



Chemical analyses of pre-Holocene rocks from Medicine Lake Volcano and vicinity, northern California

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2008

Open-File Report 2008–1094

**U.S. Department of the Interior
U.S. Geological Survey**

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1. Chemical analyses of pre-Holocene rocks

ABSTRACT

Chemical analyses are presented in an accompanying table (Table 1) for more than 600 pre-Holocene rocks collected at and near Medicine Lake Volcano, northern California. The data include major-element X-ray fluorescence (XRF) analyses for all of the rocks plus XRF trace element data for most samples, and instrumental neutron activation analysis (INAA) trace element data for many samples. In addition, a limited number of analyses of Na₂O and K₂O by flame photometry (FP) are included as well as some wet chemical analyses of FeO, H₂O+/-, and CO₂. Latitude and longitude location information is provided for all samples. This data set is intended to accompany the geologic map of Medicine Lake Volcano (Donnelly-Nolan, in press); map unit designations are given for each sample collected from the map area.

INTRODUCTION

Medicine Lake Volcano (MLV) is a large Pleistocene and Holocene volcano located in the northern California Cascade Range (Figure 1). In conjunction with geologic mapping of the volcano, rocks were collected for chemical analysis. Analyses from all Holocene map units have been published (Grove and Donnelly-Nolan, 1986; Donnelly-Nolan and Champion, 1987; Grove and others, 1988; Grove and others, 1997; Kinzler and others, 2000). Donnelly-Nolan and others (2007) present the map distribution and revised ages of Holocene and early postglacial lavas. Chemical analyses of early postglacial lavas can be found in Donnelly-Nolan and Champion (1987), Baker and others (1991), and Donnelly-Nolan and others (1991). A limited number of analyses of older lavas were published in Donnelly-Nolan and Champion (1987), Wagner and others (1995), and Elkins Tanton and others (2001); some of these analyses of older lavas are also included in Table 1, but most of the data in Table 1 have not previously been published.

Column A of Table 1 gives sample numbers in consecutive order within each section of the table. All samples were collected by the author with the exception of samples 2B193, 4B543, B8719, and 5B320 which were collected by D.E. Champion as drill cores during sampling for paleomagnetic analysis. Column B gives the designation M (for Medicine Lake) plus in some cases an additional letter. Most such samples are

pumice lumps collected from unit **dta**, with some exceptions. Samples 1101M-A and -B and 1471M-A and -B are, respectively, porphyritic and finer-grained pairs of samples collected from single outcrops. Samples 1694M-A,B,C are respectively the lower, middle, and upper tephra collected at one roadcut locality. Samples 828M-B and -C are respectively the white and black tephra collected from a quarry where samples 1501M-1503M were also collected.

The accompanying table (Table 1) is divided into three sections, first the chemical analyses of mapped units of MLV; second, units that pre-date MLV but were collected from older units identified on the geologic map; third, other volcanic rocks (all pre-MLV in age) in the area surrounding MLV. Map unit symbols (Donnelly-Nolan, in press) are given in column C.

Latitude and longitude location information is given as minutes and decimal seconds for each sample. In column D, locations are provided with respect to 41 degrees north latitude; in column E with respect to 121 degrees west longitude. All samples were collected within the area shown in Figure 1. However, only a few samples were located using GPS (Global Positioning System) technology. Most samples were located by inspection on 15-minute topographic maps, the only topographic maps available during most of the time period when the rocks were collected. Locations were then measured by hand and typed into the spreadsheet of data. Thus, locations may be only approximate, and some may be in error.

Analytical data are tabulated in columns F through BG. Most of the data were provided to the author on typewritten forms, and then typed by the author into the spreadsheet. The data have been checked, but errors in transcription are possible.

METHODS

Rock samples were collected from outcrops and typically were broken up in the field into 1 to 2 cm fragments using a hammer, and later chipped into <0.5 cm fragments in a small alumina jaw crusher. Samples were powdered in an alumina shatterbox and subsequently analyzed in USGS analytical laboratories in Lakewood, Colorado, in Menlo Park, California, and in Reston, Virginia.

Major elements were analyzed by wave-length X-Ray fluorescence (XRF; Taggart and others, 1987) in Lakewood by J.S. Wahlberg, J. Taggart, J. Baker, A.J. Bartel, K. Stewart, D.F. Siems, and J.S. Mee. In the original major element analyses, all iron was reported as Fe_2O_3 . The normalized analyses presented in columns F to O were calculated by first multiplying the iron value reported as Fe_2O_3 by 0.9 to calculate all iron as FeO. Then all major elements were summed without LOI (Loss on Ignition, column Q), resulting in the “original total” (column P). Subsequently, the major elements were recalculated to 100 percent total, volatile-free. The resulting normalized analyses (columns F to O) thus are calculated with all iron as FeO (in column H, indicated as FeO*). Note that P_2O_5 data with values of 0.00 wt. % in column N were originally reported as <0.05 wt. %.

XRF trace elements reported in Table 1 (columns X to AG) were analyzed by energy-dispersive XRF (Kevex; Siems, 2000) at the chemical analysis laboratory in Menlo Park, California, by P. Bruggman, B. King, and J. Kent, and subsequently in Lakewood by D. Siems, although some data were generated by H.J. Rose, J.R. Lindsay, R. Johnson, B. McCall, G. Sellers, and D. Burgi by wave-length dispersive XRF in Reston, Virginia. At low concentrations, some of these data (e.g. Nb) have large errors.

Flame photometry (FP) analyses of Na_2O and K_2O (columns R and S) were obtained for a subset of the samples; these analyses were performed by L. Espos, P. Klock, and T. Fries in Menlo Park, California. Also presented in Table 1 are a limited number of wet chemical analyses of FeO, H_2O^+ , H_2O^- , and CO_2 (columns T to W) in weight percent, performed in Lakewood by L. Jackson, G. Mason, and J. Ryder, and in Menlo Park by W. Updegrave, S. Neil, L. Espos, P. Klock, and T. Fries.

Instrumental Neutron Activation trace element analyses (INAA; Budahn and Wandless, 2002) are presented in columns AH to BG. The analyses were performed in Reston by C. Palmer, G. Wandless, P. Baedecker, J.S. Mee, and L. Schwarz, and in Denver by D. McKown, J. Budahn, R. Knight, and H. Millard, Jr.

Precision and accuracy of similar data generated for Crater Lake lavas are reported and discussed in Bacon and Druitt (1988).

ACKNOWLEDGMENTS

Funding for this work was provided by the USGS Geothermal and Volcano Hazards Programs. I thank W. Loskutoff and others for help with sample preparation. D. Ramsey prepared Figure 1.

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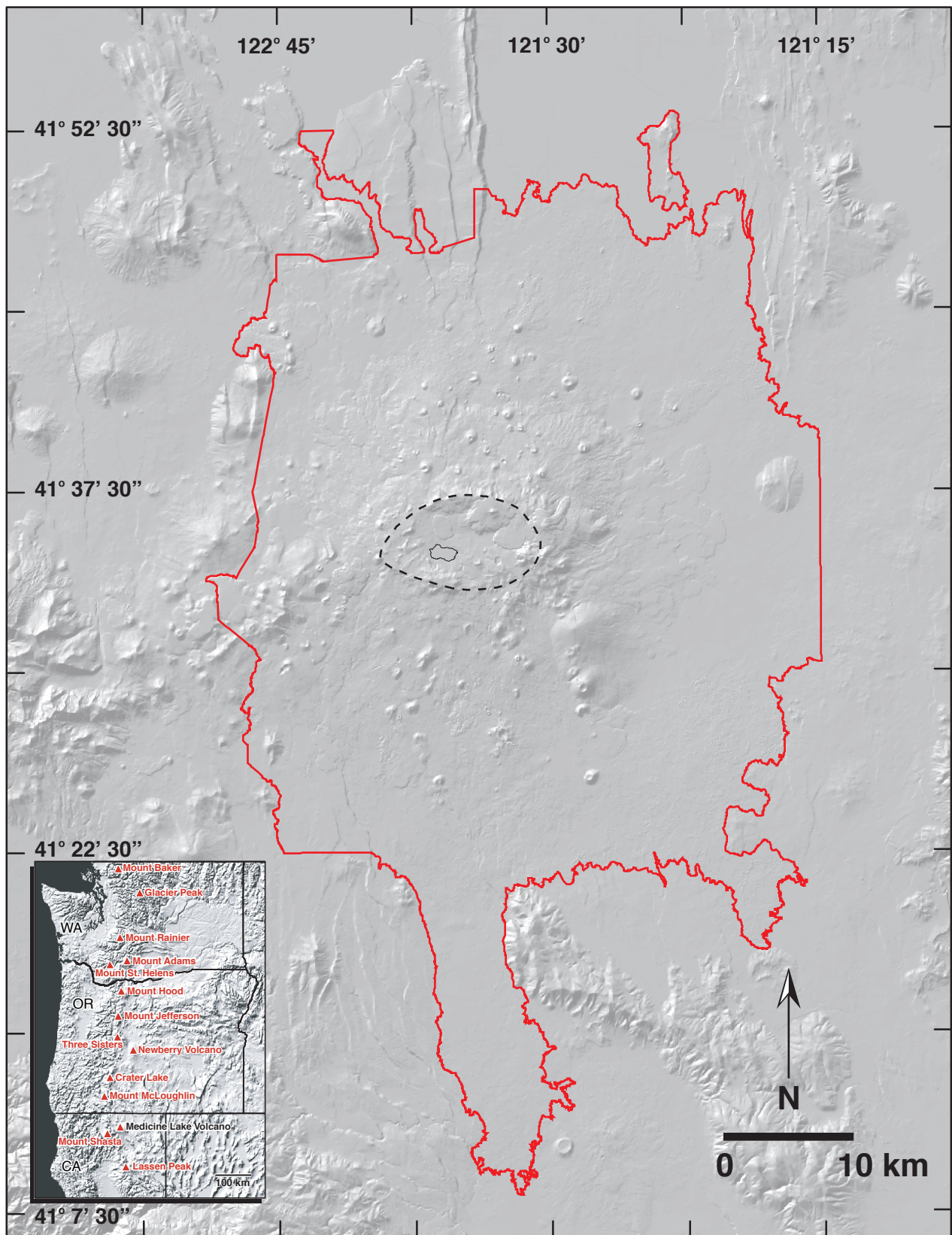


Figure 1. Map showing area in which samples reported in Table 1 were collected. Red outline indicates area of geologic mapping of Medicine Lake Volcano (Donnelly-Nolan, in press). Dashed black line is rim of Medicine Lake caldera. Inset map shows major Cascade volcanoes; Medicine Lake Volcano is indicated with black text.