

Chemical Data for Flows and
Feeder Dikes of the Yakima Basalt Subgroup,
Columbia River Basalt Group,
Washington, Oregon, and Idaho,
and their Bearing on a Petrogenetic Model

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By THOMAS L. WRIGHT, MARGARET MANGAN, and
DONALD A. SWANSON

DEPARTMENT OF THE INTERIOR
DONALD PAUL HODEL, Secretary

U.S. GEOLOGICAL SURVEY
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CONTENTS

Abstract	1
Introduction	2
Acknowledgments	2
Stratigraphic nomenclature and geochemical units	3
General geologic relations	5
Vents for flows of the Yakima Basalt Subgroup	6
Chemical study of the Yakima Basalt Subgroup	6
Methods of analysis	6
Presentation of chemical data	8
Petrography	9
Classification and naming of chemical types	9
Bulk rock chemistry: observations	10
Grande Ronde Basalt	10
Wanapum Basalt	11
Saddle Mountains Basalt	11
Comparison of dike and flow chemistry	12
Glass chemistry: observations	12
Discussion: implications for the origin of the Yakima Basalt Subgroup	13
Emplacement of flows	13
Magma storage before eruption	13
Nature of the source mantle	14
Inferences based on geochemical data	14
Inferences based on tectonic history	16
Summary and synthesis: magma generation, storage, and eruption of the Yakima Basalt Subgroup	16
Previous hypotheses	16
Synthesis	18
References cited	19

FIGURES

1. Index map of Columbia River Plateau	3
2. Stratigraphy of the Columbia River Basalt Group	5
3. Location of linear vent systems, Yakima Basalt Subgroup	7
4. MgO variation diagrams, Yakima Basalt Subgroup	46
5. Thorium variation diagrams, Yakima Basalt Subgroup	58
6. Rare-earth-element data (chondrite normalized) for Yakima Basalt Subgroup	66
7. Isotope data, Yakima Basalt Subgroup	69
8. MgO variation diagrams, comparing rock and compositions, Yakima Basalt Subgroup glass	70
9. Patterns of chondrite-normalized abundances of rare-earth elements in additional analyzed flows	71

TABLES

1. Relation between stratigraphic position and chemistry, Yakima Basalt Subgroup, Columbia River Basalt Group	4
2. Average analytical differences among three laboratories	8
3. Average chemical composition of geochemical units in the Grande Ronde Basalt	24
4. Average chemical composition of geochemical units in the Wanapum Basalt	30
5. Average chemical composition of geochemical units in the Saddle Mountains Basalt	31
6. Average chemical composition of feeder dikes in the Yakima Basalt Subgroup	34
7. Ratios of incompatible trace elements in samples of the Grande Ronde Basalt	35

8.	Ratios of incompatible trace elements in samples of Wanapum Basalt	36
9.	Ratios of incompatible trace elements in samples of Saddle Mountains Basalt	37
10.	Corrected and normalized major-oxide chemistry of naturally quenched glass from the Grande Ronde Basalt	38
11.	Corrected and normalized major-oxide chemistry of naturally quenched glass from the Wanapum Basalt	38
12.	Corrected and normalized major-oxide chemistry of naturally quenched glass from the Saddle Mountains Basalt	38
13.	Summary of chemical characteristics for geochemical units in the Yakima Basalt Sub-group	39
14.	Relation between stratigraphic position, chemistry, and additional flows, for which chemical data are available, Columbia River Plateau	41
15.	Average chemical composition of additional analyzed flows of table 14	42
16.	Ratios of incompatible trace elements in additional analyzed flows of table 14	44
17.	Summary of chemical characteristics of additional flows for which chemical data are available, Columbia River Plateau	45

Chemical Data for Flows and Feeder Dikes of the Yakima Basalt Subgroup, Columbia River Basalt Group, Washington, Oregon, and Idaho, and their Bearing on a Petrogenetic Model

By Thomas L. Wright, Margaret Mangan, and Donald A. Swanson

Abstract

Regional geologic mapping of the Yakima Basalt Subgroup of the Columbia River Basalt Group provides for a threefold division of the subgroup—from oldest to youngest, the Grande Ronde Basalt, Wanapum Basalt, and Saddle Mountains Basalt. Systems of feeder dikes and vents trending north to north-northwest have been identified for most units. Regional flow correlations, made on the basis of geologic mapping, major-oxide chemistry, and magnetic polarity, show that many flows have volumes of more than 500 km³, traveled hundreds of kilometers from their vents, and had emplacement times of a few days to a few weeks.

Samples of the subgroup were extensively analyzed for major oxides, trace elements, and isotopes of Sr, εNd, Pb, and O. Chemical analyses allow recognition of geochemical units consisting of a flow or several related flows of near-uniform composition, some of which represent chemical types repeated throughout the section. Average compositions of geochemical units have been plotted and are used in drawing petrogenetic inferences. All regionally extensive flows are tholeiitic and have MgO contents in the range of 2.5–8.0 percent. Each formation is characterized by specific chemical compositions. Flows of the Grande Ronde Basalt are aphyric and have low MgO (3–6 percent), high SiO₂ (52–58 percent), and relatively low TiO₂ (1.5–2.5 percent) and FeO (9–13 percent) contents and a relatively low initial ⁸⁷Sr/⁸⁶Sr isotopic ratio (0.703–0.705). Most flows of the Wanapum Basalt have lower SiO₂ (49–52 percent) for the same range of MgO, much higher contents of TiO₂ (>3 percent) and FeO (13–15 percent), and somewhat higher initial ⁸⁷Sr/⁸⁶Sr (0.704–0.7055). Trace elements are similar overall in both formations despite the contrasts in major-oxide chemistry. Chondrite-normalized rare-earth patterns for both formations show enrichment in light rare earths, are slightly concave upward with small or no negative europium anomalies, and at differing degrees of absolute rare-earth content are parallel to slightly convergent toward the heavy rare earths.

The Saddle Mountains Basalt contains members spanning a wide range of major-oxide composition, including some whose major-oxide compositions match those of older units. With few exceptions, incompatible trace elements are

enriched and compatible elements (particularly Sr and Sc) depleted relative to older formations, and values of the ⁸⁷Sr/⁸⁶Sr ratio are higher (>0.707). Chondrite-normalized rare-earth patterns for many of the ten members of the Saddle Mountains Basalt are steeper and show a more pronounced negative europium anomaly than those of the older formations. However, one member (the Umatilla Member) has a pronounced positive europium anomaly.

Trace-element variation, represented by interelement ratios or absolute abundances plotted against MgO content, is less coherent in the Yakima Basalt Subgroup than in oceanic basalts. Some major-oxide compositional trends suggest that crystal-liquid fractionation occurred, but several other lines of evidence, both geologic and geochemical, argue against low-pressure crustal storage and fractionation. This evidence includes the divergence of chemical trends for coexisting rock and glass from those relating one bulk rock composition to another, inconsistent incompatible trace-element enrichments, and the absence of major structural depressions such as grabens and calderas that should be associated with the virtually instantaneous eruption of the huge volumes represented by individual units. Major- and trace-element variation likewise is difficult to reconcile with any simple model of high-pressure fractionation, melting from a common source, or mixing of magma with crustal material, although some combination of these processes may have been important for some units.

Interpretation of the petrogenesis of the Yakima Basalt Subgroup requires consideration of such diverse topics as the tectonic history of the Pacific Northwest, generalized models of flood-basalt genesis, and the process of continental crustal underplating, in addition to the chemical and isotopic data. Recent geophysical studies and deep drill holes show that the Columbia River Basalt Group is underlain by upper Mesozoic and lower Cenozoic sedimentary rocks that define a large preexisting sedimentary basin. This depression may have been created as a back-arc basin when the older continental margin and associated subduction zone moved westward away from a once-continuous belt of plutonic rocks represented in the Pacific Northwest by the Idaho batholith. We postulate that large volumes of magma were generated in a mixed oceanic-continental source mantle and were emplaced

as sill-like bodies at the base of the continental crust. These bodies were then progressively tapped during deep continental rifting and supplied magma to the surface via extensive linear vent systems. All of the magma is inferred to have equilibrated with clinopyroxene during either melting or storage. Recent experimental study of the Grande Ronde Basalt at pressures to 3 GPa suggests an origin by direct partial melting of a source related to mid-ocean ridge basalt and its source mantle, possibly at a depth of 70 km.

The earliest basalt of regional extent, the Grande Ronde Basalt, began to fill the basin with lava derived from fairly young oceanic crust and mantle contaminated with subducted oceanic sediment. Such magma was then repeatedly erupted after storage near the crust-mantle boundary and minor interaction with lower-crustal rocks during ascent. The Wanapum Basalt formed in much the same way, although from a part of the oceanic mantle that had undergone Fe-Ti-P metasomatism. The similarity of the major-oxide chemistry of the Saddle Mountains Basalt to that of older units suggests a source related in some way to that of the Grande Ronde and Wanapum Basalts. However, the different trace-element abundances and ratios require additional complexity, such as melting of a 2.5-Ga continental mantle having rather high incompatible element contents or melting of mantle that had undergone a second metasomatic event.

Eruption rates generally kept pace with continued basining during formation of the Yakima Basalt Subgroup. At times, the basin became filled and large-volume flows continued through the ancient Columbia River Gorge to the Pacific Ocean. At other times, continued subsidence of the central plateau, possibly combined with uplift of the eastern margin of the plateau, created a gentle westward-dipping slope. Volcanism waned around 14 Ma and finally stopped at about 6 Ma, though the weight of the basalt pile caused continued subsidence as isostatic equilibrium was reestablished.

INTRODUCTION

The Columbia River Basalt Group forms the youngest known flood-basalt province; eruptions occurred over an 11-m.y. period (17–6 Ma) in the early to late Miocene (Watkins and Baksi, 1974; McKee and others, 1977, 1981; Hooper, 1982, fig. 2; Swanson and others, 1979a). The group contains more than 170,000 km³ of tholeiitic basalt and covers an area of 160,000 km² to a maximum depth of at least 3 km, as inferred from surface exposures (fig. 1; Tolan and others, 1987), drill-hole data (Lingley and Walsh, 1986), and geophysical surveys (Stanley, 1984; Glover and others, 1985). The basalt is underlain by sedimentary rocks 4–5 km thick (Stanley, 1984; Glover and others, 1985), which in turn overlie crystalline basement that extends to the base of the crust, estimated to be 40 km deep (Catchings and others, 1984). Mapping of the province is complete; regional map compilations are by Swanson and others (1979b, 1980, 1981), Anderson and others (in press), and Hooper and Swanson (in press) and references therein. Wright and others (1973) summarized major-oxide chemistry of important units and used those data to define several chemical types. Trace-element data for selected units were published by Nathan and Fruchter (1974) and by the NASA-sponsored Basaltic Volcanism

Study Project (1981). Subsequently, we have defined additional chemical types and analyzed trace elements for all chemical types defined by major oxides. Many of these data are contained in open-file reports (Wright and others, 1979, 1980, 1982); a summary is given by Swanson and Wright (1981). Isotopic data for Sr, ϵ Nd, Pb, and O are summarized by McDougall (1976), Nelson (1980; 1983), Carlson and others (1981), Carlson (1984a), and Church (1985).

The purposes of this paper are (1) to summarize the chemical composition of all units of the Yakima Basalt Subgroup (table 1) mapped by two of us (DAS and TLW) in Washington and northeastern Oregon and (2) to evaluate implications for the origin of the Yakima Basalt Subgroup from available geologic, petrographic, geochemical, and geophysical data. A definitive petrogenesis awaits quantitative modeling of chemical data using the results of experimental study of representative flow compositions. Such studies are currently underway by Eiichi Takahashi, and preliminary results are available (Takahashi, 1988). Ultimately, any quantitative petrogenetic model of the complex processes that gave rise to the basalt must be based on significantly improved understanding of the nature of the upper mantle and lower crust; such an understanding probably remains far in the future.

The descriptive sections of the main body of this paper give (1) summary tabulations and graphical representations of the major-oxide, trace-element, and isotope chemistry of the flows; (2) chemistry of feeder-dike systems for many units; and (3) a comparison of major-oxide chemistry of regionally extensive flows with that of glass selvages from associated dikes or pillows. Discussion sections specify the critical observations that delimit possible conditions of magma generation, storage, and eruption and summarize the tectonic and magmatic evolution of the Yakima Basalt Subgroup. Tables 14–17 summarize chemical data for additional flows collected by us that either have been mapped in more detail by other workers or are interlayered with, but not considered by other workers to be a part of, the Columbia River Basalt Group. These flows include local flows in the Clearwater embayment of Idaho mapped by Camp (1981) and local flows in northeastern Oregon mapped by P.R. Hooper and W.T. Taubeneck (unpublished work). Both groups of flows are of Saddle Mountains age (table 14). With one exception, flows of the former group are considered to belong to the Yakima Basalt Subgroup. Flows of the latter group are, again with one exception, mapped separately by Hooper (1984) as the basalts of Powder River, which he excludes from the Columbia River Basalt Group.

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STRATIGRAPHIC NOMENCLATURE AND GEOCHEMICAL UNITS

Swanson and others (1979a; table 1 and fig. 2) worked out a revised stratigraphy for the Columbia River Basalt Group (fig. 2). The Grande Ronde Basalt has no formally named members but is divided for purposes of mapping into four magnetostratigraphic units: from oldest to youngest, R_1 , N_1 , R_2 , and N_2 . The Wanapum Basalt has four formally named members: from oldest to youngest, the Eckler Mountain, Frenchman Springs, Roza, and Priest

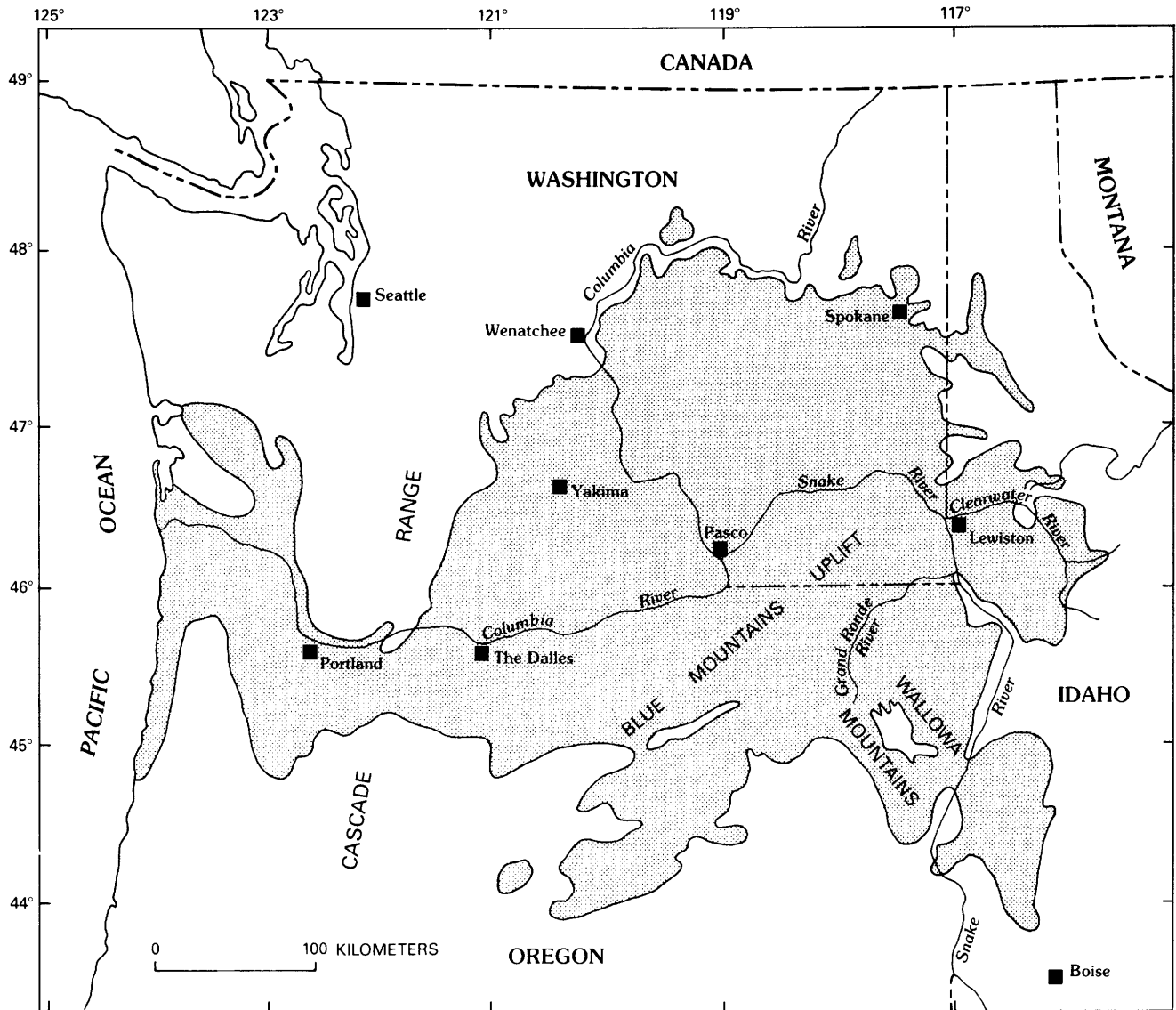


Figure 1. Index map of Columbia River Plateau. Outline of area covered by Columbia River Basalt Group (shaded) is modified from that given in Swanson and others (1979a) to include results of recent mapping west of Cascade Range (see Tolan and others, 1984, fig. 2). Compilation of mapping east of Cascade Range has resulted in slight modification of rest of boundary but not to an extent to affect material presented in this paper.

Table 1. Relation between stratigraphic position and chemistry, Yakima Subgroup, Columbia River Basalt Group

[MP, magnetic polarity as determined in the field using a fluxgate magnetometer; N, normal; R, reversed; T, transitional. Magnetic polarity units after Swanson and others (1979a). Geochemical units of the Grande Ronde Basalt are defined in Mangan and others (1985). Units marked with an asterisk (*) are defined from sections located in the western plateau]

Formation				Formation--Continued			
Member				Magnetostratigraphic unit			
Flow (Informal Name)	Age (Ma)	MP	Geochemical unit		Age (Ma)	MP	Geochemical unit
SADDLE MOUNTAINS BASALT				GRANDE RONDE BASALT	16.5-14.4		
Lower Monumental	6	N	LM			N	JOSEPH ¹ GR INC
Ice Harbor	8.5			Magnetostratigraphic unit N2		N	1A
Basalt of Goose Island		N	GOOSE				1A*
Basalt of Martindale							2A
Indian Memorial flow		N	INDIAN				UNC
Martindale flow		R	MARTIN				3A
Basalt of Basin City		N	BASIN				3A*
							UNC
Buford		N	BUFORD				4A
							4A*
Elephant Mountain	10.5	N,T	ELEPHANT				UNC*
Wenaha flow (Walker, 1973)							3B
							UNC
Pomona	12.0	R	POMONA				4B
Basalt of Weippe (Camp, 1981)		R	VC WEIPE				LH2
							MG6
Esquatzel		N	ESQUAT				2B
		N	RB ESQUAT				5A
		R	FS IC				5A*
Weissenfels Ridge							UNC
Basalt of Slippery Creek		N	SLIP				2C
Basalt of Lewiston Orchards		N	LEW ORCH				4C
Sprague Lake flow			SPRAGUE				5B
			SW SPRAG				3C
Anatone flow (Price, 1974)			NEW				3C*
			SW NEW	Magnetostratigraphic unit N2		N	UNC
Asotin		N	ASOTIN			T	5C
			VC ASOT			T	5C*
Huntzinger flow (Ward, 1976)		N	SW HUNTZ	Magnetostratigraphic unit R2		R	3D
			HUNTZ				5D
Basalt of Lapwai		N	VC LAP				5D*
Eagle Lake flow		N	EAGLE				UNC
							2D
Wilbur Creek		N	WILBUR				2D*
							UNC
Umatilla		N	UMATILLA	Magnetostratigraphic unit R2		R	AS16
			UM FRAC				
Umatilla flow			PH UMTIL			T	2E
Sopher Ridge flow			PH SOPHER				
WANAPUM BASALT				Magnetostratigraphic unit N1		N	UNC
							KB5
Priest Rapids		R	PV				3E
			LOLO INC				UNC
			ROSALIA				3F
							5E
Powatka flow		T,R	PH POWAT				AS16
							3G
							5F
Roza		N2	ROZA				UNC
							AS16
Frenchman Springs		N2	FS ANOM				5G
			FS INC				AS16
Eckler Mountain		N2					5H
Basalt of Shumaker Creek			SHUMAKER				AS16
			PH SHUM				UNC
Lookingglass flow			SW LOOK	Magnetostratigraphic unit N1		N	AS16
Basalt of Dodge			DODGE				
Basalt of Robinette Mountain			ROBIN	Magnetostratigraphic unit R1		R	2F
							5I
							AS16
				IMNAHA (C-3)	17-16	R	

¹ The Joseph volcanics unit of Kleck (1976) is composed of a series of glassy shield volcanoes sitting on top of the Grande Ronde Basalt in northeastern Oregon. Their chemistry suggests affinity with the Grande Ronde Basalt.

Rapids Members. Beeson and others (1985) have shown that there is no interbedding between these two formations, though such had been inferred by Swanson and others (1979a). Neither set of authors found evidence of a magnetic polarity reversal between Grande Ronde and Wanapum times. Accordingly, we believe that the Eckler Mountain and Frenchman Springs Members of the Wanapum Basalt fall in magnetostratigraphic unit N₂, as does the upper part of the Grande Ronde Basalt; the Roza Member is transitional from unit N₂ into unit R₃, and the Priest Rapids Member lies within unit R₃. The Saddle Mountains Basalt is divided into ten members (fig. 2), all of which are separated by erosional unconformities. The magnetostratigraphy is probably discontinuous; thus the polarity of each unit is designated in table 1 as N or R without a subscript.

Each member of the formations of the Yakima Basalt Subgroup (fig. 2) is composed of several flows, or sequences of flows, each of which has a distinctive major-oxide composition. The average chemistry of each flow sequence defines a geochemical unit of a certain chemical type¹, which is given an abbreviation related to its stratigraphic position (table 1). Some of the names of such units relate to locally mappable units defined since 1979 (see columns headed "flow" and "informal name" in table 1). Regional correlation of flows is based on geologic mapping, flow chemistry, and paleomagnetic measurements (compare Choiniere and Swanson, 1979; Hooper, 1981; Mangan and others, 1986). The stratigraphic nomenclature, chemical designation, radiometric age, and magnetic polarity of each unit in the Yakima Basalt Subgroup are given in table 1.

GENERAL GEOLOGIC RELATIONS

The regional distribution of formations and members in the Yakima Basalt Subgroup was originally given by Swanson and others (1979a, plate 1); distribution maps for individual members have been updated by Anderson and others (in press), and the volumes of the formations that make up the Yakima Basalt Subgroup have been newly estimated by Tolan and others (1987). Flows of the Grande Ronde Basalt cover most of the Columbia Plateau. Products of individual eruptions range from plateau-covering basaltic floods to smaller flows of more local distribution (Mangan and others, 1985, 1986). Flow contacts in the Grande Ronde are generally tight; a few thin and discontinuous intercalations of sedimentary and tuffaceous depos-

its occur, principally near the margins of the Plateau. The regional extent of the Wanapum Basalt is nearly as great as that of the Grande Ronde Basalt. Both formations contain flows or sequences of flows with volumes in excess of 500 km³; examples include the Roza Member, the Priest Rapids Member (LOLO geochemical unit); Grande Ronde Basalt, unit 4a*. Maximum thicknesses where flows were ponded are nearly 100 m, as estimated from surface exposures and drill holes.

A saprolite separates the Grande Ronde and Wanapum Basalts in the central part of the Columbia Plateau. The saprolite thickens toward the margins of the Plateau, where it may underlie or be replaced by sedimentary deposits (the Vantage Member of the Ellensburg Formation in the western Plateau and the Latah Formation in the eastern Plateau). The time interval during which the Grande Ronde Basalt was weathered was apparently not great, for no evidence

Group	Sub-group	Formation	Member
Columbia River Basalt Group	Yakima Basalt Subgroup	Saddle Mountains Basalt	Lower Monumental Member Ice Harbor Member Buford Member Elephant Mountain Member Pomona Member Esquatzel Member Weissenfels Ridge Member Asotin Member Wilbur Creek Member Umatilla Member
		Wanapum Basalt	Priest Rapids Member Roza Member Frenchman Springs Member Eckler Mountain Member
		Grande Ronde Basalt	No members defined (see table 1 for paleomagnetic and chemical subdivision)
	Picture Gorge Basalt	Innaha Basalt	No members defined

¹The term "geochemical unit" as used in this paper is similar to but not identical to the term "chemical type" used previously. The former term carries with it a specific stratigraphic as well as chemical meaning. The Grande Ronde Basalt is the only unit where chemical types, originally defined for one stratigraphically bound set of flows of similar chemistry are repeated throughout the stratigraphic section and thus are properly referred to as "chemical types," retaining the chemical definition while dropping the stratigraphic meaning; see, for example Mangan and others (1986).

Figure 2. Stratigraphy of the Columbia River Basalt Group, after Swanson and others (1979a).

for a polarity reversal has been recognized between Grande Ronde and Wanapum times.

The Saddle Mountains Basalt began to be erupted after an interval of erosion lasting perhaps as long as one million years. Saddle Mountains time extends from approximately 13 to 6 Ma, and significant time breaks separate the eruptions of each member of the Saddle Mountains (table 1). Many flows of the Saddle Mountains Basalt are preserved as remnants of valley-filling flows on the walls of present-day canyons and must originally have flowed down deep canyons cut into the Grande Ronde and Wanapum Basalts in the eastern part of the Plateau (Swanson and others, 1975a). Some individual flows are as voluminous as those in the Wanapum and Grande Ronde Basalts, and many traveled as far or farther because of channeling in river valleys. However, the Saddle Mountains Basalt covered less of the Plateau surface than did the earlier units.

VENTS FOR FLOWS OF THE YAKIMA BASALT SUBGROUP

All flows were erupted from fissures and fed by dikes. Several outcrops are known in which a dike merges into the flow it fed (Swanson and others, 1975b). The location of a dike thus defines the trace of a vent fissure. Only rarely are the surface expressions of vents, such as cones and spatter ramparts, preserved (Bingham, 1970; Swanson and others, 1975b).

Dikes and vents for flows of each stratigraphically bound geochemical unit define one or more narrow linear vent systems (Swanson and others, 1975b), as long as 150 km, many of which are composed of echelon segments. Most or all vent systems trend NNW. (azimuth 310°–350°) and are located in central and eastern Washington, north-eastern Oregon, and western Idaho (fig. 3), with most known dikes occurring in extreme southeastern Washington and northeastern Oregon (Taubeneck, 1970, 1978). Each linear vent system is near the eastern margin of the flows it fed, consistent with other indications that a gentle westward slope was formed and persisted during the time of eruption of the Yakima Basalt Subgroup. Maps showing the location of linear vent systems and the known extent of flows they fed are given in Swanson and others (1979a, plate 1) and in Anderson and others (1987). Locations and chemical analyses of individual dikes are given by Wright and others (1979, 1980). Locations of additional dikes and vents in Idaho are given by Camp (1981) and Swanson and others (1981).

Within the overall dike system (fig. 3) no regular progression of vent location through time has been recognized for the subgroup. Linear vent systems for the Grande Ronde Basalt occur throughout the entire area of the dike swarm. Linear vent systems for each member in the Wanapum Basalt occupy distinct regions, and there was an eastward migration of vents for the upper members of the

Wanapum: those for the Frenchman Springs Member are near the western margin of the dike swarm, those for the Roza are in eastern Washington, and those for the Priest Rapids are in Idaho within the eastern part of the dike swarm. Sources for the Saddle Mountains Basalt are scattered across the entire area, from central Washington (Ice Harbor Member) through eastern Washington and Oregon (Elephant Mountain and Umatilla Members) to Idaho (Weissenfels Ridge Member), and show no systematic variation of vent location with time.

The following members of the Saddle Mountains Basalt have no identified source vents or dikes: Lower Monumental, Pomona, Esquatzel, Weissenfels Ridge (basalt of Slippery Creek), Asotin, and Wilbur Creek Members. Until recently, the basalt of Weippe (Camp, 1981) and its feeder dike were considered correlative with the Pomona Member. However, the paleomagnetic and isotopic signatures of the Weippe dike have been shown to differ from those of the mapped Pomona Member, although the major-oxide and trace-element chemistry are identical (P.R. Hooper, oral commun., 1986). In addition, no vent is known for the Lookingglass flow in the Eckler Mountain Member of the Wanapum Basalt. Four of these units traveled to or across the central area of the Columbia Plateau, and three of these (Pomona, Asotin, and Wilbur Creek Members) each have volumes that exceed 40 km³. We made a careful search along the eastern margin of the outcrop area, after completion of the regional mapping, for dikes or vents that could have fed these units, but we found none. Therefore we conclude that the sources for these units are not presently exposed.

CHEMICAL STUDY OF THE YAKIMA BASALT SUBGROUP

Methods of Analysis

Major oxides were analyzed in the laboratories of the U.S. Geological Survey (USGS) in Reston, Virginia, (Rapid Rock) and Menlo Park, California, (X-ray fluorescence) and in the X-ray fluorescence laboratory at Washington State University (WSU) under the direction of P.R. Hooper (see Hooper, 1981). Electron microprobe analyses of natural sideromelane glass were obtained at the Smithsonian Institution in Washington, D.C. Trace-element determinations have been made by instrumental neutron activation analysis (INAA) for Ba, Co, Cr, Cs, Hf, Rb, Th, U, Zn, Zr, Sc, and the rare-earth elements La, Ce, Nd, Sm, Eu, Gd, Tb, Yb, and Lu; by X-ray fluorescence (XRF) for Ba, Rb, Sr, Zr, Nb, and Y; and by atomic absorption spectroscopy (AAS) for Cu and Ni. Ba, Rb, and Zr were therefore determined by both XRF and INAA. Analyses and methods used for major oxides and for trace-element data obtained by INAA are given by Wright and others (1979, 1980, 1982).

Analyses of trace elements by XRF were obtained from the USGS laboratory in Reston under the direction of Harry Rose; the analyst was Bob Johnson. Analyses by AAS were done by Jean Kane, also in the USGS laboratories in Reston.

The analytical errors of the various data, though difficult to estimate, are much less than the differences between defined geochemical units and therefore do not affect the discrimination of one such unit from another. Relative values for trace elements analyzed by two methods are in

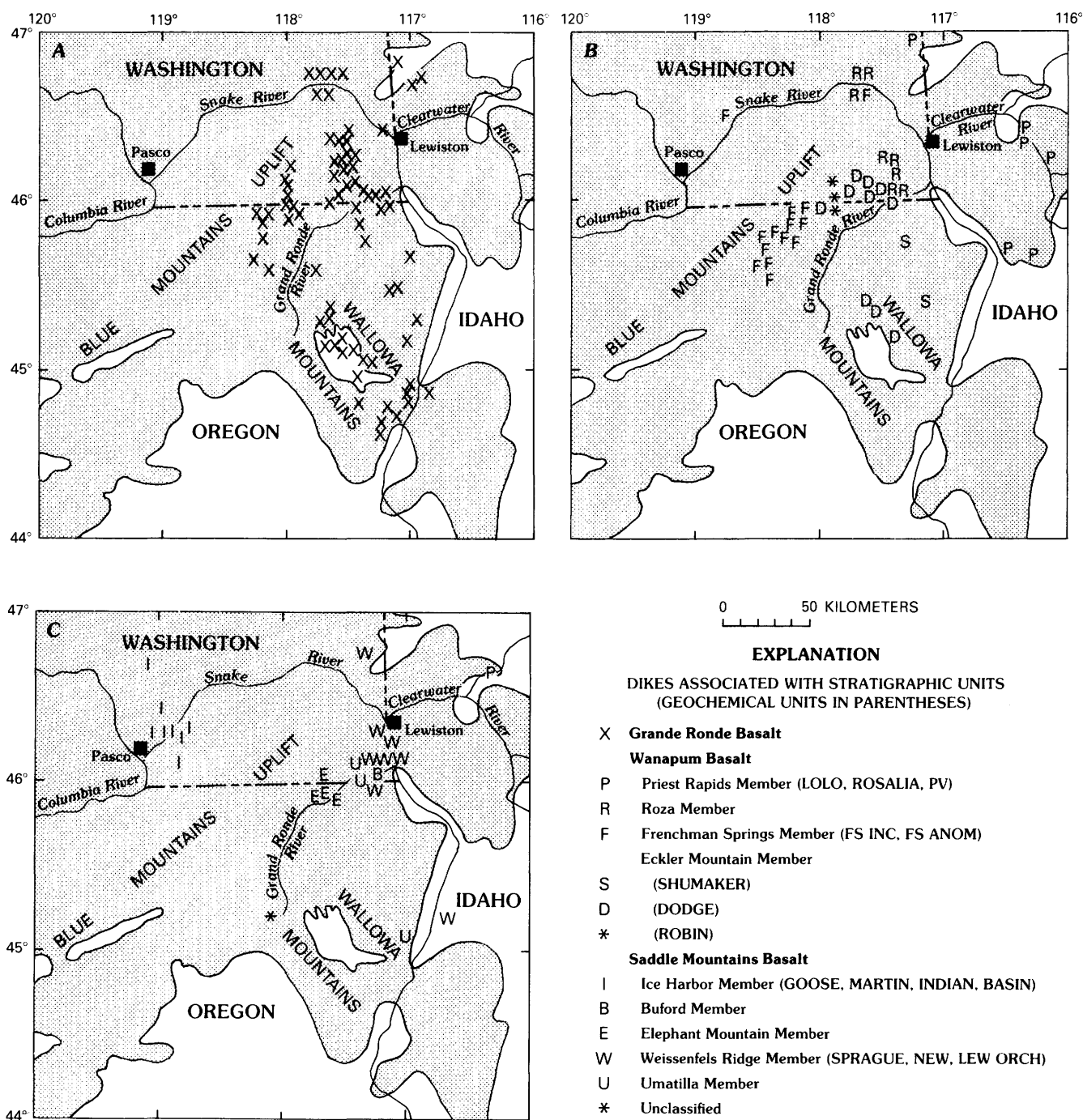


Figure 3. Location of linear vent systems for the Yakima Basalt Subgroup of the Columbia River Basalt Group. Area of Columbia River Basalt Group shaded. See table 1 for stratigraphic setting of units. A, Grande Ronde Basalt. B, Wanapum Basalt. C, Saddle Mountains Basalt.

Table 2. Average analytical differences among three laboratories

[XRF, U.S. Geological Survey, Menlo Park, X-ray fluorescence analyses; WSU, Washington State University, X-ray fluorescence analyses. Biases obtained by subtracting values obtained by reference laboratory, U.S. Geological Survey, Reston, rapid rock analyses. Biases calculated from analyses of selected samples done by at least two different laboratories, and from comparison of flow averages for major stratigraphic units calculated from analyses obtained from each laboratory in course of regional mapping. Biases vary somewhat as a function of bulk chemistry, although signs, + or -, are consistent. Analyses in tables in this paper have not been corrected]

Oxide	Bias (XRF)	Bias (WSU)
SiO ₂	-0.74	+0.33
Al ₂ O ₃	0.00	-0.48
'FeO'	+0.35	-0.30
MgO	+0.01	+0.16
CaO	-0.08	-0.17
K ₂ O	+0.07	0.00
TiO ₂	+0.04	+0.02
P ₂ O ₅	+0.09	+0.05
MnO	n.a.	n.a.

good agreement, although the precision and accuracy are probably greater for the XRF than for the INAA analyses. We have not specifically studied the chemical variation within single flows. An empirical measure of the variability within a geochemical unit is obtained from the variation shown by regional sampling of the same flow or stratigraphic unit, and this variability has been used to scale the plots (see below). Major-oxide analyses made in different laboratories show systematic differences (table 2), and only analyses from the USGS laboratories are used to discuss chemical differences and their interpretation. This is done for consistency and does not imply that these analyses are better than those from other laboratories.

Presentation of Chemical Data

The chemical data on which the discussions and inferences in this paper are based are given in tabular form. Average whole-rock and glass compositions and trace-element ratios of geochemical units and whole-rock compositions of feeder dikes appear together with the main text (tables 3–12); chemical data on some additional analyzed flows (see table 14) are given in the appendix (tables 15–17); and the individual analyses from which the average compositions were derived are published separately as a U.S. Geological Survey Open-File Report (Wright, 1988).

The average major- and trace-element compositions of flows defining each geochemical unit are listed in stratigraphic order in tables 3–5. Each average represents at least two analyses. Data for samples used to compute the averages, including the range and standard deviation for each

element, are summarized for the Grande Ronde Basalt in Mangan and others (1985) and for the Wanapum and Saddle Mountains Basalts in Wright (1988). The compositions of dikes for geochemical units that have well-defined linear vent systems are given in table 6; individual analyses are listed in Wright (1988). The averages calculated from WSU analyses are preceded by a two-letter abbreviation related to the original collectors of the sample: PH, Peter Hooper; VC, Victor Camp; WT, William Taubeneck; RB, Robert Bentley; SW, D.A. Swanson and T.L. Wright. Abbreviations for geochemical units are denoted similarly: SHUMAKER = USGS average, PH SHUM = WSU average for samples of the basalt of Shumaker Creek in table 1. Selected trace-element ratios for all formations are summarized in tables 7–9.

Selected data for all units of the Yakima Basalt Subgroup are also plotted in figures 4–6. Data are plotted by formation, with the Saddle Mountains Basalt divided into two sets of plots, one for flows whose major-oxide chemistry broadly resembles that of the Grande Ronde Basalt (relatively high SiO₂, low total iron as FeO, and low TiO₂), the other for flows whose major-oxide chemistry broadly resembles that of the upper part of the Wanapum Basalt (relatively low SiO₂, high total iron as FeO, and high TiO₂). Letter symbols given in tables 4 and 5 are related to stratigraphic position within the Saddle Mountains and Wanapum Basalts from top (A) toward bottom (Z). In the data for the Grande Ronde Basalt (table 3), symbols A–E refer to major geochemical units repeated throughout the section, 1 and 2 refer to two distinctively magnesian units of local extent, and X represents the remaining Grande Ronde flows (Mangan and others, 1985; 1986). An asterisk (*) in the name for a geochemical unit (for example, 1A*) is used to distinguish averages for units defined in the western plateau; the name used for the same unit in the eastern plateau has no asterisk (for example, 1A). The chemical data are plotted as follows:

MgO variation diagrams (fig. 4).—The chemistry of each unit is summarized on three plots showing, respectively, major oxides and compatible trace elements, incompatible trace elements, and selected trace-element ratios.

Th (Thorium) variation diagrams (fig. 5).—The same data (including MgO) are plotted against a well-determined incompatible element.

Chondrite-normalized rareearth-element (REE) data (fig. 6).—Symbols differ from those used in tables 3–5 because of the plotting program used. The plots in figure 6 show only La, Ce, Sm, Eu, Tb, Yb, and Lu.

Isotope data (fig. 7).—Previously published data for isotopes of εNd, Pb, and O are plotted against ⁸⁷Sr/⁸⁶Sr.

Except for plots of trace-element ratios, figures 4 and 5 are scaled so that the symbol size represents approximately two standard deviations in each oxide or element as estimated from statistical analysis of the geochemical units. This is necessarily somewhat subjective, as the standard

deviation for a given element or oxide varies from one geochemical unit to another. Nonetheless they are sufficiently similar to interpret separation of symbols as a petrologically significant chemical difference. Symbol separation does not have the same meaning on plots of MgO or Th against element ratios, because of difficulty in estimating quantitatively the errors in the ratios.

The chemistry of natural glasses is tabulated separately (tables 10–12). Data from which the glass averages were calculated is given in Wright (1988). The major-oxide compositions of bulk rocks and associated quenched glasses are compared for selected units in figure 8.

The chemical characteristics that distinguish each unit in the Yakima Basalt Subgroup are summarized in table 13.

Petrography

Petrographic data for most units of the Yakima Basalt Subgroup have been summarized by Wright and others (1973, table 3), although the unit names have been changed since (the names 'Upper', 'Middle', and 'Lower' Yakima Basalt of previous usage correspond, respectively, to what are now defined as the Saddle Mountains, Wanapum, and Grande Ronde Basalts). What follows is a brief recapitulation of the previously published data, with petrographic information added for units not identified in 1973.

Grande Ronde Basalt.—Rocks of this unit are mostly aphyric or sparsely microphyric with augite-plagioclase clots. Many flows have rare microphenocrysts of hypersthene. A few local flows are mappable on the basis of visible plagioclase phenocrysts but are not chemically distinctive.

Wanapum Basalt.—Rocks of this unit are commonly sparsely porphyritic with either single (as large as 1 cm) plagioclase phenocrysts (Roza Member; basalt of Dodge of the Eckler Mountain Member), large (1–3 cm) plagioclase clots (many flows of the Frenchman Springs Member; see also Beeson and others, 1985, table 1), or small (1–3 mm) olivine and plagioclase phenocrysts (Priest Rapids Member, LOLO geochemical unit). Some flows in the Frenchman Springs Member (see also Beeson and others, 1985) and a few flows in the basalt of Dodge are aphyric, as are the basalt of Robinette Mountain of the Eckler Mountain Member and other Wanapum flows not mentioned above. The basalt of Dodge is exceptionally coarse-grained (>1 mm) and is identifiable in the field, where it commonly weathers to a coarse grus.

Saddle Mountains Basalt.—Flows of the Pomona Member and the basalt of Martindale (of the Ice Harbor Member) are distinguished by an augite-plagioclase-olivine assemblage of phenocrysts and glomerocrysts. The basalt of Goose Island (of the Ice Harbor Member) is unique for the occurrence of microphenocrysts of ilmenite and titanomagnetite. The Esquatzel Member is distinguished by the presence of augite (as large as 5 mm) as the only phenoc-

ryst type. The rest of the Saddle Mountains Basalt flows are aphyric to sparsely porphyritic, with no phenocrysts large enough to be of use in mapping. Flows of the Asotin Member have a coarse subophitic texture that is distinctive on a weathered surface.

Classification and Naming of Geochemical Units

The geochemical units (tables 1 and 3–6) were classified according to normative data using procedures recommended by Irvine and Baragar (1971) and by Wilkinson (1986). The total iron (FeO of tables 3–6) was partitioned to Fe_2O_3 and FeO in a ratio of 9:1 before calculating the norm. The Irvine and Baragar procedures yield more consistency in the naming of such units than do the Wilkinson procedures. With one exception, geochemical units in the Yakima Basalt Subgroup are classified as subalkaline tholeiitic basalt using their figure 4 (OI'-Ne'-Q'), figure 5 (Cpx-OI-Opx), figure 6 (normative plagioclase An content versus Al_2O_3), and figure 7A (normative plagioclase An content versus color index). The most fractionated chemistry in the Umatilla Member (UM FRAC geochemical unit of table 1) was classified as calc-alkaline tholeiitic andesite using their figures 6 and 7A. The single sample of magnesian Imnaha Basalt (table 15) was also classified as subalkaline tholeiitic basalt on their figures 4, 6, and 7A, although it was barely alkaline using figure 5. Classification of the basalts in terms of sodic or potassic affinity using their figure 8 (An-Ab'-Or) shows that all the samples of the Grande Ronde and Wanapum Basalts, and the single sample of the Imnaha Basalt, have average K_2O contents. Some members of the Saddle Mountains Basalt (Pomona, Esquatzel, Asotin, Wilbur Creek, and Umatilla Members) have composition ranges that extend slightly into the field of K-rich basalts.

The basalts of Powder River (Hooper, 1984), whose compositions are given in table 15, were classified as a mixture of subalkaline and alkaline basalt using figures 4 and 5 of Irvine and Baragar (1971). Alkaline geochemical units include alkalic basalt (WT NEPH), hawaiite (WT SPRMT, WT SUGMT), and trachybasalt (WT WBRMT), using their figure 10 (normative plagioclase An content versus color index). WT JONES and WT ANDES were classified as calc-alkaline andesite using their figures 6 and 7A. Of the Idaho flows mapped by Victor Camp (table 15), only VC POT is classified as an alkaline basalt. The remaining geochemical units of Camp (1981), the PH EDEN unit in northeastern Oregon, and WT OBHTI and WT OBLTI units in the basalts of Powder River, are all classified as subalkaline tholeiitic basalt.

Classification using the procedures of Wilkinson (1986, table 1) resulted in all units containing a mixture of basalt and andesite varying from quartz- to olivine-tholeiitic

affinity depending on MgO content. The most magnesian geochemical units in the Saddle Mountains Basalt (ASOTIN and VC ASOT) and Wanapum Basalt (ROBIN) are classified, because of their high content of Al_2O_3 , as olivine subalkaline basalt. The UM FRAC geochemical unit is classified as a calc-alkaline tholeiitic dacite. The alkaline samples in the basalts of Powder River are classified as basanite (WT NEPH), high-Al transitional andesite (WT SUGMT and VC POT), and alkali andesite (WT SPRMT and WT WBRMT). WT ANDES and WT JONES are classified as calc-alkaline andesite and dacite, respectively. We prefer the simpler naming using the Irvine and Baragar (1971) procedures.

Bulk Rock Chemistry: Observations

Grande Ronde Basalt

The Grande Ronde Basalt (figures 4A–C, 5A–B) shows the most regular chemical variation of the three formations examined. All oxides and elements vary continuously over a limited range of MgO content (3–6 percent). Within this apparent continuum several different chemical subtypes, distinguished by small variations in MgO, TiO_2 , and P_2O_5 , have been defined (Mangan and Wright, 1984; Mangan and others, 1986). The most regular major-oxide variations are the covariance of CaO and MgO and a less strong covariance of Al_2O_3 and MgO. Among the compatible trace elements, Sc shows the most regular variation, a strong positive correlation with CaO. The Sc/CaO ratio rises as MgO decreases. The elements Co, Cr, Ni, Cu, and Sr also behave as compatible elements with decreasing MgO. The variation of Cu suggests the separation of a sulfide melt at or near the liquidus temperature of these basalts (see Skinner and Peck, 1969). One magnesian flow (LH 2 in table 3; symbol “1” in figs. 4A–C and 5A–B) has an anomalous chemistry marked by low TiO_2 , FeO, Co, and Sc and high SiO_2 and Sr. This unit has a well-defined dike system (table 6; Mangan and others, 1986) and covers a limited area in the Snake River-Blue Mountains region. A few other symbols plot off the main trend for a single oxide or element but show no correlated anomalies for other analyzed elements. These represent single analyses and perhaps reflect analytical errors.

Incompatible trace elements and oxides show different degrees of enrichment with decreasing MgO (fig. 4B) or increasing Th (fig. 5A). The elements Cs and Rb behave most incompatibly (greatest enrichment), followed by K_2O and Th. The elements Ba, La, and Hf are less enriched, followed by TiO_2 , P_2O_5 , Ta, Zr, and Nb. The least enrichment is shown by Yb, which remains nearly constant with decreasing MgO.

The spread of data for certain constituents is greater in samples of the Grande Ronde Basalt having MgO less than 3.5 percent. Particularly variable are K_2O and Th, and a group of samples with high values for those elements plots above trends defined by the rest of the Grande Ronde

Basalt samples (fig. 4A). This is expressed on the Th variation diagram (fig. 5A) by a change in slope separating the trend of samples with greater than 6 ppm Th from that defined by the rest of the data. Th continues to increase while the highly incompatible elements like Rb, Cs, and K_2O may remain constant or decrease at higher values of Th.

Trace-element ratios (figs. 4C, 5B) are highly variable, in part owing to the error amplification inherent in calculating ratios. The only ratios that remain fairly constant over a wide range of MgO and Th contents are K/Rb and Hf/Ta. The compatible nature of Sr is strikingly demonstrated by the steep positive slopes of K/Sr and Rb/Sr. The ratio Hf/Th shows a regular decrease with increasing Th or decreasing MgO, emphasizing the compatible behavior of Hf relative to Th.

The Grande Ronde Basalt also shows regular and similar chondrite-normalized rare-earth patterns (fig. 6A). Samples with the lowest MgO contents have the highest REE contents; all patterns are enriched in light rare earths and slightly concave upward. A small negative Eu anomaly, expressed as the ratio of analyzed Eu to Eu measured by linear extrapolation between Sm and Tb, is common. This Eu anomaly ranges from 0.94 to 0.77 and is not correlated with either MgO or the La/Yb ratio. The ratio La/Yb is 4.0–5.5 for chondrite-normalized La contents of 55 to 95 and increases with increasing Th (fig. 5B) or decreasing MgO (fig. 4C).

The data for the Joseph volcanic unit of Kleck (1976) plot within the range of the low-MgO compositions in the Grande Ronde Basalt (table 15; symbol # in figs. 4A–C, 5A–B, 6A). This chemistry and the stratigraphic position of the Joseph volcanic unit indicate that it is the latest outpouring of the Grande Ronde Basalt and formed a line of small shields.

The Sr-isotope data for the Grande Ronde Basalt are tightly clustered near $^{87}\text{Sr}/^{86}\text{Sr} = 0.7050$ (fig. 7A); the total range is 0.7043–0.7055, including samples of McDougall (1976) for which Sr is the only isotopic system analyzed. McDougall (1976, tables 4 and 6) demonstrated a consistent difference within the Grande Ronde Basalt between samples collected in southeastern Washington and adjacent Oregon, within the area of many feeder dikes, and samples from central Washington. We now know that all of the latter samples were from flows that were also erupted from eastern vents (Mangan and others, 1986) but are from the uppermost magnetostratigraphic unit (N_2), whereas the eastern samples are from older units (R_2 and N_1). Hooper (1984, fig. 2) generalized the relations between $^{87}\text{Sr}/^{86}\text{Sr}$ and magnetostratigraphy to draw a smooth trend showing increasing $^{87}\text{Sr}/^{86}\text{Sr}$ with decreasing age, from the Imnaha Basalt through the Grande Ronde Basalt to the top of the Wanapum Basalt. We interpret the same data to indicate a discontinuous jump near the R_2 – N_2 unit boundary, with little Sr-isotopic distinction among the regionally correlated older flows. Within parts of units R_2 and N_2 , $^{87}\text{Sr}/^{86}\text{Sr}$

increases with increasing age. The other isotopes also vary over fairly narrow ranges. Neodymium isotopic data, expressed as the difference from a chondritic value (ϵNd) generally show an inverse correlation with $^{87}\text{Sr}/^{86}\text{Sr}$. On a plot of ϵNd against $^{87}\text{Sr}/^{86}\text{Sr}$ (see fig. 7A; Carlson, 1984a, fig. 3) the samples of the Grande Ronde Basalt define a linear pattern with little scatter.

Wanapum Basalt

The Wanapum Basalt can be divided into two broad chemical groupings, which are also stratigraphically separate. The oldest stratigraphic unit (Eckler Mountain Member) is confined to the southeastern part of the province, where the flows occupy elongate outcrop belts parallel to known linear vent systems (Swanson and others, 1975b, 1979a). Except for P_2O_5 , major oxides of the Eckler Mountain Member plot near or on the extrapolated trends for the Grande Ronde Basalt on variation diagrams using either MgO or Th as a base (compare figs. 4D–F and 5C–D with figs. 4A–C and 5A–B (symbols G, H, I, and J)); one unit, the basalt of Dodge, is identical in major-oxide chemistry to some magnesian flows of the Grande Ronde Basalt. The upper stratigraphic units of the Wanapum Basalt (the Frenchman Springs, Roza, and Priest Rapids Members; symbols A–F) contain the same range of MgO (3–6 percent) as the Grande Ronde Basalt but have distinctly higher FeO, TiO_2 , and P_2O_5 and lower SiO_2 contents.

The Wanapum, like the Grande Ronde, shows a strong covariance of CaO with MgO and Sc and an increasing Sc/CaO ratio with decreasing MgO. Other chemical variations are less regular; for example, TiO_2 and P_2O_5 are poorly correlated.

Trace elements for all members of the Wanapum Basalt plot close to trends defined for the Grande Ronde Basalt, although several incompatible elements (Hf, Nb, Ta, Zr, Y, and the REE) and Ni are systematically enriched in the Wanapum at the same MgO content (compare fig. 4E with fig. 4B). The chondrite-normalized rare-earth patterns (fig. 6B) for the Wanapum have shapes similar to those for the Grande Ronde and also show an overall increase in REE abundance with decreasing MgO; the total REE content is generally higher in all but the DODGE and ROBIN geochemical units. Slightly negative Eu anomalies, as defined above, are present and range from 0.93 to 0.99. The La/Yb ratio is similar in all units, except the DODGE and ROBIN geochemical units, and ranges between 4 and 5 (figs. 4F,

5D). However, the variation of La/Yb ratio with MgO or Th is the opposite of that in the Grande Ronde: the ratio decreases with decreasing MgO or increasing Th in the Wanapum Basalt. The basalt of Robinette Mountain has the flattest REE pattern of any unit in either the Grande Ronde or Wanapum Basalt, with a La/Yb ratio of only 2.4.

Isotopic data for the Wanapum Basalt overlap on all plots with those for the Grande Ronde Basalt (compare fig. 7B with fig. 7A). Both units show a positive correlation of $^{87}\text{Sr}/^{86}\text{Sr}$ with $^{208}\text{Pb}/^{204}\text{Pb}$ and a negative correlation with $^{206}\text{Pb}/^{204}\text{Pb}$ and ϵNd ; $^{207}\text{Pb}/^{204}\text{Pb}$ is constant at about 15.6, and $\delta^{18}\text{O}$, which varies between +5 and +7, does not correlate with variations in other isotopes.

Saddle Mountains Basalt

The Saddle Mountains Basalt (figs. 4G–L, 5E–H) has a much wider range of chemical composition than either the Grande Ronde or Wanapum Basalt, yet it retains a strong positive correlation of CaO with MgO. Many members of the Saddle Mountains Basalt have major-oxide compositions similar to those of older units. We have divided the plots into those geochemical units broadly similar to ones in the Grande Ronde Basalt or the Eckler Mountain Member of the Wanapum Basalt (that is, relatively high SiO_2 coupled with low TiO_2 and total iron) and those geochemical units broadly similar to the upper units of Wanapum Basalt (that is, relatively low SiO_2 coupled with high TiO_2 and total iron). Examples of the former are BUFORD, POMONA, VC WEIPE, ASOTIN, SW HUNTZ, HUNTZ, VC LAP, and WILBUR; examples of the latter are LM, ELEPHANT, LEW ORCH, SPRAGUE, SW SPRAG, NEW, and SW NEW. However, the trace elements in all the geochemical units are quite distinct and serve to distinguish the Saddle Mountains Basalt from older units. Most incompatible elements, with the notable exception of Cs, are highly enriched, and some compatible elements (for example, Sc and Sr) are depleted, in the Saddle Mountains Basalt relative to their levels in older units.

Comparison of the Saddle Mountains Basalt with older units (Wanapum Basalt and Grande Ronde Basalt) is most easily made on an element-by-element basis at the same MgO content (fig. 4). Results of such comparison are shown in the following table (elements marginal between two categories are shown intermediate between two columns):

Saddle Mountains Basalt relative to Grande Ronde Basalt																			
	Co	Cr	Sr	Zn	Sc	Cu	Ni	Ba	Cs	Hf	Nb	Rb	Ta	Th	U	Y	Zr	La	Yb
Enriched	x	—	—	—	—	—	x	x	—	x	x	—	x	x	—	x	x	x	—
Similar	—	x	—	x	—	x	—	x	—	—	—	x	—	—	x	—	—	—	x
Depleted	—	—	x	—	x	—	—	—	x	—	—	—	—	—	—	—	—	—	—
Saddle Mountains Basalt relative to Wanapum Basalt																			
Enriched	x	—	—	—	—	—	—	—	—	x	x	—	x	x	—	—	x	x	x
Similar	—	x	—	—	x	x	x	x	x	—	—	x	—	—	x	x	—	—	x
Depleted	—	—	x	x	x	—	—	—	—	—	—	x	—	—	—	—	—	—	—

The Th variation diagrams (fig. 5E, 5G) for the Saddle Mountains Basalt show different trends for all incompatible elements relative to those for the Grande Ronde or Wanapum Basalt (fig. 5A, 5C). Chondrite-normalized rare earth patterns (fig. 6C) are also steeper for a given MgO content than those of the Grande Ronde or Wanapum, and the La/Yb ratio reaches values greater than 7 (compare fig. 4H, 4K with fig. 4B, 4E). Negative europium anomalies are more pronounced in the Saddle Mountains Basalt than in older units of similar major-oxide chemistry, particularly in flows with high total REE content. Some units of the Saddle Mountains Basalt show unique chemical characteristics, including extreme values for certain elements. Flows of the Ice Harbor Member have higher Sc, Hf, Nb, Ta, Zr, and REE and lower Th and Ni than other units in the Saddle Mountains Basalt. The Ice Harbor Member is richer in REE (fig. 6C) than many other Saddle Mountains Basalt units, but its La/Yb ratios of 5–6 (fig. 4L) are not especially high. The basalt of Goose Island (symbol B in fig. 4J–K) has an extreme composition marked by high total iron and P_2O_5 , far exceeding the range normally found in basalt. This unit also has abnormally high enrichment of Hf, Y, Nb, Ta, and REE. Significantly, Eu anomalies are absent from all geochemical units in the Ice Harbor Member, even the iron-rich GOOSE composition.

The Esquatzel Member has exceptionally high values of Rb and Th, and the Umatilla Member has exceptionally high values of Ba, Rb, Th, and Eu. The Umatilla is the only unit within the Yakima Basalt Subgroup that has a pronounced positive Eu anomaly.

Isotope data for the Saddle Mountains Basalt show that mean data for all isotope systems fall close to the trends defined by the older units extrapolated to higher values of $^{87}Sr/^{86}Sr$. For many units of the Saddle Mountains Basalt, values of $^{87}Sr/^{86}Sr$ cluster around 0.7076. The samples forming this cluster show wider scatter in Pb isotope ratios than the Wanapum or Grande Ronde Basalt. Overall, the Saddle Mountains Basalt has higher values of $^{208}Pb/^{204}Pb$ and $^{207}Pb/^{204}Pb$ than the Grande Ronde and Wanapum Basalts. It also has generally lower values of $^{206}Pb/^{204}Pb$, although the Elephant Mountain, Lower Monumental, and Esquatzel Members have values that equal or exceed those of the Grande Ronde–Wanapum sequence. Values of ϵNd and $\delta^{18}O$ are fairly constant. Data for the following members of the Saddle Mountains Basalt plot at even higher values of $^{87}Sr/^{86}Sr$, with wide and generally uncorrelated scatter in the other isotopes: Lower Monumental, Esquatzel Asotin (Huntzinger flow only), Wilbur Creek, and Umatilla.

Data for additional analyzed flows are given in the appendix (tables 14–17), summarized in a manner similar to that of tables 1, 3–9, and 13.

Comparison of Chemistry of Dikes and Flows

Major-oxide analyses of feeder dikes have been previously published (Wright and others, 1979, 1980, 1982).

Limited trace-element data were obtained for representative dikes of several geochemical units (table 6). Most dikes are uniquely identifiable from both trace elements and major oxides and correlated with specific flows or several flows of similar chemistry. Although one study (Ross, 1983) has shown individual dikes to be chemically inhomogeneous, and another (Taubeneck, 1970) identified inclusions of country rock in many dikes, the data of table 6 show general agreement between analyses of dikes and the average chemistry of flows they fed. Exceptions are (1) the Grande Ronde Basalt, where ambiguity exists for dikes whose chemistry matches that of flows occurring at several different stratigraphic levels and (2) dikes that apparently did not reach the surface to feed a flow (for example, Fairview Bar dike of Ross, 1983; analyses 78-365 and 78-366 in table 2 of Wright and others, 1980). MgO variation diagrams (not shown) indicate no systematic differences in either MgO or other elements between flows and related dikes; all differences in composition are within the range of variation shown by the flows (Mangan and others, 1985; Wright, 1988) and the dikes (Wright, 1988).

Glass Chemistry: Observations

Fresh sideromelane is common as selvages on dikes, on partially palagonitized vent spatter, in water-quenched pillows, and in hyaloclastic debris. Glass is also infrequently preserved on chilled flow bases or unoxidized pahoehoe flow tops. We systematically collected fresh glass and prepared grain mounts for analysis by electron microprobe at the Smithsonian Institution (see Wright and others, 1979, 1980, 1982 for analyses and description of method). Uncorrected analyses are given in Wright (1988), grouped by stratigraphic unit and geochemical unit. For glass-rock pairs, the Na_2O content (and, rarely, K_2O content) of the glass is almost always lower than that of the bulk rock, a relation not possible in normal basaltic crystallization. We ascribe this anomalous relation either to hydration and leaching during the first stage of palagonitization or to loss of sodium during electron bombardment (Rutherford and others, 1985). In rare instances K_2O , Na_2O , or both may have been added to the glass. Helz (1978) did not find a similar discrepancy in her study of the Ice Harbor Member, perhaps because she worked only with polished thin sections and analyzed only the freshest glass.

We adjusted the Na_2O content of glass for each geochemical unit according to preliminary petrologic mixing calculations, which match the bulk rock with phenocrysts plus glass using a representative set of phenocryst compositions (R.T. Helz, unpublished data). In practice we added less than 0.5 percent Na_2O to each glass average to bring the Na_2O in the glass to about 0.1–0.2 percent higher than that of the corresponding rock. The adjusted data, normalized to 100 percent, and the corresponding rock chemistry

are tabulated in tables 10–12 and plotted in figure 8. These glass-rock pairs demonstrate chemical trends expected from phenocryst-related, low-pressure crystallization.

The glass-rock trends of figure 8 differ systematically from the trends defined by the bulk composition of rocks in the same formation, defined by the trends of figure 4. This is particularly evident on plots of MgO against CaO and Al_2O_3 for the Grande Ronde Basalt (fig. 8A). A similar observation can be made for the Wanapum (fig. 8B) and the Saddle Mountains (fig. 8C) Basalts, although the bulk-rock trends are not as closely defined as for the Grande Ronde Basalt.

DISCUSSION: IMPLICATIONS FOR THE ORIGIN OF THE YAKIMA BASALT SUBGROUP

In this section we use information of many kinds to address the fundamental questions of the melting, storage, and mechanics of eruption of the flood basalts of the Yakima Basalt Subgroup. These include the development of a basin to hold the basalt, evolution of a tectonic setting in which rapid eruption of large volumes of basalt could be realized, and the nature of source regions and storage areas consistent with the eruptive history, chemistry, and petrology of the basalts. We draw on information gained from the following: field mapping, stratigraphic study and age dating, geochemical study of the Yakima Basalt Subgroup, and tectonic history of the Pacific Northwest.

Emplacement of Flows

Geologic mapping combined with the chemical data presented in this paper provide insights into the nature of the eruptions that produced the Yakima Basalt Subgroup. Most eruptions were in the eastern part of the province (fig. 3) and fed flows that moved westward down a gentle paleoslope estimated to have been about 1:1,000. Although ongoing subsidence beneath the central Columbia Plateau near Pasco (fig. 1) temporarily reversed the slope in the western part of the province, the largest flows, which have volumes exceeding 500 km^3 , traveled across the Pasco Basin and ponded against the opposed slope of the rising Cascade Range. Smaller eruptions produced flows of local extent that traveled only a few kilometers from their source. Some of these flows (for example, the basalts of Dodge and Robinette Mountain of the Eckler Mountain Member; the Ice Harbor Member) were confined to shallow grabens parallel to their vent systems. Emplacement of the voluminous flows was virtually instantaneous, as evidenced by (1) flat upper flow surfaces, which, together with columnar jointing, indicate ponding of a fluid, and (2) similar crystallinity of dike selvages and basaltic pillows in the same unit separated by 100 km or more (see Shaw and Swanson,

1970; unpub. data of C.A. Hodges for internal dikes in the Roza Member of the Wanapum Basalt—see Hodges, 1978; data of Mangan and others, 1985, 1986, p. 1317 for the Grande Ronde Basalt). Shaw and Swanson (1970) estimated eruption rates as high as $1 \text{ km}^3/\text{d}$ per kilometer of fissure length for the Roza Member. Our mapping and the studies of Swanson and others (1975b) and Mangan and others (1986) are consistent with this figure for the Roza and demonstrate similar relations for large flows of the Grande Ronde Basalt and flows of the Priest Rapids Member of the Wanapum Basalt. We do not know the source dikes for many of the Saddle Mountains units, but we assume from the widespread distribution of the flows that eruption rates were similar to those of the older units.

Eruption rates for the localized flows were probably less than those for the plateau-covering units but still high, as indicated by the absence of thickening of units toward their vents. The viscosities of the flows, calculated from their chemistry (Shaw, 1972), are comparable to or higher than those calculated for basalts of Kilauea, known to form shields or to build up the rift zones in the vicinity of their source vents. Therefore the rates of eruption for all but the least voluminous flows are inferred to be higher for the Yakima Basalt Subgroup, in order to produce ponded flows even at distances of only a few kilometers.

Swanson and others (1975b) emphasized that, although instantaneous eruption rates (discharges) were very high, the average rate at which magma was generated to supply the flows of the Yakima Basalt Subgroup need not have been particularly high. This supply rate for the period of maximum activity (obtained by dividing the total volume of the Grande Ronde and Wanapum Basalts by the length of time during which eruptions occurred) is about $0.075 \text{ km}^3/\text{yr}$, close to the magma supply rate of $0.1 \text{ km}^3/\text{yr}$ currently estimated for Kilauea (Swanson, 1972). If the supply rate is equated with the melting rate in the mantle, then magma that fed the most voluminous individual flows in the Yakima Basalt Subgroup required 5,000–20,000 yr to accumulate before eruption.

Magma Storage Before Eruption

The most voluminous flows in the Yakima Basalt Subgroup have volumes comparable to those of large eruptions of ash-flow tuff, yet no evidence exists for surface deformation in the form of calderas associated with vent areas. Evidence is also lacking for the existence of discrete volcanic centers or of rhyolites, both of which are associated with basaltic provinces known to have crustal magma reservoirs (for example, Iceland and the Snake River Plain). Deformation broadly contemporaneous with volcanism took place in the western part of the Columbia Plateau (compressional folding; Reidel, 1984), the central part (shallow basining), and the eastern part (formation of complex faulted basins; Hooper and Camp, 1981).

Evidence is not convincing, however, that this deformation is directly or specifically connected with large eruptions. Therefore the observations favor magma storage at deep crustal or subcrustal depths.

In general the chemical variations within the Yakima Basalt Subgroup do not fit simple phenocryst-related fractionation in shallow reservoirs (that is, total pressure ≤ 200 MPa, depth ≤ 6 km). Stratigraphically adjacent units do not show a simple chemical relation to one another, in terms of either bulk-rock chemistry or glass-rock trends. Aphyric and porphyritic flows within the same stratigraphic unit (for example, the Frenchman Springs Member or the basalt of Dodge of the Eckler Mountain Member) are chemically identical, a relation that seems to deny the possibility of crystal accumulation. Incompatible-element enrichments vary greatly with either decreasing MgO or increasing Th, inconsistent with low-pressure crystallization of olivine, pyroxene, and plagioclase. In general, Cs, Rb, K_2O , and Th show enrichment of as much as twice that shown by Hf, Zr, La, Ta, and P_2O_5 for a given decrease in MgO (fig. 4). All these observations contrast with those from basalt erupted in ocean-ridge and oceanic-island settings, where abundant evidence exists of shallow crystal fractionation and accumulation superimposed on chemical variation related to deeper processes.

One reviewer of this paper (J. Pallister) pointed out that the chemical data plot near the low-pressure cotectic in the simplified basalt system and suggested that this is evidence for crustal storage. However, this relation does not prove crustal storage; most basalts with fairly low MgO will plot near the low-pressure cotectic in the simplified basalt system, irrespective of their origin.

The chemical variations thus are not consistent with simple crystal fractionation at low pressure. They do not, however, prove the complete absence of crustal storage, because more complex models of crystallization combined with assimilation or replenishment of melt might satisfactorily explain the chemical data. For example, Helz (1978, 1980) has shown that some of the basalt in the Ice Harbor Member experienced intracrustal storage (pressure 350 MPa, depth 10 km). Of the large-volume flows, the Pomona Member is the only one that might have undergone crustal storage, because its phenocryst assemblage is similar to that of the basalt of Martindale of the Ice Harbor Member studied by Helz (1978).

Finally, there is evidence for sulfur saturation of magma in storage. Compatible behavior of sulfophile elements, particularly Cu, implies the existence at near-liquidus temperatures of an immiscible sulfide liquid in much magma of the Yakima Basalt Subgroup (see also Skinner and Peck, 1969). The former existence of such liquid has been confirmed by the identification of spherical blebs of crystalline sulfides found during electron microprobe traverses across dike selvages (R.T. Helz, unpub. data).

Nature of the Source Mantle

Inferences Based on Geochemical Data

The following geochemical observations need to be explained in any satisfactory model of magma generation for the Yakima Basalt Subgroup. In themselves they also provide some limitations on the chemistry and mineralogy of the source mantle.

1. MgO and CaO are correlated with a slope of about 1.2 for all formations in the Yakima Basalt Subgroup. This correlation holds independent of other chemical variations (fig. 4). Helz (1978, p. 211) pointed out that this ratio matches that in clinopyroxene over a considerable range of pressure and is also typical of melt increments derived from partial melting of clinopyroxene-rich sources. That Sc also varies with CaO is consistent with clinopyroxene fractionation. Therefore we propose, following Helz (1978), that the magma of the Yakima Basalt Subgroup equilibrated with clinopyroxene at some stage in its history.
2. Chondrite-normalized REE patterns are enriched in light rare earths and are concave upward. The La/Yb ratio is relatively low (4–5.5) and fairly uniform in the Grande Ronde Basalt, most of the Wanapum Basalt, and many members of the Saddle Mountains Basalt. The shape of the rare-earth patterns and, as with the CaO/MgO ratio, the similarity of all rare-earth patterns independent of other chemical variations are consistent with equilibration with clinopyroxene either as a melting residue or during storage. The absence of high values of the La/Yb ratio and of significant negative europium anomalies disallow the presence of garnet or plagioclase as significant residual or fractionating phases, assuming that the source and storage regions were not subjected to unusually high oxygen fugacities.
3. Each formation boundary involves significant chemical changes that reflect changes in the melting regime.
 - a. Grande Ronde-Wanapum.

The Frenchman Springs, Roza, and Priest Rapids Members represent a major shift to magma with less SiO_2 , more FeO and TiO_2 , and higher content of incompatible elements when compared at the same MgO content. The shift in major-oxide chemistry occurred over a short time interval (within one magnetic polarity epoch). The Eckler Mountain Member is a transition unit; it shows slight trace-element enrichment but retains a chemistry characterized by high SiO_2 and low FeO and TiO_2 , similar to that of the underlying Grande Ronde Basalt.

Isotopic ratios remain the same across the Grande Ronde-Wanapum boundary. Values of $^{87}Sr/^{86}Sr$ (fig. 7) are 0.7043–0.7055 for the Grande

Ronde Basalt and 0.7049–0.7055 for the Wanapum. The aggregate volume of the Wanapum Basalt is about 7 percent that of the Grande Ronde Basalt (Tolan and others, 1987). The length of time during which eruptions occurred is too poorly known, within the limitation of present K-Ar ages, to prove that the supply rate changed across this boundary. The overall magma production rate may have begun to decrease in Wanapum time (see Hooper, 1982, fig. 2), whereas the discharge during individual eruptions of plateau-covering flows remained about the same as in Grande Ronde time.

b. Wanapum-Saddle Mountains.

A major time break separates the last Wanapum and first Saddle Mountains eruptions, a break sufficiently long to allow the cutting of canyons in the older formations. The magma production rate decreased markedly as the time between eruptions increased, and large-volume flows of the Saddle Mountains Basalt were transported westward through deep canyons and shallow valleys cut in older formations.

A major chemical break also coincides with the Wanapum-Saddle Mountains boundary. The $^{87}\text{Sr}/^{86}\text{Sr}$ ratio jumps to over 0.707 for several Saddle Mountains units of otherwise diverse major-oxide and trace-element chemistry and widely separated vent systems. Some units have extremely high $^{87}\text{Sr}/^{86}\text{Sr}$, as high as 0.715. Major-oxide chemistry of many units is similar to that of the Wanapum or Grande Ronde Basalt. Most trace elements are enriched, particularly P, Zr, Hf,

Ta, Y, and Nb; K, Cs, and Rb are not as enriched and may in some cases be depleted relative to earlier geochemical units having similar major-oxide chemistry.

The chemical variations in the Yakima Basalt Subgroup cannot all be explained within a common mantle source. Rather, we postulate that the chemical changes seen at formation boundaries indicate significant changes in the mantle source for the magma.

4. Tectonic classification of geochemical units.

Rocks of the Yakima Basalt Subgroup are inconsistently classified on the many published diagrams that use major-oxide or trace-element chemistry to infer tectonic setting. We classified the rocks using the following sources:

- Pearce and Cann (1973, figs. 3 and 4): Zr-Ti/100-Y; Zr-Ti/100-Sr/2; categories are calc-alkaline basalt (CAB), low-potassium tholeiite (LKT), within-plate basalt (WPB), and ocean-floor basalt (OFB).
- Pearce and others (1977, fig. 1): MgO-FeO*-Al₂O₃; categories are ocean island (OI), ocean-ridge and floor (ORF), continental (C), orogenic (O), and spreading-center island (SCI).
- Wood (1980, fig. 1): Th-Hf/3-Ta; categories are N-type MORB (NOF), E-type MORB including tholeiitic within-plate basalt (EOF-WPT), alkaline within-plate (AWPT), and destructive plate-margin (DPM).

Results of the classification are shown in the following table: (–, units that plot outside of the classification boundaries):

Formation Member	Classification			
	a (fig. 3)	a (fig. 4)	b	c
Grande Ronde Basalt	Mostly WPB, some CAB, OFB	Mostly OFB, CAB	C	All DPM
Wanapum Basalt				
Priest Rapids	WPB	OFB	C	DPM
Roza	WPB	OFB	C	DPM
Frenchman Springs	WPB	OFB	C	DPM
Eckler Mountain	CAB, WPB, OFB	OFB, LKT-CAB	C, ORF(ROBIN)	DPM, WPT (ROBIN)
Saddle Mountains Basalt				
Lower Monumental	WPB	OFB	C	–
Ice Harbor	CAB(GOOSE), WPB	(GOOSE), OFB	OIB	WPT
Buford	WPB-CAB	OFB	C	DPM
Elephant Mountain	WPB	OFB	C	DPM
Pomona	WPB	OFB	OIB-ORF	DPM
Esquatzel	WPB	OFB	C	DPM
Weissenfels Ridge	WPB-OFB	OFB	OIB, C, ORF	WPT
Asotin	WPB-OFB	OFB	OIB-ORF	DPM
Wilbur Creek	CAB	CAB	C	DPM
Umatilla	–	–	SCI	DPM

The most consistent discrimination is that on the basis of major oxides using classification b (Pearce and others, 1977), which identifies most of the basalt units as belonging to a continental setting. The classification of Wood (1980) is apparently the least consistent with the known tectonic setting, for most of the units are classified as basalt associated with a destructive plate margin. This inconsistency was also noted by Prestvik and Goles (1985). These tabulations demonstrate the difficulty in blindly using chemistry to infer tectonic setting. However, the apparent failure of the classifications may be not so much a negation of the method as an indication of the mixed nature of the source regions for the Yakima Basalt Subgroup. The tectonic history of the area presently occupied by the Yakima Basalt Subgroup is complex, and the problems of classification may be related to chemical signatures inherited from a source region composed of continental, oceanic, and subducted components.

Inferences Based on the Tectonic History

The source region for the Yakima Basalt Subgroup may be further delimited by consideration of the following:

1. History of plate interactions at the western edge of the North American continent.

Hamilton and Myers (1966) suggested that the Idaho batholith and the Sierra Nevada batholith were once a continuous coastal plutonic belt, later disrupted by Tertiary extension, that formed a major basin in back of the zone of continued subduction, now represented by the Cascade Range. Consistent with this is the oceanic character of pre-Tertiary rocks exposed beneath the eastern margin of the Columbia Plateau (Vallier, 1977). Additional evidence for the existence of a back-arc basin in Tertiary time is summarized by Eaton (1984 and references cited there). The north-northwest-trending zone of linear vent systems for the Yakima Basalt Subgroup is on the northward extension of the Northern Nevada rift in the Basin and Range province to the south (Zoback and Thompson, 1978). Eaton (1984, p. 292) suggests that at the beginning of plateau volcanism about 17 Ma, shallow crustal faulting in the Basin and Range to the south gave way to wholesale lithospheric rifting to the north.

The fact that most of the presently exposed dikes of the Yakima Basalt Subgroup lie west of the pre-Tertiary continental margin implies that oceanic crust and mantle are likely to have formed a significant part of the source region for the lava. Westward migration of the continental margin provided in addition a continental component of mantle and crust to the sources for the Yakima Basalt Subgroup, particularly for the Saddle Mountains Basalt.

2. The record of paleomagnetic rotations in the Tertiary rocks of the Pacific Northwest.

The paleomagnetic history records block rotations in front of the back-arc basin (Simpson and Cox, 1977; Beck, 1980; Beck and Plumley, 1980; Magill and Cox, 1981; a recent summary is by Fox and Beck, 1985). These may have contributed to the complex distribution of oceanic crust and mantle in the source area for the basalt (see also Heller and Ryberg, 1983). The tectonic history therefore is consistent with the existence of a complexly heterogeneous source region for the Yakima Basalt Subgroup.

3. The present mantle structure beneath the Columbia Plateau.

The mantle structure has been recently reevaluated using a variety of geophysical methods. The depth to the Moho has been determined to be 40 km using seismic refraction methods (Catchings and others, 1984; Catchings and Mooney, 1988), an increase from the earlier value of 30 km obtained by Hill (1972). The higher value agrees with analysis of teleseismic *P*-wave arrivals (Michaelson and Weaver, 1986, fig. 10). Michaelson and Weaver's data also identify an east-dipping high-velocity zone extending well into the mantle at least as far east as the central plateau; they interpret this zone as the subducting Juan de Fuca plate. Catchings and Mooney (1988) consider the deep crustal and upper mantle structure beneath the central Columbia Plateau to be complex and atypical of normal deep continental crust.

We conclude that the complex chemical variation within the Yakima Basalt Subgroup is related to the presence of the following types of material in the source region: depleted oceanic mantle, subducted oceanic crust, original continental mantle, and old continental crust.

SUMMARY AND SYNTHESIS: MAGMA GENERATION, STORAGE, AND ERUPTION OF THE YAKIMA BASALT SUBGROUP

Previous Hypotheses

Discussion of the origin of the Yakima Basalt Subgroup requires an explanation for (1) the extremely high extrusion rates and large flow volumes characterizing the eruptions of flood basalt and (2) the chemical variation and petrographic character of the flows making up the Yakima Basalt Subgroup.

The earliest paper to discuss the distinctive character of flood basalt eruptions was by Shaw and Swanson (1970), who demonstrated the necessity for individual lava flows to be erupted at rates several orders of magnitude higher than those historically observed in active volcanic areas. They did not, however, address the ultimate cause of such high

eruption rates. Swanson and others (1975b) showed that the magma supply rate during Grande Ronde Basalt time (the time of maximum activity) was actually not excessive compared to that of other basaltic provinces characterized by lower discharge rates for individual eruptions. This finding indicates that the distinctive origin of flood basalts is related to the mechanics of storage and extrusion rather than to unusual conditions accompanying melting. Cox (1980) provided a general model for the origin of flood basalts that involved emplacement of thick sills at the crust-mantle boundary and occasional release of magma to the surface through deep crustal fractures; in this model the magma is erupted at high rates because of the high differential pressure that results when the lower lithosphere is fractured.

The earliest statement regarding the petrogenesis of the basalts was made by Waters (1961, p. 583), who concluded that the different units of the then-named Columbia River Basalt were "products from separate magmatic hearths, and not differentiates of a hypothetical uniform magma." Many years later, McDougall (1976) postulated that the basalt was produced by partial melting of the upper mantle following diapiric upwelling of a peridotite having low to moderate water content. He further postulated that the basalt underwent extensive olivine fractionation and some crustal contamination at low pressures before eruption in an extensional environment analogous to that of a back-arc basin. Thompson (1977; see also Thompson and others, 1983) also postulated formation of the basalt in a back-arc basin and pointed out that subducted oceanic crust could not be ruled out in considering the petrogenesis. Helz (1978) was the first worker to develop a comprehensive model for the origin of the Ice Harbor Member (of the Saddle Mountains Basalt). She postulated that the four types of basalt in the member represented separate partial melting events in a mantle source dominated by clinopyroxene. The immediate source regions for two of the basalt types (basalt of Basin City and basalt of Goose Island) were at pressures greater than 1 GPa, and the mechanism of magma release was considered to be deep continental rifting. The other two units (basalt of Martindale and basalt of Indian Memorial) underwent brief storage at crustal levels (pressures of 300–400 MPa) before erupting to the surface. This model was extended to discuss the origin of the entire Yakima Basalt Subgroup (Helz and Wright, 1982, 1987), particularly regarding the need for a pyroxene-rich source. A metasomatic event was postulated to have preceded the first eruption of the Saddle Mountains Basalt. Helz (1978, p. 215) also made a detailed criticism of the model of Thompson (1977) and showed that many of Thompson's assumptions did not fit the field evidence or chemical data.

Cox (1980), in his general model for flood-basalt genesis, postulated a peridotite source, a picritic primary magma, and extensive fractionation of olivine, augite, and plagioclase to produce the observed flood-basalt sequences

around the world. He did not specifically infer the origin of the Columbia River Basalt, but his model agrees with the earlier specific model of McDougall (1976).

Duncan (1982) proposed that eruptions of the Columbia River Basalt Group were triggered by a hot spot that is presently located beneath Yellowstone and which he inferred to have previously formed an oceanic volcanic lineament in early Tertiary time. Duncan related late Eocene and Oligocene volcanism in the Coast Range, Miocene volcanism on the Columbia Plateau, and the basalt of the Snake River Plain and Yellowstone to westward migration of the North American plate over this hot spot.

Reidel (1983) accepted the idea that a pyroxene-rich mantle produced melts of the Grande Ronde Basalt that were stored and partly crystallized in deep crustal storage areas. He explained the chemical character of the Grande Ronde Basalt by discontinuous mixing of freshly melted magma with previously stored magma undergoing fractionation of plagioclase, augite, and olivine.

Hooper (1984) postulated large (>700 km³) storage reservoirs near the crust-mantle boundary as a source for the more voluminous units of the Columbia River Basalt Group. He accepted Cox's (1980) idea that primary magma was melted from a peridotite source and that subsequent fractionation of olivine, augite, and plagioclase yielded the fundamental chemical character of the basalt. Hooper also recognized the need for crustal assimilation in an open replenished magma system and for a metasomatic component to explain some of the chemical variation.

Carlson and others (1981) and Carlson (1984a, b) have provided a comprehensive model for the origin of the Columbia River Basalt Group through analyses of four isotopic series: Sr, Nd, Pb, and O. Carlson's work builds on earlier studies by McDougall (1976), DePaolo and Wasserburg (1979), Nelson (1983), and Church (1985). Carlson's model combines initial mantle heterogeneity with fractionation and crustal contamination to explain the isotopic data. He interprets the chemistry of the Yakima Basalt Subgroup in terms of interaction of at least four end members: (C1) mantle depleted in incompatible-elements, (C2) mantle contaminated by subducted oceanic sediment, (C3) metasomatically enriched subcontinental lithosphere, and (C4) upper crust.

Isotopic values for these end members are tabulated below (Carlson, 1984a, table 4) and plotted in figure 7 using symbols 1, 2, 3, and 4, respectively.

	C1	C2	C3	C4
⁸⁷ Sr/ ⁸⁶ Sr	0.7035	0.7038	0.7076	0.715
εNd	+6.5	+4.5	-6.0	-30.0
²⁰⁶ Pb/ ²⁰⁴ Pb	18.77	19.09	18.66	17.0
²⁰⁷ Pb/ ²⁰⁴ Pb	15.51	15.65	15.65	15.52
²⁰⁸ Pb/ ²⁰⁴ Pb	38.28	28.73	39.44	38.5
δ ¹⁸ O	+5.6	+6.1	+5.8	+14.0

We consider Carlson's model to provide a good framework within which to examine the Yakima Basalt Subgroup, but we recognize the difficulties in reconciling quantitatively the isotopic data with the major-oxide and trace-element chemistry.

Synthesis

We favor a model emphasizing the following points:

1. Following Cox (1980) we postulate storage of Yakima Basalt Subgroup melt in large sill-like reservoirs at the base of the crust (about 40 km deep and at 1.2 GPa pressure). If these storage reservoirs were located directly beneath the longest linear vent systems, and if their volumes were made equal to that of a representative large-volume flow, minimum dimensions of the stored magma body would be 150 by 15 by 0.5 km. The 40-km depth would provide the energy for eruption to the surface when the crust was disrupted by deep continental rifting; draining of such deep sills during flood-basalt eruptions would not be likely to produce significant surface deformation. At the estimated magma supply rate of 0.075 km³/yr (Swanson and others, 1975b), such reservoirs would take 10,000–20,000 yr to accumulate magma for the largest eruptions. Presumably, smaller eruptions would tap reservoirs that were at an earlier stage of development after only a few hundreds or thousands of years of magma accumulation. These estimates are minimum values because we do not know what percentage of magma in storage actually was erupted.
2. Little evidence exists for fractionation of olivine, augite, and plagioclase, as postulated in the Cox (1980) and Hooper (1984) models. Helz's (1978) experimental results on the Ice Harbor Member show olivine to be barely on the liquidus at high pressure for compositions more magnesian and less silicic than most of the large-volume flows. The basalt flows are characterized by small negative europium anomalies that are not, however, correlated with MgO content, degree of incompatible-element enrichment, total rare-earth content, or other possible indicators of fractionation or degree of melting. Thus we do not interpret the presence of the europium anomalies as caused by extensive plagioclase fractionation. The shape of rare-earth patterns, their near parallelism in all units of the Yakima Basalt Subgroup, and the CaO-MgO coherence are also consistent with the evidence against extensive plagioclase fractionation and argue for melt in equilibrium with clinopyroxene and probably derived from a pyroxene-rich source. Possibly, however, picritic melt was produced initially, and ultramafic cumulate contributed to the process of underplating the lithosphere (see below) after ponding of less mafic magma at the base of the lithosphere. This hypothesis

requires a significantly higher magma production rate than that estimated from the eruption rates given above.

Our inference that the magma was primary melt from a pyroxene-rich mantle is made on the basis of the extremely large volume of olivine-rich primary magma that would be required to produce such large volumes of uniform magma, as well as on the absence of any olivine-controlled (picritic) flows in the Yakima Basalt Subgroup. This inference has received support from high-pressure experiments conducted by E. Takahashi, who postulates (Takahashi, 1988) that the chemical trends in the Grande Ronde Basalt are consistent with a high and varying degree of partial melting from a source chemically similar to midocean-ridge basalt at a pressure of 2.25 GPa corresponding to a depth of 70 km. These data are consistent with the model presented by Helz and Wright (1987).

3. Continental crustal underplating (Lachenbruch and Sass, 1978; Furlong and Fountain, 1986) is attractive as a mechanism to explain both the seismic velocity structure and the presence of large bodies of melt stored at the base of the lithosphere. Whether magma is released as flood basalt may depend critically on the tectonics of shallower regions. Lachenbruch and Sass (1978) show that rapid extension, as in the Basin and Range province in Nevada, results in emplacement of large quantities of basaltic magma within the lithosphere; this magma serves as a heat source for crustal melting and production of large-volume silicic melts. Slower extension, as across the Columbia Plateau, may allow deep crustal rifting to directly tap the basaltic magma emplaced at the base of the lithosphere, with little crustal interaction during transport to the surface.

The chemical and isotopic variations within each formation of the Yakima Basalt Subgroup are a function of small-scale source heterogeneity, as well as of differences in rate and degree of partial melting and interaction with continental crust; the latter processes are important only for certain units. The isotopic evidence for source heterogeneity and crustal interaction are neatly summarized by Carlson and others (1981) and Carlson (1984a). Major and abrupt changes in chemical composition at the top of the Eckler Mountain Member and the top of the Wanapum Basalt must, however, be ascribed to fundamental changes in the source composition, most likely owing to some form of metasomatism, as first suggested by Helz and Wright (1982) and amplified later (Helz and Wright, 1987).

Some units underwent a brief period of crustal storage, as for example the basalt of Martindale of the Ice Harbor Member (Helz, 1978, 1980). Of the large-volume units, the Pomona Member is the only one whose petrography indicates a period of crustal stor-

age. Other members of the Saddle Mountains Basalt, such as the Lower Monumental (source unknown but at least as far east as western Idaho), the Esquatzel (source unknown), and the Umatilla (source in southeasternmost Washington), must have had specific interaction, either melt-rock or rock-rock, with (Precambrian?) continental crust in order to explain their anomalous isotopic signatures.

Once eruption was initiated, most flows must have developed their present phenocryst assemblage near the surface or after ponding. Possibly some units were superheated on eruption, depending on the phase relations and on the rate of heat loss during transport to the surface. The presence of phenocrysts in dikes that fed porphyritic flows implies formation of those phenocrysts before eruption. However, even scattered large plagioclase phenocrysts show no evidence of either concentration or flow orientation, so we infer that they could have formed in a turbulent environment just before eruption or after ponding.

Some flows have an unusual posteruption history. Hooper (1985) and Reidel and Fecht (1987) make compelling cases for the mixing of two independently generated magmas before and during eruption to the surface or after ponding or both. We have also determined that at least one thick flow of the Grande Ronde Basalt shows evidence of contemporaneous eruption and mixing in a single cooling unit of lava of mixed chemistry. The flows of the Roza Member show peculiar structures similar to ring dikes (Hodges, 1978). We have determined that the glassy selvages on these dikes have the same chemical composition as that of both the fresh sideromelane at Roza vents and the unpalagonitized pillow selvages where the Roza went into water far from its source. Hodges infers that these dikes originated from within the basalt, and we infer that they may have formed over areas where magma was being fed directly into an already ponded flow.

The structural evolution of the plateau was marked throughout the flood-basalt eruptions by uplift of the Rocky Mountains to the east and the Cascades to the west and subsidence of the area between. Eruptions of the Yakima Basalt Subgroup may have indirectly accelerated basining and other structural adjustments in the Columbia Plateau. The Grande Ronde and Wanapum flows having the largest volumes temporarily masked the deformation and formed a fairly flat surface of ponded basalt. The ancestral Columbia River system was antecedent to the Cascades and carried many of these flows beyond the Plateau to or near the Pacific Ocean (Tolan and others, 1984). The longer time intervals between eruptions of the Saddle Mountains Basalt allowed a more mature drainage to develop. By the beginning of Saddle Mountains time,

the ancestral Snake-Clearwater system and most of its major tributaries had formed on the Plateau (Swanson and others, 1975a; Fecht and others, 1986; Waitt and Swanson, 1987). Flows of the Saddle Mountains Basalt were channeled, and some moved great distances, ponding against the rising Cascades. The Pomona Member is the only one that traversed both the ancestral Snake-Clearwater and Columbia River drainages, reaching the coast and pouring into the Pacific Ocean (Tolan and others, 1984). The eruptions came to a gradual end; the last three events, of decreasing size, are dated at 10.5 Ma (Elephant Mountain Member), 8.5 Ma (Ice Harbor Member), and 6.0 Ma (Lower Monumental Member). Subsidence continued in the Pasco Basin through at least the Pliocene, until isostatic equilibrium was reestablished.

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TABLES 3–17; FIGURES 4–9

Table 3. Average chemical composition of geochemical units in the Grande Ronde Basalt

[Geochemical units listed in approximate stratigraphic order, see table 1; chemical types A-E are major types repeated throughout section, 1 and 2 are distinctively magnesian units of local extent, and X refers to all other Grande Ronde flows. Trace elements followed by X are analyzed by X-ray fluorescence; all others analyzed by instrumental neutron activation. n.a., not analyzed. Data from which these average compositions were calculated are given in Mangan and others (1985)]

Geochemical unit----- chemical type-	1A A	1A* A	3A C	3A C	3A C	3A C	UNC X	4A D	4A D	4A* D	3B C	LH2 1	MG6 2
Major oxides (weight percent)													
SiO ₂	54.58	54.53	53.53	53.65	54.09	53.71	54.52	54.52	53.83	53.99	53.66	53.80	51.52
Al ₂ O ₃	14.51	14.32	14.52	14.30	14.42	14.34	14.51	14.30	13.89	14.14	14.71	15.19	14.98
FeO	10.93	11.08	11.34	11.31	11.17	11.34	10.20	12.10	11.62	11.72	11.21	9.40	11.83
MgO	4.53	4.83	5.19	5.23	4.95	5.24	5.14	4.52	4.83	4.84	5.13	6.12	5.96
CaO	8.53	8.43	9.09	9.01	8.80	8.95	8.95	7.81	8.84	8.53	8.83	10.09	9.71
Na ₂ O	2.84	3.04	2.81	2.84	2.81	2.91	2.89	2.83	2.88	3.00	2.77	2.88	2.76
K ₂ O	1.42	1.32	1.02	1.04	1.18	1.10	1.13	0.97	1.32	1.22	1.31	0.69	0.60
TiO ₂	1.80	1.73	1.74	1.83	1.81	1.75	1.91	2.05	1.97	1.86	1.72	1.13	1.70
P ₂ O ₅	0.36	0.28	0.26	0.29	0.32	0.23	0.32	0.33	0.32	0.27	0.27	0.26	0.32
MnO	0.18	0.18	0.21	0.18	0.19	0.19	0.17	0.19	0.21	0.19	0.20	0.16	0.18
Total	99.68	99.74	99.71	99.68	99.74	99.76	99.74	99.62	99.71	99.76	99.81	99.72	99.56
Trace elements (parts per million)													
Ba	579	502	464	458	537	472	507	450	494	439	518	367	407
BaX	577	537	465	469	491	454	487	433	512	439	500	n.a.	0
Co	37.30	33.60	38.80	38.50	40.0	38.60	40.80	39.80	38.60	38.10	39.50	34	40.10
Cr	43.40	39.10	52.40	49.90	60	54.90	51	31	25.30	23.80	48.90	84.50	99.80
Cs	0.90	0.85	0.70	0.50	1.10	0.65	0.92	0.39	0.80	0.50	0.70	n.a.	0.43
Hf	4.14	3.70	3.70	4.05	3.97	3.85	4.03	3.81	4.10	3.90	3.80	2.77	3.20
NbX	11	13	11	13	16	11.50	12	12	15	14	12	n.a.	0
Rb	30.60	29.50	26.50	29	30	30.50	35.50	27	33	27	35.20	17.20	26.50
RbX	32.70	34.50	28	23	26	29.50	32	24	31	29	31	26	0
SrX	334	325	333	315	363	335	322	321	337	338	328	486	0
Ta	0.75	0.72	0.70	0.67	0.76	0.73	0.74	0.71	0.78	0.72	0.69	0.43	0.62
Th	4.19	3.60	3.20	3.60	3.66	3.40	3.53	3.22	3.79	3.60	3.63	1.85	2.38
U	1.10	1.05	0.80	0.88	n.a.	0.75	1.20	n.a.	n.a.	0.80	0.80	n.a.	0.90
YX	33.30	31	31	31	40	32	30	30	33	32	30	23	0
Zn	132	114	113	122	138	118	127	120	130	112	142	122	123
Zr	163	160	115	207	n.a.	135	150	213	144	76	200	n.a.	147
ZrX	170	167	158	157	175	158	164	157	173	164	161	119	0
Sc	34.50	31	36.30	37.20	36.70	37.80	37	36.10	36.80	35.20	37.10	36.80	38.30
La	22.10	18.50	18	19.40	20.80	19.50	19.90	18.10	20.40	19	20.10	15.60	16
Ce	43.20	38.50	35.50	38.80	43.10	38	38.80	36.30	39.90	37	40.20	30.90	32.30
Nd	26.60	24	20.50	23	23.70	22.50	24.50	28	27	23	23.50	18.40	21.30
Sm	6	5.30	5.10	5.40	6	5.60	5.60	5.70	5.90	5.20	5.60	4.30	3.80
Eu	1.61	1.57	1.58	1.59	1.81	1.64	1.62	1.42	1.66	1.57	1.69	1.25	1.44
Gd	5.30	5.30	6.20	6.30	n.a.	4.60	6.10	5.40	6.10	4.80	5.90	n.a.	4.80
Tb	0.96	0.84	0.84	0.95	1.10	0.90	0.98	0.90	0.85	0.82	1	0.77	0.78
Yb	3.13	2.90	3.15	3.30	3.12	3.25	3.17	3.03	3.20	3.10	3.12	1.93	2.90
Lu	0.52	0.43	0.47	0.47	0.56	0.48	0.49	0.51	0.52	0.46	0.51	0.41	0.41
Cu	31.30	29.50	36	34	n.a.	34	36	36	28	28	33	n.a.	0
Ni	15.70	10.50	18	14	n.a.	15.50	16	17	16	8	14	n.a.	0
Normalized major-oxide contents (weight percent)													
SiO ₂	54.76	54.67	53.69	53.82	54.23	53.84	54.66	54.73	53.99	54.12	53.76	53.95	51.75
Al ₂ O ₃	14.56	14.36	14.56	14.35	14.46	14.37	14.55	14.35	13.93	14.17	14.74	15.23	15.05
FeO	10.97	11.11	11.37	11.35	11.20	11.37	10.23	12.15	11.65	11.75	11.23	9.43	11.88
MgO	4.54	4.84	5.21	5.25	4.96	5.25	5.15	4.54	4.84	4.85	5.14	6.14	5.99
CaO	8.56	8.45	9.12	9.04	8.82	8.97	8.97	7.84	8.87	8.55	8.85	10.12	9.75
Na ₂ O	2.85	3.05	2.82	2.85	2.82	2.92	2.90	2.84	2.89	3.01	2.78	2.89	2.77
K ₂ O	1.42	1.32	1.02	1.04	1.18	1.10	1.13	0.97	1.32	1.22	1.31	0.69	0.60
TiO ₂	1.81	1.73	1.75	1.84	1.81	1.75	1.91	2.06	1.98	1.86	1.72	1.13	1.71
P ₂ O ₅	0.36	0.28	0.26	0.29	0.32	0.23	0.32	0.33	0.32	0.27	0.27	0.26	0.32
MnO	0.18	0.18	0.21	0.18	0.19	0.19	0.17	0.19	0.21	0.19	0.20	0.16	0.18
Total	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00

Table 3. Average chemical composition of geochemical units in the Grande Ronde Basalt—Continued

Geochemical unit----- Chemical type - 2	2B	5A B	5A E	5A E	5A E	5A* E	2C E	2C B	5B B	3C E	3C C	3C* C	C
Major oxides (weight percent)													
SiO ₂	52.44	56.69	55.59	56.04	55.99	55.94	56.46	54.80	55.42	56.24	54.35	53.29	53.75
Al ₂ O ₃	14.67	13.56	13.77	14.06	13.91	13.46	13.98	13.35	13.80	14.13	14.33	14.56	14.54
FeO	11.45	11.01	11.95	11.16	11.62	11.63	11.17	12.90	12.13	10.92	10.87	10.89	11.27
MgO	5.51	3.43	3.74	3.67	3.72	3.70	3.62	3.61	3.41	3.73	4.62	5.44	5.12
CaO	9.76	7.07	7.20	7.24	6.95	7.31	7.09	6.93	7.01	7.15	9.04	9.70	8.68
Na ₂ O	2.92	3.00	2.91	3.02	3.23	2.90	3.31	3.36	3.15	3.02	2.94	2.82	3.08
K ₂ O	0.75	2.16	1.81	2.03	1.70	1.94	1.75	1.69	1.99	2.20	1.38	1.01	1.13
TiO ₂	1.56	2.28	1.96	1.95	1.97	2.12	1.92	2.38	2.27	1.88	1.76	1.64	1.72
P ₂ O ₅	0.29	0.41	0.34	0.34	0.35	0.31	0.28	0.42	0.38	0.32	0.31	0.24	0.27
MnO	0.18	0.20	0.19	0.19	0.18	0.20	0.18	0.20	0.20	0.18	0.20	0.19	0.18
Total	99.53	99.81	99.46	99.70	99.62	99.51	99.76	99.64	99.76	99.77	99.80	99.78	99.74
Trace elements (parts per million)													
Ba	432	843	733	759	713	777	679	736	746	707	533	413	485
BaX	393	n.a.	689	767	705	782	704	702	n.a.	715	410	414	474
Co	38.4	35.1	37.3	37.3	37.7	35.9	35.2	36.8	36.1	33.7	43.9	39.5	38.8
Cr	110	6.1	6.9	11.6	10.0	8.5	9.4	3.5	n.a.	10.8	56.2	129.6	52.7
Cs	0.89	1.39	1.62	1.37	0.81	1.60	0.94	0.85	1.37	1.33	0.86	0.68	0.66
Hf	3.12	4.90	4.96	4.88	4.90	4.97	4.75	5.00	4.80	4.61	3.83	3.38	3.58
NbX	13	n.a.	14	13	15	11	16	15	n.a.	14	24	12	14.3
Rb	19.1	56.4	57.0	53.1	51.8	53.0	51.3	48.7	46.5	56.0	30.7	22.7	28.7
RbX	26.0	n.a.	54.70	49.0	48.0	57.0	53.7	51.3	n.a.	53.0	36.0	25.5	30.0
SrX	381	n.a.	341	339	324	336	333	336	n.a.	319	345	317	354
Ta	0.62	0.64	0.93	0.99	0.93	0.90	0.90	0.95	0.86	0.87	0.79	0.64	0.72
Th	2.26	5.91	6.40	6.15	6.58	6.30	6.27	5.73	5.62	6.10	3.66	3.10	3.16
U	0.69	n.a.	1.59	1.70	1.56	1.80	1.59	1.43	n.a.	1.70	n.a.	0.82	0.79
YX	30.7	n.a.	34.0	34.0	35.0	34.0	35.0	39.3	n.a.	34.0	28.0	29.5	32.0
Zn	125	134	130	138	138	132	136	142	133	131	130	128	125
Zr	179	n.a.	194	144	n.a.	243	189	200	n.a.	214	n.a.	116	165
ZrX	131	n.a.	199	190	190	201	189	196	n.a.	184	160	137	148
Sc	37.5	32.4	31.3	31.8	32.1	31.8	30.7	33.1	32.8	29.9	36.4	37.9	36.0
La	17.3	30.0	28.0	26.9	27.0	26.7	27.6	27.8	28.1	25.8	19.8	16.8	18.7
Ce	34.2	58.5	53.7	53.7	54.2	51.0	52.0	53.8	55.2	48.9	40.2	33.7	37.0
Nd	22.3	36.9	31.8	30.5	28.3	28.0	30.0	31.0	29.9	31.0	23.0	19.0	22.3
Sm	5.1	7.5	6.8	6.7	6.8	6.3	6.5	7.1	7.0	6.3	5.8	4.9	5.5
Eu	1.55	2.20	1.83	1.89	1.99	1.80	1.84	1.98	2.19	1.69	1.76	1.51	1.58
Gd	4.6	n.a.	7.3	6.6	8.4	6.6	6.5	7.1	n.a.	7.9	n.a.	4.5	20.4
Tb	0.88	1.29	1.18	1	1.13	1.18	0.99	1.25	1.07	0.88	0.99	0.88	0.85
Yb	2.69	3.78	3.53	3.09	3.17	3.48	3.41	3.63	3.54	3.31	2.73	2.67	3.10
Lu	0.44	0.64	0.51	0.52	0.54	0.53	0.50	0.55	0.56	0.50	0.51	0.45	0.45
Cu	49.5	n.a.	13.8	15.0	13.0	17.0	n.a.	13.0	n.a.	n.a.	n.a.	n.a.	n.a.
Ni	20.0	n.a.	6.4	6.2	6.5	7.0	n.a.	5.2	n.a.	n.a.	n.a.	n.a.	n.a.
Normalized major-oxide contents (weight percent)													
SiO ₂	52.69	56.80	55.89	56.21	56.20	56.22	56.60	55.00	55.55	56.37	54.46	53.41	53.89
Al ₂ O ₃	14.74	13.59	13.84	14.10	13.96	13.53	14.01	13.40	13.83	14.16	14.36	14.59	14.58
FeO	11.50	11.03	12.01	11.19	11.66	11.69	11.20	12.95	12.16	10.95	10.89	10.91	11.30
MgO	5.54	3.44	3.76	3.68	3.73	3.72	3.63	3.62	3.42	3.74	4.63	5.45	5.13
CaO	9.81	7.08	7.24	7.26	6.98	7.35	7.11	6.96	7.03	7.17	9.06	9.72	8.70
Na ₂ O	2.93	3.01	2.93	3.03	3.24	2.91	3.32	3.37	3.16	3.03	2.95	2.83	3.09
K ₂ O	0.75	2.16	1.82	2.04	1.71	1.95	1.75	1.70	1.99	2.21	1.38	1.01	1.13
TiO ₂	1.57	2.28	1.97	1.96	1.98	2.13	1.92	2.39	2.28	1.88	1.76	1.64	1.72
P ₂ O ₅	0.29	0.41	0.34	0.34	0.35	0.31	0.28	0.42	0.38	0.32	0.31	0.24	0.27
MnO	0.18	0.20	0.19	0.19	0.18	0.20	0.18	0.20	0.20	0.18	0.20	0.19	0.18
Total	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00

Table 3. Average chemical composition of geochemical units in the Grande Ronde Basalt—Continued

Geochemical unit-----	3D	5D	5D	UNC	2D	2D	2D	2D	2D	2D	2D	2D	2D*
Chemical type - C	E	E	X	B	B	B	B	B	B	B	B	B	B
Major oxides (weight percent)													
SiO ₂	53.43	54.90	55.17	54.97	54.45	54.65	54.22	54.71	56.07	54.01	55.44	55.56	55.39
Al ₂ O ₃	14.60	14.04	14.01	13.46	13.93	14.18	14.12	14.09	13.61	13.93	13.85	13.75	13.95
FeO	11.22	11.98	11.52	12.33	12.43	12.04	12.81	12.03	11.66	12.68	11.95	11.82	11.92
MgO	5.25	4.16	4.37	3.81	3.89	3.79	3.66	3.55	3.23	3.80	3.46	3.50	3.54
CaO	9.10	7.79	7.74	7.45	7.60	7.48	7.03	7.29	6.78	7.24	6.97	7.09	7.09
Na ₂ O	2.89	2.82	3.09	3.03	3.10	3.18	3.27	2.95	2.99	3.16	3.11	3.19	3.15
K ₂ O	1.11	1.66	1.54	1.73	1.65	1.73	1.68	2.10	2.39	1.84	2.01	2.01	1.87
TiO ₂	1.72	1.96	1.87	2.30	2.13	2.12	2.25	2.43	2.31	2.46	2.33	2.22	2.31
P ₂ O ₅	0.26	0.29	0.28	0.50	0.39	0.37	0.43	0.44	0.52	0.44	0.46	0.43	0.37
MnO	0.19	0.18	0.19	0.20	0.21	0.19	0.20	0.20	0.22	0.20	0.21	0.20	0.18
Total	99.77	99.78	99.78	99.78	99.78	99.73	99.67	99.79	99.78	99.76	99.79	99.77	99.77
Trace elements (parts per million)													
Ba	495	644	632	727	692	630	693	701	824	714	802	730	685
BaX	485	n.a.	n.a.	n.a.	645	649	683	n.a.	827	706	619	715	746
Co	41.1	41.5	41.9	41.1	43.4	38.0	38.6	40.2	38.4	38.7	37.1	34.6	35.9
Cr	100.0	19.1	22.3	9.2	14.0	8.3	10.3	13.5	9.4	14.5	11.7	12.5	7.9
Cs	0.71	1.29	0.82	1.08	1.12	0.65	1.01	1.28	1.54	1.14	1.50	1.32	1.38
Hf	3.72	4.30	4.17	4.79	4.82	4.48	4.67	5.14	4.91	4.99	5.27	4.74	4.88
NbX	13.3	n.a.	n.a.	n.a.	16.0	13.0	10.0	n.a.	12.0	12.0	14.0	13.0	14.0
Rb	27.7	41.0	41.3	39.5	37.5	50.0	36.4	57.5	69.2	46.3	48.6	45.4	55.0
RbX	36.3	n.a.	n.a.	n.a.	48.0	41.0	29.0	n.a.	66.0	46.0	54.0	51.0	52.0
SrX	308	n.a.	n.a.	n.a.	313	312	322	n.a.	321	312	340	359	324
Ta	0.74	0.9	0.95	0.99	0.85	0.91	0.95	1.02	0.91	0.88	0.95	0.90	0.92
Th	3.46	4.82	4.99	5.33	4.58	5.0	4.96	5.33	7.34	5.16	5.96	6.38	6.50
U	1.02	n.a.	n.a.	n.a.	n.a.	1.27	1.44	n.a.	n.a.	1.18	n.a.	1.78	1.63
YX	33.3	n.a.	n.a.	n.a.	40.0	32.0	32.0	n.a.	34.0	33.0	33.0	37.0	35.0
Zn	131	141	128	150	151	149	139	147	137	147	147	140	145
Zr	120	n.a.	n.a.	n.a.	n.a.	159	151	n.a.	n.a.	198	n.a.	178	157
ZrX	164	n.a.	n.a.	n.a.	194	176	183	n.a.	198	194	231	183	191
Sc	37.1	34.2	33.0	33.5	35.8	32.6	32.6	33.4	29.3	32.5	31.0	31.0	30.5
La	18.3	22.9	21.6	26.2	25.1	24.0	25.9	27.6	28.9	26.0	28.7	27.6	27.9
Ce	37.5	44.8	42.6	54.0	51.3	46.2	53.1	58.7	57.5	54.6	58.5	54.2	56.0
Nd	20.5	26.2	21.8	29.8	30.3	27.0	28.8	32.7	37.1	31.7	33.5	29.9	28.0
Sm	5.3	6.0	5.9	7.3	7.1	6.5	7.0	7.8	8.0	7.3	7.8	7.3	7.3
Eu	1.66	1.90	1.77	2.13	2.11	1.83	2.05	2.34	2.08	2.02	2.19	2.06	1.92
Gd	5.0	n.a.	n.a.	n.a.	n.a.	6.6	6.6	n.a.	n.a.	7.0	n.a.	7.0	6.2
Tb	0.96	1.17	1.04	1.07	1.42	0.93	1.05	1.35	1.22	1.21	1.2	1.08	1.20
Yb	2.72	2.96	2.79	3.37	3.41	3.54	3.56	3.65	3.40	3.54	3.67	3.13	3.40
Lu	0.50	0.54	0.50	0.63	0.62	0.53	0.58	0.61	0.55	0.59	0.63	0.55	0.52
Cu	41.0	n.a.	n.a.	n.a.	62.0	n.a.	n.a.	n.a.	18.0	n.a.	n.a.	n.a.	n.a.
Ni	16.0	n.a.	n.a.	n.a.	17.0	n.a.	n.a.	n.a.	6.0	n.a.	n.a.	n.a.	n.a.
Normalized major-oxide contents (weight percent)													
SiO ₂	53.55	55.02	55.29	55.09	54.57	54.80	54.40	54.83	56.19	54.14	55.56	55.69	55.52
Al ₂ O ₃	14.63	14.07	14.04	13.49	13.96	14.22	14.17	14.12	13.64	13.96	13.88	13.78	13.98
FeO	11.25	12.01	11.55	12.36	12.46	12.07	12.85	12.06	11.69	12.71	11.98	11.85	11.95
MgO	5.26	4.17	4.38	3.82	3.90	3.80	3.67	3.56	3.24	3.81	3.47	3.51	3.55
CaO	9.12	7.81	7.76	7.47	7.62	7.50	7.05	7.31	6.79	7.26	6.98	7.11	7.11
Na ₂ O	2.90	2.83	3.10	3.04	3.11	3.19	3.28	2.96	3.00	3.17	3.12	3.20	3.16
K ₂ O	1.11	1.66	1.54	1.73	1.65	1.73	1.69	2.10	2.40	1.84	2.01	2.01	1.87
TiO ₂	1.72	1.96	1.87	2.31	2.13	2.13	2.26	2.44	2.32	2.47	2.33	2.23	2.32
P ₂ O ₅	0.26	0.29	0.28	0.50	0.39	0.37	0.43	0.44	0.52	0.44	0.46	0.43	0.37
MnO	0.19	0.18	0.19	0.20	0.21	0.19	0.20	0.20	0.22	0.20	0.21	0.20	0.18
Total	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00

Table 3. Average chemical composition of geochemical units in the Grande Ronde Basalt—*Continued*

Geochemical unit-----	2D*	2D*	2D	2D	UNC	AS16	AS16	AS16	AS16	KB5	KB5	3E	3E
Chemical type - B	B	B	B	B	X	4	4	4	4	3	3	C	C
Major oxides (weight percent)													
SiO ₂	55.25	55.08	55.81	56.27	56.04	54.15	54.09	53.83	54.56	53.66	53.56	54.67	53.96
Al ₂ O ₃	13.93	13.89	14.20	14.12	14.10	13.70	14.27	14.08	13.71	13.77	14.17	14.13	14.42
FeO	12.02	12.24	11.76	11.43	10.26	12.26	11.95	12.44	12.03	12.86	12.64	11.26	11.17
MgO	3.60	3.50	3.40	3.50	3.76	4.59	4.48	4.40	4.49	4.13	4.16	5.05	5.16
CaO	7.11	6.99	6.95	6.44	7.59	8.19	8.26	7.99	8.01	7.82	7.79	8.14	8.79
Na ₂ O	3.28	3.39	3.01	3.32	3.07	2.96	2.90	3.09	3.19	3.15	3.11	3.28	3.07
K ₂ O	1.76	1.82	1.91	1.91	2.04	1.43	1.34	1.33	1.19	1.50	1.52	1.16	1.15
TiO ₂	2.24	2.30	2.16	2.23	2.14	2.00	2.00	2.09	2.06	2.24	2.21	1.71	1.66
P ₂ O ₅	0.35	0.37	0.39	0.41	0.43	0.32	0.30	0.33	0.35	0.43	0.41	0.25	0.24
MnO	0.18	0.19	0.18	0.17	0.21	0.20	0.19	0.20	0.21	0.22	0.21	0.16	0.19
Total	99.72	99.77	99.77	99.80	99.64	99.80	99.78	99.78	99.80	99.78	99.78	99.81	99.81
Trace elements (parts per million)													
Ba	689	755	730	722	844	491	504	609	549	585	530	427	480
BaX	700	710	n.a.	614	873	n.a.	n.a.	n.a.	n.a.	597	n.a.	n.a.	346
Co	35.3	33.3	36.3	40.7	35.7	42.2	42.7	41.9	42.4	39.6	40.7	39.6	39.8
Cr	10.7	7.9	11.8	17.3	10.6	23.4	19.8	19.3	21.9	22.5	22.6	25.5	24.2
Cs	1.40	1.50	1.37	1.10	1.34	1.80	1.13	0.94	n.a.	1.12	1.18	0.95	0.97
Hf	4.71	5.20	4.93	4.77	4.86	3.94	3.98	4.08	4.14	4.27	4.14	3.65	3.59
NbX	14.5	16.0	n.a.	13.0	14.0	n.a.	n.a.	n.a.	n.a.	12.0	n.a.	n.a.	11.0
Rb	49.0	51.0	41.2	38.2	61.0	28.9	33.8	43.9	23.5	30.8	39.5	n.a.	25.8
RbX	53.5	49.0	n.a.	50.0	57.0	n.a.	n.a.	n.a.	n.a.	40.5	n.a.	n.a.	30.0
SrX	332	337	n.a.	380	330	n.a.	n.a.	n.a.	n.a.	342	n.a.	n.a.	298
Ta	0.84	0.96	0.90	0.84	0.88	0.68	0.73	0.67	0.74	0.83	0.83	0.62	0.72
Th	6.05	5.80	6.80	6.71	6.50	3.84	3.77	3.96	3.73	3.95	4.33	3.84	3.48
U	1.50	1.45	n.a.	n.a.	1.50	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.
YX	36.0	39.0	n.a.	38.0	35.0	n.a.	n.a.	n.a.	n.a.	36.0	n.a.	n.a.	31.0
Zn	137	146	146	137	138	136	138	143	145	146	146	122	129
Zr	179	195	n.a.	n.a.	125	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.
ZrX	190	207	n.a.	201	191	n.a.	n.a.	n.a.	n.a.	177	n.a.	n.a.	145
Sc	31.3	30.8	31.9	31.3	30.2	35.0	36.0	35.7	35.2	35.9	36.5	34.2	34.6
La	26.3	27.9	27.7	27.3	27.6	18.6	18.7	19.1	19.1	21.3	20.6	16.5	16.7
Ce	50.3	56.0	58.1	55.8	54.0	38.9	38.8	41.7	39.1	44.6	43.0	34.2	36.5
Nd	27.5	31.0	32.5	32.1	30.0	23.0	23.9	26.8	23.2	26.6	25.8	21.3	20.50
Sm	6.6	7.3	7.6	7.3	7.0	6.2	6.1	6.2	6.1	6.6	6.8	5.4	5.3
Eu	1.86	2.09	2.17	2.15	1.94	1.71	1.77	1.90	1.75	1.99	1.99	1.65	1.66
Gd	6.3	8.2	n.a.	n.a.	6.0	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.
Tb	1.00	1.08	1.23	1.21	1.02	1.07	1.18	1.25	1.00	1.19	1.10	1.14	0.85
Yb	3.42	3.94	3.58	3.1	3.47	2.83	3.10	3.29	2.94	3.35	3.52	2.59	2.41
Lu	0.52	0.59	0.55	0.53	0.51	0.51	0.53	0.54	0.55	0.58	0.59	0.44	0.48
Cu	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.	27.0	n.a.	n.a.	n.a.
Ni	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.	10.5	n.a.	n.a.	n.a.
Normalized major-oxide contents (weight percent)													
SiO ₂	55.41	55.21	55.94	56.38	56.24	54.26	54.21	53.95	54.67	53.78	53.68	54.77	54.06
Al ₂ O ₃	13.97	13.92	14.23	14.15	14.15	13.73	14.30	14.11	13.74	13.80	14.20	14.16	14.45
FeO	12.05	12.27	11.79	11.45	10.30	12.28	11.98	12.47	12.05	12.89	12.67	11.28	11.19
MgO	3.61	3.51	3.41	3.51	3.77	4.60	4.49	4.41	4.50	4.14	4.17	5.06	5.17
CaO	7.13	7.01	6.97	6.45	7.62	8.21	8.28	8.01	8.03	7.84	7.81	8.16	8.81
Na ₂ O	3.29	3.40	3.02	3.33	3.08	2.97	2.91	3.10	3.20	3.16	3.12	3.29	3.08
K ₂ O	1.76	1.82	1.91	1.91	2.05	1.43	1.34	1.33	1.19	1.50	1.52	1.16	1.15
TiO ₂	2.25	2.31	2.16	2.23	2.15	2.00	2.00	2.09	2.06	2.24	2.21	1.71	1.66
P ₂ O ₅	0.35	0.37	0.39	0.41	0.43	0.32	0.30	0.33	0.35	0.43	0.41	0.25	0.24
MnO	0.18	0.19	0.18	0.17	0.21	0.20	0.19	0.20	0.21	0.22	0.21	0.16	0.19
Total	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00

Table 3. Average chemical composition of geochemical units in the Grande Ronde Basalt—*Continued*

Geochemical unit-----	AS16	3F	5E	AS16	AS16	AS16	AS16	3G	3G	5F	5F	AS16	AS16
Chemical type -	4	C	E	4	4	4	4	C	C	E	E	4	4
Major oxides (weight percent)													
SiO ₂	53.90	54.21	54.46	54.39	54.30	53.97	54.43	55.01	54.41	55.68	55.40	54.35	54.47
Al ₂ O ₃	13.97	14.59	14.43	14.41	14.05	14.03	14.17	14.87	14.50	14.38	13.81	13.93	14.27
FeO	12.52	11.01	11.53	11.64	12.15	12.43	12.62	10.11	11.21	11.01	12.02	12.48	11.83
MgO	4.27	4.99	4.49	4.23	3.94	4.43	4.07	4.72	4.79	3.93	3.77	4.05	4.29
CaO	7.89	8.84	8.14	8.13	7.77	7.71	7.27	8.59	8.53	7.56	7.22	7.51	7.93
Na ₂ O	3.13	2.94	2.99	3.12	3.08	3.26	3.22	3.12	2.94	3.22	3.43	3.51	3.30
K ₂ O	1.54	1.12	1.45	1.39	1.72	1.38	1.40	1.19	1.25	1.57	1.57	1.43	1.29
TiO ₂	1.95	1.67	1.79	1.94	2.18	2.01	2.05	1.73	1.70	1.95	2.04	1.98	1.93
P ₂ O ₅	0.41	0.25	0.30	0.34	0.40	0.39	0.34	0.29	0.26	0.30	0.34	0.33	0.27
MnO	0.21	0.18	0.20	0.20	0.20	0.19	0.20	0.19	0.19	0.19	0.18	0.22	0.20
Total	99.79	99.80	99.78	99.79	99.79	99.80	99.77	99.82	99.78	99.79	99.78	99.79	99.78
Trace elements (parts per million)													
Ba	574	497	542	550	566	607	636	611	479	676	547	543	543
BaX	n.a.	528	n.a.	n.a.	614	n.a.	n.a.	n.a.	n.a.	647	n.a.	564	n.a.
Co	38.7	39.1	40.0	38.4	38.7	39.3	39.5	36.4	38.9	39.5	38.5	39.5	38.4
Cr	16.3	22.9	17.2	16.3	21.6	14.5	13.7	24.0	30.6	10.0	5.3	13.8	13.5
Cs	1.01	1.00	1.08	0.90	1.35	0.90	1.18	0.85	1.08	1.19	0.80	1.41	0.70
Hf	4.22	3.61	4.10	4.00	4.15	4.23	4.45	3.89	3.76	4.74	4.95	4.25	4.22
NbX	n.a.	8.0	n.a.	n.a.	12.0	n.a.	n.a.	n.a.	n.a.	12.0	n.a.	13.0	n.a.
Rb	30.1	26.8	35.4	33.9	36.6	37.6	56.4	23.1	39.8	37.7	47.9	40.5	21.4
RbX	n.a.	29.0	n.a.	n.a.	41.0	n.a.	n.a.	n.a.	n.a.	38.0	n.a.	39.0	n.a.
SrX	n.a.	346	n.a.	n.a.	333	n.a.	n.a.	n.a.	n.a.	324	n.a.	306	n.a.
Ta	0.74	0.68	0.69	0.83	0.70	0.72	0.81	0.71	0.64	0.84	0.82	0.64	0.78
Th	3.82	3.58	3.72	4.19	4.59	4.22	4.49	3.95	3.49	4.53	3.96	3.78	3.87
U	0.0	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.
YX	0.0	35.0	n.a.	n.a.	33.0	n.a.	n.a.	n.a.	n.a.	35.0	n.a.	33.0	n.a.
Zn	152	132	135	134	144	137	140	142	132	147	134	127	146.00
Zr	0	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.
ZrX	n.a.	154	n.a.	n.a.	179	n.a.	n.a.	n.a.	n.a.	197	n.a.	178	n.a.
Sc	36.5	35.2	35.8	34.9	35.5	35.5	34.5	35.6	34.7	33.2	32.6	32.8	32.8
La	21.1	17.4	19.5	20.7	22.2	20.2	21.3	19.0	18.1	21.3	22.3	20.1	19.2
Ce	44.6	36.8	40.8	42.5	46.2	41.2	44.8	40.3	36.8	45.8	45.9	42.0	39.3
Nd	26.3	20.7	24.1	24.9	25.5	21.2	26.8	24.1	21.2	25.9	26.0	24.2	22.1
Sm	6.6	5.4	6.1	5.9	6.2	5.8	6.9	5.7	5.3	6.7	6.5	6.0	5.8
Eu	2.00	1.64	1.84	1.82	1.97	1.88	2.04	1.74	1.68	2.06	1.99	1.91	1.75
Gd	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.
Tb	1.03	0.96	1.04	0.97	1.02	1.02	1.17	0.98	0.94	1.12	1.08	1.19	1.24
Yb	3.26	2.51	3.15	2.99	2.95	3.03	3.25	2.84	2.63	3.23	3.19	3.02	2.92
Lu	0.62	0.48	0.54	0.52	0.53	0.59	0.58	0.54	0.49	0.57	0.57	0.57	0.50
Cu	n.a.	37.0	n.a.	n.a.	25.0	n.a.	n.a.	n.a.	n.a.	32.5	n.a.	52.0	n.a.
Ni	n.a.	16.0	n.a.	n.a.	10.0	n.a.	n.a.	n.a.	n.a.	12.0	n.a.	10.0	n.a.
Normalized major-oxide contents (weight percent)													
SiO ₂	54.01	54.32	54.58	54.50	54.41	54.08	54.56	55.11	54.53	55.80	55.52	54.46	54.59
Al ₂ O ₃	14.00	14.62	14.46	14.44	14.08	14.06	14.20	14.90	14.53	14.41	13.84	13.96	14.30
FeO	12.55	11.03	11.56	11.66	12.18	12.45	12.65	10.13	11.23	11.03	12.05	12.51	11.86
MgO	4.28	5.00	4.50	4.24	3.95	4.44	4.08	4.73	4.80	3.94	3.78	4.06	4.30
CaO	7.91	8.86	8.16	8.15	7.79	7.73	7.29	8.61	8.55	7.58	7.24	7.53	7.95
Na ₂ O	3.14	2.95	3.00	3.13	3.09	3.27	3.23	3.13	2.95	3.23	3.44	3.52	3.31
K ₂ O	1.54	1.12	1.45	1.39	1.72	1.38	1.40	1.19	1.25	1.57	1.57	1.43	1.29
TiO ₂	1.95	1.67	1.79	1.94	2.18	2.01	2.05	1.73	1.70	1.95	2.04	1.98	1.93
P ₂ O ₅	0.41	0.25	0.30	0.34	0.40	0.39	0.34	0.29	0.26	0.30	0.34	0.33	0.27
MnO	0.21	0.18	0.20	0.20	0.20	0.19	0.20	0.19	0.19	0.19	0.18	0.22	0.20
Total	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00

Table 3. Average chemical composition of geochemical units in the Grande Ronde Basalt—*Continued*

Geochemical unit----- Chemical type -	AS16 4	AS16 4	5H E	AS16 4	UNC X	AS16 4	AS16 4	2F B	2F B	5I E	AS16 4
Major oxides (weight percent)											
SiO ₂	54.15	53.75	55.24	54.54	56.66	56.60	54.60	54.74	54.68	55.82	54.21
Al ₂ O ₃	14.00	14.55	14.21	14.54	13.88	13.52	14.67	13.61	13.58	14.02	13.75
FeO	12.46	12.61	11.50	11.60	11.67	11.87	11.61	12.86	12.81	11.33	12.13
MgO	4.07	4.40	4.00	4.24	3.12	3.32	4.20	3.82	3.62	3.96	4.39
CaO	7.88	7.98	7.66	7.83	6.65	6.88	7.71	7.29	7.13	7.49	8.34
Na ₂ O	3.04	2.88	3.03	3.13	3.18	3.09	3.09	3.17	3.46	3.09	2.85
K ₂ O	1.57	1.06	1.67	1.52	1.99	1.94	1.44	1.54	1.58	1.61	1.44
TiO ₂	2.07	2.11	1.99	1.93	2.14	2.09	1.99	2.16	2.27	2.00	2.15
P ₂ O ₅	0.36	0.26	0.29	0.29	0.32	0.30	0.32	0.37	0.44	0.31	0.34
MnO	0.20	0.18	0.19	0.19	0.18	0.19	0.18	0.23	0.20	0.19	0.20
Total	99.80	99.78	99.78	99.81	99.79	99.80	99.81	99.79	99.77	99.82	99.80
Trace elements (parts per million)											
Ba	536	496	642	639	710	641	597	557	644	574	534
BaX	n.a.	n.a.	n.a.	n.a.	634	681	n.a.	583	n.a.	n.a.	n.a.
Co	37.9	39.5	40.3	38.3	37.0	36.5	38.5	38.5	38.8	39.0	40.5
Cr	10.9	13.0	12.3	14.6	9.0	10.2	16.7	17.2	14.5	11.3	35.3
Cs	1.31	1.12	1.30	1.20	1.86	1.44	1.46	0.80	1.22	1.33	1.57
Hf	4.55	4.67	4.62	4.43	5.22	5.05	4.80	4.94	5.33	4.82	4.64
NbX	n.a.	n.a.	n.a.	n.a.	15.5	15.0	n.a.	16.0	n.a.	n.a.	n.a.
Rb	47.8	25.5	52.4	30.8	53.0	49.2	34.9	50.1	39.6	47.2	40.4
RbX	n.a.	n.a.	n.a.	n.a.	64.0	54.0	n.a.	42.0	n.a.	n.a.	n.a.
SrX	n.a.	n.a.	n.a.	n.a.	337	327	n.a.	305	n.a.	n.a.	n.a.
Ta	0.69	0.93	0.90	0.83	0.95	0.90	0.77	0.85	0.81	0.87	0.88
Th	3.83	4.73	4.71	4.39	5.81	5.54	4.87	4.30	4.49	5.26	3.79
U	0.0	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.
YX	n.a.	n.a.	n.a.	n.a.	40.0	35.0	n.a.	39.0	n.a.	n.a.	n.a.
Zn	132	137	128	146	145	135	147	137	143	117	125
Zr	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.
ZrX	n.a.	n.a.	n.a.	n.a.	217	206	n.a.	200	n.a.	n.a.	n.a.
Sc	33.7	33.3	32.7	34.0	30.1	30.6	31.5	33.1	33.2	31.0	34.1
La	21.1	21.1	21.7	20.3	24.8	23.3	23.1	22.1	23.7	22.2	21.2
Ce	44.5	44.0	45.5	41.9	52.4	48.6	48.7	47.4	51.6	46.5	46.1
Nd	26.9	26.5	27.7	25.3	31.6	29.0	27.4	30.6	34.7	26.6	26.7
Sm	6.4	6.7	6.6	6.4	7.2	6.7	7.0	6.9	7.5	6.4	6.7
Eu	1.97	1.96	1.87	1.91	1.98	2.00	1.95	2.11	2.26	1.87	1.96
Gd	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.
Tb	1.01	0.98	1.16	1.03	1.16	1.15	1.11	1.21	1.23	1.00	1.17
Yb	3.06	3.04	3.09	3.19	3.39	3.19	3.07	3.45	3.84	2.87	3.40
Lu	0.58	0.56	0.52	0.57	0.59	0.56	0.54	0.57	0.63	0.51	0.55
Cu	n.a.	n.a.	n.a.	n.a.	19.0	21.0	n.a.	42.0	n.a.	n.a.	n.a.
Ni	n.a.	n.a.	n.a.	n.a.	10.0	10.0	n.a.	14.0	n.a.	n.a.	n.a.
Normalized major-oxide contents (weight percent)											
SiO ₂	54.26	53.87	55.36	54.64	56.78	56.71	54.70	54.86	54.81	55.92	54.32
Al ₂ O ₃	14.03	14.58	14.24	14.57	13.91	13.55	14.70	13.64	13.61	14.05	13.78
FeO	12.48	12.64	11.53	11.62	11.69	11.89	11.63	12.89	12.84	11.35	12.15
MgO	4.08	4.41	4.01	4.25	3.13	3.33	4.21	3.83	3.63	3.97	4.40
CaO	7.90	8.00	7.68	7.84	6.66	6.89	7.72	7.31	7.15	7.50	8.36
Na ₂ O	3.05	2.89	3.04	3.14	3.19	3.10	3.10	3.18	3.47	3.10	2.86
K ₂ O	1.57	1.06	1.67	1.52	1.99	1.94	1.44	1.54	1.58	1.61	1.44
TiO ₂	2.07	2.11	1.99	1.93	2.14	2.09	1.99	2.16	2.28	2.00	2.15
P ₂ O ₅	0.36	0.26	0.29	0.29	0.32	0.30	0.32	0.37	0.44	0.31	0.34
MnO	0.20	0.18	0.19	0.19	0.18	0.19	0.18	0.23	0.20	0.19	0.20
Total	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00

Table 4. Average chemical composition of geochemical units in the Wanapum Basalt

[Chemical types listed in approximate stratigraphic order, see table 1. Trace elements followed by X analyzed by X-ray fluorescence; all others analyzed by instrumental neutron activation. n.a., not analyzed; Data from which these average compositions are calculated are given in Wright (1988)]

Geochemical unit----- Chemical type-	PV A	LOLO INC B	ROSALIA C	PH POWAT D	ROZA E	FS ANOM F	FS INC F	SHUMAKER G	PH SHUM G	SW LOOK H	DODGE I	ROBIN J
Major oxides (weight percent)												
SiO ₂	51.27	49.98	49.78	53.82	50.74	52.68	51.51	54.36	54.29	54.48	52.15	49.78
Al ₂ O ₃	15.41	14.13	13.39	14.59	14.21	13.94	13.43	13.86	14.38	14.66	15.41	16.78
FeO	10.23	13.89	14.73	12.93	13.45	11.16	14.30	13.42	13.46	13.28	10.73	10.13
MgO	3.75	5.15	4.48	3.23	4.55	4.29	4.42	2.89	2.97	2.91	5.89	8.19
CaO	10.42	8.80	8.64	7.00	8.53	8.68	8.05	6.39	6.55	6.54	9.86	10.80
Na ₂ O	3.13	2.58	2.79	3.03	2.84	2.86	2.73	3.23	3.41	3.05	2.90	2.42
K ₂ O	1.18	1.05	1.21	1.93	1.25	1.38	1.35	1.87	1.81	1.96	0.68	0.37
TiO ₂	3.50	3.22	3.59	2.67	3.11	3.52	3.01	2.45	2.47	2.71	1.45	0.97
P ₂ O ₅	0.79	0.73	0.84	1.15	0.68	0.71	0.61	0.90	0.89	0.75	0.36	0.17
MnO	0.15	0.19	0.20	0.24	0.25	0.23	0.22	0.30	0.26	0.22	0.15	0.14
Total	99.83	99.72	99.65	100.59	99.61	99.45	99.63	99.67	100.49	100.56	99.58	99.75
Trace elements (parts per million)												
Ba	677	539	600	1043	556	812	549	991	1024	762	348	175
BaX	650	481	524	955	572	807	457	1001	972	761	327	135
Co	28.8	40.2	37.7	n.a.	38.5	52.9	38.8	23.9	n.a.	n.a.	36.9	46.6
Cr	102.9	82	13.6	n.a.	44.3	24.4	39.5	4.4	4.9	6.8	159.7	151.4
Cs	0.62	0.70	0.89	1.15	0.78	0.95	0.93	1.02	0.86	1.21	0.50	n.a.
Hf	5.02	4.66	5.43	6.40	4.64	5.08	4.42	6.16	6.33	5.63	2.86	1.48
NbX	19.0	15.8	17.0	20.0	17.0	17.5	13.0	19.3	24.0	18.5	8.7	6.0
Rb	28.2	26.6	34.6	38.9	31.4	37.5	35.4	35.5	36.0	36.7	15.8	n.a.
RbX	31.5	27.3	31.7	42.0	37.8	45.0	32.3	42.7	44.0	49.0	13.6	n.a.
SrX	330	302	286	354	313	353	317	335	308	331	397	353
Ta	1.18	1.09	1.22	1.26	1.05	1.22	0.98	1.33	1.29	1.29	0.51	0.31
Th	3.89	3.56	4.04	5.44	3.98	4.91	3.88	5.27	5.52	5.47	1.16	0.37
U	1.11	0.88	0.93	1.64	1.22	1.56	0.85	1.66	1.45	1.56	0.52	n.a.
YX	50.5	41.8	43.7	56.0	44.3	58.0	41.3	55.0	52.0	48.5	31	20.3
Zn	208	170	173	152	163	216	172	165	153	141	123	112
Zr	133	223	238	270	223	202	196	307	255	239	136	96
ZrX	207	198	224	260	200	219	185	256	260	235	124	68
Sc	39.9	36.0	37.3	31.5	36.1	38.6	35.8	33.4	33.6	30.6	40.1	36.7
La	30.0	28.0	31.1	37.1	27.0	31.1	25.4	36.1	35.7	30.0	15.2	7.6
Ce	58.9	57.4	62.7	82.0	54.2	57.7	51.0	75.2	77.7	64.4	32.3	16.3
Nd	37.5	34.4	35.8	49.9	34.8	35.9	30.9	44.7	47.0	38.4	20.8	12.8
Sm	9.4	7.9	7.6	11.7	6.9	9.0	6.1	11.1	11.2	9.0	4.2	2.7
Eu	2.69	2.56	2.66	3.35	2.25	2.34	2.16	3.24	3.28	2.48	1.41	1.00
Gd	9.2	8.6	9.4	11.5	7.9	8.9	7.9	10.1	10.5	8.7	5.0	3.6
Tb	1.47	1.37	1.44	1.87	1.19	1.39	1.21	1.60	1.67	1.36	0.81	0.63
Yb	4.62	3.92	4.62	5.69	3.89	4.63	3.52	5.55	5.62	4.65	3.08	2.10
Lu	0.73	0.62	0.69	0.92	0.64	0.78	0.59	0.86	0.91	0.78	0.47	0.35
Cu	45.0	37.3	27.7	n.a.	39.6	29.5	35.3	29.5	n.a.	15.0	105.7	78.7
Ni	31.5	35.0	27.0	n.a.	33.6	46.5	39.3	2.5	n.a.	1	42.7	117
Normalized major-oxide contents (weight percent)												
SiO ₂	51.36	50.12	49.95	53.50	50.94	52.97	51.70	54.54	54.03	54.18	52.37	49.90
Al ₂ O ₃	15.44	14.17	13.44	14.50	14.27	14.02	13.48	13.91	14.31	14.58	15.47	16.82
FeO	10.25	13.93	14.78	12.85	13.50	11.22	14.35	13.46	13.39	13.21	10.78	10.16
MgO	3.76	5.16	4.50	3.21	4.57	4.31	4.44	2.90	2.96	2.89	5.91	8.21
CaO	10.44	8.82	8.67	6.96	8.56	8.73	8.08	6.41	6.52	6.50	9.90	10.83
Na ₂ O	3.14	2.59	2.80	3.01	2.85	2.88	2.74	3.24	3.39	3.03	2.91	2.43
K ₂ O	1.18	1.05	1.21	1.92	1.25	1.39	1.36	1.88	1.80	1.95	0.68	0.37
TiO ₂	3.51	3.23	3.60	2.65	3.12	3.54	3.02	2.46	2.46	2.69	1.46	0.97
P ₂ O ₅	0.79	0.73	0.84	1.14	0.68	0.71	0.61	0.90	0.89	0.75	0.36	0.17
MnO	0.15	0.19	0.20	0.24	0.25	0.23	0.22	0.30	0.26	0.22	0.15	0.14
Total	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00

Table 5. Average chemical composition of geochemical units in the Saddle Mountains Basalt

[Chemical types listed in approximate stratigraphic order, see table 1. Trace elements followed by X analyzed by X-ray fluorescence; all others analyzed by instrumental neutron activation. n.a., not analyzed. Data from which these average compositions are calculated are given in Wright (1988)]

Geochemical unit----- Chemical type-	LM A	GOOSE B	INDIAN C	MARTIN D	BASIN E	BUFORD F	ELEPHANT G	POMONA H	VC WEIPE I	ESQUAT J	RBESQUAT J	FS IC K	SLIP L
Major oxides (weight percent)													
SiO ₂	50.01	47.54	49.17	48.58	47.35	53.82	51.19	51.94	51.72	54.14	53.51	52.47	52.00
Al ₂ O ₃	14.12	12.34	13.76	13.69	13.74	14.17	13.26	15.02	15.30	13.82	14.35	13.48	14.29
FeO	14.16	17.36	14.36	14.40	14.98	11.34	14.77	10.44	11.26	12.62	13.33	13.90	11.69
MgO	4.87	4.50	5.40	5.87	6.03	4.98	4.17	7.18	6.49	3.97	3.70	4.20	5.47
CaO	8.60	8.76	9.13	9.88	9.72	8.47	8.34	10.38	10.39	7.38	7.53	7.60	9.60
Na ₂ O	2.68	2.42	2.41	2.36	2.27	2.77	2.44	2.23	2.53	2.57	2.54	2.66	2.64
K ₂ O	1.43	1.21	1.00	0.74	0.78	1.41	1.25	0.65	0.49	1.71	1.65	1.61	0.91
TiO ₂	2.92	3.80	3.40	3.31	3.69	2.21	3.60	1.62	1.83	2.91	3.15	3.16	2.43
P ₂ O ₅	0.64	1.56	0.84	0.74	0.87	0.38	0.60	0.24	0.29	0.44	0.42	0.52	0.53
MnO	0.20	0.27	0.23	0.21	0.24	0.15	0.23	0.18	0.19	0.22	0.19	0.21	0.15
Total	99.63	99.76	99.70	99.78	99.67	99.70	99.85	99.88	100.49	99.78	100.37	99.81	99.71
Trace elements (parts per million)													
Ba	541	852	701	638	611	425	485	243	292	592	611	592	423
BaX	523	830	687	612	603	444	320	173	265	577	658	481	308
Co	41.6	38.1	41.6	43.1	46.5	39.2	45.0	43.5	n.a.	40.7	n.a.	46.1	43.8
Cr	24.0	41.3	117.5	194.9	148.8	44.5	18.5	109.0	51.8	18.9	7.1	21.4	38.3
Cs	1.23	0.45	0.50	n.a.	n.a.	0.85	0.70	0.90	n.a.	0.98	0.89	0.78	n.a.
Hf	4.63	10.74	6.74	5.90	6.59	5.30	6.41	3.23	3.28	5.57	5.81	5.38	4.40
NbX	25.7	48.5	29.0	21.3	23.7	21.3	24.5	13.0	12.5	21.80	23.0	24.0	18.0
Rb	18.0	30.1	28.3	20.6	19.8	35.5	27.6	10.0	9.0	47.3	32.0	35.30	19.0
RbX	35.3	22.5	16.0	10.0	10.0	43.3	24.0	18.0	26.5	51.5	50.0	46.5	25.0
SrX	350	220	233	235	227	230	213	227	235	257	269	245	244
Ta	1.68	3.07	1.92	1.66	1.91	1.53	1.74	0.82	0.81	1.63	1.67	1.79	1.24
Th	4.98	4.34	3.53	2.10	2.19	7.10	5.91	2.55	2.47	8.67	8.93	7.23	4.70
U	1.38	1.12	0.90	0.65	0.75	1.70	1.42	0.77	0.47	2.08	1.61	1.80	1.10
YX	35.3	89.5	52.0	45.0	52.3	44.0	42.0	26.5	27.5	41.3	38.0	35.5	28.0
Zn	147	226	173	161	177	129	179	118	102	134	127	164	135
Zr	350	502	297	304	293	n.a.	316	n.a.	170	240	250	250	n.a.
ZrX	189	436	273	238	273	216	229	137	140	226	223	219	180
Sc	22.6	37.5	38.5	39.3	39.0	31.2	31.7	34.4	35.8	27.7	28.4	29.3	33.4
La	35.3	74.1	46.0	36.8	42.9	33.0	33.7	17.0	17.1	38.2	38.6	35.8	28.0
Ce	67.8	149	90.9	73.6	85.6	63.0	68.7	35.3	35.4	73.8	74.6	69.3	57.5
Nd	34.0	86.9	53.3	47.3	53.7	33.5	35.3	20.5	21.1	36.7	38.5	34.3	31.0
Sm	7.1	18.7	11.0	9.1	10.7	7.3	9.4	4.7	5.0	8.4	8.3	8.3	7.0
Eu	2.10	5.41	3.22	2.82	3.31	1.83	2.44	1.41	1.47	2.18	2.24	2.15	1.94
Gd	8.2	18.4	10.1	9.0	9.7	6.9	8.7	5.0	5.1	8.0	8.2	8.2	7.5
Tb	1.34	2.66	1.21	1.11	1.24	1.41	1.39	0.68	0.82	1.49	0.98	1.42	1.12
Yb	3.13	9.08	5.55	5.01	5.64	3.90	4.58	2.73	2.66	3.65	3.58	3.75	3.40
Lu	0.46	1.35	0.82	0.74	0.82	0.57	0.66	0.39	0.44	0.51	0.53	0.53	0.48
Cu	17.5	27.0	39.0	33.0	46.0	32.5	39.7	63.0	n.a.	34.5	n.a.	28.0	n.a.
Ni	28.5	9.8	13.0	34.0	42.0	40.5	30.7	51.3	n.a.	16.0	n.a.	16.0	n.a.
Normalized major-oxide contents (weight percent)													
SiO ₂	50.20	47.65	49.32	48.69	47.51	53.98	51.27	52.00	51.47	54.26	53.31	52.57	52.15
Al ₂ O ₃	14.17	12.37	13.80	13.72	13.79	14.21	13.28	15.04	15.23	13.85	14.30	13.51	14.33
FeO	14.21	17.40	14.40	14.43	15.03	11.37	14.79	10.45	11.21	12.65	13.28	13.93	11.72
MgO	4.89	4.51	5.42	5.88	6.05	4.99	4.18	7.19	6.46	3.98	3.69	4.21	5.49
CaO	8.63	8.78	9.16	9.90	9.75	8.50	8.35	10.39	10.34	7.40	7.50	7.61	9.63
Na ₂ O	2.69	2.43	2.42	2.37	2.28	2.78	2.44	2.23	2.52	2.58	2.53	2.67	2.65
K ₂ O	1.44	1.21	1.00	0.74	0.78	1.41	1.25	0.65	0.49	1.71	1.64	1.61	0.91
TiO ₂	2.93	3.81	3.41	3.32	3.70	2.22	3.61	1.62	1.82	2.92	3.14	3.17	2.44
P ₂ O ₅	0.64	1.56	0.84	0.74	0.87	0.38	0.60	0.24	0.29	0.44	0.42	0.52	0.53
MnO	0.20	0.27	0.23	0.21	0.24	0.15	0.23	0.18	0.19	0.22	0.19	0.21	0.15
Total	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00

Table 5. Average chemical composition of geochemical units in the Saddle Mountains Basalt—Continued

Geochemical unit-----	LEW ORCH	SPRAGUE	SW SPRAG	NEW	SW NEW	ASOTIN	VC ASOT	SW HUNTZ	HUNTZ	VC LAP	EAGLE	WILBUR	UMATILLA
Chemical type- M	N	N	N	O	O	P	P	Q	Q	R	R	S	T
Major oxides (weight percent)													
SiO2	49.63	50.76	49.65	49.22	50.11	50.76	51.11	52.37	52.50	53.08	52.86	54.23	54.51
Al2O3	15.22	13.72	14.54	15.64	15.95	16.02	16.54	16.05	15.39	15.85	15.35	14.46	14.18
FeO	11.80	12.60	13.41	10.85	10.42	9.52	9.83	9.56	10.37	10.47	10.52	11.16	12.23
MgO	7.09	5.07	6.39	7.97	7.41	8.23	7.88	7.56	6.52	5.84	5.99	4.50	2.74
CaO	10.08	9.83	9.92	11.00	11.50	10.77	10.78	10.18	9.38	9.26	9.16	8.40	6.18
Na2O	2.32	2.64	2.63	2.19	2.51	2.29	1.99	2.43	2.34	2.33	2.45	2.64	3.37
K2O	0.46	0.87	0.61	0.40	0.29	0.39	0.46	0.65	1.22	1.32	1.15	1.76	2.55
TiO2	2.42	3.17	2.73	1.89	1.93	1.48	1.48	1.47	1.63	1.71	1.70	1.90	2.78
P2O5	0.55	0.79	0.64	0.47	0.47	0.19	0.24	0.28	0.34	0.41	0.41	0.51	0.82
MnO	0.21	0.18	0.22	0.15	0.18	0.14	0.20	0.20	0.20	0.17	0.13	0.23	0.19
Total	99.78	99.63	100.74	99.78	100.77	99.79	100.51	100.75	99.89	100.44	99.72	99.79	99.55
Trace elements (parts per million)													
Ba	410	767	506	320	342	303	277	352	410	665	570	876	3313
BaX	351	598	486	329	336	249	278	262	n.a.	598	593	757	3184
Co	43.8	46.2	n.a.	45.0	n.a.	46.9	n.a.	n.a.	42.7	n.a.	41.2	42.5	28.1
Cr	267.0	48.6	150.5	300.5	307.5	283.7	272.5	222.0	172.0	115.4	98.1	36.6	3.6
Cs	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.	0.34	0.50	0.63	0.70	0.80	0.73
Hf	3.90	6.35	5.23	3.05	3.05	2.78	2.90	3.23	4.20	4.89	5.13	5.90	10.22
NbX	18.0	28.0	23.0	18.0	16.5	13.0	12.0	13.0	n.a.	18.30	16	16.70	21.30
Rb	n.a.	n.a.	17.5	n.a.	13.4	10.0	17.5	22.0	26.5	31.5	38.8	38.2	47.0
RbX	n.a.	6.0	n.a.	13.0	9.0	14.5	10.0	17.0	n.a.	34.30	24.0	43.3	43.0
SrX	242	253	229	252	259	252	249	240	n.a.	274	296	277	276
Ta	1.23	1.64	1.49	0.96	0.88	0.65	0.62	0.80	0.72	0.91	1.18	1.06	1.49
Th	1.53	3.00	2.34	1.40	1.35	2.05	2.15	2.63	3.90	5.13	5.04	6.65	7.25
U	0.95	1.45	0.61	n.a.	1	n.a.	0.70	0.61	0.50	1.08	1.17	1.10	1.43
YX	34.5	47.0	46.5	35.0	33.0	26.7	25.0	24.0	n.a.	43.0	37.0	41.7	49.7
Zn	131	162	128	116	111	107	103	87	103	117	126	131	135
Zr	203	360	255	n.a.	160	n.a.	126	140	195	230	234	282	520
ZrX	172	274	228	141	134	121	120	141	n.a.	216	164	249	482
Sc	37.3	37.4	38.5	34.6	35.7	30.8	30.9	29.2	28.7	27.8	29.0	26.6	26.2
La	27.8	42.5	34.7	20.0	20.5	15.7	15.0	17.7	27.0	33.3	31.6	43.7	46.7
Ce	56.0	87.0	73.9	41.0	41.5	31.5	30.5	36.2	52.0	62.1	61.2	80.5	88.2
Nd	29.3	48.0	42.4	24.0	27.0	18.0	19.0	19.5	27.5	33.8	32.3	39.5	49.2
Sm	6.8	10.3	9.2	5.3	5.1	4.3	4.0	4.5	5.9	6.6	6.5	7.8	9.8
Eu	2.03	2.69	2.44	1.61	1.71	1.28	1.29	1.30	1.56	1.71	1.92	1.93	3.95
Gd	6.9	9.5	8.2	4.2	6.4	4.7	4.5	4.4	5.6	6.8	7.2	7.6	10.2
Tb	1.11	1.63	1.32	0.86	0.93	0.74	0.80	0.73	0.82	0.99	1.04	1.32	1.69
Yb	3.53	4.80	4.17	2.50	2.85	2.37	2.45	2.30	2.85	3.59	3.85	4.23	4.52
Lu	0.53	0.69	0.75	0.40	0.42	0.35	0.35	0.49	0.55	0.56	0.56	0.63	0.66
Cu	59.5	46.0	57.0	67.0	89.0	111.7	100.0	96.0	n.a.	49.0	52.5	25.0	19.0
Ni	84.5	70.0	85.0	140.0	140.0	203.3	140.0	170.0	n.a.	54.0	93.0	47.0	2.7
Normalized major-oxide contents (weight percent)													
SiO2	49.74	50.95	49.29	49.33	49.73	50.87	50.85	51.98	52.56	52.85	53.01	54.34	54.76
Al2O3	15.25	13.77	14.43	15.67	15.83	16.05	16.46	15.93	15.41	15.78	15.39	14.49	14.24
FeO	11.83	12.65	13.31	10.87	10.34	9.54	9.78	9.49	10.38	10.42	10.55	11.18	12.29
MgO	7.11	5.09	6.34	7.99	7.35	8.25	7.84	7.50	6.53	5.81	6.01	4.51	2.75
CaO	10.10	9.87	9.85	11.02	11.41	10.79	10.73	10.10	9.39	9.22	9.19	8.42	6.21
Na2O	2.33	2.65	2.61	2.19	2.49	2.29	1.98	2.41	2.34	2.32	2.46	2.65	3.39
K2O	0.46	0.87	0.61	0.40	0.29	0.39	0.46	0.65	1.22	1.31	1.15	1.76	2.56
TiO2	2.43	3.18	2.71	1.89	1.92	1.48	1.47	1.46	1.63	1.70	1.70	1.90	2.79
P2O5	0.55	0.79	0.64	0.47	0.47	0.19	0.24	0.28	0.34	0.41	0.41	0.51	0.82
MnO	0.21	0.18	0.22	0.15	0.18	0.14	0.20	0.20	0.20	0.17	0.13	0.23	0.19
Total	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00

Table 5. Average chemical composition of geochemical units in the Saddle Mountains Basalt—*Continued*

Geochemical unit-----UM Chemical type- T	FRAC T	PH UMTIL T	PH SOPHER U
Major oxides (weight percent)			
SiO ₂	59.09	55.56	54.49
Al ₂ O ₃	16.62	15.04	15.08
FeO	5.79	11.77	11.76
MgO	0.60	2.49	2.89
CaO	5.32	6.01	6.63
Na ₂ O	4.10	3.16	3.06
K ₂ O	3.51	2.60	2.41
TiO ₂	3.27	2.71	3.06
P ₂ O ₅	1.20	0.97	0.86
MnO	0.06	0.19	0.23
Total	99.56	100.50	100.47
Trace elements (parts per million)			
Ba	4410	3585	3153
BaX	4161	3478	3281
Co	12.7	n.a.	n.a.
Cr	3.2	1.7	1.4
Cs	0.69	0.63	0.65
Hf	12.48	10.55	9.83
NbX	25.0	20.0	17.5
Rb	59.8	56.3	42.3
RbX	56.0	44.0	35.0
SrX	346	285	338
Ta	1.58	1.62	1.44
Th	8.89	7.45	7.00
U	1.74	1.42	1.02
YX	54.0	49.0	43.0
Zn	199	135	139
Zr	631	543	494
ZrX	546	517	462
Sc	29.9	25.8	27.4
La	57.0	48.8	45.0
Ce	114.4	89.3	85.5
Nd	57.3	50.5	43.3
Sm	12.6	9.5	8.5
Eu	5.15	4.12	3.68
Gd	12.0	10.70	9.2
Tb	1.74	1.49	1.29
Yb	4.39	4.55	4.28
Lu	0.66	0.66	0.65
Cu	5.0	26.0	15.0
Ni	n.a.	75.0	5.0
Normalized major-oxide contents (weight percent)			
SiO ₂	59.35	55.28	54.24
Al ₂ O ₃	16.69	14.97	15.01
FeO	5.82	11.71	11.70
MgO	0.60	2.48	2.88
CaO	5.34	5.98	6.60
Na ₂ O	4.12	3.14	3.05
K ₂ O	3.53	2.59	2.40
TiO ₂	3.28	2.70	3.05
P ₂ O ₅	1.21	0.97	0.86
MnO	0.06	0.19	0.23
Total	100.00	100.00	100.00

Table 6. Average chemical composition of feeder dikes in the Yakima Basalt Subgroup

[Chemical types listed in approximate stratigraphic order, see table 1. Trace elements followed by X analyzed by X-ray fluorescence; all others analyzed by instrumental neutron activation. n.a., not analyzed. Data from which these average compositions are calculated are given in Wright (1988). GRB CT1 and GRB CT3 refer to geochemical units 1 and 3 in table 3]

Geochemical unit-----	ELEPHANT G	UMATILLA T	LOLOINC B	ROSALIA C	ROZA E	FSINC F	SHUMAKER G	DODGE I	ROBIN J	GRB CT1 A	GRB CT3 C	GRB CT3 C	GRB LH2 1	GRB MG6 2
Chemical type-	G	T	B	C	E	F	G	I	J	A	C	C	1	2
Major oxides (weight percent)														
SiO ₂	51.25	54.68	50.75	50.23	51.07	51.43	54.43	51.94	49.43	54.29	53.22	54.54	54.48	52.56
Al ₂ O ₃	13.22	13.07	13.57	12.76	13.46	13.49	13.06	14.56	16.58	14.49	14.44	14.21	14.83	14.38
FeO	14.75	13.34	13.77	15.35	14.05	14.19	13.42	10.70	10.39	11.15	11.47	11.39	9.48	11.63
MgO	4.07	3.03	5.26	4.53	4.59	4.54	2.91	6.42	8.59	4.90	5.49	4.57	6.15	5.82
CaO	8.10	5.66	8.47	8.19	8.33	8.12	6.65	10.16	11.03	8.53	9.11	8.09	9.39	9.29
Na ₂ O	2.64	3.24	2.73	2.47	2.75	2.73	3.51	3.16	2.35	2.86	2.75	3.15	2.80	2.72
K ₂ O	1.22	2.51	0.84	1.13	1.43	1.34	2.01	0.61	0.23	1.22	1.11	1.22	0.96	1.03
TiO ₂	3.76	2.72	3.25	3.71	3.26	2.94	2.47	1.53	0.92	1.73	1.73	1.93	1.27	1.78
P ₂ O ₅	0.49	1.02	0.83	0.84	0.60	0.62	0.83	0.36	0.18	0.35	0.32	0.33	0.29	0.36
MnO	0.20	0.17	0.18	0.19	0.23	0.25	0.28	0.16	0.19	0.19	0.17	0.17	0.16	0.17
Total	99.70	99.44	99.65	99.40	99.77	99.65	99.57	99.60	99.89	99.71	99.81	99.60	99.81	99.74
Trace elements (parts per million)														
Ba	447	3510	460	531	560	537	1070	357	145	487	401	524	390	419
BaX	480	3460	490	544	559	524	1050	345	151	498	427	525	433	434
Co	41.3	25.3	36.1	36.5	39.1	38.7	23.9	36.1	45.9	38.1	38.7	35.3	32.1	38.7
Cr	18.3	4.1	91.5	18.5	33.6	46.6	4.4	161.0	152.5	44.1	83.6	26.6	86.3	112.1
Cs	0.50	n.a.	1.20	n.a.	0.80	0.95	0.60	n.a.	n.a.	0.90	0.65	0.90	0.40	0.68
Hf	6.10	9.80	3.90	5.00	4.90	4.50	6.50	3.00	1.55	3.50	3.20	4.30	2.70	3.18
NbX	10.0	5.0	17.5	19.0	19.0	15.0	20.0	10.0	9.0	12.0	12.5	12.5	11.0	12.0
Rb	35.0	42.0	28.5	19.0	38.0	41.3	51.0	13.0	4.0	32.0	27.5	41.0	20.5	31.8
RbX	15.0	44.0	31.0	20.0	39.0	39.5	53.0	14.0	6.0	32.0	27.5	40.5	23.5	31.0
SrX	382	291	330	345	322	318	342	377	332	346	343	329	392	386
Ta	1.71	1.40	3.00	3.60	4.20	4.10	5.70	1.10	0.50	3.20	2.55	4.20	1.75	2.25
U	1.20	1.20	0.75	0.80	1.10	0.93	1.50	n.a.	n.a.	0.80	0.80	1.20	0.45	0.65
YX	29.0	48.0	40.5	45.0	40.0	36.5	59.0	28.0	21.5	31.0	29.5	32.0	25.5	29.3
Zn	137	134	131	154	140	146	165	110	89	120	105	118	90	114
Zr	190	519	150	210	200	145	270	90	n.a.	120	170	230	125	160
ZrX	125	484	180	216	200	185	263	124	74	151	143	172	126	140
Sc	28.9	25.0	32.7	36.7	37.4	35.4	36.0	46.0	36.5	33.6	36.3	33.0	36.4	38.8
La	31.0	45.0	24.0	28.0	27.0	24.3	38.0	14.0	7.0	18.0	16.0	20.5	14.5	16.5
Ce	60.0	86.0	49.0	55.0	56.0	49.3	79.0	30.0	15.0	36.0	33.0	39.5	30.0	32.8
Nd	34.0	47.0	30.5	33.0	33.0	27.8	47.0	21.0	10.5	21.0	22.0	22.5	17.5	19.5
Sm	8.3	10.2	7.6	7.5	8.4	5.6	10.8	4.9	2.9	49.0	4.6	5.7	4.1	13.4
Eu	2.41	4.24	2.34	2.45	2.42	2.02	3.58	1.40	0.94	1.48	1.50	1.68	1.22	1.52
Gd	8.8	10.5	6.5	7.8	8.2	7.9	12.7	6.4	2.3	55.0	4.8	7.0	4.0	4.4
Tb	1.22	1.39	1.09	1.17	1.21	1.10	1.72	0.73	0.55	0.81	0.71	0.95	0.60	0.70
Yb	4.30	4.20	3.65	4.40	4.10	3.73	5.80	3.10	2.35	3.00	2.85	3.25	2.40	3.00
Lu	0.63	0.63	0.54	0.63	0.60	0.54	0.86	0.48	0.35	0.44	0.45	0.47	0.37	0.44
Cu	25.0	2.0	43.5	30.0	32.0	29.5	17.0	110.0	92.0	50.0	55.0	35.5	67.0	54.7
Ni	14.0	1.0	44.0	14.0	19.0	10.0	1.0	41.0	93.0	21.0	23.5	12.0	23.5	20.3
Normalized major-oxide contents (weight percent)														
SiO ₂	51.40	54.99	50.93	50.53	51.19	51.61	54.67	52.15	49.48	54.45	53.32	54.76	54.58	52.70
Al ₂ O ₃	13.26	13.14	13.62	12.84	13.49	13.54	13.12	14.62	16.60	14.53	14.47	14.27	14.86 ₃	14.42
FeO	14.79	13.42	13.82	15.44	14.08	14.24	13.48	10.74	10.40	11.18	11.49	11.44	9.50	11.66
MgO	4.08	3.05	5.28	4.56	4.60	4.56	2.92	6.45	8.60	4.91	5.50	4.59	6.16	5.84
CaO	8.12	5.69	8.50	8.24	8.35	8.15	6.68	10.20	11.04	8.55	9.13	8.12	9.41	9.31
Na ₂ O	2.65	3.26	2.74	2.48	2.76	2.74	3.53	3.17	2.35	2.87	2.76	3.16	2.81	2.73
K ₂ O	1.22	2.52	0.84	1.14	1.43	1.34	2.02	0.61	0.23	1.22	1.11	1.22	0.96	1.03
TiO ₂	3.77	2.74	3.26	3.73	3.27	2.95	2.48	1.54	0.92	1.74	1.73	1.94	1.27	1.78
P ₂ O ₅	0.49	1.03	0.83	0.85	0.60	0.62	0.83	0.36	0.18	0.35	0.32	0.33	0.29	0.36
MnO	0.20	0.17	0.18	0.19	0.23	0.25	0.28	0.16	0.19	0.19	0.17	0.17	0.16	0.17
Total	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00

Table 7. Ratios of incompatible trace elements in samples of the Grande Ronde Basalt

[Geochemical units listed in approximate stratigraphic order, see table 1. n.a., not analyzed]

Geochemical unit----- Chemical type-	1A N2 A	1A* N2 A	2A N2 B	UNC N2 X	3A N2 C	3A* N2 C	UNC N2 X	4A N2 D	4A* N2 D	UNC N2 X	3B N2 C	UNC N2 X	4B N2 D
K/Rb	3.48	3.18	n.a.	n.a.	3.28	3.10	3.19	3.20	3.58	n.a.	3.16	n.a.	n.a.
K/Th	27.14	30.44	n.a.	n.a.	24.53	26.86	28.93	25.07	28.82	n.a.	26.99	n.a.	n.a.
K/P	7.04	8.96	5.58	4.99	6.82	9.09	7.09	5.93	7.92	13.03	8.01	6.45	5.33
K/Ba	19.71	20.41	n.a.	n.a.	18.25	20.11	20.97	18.60	23.64	n.a.	19.59	n.a.	n.a.
K/Sr	34.05	33.72	n.a.	n.a.	25.70	27.26	31.71	26.75	30.70	n.a.	29.87	n.a.	n.a.
Rb/Sr	9.79	10.62	n.a.	n.a.	7.83	8.81	9.94	8.36	8.58	n.a.	9.45	n.a.	n.a.
Hf/Th	0.99	1.03	n.a.	n.a.	1.11	1.13	1.14	1.13	1.08	n.a.	1.05	n.a.	n.a.
Hf/Ba	7.20	6.90	n.a.	n.a.	8.20	8.50	8.30	8.40	8.90	n.a.	7.60	n.a.	n.a.
Hf/Zr	24.40	22.20	n.a.	n.a.	24.10	24.40	24.60	24.00	23.80	n.a.	23.60	n.a.	n.a.
Hf/Ta	5.52	5.14	n.a.	n.a.	5.42	5.27	5.45	5.28	5.42	n.a.	5.51	n.a.	n.a.
Hf/La	18.73	20.00	n.a.	n.a.	19.70	19.74	20.25	20.52	20.53	n.a.	18.91	n.a.	n.a.
Th/Ba	7.30	6.70	n.a.	n.a.	7.40	7.50	7.20	7.40	8.20	n.a.	7.30	n.a.	n.a.
La/Sm	2.25	2.13	n.a.	n.a.	2.16	2.12	2.17	2.03	2.23	n.a.	2.19	n.a.	n.a.
La/Yb	4.66	4.21	n.a.	n.a.	4.15	3.96	4.15	4.08	4.05	n.a.	4.25	n.a.	n.a.

Geochemical unit----- Chemical type-	LH2 N2 1	MG6 N2 2	2B N2 B	5A N2 E	5A* N2 E	UNC N2 X	2C N2 B	4C N2 D	5B N2 E	3C N2 C	3C* N2 C	UNC N2 X	5C N2R2 E
K/Rb	2.49	2.14	n.a.	2.99	2.69	n.a.	2.95	n.a.	2.99	3.46	3.13	n.a.	n.a.
K/Th	32.31	23.97	27.67	24.89	23.04	n.a.	26.55	n.a.	25.99	29.29	29.69	n.a.	n.a.
K/P	4.72	4.25	9.14	11.00	11.41	5.81	8.65	5.50	11.35	8.22	7.96	5.95	9.45
K/Ba	16.11	14.15	n.a.	22.02	20.52	n.a.	21.52	n.a.	22.18	24.38	19.79	n.a.	n.a.
K/Sr	13.80	14.60	n.a.	47.05	43.38	n.a.	44.97	n.a.	49.70	30.81	26.50	n.a.	n.a.
Rb/Sr	5.54	6.82	n.a.	15.73	16.13	n.a.	15.27	n.a.	16.61	8.90	8.47	n.a.	n.a.
Hf/Th	1.50	1.36	0.83	0.77	0.76	n.a.	0.86	n.a.	0.76	1.06	1.13	n.a.	n.a.
Hf/Ba	7.50	8.00	n.a.	6.80	6.70	n.a.	7.00	n.a.	6.40	8.90	7.60	n.a.	n.a.
Hf/Zr	22.20	24.10	n.a.	25.10	25.10	n.a.	25.10	n.a.	25.10	25.20	24.20	n.a.	n.a.
Hf/Ta	6.44	5.10	7.66	5.23	5.28	n.a.	5.41	n.a.	5.30	5.00	4.97	n.a.	n.a.
Hf/La	17.76	19.04	16.33	18.02	17.21	n.a.	17.63	n.a.	17.87	19.62	19.14	n.a.	n.a.
Th/Ba	5.00	5.90	n.a.	8.80	8.90	n.a.	8.10	n.a.	8.50	8.30	6.70	n.a.	n.a.
La/Sm	2.21	2.30	2.44	2.48	2.59	n.a.	2.40	n.a.	2.50	2.10	2.11	n.a.	n.a.
La/Yb	5.34	3.91	5.24	5.46	5.34	n.a.	5.13	n.a.	5.15	4.53	3.98	n.a.	n.a.

Geochemical unit-----5C* N2R2 Chemical type-	3D R2 C	5D R2 E	5D* R2 E	5D* R2 E	UNC R2 X	2D R2 B	2D* R2 B	UNC R2 X	AS16 R2 3	2E R2N1 B	UNC* N1 X	KB5 N1 4
K/Rb	n.a.	2.52	n.a.	n.a.	n.a.	3.28	2.89	2.81	n.a.	n.a.	n.a.	3.10
K/Th	n.a.	26.39	26.71	n.a.	n.a.	27.57	27.72	24.63	24.65	28.76	n.a.	31.18
K/P	12.83	8.04	10.36	10.53	11.06	7.32	8.40	9.30	7.64	7.84	10.46	6.84
K/Ba	n.a.	18.83	n.a.	n.a.	n.a.	23.13	21.04	18.35	n.a.	n.a.	n.a.	21.00
K/Sr	n.a.	29.65	n.a.	n.a.	n.a.	47.51	45.39	48.55	n.a.	n.a.	n.a.	36.65
Rb/Sr	n.a.	11.79	n.a.	n.a.	n.a.	14.49	15.71	17.27	n.a.	n.a.	n.a.	11.84
Hf/Th	n.a.	1.08	0.86	n.a.	n.a.	0.90	0.87	0.80	0.75	1.06	n.a.	1.06
Hf/Ba	n.a.	7.70	n.a.	n.a.	n.a.	7.20	6.80	5.60	n.a.	n.a.	n.a.	7.10
Hf/Zr	n.a.	22.70	n.a.	n.a.	n.a.	25.30	25.10	25.40	n.a.	n.a.	n.a.	24.00
Hf/Ta	n.a.	5.03	4.55	n.a.	n.a.	4.84	5.42	5.47	5.52	5.66	n.a.	5.12
Hf/La	n.a.	20.33	19.05	n.a.	n.a.	18.28	18.33	17.97	17.61	21.38	n.a.	20.05
Th/Ba	n.a.	7.10	n.a.	n.a.	n.a.	8.30	8.50	7.40	n.a.	n.a.	n.a.	6.70
La/Sm	n.a.	2.10	2.29	n.a.	n.a.	2.19	2.22	2.36	2.40	1.88	n.a.	1.96
La/Yb	n.a.	4.44	5.11	n.a.	n.a.	5.13	5.12	5.04	5.25	4.07	n.a.	4.14

Geochemical unit----- Chemical type-	3E N1 C	UNC N1 X	3F N1 C	5E N1 E	AS16 N1 3	3G N1 C	5F N1 E	UNC N1 X	AS16 N1 3	5G N1 E	AS16 N1 3	5H N1 E	AS16 N1 3
K/Rb	3.18	n.a.	3.21	n.a.	2.96	n.a.	3.43	n.a.	2.89	n.a.	n.a.	n.a.	n.a.
K/Th	26.52	33.47	25.97	32.36	27.61	28.59	29.69	n.a.	29.56	n.a.	25.60	29.43	28.74
K/P	9.11	7.14	8.52	9.19	7.71	8.73	9.63	5.50	8.62	9.87	8.10	10.95	9.97
K/Ba	27.59	n.a.	17.61	n.a.	19.74	n.a.	20.14	n.a.	20.02	n.a.	n.a.	n.a.	n.a.
K/Sr	32.04	n.a.	26.87	n.a.	36.40	n.a.	40.23	n.a.	36.90	n.a.	n.a.	n.a.	n.a.
Rb/Sr	10.07	n.a.	8.38	n.a.	12.31	n.a.	11.73	n.a.	12.75	n.a.	n.a.	n.a.	n.a.
Hf/Th	1.00	1.10	1.01	1.10	0.97	1.05	1.09	n.a.	1.11	n.a.	1.08	0.98	1.01
Hf/Ba	10.40	n.a.	6.80	n.a.	6.90	n.a.	7.40	n.a.	7.50	n.a.	n.a.	n.a.	n.a.
Hf/Zr	24.90	n.a.	23.40	n.a.	23.80	n.a.	24.40	n.a.	23.80	n.a.	n.a.	n.a.	n.a.
Hf/Ta	5.23	5.70	5.31	5.94	5.46	5.74	5.71	n.a.	5.96	n.a.	5.69	5.13	5.34
Hf/La	21.75	20.00	20.75	21.03	20.19	20.71	22.22	n.a.	21.58	n.a.	21.85	21.29	21.82
Th/Ba	10.40	n.a.	6.80	n.a.	7.10	n.a.	6.80	n.a.	6.80	n.a.	n.a.	n.a.	n.a.
La/Sm	1.91	1.95	1.96	1.95	2.04	2.07	1.97	n.a.	2.02	n.a.	1.98	2.00	1.93
La/Yb	4.44	4.27	4.58	4.09	4.51	4.51	4.43	n.a.	4.36	n.a.	4.57	4.64	4.20

Table 7. Ratios of incompatible trace elements in samples of the Grande Ronde Basalt—Continued

Geochemical unit----- Chemical type-	UNC N1 X	AS16 N1 3	2F R1 B	5I R1 E	AS16 R1 3
K/Rb	2.58	2.60	3.08	n.a.	n.a.
K/Th	28.43	26.98	29.50	25.41	31.54
K/P	11.82	10.37	7.42	9.87	8.05
K/Ba	26.06	20.60	22.21	n.a.	n.a.
K/Sr	49.02	42.90	42.46	n.a.	n.a.
Rb/Sr	18.99	16.51	13.77	n.a.	n.a.
Hf/Th	0.90	0.95	1.17	0.92	1.22
Hf/Ba	8.20	7.20	8.80	n.a.	n.a.
Hf/Zr	24.10	23.90	25.70	n.a.	n.a.
Hf/Ta	5.49	5.93	6.18	5.54	5.27
Hf/La	21.05	21.21	22.40	21.71	21.89
Th/Ba	9.20	7.60	7.50	n.a.	n.a.
La/Sm	2.10	2.05	1.94	2.11	1.93
La/Yb	4.83	4.89	4.14	5.11	4.12

Table 8. Ratios of incompatible trace elements in samples of the Wanapum Basalt

[Geochemical units listed in approximate stratigraphic order, see table 1. n.a., not analyzed]

Geochemical unit----- Chemical type-	PV A	LOLO INC B	ROSALIA C	PH POWAT D	ROZA E	FS ANOM F	FS INC F	SHUMAKER G	PH SHUM G	SW LOOK H	DODGE I	ROBIN J
K/Rb	3.11	3.19	3.17	3.81	2.75	2.55	3.47	3.64	3.41	3.32	4.15	n.a.
K/Th	25.18	24.48	24.86	29.45	26.07	23.33	28.88	29.46	27.22	29.75	48.66	48.23
K/P	2.84	2.73	2.74	3.19	3.50	3.70	4.21	3.95	3.87	4.97	3.59	2.40
K/Ba	15.07	18.12	19.17	16.78	18.14	14.20	24.52	15.51	15.46	21.38	17.26	11.86
K/Sr	29.68	28.86	35.12	45.26	33.15	32.45	35.35	46.34	48.78	49.16	14.22	5.03
Rb/Sr	9.55	9.04	11.08	11.86	12.08	12.75	10.19	12.75	14.29	14.80	3.43	n.a.
Hf/Th	1.29	1.31	1.34	1.18	1.17	1.03	1.14	1.17	1.15	1.03	2.47	4.00
Hf/Ba	7.70	9.70	10.40	6.70	8.10	6.30	9.70	6.20	6.50	7.40	8.70	11.00
Hf/Zr	24.30	23.50	24.20	24.60	23.20	23.20	23.90	24.10	24.30	24.00	23.10	21.80
Hf/Ta	4.25	4.28	4.45	5.08	4.42	4.16	4.51	4.63	4.91	4.36	5.61	4.77
Hf/La	16.73	16.64	17.46	17.25	17.19	16.33	17.40	17.06	17.73	18.77	18.82	19.47
Th/Ba	6.00	7.40	7.70	5.70	7.00	6.10	8.50	5.30	5.70	7.20	3.50	2.70
La/Sm	1.95	2.16	2.49	1.93	2.39	2.11	2.54	1.98	1.94	2.03	2.21	1.72
La/Yb	4.29	4.72	4.44	4.31	4.58	4.44	4.76	4.30	4.19	4.26	3.26	2.39

Table 9. Ratios of incompatible trace elements in samples of the Saddle Mountain Basalt

[Geochemical units listed in approximate stratigraphic order, see table 1. n.a., not analyzed]

Geochemical unit----- Chemical type-	IM A	GOOSE B	INDIAN C	MARTIN D	BASIN E	BUFORD F	ELEPHANT G	POMONA H	VC WEIPE I	ESQUAT J	RBESQUAT J	FS IC K	SLIP L
K/Rb	3.36	4.46	5.19	6.14	6.48	2.70	4.32	3.00	1.53	2.76	2.74	2.87	3.02
K/Th	23.84	23.14	23.52	29.25	29.57	16.49	17.56	21.16	16.47	16.37	15.34	18.49	16.07
K/P	4.25	1.47	2.26	1.90	1.70	7.05	3.96	5.15	3.21	7.39	7.47	5.89	3.26
K/Ba	22.70	12.10	12.08	10.04	10.74	26.36	32.43	31.19	15.35	24.60	20.82	27.79	24.53
K/Sr	33.92	45.66	35.63	26.14	28.52	50.89	48.72	23.77	17.31	55.24	50.92	54.55	30.96
Rb/Sr	10.09	10.23	6.87	4.26	4.41	18.83	11.27	7.93	11.28	20.04	18.59	18.98	10.25
Hf/Th	0.93	2.47	1.91	2.81	3.01	0.75	1.08	1.27	1.33	0.64	0.65	0.74	0.94
Hf/Ba	8.90	12.90	9.80	9.60	10.90	11.90	20.00	18.70	12.40	9.70	8.80	11.20	14.30
Hf/Zr	24.50	24.60	24.70	24.80	24.10	24.50	28.00	23.60	23.40	24.60	26.10	24.60	24.40
Hf/Ta	2.76	3.50	3.51	3.55	3.45	3.46	3.68	3.94	4.05	3.42	3.48	3.01	3.55
Hf/La	13.12	14.49	14.65	16.03	15.36	16.06	19.02	19.00	19.18	14.58	15.05	15.03	15.71
Th/Ba	9.50	5.20	5.10	3.40	3.60	16.00	18.50	14.70	9.30	15.00	13.60	15.00	15.30
La/Sm	3.03	2.42	2.55	2.46	2.44	2.76	2.19	2.20	2.08	2.77	2.83	2.63	2.44
La/Yb	7.45	5.39	5.47	4.85	5.02	5.59	4.86	4.11	4.24	6.91	7.12	6.30	5.44

Geochemical unit-----LEW ORCH Chemical type-	M	SPRAGUE N	SW SPRAG N	NEW O	SW NEW O	ASOTIN P	VC ASOT P	SW HUNTZ Q	HUNTZ Q	VC LAP R	EAGLE R	WILBUR S	UMATILLA T
K/Rb	n.a.	12.04	0.00	2.55	2.67	2.23	3.82	3.17	0.00	3.19	3.98	3.37	4.92
K/Th	4.96	24.07	21.64	23.72	17.83	15.79	17.76	20.52	25.97	21.36	18.94	21.97	29.20
K/P	.59	2.09	1.81	1.62	1.17	3.90	3.64	4.41	6.82	6.12	5.33	6.56	5.91
K/Ba	0.88	12.08	10.42	10.09	7.16	13.00	13.74	20.60	0.00	18.32	16.10	19.30	6.65
K/Sr	5.78	28.55	22.11	13.18	9.30	12.85	15.34	22.48	0.00	39.99	32.25	52.75	76.70
Rb/Sr	.00	2.37	0.00	5.16	3.47	5.75	4.02	7.08	0.00	12.52	8.11	15.63	15.58
Hf/Th	.55	2.12	2.24	2.18	2.26	1.36	1.35	1.23	1.08	0.95	1.02	0.89	1.41
Hf/Ba	1.10	10.60	10.80	9.30	9.10	11.20	10.40	12.30	0.00	8.20	8.70	7.80	3.20
Hf/Zr	2.70	23.20	22.90	21.60	22.80	23.00	24.20	22.90	0.00	22.60	31.30	23.70	21.20
Hf/Ta	.17	3.87	3.51	3.18	3.47	4.28	4.68	4.04	5.83	5.37	4.35	5.57	6.86
Hf/La	4.03	14.94	15.07	15.25	14.88	17.71	19.33	18.25	15.56	14.68	16.23	13.50	21.88
Th/Ba	.40	5.00	4.80	4.30	4.00	8.20	7.70	10.00	0.00	8.60	8.50	8.80	2.30
La/Sm	.49	2.52	2.30	2.30	2.45	2.23	2.29	2.40	2.79	3.08	2.96	3.41	2.90
La/Yb	.20	5.85	5.49	5.28	4.75	4.37	4.04	5.08	6.26	6.12	5.42	6.82	6.82

Geochemical unit-----UM FRAC Chemical type-	T	PH UMTIL T	PHSOPHER U
K/Rb	5.20	4.91	5.72
K/Th	32.78	28.97	28.58
K/P	5.56	5.10	5.33
K/Ba	7.00	6.21	6.10
K/Sr	84.21	75.73	59.19
Rb/Sr	16.18	15.44	10.36
Hf/Th	1.40	1.42	1.40
Hf/Ba	3.00	3.00	3.00
Hf/Zr	22.90	20.40	21.30
Hf/Ta	7.90	6.51	6.83
Hf/La	21.89	21.62	21.84
Th/Ba	2.10	2.10	2.10
La/Sm	2.76	3.13	3.23
La/Yb	8.57	7.08	6.94

Table 10. Corrected and normalized major-oxide chemistry of naturally quenched glass from the Grande Ronde Basalt

[Geochemical units listed in approximate stratigraphic order, see table 1. Chemical type names for glasses correspond to the chemical types for whole rocks. Data are averages computed from original data given in Wright (1988)]

Geochemical unit-----	1A* GL	3A* GL	4A* GL	LH2N2 GL	MG6N2 GL	5A* GL	3C* GL	5C* GL	5D* GL	2D* GL
Chemical type-	A	C	D	1	2	E	C	E	E	B
SiO ₂	54.82	54.36	54.30	55.41	52.47	56.26	54.51	57.37	55.98	55.74
Al ₂ O ₃	13.89	13.59	13.43	14.38	14.08	13.24	13.45	13.10	12.92	12.96
FeO	11.90	12.27	12.77	10.42	12.57	12.51	12.79	12.24	12.98	12.98
MgO	4.08	4.39	4.08	5.00	5.25	3.18	3.96	2.83	3.03	2.80
CaO	8.18	8.71	8.21	9.21	9.67	7.04	7.93	6.69	6.89	6.75
Na ₂ O	3.15	3.04	3.10	3.01	2.97	3.39	3.30	3.21	3.41	3.44
K ₂ O	1.60	1.35	1.50	0.88	0.81	1.90	1.45	2.01	2.05	2.13
TiO ₂	2.04	2.03	2.26	1.41	1.87	2.16	2.26	2.22	2.38	2.77
P ₂ O ₅	0.34	0.27	0.36	0.28	0.31	0.32	0.35	0.33	0.35	0.44
Total	100.00	100.01	100.01	100.00	100.00	100.00	100.00	100.00	99.99	100.01

Table 11. Corrected and normalized major-oxide chemistry of naturally quenched glaass from the Wanapum Basalt

[Geochemical units listed in approximate stratigraphic order, see table 1. Chemical type names for glasses correspond to the chemical types for whole rocks. Data are averages computed from original data given in Wright (1988)]

Geochemical unit-----	LOLO GL	ROSAL GL	ROZA GL	FRSP GL	DODGE GL	ROBIN GL
Chemical type-	2	3	4	5	6	7
SiO ₂	50.31	50.42	51.32	52.36	52.54	49.72
Al ₂ O ₃	12.24	12.08	12.77	12.44	14.07	16.06
FeO	15.37	15.91	14.95	15.08	12.39	10.83
MgO	4.31	3.85	3.90	3.51	5.31	7.71
CaO	9.13	8.56	8.39	8.05	9.74	11.39
Na ₂ O	2.73	2.88	2.95	2.89	3.04	2.51
K ₂ O	1.19	1.35	1.44	1.56	0.76	0.44
TiO ₂	3.92	4.18	3.58	3.51	1.76	1.12
P ₂ O ₅	0.80	0.78	0.69	0.60	0.39	0.20
Total	100.00	100.01	99.99	100.00	100.00	99.98

Table 12. Corrected and normalized major-oxide chemistry of naturally quenched glass from the Saddle Mountains

[Geochemical units listed in approximate stratigraphic order, see table 1. Chemical type names for glasses correspond to the chemical types for whole rocks. Data are averages computed from original data given in Wright (1988)]

Geochemical unit-----	LM GL	GOOSE GL	INDIAN GL	MARTIN GL	BASIN GL	BUFORD GL	EM GL	POMONA GL	WEIPE GL	ESQUAT GL	SPRAGE GL	NEW GL	ASOTIN GL
Chemical type-	A	B	C	D	E	F	G	H	I	K	N	O	P
SiO ₂	51.68	48.17	49.55	48.25	47.47	55.15	51.71	52.98	52.17	54.88	49.63	49.76	53.60
Al ₂ O ₃	11.61	10.96	11.54	12.62	11.55	12.47	11.81	13.74	13.52	12.72	11.99	14.96	14.09
FeO	15.96	18.33	16.59	15.88	17.36	13.71	16.12	11.90	12.66	14.02	15.99	11.38	11.66
MgO	3.35	3.77	3.84	5.29	4.61	3.11	3.56	5.46	5.71	3.07	4.46	7.04	5.00
CaO	8.16	9.02	8.95	10.13	9.90	7.57	8.29	10.26	10.31	7.07	9.43	11.58	9.75
Na ₂ O	2.86	2.55	2.68	2.47	2.50	3.06	2.54	2.46	2.68	2.72	2.73	2.37	2.80
K ₂ O	1.92	1.34	1.39	0.81	1.00	1.84	1.43	0.85	0.58	1.92	1.10	0.44	0.70
TiO ₂	3.88	4.09	4.56	3.80	4.62	2.73	4.01	2.10	2.13	3.24	3.91	2.04	2.04
P ₂ O ₅	0.59	1.77	0.90	0.76	0.99	0.37	0.53	0.25	0.23	0.37	0.75	0.44	0.35
Total	100.01	100.00	100.00	100.01	100.00	100.01	100.00	100.00	99.99	100.01	99.99	100.01	99.99

Geochemical unit-----	LAP GL	WILBUR GL	UM GL
Chemical type-	R	S	T
SiO ₂	54.69	55.41	56.10
Al ₂ O ₃	13.48	13.45	13.23
FeO	12.20	12.26	12.41
MgO	4.07	3.30	2.39
CaO	8.28	7.53	6.05
Na ₂ O	2.69	2.83	3.21
K ₂ O	1.77	2.26	3.03
TiO ₂	2.34	2.42	2.56
P ₂ O ₅	0.48	0.53	1.02
Total	100.00	99.99	100.00

Table 13. Summary of chemical characteristics for geochemical units in the Yakima Basalt Subgroup

[Geochemical units listed in approximate stratigraphic order, see table 1; differences from most similar units indicated by: >, greater content in unit labeled in column 1; <, smaller content in unit labeled in column 1; differences based on MgO variation diagrams. LREE, light rare-earth elements; HREE, heavy rare-earth elements; Eu*, europium anomaly (+ or -) Diagnostic oxides and elements are those that stand out on the MgO variation diagrams. Sources of data for $^{87}\text{Sr}/^{86}\text{Sr}$ ratio: a, Carlson (1984a); b, Carlson and others (1981); c, Nelson (1980); d, J. Arth (unpublished data); e, McDougall (1976)]

Geochemical unit	Most similar unit (differences)		Diagnostic oxides and elements		⁸⁷ Sr/ ⁸⁶ Sr
	By major oxides	By trace elements	elements		
			High	Low	
		Saddle Mountains	Basalt		
LM	LOLO INC (>K ₂ O,Cs,Nb,Ta, Th,U,LREE; <Sc,HREE)	LOLO INC (<Cr)	Nb	Sc	.71087(a) .71076-.71094(c)
GOOSE	Unique	Unique	FeO,Hf,Ta,P ₂ O ₅ , Nb,Y,Zn,REE	SiO ₂ ,Co	.70769(d)
INDIAN	MARTIN (<MgO)	MARTIN			.70780(a),.70762(d)
MARTIN	Unique	Unique			.70764(a),.70770(b), .70785,.7076(d)
BASIN	MARTIN (<SiO ₂ ; >TiO ₂ ,P ₂ O ₅)		SiO ₂		.70782(a),.70774(d)
BUFORD	GR INC (>K ₂ O,TiO ₂ ,P ₂ O ₅)	GR INC (>Hf,Ta,Th,U, LREE,HREE; <Sc,Sr)	Th		
ELEPHANT	ROSALIA (<Na ₂ O,P ₂ O ₅)	ROSALIA (>Co,Hf,Nb,Cu,Ni; <Ba,Rb,Sr,HREE	FeO,TiO ₂	Ba,Rb	.70762(b) .70807,.70772(c) .70788-.70829(e)
POMONA VC WEIPE	DODGE (<Na ₂ O; >TiO ₂ ,LREE) ROBIN (<MgO)	DODGE (<Cr,Sr,Sc; >Ta,Hf) ROBIN (<Sr,Sc; >Ta,Th, Hf,LREE)		Ba,Cr	.70750(b),.70780(c) .70761-.70784(e)
ESQUAT RBESQUAT	FS IC	FS IC (>Th)	Th, Ta		.71317(a),.71448(b) .71471,.71449(e)
FS IC	FS INC (<P ₂ O ₅)	FS INC (>Co,Nb)	Ta		.71310(e)
SLIP	GR INC,(<Na ₂ O,Sc; >TiO ₂ ,P ₂ O ₅ ,LREE)	GR INC (>Co,Hf,Ta,Th; <Cr,<Sr)		Ba,Cr	
LEW ORCH SPRAGUE SW SPRAG NEW SW NEW	LOLO INC (>MgO; <K ₂ O)	LOLO INC (<Cr,Rb,Sr; >Ba,Hf,Nb,Ta,Zr,LREE, HREE,-Eu*)	FeO, TiO ₂ ,P ₂ O ₅ , Nb,Ta,Sc,REE		
ASOTIN VC ASOT	ROBIN (>SiO ₂ ; <TiO ₂ ,)	ROBIN (>Hf,Nb,Sc,LREE; <Rb,Th)		Sc	.70755(a)
SW HUNTZ HUNTZ	POMONA (<CaO; >K ₂ O)	POMONA (>Ba,Cr,Th, LREE; <Cs,Sc)		Sc	.70884(a)
VC LAP EAGLE	DODGE (<CaO,Na ₂ O; >K ₂ O)	DODGE (>Ba,Nb,Rb,Ta, Th,LREE,HREE,Ni; <Cr, <Hf<Sc,<Zr,<Cu)		Sc	
WILBUR	GR INC (<Na ₂ O; >K ₂ O,P ₂ O ₅)	GR INC (>Ba,Hf,Ta,Th, Zr,LREE,HREE; <Sr,Sc)		Sc	.71047,.71063(a)
UMATILLA UM FRAC PH UMTIL	SHUMAKER (<FeO; >K ₂ O)	SHUMAKER (>Ba,Co,Hf, Ta,Th,Zr,LREE; <Cs,Sr, Sc,HREE)	Ba,Hf,Zr,Eu	Co,Cr	.70892(b) .70935(e)
PH SOPHER	PH UMTIL (<TiO ₂ ,P ₂ O ₅)	PH UMTIL (<Sc,Rb)			

Table 13. Summary of chemical characteristics for geochemical units in the Yakima Basalt Subgroup—*Continued*

Geochemical unit	Most similar unit (differences)		Diagnostic oxides and elements		⁸⁷ Sr/ ⁸⁶ Sr
	By major oxides	By trace elements			
			High	Low	
Wanapum Basalt					
PV	Unique		CaO,Cr,Sc	FeO,K ₂ O,Co	.70521(a)
LOLO INC	LM (see above)	ROZA (>Cr)		Sc	.70506(a),.70439(c)
ROSALIA	LOLO INC (<MgO; >TiO ₂)		TiO ₂	Cr	.70507(a), .70538,.70558(e)
	ELEPHANT (>P ₂ O ₅)	ELEPHANT (>Ba,Rb,Sc; <Nb,Ta)			
PH POWAT	SHUMAKER (>P ₂ O ₅)	SHUMAKER (<Sc; >REE)		Cr	
ROZA	FS INC (>MgO)	FS INC			.70517(b),.70498(c) .70545(e)
FS ANOM	FS INC (<FeO; >CaO,TiO ₂ ,P ₂ O ₅)	FS INC (>Ba,Ta,Hf,Th,Zn, Sc,REE)	Co		
FS INC	ROZA (<MgO)	ROZA			.70510,.70520(b) .70517(a),.70530 .70559(e)
SHUMAKER PH SHUM	GR INC (<SiO ₂ ,K ₂ O; >P ₂ O ₅)	GR INC (>Ba,Hf,Nb,Ta,Y,Sc, HREE; <Cr,Cs,Rb,Th)		Cr	.70481(a)
SW LOOK	SHUMAKER (<P ₂ O ₅)	SHUMAKER (<Ba,Hf,Nb,Sc; >HREE)			.70546(a)
DODGE	GR INC	GR INC (>Cr,Sc; <Th)			.70435(a)
ROBIN	ASOTIN (see above)		MgO	Rb,REE,Cr	.70438(a)
Grande Ronde Basalt					
GR INC			SiO ₂ ,Cs,Rb	TiO ₂ ,P ₂ O ₅ , Ta,REE	.70489-.70504(a) .70438-.70507(b) .70440-.70606(e) .70454-.70552(c)

Table 14. Relation between stratigraphic position and chemistry, additional flows for which chemical data are available, Columbia River Plateau

[MP, magnetic polarity as determined in the field using a fluxgate magnetometer: N, normal; R, reversed, T, transitional]

Formation				
Member				
Flow	Age	MP	Geochemical	
Informal name	(Ma)		unit	
SADDLE MOUNTAINS BASALT				
Unclassified flows in Idaho		N	VC CRAIG ¹	
[Clearwater Embayment; Craigmont,		N	VC SWAMP ¹	
Swamp Creek, Icicle Flat, and		N	VC IFLAT ¹	
Grangeville Members of Camp		R	VC GRNGE ¹	
(1981)]				
WANAPUM BASALT				
Basalt of Feary Creek ³ (Camp, 1981)		N	VC FEARY ¹	
Basalt of Potlatch ⁴ (Camp, 1981)		N	VC POT ²	
UNCLASSIFIED				
Local flows in NE Oregon (basalts				
of Powder River of Hooper(1984);	11.6	N	WT NEPH ²	
mapping of W.T. Taubeneck, D.A.	13.8-9.5	N	WT ANDES ²	
Swanson, T.L. Wright, and P.R.		N	WT JONES ²	
Hooper)		N	WT SUGMT ²	
		N	WT SPRMT ²	
		N	WT WBRMT ²	
		N	PH EDEN ¹	
		N	WT OBHTI ¹	
		N	WT OBLTI ¹	

¹ Chemistry falls within the fields defined for the Yakima Basalt Subgroup

² Chemistry falls outside of the fields defined for the Yakima Basalt Subgroup

³ Included in Frenchman Springs Member

⁴ Included in Onaway Member of Camp (1981)

Table 15. Average chemical composition of additional analyzed flows of table 14

[Chemical types listed in approximate stratigraphic order, see table 14. Trace elements followed by X analyzed by X-ray fluorescence; all others analyzed by instrumental neutron activation. n.a., not analyzed Data from which these average compositions are calculated are given in Wright (1988)]

Geochemical unit-----	WT NEPH	WT ANDES	WT JONES	WT SUGMT	WT SPRMT	WT WBRMT	PH EDEN	WT OBHTI	WT OBLTI	VC CRAIG	VC SWAMP	VC IFLAT	VC GRNGE
Chemical type- A	B	C	D	E	F	G	H	I	J	K	L	M	
Major oxides (weight percent)													
SiO ₂	43.71	59.31	62.89	53.68	51.14	53.56	52.07	50.80	50.69	52.77	51.74	52.06	53.38
Al ₂ O ₃	15.26	18.01	19.32	17.09	16.10	17.45	14.28	16.76	17.41	14.73	14.91	16.01	16.21
FeO	15.58	5.97	3.36	7.99	9.18	7.63	13.96	9.87	9.37	13.82	12.93	11.54	8.78
MgO	7.97	3.12	2.43	3.78	4.98	4.43	3.52	8.86	8.11	3.83	4.89	5.36	6.62
CaO	9.00	6.31	6.56	8.39	9.32	8.69	7.44	9.34	10.78	7.12	8.53	9.56	10.57
Na ₂ O	4.32	4.63	4.29	4.33	4.24	4.66	2.58	2.47	2.36	2.73	2.96	3.12	2.55
K ₂ O	0.71	1.78	1.09	2.39	2.29	1.70	2.24	0.47	0.37	1.74	1.15	0.78	0.68
TiO ₂	3.06	0.97	0.45	1.85	2.08	1.56	2.75	1.44	1.02	2.96	2.73	1.46	1.36
P ₂ O ₅	0.84	0.49	0.25	1.05	1.16	0.83	1.39	0.42	0.28	0.53	0.39	0.37	0.21
MnO	0.22	0.13	0.11	0.16	0.15	0.15	0.25	0.18	0.17	0.19	0.22	0.26	0.17
Total	100.67	100.72	100.75	100.71	100.64	100.66	100.48	100.61	100.56	100.42	100.45	100.52	100.53
Trace elements (parts per million)													
Ba	294	848	592	1234	1088	776	1788	334	272	715	549	414	311
BaX	198	821	n.a.	n.a.	711	n.a.	1774	428	245	n.a.	n.a.	n.a.	n.a.
Co	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.	46.5	n.a.	n.a.	n.a.	n.a.	n.a.
Cr	85.1	46.8	24.1	33.0	60.0	104.4	34.8	363.5	32.0	7.4	38.1	82.2	133.0
Cs	n.a.	0.42	0.82	0.60	0.36	0.58	0.81	0.20	n.a.	0.68	0.67	n.a.	n.a.
Hf	1.73	3.05	2.05	4.03	3.11	3.00	10.80	2.18	1.70	6.08	5.41	2.60	3.21
NbX	6.0	n.a.	n.a.	n.a.	20.0	n.a.	34.0	14.0	n.a.	n.a.	n.a.	n.a.	n.a.
Rb	n.a.	16.0	17.5	20.8	13.0	14.0	52.8	15.7	23.9	41.7	23.7	n.a.	24.0
RbX	n.a.	12.5	n.a.	n.a.	36.0	n.a.	49.0	10.0	8.0	n.a.	n.a.	n.a.	n.a.
SrX	980	1012	n.a.	n.a.	2443	n.a.	238	479	467	n.a.	n.a.	n.a.	n.a.
Ta	0.45	0.57	0.46	0.60	0.52	0.45	2.37	0.51	0.37	1.67	1.53	0.52	0.97
Th	n.a.	1.75	2.05	3.07	1.83	2.58	7.73	0.85	0.61	7.15	3.15	1.05	2.85
U	n.a.	0.75	0.73	0.83	0.63	0.65	1.67	0.37	n.a.	1.47	1.03	0.49	0.82
YX	13.5	20.5	n.a.	n.a.	n.a.	n.a.	59.0	27.0	21.0	n.a.	n.a.	n.a.	n.a.
Zn	133	86	65	121	138	110	152	95	88	149	160	156	123
Zr	n.a.	142	117	163	135	173	487	98	81	243	420	160	147
ZrX	67	125	n.a.	n.a.	122	n.a.	480	136	81	n.a.	n.a.	n.a.	n.a.
Sc	25.8	13.5	8.6	16.1	15.9	16.6	27.1	28.1	32.6	26.1	33.8	39.5	33.0
La	16.7	30.5	14.5	52.3	46.7	37.9	58.7	13.2	8.4	42.9	32.3	13.3	19.9
Ce	44.3	51.1	25.8	112.8	105.0	80.4	116.8	27.2	18.4	81.5	56.4	26.7	36.7
Nd	32.4	28.3	12.5	62.2	57.6	43.2	63.5	17.2	12.5	44.9	34.5	19.5	19.2
Sm	6.9	5.3	2.3	8.8	8.8	7.0	13.7	3.8	2.9	8.8	7.8	4.5	4.7
Eu	2.11	1.52	0.75	2.47	2.33	1.86	4.42	1.26	0.95	2.41	2.24	1.32	1.25
Gd	4.50	4.50	2.20	6.40	5.80	5.30	14.10	3.70	2.70	8.90	8.00	4.70	4.70
Tb	0.83	0.60	0.28	0.70	0.68	0.58	2.08	0.69	0.57	1.25	1.36	0.84	0.72
Yb	1.12	1.76	0.90	1.32	1.10	1.22	5.79	2.22	2.03	3.99	4.15	3.34	2.50
Lu	0.14	0.25	0.12	0.18	0.17	0.19	0.83	0.33	0.33	0.67	0.74	0.67	0.46
Cu	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.
Ni	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.
Normalized major-oxide contents (weight percent)													
SiO ₂	43.42	58.89	62.42	53.30	50.81	53.21	51.82	50.49	50.41	52.55	51.51	51.79	53.10
Al ₂ O ₃	15.16	17.88	19.18	16.97	16.00	17.34	14.21	16.66	17.31	14.67	14.84	15.93	16.12
FeO	15.48	5.93	3.33	7.93	9.12	7.58	13.89	9.81	9.32	13.76	12.87	11.48	8.73
MgO	7.92	3.10	2.41	3.75	4.95	4.40	3.50	8.81	8.06	3.81	4.87	5.33	6.59
CaO	8.94	6.26	6.51	8.33	9.26	8.63	7.40	9.28	10.72	7.09	8.49	9.51	10.51
Na ₂ O	4.29	4.60	4.26	4.30	4.21	4.63	2.57	2.46	2.35	2.72	2.95	3.10	2.54
K ₂ O	0.71	1.77	1.08	2.37	2.28	1.69	2.23	0.47	0.37	1.73	1.14	0.78	0.68
TiO ₂	3.04	0.96	0.45	1.84	2.07	1.55	2.74	1.43	1.01	2.95	2.72	1.45	1.35
P ₂ O ₅	0.83	0.49	0.25	1.04	1.15	0.82	1.38	0.42	0.28	0.53	0.39	0.37	0.21
MnO	0.22	0.13	0.11	0.16	0.15	0.15	0.25	0.18	0.17	0.19	0.22	0.26	0.17
Total	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00

Table 15. Average chemical composition of additional analyzed flows of table 14—*Continued*

Geochemical unit-----	VC FEARY	VC POT	JOSEPH	IMNAHA
Chemical type- N	O	‡	*	
Major oxides (weight percent)				
SiO ₂	51.70	48.41	55.40	47.80
Al ₂ O ₃	14.41	16.82	15.05	15.32
FeO	14.12	12.96	12.43	10.61
MgO	4.08	5.04	3.24	12.02
CaO	7.66	7.45	6.73	10.19
Na ₂ O	3.00	3.52	2.82	2.14
K ₂ O	1.60	1.69	1.72	0.22
TiO ₂	3.25	3.84	2.36	1.25
P ₂ O ₅	0.40	0.72	0.47	0.09
MnO	0.18	0.18	0.20	0.18
Total	100.40	100.63	100.42	99.82
Trace elements (parts per million)				
Ba	550	367	857	140
BaX	521	360	n.a.	89
Co	n.a.	42.9	n.a.	49.1
Cr	20.7	8.9	6.5	861.5
Cs	0.83	0.37	1.27	n.a.
Hf	5.38	6.84	4.85	1.73
NbX	24.0	42.0	n.a.	10.0
Rb	42.0	29.8	43.5	n.a.
RbX	45.0	26.0	n.a.	n.a.
SrX	245	540	n.a.	384
Ta	1.70	3.61	0.90	0.26
Th	7.10	3.13	5.39	0.29
U	1.76	1.21	1.64	n.a.
YX	45.0	27.0	n.a.	17.0
Zn	152	129	134	89
Zr	237	281	227	n.a.
ZrX	190	298	n.a.	82
Sc	29.1	19.1	30.6	26.9
La	35.9	37.4	26.9	5.1
Ce	64.2	74.8	52.3	12.9
Nd	36.2	41.3	30.1	10.5
Sm	7.5	9.2	7.5	2.9
Eu	2.11	2.69	1.96	0.93
Gd	8.3	7.8	7.5	2.9
Tb	1.09	1.19	1.07	0.43
Yb	3.73	2.29	3.83	1.69
Lu	0.66	0.39	0.58	0.27
Cu	n.a.	n.a.	n.a.	n.a.
Ni	n.a.	n.a.	n.a.	n.a.
Normalized major-oxide contents (weight percent)				
SiO ₂	51.49	48.11	55.17	47.89
Al ₂ O ₃	14.35	16.71	14.99	15.35
FeO	14.06	12.88	12.38	10.63
MgO	4.06	5.01	3.23	12.04
CaO	7.63	7.40	6.70	10.21
Na ₂ O	2.99	3.50	2.81	2.14
K ₂ O	1.59	1.68	1.71	0.22
TiO ₂	3.24	3.82	2.35	1.25
P ₂ O ₅	0.40	0.72	0.47	0.09
MnO	0.18	0.18	0.20	0.18
Total	100.00	100.00	100.00	100.00

Table 16. Ratios of incompatible trace elements in additional analyzed flows of table 14

[Geochemical units listed in approximate stratigraphic order, see table 14. n.a., not analyzed]

Geochemical unit-----	WT NEPH A	WT ANDES B	WT JONES C	WT SUGMT D	WT SPRMT E	WT WBRMT F	PH EDEN G	WT OBHTI H	WT OBLTI I	VC CRAIG J	VC SWAMP K	VC IFLAT L	VC GRNGE M
Chemical type-													
K/Rb	n.a.	11.82	n.a.	n.a.	5.28	n.a.	3.79	3.90	3.84	n.a.	n.a.	n.a.	n.a.
K/Th	n.a.	84.44	44.14	64.63	99.99	54.70	24.06	45.90	50.35	20.20	30.31	61.67	19.81
K/P	1.61	6.91	8.29	4.33	3.75	3.89	3.06	2.13	2.51	6.24	5.61	4.01	6.16
K/Ba	29.77	18.00	n.a.	n.a.	26.74	n.a.	10.48	9.12	12.54	n.a.	n.a.	n.a.	n.a.
K/Sr	6.01	14.60	n.a.	n.a.	7.78	n.a.	78.13	8.15	6.58	n.a.	n.a.	n.a.	n.a.
Rb/Sr	n.a.	1.24	n.a.	n.a.	1.47	n.a.	20.59	2.09	1.71	n.a.	n.a.	n.a.	n.a.
Hf/Th	n.a.	1.74	1.00	1.31	1.70	1.16	1.40	2.56	2.79	0.85	1.72	2.48	1.13
Hf/Ba	8.70	3.70	n.a.	n.a.	4.40	n.a.	6.10	5.10	6.90	n.a.	n.a.	n.a.	n.a.
Hf/Zr	25.80	24.40	n.a.	n.a.	25.50	n.a.	22.50	16.00	21.00	n.a.	n.a.	n.a.	n.a.
Hf/Ta	3.84	5.35	4.46	6.72	5.98	6.67	4.56	4.27	4.59	3.64	3.54	5.00	3.31
Hf/La	10.36	10.00	14.14	7.71	6.66	7.92	18.40	16.52	20.24	14.17	16.75	19.55	16.13
Th/Ba	n.a.	2.10	n.a.	n.a.	2.60	n.a.	4.40	2.00	2.50	n.a.	n.a.	n.a.	n.a.
La/Sm	1.48	3.51	3.84	3.62	3.23	3.30	2.61	2.12	1.77	2.97	2.52	1.80	2.58
La/Yb	9.85	11.44	10.64	26.16	28.03	20.51	6.69	3.93	2.73	7.10	5.14	2.63	5.26

Geochemical unit-----	VC FEARY N	VC POT O	JOSEPH #	IMNAHA *
Chemical type-				
K/Rb	2.95	5.40	n.a.	n.a.
K/Th	18.71	44.82	26.34	62.98
K/P	7.61	4.46	6.92	4.65
K/Ba	25.49	38.97	n.a.	20.52
K/Sr	54.21	25.98	n.a.	4.76
Rb/Sr	18.37	4.81	n.a.	n.a.
Hf/Th	0.76	2.19	0.90	5.97
Hf/Ba	10.30	19.00	n.a.	19.40
Hf/Zr	28.30	23.00	n.a.	21.10
Hf/Ta	3.16	1.89	5.39	6.65
Hf/La	14.99	18.29	18.03	33.92
Th/Ba	13.60	8.70	n.a.	3.30
La/Sm	2.92	2.48	2.19	1.07
La/Yb	6.36	10.78	4.64	1.99

Table 17. Summary of chemical characteristics of additional flows for which chemical data are available, Columbia River Plateau

[Chemical analyses on which this table is based are given in table 15; geochemical units listed in approximate stratigraphic order, see table 14. The chemistries of these flows are compared with chemistries of basalts erupted in Saddle Mountains, Wanapum, or Grande Ronde time, using MgO variation diagrams (Fig. 4). Differences from most similar units indicated by: >, greater content in unit labeled in column 1; <, smaller content in unit labeled in column 1; differences based on MgO variation diagrams. LREE, light rare-earth elements; HREE, heavy rare-earth elements; Eu*, europium anomaly (+ or -). Diagnostic oxides and elements are those that stand out on the MgO variation diagrams. Sources of data for $^{87}\text{Sr}/^{86}\text{Sr}$ ratio: a, Carlson (1984a); b, Carlson and others (1981); c, Nelson (1980); d, J. Arth (unpublished data); e, McDougall (1976). The column labeled 'comment' gives a qualitative assessment of chemical classification and possible petrologic connections to units closely related in time. See text for discussion of specific examples]

Geochemical unit	Most similar unit (differences)		Diagnostic oxides and elements		⁸⁷ Sr/ ⁸⁶ Sr	comment
	By major oxides	By trace elements	elements			
			High	Low		
WT NEPH	unique	unique	Na ₂ O,Sr	SiO ₂ ,Cr,Nb, Rb (n.d.)	0.70343(b)	The only nepheline-normative basalt (basanite) in the Columbia Plateau.
WT ANDES	unique	unique	SiO ₂ ,Sr Na ₂ O	Sc	0.70354(b)	The only true andesites in the Columbia River Plateau
WT JONES	WT ANDES (<MgO)	WT ANDES	SiO ₂ ,Na ₂ O	Sc		
WT SUGMT	unique	unique	Na ₂ O,K ₂ O	Sc		More evolved andesite, related to WT ANDES
WT SPRMT	unique	unique	Na ₂ O,K ₂ O,Sr	Hf,Sc		Parental to WT ANDES?
WT WBRMT	WT SUGMT (<K ₂ O)	WT SUGMT (<Hf,REE)	Na ₂ O	Sc		Low silica 'andesite'-related to WT NEPH?
PH EDEN	UMATILLA (>MgO, P ₂ O ₅ , Y,REE)	UMATILLA (<Ba; >Hf, Nb,Ta, Y,REE)	'FeO',P ₂ O ₅ Ba,Hf,Rb,Ta Th,Zr,REE	SiO ₂ ,Al ₂ O ₃ , Na ₂ O		
WT OBLTI	WT OBHTI (>TiO ₂)					Olivine basalt
WT OBLTI	ROBIN (>P ₂ O ₅)	ROBIN (>Ba,Cr,Rb,Th)			0.70353(b)	Olivine basalt
VC CRAIG	ESQUAT	ESQUAT (<Th,<Sc; >LREE,HREE)		Sc		
VC SWAMP	unique	unique		Th		
VC IFLAT	GR INC (<SiO ₂ ; >Al ₂ O ₃ ,P ₂ O ₅)	GR INC (<Hf,Ta,Th,U, LREE;>HREE	Sc	TiO ₂ ,Hf,Ta, Th,Zr,LREE		Flatter REE pattern than other units of Saddle Mountains Basalt.
	DODGE (<MgO)	DODGE,(<Hf,LREE; >HREE)				
VC GRANGE	GR INC (>MgO,TiO ₂ ; <Na ₂ O)	GR INC (>Hf,Rb,Th,Ta, REE; <Sc)	SiO ₂	'FeO',P ₂ O ₅		Most like GR LH 2 N2 type
VC FEARY	FS IC	FS IC				Magnetic polarity opposite to that of FS IC.
VC POT	unique	unique	TiO ₂ ,Nb,Ta	SiO ₂ ,CaO, Ba,Cr,Sc		Columbia River(?) Basalt Group. Resembles ROZA in field appearance.

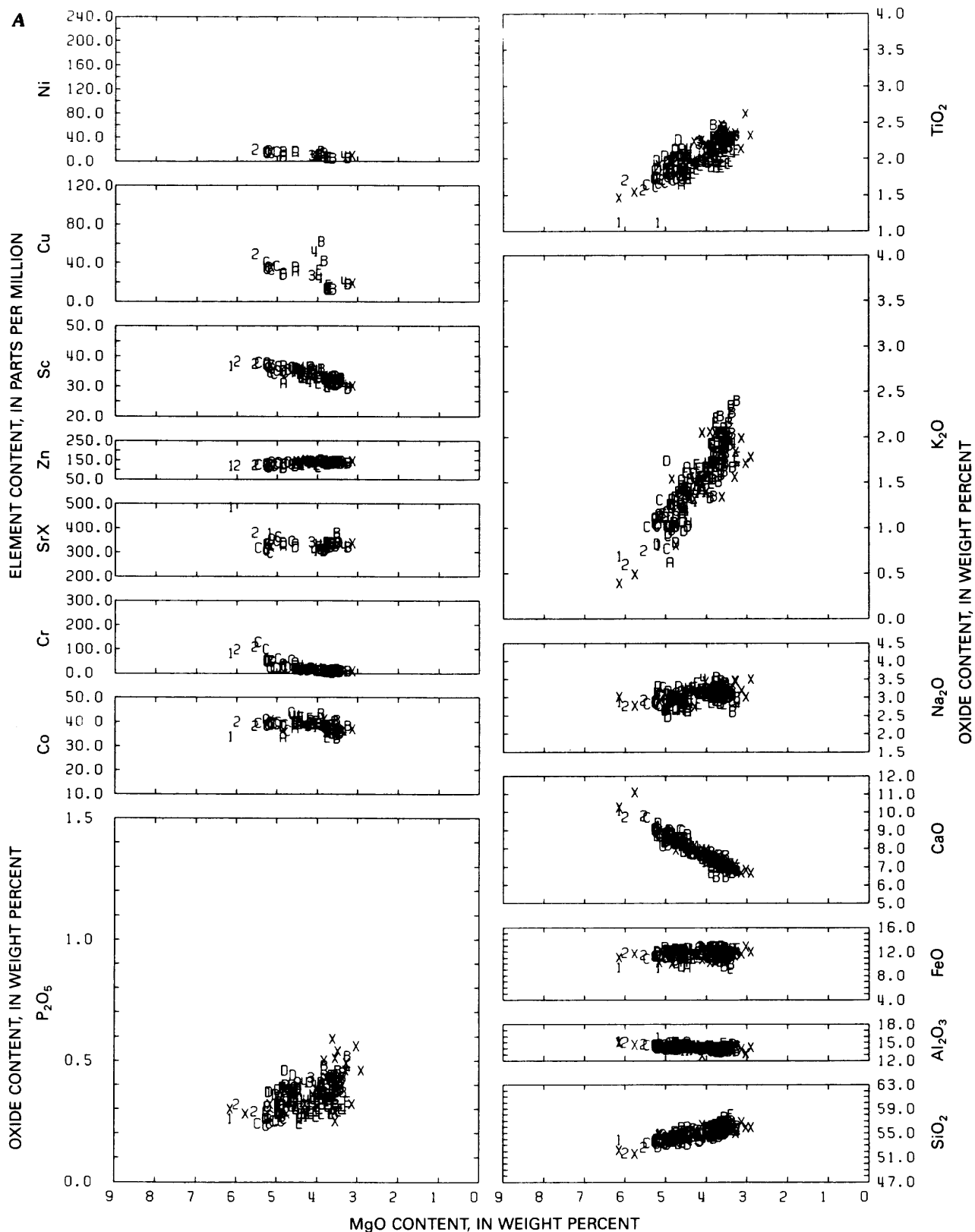
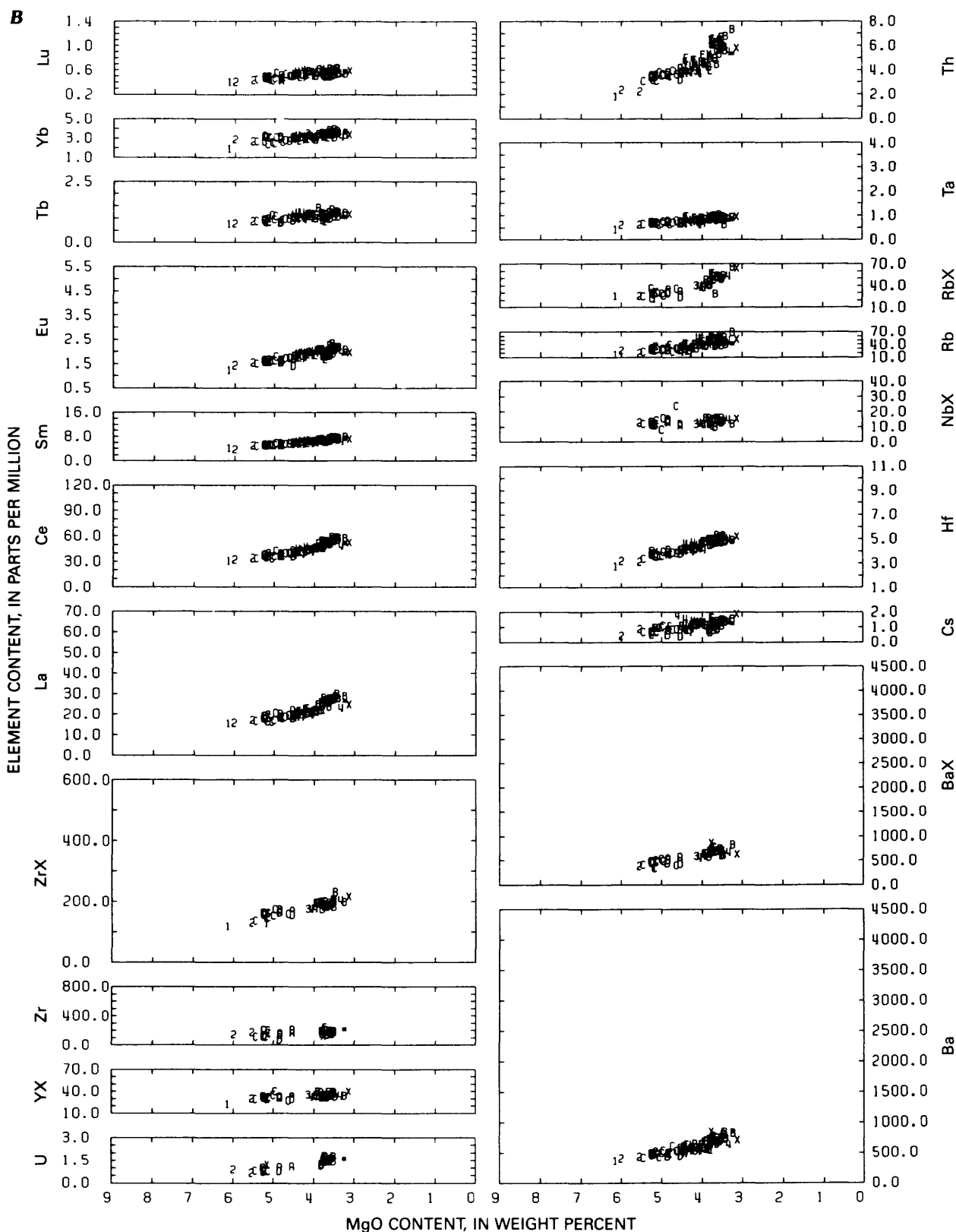


Figure 4. Major-oxide and trace-element chemistry of units of the Yakima Basalt Subgroup, presented as MgO variation diagrams. Diagrams scaled so that symbol size reflects approximately two standard deviations. Plotted data are averages of at least two samples from each geochemical unit; analyses recalculated to 100 percent dry weight with total iron expressed as FeO (see tables 3–5). Except where

otherwise noted, symbols refer to chemical types identified in table 3 (Grande Ronde Basalt), 4 (Wanapum Basalt), and 5 (Saddle Mountains Basalt). A, Grande Ronde Basalt, major oxides and compatible trace elements. Symbol # is the Joseph volcanics of Kleck (1976). B, Grande Ronde Basalt, incompatible trace elements. C, Grande Ronde Basalt, selected trace-element ratios. D, Wanapum Basalt,



major oxides and compatible trace elements. *E*, Wanapum Basalt, incompatible trace elements. *F*, Wanapum Basalt, selected trace-element ratios. *G*, Saddle Mountains Basalt, major oxides and compatible trace elements for geochemical units with relatively high SiO_2 and low FeO and TiO_2 (that is, of Grande Ronde affinity). *H*, Saddle Mountains Basalt, incompatible trace elements for units of Grande Ronde affinity. *I*, Saddle Mountains Basalt, se-

lected trace-element ratios for units of Grande Ronde affinity. *J*, Saddle Mountains Basalt, major oxides and compatible trace elements for geochemical units with relatively low SiO_2 and high FeO and TiO_2 (that is, of Wanapum affinity). *K*, Saddle Mountains Basalt, incompatible trace elements for units of Wanapum affinity. *L*, Saddle Mountains Basalt, selected trace-element ratios for units of Wanapum affinity.

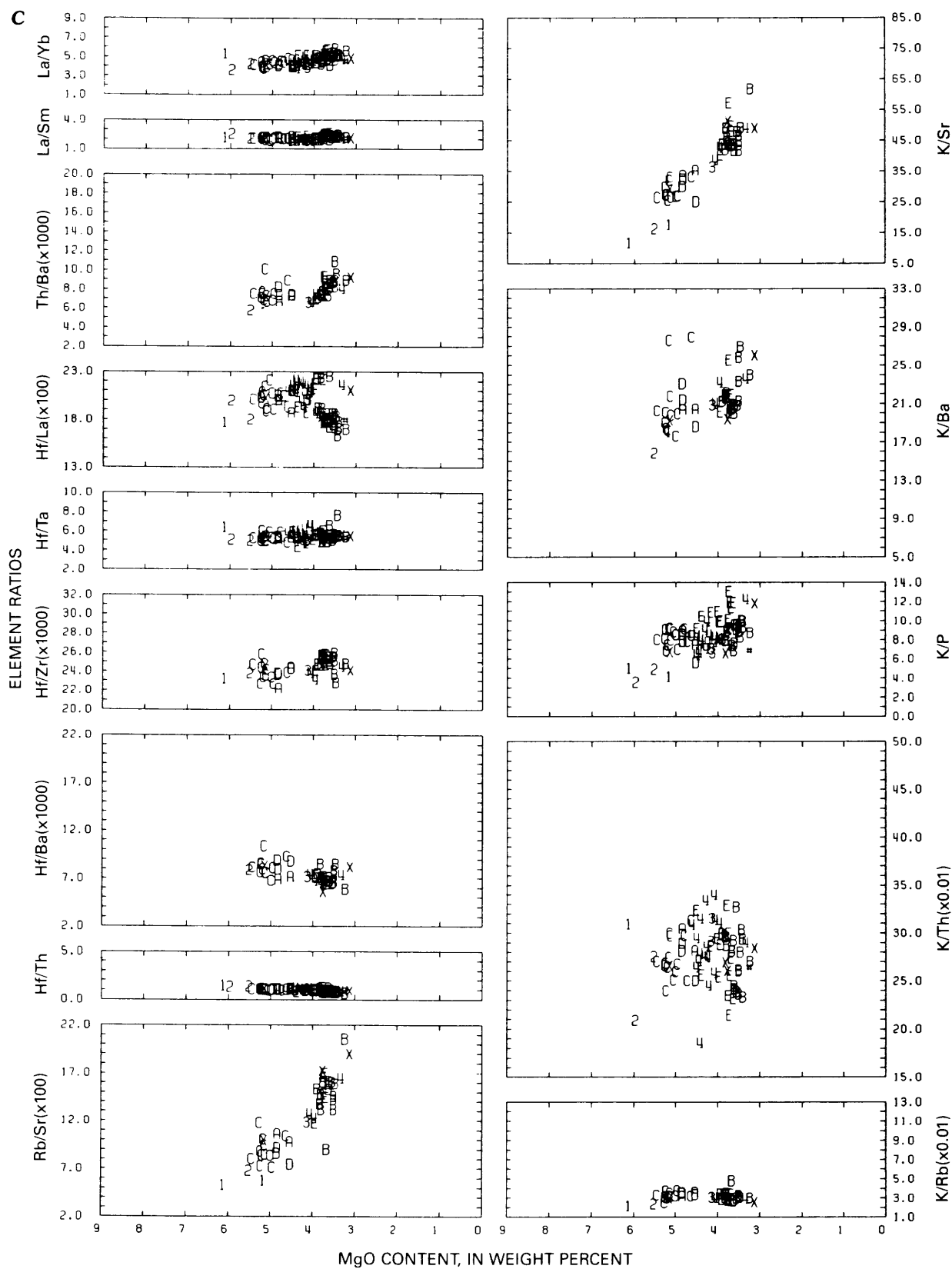


Figure 4. Continued.

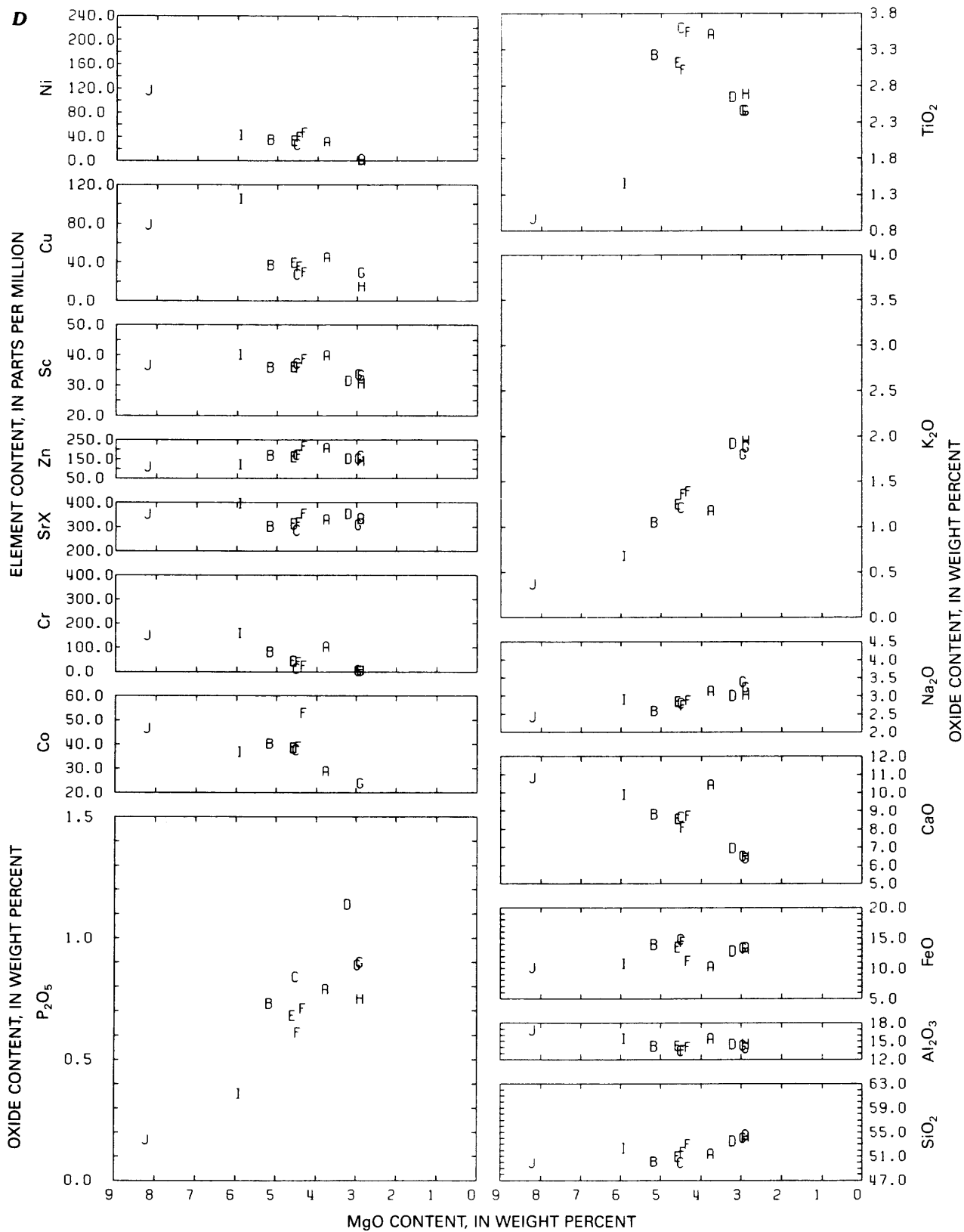


Figure 4. Continued.

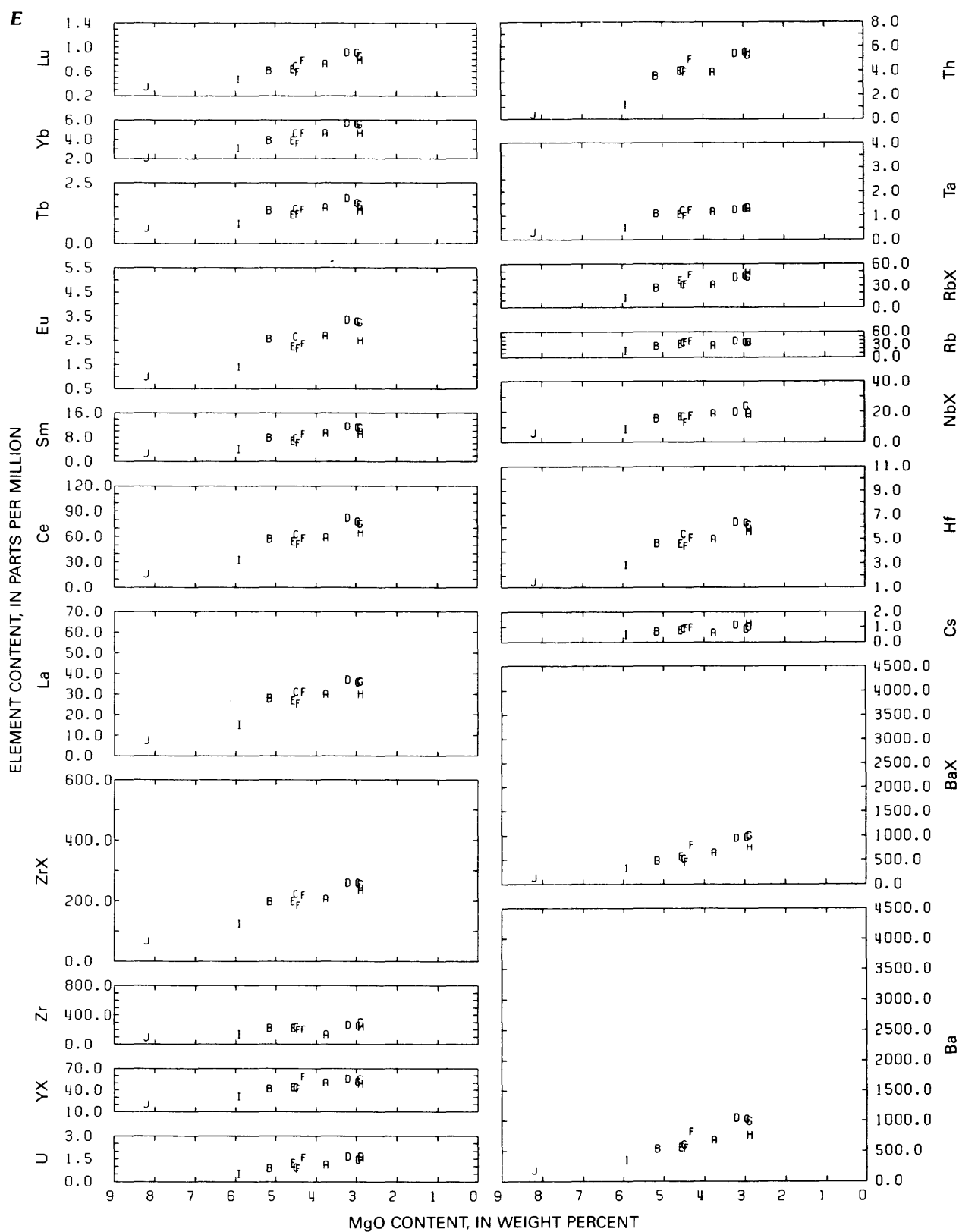


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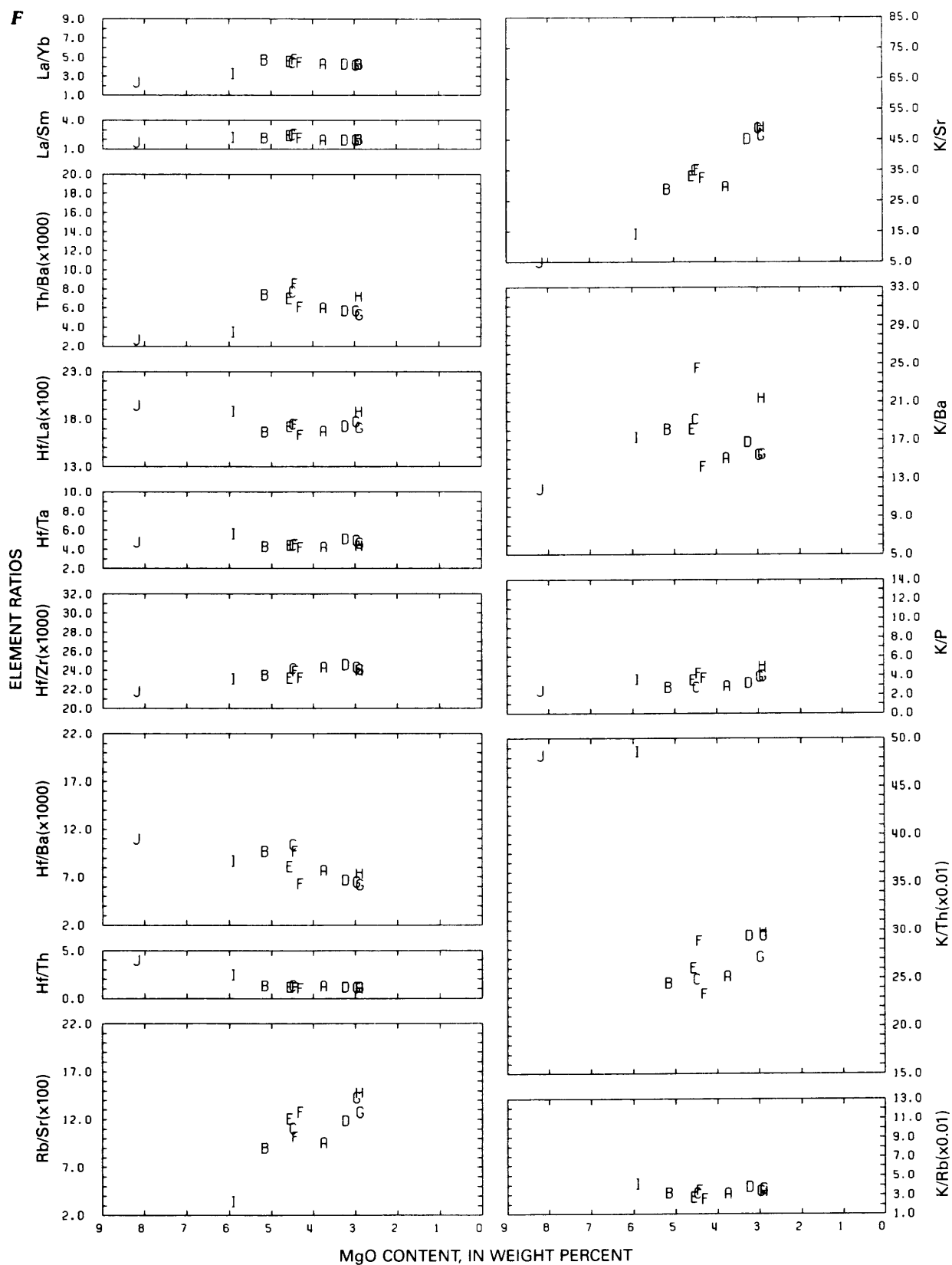


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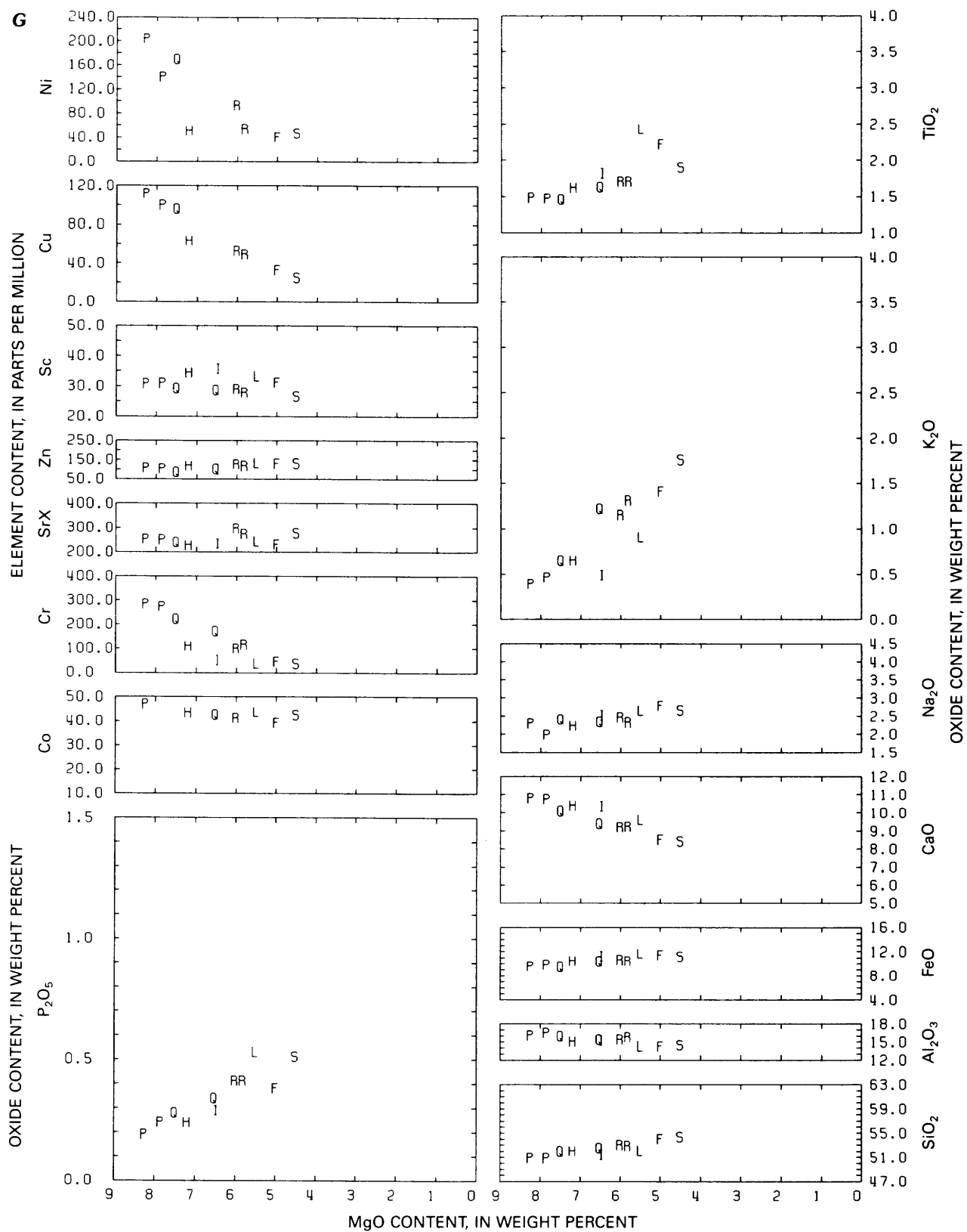


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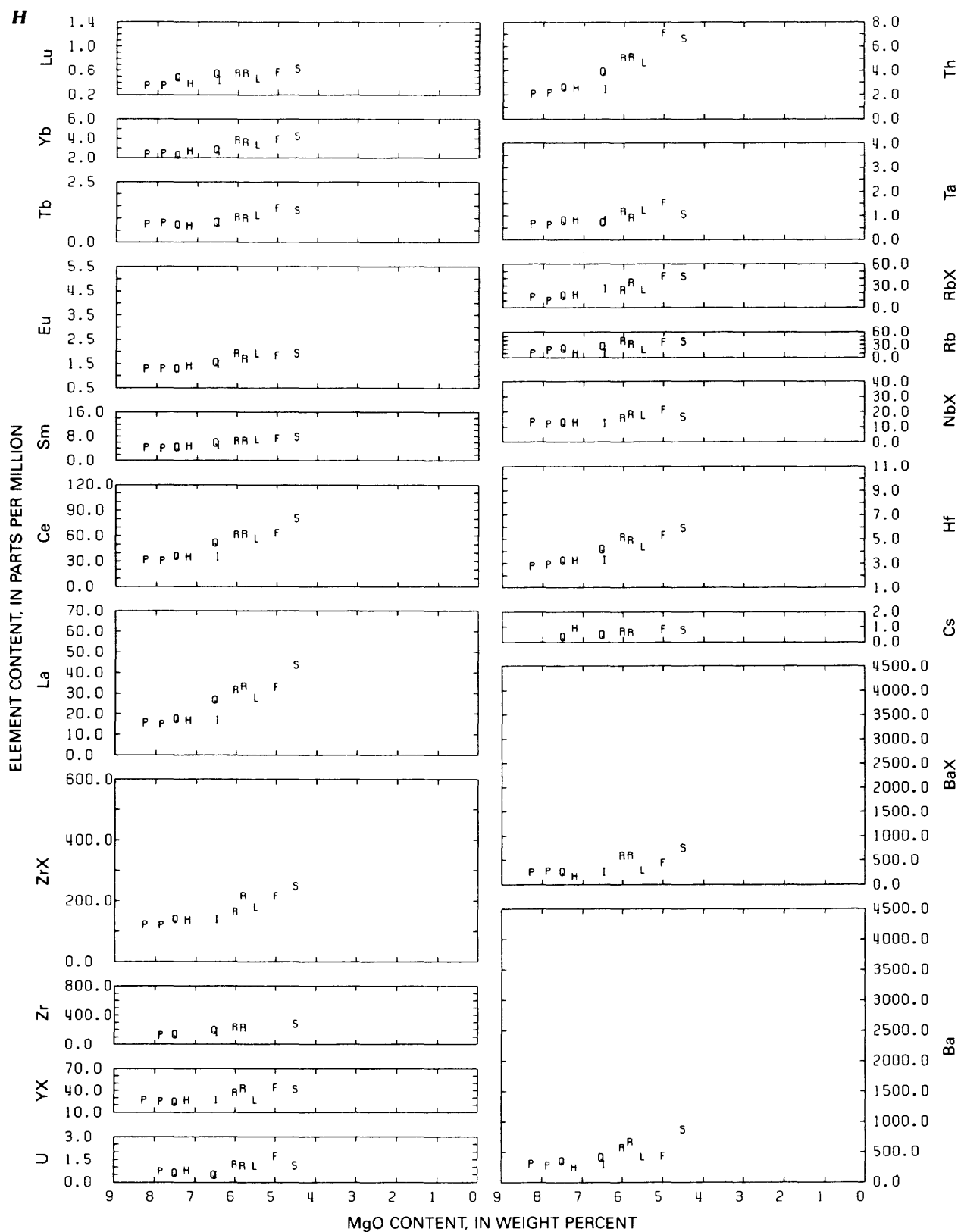


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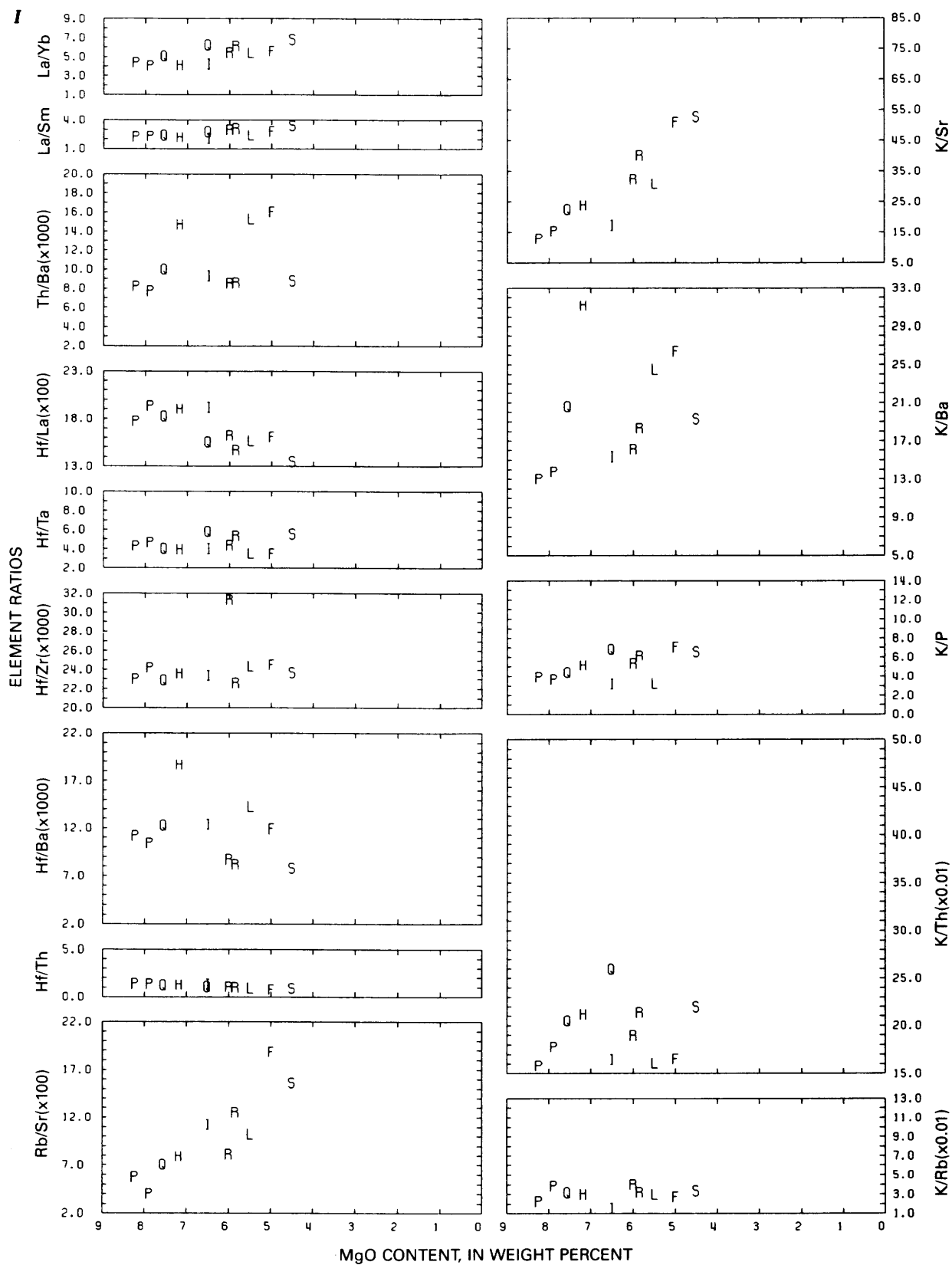


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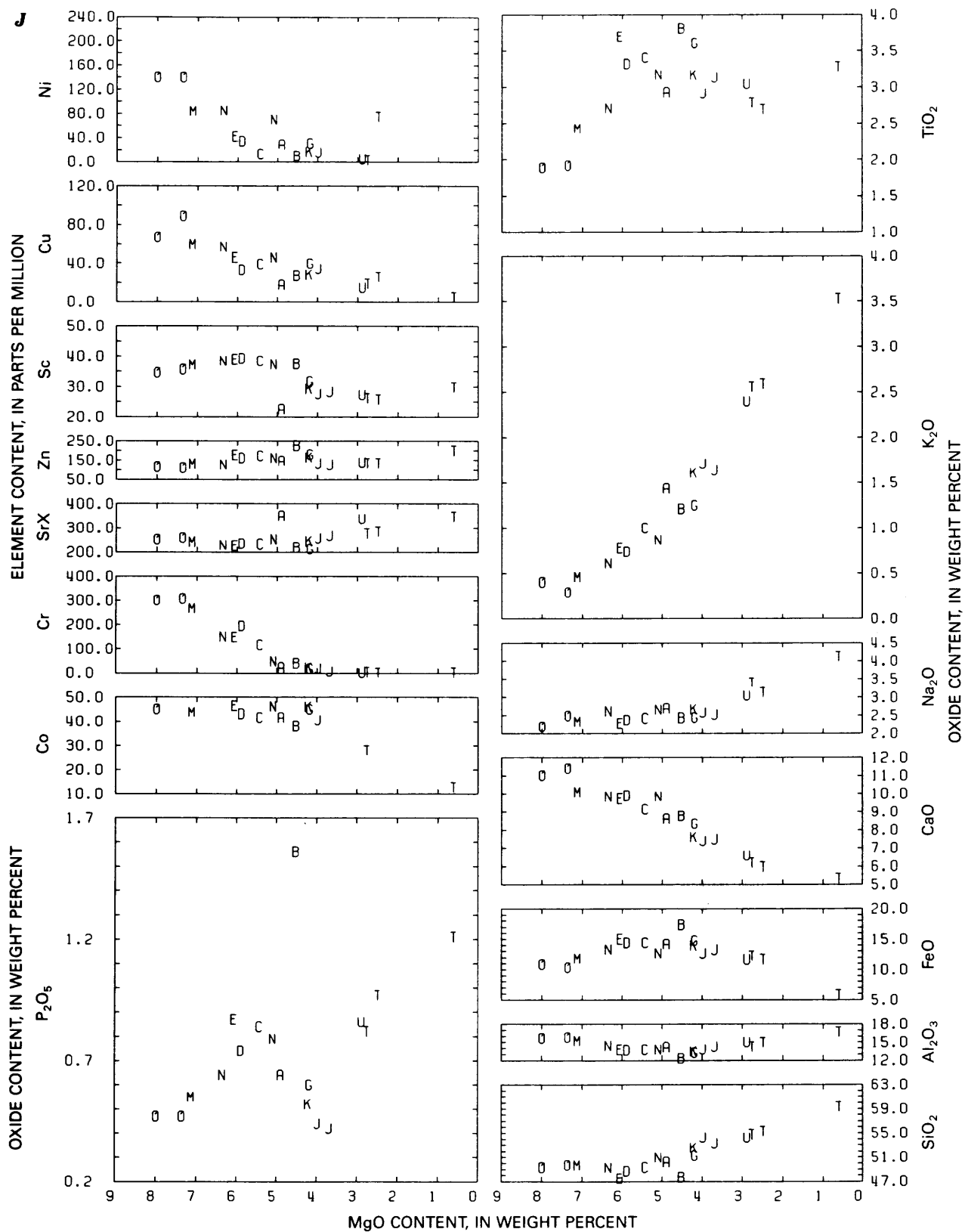


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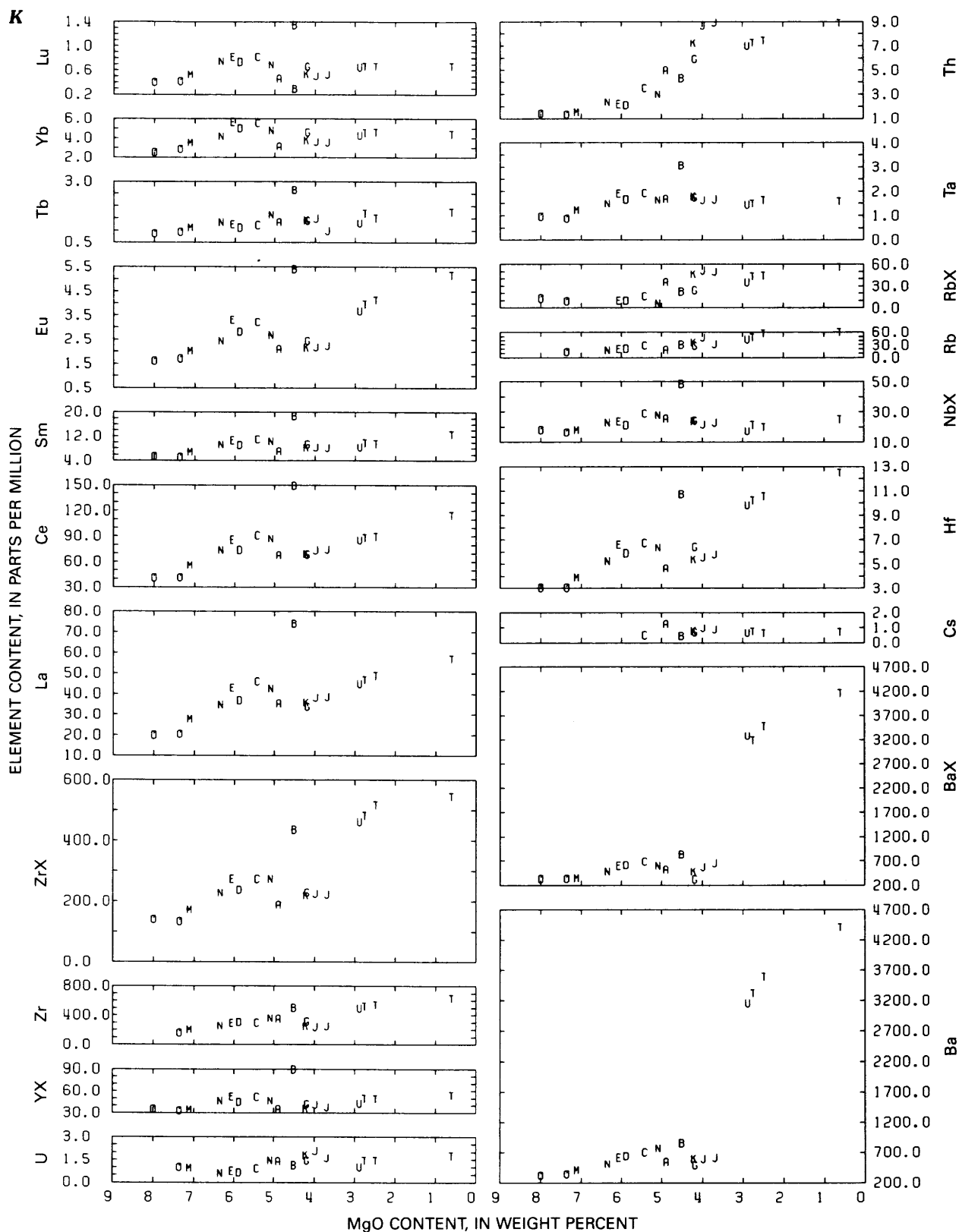


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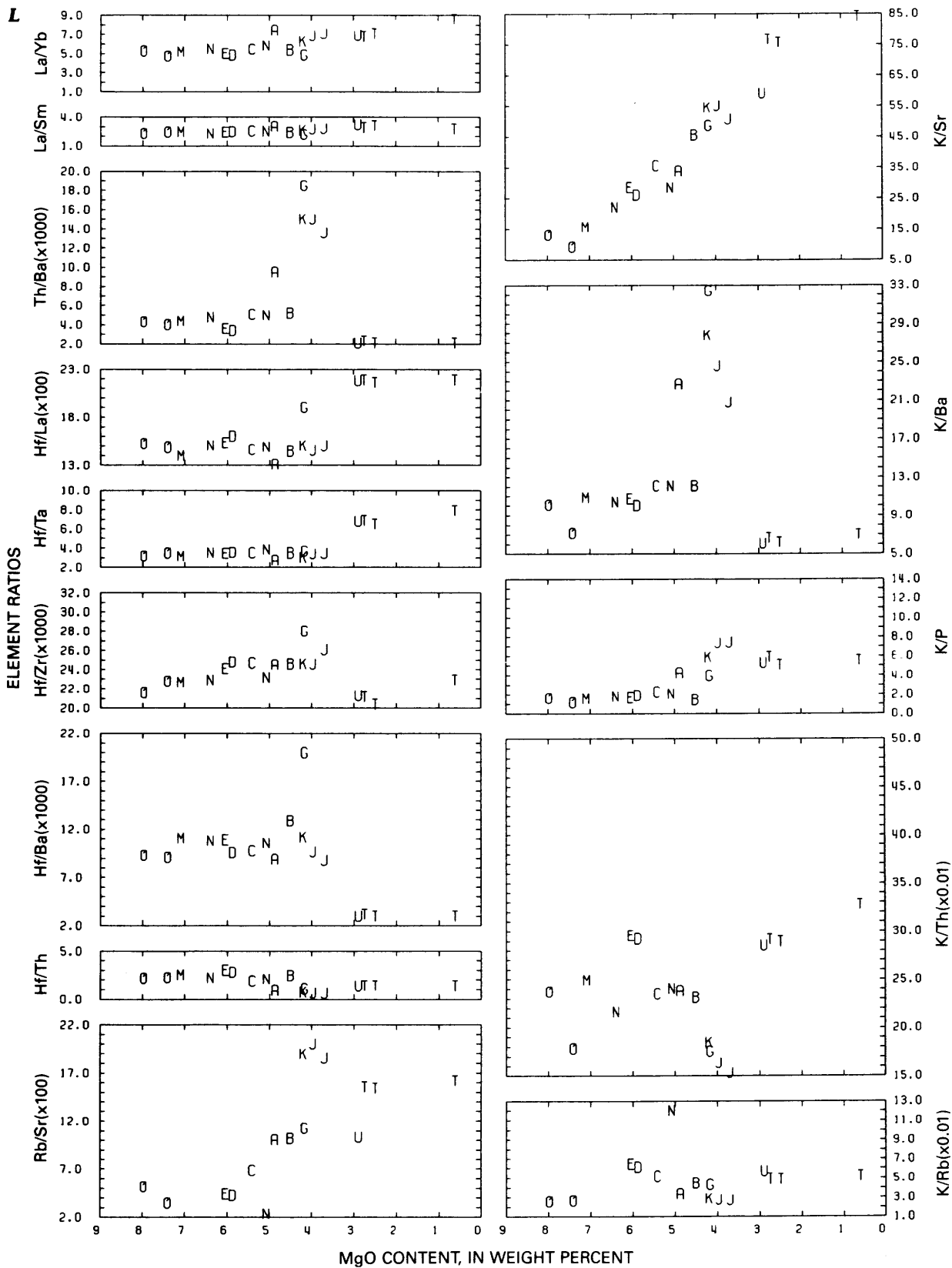


Figure 4. Continued.

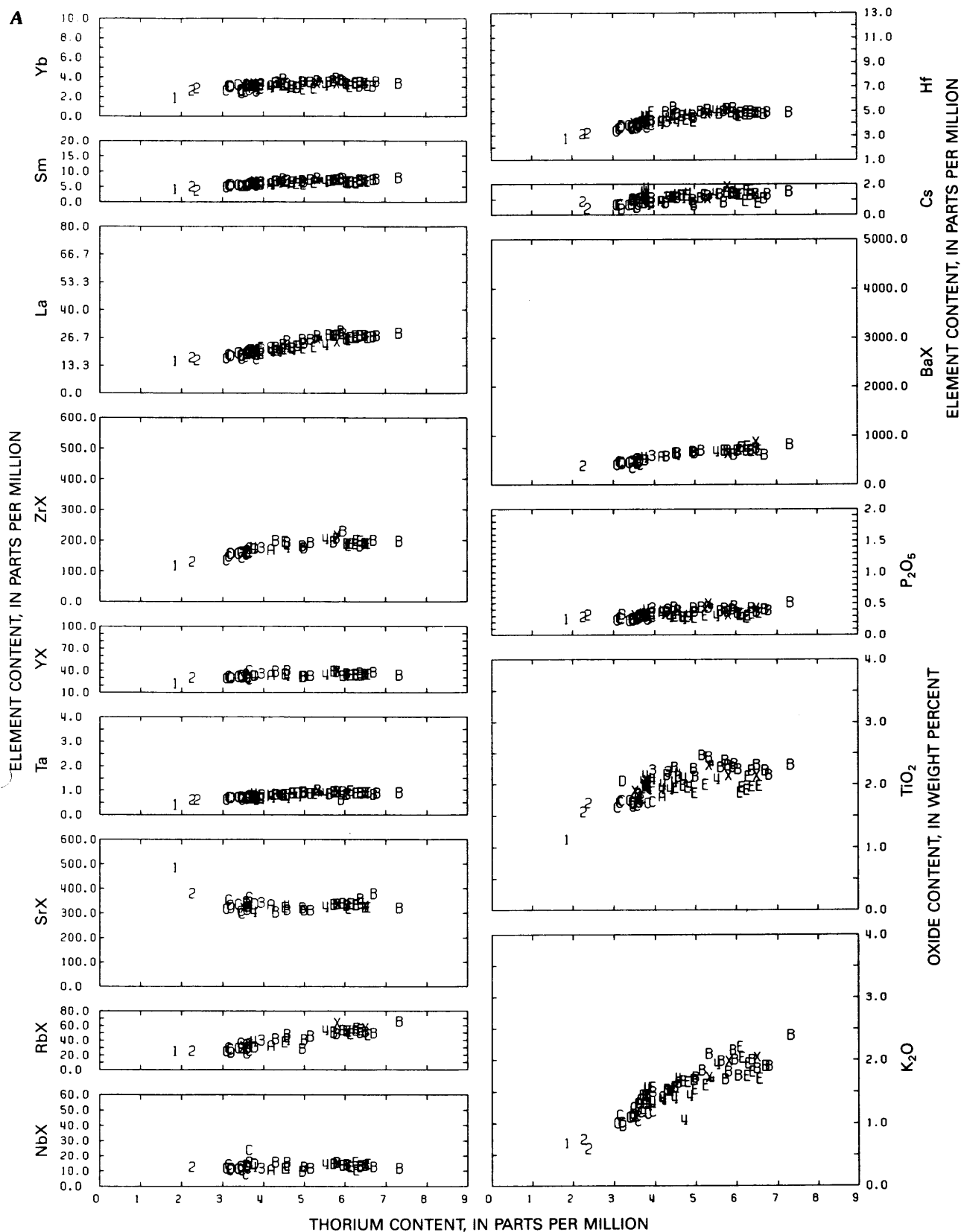
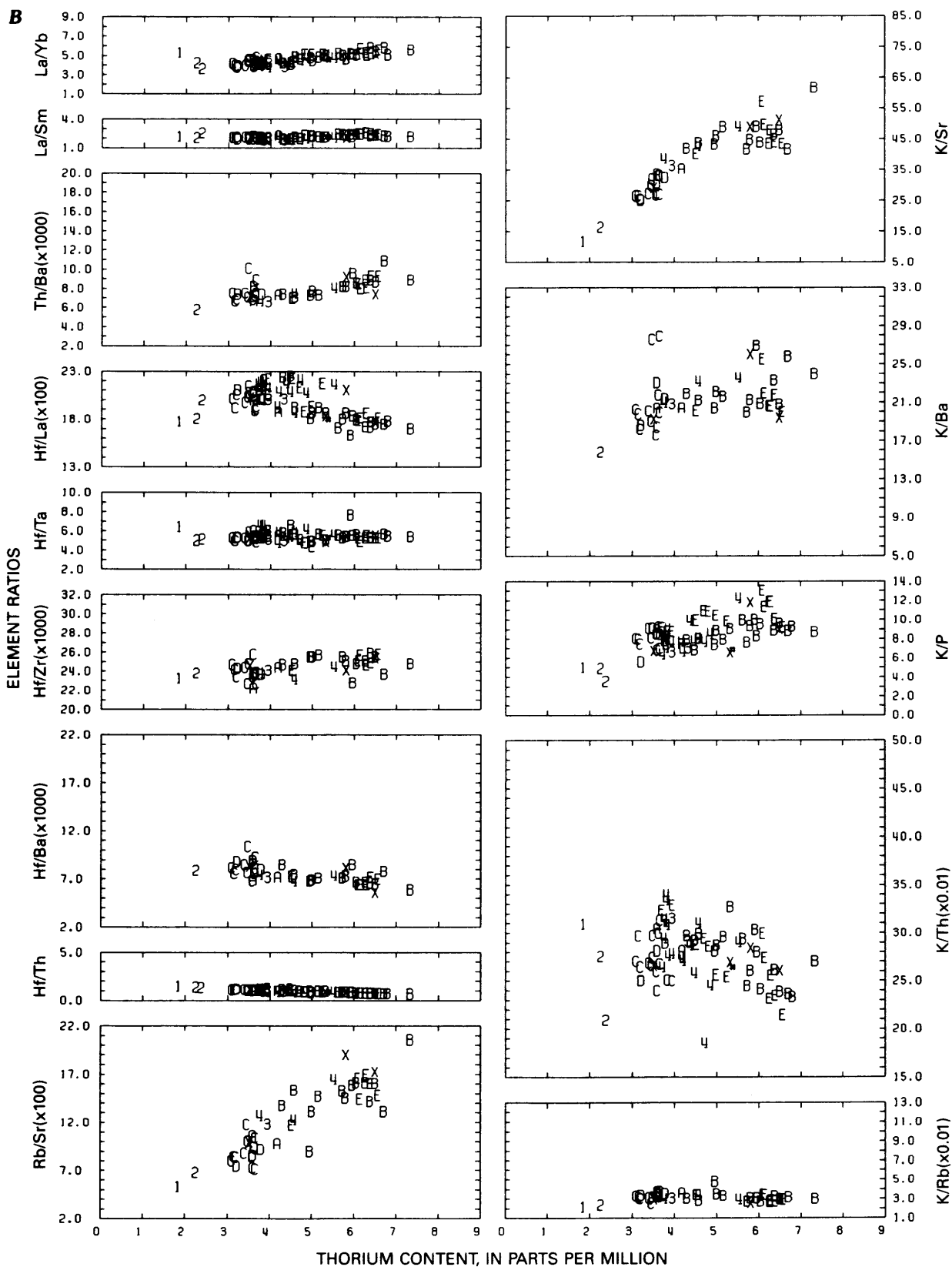


Figure 5. Chemistry of selected oxides and elements in units of the Yakima Basalt Subgroup, presented as thorium variation diagrams. Data from tables 3–5; symbols as in figure 4. *A*, Grande Ronde Basalt, minor and trace incom-

patible oxides and elements. *B*, Grande Ronde Basalt, trace-element ratios. *C*, Wanapum Basalt, minor and trace incompatible oxides and elements. *D*, Wanapum Basalt, trace-element ratios. *E*, Saddle Mountains Basalt (of Grande



Ronde affinity), minor and trace incompatible oxides and elements. *F*, Saddle Mountains Basalt (of Grande Ronde affinity), trace-element ratios. *G*, Saddle Moun-

tains Basalt (of Wanapum affinity), minor and trace incompatible oxides and elements. *H*, Saddle Mountains Basalt (of Wanapum affinity), trace-element ratios.

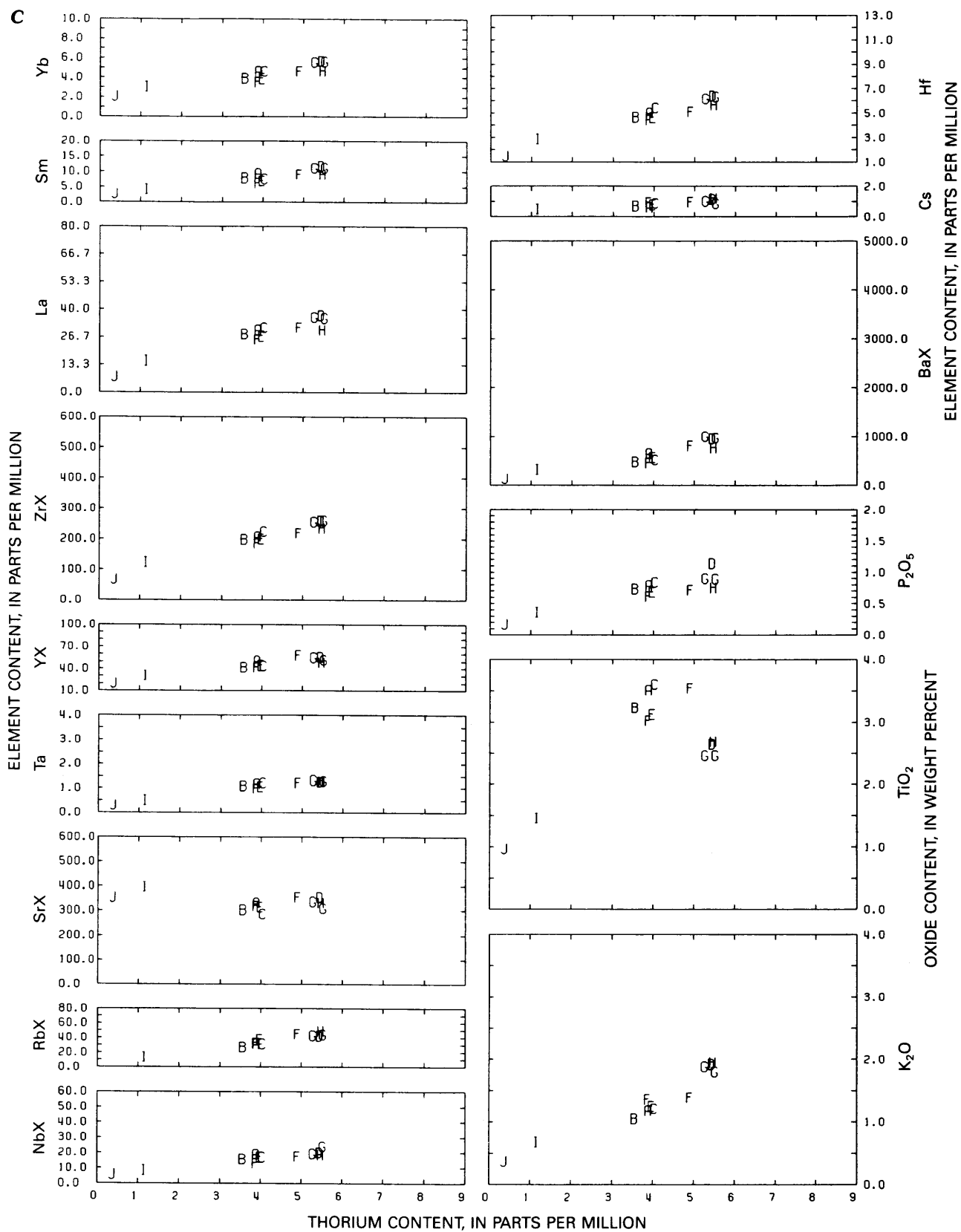


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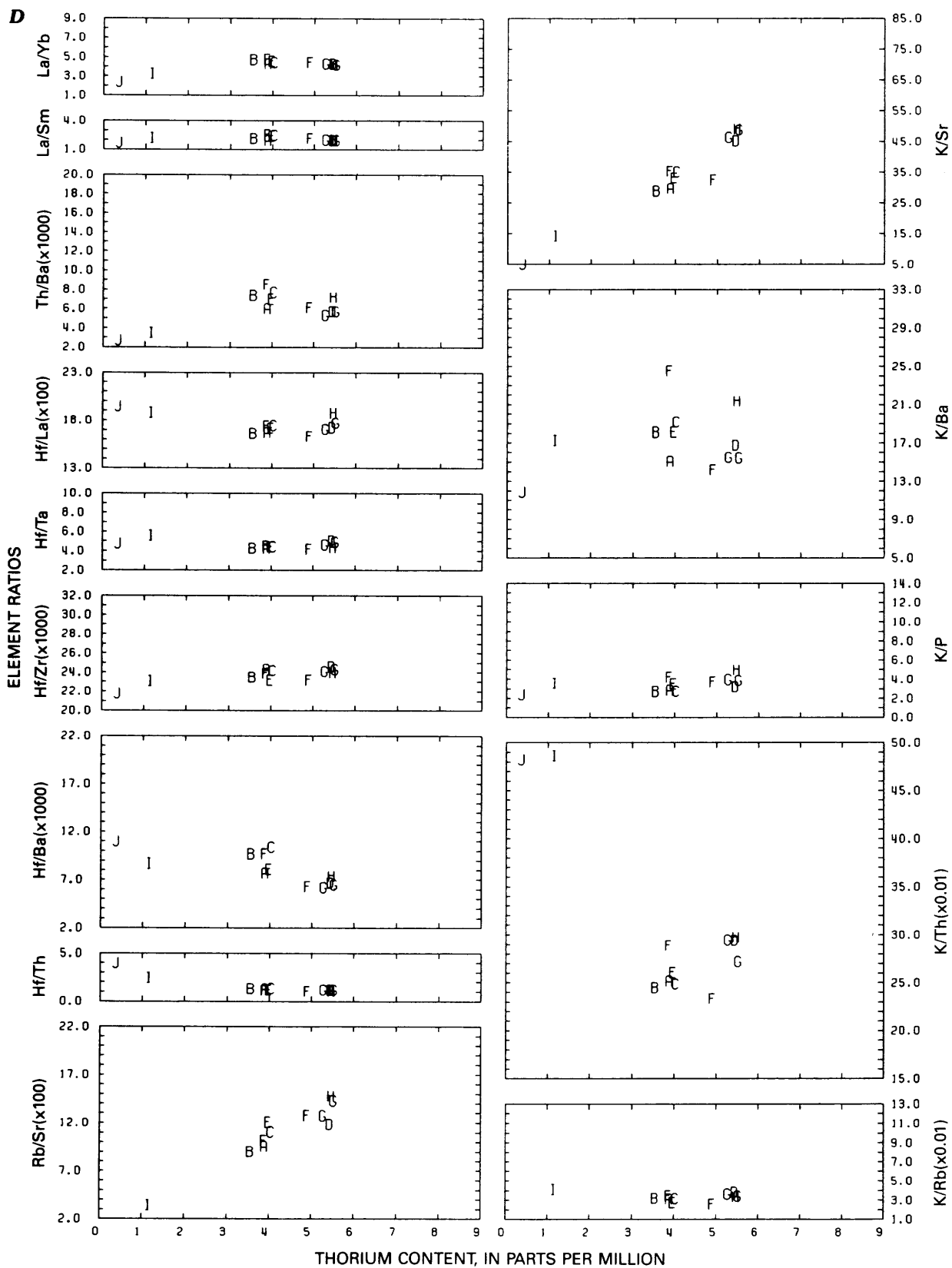


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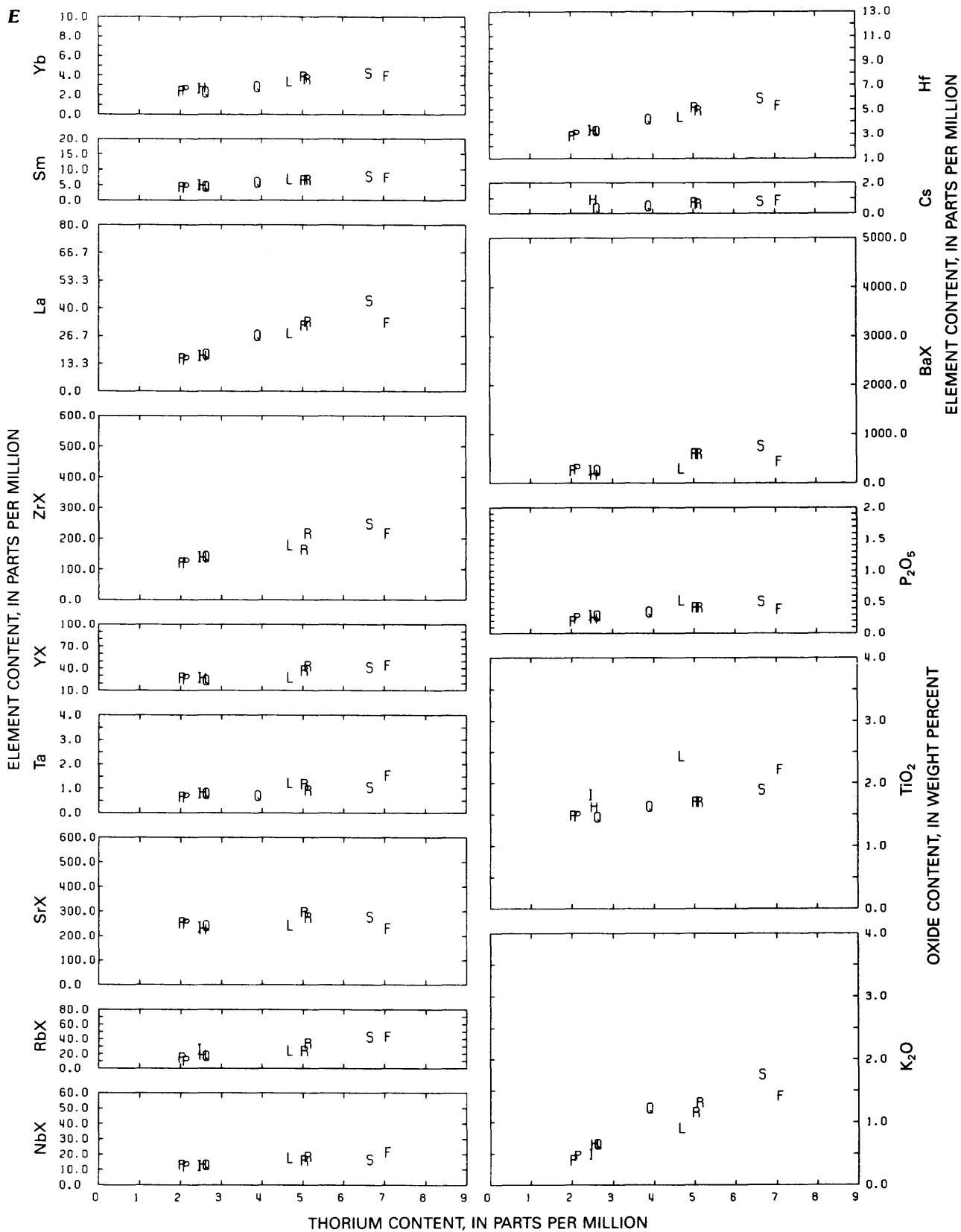


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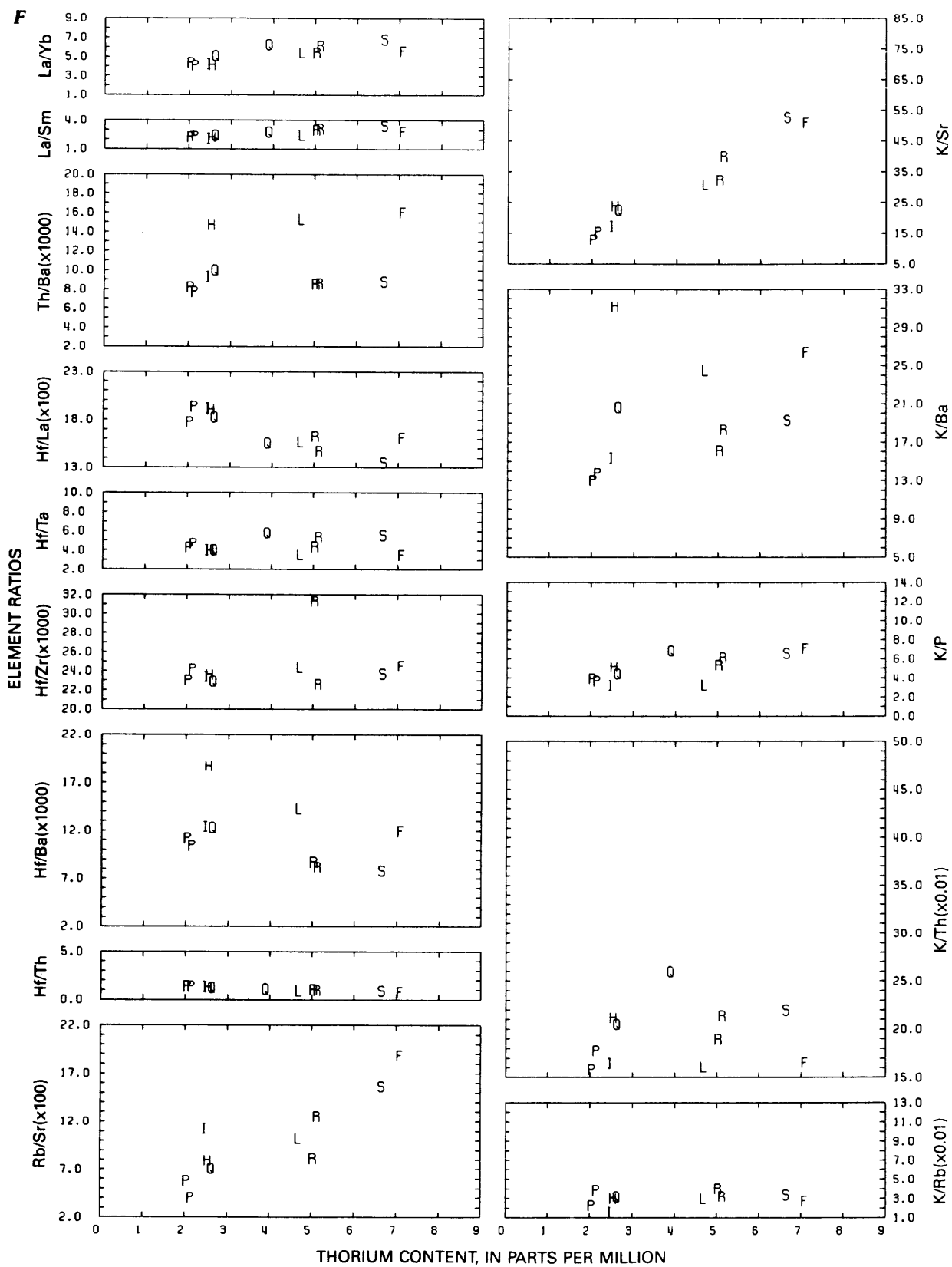


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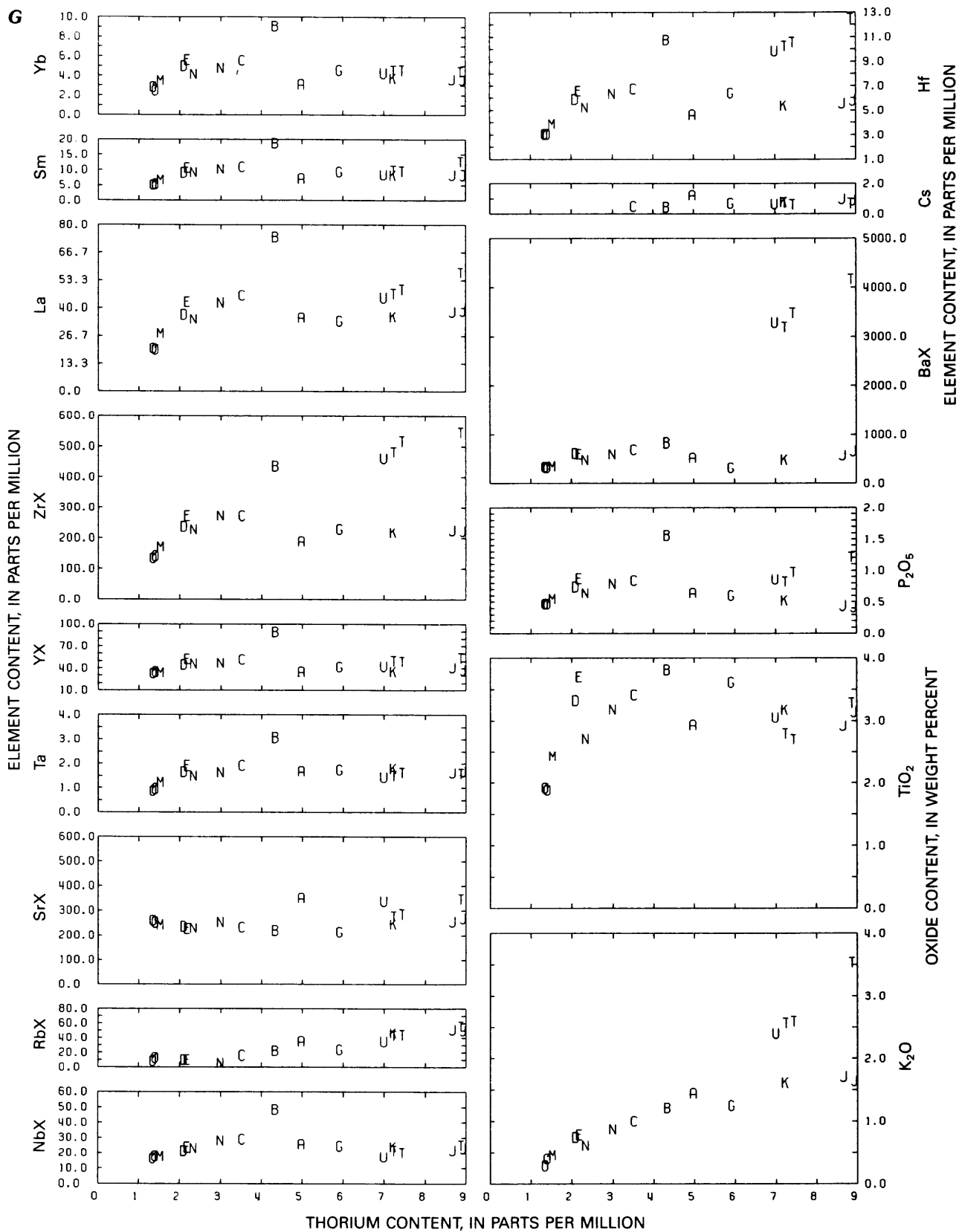


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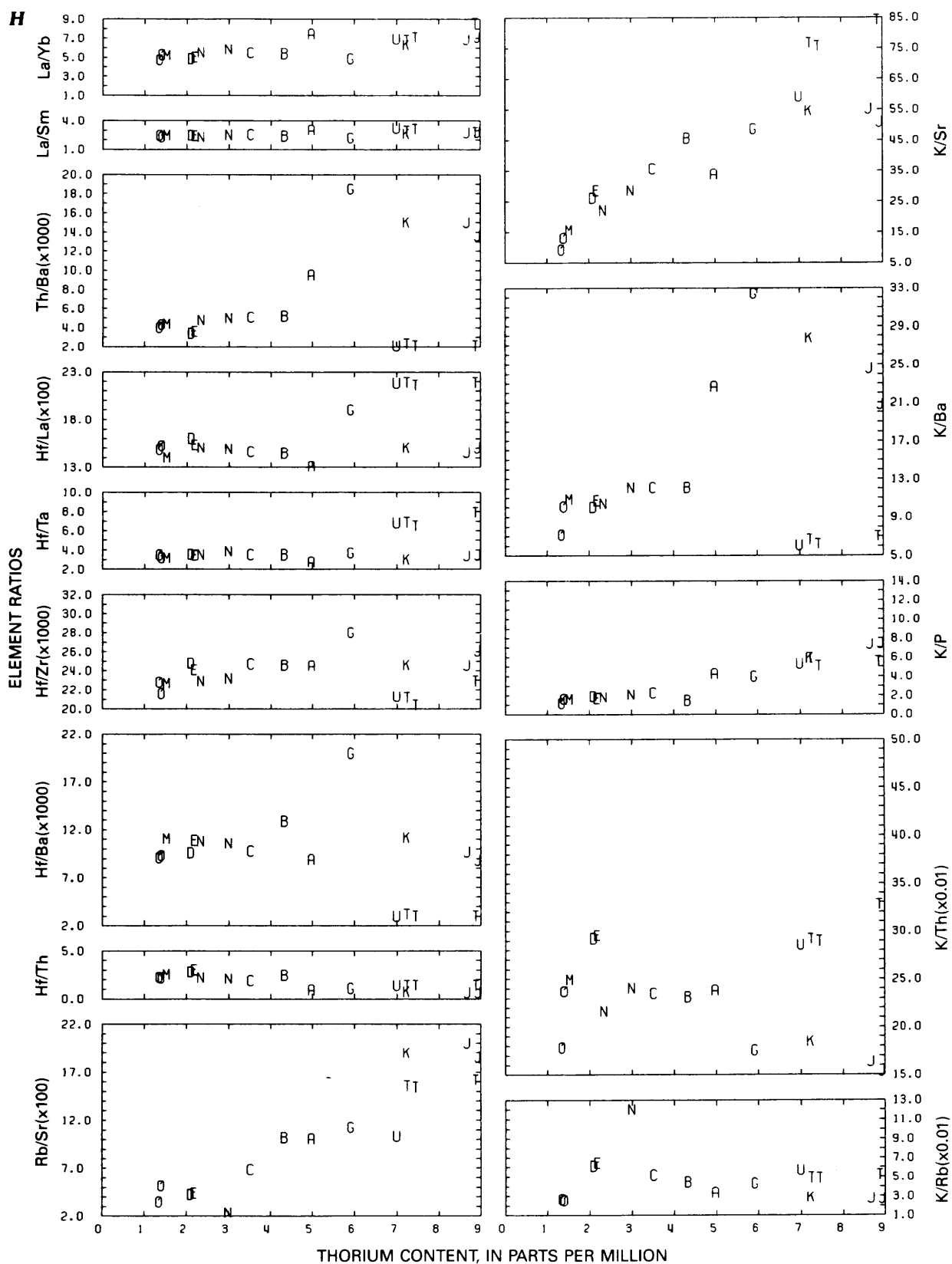


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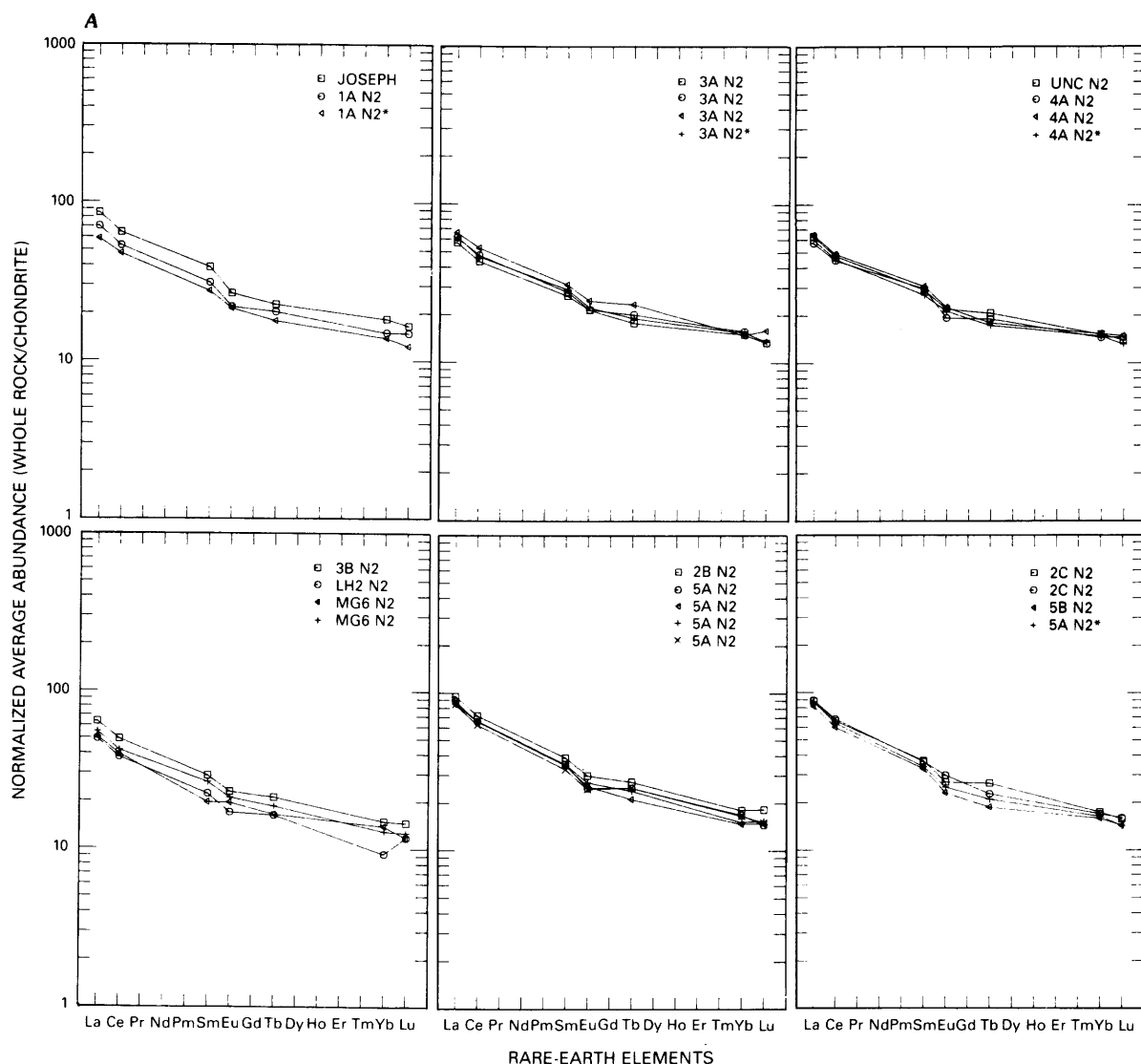


Figure 6. Chondrite-normalized abundances of selected rare-earth elements (La, Ce, Sm, Eu, Tb, Yb, and Lu) in flows of the Yakima Basalt Subgroup. Values used in normalizing are those of Haskin and others (1968). Names of geochemical units as in tables 3–5 and figures 4–5. See table 1 for stratigraphic association of units. **A**, Grande Ronde Basalt. Data arranged in stratigraphic order from youngest (1A N2) to oldest (AS 16 R1), using terminology of Mangan and others (1985, 1986); data for the Joseph volcanics unit of Kleck (1976) (JOSEPH in upper left box) and one sample of the Imnaha Basalt (IMNAHA in lowermost right-hand box) shown for

comparison. **B**, Wanapum Basalt. Data shown in two boxes at upper left, arranged in stratigraphic order from youngest to oldest within members. Priest Rapids Member (PV, LOLO INC, ROSALIA); Powatka flow (PH POWAT); Roza Member (ROZA); Frenchman Springs Member (FS INC; FS ANOM); Eckler Mountain Member (SHUMAKER=PH SHUM; SW LOOK; DODGE; ROBIN). **C**, Saddle Mountains Basalt. Data arranged by member, from youngest to oldest, from Lower Monumental Member (LM at upper right) to Umatilla Member (UMATILLA at lower right).

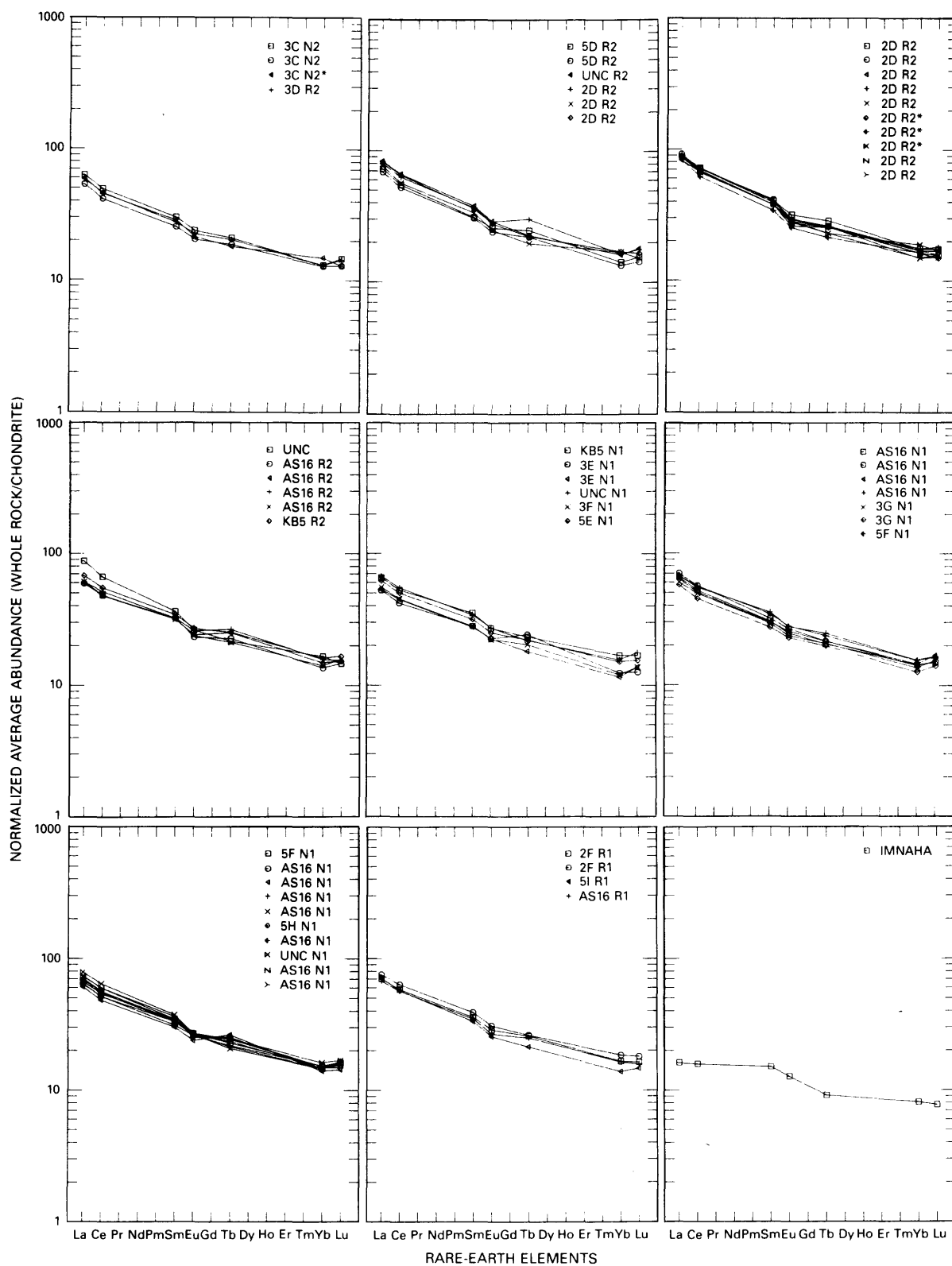


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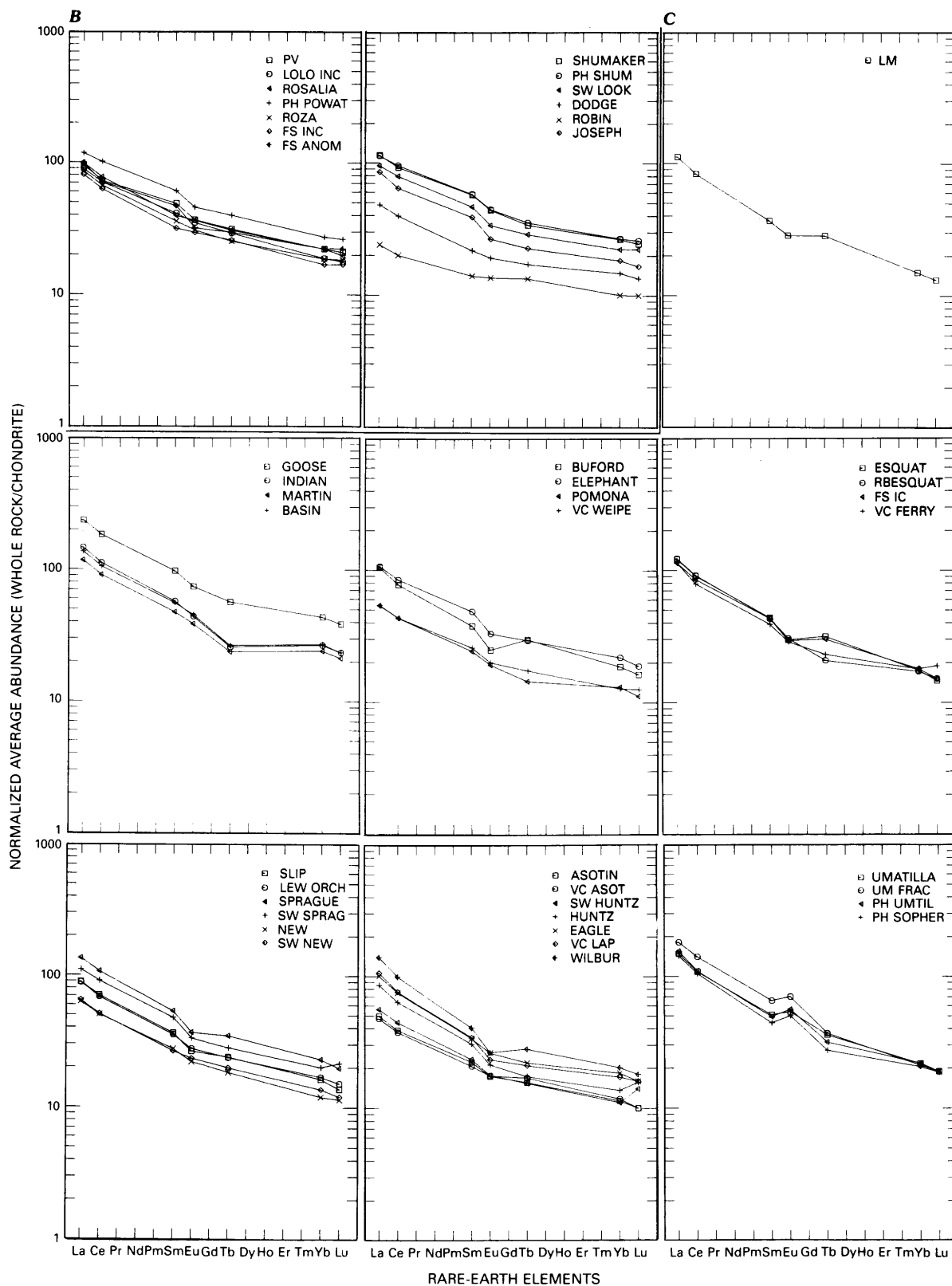


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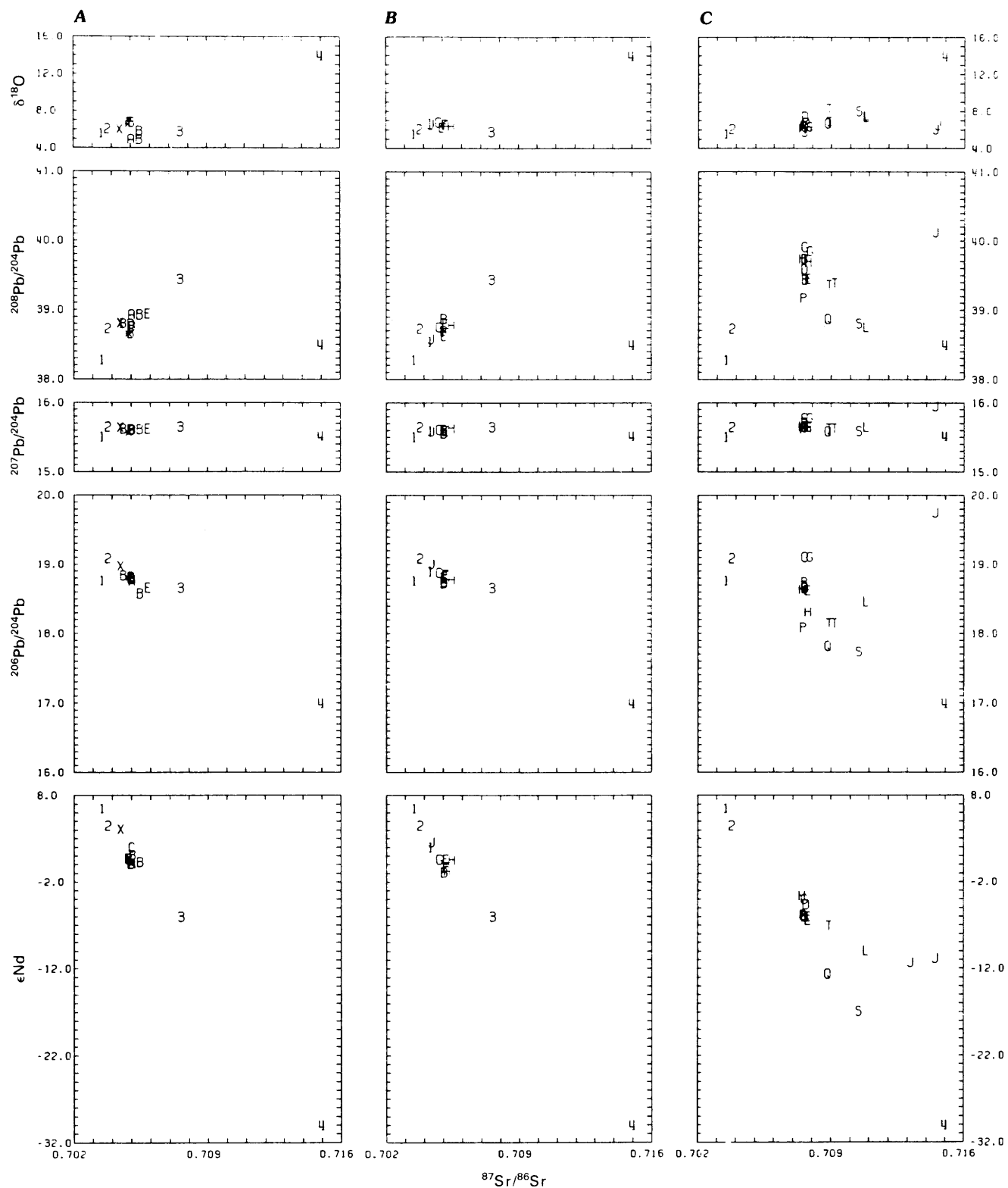


Figure 7. Isotope data for units of the Yakima Basalt Subgroup. Symbols 1, 2, 3, and 4 represent Carlson's (1984a, table 3) end-member compositions C1, C2, C3, and C4. See text for further discussion. A, Grande Ronde Basalt. *, LH2 N2 geochemical unit; #, MG6 N2 geochemical unit; other symbols as in table 3. B, Wanapum Basalt. Symbols as in table 4. C, Saddle Mountains Basalt. Symbols as in table 5.

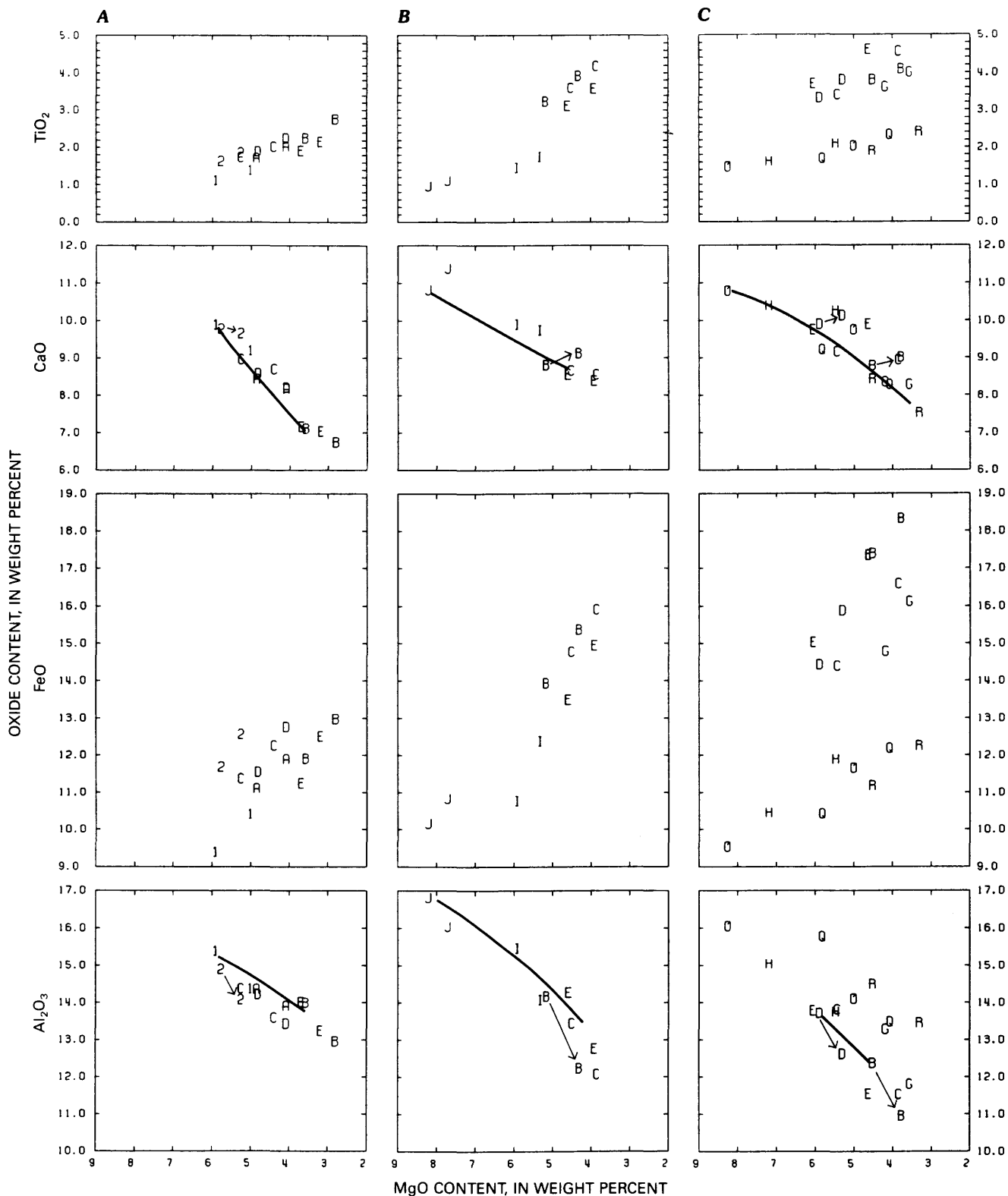


Figure 8. MgO variation diagrams comparing major-oxide chemistry of representative rocks and coexisting glasses in units of the Yakima Basalt Subgroup. Glass analyses adjusted before plotting (see text). Plots of CaO-MgO and Al_2O_3 -MgO are annotated to show average trends of bulk-rock chemistry (heavy solid lines) and representative trends

connecting coexisting rock and glass (short solid lines, arrows point toward glass composition). A, Grande Ronde Basalt. Symbols as in table 3. B, Wanapum Basalt. Symbols as in table 4. C, Saddle Mountains Basalt. Symbols as in table 5, except O, SW NEW only; Q, HUNTZ only; R, VC LAP only.

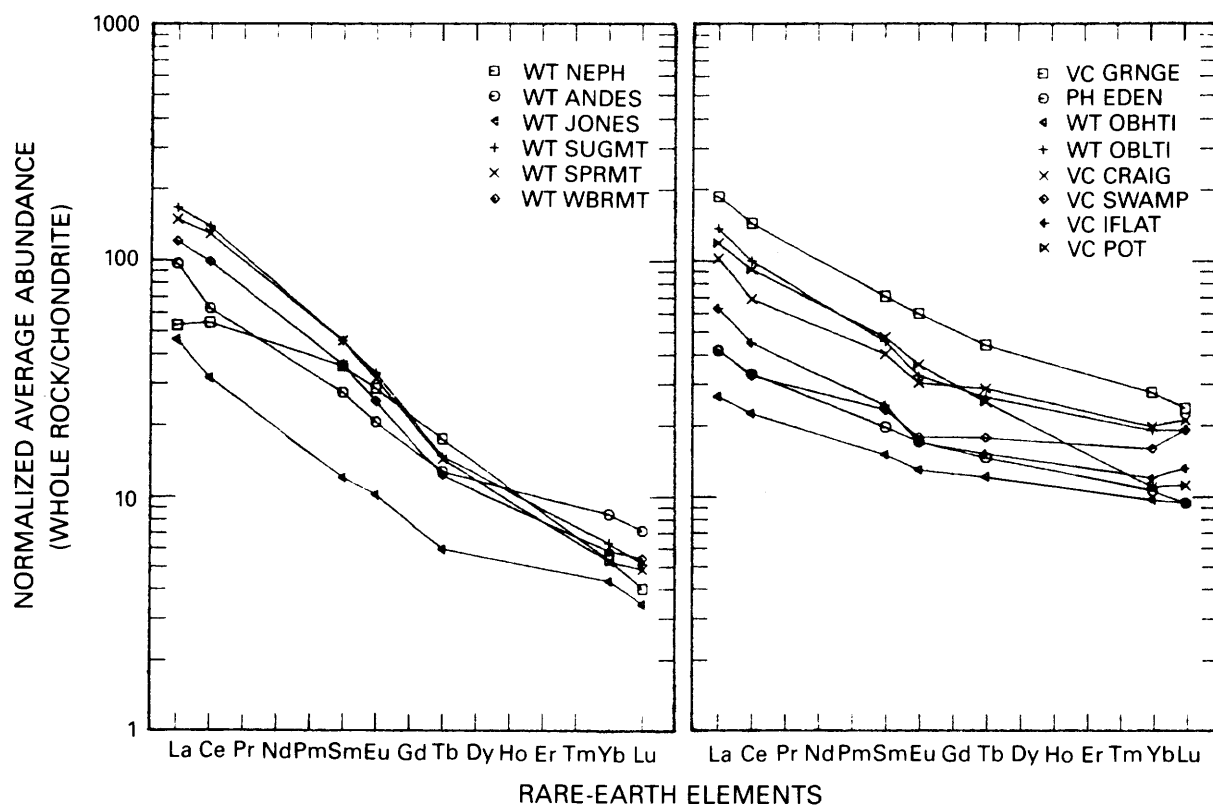


Figure 9. Patterns of chondrite-normalized abundances of rare-earth elements in additional analyzed flows. Symbols for flows as in tables 14 and 15.

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