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SOURCE PROVENANCE OF OBSIDIAN CRUCIFORMS FROM THE SOUTHERN U.S. SOUTHWEST

by

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Report Prepared for

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INTRODUCTION

The analysis here of 21 obsidian cruciforms, and one cruciform produced from a volcanic or plutonic rock, from archaeological sites presumably from southern New Mexico and northern Chihuahua indicates a diverse provenance assemblage dominated by the northern Chihuahuan obsidian source at Sierra Fresnal (Shackley 2005). The remainder of the artifacts are produced from one of the sources in the Jemez Mountains of northern New Mexico, but likely procured from Rio Grande Quaternary alluvium (see Church 2000; Shackley 2021). A brief discussion is offered below.

ANALYSIS AND INSTRUMENTATION

 All archaeological samples are analyzed whole. The results presented here are quantitative in that they are derived from "filtered" intensity values ratioed to the appropriate xray continuum regions through a least squares fitting formula rather than plotting the proportions of the net intensities in a ternary system (McCarthy and Schamber 1981; Schamber 1977). Or more essentially, these data through the analysis of international rock standards, allow for inter-instrument comparison with a predictable degree of certainty (Hampel 1984; Shackley 2011).

 All analyses for this study were conducted on a ThermoScientific *Quant'X* EDXRF spectrometer, located at the Geoarchaeological XRF Laboratory, Albuquerque, New Mexico. It is equipped with a thermoelectrically Peltier cooled solid-state Si(Li) X-ray detector, with a 50 kV, 50 W, ultra-high-flux end window bremsstrahlung Rh target X-ray tube and a 76 μ m (3 mil) beryllium (Be) window (air cooled), that runs on a power supply operating from 4-50 kV/0.02- 1.0 mA at 0.02 increments. The spectrometer is equipped with a 200 l min −1 Edwards vacuum pump, allowing for the analysis of lower-atomic-weight elements between sodium (Na) and titanium (Ti). Data acquisition is accomplished with a pulse processor and an analogue-todigital converter. Elemental composition is identified with digital filter background removal, least squares empirical peak deconvolution, gross peak intensities and net peak intensities above background.

Trace Element Analysis

 The analysis for mid Zb condition elements Ti-Nb, Pb, Th, the x-ray tube is operated at 30 kV, using a 0.05 mm (medium) Pd primary beam filter in an air path at 100 seconds livetime to generate x-ray intensity $K\alpha_1$ -line data for elements titanium (Ti), manganese (Mn), iron (as $Fe₂O₃^T$), cobalt (Co), nickel (Ni), copper, (Cu), zinc, (Zn), gallium (Ga), rubidium (Rb), strontium (Sr), yttrium (Y), zirconium (Zr), niobium (Nb), lead (Pb), and thorium (Th). Not all these elements are reported since their values in many volcanic rocks are very low. Trace element intensities were converted to concentration estimates by employing a linear calibration line ratioed to the Compton scatter established for each element from the analysis of international rock standards certified by the National Institute of Standards and Technology (NIST), the US. Geological Survey (USGS), Canadian Centre for Mineral and Energy Technology, and the Centre de Recherches Pétrographiques et Géochimiques in France (Govindaraju 1994). Line fitting is linear (XML) for all elements. When barium (Ba) is analyzed in the High Zb condition, the Rh tube is operated at 50 kV and up to 1.0 mA, ratioed to the bremsstrahlung region (see Davis 2011; Shackley 2011a). Further details concerning the petrological choice of these elements in Southwest obsidians is available in Shackley (1988, 1995, 2005; also Mahood and Stimac 1991; and Hughes and Smith 1993). Nineteen specific pressed powder standards are used for the best fit regression calibration for elements Ti-Nb, Pb, Th, and Ba, and include G-2 (basalt), AGV-2 (andesite), GSP-2 (granodiorite), SY-2 (syenite), BHVO-2 (hawaiite), STM-1 (syenite), QLO-1 (quartz latite), RGM-1 (obsidian), W-2 (diabase),

BIR-1 (basalt), SDC-1 (mica schist), TLM-1 (tonalite), SCO-1 (shale), NOD-A-1 and NOD-P-1 (manganese) all US Geological Survey standards, NIST-278 (obsidian), U.S. National Institute of Standards and Technology, BE-N (basalt) from the Centre de Recherches Pétrographiques et Géochimiques in France, and JR-1 and JR-2 (obsidian) from the Geological Survey of Japan (Govindaraju 1994).

Major and Minor Oxide Analysis

 Analysis of the major oxides of Si, Al, Ca, Fe, K, Mg, Mn, Na, and Ti is performed under the multiple conditions elucidated below. This fundamental parameter analysis (theoretical with standards), while not as accurate as destructive analyses (pressed powder and fusion disks) is usually within a few percent of actual, based on the analysis of USGS RGM-1 obsidian or in this study the USGS AGV-1 andesite standard (see also Shackley 2011b). The fundamental parameters (theoretical) method is run under conditions commensurate with the elements of interest and calibrated with 11 USGS standards (RGM-1, rhyolite; AGV-2, andesite; BHVO-1, hawaiite; BIR-1, basalt; G-2, granite; GSP-2, granodiorite; BCR-2, basalt; W-2, diabase; QLO-1, quartz latite; STM-1, syenite), and one Japanese Geological Survey rhyolite standard (JR-1). See Lundblad et al. (2011) for another set of conditions and methods for oxide analyses. Conditions Of Fundamental Parameter Analysis¹:

Low Za (Na, Mg, Al, Si, P)

Mid Zb (K, Ca, Ti, V, Cr, Mn, Fe)

¹ Multiple conditions designed to ameliorate peak overlap identified with digital filter background removal, least squares empirical peak deconvolution, gross peak intensities and net peak intensities above background. 2 Current is set automatically based on the mass absorption coefficient.

 The data from the WinTrace software were translated directly into Excel for Windows and into SPSS ver. 27 and JMP 12.0.1 for statistical manipulation. The USGS rhyolite standard RGM-1 is analyzed during each sample run of \leq 19 samples for obsidian artifacts to evaluate machine calibration (Table 1). Source assignments were made by reference to source data at http://swxrflab.net/swobsrcs.htm, (see also Shackley 1995, 2005, 2021; Shackley et al. 2018).

DISCUSSION

 Artifacts produced from sources present in the collection while seemingly from a large area of the North American Southwest, are likely all from the southern portion of New Mexico or the northern portion of Chihuahua. All the sources from the Jemez Mountains and Mount Taylor volcanic fields in northern New Mexico are present in the Quaternary alluvium of the Rio Grande as far south as Chihuahua (Church 2000; Shackley 2021). While the primary source for Sierra Fresnal obsidian is the mountain range by the same name in north-central Chihuahua, it is also available as secondary deposits in the Rio Nuevo Casas Grandes nearer the border (Shackley 2005). It is always difficult to determine whether the raw material for a given artifact was procured from the primary source through exchange or directly, the first approximation is that the nearest available source is the most likely, as shown recently in archaeological contexts between Pojoaque, and Las Cruces New Mexico (Shackley 2021).

Sierra Fresnal

 Nearly 50% (47.6%) of the assemblage was produced from the Sierra Fresnal source, a large set of coalesced rhyolite domes near Lago Fresnal and Lago Guzman (see Shackley 2005:83-84; Tables 1 and 2 here). This is one of the few known sources in Chihuahua with a known primary location, and its elemental composition is calc-alkalic and not peralkaline, and appears to not be part of the Sierra Madre volcanic province (Murray et al. 2014; Ferrari et al. 2007; Figure 1 here). The nodules have eroded north at least as far as Nuevo Casas Grandes and into Lago Fredrico to the west of Sierra Fresnal. Five of the nodules collected by Alan Phelps at Lago Fredrico are now considered part of the Sierra Fresnal source. It does occur in various archaeological contexts north of the border, and appears to be one of the major sources in the region (Dolan et al. 2017a, 2017b, 2019; Shackley 2005).

Jemez Lineament Sources

 The four sources from the two Jemez Lineament volcanic fields (Jemez and Mount Taylor) including Cerro Toledo Rhyolite (23.8%), El Rechuelos Rhyolite (9.5%), Canovas Canyon Rhyolite (4.8%), and Grants Ridge at Mount Taylor (4.8%) are commonly recovered in Rio Grande Quaternary alluvium (Church 2000; Shackley 2021; Tables 1 and 2 and Figure 1 here). The Mount Taylor sources only enter the Rio Grande from the Rio Puerco near Socorro, New Mexico. Cerro Toledo Rhyolite obsidian is the most common in Rio Grande gravels due to the violent nature of the Cerro Toledo eruptive event and entering the Rio Grande at multiple points along the river basin (Shackley 2021).

Other Unlocated Source

 Nearly 10% of the artifacts were produced from an as yet unlocated source likely somewhere in Chihuahua (Tables 1 and 2; see Fralick et al. 1998; Figure 1 here). This source has not appeared in archaeological contexts to my knowledge including those north of the border. It does not match any of the 160,000+ sources in the Skinner/Shackley database of North American obsidian sources.

The Silicic Plutonic or Volcanic Rock Sample

 Sample DA9 appears to be a high silica volcanic or plutonic rock (Figure 2). Given the highly polished character it is difficult to determine the exact rock type, but the size of the mineral components argue for a plutonic rock likely granite (see Figure 2). The composition is similar to most grantitoid rocks common in the North American Southwest. It is definitely not a high calcium sediment.

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Table 1. Elemental concentrations and source assignments for the obsidian artifacts and USGS RGM-1 rhyolite standard. All measurements in weight percent (%) or parts per million (ppm) as noted.

Table 2. Frequency distribution of obsidian sources in the assemblage.

Figure 1. Ba/Zr and Zr/Rb bivariate plots of the archaeological samples. Confidence ellipses at 90%.

Figure 2. TAS plot of sample DA9 and USGS RGM-1 rhyolite standard.