Gem-quality rubies and fancy-color sapphires have been recovered from dolomite marble lenses located high in the Himalayan mountains of east-central Nepal (Ganesh Himal). First discovered in the early 1980s, these deposits have had only limited and sporadic production because of their isolated locations, high altitudes, and harsh seasonal weather conditions. The precise locations and geology of the two best-known deposits are described. In addition, 27 polished samples and 19 rough crystals and mineral specimens were thoroughly investigated to document their gemological characteristics and crystal morphology. The most distinctive internal features are their color-zoning characteristics, various cloud-like and other patterns, as well as a wide variety of mineral inclusions, some of which have not been seen in rubies from other sources.

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Gems & Gemology, Vol. 33, No. 1, pp. 24–41
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Nepal is a small mountainous country situated along the Himalayan mountain chain in southern Asia. Landlocked by Tibet (China) to the north and India to the east, south, and west, Nepal’s surface area covers only 140,000 km²; it has approximately 20 million inhabitants. Southern Nepal is the birthplace, more than 2,500 years ago, of Siddartha Gautama, the founder of Buddhism. For many centuries, Nepal was closed off from the rest of the world. Only since it emerged from a feudal state in 1950, have the minerals and gemstones of Nepal begun to enter the world economy. Although Nepal has a reigning monarch, a multi-party democracy was established in 1990.

Most widely recognized for having the highest elevation of any country in the world, Nepal is the home of Mount Everest (8,848 m/29,028 feet above sea level). In all, seven of the eight highest peaks in the world can be found in this country’s rugged terrain. Although Nepal has limited industrial development and relatively meager mineral wealth, it is blessed with a broad range of gem minerals that includes tourmaline, beryl, garnet, quartz, spinel, danburite, hambegrite, kyanite, apatite, sodalite, zircon, sphalerite, epidote, diopside, iolite, feldspar, pyrite, hematite,andalusite, lepidolite, and corundum (Bassett, 1979, 1984, 1985a and b, 1987). Gems have been recovered in Nepal since 1934, when tourmaline and aquamarine were first discovered there (Bassett, 1979). However, not until the early 1980s did rubies and sapphires begin to appear (figure 1).

As the story was told to authors AMB and MNM, goat-herders of the Tamang ethnic group first spotted red crystals in 1981, high in the Ganesh Himal massif of the Dhading District in east-central Nepal. These stones were brought to the capital city (Katmandu) to sell for shop displays, where they were identified as rubies. The first official report of corundum in Nepal was made by Baba (1982), who erroneously described
rubies and fancy-color sapphires have been recovered in the Himalayan Mountains of east-central Nepal. The Nepalese rubies shown here range from 0.50 to 2.50 ct. Jewelry courtesy of the Gold Rush, Northridge, California; photo by Shane F. McClure.

None of these mining ventures lasted long, however, because of the isolated locations, harsh weather conditions, and difficulties encountered in efforts to develop larger-scale mining. All of the claims were eventually abandoned by their owners. Today, local Tamang farmers continue rudimentary mining; they bring their production to Katmandu for sale and distribution. As a result, throughout the mid- to late 1980s and early 1990s, the production of gem-quality rubies and fancy-color sapphires was sporadic. Although Himalayan Gems Nepal also ceased active mining after about two years, they continue to buy gems from the local farmers and cut them in their factory.

Since mid-1996, two other deposits—Shelghar and Shongla—have produced corundum that is comparable in quantity and quality to Chumar and Ruyil. Himalayan Gems Nepal began officially recognized mining activities during 1985. They also set up a cutting factory to bring these rubies and fancy-color sapphires to the gem and jewelry markets. Several other parties also mined unofficially in the surrounding region.
but they have not yet been mined. However, Chumar and Ruyil remain the most recognized deposits of commercial-grade and finer quality rubies and pink sapphires in Nepal.

One of the authors (AMB) was told that blue sapphires are currently being recovered from the Taplejung District of eastern Nepal. The authors have examined a few non-gem-quality specimens, but there is no additional information on the amount or quality of the material being unearthed. The claim holder reports, however, that the sapphire crystals occur in outcrops on the slopes of the mountain known as Sapphire Hill, approximately 1.5 km northwest of the small village called Khupatal, north of the Tamur River in the Taplejung District. Contrary to Baba's 1982 report, our investigations yielded no evidence that rubies have ever been found in this district.

The first in-depth study of ruby from Nepal was provided by Harding and Scarratt (1986). Soon after, researchers in Germany also described fancy-color (pink and violet) sapphires from unspecified deposits in Nepal (Kiefert and Schmetzer, 1986, 1987). A detailed investigation of one very fine gem-quality ruby from Nepal was reported in 1988 (Bank et al., 1988). Robinson et al. (1992) were the first to mention a specific deposit, named Chumar, for the corundum occurrences. Overviews on gems from Nepal also described corundum deposits [e.g., Niedermayr et al., 1993]. The most recent observations on the internal features of this material were provided by Henn and Milisenda (1994). The present study supports the findings of these previous researchers, and adds both new characteristics and specific source information to these earlier reports.

**LOCATION AND ACCESS**

The corundum deposits occur on the southwest flank of the Ganesh Himal (a high mountain massif consisting of several major peaks clustered together), 68 km north-northwest of the capital city of Katmandu, and 40 km west of Trisuli Bazar. An overview of the region reveals rice paddies in the lowlands, with primarily corn and barley crops in the hills. Above approximately 1,830 m (6,000 feet), one finds forests of rhododendron trees, then grass, and, eventually, bare rock. The landscape at the deposits is that of extremely steep, cliff-like domes of dolomitic marble about 150 m (500 feet) high lying in lenticular bodies that are surrounded by steep slopes of schist. There are deeply entrenched, rocky gorges, choked with dense vegetation that
reaches almost to the corundum deposits, which crop out in the dangerously craggy cliffs.

Travel to the deposits from Katmandu usually takes about six days. The trek begins with a circuitous bus ride from Katmandu to Trisuli Bazar, and then five days of rather strenuous hiking up the Ankhu River Valley and its tributaries. Travel to the deposits is hampered not only by the high elevations, remote locations, and rugged terrain, but also by the severe seasonal weather conditions. Winter snow usually covers the higher trails from mid-November to the end of April, and monsoons buffet the countryside from mid-June through September. Therefore, the best times to reach the deposits are usually from early May to mid-June and from early October to mid-November. A knowledgeable guide is essential.

Both the Chumar and Ruyil deposits lie within what was formerly known as the Laba Panchayat of the northern Dhading District, Bagmati Zone, in central Nepal (figure 2). The Chumar deposit lies at 28° 13' 20" N, 84° 58' 52" E, at an altitude of 3,800 m (12,500 feet); it is approximately 1.2 km south of the Mandra Danda mountain peak (elevation 4,358 m/14,300 feet), near the Tamang village of Burang. The Ruyil deposit lies about 6 km northeast of Chumar, at 28° 15' 05" N, 85° 02' 07" E, and an elevation of about 4,200 m (13,800 feet).

**GEOLOGY AND OCCURRENCE**

The geology of Nepal is dominated by the Himalayan Mountains, which formed as a result of the collision between the northward-drifting Indian plate and the Asian plate starting about 30 to 50 million years ago. The initial line of collision (called the Indus Suture Zone) is now in Tibet. The Indian plate continued in a northward drift, pushing beneath the Asian plate, whereupon the Indian plate broke into northern and southern segments, as the southern segment was forced beneath the northern segment. This major structural breakage is known as the Main Central Thrust (MCT) and extends east-west, the full length of Nepal (figure 3). At least twice subsequently, the continuing northward drift of the Indian plate caused a major breakage along progressively younger and more southerly thrusts, known as the Mahabharat Thrust and the Main Boundary Thrust (MBT). Each continental slice was pushed beneath the higher and more northerly slices of the Indian plate to form the imbricate (shingle-like) stack that is responsible for the elevation of the Himalayan Mountains. Each of these various continental slices has a different lithology and grade of metamorphism; yet they are all roughly the same geologic age (Precambrian and lower Paleozoic), with the highest grade above the MCT and lower grades below the MCT [there is no metamorphism below the MBT]. This sequence is inverted from what is normally encountered, because of the subduction thrusts.

The Ganesh Himal corundum occurs near the top of the lower-grade metasediments of the Nawakot Series (again, see figure 3), below the MCT. The gems are found in what was a limestone formation (the Malekhu Limestone, lower Paleozoic) that has

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Figure 3. This simplified geotectonic map of Nepal indicates the main geologic structures (tectonic zones) and the main rock-forming series and groups in Nepal. The study area is near the top of the Nawakot series, below the MCT. Source: Mineral Resources Map of Nepal, United Nations publication, 1993.
been converted to a dolomite marble because of its proximity to the tectonic forces of the MCT. This once-continuous dolomite marble layer has been torn into a dozen isolated bodies, about 1 km from one another. These dolomite bodies occur as thickened beds and isolated pods 60–150 m thick and up to a kilometer long. The bodies are separated by black schist with quartzite interlayers.

Within these isolated dolomite pods are mineral seams that were originally aluminous clay interlayers in the limestone, these have been converted, again due to the intense shearing pressures of the MCT, into a suite of metamorphic minerals that includes corundum. Each seam has a slightly different suite of minerals, depending on variations in the original composition of the clay interlayers. In one notable variation (the Ruyil deposit), abundant graphite has formed, presumably due to a local abundance of marine organic material in the original interlayer. The dolomite marble host rock is consistent throughout the dozen pods.

At the western end of these dozen dolomite bodies, which extend for 15 km, they are more distinctively pod-like lenses, in the central section, the dolomite bodies are more like a thickened stratigraphic bed with parallel walls less than 100 m thick. The continuation eastward has not been explored in the very deep, densely forested chasms, but corundum is known to occur at the Sublay claim at the far eastern end. Of the dozen dolomite bodies, the westernmost four—Pola, Chumar, Shongla, and Shelghar—have produced corundum, only Ruyil in the central part, and only Sublay at the eastern end, are corundum bearing. Of these six corundum-bearing bodies, Chum and Ruyil have been the most productive, with Shongla and Shelghar only recently developed and Pola and Sublay not yet consistently worked. This study focuses primarily on the Chumar and Ruyil deposits.

Chumar Deposit. First studied in 1984 by AMB, the Chumar deposit extends east-west for about 550 m, with a central thickness of approximately 150 m (figure 4). At the west end, the dolomitic body is bounded by a sharp fault trending N60°W, dipping steeply to the northeast. The east end of the body tapers down to a tail, with no fault line at the boundary. Distinct bedding in the dolomite body, which now consists of large eroded domes, strikes east-west throughout all but the extremities, where it is bent to the northwest, creating a sigmoid shape. The seams, which are about 8 m apart and dip 32°, vary widely in thickness but average about 20 cm; they are sometimes folded on a small scale.

Ruyil Deposit. The first studies of the Ruyil deposit were conducted in 1985 by AMB. This deposit extends east-west for approximately 128 m, with a general thickness of 60 m. The western portion of the dolomite body appears to end abruptly in a curved outline of shattered dolomite without distinct bedding, yet there is no discernible fault line. No corundum has been found in the continuous outcrops of bedded dolomite on the eastern side of this body.
ASSOCIATED MINERALS OF THE CORUNDUM DEPOSITS

Six specimens of corundum in host rock from the Chumar mine (ranging from 2.3 × 1.3 × 1.8 cm to 7.3 × 3.5 × 4.6 cm) were examined. We identified the seam material as primarily calcite and dolomite. Associated minerals include blue tabular blades of kyanite, small brownish orange euhedral crystals of rutile, different micas (stacked sheets of transparent green fuchsite, orange-brown phlogopite, and whitish margarite), and colorless apatite and scapolite, in addition to the violetish blue-to-red crystals of corundum (figure 5). One of the authors also identified brucite, tremolite, talc, pyrite, and sphalerite during his geologic studies at the Chumar mine (Bassett, 1985a). This list expands on the ones published previously; other researchers have additionally identified zoisite, epidote, muscovite, and analphite in mineral samples containing corundum from Nepal (Harding and Scarratt, 1986). Mineral specimens from this locality are often very colorful, with green fuchsite, orange phlogopite, blue kyanite, and red corundum, all occurring together in a white calcite matrix.

Rubies from the Ruyil deposit, unlike those from Chumar and the other deposits in Ganesh Himal, are embedded in abundant graphite with a less complete suite of associated minerals. In all other respects, though, they appear to be similar to the rubies from elsewhere in the district. Well-formed gem-quality corundum crystals are more common from Ruyil than from Chumar.

MINING METHODS

The actual mining for gemstones is usually performed by local Tamang villagers using crowbars, picks, and shovels (figure 6), with occasional blasting. Blasting followed by hammering is used more commonly at the Chumar deposit than at Ruyil, and incline tunnels have been started at the latter deposit (Chakrabarti, 1994). All of the ruby and fancy-color sapphire deposits are primary (i.e., the stones are found in the host rock); no secondary deposits have been located to date. The miners usually excavate where the marble outcrops can be seen at the surface; the corundum itself is the prime indicator mineral for new deposits.

PRODUCTION, QUALITY, AND SIZES

It is very difficult to estimate how many tens of thousands of carats of corundum have been produced from these deposits over the past 15 years or so. Except for
of testing 27 polished samples (0.21 to 9.47 ct), none of which had been heat treated. Seven of these were from the Chumar deposit and 11 were from Ruyil; the exact deposits for the remaining nine are not known. Also included in this study were 13 crystals and crystal fragments, ranging from 0.35 to 106.42 ct, and the six Chumar mineral specimens described earlier. This collection of gemstones and crystals represents the full range of colors—tone and saturation, as well as hue—observed in Nepalese corundums to date.

We used standard gemological instrumentation to record the refractive indices, birefringence, optic character, pleochroism, optical absorption spectra (desk-model spectroscope), and reaction to ultraviolet radiation (365 nm long-wave, 254 nm shortwave) on all fashioned samples. Specific gravity was determined by hydrostatic weighing with an electronic balance equipped with the appropriate attachments. A binocular microscope, incorporating fiber-optic and other lighting techniques, was used to document the internal features of all fashioned samples and single crystals. The results are given in table 1. We used a Perkin Elmer Lambda 9 spectrometer, with a beam condenser and polarizing filters, for polarized spectroscopy in the UV-visible

Figure 7. Many well-formed crystals have been recovered from the corundum deposits in east-central Nepal. While most crystals are small or medium (usually 5 ct or less) in size, some—such as this 106.42 ct crystal—exceed 100 ct. Photo by Shane F. McClure.

Figure 8. The Nepal deposits have produced some very high quality rubies, such as the fine 1.40 ct faceted ruby and 3.08 ct ruby crystal shown here. The crystal has a habit of c, r, n and z crystal faces. Also note the narrow, blade-like form which is characteristic of much of the rough from these unique deposits. Photo by Shane F. McClure.

MATERIALS AND METHODS

All of the samples in this study were collected by two of the authors (MNM and AMB) in Nepal, from independent miners and the mines managed directly by Himalayan Gems Nepal. We originally selected a total of 29 polished gemstones (22 faceted and seven cut en cabochon) that were fashioned from rough at the cutting factory of Himalayan Gems Nepal. Two faceted samples were subsequently removed from the study. We identified one as a flame-fusion synthetic ruby, and the second yielded properties and characteristics that were not totally consistent with the other samples. Thus, this article presents the results of testing 27 polished samples (0.21 to 9.47 ct), none of which had been heat treated. Seven of these were from the Chumar deposit and 11 were from Ruyil; the exact deposits for the remaining nine are not known. Also included in this study were 13 crystals and crystal fragments, ranging from 0.35 to 106.42 ct, and the six Chumar mineral specimens described earlier. This collection of gemstones and crystals represents the full range of colors—tone and saturation, as well as hue—observed in Nepalese corundums to date.

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through near-infrared region (between 280 and 880 nm) on 17 fashioned samples. For the region between 400 and 6000 wavenumbers (cm⁻¹), we used a Pye-Unicam Fourier-transform infrared (FTIR) 9624 spectrometer with a diffuse reflectance unit for sample measurement. We performed a total of 49 analyses on different regions, color zones, and orientations of 27 samples. Energy-dispersive X-ray fluorescence (EDXRF) chemical analyses were performed on 25 samples (and on different color zones on some samples, for a total of 34 analyses) on a Spectrace TN5000 system, using a program specially developed by Prof. W. B. Stern for the semi-quantitative analysis of corundum. Prof. Stern’s program uses chemically pure element standards and three spectra focusing on the light, medium, and heavy elements, so that the results can be interpreted to three decimal places (i.e., 0.001). He also used a beam condenser to measure small areas or zones of an individual stone. To analyze the internal growth structures of the polished stones, we used a horizontal microscope, a specially designed stone holder, and a mini-goniometer attached to one of the oculars on the microscope, employing the methods described by Schmetzer (1986a and b), Kiefert and Schmetzer (1991), and Smith (1996). We analyzed more than 45 inclusions using a scanning electron microscope with an energy-dispersive X-ray spectrometer (SEM-EDS), X-ray diffraction analysis, and a Raman micro-spectrometer. Some of the inclusions were identified by a single method and others by a combination of techniques (see table 1).

**TABLE 1. Gemological characteristics of the rubies and fancy-color sapphires from Nepal.**

<table>
<thead>
<tr>
<th>Properties</th>
<th>No. samples</th>
<th>Observations</th>
</tr>
</thead>
<tbody>
<tr>
<td>Color</td>
<td>27 polished</td>
<td>Ranging from red to pink to purplish pink, often with visible violetish blue color banding</td>
</tr>
<tr>
<td>Clarity</td>
<td>27 polished</td>
<td>Very clean to heavily included, most in the range slightly to heavily included</td>
</tr>
<tr>
<td>Refractive Index</td>
<td>20 faceted</td>
<td>nₑ = 1.760-1.762</td>
</tr>
<tr>
<td>Birefringence</td>
<td>7 cabochon</td>
<td>n = 1.76-1.77 (spot method)</td>
</tr>
<tr>
<td>Specific gravity</td>
<td>27 polished</td>
<td>3.94-4.01</td>
</tr>
<tr>
<td>Pleochroism</td>
<td>27</td>
<td>Moderate to strong dichroism</td>
</tr>
</tbody>
</table>

**Red zones:** yellowish orange to orangy red parallel to the c-axis, and reddish purple to purple-red, or purplish pink to purple-pink, perpendicular to the c-axis

**Violetish blue zones:** greenish blue to blue parallel to the c-axis, and violetish blue to violet-blue perpendicular to the c-axis.

**UV luminescence** 27

**Red to pink zones.** Long-wave (365 nm): Moderate to very strong red Short-wave (254 nm): Faint to moderate red

**Violetish blue zones:** Generally inert

**Optical absorption spectrum (nm)** 27

General absorption up to approximately 450 nm, 468 (sharp, narrow), 475 nm (sharp, weak to moderate), 476 nm (sharp, narrow), 525-585 nm (broad band), 659 nm (faint, narrow), 668 nm (faint, narrow), 675 nm (very faint, narrow, when present), 692 nm (sharp, narrow), 694 nm (sharp, narrow)

**Internal features** 27

Short rutile needles, various cloud patterns, stringer formations, strong color zoning, weak to prominent growth structures, laminate twinning and parting, “fingerprints,” negative crystals and crystalline inclusions of (with elements identified and mode of identification in parentheses):

- margarite mica (Al, Si, Ca; SEM-EDS and XRD)
- apatite (Ca, P; SEM-EDS, XRD and Raman)
- rutile (Ti; SEM-EDS)
- diaspore (XRD, Raman, and FTIR)
- dolomite (Raman)
- phlogopite mica (K, Mg, Al, Si, Fe; SEM-EDS and XRD)
- calcite (Ca, C; SEM-EDS and FTIR)
- uvite tourmaline (XRD)
- anorthite feldspar (Raman)
- boehmite (FTIR)

*Often containing black inclusions that presumably are graphite.*

The data revealed no significant differences from one sample to the next-including those samples identified as originating from Chumar and Ruyil—
in terms of standard gemological properties, crystal morphology, inclusion patterns, UV-Vis-NIR spectra, or chemical composition. Therefore, the complete data collected on all of the samples—those with specific deposit designations, as well as those without—will be presented as a single group.

**Crystal Morphology.** The rough crystals were predominantly euhedral, with little or no evidence of chemical dissolution on their surfaces. Two main crystal forms dominate the morphology of the corundum found in Nepal (figure 9). The first are dipyramidal crystal habits composed of larger, dominant hexagonal dipyramid z (2241) faces and smaller, subordinate basal pinacoid c (0001) and positive rhombohedron r (1011) faces. The second is a modification of this basic habit, where there is an addition of subordinate hexagonal dipyramid n (2243) faces. Rarely, crystal forms consisting of dominant hexagonal dipyramid r (14 14 28 3) faces, with subordinate c (0001), r (1011), and occasionally n (2243) faces, were also encountered. Many of the crystals had a blade-like appearance when viewed parallel to the c-axis (again, see figure 8). Groups of intergrown crystals were also frequently encountered.

**Visual Appearance.** Face up, most of the polished samples ranged from a “pure” red or pink to a purplish pink. Both the rough and polished samples commonly had strong, eye-visible color zoning (see Growth Characteristics below). A few could even be better described as bicolored (red and very dark violetish blue) corundum; some had additional colorless or pink zones (figure 10). The diaphaneity of the crystals and polished samples ranged from transparent to translucent, depending on the nature and number of inclusions present, as well as on the color saturation and tone of the violetish blue zones.

**Refractive Indices, Birefringence, Optic Character, and Specific Gravity.** These standard gemological properties were found to be consistent with corundum in general (see, e.g., Webster, 1983; Liddicoat, 1989; Hughes, 1990; Hurlbut and Kammerling, 1991) and, more specifically, with the rubies and fancy-color sapphires from Nepal described by past researchers [Harding and Scarratt, 1986; Kiefert and Schmetzer, 1986, 1987; Henn and Milisenda, 1994; see table 1].

**Reaction to Ultraviolet Radiation.** The various color zones of the sample corundums had different reactions to UV radiation. The red to pink zones fluoresced red to both long-wave [moderate to very strong intensity] and short-wave [faint to moderate intensity] UV. The dark violetish blue zones were generally inert to both long- and short-wave UV radiation.

**Pleochroism.** All samples exhibited moderate to strong dichroism when viewed perpendicular to the c-axis with a dichroscope. Within the red-to-pink color zones, we observed yellowish orange to orangy red or pink parallel to the c-axis and reddish purple to purple-red or purplish pink to purple-pink [i.e., for rubies or pink sapphires, respectively] perpendicular to the c-axis. In the dark violetish blue color zones, we noted greenish blue to blue parallel to the c-axis and violet-blue to violet-blue perpendicular to the c-axis.

**Growth Characteristics.** Internal Growth Structures. Weak-to-prominent growth structures were seen in essentially all the polished gemstones examined. Most common were straight and angular sequences of the dipyramid z planes (figure 11). Less common were sequences of the dipyramids n or r, as well as the basal pinacoid c and the positive rhombohedron r.
Figure 10. A wide range of gem-quality rubies and fancy-color sapphires have been recovered from Nepal. Distinctly bicolored—red or pink and violetish blue—stones illustrate some of the unusual zoning in this material. The non-heat-treated Nepalese rubies and sapphires shown here range from 0.87 to 3.86 ct. Photo by Shane F. McClure.

Figure 11. Weak to prominent internal growth structures were frequently observed in the sample rubies and sapphires. This 4.81 ct ruby reveals moderate zonal structures parallel to two series of dipyramid z (2241) planes, along with distinct blue color banding. Immersion, magnified 8 x.

Observation of the gem perpendicular to the c-axis often enabled us to trace the progression of crystal-habit formation (figure 12).

**Color Zoning.** Chemical fluctuations in the growth environment produced obvious color zoning in most of the samples examined. Oscillations between consecutive periods of crystal growth, as well as a preferential crystallographic orientation of the color-causing mechanisms Cr³⁺ (ruby) or Fe²⁺ ? Ti⁴⁺ (blue sapphire), are responsible for the inhomogeneity of color observed in the samples (also refer to UV-Vis-NIR Spectroscopy). When the color was not homogeneous, red-to-pink and near-colorless zones were typically noted parallel to the dipyramid z planes, whereas the dark violetish blue zones were concentrated along the dipyramid z, n, or ?? planes, in addition to the positive rhombohedron r (again, see figure 12).

The most distinctive color-zoning characteristics were related to the dark violetish blue zones. The dark appearance of these zones is attributed to the presence of red (i.e., chromium) and blue (i.e., iron and titanium) chromophores in the same growth phases. These color zones could be very narrow or very thick; the latter were observed only parallel to the dipyramid z planes [see figures 11 and 13]. They also appeared as distinct “wedges” cutting into the gemstone or even dominating it to the point of a bicolor [again, see figure 13]. In these zones, we also observed a texture that might best be described as wispy or smoke-like. In some samples, an irregular color concentration had an undulating nature that resulted from near-colorless areas, or halos, surrounding very small, black-appearing mineral grains of presumably rutile (figure 14). In the case of rutile (TiO₂) inclusions, Ti would be absorbed from the host corundum, thereby depleting an essential component of the Fe²⁺ ? Ti⁴⁺ charge-transfer necessary for the blue coloration.

**Twinning.** We saw twinning parallel to the positive rhombohedron r (1011) in several of the stones. Typically, we noted only one direction of laminated twinning, parallel to a single series of r (1011) planes; occasionally, however, there were as many as three twinning systems, parallel to additional positive rhombohedral planes. Parting parallel to r (1011) was also prominent in a few samples (figure 15).

**Inclusions.** A rich diversity of inclusions were noted in the sample rubies and fancy-color sapphire.
Most commonly, clouds of very fine, short rutile needles, present throughout, gave some of the gemstones a slightly "hazy" appearance to the unaided eye (figure 16). Bright orange-to-black crystals of rutile were observed singly or in small irregular clusters (figure 17); these displayed a metallic luster when they were polished at the surface. Other stringer-type inclusion patterns consisted of nearly parallel, slightly diverging "sprays," extending essentially perpendicular to growth planes or in an antenna-like pattern (figure 18).

Apatite took on a variety of forms, including euhedral hexagonal columns (figure 19, left) and slightly curved rods (figure 19, right). Although not commonly noted in apatite, basal cleavage was
observed in several of the apatites we identified, especially in the rod-shaped forms. The calcium-rich mica margarite was often present in irregular masses within which additional inclusions were noted, such as individual crystals of apatite or rutile, or masses of graphite, sometimes forming one complex mineral assemblage (figure 20). Light brown crystals of phlogopite mica occurred in small, mostly irregular, rounded forms (figure 21). Transparent colorless crystals of calcite and dolomite were seen infrequently (most of the transparent colorless crystals in these stones proved to be apatite or margarite). Although rare, black uvite tourmaline (figure 22) and transparent colorless anorthite feldspar (figure 23) were identified. Also

unusual was the "halo" of minute rutile needles surrounding an unidentified small black mineral grain (figure 24).

We observed a wide range of fingerprint-like inclusions, all involved in various stages of the “healing” process. Two-phase (liquid and gas) inclusions were common. "Intersection tubules" at the junction of two or three twin planes were frequently penetrated by alteration products such as boehmite. Boehmite was also identified lining the parting planes. Irregular "veins" of AlO(OH)—mostly boehmite, but also diaspore—were also noted traversing several of the polished gemstones. In reflected light, the reduced luster of the AlO(OH) "vein," as compared to the higher luster of the host corundum, could be mistaken for the glass-like

Figure 15. Parting planes parallel to one, two, or three series of positive rhombohedron r (1011) crystal faces, were seen in a few samples. In this stone, the parting planes—which form a checkerboard pattern—were lined with the aluminum oxy-hydroxide boehmite, AlO(OH). Fiber-optic illumination, magnified 20×.

Figure 16. Clouds of very fine, short rutile needles were present in nearly all of the rubies and fancy-color sapphires from Nepal. However, none of these stones revealed the nest-like concentrations that are typical of rubies from Mogok. Fiber-optic illumination, magnified 22×.

Figure 17. Clusters of small rutile crystals were also common in the rubies from Nepal. Typically, they were bright orange to black and displayed a metallic luster where polished at the surface. Fiber-optic illumination, magnified 50×.

Figure 18. Antenna-like stringer formations were frequently seen in the Nepalese samples. Although such inclusions have been noted in rubies from Luc Yen (Vietnam) and Mong Hsu (Myanmar), they tended to be more densely concentrated in the Nepalese samples. Fiber-optic illumination, magnified 32×.
fillings observed in some heat-treated rubies, but careful examination will establish that the gem has not been heated.

**Absorption Spectra.** All spectra were dominated by Cr\(^{3+}\) absorption features, with the bands being weaker in the lighter red (i.e., pinkish) stones and more intense in the deeper red stones. Occasionally a secondary absorption influence was seen as a result of the Fe\(^{2+}\) ? Ti\(^{4+}\) intervalence charge transfer responsible for the blue color component in the dark violetish blue zones. These results are consistent with those reported earlier by Kiefert and Schmetzer (1986, 1987).

**Desk-Model Spectroscope.** In the visible range, a general absorption to approximately 450 nm was apparent, along with weak to distinct lines at 468 nm and at 475 and 476 nm (a doublet). The width of a moderate-to-distinct absorption band from approximately 525 to 585 nm was related to the chromium content of the gemstone. We also noted faint lines at 659 and 668 nm, plus two strong lines at 692 and 694 nm, which appear as a bright emission line at 693 nm.

**UV-Vis-NIR Spectroscopy.** The general shape of the spectral curve also varied considerably depending on the chromium content of the zones measured. The two broad bands at about 405 and 550 nm, as well as the weak to distinct sharp peaks recorded at 468, 475, 476, 659, 668, 692 and 694, are all ascribed to Cr\(^{3+}\). A faint absorption peak observed at 675 nm in some stones has also been recorded in rubies from other sources (e.g., Mong Hsu, Myanmar, and certain deposits in east Africa), but the cause is still unclear (Peretti et al., 1995).

**Infrared Spectroscopy.** In addition to the dominant absorption characteristics of corundum, between 300 and 1000 cm\(^{-1}\) (peak positions at about 760, 642, 602, and 450 cm\(^{-1}\); Wefers and Bell, 1972), the rubies...
and sapphires in this study revealed two dominant bands at about 3320 cm\(^{-1}\) and 3085 cm\(^{-1}\), with an additional pair of weaker bands at 2100 and 1980 cm\(^{-1}\) (figure 25). These absorption bands are related to OH stretching frequencies and identify the presence of boehmite (Farmer, 1974; Wefers and Misra, 1987). Although different color zones did not reveal any statistical differences in the presence or absence of boehmite, different areas and orientations of the same stone did show variations in the absolute and relative intensities of the absorption bands, as well as a slight shift in the position of the absorption maximum. To a much lesser degree, diaspore was also indicated in the infrared spectra of some samples, with bands at approximately 1990, 2040, 2885, and 3025 cm\(^{-1}\) (Farmer, 1974; Wefers and Misra, 1987; Smith, 1995).

The presence of AlO(OH)—boehmite and diaspore—was generally traced to locations along parting planes or irregular seams. Not all samples showed AlO(OH)-related absorption bands, while several displayed such strong AlO(OH) absorption features that a distinction between boehmite and diaspore was not possible. Absorption bands associated with mica and calcite were also occasionally recorded.

**Chemical Analysis.** The most significant variations were recorded in Cr concentration, which again correlated to the depth of red-to-pink color in the area measured. Titanium [Ti] and iron [Fe] were the next most significant trace elements recorded, followed by measurable amounts of vanadium [V] and gallium [Ga], as shown in table 2.

One interesting observation related to the trace-element concentrations of areas that were "pure" red as opposed to ones that were "pure" dark violetish blue. In every instance, the concentrations in absolute and relative values showed no consistent variation between Cr, Fe, or Ti within the two color zones.

**DISCUSSION**

Corundum from Nepal has received sporadic mention in the gemological literature over the past decade or so (see, e.g., Harding and Scarratt, 1986; Kiefert and Schmetzer, 1986 and 1987; Bank et al., 1988; Niedermayr et al., 1993). The results of this current investigation support and expand on the findings of such earlier researchers and, for the first time, provide detailed locality information. We identified three inclusions in the rubies from Nepal that had not been described before—uvite tourmaline, anorthite feldspar, and diaspore—as well as several distinctive inclusion and color-zoning patterns. However, we did not observe the three-phase inclusions described by Kiefert and Schmetzer (1986, 1987) in any of our samples. Also, we did not encounter the identification difficulties experienced by Bank et al. (1988); that is, all of our samples were easily identified as natural corundum.

In general, the rubies and fancy-color sapphires from Nepal are similar to corundum from other marble-type sources found around the world, including Myanmar [Burma], Vietnam, Afghanistan, Pakistan, and Tanzania. The most distinctive features of these Nepalese corundums are their inclusions.
The dense concentrations of very fine, short rutile needles are unlike the long, highly iridescent rutile needles observed in most rubies from marble-type deposits. Dense "nests" of short rutile needles, so typical of rubies from the historic Mogok stone tract in upper Myanmar, were not seen in our Nepalese study samples. The various stringer patterns we saw in these Nepalese stones have also been seen in rubies from other deposits, including Mong Hsu in Myanmar (Smith and Surdez, 1994; Peretti et al., 1995) and Vietnam (Kane et al., 1991; Smith, 1996). However, the Nepalese stones appeared to have much denser concentrations of such patterns.

Apatite crystals of various forms may be seen in rubies from many sources, including Mogok [e.g., Gübelin and Koivula, 1986], Vietnam [Kane et al., 1991], and various deposits in East Africa. However, the rod-shaped apatites observed in the samples from Nepal have not been described in rubies or fancy-color sapphires from any other source. Kane et al. (1991) described similar transparent colorless rod-shaped minerals in rubies from Vietnam, but these were identified as calcite. A common mineral inclusion in rubies from marble-type deposits, calcite was encountered only rarely in the rubies from Nepal.

Mica is common in corundum from other deposits [e.g., Myanmar, Sri Lanka, and Tanzania], but the sheer number of inclusions of the variety margarite in our test samples is unlike anything we have seen in rubies from other sources. Anorthite feldspar and uvite tourmaline, which were observed in a few of our sample rubies and fancy-color sapphires, have not been identified in rubies from any other locality. Therefore, they may also be instrumental in establishing Nepalese origin. Nor have the authors seen inclusion patterns such as the small black mineral surrounded by a spherical "halo" of minute rutile needles in rubies from other sources. On the other hand, inclusions such as pyrrhotite, zircon, pyrite, and spinel have not been identified to date in rubies from Nepal; these are frequently seen in rubies from East Africa, Vietnam, and Myanmar, as well as other sources.

In their natural [not heat-treated] state, these rubies, pink sapphires, and bicolored sapphires should not be difficult to distinguish from rubies, including those with blue color zones, from other sources. No heat-treatment studies were conducted on these samples. However, when such stones are heat treated, the inclusion features and color zoning will be greatly altered, making it more difficult to identify corundum from Nepal [see, e.g., Peretti et al., 1995]. Although it is usually difficult to detect a gemstone’s source by the inclusion features, observation of any of the above-described mineral inclusions will clearly separate a natural ruby or fancy-color sapphire from a synthetic counterpart produced by any of the known manufacturing processes.

The linear and angular sequences of non-color-zoned growth structures in our samples—following the crystal planes z, n, ?, r, and c—did not reveal any unique or diagnostic features or patterns. Similar sequences of growth structures have been observed by one of the authors (CPS) in natural rubies from other sources (e.g., Vietnam or East Africa). However, they can be useful in separating

<table>
<thead>
<tr>
<th>Oxide</th>
<th>Wt.%</th>
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<tbody>
<tr>
<td>Al₂O₃</td>
<td>98.9-99.8</td>
</tr>
<tr>
<td>Cr₂O₃</td>
<td>0.013-0.383</td>
</tr>
<tr>
<td>TiO₂</td>
<td>0.016-0.224</td>
</tr>
<tr>
<td>Fe₂O₃</td>
<td>0.004-0.069</td>
</tr>
<tr>
<td>V₂O₅</td>
<td>0.004-0.034</td>
</tr>
<tr>
<td>Ga₂O₃</td>
<td>0.010-0.023</td>
</tr>
</tbody>
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Figure 24. Still another feature that has not been noted in rubies from other sources was present in many of the rubies and fancy-color sapphires from Nepal: small black mineral grains surrounded by halos of very fine, minute rutile needles. Fiber-optic illumination, magnified 50x.
Figure 25. The non-heat-treated rubies and fancy-color sapphires from Nepal revealed additional absorption features in the infrared region of the spectrum. Distinct absorption bands at 3320 and 3085 cm\(^{-1}\), and weaker peaks at 2100 and 1980 cm\(^{-1}\), indicated the presence of boehmite, which was seen concentrated along veins or lining parting planes. Such absorption characteristics are helpful not only in identifying foreign mineral phases that may be present, but also for indicating that the gem has not been heat treated.

Chemically, these rubies and fancy-color sapphires are also similar to their counterparts from other marble-type sources (Tang et al., 1988; Hughes, 1990). Nevertheless, their chemistry will provide a ready means of separating them from high-iron natural rubies from sources such as Thailand (Tang et al., 1988), Cambodia (Jobbins and Berrange, 1981), Madagascar (Smith, 1996), and certain deposits in East Africa (see, e.g., Hänni and Schmetzer, 1991). In addition, the collection of chemical data is also valuable in separating natural from synthetic corundum (e.g., Stern and Hänni, 1982; Muhlmeister and Devouard, 1991).

Infrared spectroscopy may provide additional proof that the gemstone was not heat treated, when AlO(OH) is present, as well as a very good indication of whether it is natural or synthetic (Volynets et al., 1972; Beran, 1991; Smith, 1995, pp. 326–328). One of the authors (CPS) has observed that rubies from other natural sources, such as certain deposits in Vietnam or East Africa, sometimes also reveal dominant absorption features relating to boehmite, as well as other minerals.
In recent years, a number of ruby and sapphire deposits have been discovered in a wide variety of sources around the world. While Nepal is neither the newest nor most significant corundum source in the trade today, it offers an interesting array of ruby and fancy-color sapphires for jewelers, gemologists, and consumers alike. These gems from the Ganesh Himal appear to be concentrated along a single geologic "belt" within the northern Dhading District of east-central Nepal, and in two mines in particular, Chumar and Ruyil. Since they were first discovered in the early 1980s, these gemstones have been entering the world markets. The isolated locations, high altitudes, harsh seasonal weather conditions, and other difficulties have contributed to the sporadic mining activities and the relatively small amounts of gem material produced to date. However, research by two of the authors [MNM and AMB] indicates that larger reserves of these rubies and fancy-color sapphires are yet to be discovered. Modernized mining equipment and methods [including tunneling or ‘benching’ techniques] will be necessary for these deposits to reach their full potential.

Although similar in general to rubies and fancy-color sapphires from other marble-type deposits, the Nepal corundums may be distinguished on the basis of the entire collection of the gem’s individual properties and characteristics. Certain features that may prove helpful in the identification include dense concentrations of very fine, short rutile needles throughout the stone; rod-shaped forms of apatite; masses of transparent colorless margarite, opaque black masses of uvite tourmaline; transparent colorless anorthite feldspar; and small black mineral grains surrounded by "halos" of minute rutile needles. Nepal may also be indicated for a particular ruby if any of the transparent mineral inclusions are associated with black masses of graphite. Nepalese rubies and fancy-color sapphires may also have distinctive color-zoning characteristics, such as large red and dark violetish blue portions in a single stone (i.e., bicolor), or dark violetish blue zones in thick bands or wedge shapes. Especially distinctive is the presence of a wispy or smoke-like texture or near-colorless "halos" surrounding a mineral inclusion.

Such an ensemble of internal features also will help separate a Nepalese corundum from a synthetic corundum of any of the various production techniques. As a reminder, even in the remotest regions of the world, one should never take for granted that a gemstone is natural, as the synthetic ruby crystal fragment purchased in Nepal by one of the authors [MNM] illustrates.

Nepal’s mineral wealth—consisting of tourmaline, beryl, garnet, quartz, danburite, kyanite, apatite, sodalite, zircon, sphalerite, epidote, diopside, iolite, and andalusite, among others—is already recognized widely in the trade. Now rubies, pink sapphires, and bicolored corundum can also be added to the list.

Acknowledgments: The authors are grateful to Dr. L. Kiefert and Dr. H. A. Hänni of the Swiss Gemmological Institute (SSEF), Basel, Switzerland, for performing Raman spectral analyses; to Dr. H.-D. Von Schulz of SUVA, Lucerne, Switzerland, for performing scanning electron microscopic analyses; to Professor S. Graeser of the Institute of Mineralogy and Petrography, University of Basel, for X-ray diffraction analyses; and to Prof. W. B. Stern of the Institute of Mineralogy and Petrography, University of Basel, Switzerland and Ms. N. Surdez of the Gübelin Gemmological Laboratory, Lucerne, for providing the semi-quantitative chemical analyses. Unless otherwise noted, photomicrographs are by Christopher P. Smith.
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