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Geochemistry of Volcanic Rocks in the Marysvale Volcanic Field, West-Central Utah

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ABSTRACT

Igneous activity in the Marysvale volcanic field of west-central Utah occurred in three episodes. The first took place 34–22 Ma and was dominated by calc-alkaline, intermediate composition rocks of the Bullion Canyon Volcanics and Mount Dutton Formations. The e_{Nd} increased with time, reflecting decreasing crustal interaction with time.

The second episode, 23–14 Ma, was dominated by an alkalirhyolite and basalt (bimodal) assemblage. The similarity in isotopic ratios, together with a change to alkali rhyolites, is interpreted to be due to derivation of the nearly eutectic rhyolites by remelting the batholithic rocks that previously fed the Bullion Canyon volcanoes. Potassium-rich mafic rocks were erupted early in the second episode and were followed by more normal basalts. The third episode, 9–5 Ma, was also characterized by a bimodal assemblage of volcanic rocks. The highly variable isotopic ratios of the rhyolites indicate derivation from variable sources, reflecting episodic intervals of extensional tectonics.

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Igneous activity and associated mineralization in the Marysvale volcanic field along the west side of the Colorado Plateau in west-central Utah occurred mainly during three episodes: \approx 34–22 Ma, 23–14 Ma, and 9–5 Ma. In the first episode, during a regime of tectonic convergence, two contrasting suites of rocks were erupted concurrently: (1) poorly evolved, crystal-poor, pyroxene-plagioclase-bearing andesitic rocks, called the Mount Dutton Formation, from deep sources along the southern flank of the volcanic field, and (2) crystal-rich, biotite-hornblende-plagioclase-bearing dacitic rocks, called the Bullion Canyon Volcanics, from an east-northeast-trending shallow batholith under the middle of the field (Rowley and others, 1979, 1994; Steven and others, 1979, 1990; Steven, Rowley, and Cunningham, 1984; Cunningham and others, 1983; Steven and Morris, 1987).

Eruptions of crystal-rich ash-flow tuff from centers genetically related to the Bullion Canyon Volcanics (fig. 1) resulted in the formation of the Three Creeks caldera at 27.5 Ma (Steven, 1981), the Big John caldera at about 24 Ma (Steven, Cunningham, and Anderson, 1984), and the Monroe Peak caldera at 23 Ma (Rowley and others, 1988a,b). The Monroe Peak caldera is the largest caldera in the Marysvale volcanic field and was formed at the end of Bullion Canyon volcanism (Steven, Rowley, and Cunningham, 1984).

Isotopic data for Bullion Canyon Volcanics and related rocks (table 1, fig. 2) show a weak trend of increasing ϵ_{Nd} (-10 at 29 Ma to -5 at 21–22 Ma) with decreasing age. Initial strontium isotopic values are nearly constant (87 Sr/

⁸⁶Sr=0.7052–0.7062), whereas Pb isotopic compositions are variable. With the exception of one sample obviously contaminated by upper crustal material (No. 78-958, initial ⁸⁷Sr/ ⁸⁶Sr = 0.716, ²⁰⁶Pb/²⁰⁴Pb=19.8; table 1), Mount Dutton samples tend to have lower ε_{Nd} (–11) and ²⁰⁶Pb/²⁰⁴Pb (17.95–18.15), but similar initial ⁸⁷Sr/⁸⁶Sr to Bullion Canyon samples (fig. 2). Potassium, Rb, and U contents in samples from the Bullion Canyon Volcanics also increased with time, but remained relatively constant in Mount Dutton rocks.

At about 23-22 Ma, the composition of the igneous rocks changed to a bimodal suite of alkali rhyolite and basaltic rocks. These rocks are the products of the combined second and third igneous episodes, which continued to be erupted into the Quaternary. This change, from intermediate to bimodal volcanism at 23-22 Ma, is interpreted to be a result of the change from subduction to initial extension along the eastern margin of the Basin and Range province. The oldest basaltic lavas, erupted at 23-21 Ma (potassiumrich mafic lavas in table 1), contain 3.85 to 4.07 weight percent K₂O and have initial 87 Sr/ 86 Sr =0.7057–0.7061, ε_{Nd} = -6.7 to -8.5, and 206 Pb/ 204 Pb=18.47-19.03. The high initial 87 Sr/ 86 Sr and low ε_{Nd} of these samples relative to similar age Bullion Canyon samples may reflect either crustal contamination or input from a subducted slab (Best and others, 1980; Walker and others, 1991).

Voluminous but locally erupted rhyolites were prevalent during the second and third episodes, from 22 to 14 Ma and from 9 to 5 Ma. The largest volume of alkali rhyolite,

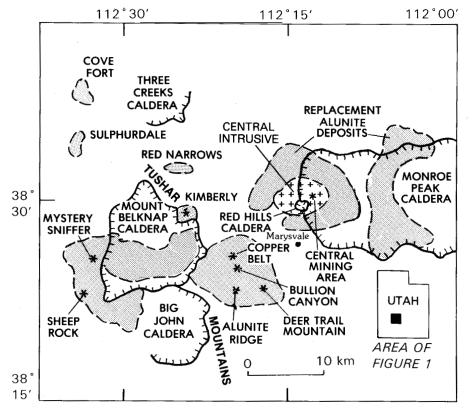


Figure 1. Location of calderas in central part of Marysvale volcanic field, Utah. Also shown are areas of altered rocks (shaded) and significant mineral deposits.

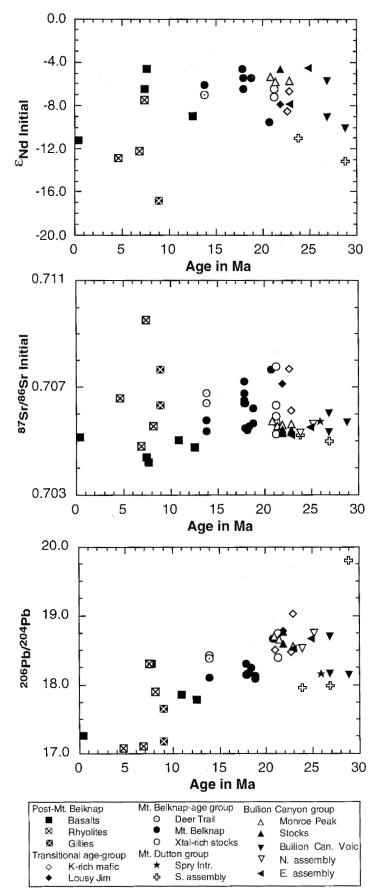


Figure 2. Plots of initial ϵ_{Nd} , ${}^{87}Sr/{}^{86}Sr$, and ${}^{206}Pb/{}^{204}Pb$ as functions of age for Marysvale samples.

called the Mount Belknap Volcanics (Cunningham and Steven, 1979; Budding and others, 1987), was erupted during the second episode (22–14 Ma, table 1). The Mount Belknap Volcanics was erupted from two areas (fig. 1): (1) the Mount Belknap caldera in the west, which formed at 19 Ma, and (2) a 12×12 -km area, about 20 km east of the Mount Belknap caldera, that was active between 21 Ma at the northeast end of the area and 14 Ma at the southwest end (Cunningham and others, 1982).

The Mount Belknap Volcanics have higher silica contents (71.4–76.8 weight percent) than the older Bullion Canyon Volcanics (58.1–65.7 weight percent; table 1), as well as generally higher Rb contents (269–564 ppm) and lower Sr contents (23–100 ppm) than most of the Bullion Canyon Volcanics (69–142 ppm Rb, 473–928 ppm Sr).

The Mount Belknap Volcanics, in addition to having major-element compositions that contrast with the Bullion Canyon Volcanics and Mount Dutton Formation, have e_{Nd} values similar to the latest Bullion Canyon Volcanics and among the highest ε_{Nd} values (-4.6 to -6.7, with one outlier; table 1, fig. 2) of any of the samples analyzed in this study. These data suggest that the processes involved in the production of these rhyolites involved comparatively little crustal contamination. Lead isotopic compositions of these samples (again with one exception) are tightly clustered with ²⁰⁶Pb/²⁰⁴Pb=18.08-18.24 (table 1; fig. 2). In contrast, the initial ⁸⁷Sr/⁸⁶Sr values of the Mount Belknap Volcanics (0.7053-0.7072) are somewhat higher and more variable than those of the Bullion Canyon Volcanics (fig. 2). Because the Sr abundances of the Mount Belknap Volcanics are comparatively low, the Sr isotopic compositions of these samples were more susceptible to modification by relatively small amounts of crustal contamination than were those rocks of Bullion Canyon Volcanics, which have higher Sr contents.

During the third episode of igneous activity (9–5 Ma), sparse basaltic lavas had normal K₂O contents (1.82–2.97 weight percent), lower initial ⁸⁷Sr/⁸⁶Sr (0.7042–0.7051), highly variable ε_{Nd} (-4.7 to -11.2), and low but variable ²⁰⁶Pb/²⁰⁴Pb (17.3–18.3, table 1; fig. 2) when compared to the older, potassium-rich, basaltic lavas. Low initial Sr values in association with low ϵ_{Nd} and low $^{206}\text{Pb}/$ ²⁰⁴Pb have also been found among some late Tertiary basalt and basaltic andesite on the Markagunt Plateau approximately 80 km south of this study area (fig. 3), also along the western margin of the Colorado Plateau, by Nealey and others (in press). Although we have a rather limited data set, we note that the lowest $^{206}\text{Pb}/^{204}\text{Pb}$ and ϵ_{Nd} values are found in the sample (79-S-44) with the highest silica content, whereas the highest values are found in the only true basalt analyzed (78-164). This situation is analogous to that observed by

Table 1. Lead, Sr and Nd isotopic compositions of selected samples from the Marysvale Volcanic Field.

					<u> </u>					, marysvan	•	rieiu.								
Sample	age (Ma)	SiO ₂	<u>206Pb</u> 1 204Pb	<u>207Pb</u> 1 204Pb	<u>208Pb</u> 1 204Pb	<u>87Rb</u> 86Sr	<u>87Sr</u> 2 86Sr	<u>87Sr</u> 86Sr Ini.	<u>147Sm</u> 144Nd	<u>143Nd³</u> 144Nd	<u>143Nd</u> 144Nd Ini.	٤Nd I								
Basaltic lavas																				
79-S-44	0.5		17.259	15.489	37.439	0.250	0.705122	0.70512	0.0998	0.512060	0.512060	-11.23								
M848	7.6		18.302	15.589	37.434	0.257	0.704388	0.70436		0.512297	0.512291	-6.53								
78-164 M672	7.8		18.297	15.568	37.907	0.230	0.704195	0.70417	0.1033	0.512390	0.512385	-4.71								
M673 M800	11 12.7		17.848 17.783	15.519 15.508	37.765 37.489	0.105	0.705026	0.70501	0.0014	0 540407										
Rhyolites	12.7	54.67	17.703	15.506	37.409	0.098	0.704748	0.70473	0.0914	0.512167	0.512159	-8.98								
78-115	4.8	76.49	17.073	15.449	36.925	42.440	0.709433	0.7065	0.2048	0 511973	0.511967	-12.94								
78-732	7		17.103	15.439	36.972	0.290	0.704789			0.512002		-12.34								
78-175	7.6		18.303	15.579	37.655	35.580	0.713320	0.7095	0.1134	0.512244		-7.57								
78-1695A	8.3	75.17	17.888	15.519	37.676	8.270	0.706495	0.70552												
Gillies Hill	0.1	00.01	47 474	15 470	07.050	0.040	0													
79-S-8A 79-S-9A	9.1 9.1		17.171	15.479 15.499	37.652	0.919	0.707749		0.0892	0.511762	0.511757	-16.92								
.79-S-9A 9.1 77.84 17.650 15.499 37.576 14.800 0.708223 0.70631 MT. BELKNAP-AGE GROUP																				
Deer Trail Rhyolite																				
M945	14	73.2	18.417	15.586	38.537	0.628	0.706864	0.70674	0.0868	0.512296	0.512288	-6.43								
Alunite												0.10								
M341	14		18.377	15.571	38.332	0.008	0.706362	0.70636	0.0503	0.512259	0.512254	-7.09								
Mt. Belknap Volcar		70 45	10.007																	
M779A M779B	14 14	76.15 75.31	18.097	15.568	38.080	24.120	0.710116	0.7053	0.1094	0.512315	0.512305	-6.11								
M687A	14	76.7	18.135	15.567	38.074	23.300 13.080	0.710383 0.709814	0.7058 0.70647												
M840A	18		18.303	15.596	38.371	45.230	0.717922	0.70647	0.1369	0.512391	0.512375	-4.64								
M19A	18	75.64				24.340	0.712962	0.7067	0.1000	0.012001	5.512070	7.04								
M672A	18.5	71.44	18.239	15.577	38.248	6.807	0.707308	0.70552												
M408A	18.1		18.143	15.567	38.234	1.945	0.706880		0.0937	0.512290	0.512279	-6.51								
M841 M502A	18	75.68	10 1 10	15 507	00 4 40	33.610	0.715782	0.7072	0.400.	0 6 4 0 0 4 1										
M833	18.1 18.3		18.148 18.165	15.567 15.567	38.149 38.076	15.350 19.120	0.709396 0.710339	0.70545 0.70537	0.1091	0.512344	0.512331	-5.49								
M820A	19		18.123	15.566	38.085	18.750	0.710689	0.70563	0 0965	0.512343	0.512331	-5.47								
M119A	19	76.41	18.075	15.577	38,107	20.730	0.711794	0.7062	0.0000	0.512545	0.012001	-3.47								
M50	20.9	71.5	18.662	15.638	38.999	5.180	0.709188	0.70765	0.1028	0.512131	0.512117	-9.60								
Crystal-rich stocks																				
M49A	21.4		18.380	15.605	38.494	2.293	0.705907	0.70521	0.0972	0.512251	0.512237	-7.24								
M837A M837B		74.76				11.060	0.709031	0.70567												
M838A	21.4 21.4	76.52	18.730	15 623	38.910	14.670 5.280	0.712219 0.707505	0.70776	0 1012	0.512287	0 510070	0.55								
M838B		76.62	10.700	10.020	50.310	17.730	0.711689	0.70630	0.1013	0.512267	0.512273	-6.55								
TRANSITIONAL-AG						17.700	0.711000	0.70000												
K-rich mafic lavas																				
M798	21.1		18.502	15.607	38.226	0.215	0.705754	0.70569												
M657	22.8		18.469	15.617	38.506	0.326	0.707785	0.70768		0.512187		-8.53								
M856	23	53,39	19.027	15.667	39.001	0.220	0.706182	0.70611	0.1031	0.512278	0.512262	-6.71								
Lousy Jim M822	22	69.4				1.991	0.707752	0.70713												
M823A	22	67.2	18.775	15.664	38.876	2.671	0.707955	0.70712	0 1041	0.512217	0.512202	-7.91								
BULLION CANYON	VOLC							011 07 12	011011	01012217	0.012202	1.01								
Monroe Peak																				
M678	21		18.694	15.626	38.559	0.626	0.705887		0.1052			-5.43								
79-1664A	23		18.551	15.624	38.410	1.897	0.706210	0.70559		0.512327		-5.75								
M569 M608	21.5 22	64.87 65.7	18.640	15.626	38.446	1.448 1.533	0.705962 0.706049		0.0937	0.512319	0.512306	-5.90								
Monzonite stocks	<u> </u>	00.7				1.000	0.700049	0.70557												
M867	22	58.25	18.583	15.614	38.430	0.949	0.705566	0.70527												
M621	22		18.758		38.521	0.841	0.705583		0.1060	0.512386	0.512308	-4.62								
M845	22	59.58				1.238	0.705807	0.70542												
M851	22	58.69				0.941	0.705574	0.70528												
M869	23	58.1				0.833	0.705552	0.70528												
M870 Bullion Canyon Vo	23 Icanic	58.67				0.945	0.705659	0.70535												
Bullion Canyon Vo M90	27		18.161	15.586	38.343	0.381	0.706156	0 70601	0 1026	0.512152	0 512134	-9.12								
79-1666	27		18.685	15.655	38.343	0.557	0.705493	0.70528		0.512152		-9.12								
79-S-1	29		18.146	15.577	38.088	0.190	0.705728	0.70565		0.512100		-10.14								
Northern Assembla	9																			
79-S-3	24		18.513	15.617	38.154	0.302	0.705353	0.70525												
M739		61.75	18.736	15.616	38.163	0.830	0.705898	0.70560												
Eastern Assemblag 79-1517	ge 23	56 56	18.523	15.588	38.260	0.170	0.705255	0 70520	0 1052	0 510015	0 512100	-7.94								
79-1518	23 25		18.661		38.232		0.705255			0.512215 0.512388		-7.94 -4.55								
MOUNT DUTTON F					00.202	0.000	0.,00014	0.70048	0.1073	0.012000	5.512570									
Mount Dutton Fmn. (southern assemblage)																				
79-1665	24	56.55	17.953	15.558	38.324	0.151	0.705241		0.1060	0.512056	0.512039	-11.04								
78-698	27		17.987	15.548	38.171	0.343	0.705092	0.70496												
78-958	29	57.51	19.796	15.787	39.983	0.850	0.716530	0.71618	0.1216	0.511945	0.511922	-13.20								
Spry Intrusion R2106	26.1	66 73	18 152	15.549	38 1/19	0.360	0.705833	0 70570												
									andard C	M-981 Un	ortainties in +	he data								
VUINCUEU IUL IIIdas III	uuuuid	ฉบา นนเย	iu andivst	/ J UI U. I Z I	- u.ua /o DE	/ mass Da	agu ur dridiV	อธอ บเทีย รีเ	anuaru of	11VI-201. UN	¹ Corrected for mass fractionation during analyses of 0.12 ± 0.03% per mass based on analyses of Pb standard SRM-981. Uncertainties in the data									

¹ Corrected for mass fractionation during analyses of 0.12 ± 0.03% per mass based on analyses of Pb standard SRM-981. Uncertainties in the data generally correspond to the fractionation uncertainties.
² Measured uncertainties are generally 0.003% to 0.004% and are relative to a value of 0.710254 ± 15 (6 analyses) of Sr standard SRM-987.
³ Measured uncertainties are generally 0.003% to 0.004% and are relative to a value of 0.511854 ± 12 (8 analyses) of the La Jolla Nd Standard.

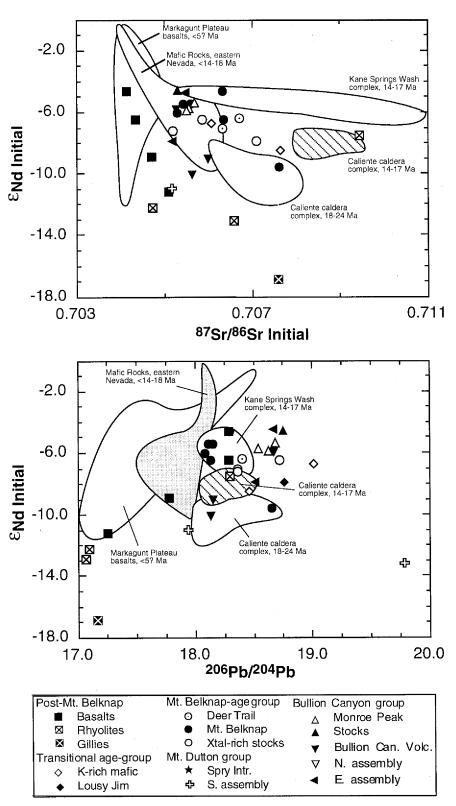


Figure 3. Plots of ε_{Nd} vs. ⁸⁷Sr/⁸⁶Sr, and ²⁰⁶Pb/²⁰⁴Pb for Marysvale samples. Shown for reference are fields of data for the Kane Springs Wash and Caliente caldera complexes in southeastern Nevada and the Markagunt Plateau in southwestern Utah.

Nealey and others (in press), who ascribed this rather unusual isotopic signature to interaction of ascending mantle-derived magmas with the lower crust. This interpretation was based primarily on two factors: (1) a suite of deep-crustal xenoliths from the San Francisco volcanic field, central Arizona (along

the southern margin of the Colorado Plateau) have isotopic characteristics that represent the end-member of this trend ($^{206}Pb/^{204}Pb \approx 16.0$, $^{87}Sr/^{86}Sr \approx 0.7025$, $\varepsilon_{Nd} \approx -20$; Nealey and Unruh, 1991), and (2) evidence for this isotopic signature is not found west of the approximate Colorado Plateau

boundary, even among similar-aged basalts in the St. George basin immediately southwest of the Markagunt Plateau (Unruh and others, in press).

Rhyolites that were erupted during the third episode of igneous activity had major-element compositions similar to those of the 22–14 Ma group but had markedly different radiogenic-isotope characteristics. Lead, Sr, and Nd isotopic ratios all show much more variation than ratios observed among the other age groups (fig. 2), and the lowest ε_{Nd} values(–12 to –16) and ²⁰⁶Pb/²⁰⁴Pb (17.0–17.2) are found among these samples. However, initial ⁸⁷Sr/⁸⁶Sr values do not appear to be well correlated with Nd or Pb isotopic compositions (figs. 2 and 3), and rhyolites from the third episode show the widest variation in initial Sr and Nd values found within the entire sample suite (again with the exception of the highly contaminated Mount Dutton sample previously mentioned).

The isotopic trends exhibited by the Bullion Canyon and Mount Belknap Volcanics show similarities to those exhibited by the similar-age Caliente and Kane Springs Wash caldera complexes in southeastern Nevada (fig. 3; data from Scott and others, 1995; Unruh and others, 1995; D.M. Unruh, unpub. data, 1995). In both areas, ϵ_{Nd} values generally increase with decreasing age (at least from 29-18 Ma in Marysvale), whereas initial ⁸⁷Sr/⁸⁶Sr values tend to become more variable with decreasing age (figs. 2 and 3). In both areas these features may be attributed to different processes and sources involved during the change from subductionrelated to extension-related volcanism. In the Caliente and Kane Springs Wash complexes, this transition is also reflected in the major-element chemistry of the rhyolite tuffs: Kane Springs Wash and post-17 Ma Caliente tuffs become metaluminous to weakly peralkaline, whereas the older tuffs are generally peraluminous (Rowley and others, 1995; Scott and others, 1995).

The Mount Belknap and Bullion Canyon Volcanics show a larger range of Pb isotopic compositions than do the tuffs of the Caliente and Kane Springs Wash complexes (figs. 3 and 4), although Pb isotopic data for the individual groups show fairly tight clusters (fig. 4). Lead isotopic data for the post-Mount Belknap samples are more similar to those for late Tertiary Markagunt Plateau basalts than to data for either Marysvale or Caliente-Kane Springs Wash samples.

All of the data on the ²⁰⁶Pb/²⁰⁴Pb vs. ²⁰⁷Pb/²⁰⁴Pb diagram plot significantly above the "northern hemisphere reference line" (NHRL; Hart, 1984), that represents the average trend of oceanic basalts in the northern hemisphere. This suggests that the Pb isotopic characteristics of these samples are derived primarily from lithospheric sources. Furthermore, Pb isotopic data for Marysvale samples in general show a pronounced average decrease in ²⁰⁶Pb/²⁰⁴Pb with decreasing age from about 22 Ma to the present (fig. 2), which may indicate a progressive change in the source region for these samples. A best-fit line through the ²⁰⁶Pb/²⁰⁴Pb vs. ²⁰⁷Pb/²⁰⁴Pb data in figure 4 yields an apparent age of 1.9 in view of the evidence for multiple sources for these samples, this apparent age may have no rigorous significance.

The isotopic data presented here for Marysvale samples are consistent with an interpretation of having been derived from three primary sources: the lithospheric mantle (his context, the terms "upper" and "lower" crust are not meant to imply that these represent exactly two discreet physical entities; rather, these terms are used in the context of two end-member isotopic signatures). The upper-crustal endmember, most evident in Mount Dutton sample 78-958 (table 1), is characterized by radiogenic Pb and Sr (²⁰⁶Pb/ $^{204}\text{Pb}{>}19.8;~^{87}\text{Sr}{/}^{86}\text{Sr}{>}0.715)$ and nonradiogenic Nd (ϵ_{Nd} <-13). The lower-crustal end-member, most evident in Gillies Hill sample 79-S8A, is also characterized by low ε_{Nd} (<-18) but perhaps only moderately radiogenic Sr (87Sr/ ⁸⁶Sr>0.708) and nonradiogenic Pb (²⁰⁶Pb/²⁰⁴Pb<17.0). This lower-crustal end-member is similar to that inferred for the Markagunt Plateau except that Sr appears to have been more radiogenic in the Marysvale area.

The isotopic characteristics of the lithospheric mantle are probably most-closely approximated by the only true basalt analyzed (sample 78-164, table 1; $\epsilon_{Nd} \approx -4.7$, $^{206}Pb/$ 204 Pb ≈ 18.3 , and 87 Sr/ 86 Sr ≈ 0.704). These values are very similar to those estimated for the lithospheric mantle under the Kane Springs Wash and Caliente caldera complexes (Scott and others, 1995; Unruh and others, 1995), and surprisingly close to those estimated by Johnson and Thompson (1991; $\epsilon_{Nd} \approx -4$, ${}^{206}Pb/{}^{204}Pb \approx 18.2$, ${}^{87}Sr/{}^{86}Sr \approx 0.705$) for the isotopic composition of the lithospheric mantle in the northern Rio Grande Rift on the opposite side of the Colorado Plateau. This isotopic component may represent the lithospheric mantle directly or the lithospheric mantle modified by a "subduction component" (Fitton and others, 1991) in the form of fluids or melts derived from the subducted slab. The fact that basalts with $\varepsilon_{Nd} = -4$ to 0 are relatively common among late Tertiary and Quaternary basalts on the Markagunt Plateau (Nealey and others, in press) suggests that the ε_{Nd} value of unmodified lithosphere in this area may be closer to zero (Farmer and others, 1989).

During eruption of the Mount Dutton Formation, the Bullion Canyon Volcanics, and the 20–18 Ma Mount Belknap Volcanics, crustal interaction progressively decreased, as evidenced by progressively increasing ε_{Nd} (fig. 2). The variation in Pb isotopic data during this interval suggests that both the Bullion Canyon and Mount Dutton magmas interacted with both upper and lower crust (figs. 2 and 3). The apparent progressive decrease in crustal interaction may reflect either thinning of the lithosphere during the onset of extension, providing a thinner column of crust for rising magmas to interact with, or production of progressively larger volumes of magma within the lithospheric mantle: all of these in effect dilute the crustal influence.

The similarity of neodymium, lead, and strontium isotopic values, together with the contrasting major element

15.8

15.7

15.6

15.5

15.4

40.5

39.5

38.5

37.5

36.5

 \boxtimes

X

 \diamond

Post-Mt. Belknap

Basalts

Gillies

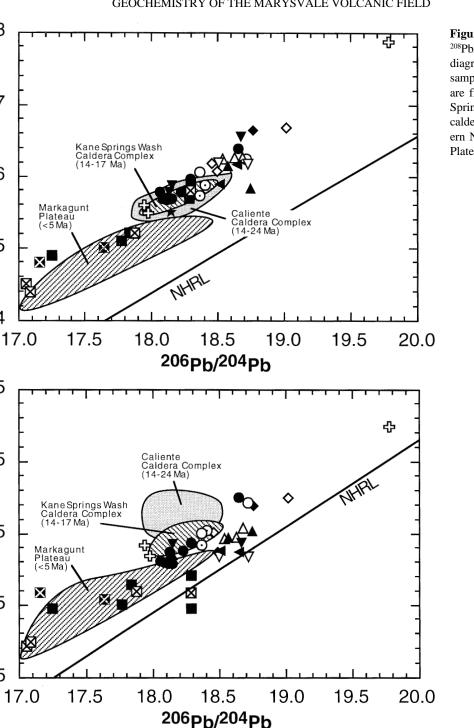
Transitional age-group

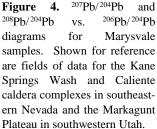
Rhyolites

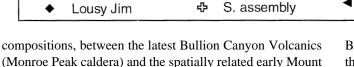
K-rich mafic

208pb/204pb

207pb/204pb







Mt. Belknap-age group

Deer Trail

Spry Intr.

Mt. Belknap

Xtal-rich stocks

0

0

★

Mt. Dutton group

Bullion Canyon group

Stocks

Δ

▲

▼

 ∇

◀

Monroe Peak

N. assembly

E. assembly

Bullion Can. Volc.

Belknap Volcanics (crystal-rich stocks), is consistent with the interpretation that the Mount Belknap Volcanics was derived from partial melting of Bullion Canyon batholithic rocks. This could be achieved as rising isotherms from the onset of extensional tectonics melted nearly eutectic magmas from the former magma chambers that fed Bullion Canyon volcanoes.

The Sr, Nd (fig. 3), and particularly the Pb isotopic compositions (fig. 4) of the samples formed during the third episode of volcanic activity appear to be strongly influenced by interaction with the lower crust (although rhyolites from this group have highly variable isotopic compositions). The fact that both mafic and silicic rocks have this signature may imply that the site of primary melt generation may have risen to the crust-mantle boundary during this period, particularly if this boundary had been modified by numerous earlier injections of mantle-derived magmas (Johnson and others, 1990). This interpretation is consistent with episodic intervals of increased extension and derivation of the late rhyolitic magmas from a variety of levels in the crust as isotherms rose in response to extension.

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