NOTES AND NEW TECHNIQUES

GEM-QUALITY HAÜYNE FROM THE EIFEL DISTRICT, GERMANY

By Lore Kiefert and H. A. Hänni

Haüyne is a rare mineral and an extremely rare gemstone. Recently, the authors studied a large number of faceted haüynes from the Eifel district of Germany. The R.I. and S.G. data were consistent with those reported in the literature, and the samples’ identity was confirmed by Raman spectrometry, with the key maxima at 543 and 988 cm⁻¹. EDXRF analyses revealed potassium and iron, as well as the major and minor elements expected in haüyne. Although mineral inclusions were uncommon, apatite and augite were identified, and negative crystals (often surrounded by healed fractures) were seen in approximately one-third of the stones. Short needles and fine, dust-like particles were present in about half the samples. Paraffin wax was identified in some open fissures.

In the summer of 1999, the authors were surprised to receive 100 faceted haüynes (pronounced “how-een”) for analysis. The client who submitted these stones subsequently fashioned most of them into a brooch set with diamonds and a pink sapphire (figure 1). This butterfly brooch sold at the Sotheby’s November 1999 Geneva auction for 45,000 SFr (approximately US$30,000).

In spite of its attractive color, however, haüyne is rarely seen in jewelry. Not only is the mineral itself rare, but it also has a relatively low hardness [5.5–6 on Mohs scale].

Transparent haüyne primarily occurs as small crystals [see also Mertens, 1984] of an unusual bright “apatite” to “sapphire” blue color. Mineralogically, it is a feldspathoid that belongs in the sodalite group (which also includes sodalite, lazurite, and nosean), and is often one of the components of lapis lazuli. The chemical formula for haüyne is ideally [Na,Ca]₄₋₈Al₆Si₆(O,S)₂₄(SO₄,Cl)₁₋₂ (Mandarino, 1999). The crystal system for this silicate is cubic, the crystal class is 4₃m. Cleavage planes are distinct in the [110] direction, and twinning is common along [111]. Haüyne has been reported as white to gray, green, yellow, and red (Arem, 1987), but only the blue color has been noted thus far as faceted material.

GEOLOGY AND OCCURRENCE

Haüyne is found in association with alkaline volcanic rocks (mainly phonolites, which are composed of alkali feldspar, mafic minerals, and felds-
pathoids; figure 2). It has been reported from many countries, including the U.S., Canada, France, Italy, Spain (Tenerife), Morocco, and Germany (Arem, 1987). In Germany, it occurs in relative abundance near Laacher See in the Eifel Mountains as gem-quality crystals of an unusual blue color and transparency. Gem-quality material has not been reported from other sources (Fischer and Bürger, 1976; Arem, 1987).

In the Eifel Mountains, haüyne formed in a magma chamber approximately 2–4 km below the surface, together with a suite of minerals including sanidine, nosean, nepheline, leucite, plagioclase, amphibole, augite, magnetite, titanite, phlogopite, apatite, and olivine (Matthes, 1983; Wörner and Schmincke, 1984). This phonolitic magma was volatile-rich and chemically zoned, with a strong decrease in sulfur during progressive magmatic differentiation, which is interpreted to be partially caused by crystallization of haüyne (Harms and Schmincke, 2000). This magma erupted approximately 12,900 years ago, and the resulting volcanic rocks were deposited in three zoned layers. The bottom layer, which corresponds to the top of the magma chamber, is relatively crystal-poor and consists of a nearly aphyric, highly differentiated phonolite. The top layer, which transported the contents of the bottom of the magma chamber with crystal enrichment, consists of a relatively crystal-rich mafic phonolite. Haüyne is found throughout all three layers (Schmincke, 2000; Harms and Schmincke, 2000).

We know of no attempts to mine haüyne commercially. Most of the crystals are found by amateur collectors (see, e.g., Linde, 1998). In the Eifel district,
The majority of the stones come from a commercial pumice mine. Whenever a new layer is blasted from the high wall of pumice stone, collectors arrive to search for these rare blue crystals. On the market, even the rough stones are sold by carat weight rather than by grams (C. Wild, pers. comm., 2000).

MATERIALS AND METHODS
The 100 faceted round and oval haüynes that were submitted to our laboratory for examination (figure 3) ranged from 0.095 to 0.173 ct (and from 3.11–3.16 × 2.02 mm to 3.68–3.74 × 2.56 mm for the round stones; 3.50 × 2.81 × 1.66 mm to 4.15 × 3.92 × 2.20 mm for the ovals). Additional haüynes from the Eifel district were supplied by the companies Gebrüder Bank and W. Constantin Wild, both of Idar-Oberstein, Germany: approximately 80 crystal fragments between 0.01 and 0.15 ct, 15 small faceted stones (0.02–0.10 ct), and six larger faceted stones (0.15–0.83 ct). A 1.5 mm [diameter] crystal in pumice matrix (again, see figure 2) and a faceted haüyne of 0.15 ct from the SSEF collection completed the samples. All haüyne in this study was mined in the Eifel district (C. Wild and G. Bank, pers. comm., 2000).

We measured the refractive indices of 10 samples with an Eickhorst GemLED refractometer with an LED monochromatic light source (equivalent to NaD light, \( \lambda = 589 \) nm). Specific gravity was determined hydrostatically for 31 faceted samples. For all samples, we observed reaction to long-wave (365 nm) and short-wave (254 nm) ultraviolet radiation in a darkened room. Internal features of all samples were examined using a standard gemological microscope in conjunction with brightfield, darkfield, and oblique fiber-optic illumination. Photomicrographs were taken with a Wild M8/MPS55 stereozoom microscope.

We recorded visible-range spectra for three faceted samples of variable color intensity using a Hitachi U4001 spectrophotometer in the 290–800 nm range. We recorded infrared spectra for three stones in the range 500–6000 cm\(^{-1}\) with a Philips PU 9800 Fourier transform infrared (FTIR) spectrometer to check for any treatment. Qualitative chemical analyses by energy dispersive X-ray fluorescence (EDXRF) of four faceted samples were performed using a Tracor Spectrace 5000 instrument, optimized for the detection of medium-weight [Ca-Ga] elements. We recorded Raman spectra on four samples (three faceted and one crystal), plus the

<table>
<thead>
<tr>
<th>Ideal formula</th>
<th>((\text{Na,Ca})_4\text{Al}_6\text{Si}<em>6(O,\text{S})</em>{24}(\text{SO}<em>4,\text{Cl})</em>{1–2})</th>
</tr>
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<tr>
<td>Color</td>
<td>Light blue to dark blue</td>
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<tr>
<td>Clarity</td>
<td>Transparent to translucent</td>
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<tr>
<td>Refractive index</td>
<td>1.498–1.507</td>
</tr>
<tr>
<td>Birefringence</td>
<td>None, sometimes slight anomalous birefringence</td>
</tr>
<tr>
<td>Optical character</td>
<td>Isotropic</td>
</tr>
<tr>
<td>Specific gravity</td>
<td>2.46–2.48</td>
</tr>
<tr>
<td>Hardness</td>
<td>5.5–6 (reported)</td>
</tr>
<tr>
<td>UV fluorescence</td>
<td>Inert to bright orange</td>
</tr>
<tr>
<td>Long-wave (365 nm)</td>
<td>Inert to slightly red</td>
</tr>
<tr>
<td>Short-wave (254 nm)</td>
<td>Apatite, augite, unidentified small needles and particles, unidentified dark opaque hexagonal mineral, negative crystals, partially healed fissures, and fissures treated with paraffin wax</td>
</tr>
<tr>
<td>Inclusions</td>
<td>UV-Vis absorption Spectroscope: Broad weak band in the yellow range</td>
</tr>
<tr>
<td></td>
<td>Spectrophotometer: Broad band at 600 nm, small band at 380 nm, absorption edge at 300 nm</td>
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<tr>
<td></td>
<td>FTIR spectral features Major absorption bands at 3593 and 3697 cm(^{-1})</td>
</tr>
<tr>
<td></td>
<td>Raman spectral features Major peaks at 440, 543, 988, 1089, and 1635 cm(^{-1})</td>
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*From Arem (1987).
inclusions in several stones, with a Renishaw Raman System 1000 spectrometer equipped with a CCD Peltier detector and an argon ion laser (514 nm) with a power of 25 mW.

RESULTS AND DISCUSSION

The properties determined for these samples are summarized in table 1 and discussed below.

Physical Properties. The samples ranged from a light blue similar to that of Paraiba tourmaline or apatite, to a dark blue similar to that described for fine Kashmir sapphire (figure 4). Most of the samples, however, were an evenly distributed medium blue (see, e.g., figure 3 and the third and fourth stones from the left in figure 4). We did not observe color zoning in any of our samples.

The samples were very consistent in specific gravity (2.46–2.48), which corresponds to the range of 2.40–2.50 reported in the literature (Arem, 1987; Deer et al., 1992). Refractive index results, between 1.498 and 1.507, also were consistent with the range cited in the literature (1.490–1.508; Bank, 1977, 1978–1979; Arem, 1987; Deer et al., 1992).

Approximately one-third of the 100 samples submitted to our laboratory (and a smaller proportion of the other samples) showed orange fluorescence to long-wave UV radiation; the remaining samples were inert. When fluorescence is observed, it is considered characteristic of haüyne from the Eifel district (Webster, 1994). The inconsistency in fluorescence reaction that we observed was also mentioned by Bank (1977). Most of the haüynes showed a very weak reddish fluorescence to short-wave UV.

Microscopic Properties. In approximately half of the samples, we observed short needles and fine dust-like particles arranged in lines, similar to rutile needles or partially dissolved rutile in sapphires (figure 5). Elongate, transparent, prismatic crystals (figure 6), identified as apatite by Raman microspectrometry, were seen in two samples. In contrast to our findings, Wörner and Schmincke (1984) stated that apatite never occurs in haüyne from the Eifel district. Another mineral, which was exposed at the surface of one of the samples shown in figure 4, was identified as augite, which is also a common xenocryst in the Eifel district phonolite (Wörner and Schmincke, 1984). A dark opaque hexagonal crystal exposed at the upper left in figure 6.
the surface of another stone could not be identified (figure 7). This inclusion may be metallic or too decomposed to give a useful Raman signal.

Negative crystals were observed in a third of the stones; some were well formed (figure 8), and some were rounded with a frosted surface (figure 9). They were frequently surrounded by healing fissures that resembled those seen around negative crystals in sapphires from Sri Lanka. In some cases, however, the partially healed fissures had an appearance similar to that of glass fillings in rubies (figure 10). Glass inclusions outlining growth zones in haüyne from the Eifel district were described by Harms and Schmincke (2000). Five of the samples examined contained remnants of the host rock [see, e.g., figure 6]. Where the rough surface of the original crystal face was still visible, corrosion was evident.

Fissures filled with an oily to waxy substance were common in many of the 100 haüynes that were originally submitted to the laboratory for identification. This substance was identified as paraffin wax by FTIR and Raman analyses [see below]. Because haüyne has a low refractive index, which is close to that of paraffin wax or oil, the filled frac-

Figure 9. This rounded negative crystal has a frosted surface and, like the smaller negative crystal on the lower right, a rosette-like healing fissure. Photomicrograph by H. A. Hänni; magnified 40×.

Figure 10. In a number of the haüynes, the partially healed fissures resembled the glass fillings seen in some rubies. Photomicrograph by H. A. Hänni; magnified 30×.
features sometimes showed orange to pinkish flashes; these resembled the orange flashes observed in emeralds with resin-filled fractures (Kiefert et al., 1999; Johnson et al., 1999).

Spectral Features. UV-Vis spectrometry showed a major absorption band centered at approximately 600 nm, maximum transmission at 476 nm (figure 11), a small absorption band centered at 380 nm, and an absorption edge at 300 nm in the three stones tested. This is in agreement with the absorption spectrum reported for haüyne by Henn and Bank (1990). Note in figure 11 that the absorption band at 600 nm increases in intensity with increasing depth of color, while the other spectroscopic features remain the same.

An absorption band at 600 nm, measured with electron paramagnetic resonance (EPR) spectroscopy, also has been described for sodalite and lazurite. This feature was ascribed to a color center associated with $S_2^-$ (Marfunin, 1979), which has been attributed to radiation damage (Vassilikou-Dova and Lehmann, 1990) and may be responsible for the blue color. Henn and Bank (1990) relate the orange fluorescence of this mineral group to the presence of $S_2^-$, which causes the 380 nm absorption band. Note that only a weak, broad absorption band in the yellow region is visible with a handheld type of spectroscope.

The FTIR spectra in the region between 4000 and 2400 cm$^{-1}$ of two stones in which magnification had revealed evidence of a waxy filler showed two major groups of peaks (figure 12): One is typical for haüyne and lies between 3000 and 3800 cm$^{-1}$ (comparison with a “clean” sample); the other group of peaks (between 2840 and 2960 cm$^{-1}$) is characteristic of paraffin wax. This latter group is attributable to the artificial filling of fissures with wax.

Chemical Properties. As noted above, the ideal chemical formula of haüyne is $[\text{Na, Ca}]_{4-8} \text{Al}_6\text{Si}_6\left(\text{O, S}\right)_{24}\left(\text{SO}_4, \text{Cl}\right)_{1-2}$. Qualitative EDXRF chemical analysis of four stones showed, besides the detectable elements given in this formula (i.e., calcium, silicon, aluminum, and sulfur), small but significant amounts of potassium [K] and iron [Fe], as illustrated in figure 13.

According to Wörner and Schmincke (1984), it is difficult to perform microprobe analysis of haüyne because of the large sodium content, the decomposition of the haüyne under the electron beam, and the problem in assigning SO$_3$ (as analyzed) to SO$_4$ and S. Therefore, those authors considered their microprobe analyses of seven samples to be qualitative at best. The elements measured by our qualitative EDXRF analysis (including the significant amounts of K and Fe) are consistent with the chemical data provided by these and other researchers for haüyne from Laacher See and from Italy (see also Deer et al., 1963; Xu and Veblen, 1995; Sapozhnikov et al., 1997).

Figure 12. Shown here are the FTIR spectra measured between 4000 and 2400 cm$^{-1}$ of (A) haüyne, (B) wax-treated haüyne, and (C) paraffin wax. The peaks at 3697 and 3593 cm$^{-1}$ are characteristic for haüyne.
Raman Spectrometry. Raman analysis was performed on the haüyne itself and on all inclusions that were large enough to analyze. The Raman spectra of the four haüyne samples tested (see, e.g., figure 14) are in agreement with the results given by Maestrati (1989), but they differ significantly from those provided in the Renishaw Raman database.

Raman analysis identified both apatite and augite as inclusions in some of the samples. In addition, in one of the several samples that revealed fissure fillings with magnification, the filler was exposed on the surface of the stone, so we were able to get a particularly good Raman signal. The pattern matched that of paraffin wax (figure 14), which confirmed the results recorded with the FTIR spectrometer (again, see figure 12).

Separation from Possible Imitations. The bright blue color of haüyne might be duplicated by cobalt glass, blue cubic zirconia, or cobalt spinel. Lazurite and blue apatite also may resemble haüyne. Careful determination of the optic character (haüyne is cubic), R.L., S.G., and absorption spectrum will, however, enable a firm identification of haüyne. This mineral is clearly distinguishable from other minerals of the sodalite group by its Raman spectrum.

CONCLUSIONS
Although usually considered a collector’s stone because of its rarity, haüyne may be found in expensive jewelry, as was the case with several of the stones tested for this article (again, see figure 1). The use of haüyne in a brooch is appropriate given its low hardness.

The geology and petrology of the deposit at Laacher See (Eifel district) in Germany have been well studied. It is interesting to note that similar volcanic deposits containing haüyne are rare, and we know of no other deposits of the blue gem-quality material. Although the German source has been productive for a long time—haüyne was first described from Germany in 1807 (Clark, 1993)—an accumulation of 100 faceted stones is surprising due to the relative scarcity of the material.

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REFERENCES


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