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Clinoptilolite resources in the Tushar Mountains
west-central Utah

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Two potentially economic deposits and one possibly economic deposit of clinoptilolite have been found in the Tushar Mountains of west-central Utah. All three deposits are near the nonwelded distal ends of rhyolitic ash-flow tuff sheets, and appear to have formed mainly by reaction of glassy volcanic ash with ground water. Available data are too few and too preliminary to support tonnage and grade estimates, but scattered samples combined with visual estimates suggest an ultimate resource of about several billion tons. A significant part of this resource may contain rock of 80 percent or more clinoptilolite in deposits easily amenable to open-pit mining. The main dilutants are feldspar, biotite, and quartz phenocrysts and foreign lithic rock fragments, many of which possibly could be removed mechanically. Clay minerals seem to be minor constituents in most of the deposits.

What is clinoptilolite?

Clinoptilolite is a common zeolite that occurs widely in sedimentary and pyroclastic volcanic rocks. Zeolites are crystalline hydrated aluminosilicates of alkali and alkaline earth metals, and form most commonly by the reaction of silicic volcanic glass fragments and pore water (such as ground water, lake water, and ocean water). The specific zeolite mineral formed depends largely on the chemical composition of the water involved and on the activity ratio of alkali ions to hydrogen ions. Clinoptilolite commonly forms by reaction of

volcanic glass with ground water or with relatively dilute lacustrine brines; more strongly saline or alkaline lake waters result in other zeolite species or even authigenic feldspar.

Clinoptilolite has a theoretical composition of $(\text{Na},\text{K})_2\text{O}\cdot\text{Al}_2\text{O}_3\cdot 10\text{SiO}_2\cdot 6\text{H}_2\text{O}$, but it forms an isomorphous compositional series with heulandite by Ca and Mg substituting in different degrees for Na and K, and the actual composition varies considerably. Heulandite is commonly less siliceous than clinoptilolite. Clinoptilolite, as do most zeolite minerals, can gain or lose water and exchange cations without major changes in crystal structure, and thus is especially amenable to ion-exchange reactions. In addition, zeolite crystalline structures contain "holes" of specific dimensions that serve as molecular sieves in trapping other cations--the size of the hole determining which cation or molecule can be trapped. Both the ion-exchange and molecular sieve characteristics have wide industrial applications. Clinoptilolite has the specific property of trapping (NH_4) ions, and thereby is especially useful in treating sewage effluents and agricultural waste waters, as well as for other agricultural and industrial purposes. Clinoptilolite has been used extensively in Japan, but only to a minor extent in the United States. With expanding emphasis on preventing man-made pollution, a much more extensive use of clinoptilolite seems assured in the United States in the future.

Clinoptilolite is a relatively common mineral where Tertiary or Quaternary silicic volcanic ash accumulated in a watery environment--a combination found at many places throughout Western United States. A low unit price for clinoptilolite probably will prevail after widespread

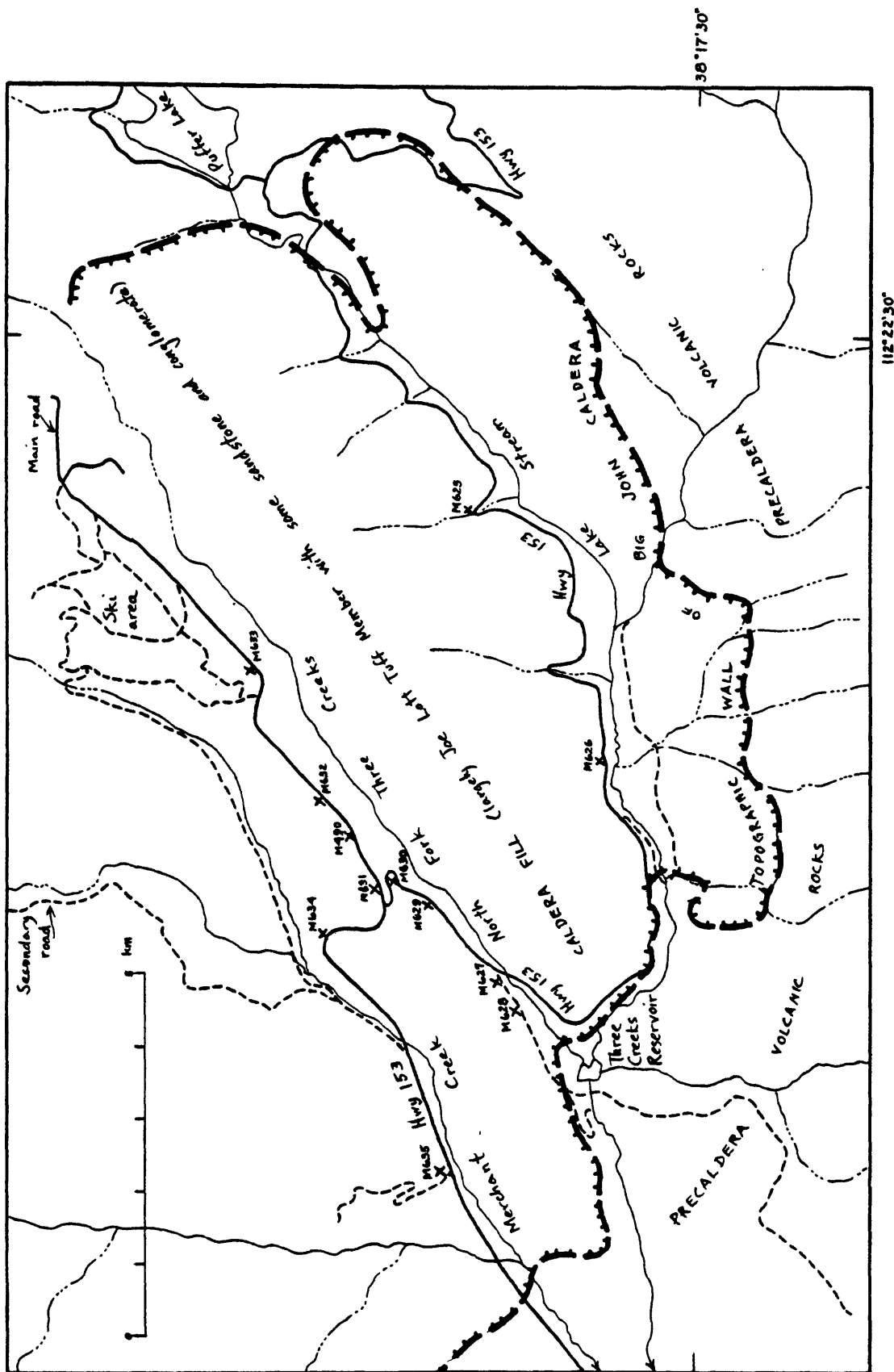
use has been achieved. A major economic consideration, once a sizeable body of good grade material has been identified therefore, will be proximity to a potential market. Mining costs also will be an important economic factor. In this report, we identify 3 potential sources of clinoptilolite, but leave most of these economic factors for potential developers to consider.

Location

The Tushar Mountains (fig. 1) are a major topographic element in the High Plateaus of west-central Utah, and are located along the eastern border of the Great Basin 240 to 290 km south of Salt Lake City. The mountains are high and in part very rugged and inaccessible. The clinoptilolite deposits we have identified, however, are either at the northwestern margin of the mountains near Cove Fort (fig. 2), or near Highway 153 that extends east from Beaver across the mountains to Junction (fig. 3), and are easily accessible by automobile.

General geology

The Tushar Mountains are in the heart of the Marysvale volcanic field and consist chiefly of volcanic or related intrusive rocks. As outlined by Steven, Rowley, and Cunningham (1978), volcanism began 35 to 30 million years (m.y.) ago, and since then and about 21 m.y. ago calc-alkaline intermediate-composition lavas were erupted from many scattered and in part clustered stratovolcanoes. Coalescing volcanoclastic aprons surrounded the volcanoes and built a widespread volcanic plateau. Ash-flow tuffs were erupted episodically through this span of time, some from centers within the Marysvale field, and some from centers to the west and southwest. The resulting volcanic assemblage is exceedingly



complex, but most details of this complexity have little relevance to this report. The area was tectonically stable during this period of volcanism, and only minor local faulting can be documented.

About 21 m.y. ago the composition of the erupted volcanic rocks changed from calc-alkaline intermediate-composition to a bimodal assemblage of basalts and high-silica alkali rhyolites that was erupted episodically until quite recently. The rhyolites were erupted in greatest volume early during this period--between 21 and 17 m.y. ago--when the Mount Belknap Volcanics accumulated around two concurrently active source areas (Cunningham and Steven, 1977). The western source area was in the northern Tushar Mountains where the Joe Lott Tuff Member erupted 19 m.y. ago and the major Mount Belknap caldera (fig. 1), approximately 14 km across, subsided as the source magma chamber was emptied. The Joe Lott Tuff Member spread radially from the caldera and accumulated in low areas on the surrounding volcanic plateau. Bimodal volcanism was broadly contemporaneous with Basin-Range block faulting that disrupted the entire Great Basin and High Plateaus in late Cenozoic time.

Clinoptilolite deposits formed from silicic ash-flow tuffs erupted during both the calc-alkalic and the bimodal periods of activity. The large deposits near Cove Fort (fig. 2) formed in the distal part of an unnamed calc-alkalic ash-flow tuff sheet derived from an unknown source in the Great Basin to the west shortly before 22 m.y. ago. The deposits near Three Creeks Reservoir in the headwaters of Beaver River (fig. 3) formed in the marginal part of an accumulation of ash-flow tuffs in the Mount Belknap Volcanics, a major rhyolite unit in the bimodal

assemblage. The possibly important clinoptilolite deposits in the western Tushar Mountains near the Beaver River are near the northern end of a local ash flow sheet that was erupted late during the period of calc-alkalic eruptions.

Cove Fort occurrence

The clinoptilolite occurrence southeast of Cove Fort (fig. 2) is near the eastern margin of a sheet of nonwelded ash-flow tuff that passes laterally eastward into tuffaceous sandstone and related sedimentary rocks. The source of the sheet appears to have been in the Great Basin to the west. The zeolitic tuff underlies the 22 m.y.-old Osiris Tuff (Fleck and others, 1975, p. 59), a major-ash flow unit to the east and southeast, and overlies in turn the tuff of Albinus Canyon and the 27 m.y.-old Three Creeks Tuff Member of the Bullion Canyon Volcanics (Steven and others, 1977), both derived from sources in the southern Pavant Range east and northeast of figure 1. The zeolitic tuff is cut by a hypabyssal pluton of monzonite that had been exposed by erosion 22 m.y. ago so that the Osiris Tuff was deposited unconformably across it. The zeolitic tuff, therefore, is presumed to have been deposited sometime during the middle of the 5 m.y. interval between eruption of the Three Creeks Tuff Member and the Osiris Tuff--perhaps about 24 m.y. ago. The zeolitic tuff has no known stratigraphic counterparts in the Pavant Range or Tushar Mountains to the northeast, east, and southeast. The area toward its probable source in the Great Basin is too poorly known at present for possible stratigraphic equivalents to be determined.

The zeolitic tuff is exposed in a series of fault blocks that form a west-extending ridge at the northwest end of the Tushar Mountains (figs. 1 and 2). Away from this spur, the tuff is exposed only in a few low knolls south of Sulphurdale. Stratigraphically equivalent tuffaceous sandstones are exposed in roadcuts along highway I-70 at the pass between Cove Creek and Clear Creek (fig. 1) and in a few natural exposures less than a kilometer to the east. Thus almost the full known extent of clinoptilolite at the Cove Fort occurrence is shown on figure 2.

The zeolitic rock is a massive white ash-flow tuff 100 m or more thick. Zeolitized matrix commonly constitutes 80 percent or more of the rock, and the remainder consists of fresh phenocrysts of feldspar, biotite, and quartz, and of irregularly distributed foreign lithic fragments. Visual estimates suggest that the rock was uniformly altered to clinoptilolite by diagenetic processes. Inasmuch as neither the ash-flow tuff nor the related tuffaceous sandstone show evidence of having been deposited in standing water, or that any lake developed in the area after deposition, it is presumed that diagenesis was accomplished by ground water. The result is a compact but soft lithified white rock that has a distinctive greasy luster on fresh surfaces. This luster, which can be compared to that on a broken surface of a cake of white laundry soap, is characteristic of zeolitic tuffs.

Qualitative X-ray diffraction studies of the matrix of 5 random samples (localities shown on fig. 2) of the zeolitic tuff gave the following results:

78-S-24A Clinoptilolite, major; quartz, significant; clay, minor
 78-S-25 Clinoptilolite, major; quartz minor
 78-S-27A Clinoptilolite, major. quartz, significant; clay, minor
 78-S-27B Clinoptilolite, major; quartz, significant; clay, minor
 78-S-28 Clinoptilolite, major; quartz, present; clay, minor

Petrographic modes of thin sections from the same samples in percent are the following:

Sample no.	Matrix	K-spar	Plag.	Qtz.	Bio.	Opaq.	Lithics
78-S-24A	85	3	4	Tr.	1	1	6
78-S-25	84	7	3	-	1	Tr.	5
78-S-27A	87	3	6	-	1	Tr.	5
78-S-27B	82	6	3	-	1.5	Tr.	7.5
78-S-28	78	12	6	1	1	Tr.	2

The quartz noted in the X-ray diffraction patterns is alpha quartz and probably represents fragments of phenocrysts and some of the residual silica left over from conversion of siliceous glass to clinoptilolite. Much of the diagenetically derived silica seems to have been removed in solution in the generally open system ground water environment of alteration, as the local sparse secondary concentrations of chalcedony are not abundant enough to account for the diagenetically available silica. The feldspar, biotite, and quartz phenocrysts, and possibly the foreign lithic fragments, are sufficiently distinctive in physical properties from the zeolitic matrix that they possibly could be separated mechanically. Diagenetic clay minerals or silica, on the other hand, probably are too finely dispersed in the matrix to be easily separated.

Although the area of occurrence of the clinoptilolite near Cove Fort is highly faulted (fig. 2), the net result is economically advantageous, inasmuch as the faulting tended to keep the zeolitic tuff unit near the surface where it could be mined easily. Open-cut mines could be established at many places where virtually no overburden needs to be stripped. Most of these places can be reached readily by short, easily maintained roads. Surface disturbance for development or mining need be minimal.

The main area of exposure of the zeolitic tuff is somewhat more than 3 km across both N-S and E-W. Outlying areas add to the total area of occurrence. Considering a square block 3 km across and 100 m thick, a minimum volume of 9×10^8 cubic meters of zeolitic tuff was computed. Again using a minimal figure of 2.0 for the density of the clinoptilolite-bearing rock, a resource of at least 1.8 billion metric tons is indicated. A metric ton is about the same as a long ton in English units. This specific figure has very little meaning in itself, other than to indicate that a tremendous tonnage of potentially valuable zeolitic tuff exists near Cove Fort, and that large amounts of this are easily available. Careful sampling and analysis of many local fault blocks are needed, however, before the true economic potential of the deposits can be established.

Three Creeks Reservoir occurrence

South of their source in the Mount Belknap caldera (fig. 1), ash flows of the Mount Belknap Volcanics accumulated chiefly in a depression marking the earlier Big John caldera. Most of the ash-flow tuffs are slightly to moderately welded and are completely devitrified. Near the

southern topographic wall of the Big John caldera (fig. 3), the tuffs locally remained glassy and unwelded, and some of these glassy tuffs subsequently were altered diagenetically to clinoptilolite and montmorillonite. Through 1978, our detailed geologic mapping was limited to the northeastern part of the area shown on figure 3; the remainder, including the part with the best-grade clinoptilolite, had been covered in broad reconnaissance only, and detailed geologic relations are poorly known.

The Big John caldera formed in response to eruption of the Delano Peak Tuff Member of the Bullion Canyon Volcanics; the Delano Peak Member has been dated by the K-Ar method as about 22 m.y. (Steven and others, 1977), the same as the stratigraphically older Osiris Tuff; analytical uncertainties, however, easily permit a million years or more to have intervened. The Big John caldera was partly filled by mafic lava flows shortly after it formed, but much of the same area was still a depression several million years later when the 19 m.y. old ash flows of the Mount Belknap Volcanics were erupted. Retention of a local depression this long seems unlikely unless it were drained, but to date our reconnaissance has not located the required outlet. A sequence of fluviatile sandstones and conglomerates several tens of meters thick covered the bottom of the depression near its southern margin.

About 19 m.y. ago, great pyroclastic eruptions from the Mount Belknap caldera area to the north (fig. 1) spread voluminous crystal-poor ash flows radially outward to accumulate in topographically low areas as the Joe Lott Tuff Member (Steven and others, 1977; Cunningham and Steven, 1977). This member filled the southern part of the Big John

caldera to depths of 150 m or more. No evidence was seen to indicate that any of these ash flows were deposited in water, again suggesting that the Big John depression was drained.

Subsequent smaller-volume eruptions of more crystal-rich ash covered the massive Joe Lott Member with thinner beds of tuff. Although most of these layers appear to have spread as ash flows and to have been deposited subaerially, local exposures of well-bedded water-laid tuff indicate that local ponds existed at least part of the time.

Whereas most of the Mount Belknap ash-flow tuff in the Big John caldera is slightly to moderately welded and devitrified, some near the southern margin remained glassy and unwelded. This glassy tuff was chiefly altered by diagenetic processes to clinoptilolite or montmorillonite. The altered tuff was sampled to determine the mineralogy, but no attempt was made at this early stage of the investigations to determine thickness or lateral extent.

Qualitative X-ray diffraction studies of 11 samples of Mount Belknap tuffs near Three Creeks Reservoir (localities shown in figure 3) gave the following results:

M 490	clinoptilolite, major; feldspar and quartz, minor	(altered crystal-rich tuff, matrix only)
M 626	clinoptilolite, predominant	(altered Joe Lott Member)
M 627	clinoptilolite, predominant	(altered Joe Lott Member)
M 628	clinoptilolite, predominant	(altered Joe Lott Member)
M 629	No zeolite, mostly sanidine, albite, and quartz	(devitrified Joe Lott Member)
M 630	clinoptilolite, major	(altered crystal-rich tuff, matrix only)
M 631	montmorillonite, predominant	(altered crystal-rich tuff, matrix only)
M 632	clinoptilolite, minor; mostly sanidine, albite, and quartz	(devitrified Joe Lott Member)
M 633	No zeolite; mostly sanidine, albite, and quartz	(devitrified Joe Lott Member)
M 634	clinoptilolite, present; quartz major	(devitrified Joe Lott Member)
M 635	No zeolite; mostly sanidine, albite, and quartz	(devitrified Joe Lott Member)

Petrographic modes of thin sections from the more altered of these samples are shown in percent:

Sample

number	Matrix	K-spar	Plag.	Qtz.	Bio.	Pyrox.	Opaques	Lithics	Unit ¹
M 490	65	10	7	2	1	2	3	10	x-r
M 626	90	1	.5	.5	Tr.	Tr.	Tr.	8	JL
M 627	96	1	1	-	1	-	Tr.	1	JL
M 628	87	5	1	Tr.	1	-	Tr.	6	JL
M 630	78	13	Tr.	Tr.	Tr.	-	Tr.	8	x-r
M 631	81	T2	Tr.	-	-	-	1	6	x-r

¹x-r Crystal-rich tuff in upper part of Mount Belknap Volcanics

JL Crystal-poor Joe Lott Tuff Member, Mount Belknap Volcanics

High-grade clinoptilolite was obtained from 3 samples (M 626, M 627, and M 628) of crystal-poor tuffs of the Joe Lott Member near the Three Creeks Reservoir (fig. 3). The zeolitic matrix in these samples averages more than 90 percent of the rock. M 626 was from near the base of the unit, and M 627 and M 628 from undetermined vertical positions within the main body of the unit. About 4 km separate the easternmost from the westernmost of these samples, and brief inspection of float and outcrops between these suggest that much of the intervening area is underlain by diagenetically altered tuff of the same general aspect.

Moderate-grade clinoptilolite was determined in samples M 490 and M 630 from crystal-rich layers near the top of the Mount Belknap sequence 3-4 km northeast of the Three Creeks Reservoir (fig. 3). These samples are sufficiently near one another to suggest a potential

resource in this area, but another nearby sample of the same type of tuff, M 631, consisted largely of montmorillonite. Some of the tuff in this vicinity is well-bedded, indicating deposition in a local pond, so the diagenetic environments here could have varied considerably and have led to heterogeneously altered products.

Samples of the Joe Lott Member peripheral to the area of zeolite occurrence were largely devitrified (M 620, M 633, M 635) and contain no clinoptilolite or montmorillonite. Diagenesis in the general Three Creeks Reservoir area thus appears to have varied considerably in degree and distribution.

In summary, we can report an area at least 4 km across near the Three Creeks Reservoir in which high-grade clinoptilolite has been determined in at least 5 random samples spread over a vertical stratigraphic range of 150 meters or more. The vertical or lateral extent of high grade material has not been determined at any local area, but visual estimates suggest that potentially valuable local concentrations could exist. The area is easily accessible by good roads, and additional testing can be accomplished readily. The tonnage of diagenetically altered tuff in this vicinity is estimated in terms of millions of tons; the percentage of good grade clinoptilolite within this tonnage is unknown but could be substantial.

Occurrence near Telluride Power Plant No. 2

A layer of soft white relatively crystal-rich ash-flow tuff is exposed on the north slope of the canyon of Beaver River a few kilometers northwest of Telluride Power Plant No. 2 and above a sharp switchback on Highway 153 (fig. 1). This area had not been mapped

geologically by us through 1978, and our knowledge of the geology is minimal. The tuff appears to be stratigraphically high in the section of calc-alkalic volcanic rocks comprising this part of the Marysvale volcanic field, but its precise stratigraphic position or lateral correlatives are not known.

Two grab samples taken at the switchback (M 636, M 636A) contain significant clinoptilolite according to X-ray diffraction patterns. A petrographic mode of a thin section of one of these samples showed 75 percent zeolitic matrix, 12 percent feldspar phenocrysts, trace to 1 percent opaque, biotite, quartz, and pyroxene phenocrysts, and 11 percent lithic rock fragments.

Certainly the clinoptilolite resource potential of this ash-flow tuff sheet merits more investigation.

Cation exchange capacity

The cation exchange capacity of six zeolite samples was determined by Harry Starkey of the U.S. Geological Survey, and is tabulated below. The procedure followed is described in Starkey (1964). Samples were crushed and soaked overnight at room temperature in an ammonium chloride solution. The amount of ammonia adsorbed on the sample was determined by distillation. The results are reported in milliequivalents per 100 grams of sample which is a standard format. The results indicate that four of the six samples tested are of average quality, and two samples (M 626 and M 628) are of excellent quality.

Sample Number	Exchange capacity		
	(milliequivalents/100 grams)		
M 490	62.1	Crystal-rich tuff	} Three Creeks Reservoir occurrence
M 626	146.4	Joe Lott Tuff Member	
M 628	117.3	Joe Lott Tuff Member	
78-S-24A	54.2	} Cove Fort occurrence	
78-S-27B	63.5		
78-S-28B	62.6		

Analytical results

The six samples of clinoptilolite-bearing tuff tested for exchange capacity were also analyzed for their contents of CaO , Na_2O , and K_2O . The results tabulated below fall into 3 groups that correspond to the different tuffs sampled. Sample M 490 of a crystal-rich tuff near the top of the Mount Belknap Volcanics near Three Creeks Reservoir is relatively high in CaO , Na_2O , and K_2O for a total of 8.06 percent. The underlying Joe Lott Tuff Member (M 626, M 628) is relatively low in these constituents (5.75-5.78 percent), and of this total, CaO is predominant. The zeolitic tuff near Cove Fort, on the other hand, tends to be high in total CaO , Na_2O , and K_2O content (6.59-8.65, percent), with K_2O predominating over CaO and Na_2O . Comparing these results with the exchange capacity data, it can be seen that the tuffs with high CaO content and relatively low Na_2O and K_2O contents (M 626, M 628) have the highest cation exchange capacity. The more sodic and potassic tuffs have about average exchange capacity.

Sample number	Content (percent)				Sample description	
	CaO	Na ₂ O	K ₂ O	Total		
M 490	3.25	1.75	3.06	8.06	Crystal-rich tuff	} Three Creeks Reservoir occurrence
M 626	3.00	0.81	1.94	5.75	Joe Lott Tuff Member	
M 628	3.31	0.34	2.13	5.78	Joe Lott Tuff Member	
78-S-24A	2.38	0.79	3.41	6.59	} Cove Fort occurrence	
78-S-27B	1.69	1.46	5.50	8.65		
78-S-28	1.81	1.63	4.94	8.38		

Acknowledgments

X-ray diffraction determinations for this report were made by Michael G. Nelson, and petrographic modes by James G. Brophy. Harry Starkey measured the cation exchange capacity of six of the samples, and Violet M. Merritt analyzed the same samples for CaO, Na₂O, and K₂O.

References cited

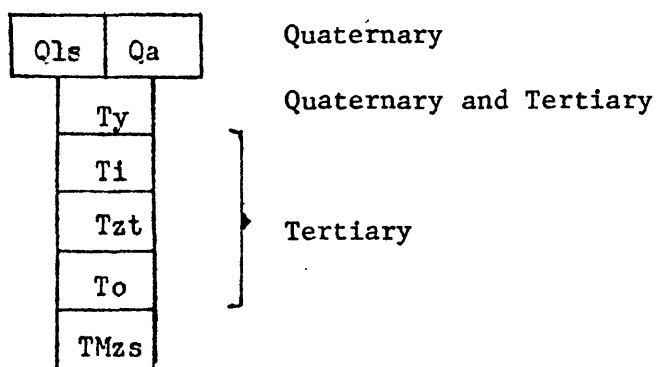
- Cunningham, C. G., and Steven, T. A., 1977, Mount Belknap and RedHills calderas and associated rocks, Marysvale volcanic field, west-central Utah: U.S. Geological Survey Open-file Report 77-568.
- Fleck, R. J., Anderson, J. J., and Rowley, P. D., 1975, Chronology of mid-Tertiary volcanism in the High Plateaus region of Utah: Geological Society of America Special Paper 160, p. 53-60.
- Starkey, H. C., 1964, Determination of the ion-exchange capacity of a zeolitic tuff: U.S. Geological Survey Prof. Paper 475-D, p. 93-D95.
- Steven, T. A., Cunningham, C. G., Naeser, C. W., and Mehnert, H. H., 1977, Revised stratigraphy and radiometric ages of volcanic rocks and mineral deposits in the Marysvale area, west-central Utah: U.S. Geological Survey Open-file Report 77-679.
- Steven, T. A., Rowley, P. D., and Cunningham, C. G., 1978, Geology of the Marysvale volcanic field, west-central Utah: Brigham Young University Geology Studies, v. 25, pt.1, p. 67-70.

Figure 2

(MAP AT BACK)

EXPLANATION

Qa	ALLUVIAL DEPOSITS (QUATERNARY)
Qls	LANDSLIDE DEBRIS (QUATERNARY)
Ty	VOLCANIC UNITS YOUNGER THAN THE ZEOLITIC TUFF (QUATERNARY AND TERTIARY)--Includes basalt flows, Joe Lott Tuff Member of Mount Belknap Volcanics, and Osiris Tuff
Ti	MONZONITE INTRUSIVE ROCK (TERTIARY)
Tzt	ZEOLITIC TUFF UNIT (TERTIARY)
To	VOLCANIC UNITS OLDER THAN THE ZEOLITIC TUFF (TERTIARY)--Includes tuff of Albinus Canyon, Three Creeks Tuff Member, volcanics of Wales Canyon, and volcanics of Dog Valley, all of Bullion Canyon Volcanics
TMzs	UNDIVIDED TERTIARY AND MESOZOIC SEDIMENTARY ROCKS



—•— FAULT--Bar and ball on downthrown side. Dotted where covered

— CONTACT

× SAMPLE LOCALITY