Council Directive Annex A Chapter I Paragraph 5	Has the centre be so designed that the animal accommodation is physically separated from the semen processing room and both are separated from the semen storage room.	

	onditions to be applied for semen collect		
3.1	90/429/EEC Council Directive	Are only animals of the species whose	
	Annex A Chapter II Paragraph 1	semen is to be collected kept at the centre	
3.2	90/429/EEC Council Directive	Is a record, file or computer record kept of	
	Annex A Chapter II Paragraph2	all porcine animals at the centre, giving	
		details of the breed, date of birth and	
		identification of each of the animals, and	
		also a record, file or computer record of all	
		checks for diseases and all vaccinations	
		carried out, giving also information from	
		the disease/health file of each animal	
3.3	90/429/EEC Council Directive	Is the centre inspected by an official	
	Annex A Chapter II Paragraph 3	veterinarian, at least twice a year, at	
	Times II Chapter II I diagraph 3	which time checks on the conditions of	
		approval and supervision shall be carried	
		out	
3.4	90/429/EEC Council Directive		
3.4		Is the entry of unauthorized persons	
	Annex A Chapter II Paragraph 4	prevented? Furthermore, authorized visitors	
		must be required to comply with the	
		conditions laid down by the centre	
2.5	00/400/EEG.G. "1.5"	veterinarian	
3.5	90/429/EEC Council Directive	Has the staff been suitably trained in	
	Annex A Chapter II Paragraph 5	disinfection procedures and hygiene	
		techniques relevant to the control of the	
		spread of disease	
3.6	90/429/EEC Council Directive	Has all semen been collected, processed and	
	Annex A Chapter II Paragraph 6a	stored in approved centres, without coming	
		into contact with any other consignment	
		of semen	
3.7	90/429/EEC Council Directive	Is the collection, processing and storage of	
	Annex A Chapter II Paragraph 6b	semen carried out only in the premises set	
		aside for that purpose and under conditions	
		of the strictest hygiene	
3.8	90/429/EEC Council Directive	Are all implements which come into contact	
	Annex A Chapter II Paragraph 6c	with the semen or the donor animal during	
		collection and processing are properly	
		disinfected or sterilized prior to use	
3.9	90/429/EEC Council Directive	Are products of animal origin used in the	
	Annex A Chapter II Paragraph 6d	processing of semen — including additives	
	Times II chapter II I diagraph 00	or diluents obtained from sources which	
		present no animal health risk or are so	
		treated prior to use that such risk is	
3.10	90/429/EEC Council Directive	Are storage flacks and transport flacks	
3.10		Are storage flasks and transport flasks	
	Annex A Chapter II Paragraph 6e	properly disinfected or sterilized before the	
2 11	00/420/EEG G 'I D' '	beginning of each filling operation	
3.11	90/429/EEC Council Directive	Had cryogenic agents been previously used	
2.12	Annex A Chapter II Paragraph 6h	for other products of animal origin	
3.12	90/429/EEC Council Directive	Is each collection of semen, whether or	
	Annex A Chapter II Paragraph 6g	not it is separated into individual doses,	
		clearly marked as follows: date of	
		collection, breed, identity of the donor	
		animal, name and registration number of	
		centre	
3.13		A (11) 1 11 14 (1 111)	
3.13	90/429/EEC Council Directive	Are antibiotics added to the diluent or	

11		Number of boars in the center:	
4.1			
4.2		All animals admitted to the centre:	
4.2.1	00/400/EPG G	Have been subjected to a period of	
	90/429/EEC Council Directive	quarantine of at least 30 days in	
	Annex B Chapter I Paragraph 1a	accommodations specifically approved for	
		the purpose by the competent authority of	
		the Member State, and where only animals	
		having at least the same health status are	
4.3		Prior to their entering the quarantine	
4.3		accommodation, all animals have been	
		chosen from herds or holdings:	
4.3.1		which are free of brucellosis in accordance	
4.3.1		with the OIE Terrestrial Animal Health	
		Code	
4.3.2		in which no animal vaccinated against foot	
7.0.2		and-mouth disease has been present in the	
	90/429/EEC Council Directive	preceding 12 months,	
4.3.3	Annex B Chapter I Paragraph 1b	in which no clinical, serological or	
11010		virological evidence of Aujeszky's	
		disease has been detected in the preceding	
		12 months,	
4.3.4		which are not situated in a restricted area	
		defined under the provisions of the	
		Community legislation due to the	
		emergence of a disease in domestic pigs.	
4.4		Before the period of quarantine specified	
		in 4.2.1 (above), and within the 30 days	
		previous to entering quarantine, have	
		been subjected to the following tests,	
		performed in accordance with standards	
		laid down in relevant Directives, with	
	90/429/EEC Council Directive	negative results:	
4.4.1	Annex B Chapter I Paragraph 1c	a complement fixation test or a buffered	
		brucella antigen test in respect of	
		brucellosis (from 1 January 2001, the	
		buffered brucella antigen test will be the	
4.4.2		only authorized test)	
4.4.2		a serum neutralization or an ELISA test	
		using all the Aujeszky's disease viral	
		antigens in the case of non-vaccinated pigs	
		or an ELISA test for Aujeszky's disease G1	
		antigens in the case of pigs vaccinated with a G1 deleted vaccine	
112		an ELISA test or a serum neutralisation test	
4.4.3.			
		for the presence of classical swine fever OR the US is free of classical swine fever	

4.5		During the last 15 days of the period of quarantine specified in 4.2.1 (above) of at least 30 days the animals have been subjected to the following tests with negative results:	
4.5.1	90/429/EEC Council Directive Annex B Chapter I Paragraph 1d	in respect of brucellosis, a complement fixation test or a buffered brucella antigen test (from the l January 2001 the buffered brucella antigen test will be the only authorized test)	
4.5.2		a serum neutralization or an ELISA test using all the Aujeszky's disease viral antigens in the case of non-vaccinated pigs, or an ELISA test for Aujeszky's disease G1 antigens in the case of pigs vaccinated with a G1 deleted vaccine	
4.6	90/429/EEC Council Directive Annex B Chapter I Paragraph 2	Are all test carried out in a laboratory approved by the competent authority	
4.7	90/429/EEC Council Directive Annex B Chapter I Paragraph 3	Are animals only admitted to the semen collection centre with the express permission of the centre veterinarian and all animal movements, both in and out, are recorded	
4.8	90/429/EEC Council Directive Annex B Chapter I Paragraph 4	No animal admitted to the semen collection centre may show any clinical sign of disease on the day of admission; all animals must, without prejudice to paragraph 5, have come directly from quarantine accommodation as referred to in 4.2.1 (above) which, on the day of consignment, officially fulfils the following conditions: (a) it is not situated in a restricted area defined under the provisions of the Community legislation due to the emergence of a disease in domestic pigs; (b) no clinical, pathological or serological evidence of Aujeszkys disease has been recorded for the past 30 days	