CHRYSOPRASE AND PRASEopal FROM
HANETI, CENTRAL TANZANIA

James E. Shigley, Brendan M. Laurs, and Nathan D. Renfro

Commercial quantities of gem-quality chrysoprase and green prase opal (nickel-bearing chalcedony and common opal, respectively) have been recovered from altered serpentinite deposits near Haneti, Tanzania. Material studied for this report came from the largest mine, located on top of Iyobo Mountain, which has been actively exploited for two decades. Although somewhat similar in color and appearance, chrysoprase and prase opal can easily be distinguished from one another by differences in refractive index (~1.55 vs. ~1.45) and specific gravity (~2.60 vs. ~2.11). Chrysoprase makes up the vast majority of the output, and the mine shows good potential for continued production.

Chrysoprase and prase opal are nickel-containing green varieties of chalcedony and non-play-of-color [common] opal, respectively. Both have been used as gem materials for thousands of years. Chrysoprase was described as being the “most prized” type of chalcedony by Webster [1994, p. 233]. Central Tanzania is an important contemporary source of both these gems. The Tanzanian materials were first briefly described by Gübelin [1975, pp. 76–78] and Schmetzer et al. [1976]. These reports were followed by a characterization of the green prase opal by Koivula and Fryer [1984], and a more detailed study of the chrysoprase by Kinnunen and Malisa [1990].

This article briefly describes the geology, mining, and gemological characteristics of high-quality Tanzanian chrysoprase and prase opal from the Iyobo Mountain mine near Haneti (figure 1). This mine is operated by Dimitri Mantheakis, who hosted two of the authors (JES and BML) there in May 2008. Although both materials have also been found in several other parts of the world (table 1), Tanzania is probably the most important source of chrysoprase after Australia.

BACKGROUND

The green color of chrysoprase has long been thought to be caused by one or more submicroscopic green nickel compounds (as Ni$^{2+}$). While this idea has been discussed for more than two centuries, the exact nature of these nickel compounds continues to be debated [Natkaniec-Nowak et al., 1989]. Some researchers suggested it was a nickel oxide (possibly the mineral bunsenite; see Heflik et al., 1989), but subsequent spectroscopic and high-magnification imaging studies disproved this idea [see Rossman, 1994; Gawel et al., 1997]. Most others have ascribed the coloring agent to layered or framework hydrated nickel silicates (such as kerolite) or nickel-containing clay minerals (such as garnierite, lizardite, or saponite; see Rossman, 1994, pp. 458–459; Nagase et al., 1997; Dyrek et al., 2001; Sachanbiński et al., 2001; Sojka et al., 2004).

The nickel compound(s) occur as tiny, colloidal particles that are dispersed as inclusions throughout the host silica matrix. The presence of a small amount of iron (as Fe$^{3+}$) can modify the color of chrysoprase to be more yellowish green, whereas light scattering from microdefects or small particles in the translucent material (the “Tyndall Effect”) has been suggested as the cause of the more bluish green
appearance seen in reflected light (Sachanbiński et al., 2001). Gawel et al. (1997) studied chrysoprase from several localities, and concluded that those samples with a lower degree of crystallinity tended to have higher nickel contents and a more intense green color. Presumably, dispersed particles of nickel-containing minerals also color the green prase opal from Tanzania, since the visible spectra of both materials are similar (see Results section). Schmetzer et al. (1976) described this material as being opal-CT.

Because of their opaque-to-translucent green appearance and ability to take a good polish, both chrysoprase and prase opal have been used as substitutes for jade. Dyed green agate and green glass have occasionally been used to imitate chrysoprase (O’Donoghue, 2006). Also known is a chromium-colored green chalcedony found in Australia, Bolivia, and Zimbabwe (material from the last has been referred to as mtorolite; see Phillips and Brown, 1989; Hyršl and Petrov, 1998; Hyršl, 1999; Willing and Stocklmayer, 2003). An opaque-to-semitranslucent yellow-green carving material from Australia, sold under the name “lemon chrysoprase,” has been described as a rock consisting of magnesite and quartz (Johnson and Koivula, 1996) or of both minerals along with chrysoprase (Henn and Milisenda, 1997). All these other materials have gemological properties that allow them to be distinguished from chrysoprase and prase opal.

OCCURRENCE AND DESCRIPTION OF THE TANZANIAN MATERIAL

Location. Chrysoprase has been found on three adjacent hills that lie ~12 km southeast of the village of Haneti and 12 km north of the village of Itiso. Haneti is 75 km north of Dodoma along Highway A104 (figure 2). The mine operated by Mr. Mantheakis is located near the top of the southernmost of the three hills (known as Iyobo Mountain; figure 3), at 5° 31.90’ S, 35° 59.33’ E and at an elevation of 1,452 m. The area is accessible during the dry season (which runs April through November) by a dirt road from Dodoma to Haneti, and from there to the mine site by a rough dirt track.
**Geology.** According to Kinnunen and Malisa (1990), rocks in the area of the Haneti deposit are of Archean age and consist of metamorphosed ultramafics that form the three chrysoprase-bearing hills. These are aligned in a northwesterly direction, which marks the orientation of regional fracture/shear zones. They described the dominant rock type hosting the chrysoprase as a silicified and ferruginized (silica- and iron-altered) serpentinite.

### Table 1. Other world sources of chrysoprase and prase opal.

<table>
<thead>
<tr>
<th>Locality</th>
<th>Geologic setting</th>
<th>Reference</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Chrysoprase</strong></td>
<td></td>
<td></td>
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<tr>
<td>Warrawanda, Western Australia</td>
<td>Weathered and altered serpentinites associated with granites</td>
<td>Nagase et al. (1997)</td>
</tr>
<tr>
<td>Yerilla Station, Western Australia</td>
<td>Veins in weathered siliceous ironstones</td>
<td>Jones (1994a,b), Brown (2000)</td>
</tr>
<tr>
<td>Niquelandia, Goiás, Brazil</td>
<td>A galena mine (?)</td>
<td>Kammerling et al. (1990)</td>
</tr>
<tr>
<td>Saxony, Germany</td>
<td>Not reported</td>
<td>Witting (2001)</td>
</tr>
<tr>
<td>Niigata Prefecture, Honshu, Japan</td>
<td>Not reported</td>
<td>Mindat.org (2009)</td>
</tr>
<tr>
<td>Sarykul Boldy, Kazakhstan</td>
<td>Nickel deposit in weathered serpentinite</td>
<td>Sachanbin’ski et al. (2001)</td>
</tr>
<tr>
<td>Ambatondrazaka, Madagascar</td>
<td>Not reported</td>
<td>Behler (1963)</td>
</tr>
<tr>
<td>Szkiry, Lower Silesia, Poland</td>
<td>Nickel deposit in weathered serpentinite</td>
<td>Drzymala and Serkies (1973), Niskiewicz (1982), Helflik et al. (1989), Sachanbin’ski et al. (2001)</td>
</tr>
<tr>
<td>Southern Urals, Russia</td>
<td>Not reported</td>
<td>Mikhailov (2000)</td>
</tr>
<tr>
<td>Tulare County, California, United States</td>
<td>Not reported</td>
<td>O’Donoghue (1995)</td>
</tr>
<tr>
<td>Various sites in Arizona, California, Colorado, Massachusetts, New York, North Carolina, Oregon, Rhode Island, and Vermont</td>
<td>Not reported</td>
<td>Mindat.org (2009)</td>
</tr>
<tr>
<td><strong>Prase opal</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>South Bohemia, Czech Republic</td>
<td>Not reported</td>
<td>Duda et al. (1991)</td>
</tr>
<tr>
<td>Silesia, Poland</td>
<td>Not reported</td>
<td>Drymala and Serkies (1973), Webster (1994)</td>
</tr>
<tr>
<td>Southern Urals, Russia</td>
<td>Small vein at a “chrome mine”</td>
<td>Mikhailov (2000)</td>
</tr>
<tr>
<td>Napa County, California, United States</td>
<td>Not reported</td>
<td>“Prase opal in California,” 1996</td>
</tr>
</tbody>
</table>

Both the chrysoprase and prase opal occur in discontinuous veins up to several centimeters thick and several meters long in the weathered host rock. According to Mr. Mantheakis, the chrysoprase veins appear to be more abundant on the western side and upper portions of Iyobo Mountain; some areas have almost no such mineralization. Within a vein and between different veins, the material can vary from high-quality translucent to low-quality opaque. The best chrysoprase occurs in those areas of a vein that are surrounded by red clay or soil, while poorer-quality material is found where the vein is enclosed by yellowish material. The best-quality prase opal is found in veins within very brittle host rock.

### Mining.

Although at one time some Ni-rich magnesite veins were prospected in the region, the only mining in the Haneti area has been for chrysoprase-
and prase opal for both gem and ornamental purposes (Kinnunen and Malisa, 1990). According to Mr. Mantheakis, chrysoprase was first found there in the early 1960s, and limited production took place before the concession was nationalized by the government in 1973, after which the mine lay dormant. Mr. Mantheakis, a Tanzanian citizen, reclaimed the mine in 1986, and in 1997 was given full mining rights on the property.

Mining is currently carried out in an open pit (figure 4) at the top of Iyobo Mountain. As many as 35 local people are employed during the dry season. So far, 28 pits have been mined on the mountain, ranging from ~20 to ~250 m². Mechanized equipment is used to expose areas containing the gem-bearing veins, which are then excavated with hand tools (figure 5). The miners follow the veins until they pinch out (figure 6), and the chrysoprase and prase opal are collected by hand (figure 7) and placed into bags for transport to a sorting facility on site.

The chrysoprase production is highly variable, ranging from as little as 100 kg to as much as 1,000 kg per month; the typical annual production is 7–10 tonnes. These quantities are for “mine-run” rough, which includes some chrysoprase in matrix. After sorting and trimming, ~20–30% of the chrysoprase is marketable, but only 3–5% is of high quality [i.e., translucent, with even, intense color, and suitable for cutting attractive cabochons]. The prase opal is much less abundant: In the past decade it has been recovered from only two areas of the mine, which yielded a total of ~1.5 tonnes of opal in matrix.

The better-quality material is sent to Dar es Salaam for trimming and polishing into cabochons. Some of the lower-quality rough is worked into beads. The material is sold through wholesale and retail outlets in Dar es Salaam, as well as through international trade shows.

Figure 3. The mine workings are situated on top of Iyobo Mountain and are marked by a row of tailings. Photo by B. M. Laurs.
Mr. Mantheakis (pers. comm., 2008) reported that his mining plan is sensitive to the local environment. He places roads to preserve as many of the large trees as possible, and most of the pits are buried as mining proceeds. He estimates that 80% of the deposit remains to be mined.

**MATERIALS AND METHODS**

For this study, we examined nine chrysoprase samples (five cabochons [3.72–18.42 ct] and four pre-formed or rough pieces weighing up to 8.6 g), and five prase opal samples (one 35.82 ct cabochon and four pre-formed or rough pieces weighing up to 7 g). All are representative of the better-quality material being produced from Mr. Mantheakis’s mine. We documented all the samples with a refractometer, gemological microscope, polariscope, desk-model spectroscope, Chelsea color filter, and standard ultraviolet (UV) lamps.

All the samples were also analyzed by each of the following methods. Qualitative chemical analyses were performed using a Thermo ARL Quant-X energy-dispersive X-ray fluorescence (EDXRF) system with multiple filter, voltage, and current settings appropriate to the elements of interest. Ultraviolet-visible–near infrared (UV-Vis-NIR) absorption spectra were recorded over an interval of 250–2500 nm with a Perkin Elmer Lambda 950 spectrometer, using a slit width of 2 nm and a data sampling interval of 0.5 mm. (Only the 400–700 nm region is shown in the results, since this is the region of greatest interest for investigating the origin of color.) Infrared spectra were recorded over the 6000–400 cm\(^{-1}\) range with a Thermo Nicolet Magna-IR 760 spectrometer, using a 6× beam condenser at a resolution of 4 cm\(^{-1}\) and 128 scans per sample to improve the signal-to-noise ratio. We obtained Raman spectra for both materials using a Renishaw InVia Raman microscope with 514.5 nm laser excitation.
RESULTS AND DISCUSSION

Gemological Characteristics. Samples of both materials were light to medium-dark slightly bluish green and opaque to translucent; all displayed a vitreous polish luster. In general, the prase opal was slightly more translucent than the chrysoprase. The color distribution in both materials appeared uniform or very slightly blotchy. In addition, the prase opal displayed small, dark, granular brownish areas along fractures that Raman analysis indicated were most likely goethite; each of these areas tended to be surrounded by a small halo where the adjacent prase opal was yellower (figure 8). The observation that the outline of the halos followed the shape of the dark areas suggests that iron from these inclusions permeated the surrounding prase opal.

The RI and SG values of all tested samples fell within the following ranges, which readily separate the two materials from the Haneti locality:

<table>
<thead>
<tr>
<th>Material</th>
<th>RI</th>
<th>SG</th>
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<tr>
<td>Chrysoprase</td>
<td>1.549 (+0.002)</td>
<td>2.57–2.63</td>
</tr>
<tr>
<td>Prase opal</td>
<td>1.455 (+0.004)</td>
<td>2.09–2.13</td>
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</table>

Kinnunen and Malisa (1990) reported similar RI and SG values for the chrysoprase from the Haneti area. These RI values are slightly above those of chalcedony in general, but the SG ranges are consistent with those reported for chrysoprase in the literature (Webster, 1994, pp. 252–253). We could not find properties for prase opal in the literature, but the RI and SG values of our samples fell within the ranges given for opal in general by O’Donoghue (2006).

All samples were inert to both long- and short-wave UV radiation. They exhibited an aggregate reaction when viewed with a polariscope; this reaction was more pronounced for the prase opal. Neither material displayed any reaction when viewed with a Chelsea color filter (i.e., all samples remained green). The desk-model spectroscope revealed broad absorption below ~450 nm and above ~650 nm in both materials. These two regions of absorption were more intense, and the intervening region of light transmission narrowed in wavelength range, for the more saturated green samples of both materials. No sharp absorption lines were observed in the spectra of either material.

The gemological properties of both the chrysoprase and prase opal were generally consistent with reports in the literature for samples from the Haneti area and elsewhere. Kinnunen and Malisa (1990) characterized the chrysoprase as displaying a distinct microtexture consisting of disordered silica spherules (~40–80 μm diameter), formed of concentric layers of quartz, chalcedony, and opal-A, within a silica...
groundmass. In contrast, they found that the silica spherules in the Tanzanian prase opal were an order of magnitude smaller (~5–6 μm). They described the chrysoprase as also containing “cloudy color distributions, clear chalcedony veinlets, whitish dots [microcavities], small fluid inclusions in bipyramidal quartz, and brownish inclusions of host rock” (p. 162) based on microscopic observations (10×–60× magnification and various lighting conditions). They interpreted the lack of a vapor phase in the liquid inclusions as implying that the chrysoprase formed at a very low temperature and was deposited by repeated evaporation of Si-rich hydrothermal solutions or surface waters along open fractures in the altered serpentinite host rock.

**Chemical Analysis.** Our EDXRF analyses indicated major amounts of Si and minor Ni, Zn, and Fe in both materials. Using atomic absorption spectroscopy, Kinnunen and Malisa (1990) measured 0.55 wt. % Ni in the chrysoprase, along with trace amounts of other elements (including the transition metals Co, Zn, Fe, Mn, and Cu at values up to 120 ppm, and rare-earth elements at 1 ppm or less).

**Spectroscopy.** The visible-range spectra of both materials showed increasing absorption below about 500 nm and a broad peak centered near 650 nm (figure 9), corresponding to the absorption patterns seen with the spectroscope (see also Schmetzer et al., 1976; Sachanbiński et al., 2001). This is consistent with the spectrum for chrysoprase from Tulare County, California (Rossman, 2009). The region of absorption above 650 nm has been attributed to Ni$^{2+}$ in octahedral coordination in a silicate mineral (Rossman, 1994, pp. 458–459; there is no indication that this element substitutes for either silicon or oxygen in either the chrysoprase or the prase opal).

The near-infrared spectra were similar for both materials (figure 10), with a broad band at ~5245 cm$^{-1}$ (more intense in the prase opal) and a group of similar features at 4517, 4326, and 4300 cm$^{-1}$ due to the presence of water molecules or hydroxyl groups in both materials (similar spectra for opals are illustrated in Langer and Flörke, 1974).

In contrast, the Raman spectra of the two materials were very different from one another (figure 11). In the chrysoprase, there were sharp Raman peaks at 1160, 807, 463 [much more intense than the other peaks], 398, 354, and 264 cm$^{-1}$. All of these features were reported previously in a study of the Raman spectra of microcrystalline silica [including chalcedony] by Kingma and Hemley (1994). The Raman spectra of the prase opal exhibited peaks at 783, 671, and a broader feature at 325 cm$^{-1}$. Similar features at ~800–780 and ~325 cm$^{-1}$ were reported in opals by Ostrooumov et al. (1999) and by Smallwood (2000).
CONCLUSION

Jewelry-quality chrysoprase and prase opal (figure 12) are mined from altered serpentinite near Haneti, Tanzania. The gemological properties of both materials are generally consistent with previous reports for chrysoprase and opal, and we did not find any features that distinguish them from the same materials from other localities. Chrysoprase can be separated from chrome chalcedony by the latter’s yellow UV fluorescence, red Chelsea color filter reaction, and sharp absorption line at 684 nm seen with a spectroscopic (Hyrsl, 1999). Although similar in color and visual appearance, chrysoprase and prase opal from the Haneti area can be readily distinguished from one another on the basis of RI and SG values, as well as by their Raman spectra. Field observations of the mining site on Iyobo Mountain suggest good potential for future production.

Figure 11. The Raman spectra for the chrysoprase and prase opal differ significantly.

Figure 12. These Tanzanian prase opal samples consist of a 10.22 ct cabochon and a 5.15 ct faceted stone, together with a vein of rough material (GIA Collection nos. 32590, 32589, and 32587, respectively; photo by Robert Weldon). Tanzanian chrysoprase has been set into a variety of jewelry styles, as shown by the earrings in the inset (~9 ct total weight; courtesy of Dimitri Mantheakis).

ABOUT THE AUTHORS

Dr. Shigley is distinguished research fellow, and Mr. Renfro is a staff gemologist, at the GIA Laboratory in Carlsbad, California. Mr. Laurs is editor of Gems & Gemology at GIA in Carlsbad.

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