The East Kemptville tin deposit, Yarmouth County, southwestern Nova Scotia: a lithogeochemical study of the wallrock metasedimentary rocks

Daniel J. Kontak
Nova Scotia Department of Natural Resources, P.O. Box 698, Halifax, Nova Scotia B3J 2T9, Canada
and
Jarda Dostal
Department of Geology, St. Mary's University, Halifax, Nova Scotia B3H 3C3, Canada

Date Received November 27, 1991
Date Accepted March 2, 1992

The results of a petrographic and lithogeochemical study of metasedimentary rocks (Lower Paleozoic Meguma Group) surrounding the northern part of the East Kemptville tin deposit, southwestern Nova Scotia are presented. The metasedimentary rocks, including interbedded psammites and pelites, are fine-grained and highly indurated. Pelitic horizons contain hornfelsic textures (i.e., ≤5 mm cordierite grains) near the granite; sparse veining includes some cassiterite and sulphides. Hydrothermal alteration is manifested by disseminated, fine-grained, secondary muscovite and/or muscovite and chlorite infilling and lining microfractures with bleached haloes and associated cassiterite, sulphides, tourmaline, fluorite and carbonate.

The major and trace element chemistry, including the rare-earth elements, of the metasedimentary rocks are compared to different data bases that comprise: (1) averages for pelites and psammites of Phanerozoic age, (2) nonmineralized Meguma Group rocks, and (3) fields in discriminant binary element diagrams. For most of the major and trace elements analyzed an apparently unaltered, normal chemistry is indicated, except for some LILE (e.g., Li, Rb, U) and ore-forming elements (e.g., Sn, Zn, Bi) where enrichment is considered to reflect infiltration of mineralizing fluids from the proximal granite/greisen of the main deposit area. The data indicate that infiltration of fluids was minimal and that, for the most part, the metasedimentary rocks acted as an impermeable cap to the ore-forming solutions.

Les résultats d'une étude pétrographique et lithogéochimique des roches métasédimentaires (Groupe de Meguma du Paléozoique inférieur) entourant la partie nord du gisement d'étain de East Kemptville, dans le sud-ouest de la Nouvelle-Écosse, sont présentés. Les roches métasédimentaires, comprenant des grès et des péries, sont à grain fin et fortement indurées. Les horizons pelitiques montrent des textures de corneéennes (i.e., des grains de cordierite ≤5 mm) près du granite; des veines épaisses contiennent de la cassiterite et des sulfures. L'altération hydrothermale se manifeste par de la muscovite secondaire à grain fin disséminée ou par de la muscovite et de la chlorite en remplissage et en tapisage de microfractures aux épontes lessivées et associées à de la cassiterite, des sulfures, de la tourmaline, de la fluorite et des carbonates.

La composition en éléments majeurs et traces, incluant les terres rares, des roches métasédimentaires sont comparées à diverses bases de données, incluant: (1) les composition moyennes des péries et des grès d'âge Phanérozoïque, (2) les roches non minéralisées du Groupe de Meguma et (3) les champs des diagrammes de discrimination binaires. Les teneurs de la plupart des éléments majeurs et traces analysés indiquent une composition normale, apparemment inaltérée, à part pour certains éléments à grand rayon ionique (e.g., Li, Rb, U) et certains éléments économiques (e.g., Sn, Zn, Bi) où l'enrichissement est considéré comme représentant l'infiltration de fluides minéralisateurs provenant du greisen proximal de la région du gisement principal. Les données indiquent que l'infiltration des fluides était minimale et que les roches métasédimentaires ont agi principalement comme une enveloppe imperméable pour les solutions minéralisatrices.

Introduction

The East Kemptville Sn-(Cu-Zn-Ag) deposit, located in Yarmouth County of southwestern Nova Scotia (Fig. 1), contained initial reserve estimates of 56 million tonnes of 0.165% Sn (Moyle, 1985). A significant resource of tin on a global scale, the deposit was discovered in the late 1970's and mining commenced in the fall of 1985 as a 18,000 tonne/day (9000 tonnes ore) open pit operation producing ca. 4000 tonnes of Sn concentrate per annum. The deposit ceased production on January 3, 1992, in part reflecting the depressed price of tin.

The East Kemptville Sn-base metal deposit is entirely contained within a topaz-muscovite leucogranite body...
(Kontak, 1990a) which intruded metaturbiditic sedimentary rocks of the Lower Paleozoic Meguma Group. While considerable attention has been focused on the host leucogranite and mineralized greisens (Chatterjee and Strong, 1984; Chatterjee et al., 1983; Kontak, 1988, 1990a, 1990b, 1990c, 1991; Kontak and Cormier, 1991; Kontak and Chatterjee, in press; Richardson, 1983, 1988; Richardson et al., 1982, 1988, 1990a), the metasedimentary rocks of the wallrock have received limited attention. Richardson (1983) reported two chemical analyses each of mineralized metasedimentary rocks at the deposit site and unaltered rock away from the deposit. Kontak (1990c, in press) presented preliminary results of a lithogeochemical study of metasedimentary rocks collected from the northern end of the deposit area near the granite-metasedimentary rock contact.

In this paper, the more detailed results of this lithogeochemical study of the metasedimentary rocks are presented, together with petrographic observations. The primary concern of the study is to evaluate the role of the wallrocks in terms of the evolution of the mineralization at East Kemptville. As first noted by Richardson et al. (1982), the highly indurated nature of the metasedimentary rocks and general lack of significant alteration and associated mineralization within them suggests that they may have acted as an impermeable barrier to the mineralizing fluids. Thus, a lithogeochemical study may provide evidence by which to evaluate this hypothesis or, alternatively, determine if in fact some fluid migration out of the granite and into the country rock did occur. A corollary of this study is to determine the nature and extent of a primary dispersion halo about the deposit which may be of significance in terms of exploration for additional reserves and other mineralized centres.

**Regional Geological Setting**

The study area is located in the southern part of the Meguma Terrane (MT in Fig. 1) which is underlain predominantly by Lower Paleozoic metasedimentary and metavolcanic rocks that were deformed by the mid-Devonian Acadian Orogeny (Poole, 1967; Reynolds and Muecke, 1978; Keppie and Dallmeyer, 1987; Muecke et al., 1988) and subsequently intruded by meta- to peraluminous 370 Ma granites (Clarke and Halliday, 1980; Reynolds et al., 1981). The northern part of the Meguma Terrane, south of the east-west trending Cobequid-Chedabucto boundary fault (CCFS in Fig. 1; Keppie, 1982; Mawer and White, 1987), is underlain by Carboniferous and younger sedimentary rocks, part of the regional Maritime Basin of Atlantic Canada, that infilled successor basins after the regional Acadian deformation and plutonism.

The East Kemptville deposit area is located at the western margin of the large, 370 Ma, composite South Mountain Batholith (MacDonald et al., 1989, 1992). More specifically the deposit is located at the contact of the Davis Lake pluton...
and metasedimentary rocks of the Meguma Group (Fig. 2).
The deposit area lies within the northeast-trending polymetallic Sn-base metal domain of Chatterjee (1983; see also Kontak et al., 1990) which includes several centres of variable mineralization associated with either granitic rocks (e.g., East Kemptville; Kempt Snare Lake [Soehl et al., 1989]) or metasedimentary rocks of the Meguma Group (e.g., Duck Pond deposit; Hattie 1989; Pitre and Richardson, 1989) (Fig. 2). Also of importance is the presence of a metalliferous-rich stratigraphic horizon referred to as the GHT or Goldenville-Halifax Transition Zone (Zentilli et al., 1986) which has been suggested to be important in terms of metallogeny both within the southwestern Nova Scotia tin domain and the auriferous Lower Paleozoic Meguma Group of the eastern Meguma Terrane (Sangster, 1990).

**Local Geological Setting**

The inset map in Figure 2 shows the outline of the deposit area which coincides with the leucogranite-metasedimentary rock. The area is shown in more detail in Figure 3 where the ore zones are indicated. The presence of roof pendants within the granite and the plunging nature of the granite-sediment contact to the northeast (about 30°) together suggest that the roof zone of the intrusion (i.e., magma chamber) is probably exposed. The northern contact of the granite and metasedimentary rocks is steeply dipping; in fact in most cases it is subvertical and in some locations coincides with a fault. This fault, referred to as the Contact fault by Richardson et al. (1988), hugs the northern contact and although coinciding with the lithological contact, may also cross-cut it, thus postdating the time of intrusion and mineralization. The fault probably formed during the main phase of post-plutonism deformation which affected most of the deposit area (Kontak, 1990a; Kontak et al., 1986; Kontak and Cormier, 1991).

The leucogranitic host rock has been described in detail by Kontak (1990a, 1991) and only a brief summary is presented here. It is a fine- to medium-grained, seriate textured topaz-muscovite leucogranite with variably developed penetrative deformational fabrics. There is a conspicuous absence of miarolitic cavities and related features (i.e., breccias, tuffisites, etc.), and pegmatites (<1 m) are restricted to a few rare localities. Thus, the leucogranite is considered to have intruded as a H₂O-undersaturated melt. Rare dykelets of leucogranite (<2-3 cm width) cut the metasedimentary rocks.

Mineralization consists of an early stage of massive and zoned topaz-cassiterite-bearing greisens which are cut by base-metal-rich quartz veins. The massive greisens are fine-grained, dark grey to black and account for the majority of the tin mineralization; they contain approximately 0.2 to 0.4% Sn and variable quantities of base and precious (Ag) metals. In contrast, the zoned greisens consist of <5 cm to 1 m wide...
zoned greisens cored by massive cassiterite- and topaz-rich zones. Further details of the mineralogy and geochemistry of the greisens, as well as the later base-metal rich quartz veins, can be found in Kontak et al. (1986), Kontak (1990c) and Richardson (1988). Mineralization in the metasedimentary rocks is rare, and consists of tin-bearing and sulphide-bearing veins of small (<1 cm width) dimensions and extent (i.e., near the contact).

As Figure 3A indicates, there is a general northeasterly trend to the ore zones and in a broad sense they generally coincide with the granite-metasediment contact. For example, as mining progressed at the deposit the extent of the Main Ore Zone decreased as the extent of the metasedimentary keel extending into the granite diminished in size; a similar situation would apply equally to the North and South Extension Ore zones.

The metasedimentary rocks enveloping the deposit are fine-grained, mixed psammites to pelites with the former dominating, and all rocks are generally highly indurated. Of the 36 samples collected, 50% are psammites, 34% pelites and the remainder semipelites; this is considered a good approximation of the proportion of lithologies present. Where Ș0 is observed it is moderately to steeply dipping and indicates a northeasterly-trending anticline extending through the area. The pelitic lithologies have a well-developed, sub-vertical slaty cleavage which may be cut by a second, flat-lying spaced cleavage. In a few instances, narrow zones (i.e., 6-10 cm) of intense deformation were observed, similar to the
mylonitic zones in the granite (Kontak, 1990a; Kontak et al., 1986). Near the granite, hornfelsic textures are present in the pelites with abundant, elliptical-shaped, 5 mm-size cordierite grains. Although bleaching and veining are not obvious in hand specimens, such features are observed in thin section and will be discussed below.

**Sampling of Metasedimentary Rocks**

Thirty-six samples of metasedimentary rock were collected from the northern part of the East Kemptville deposit area during detailed mapping in the summer of 1986 (Kontak et al., 1986). Sampling was designed to test for the possible presence and extent of a primary metal dispersion halo around the mineralized granite. Thus, the samples were collected along the granite-metasedimentary contact and along traverses away from and perpendicular to the northernmost contact (labelled A and B, respectively, in Fig. 3). Two of the samples collected (EK-86-1021 and 1079) came from a roof pendant in the southeastern part of the study area. All the samples collected were free of megascopically visible signs of veining and alteration.

In addition to the metasedimentary rocks, two greisenized leucogranites (EK-1154, 1227) were included in the study to compare the alteration geochemistry in the greisenized granite with that in the metasedimentary rocks.

**Petrography of Metasedimentary Rocks**

The metasedimentary rocks are fine-grained, dark grey to black and generally highly indurated. Hornfelsic textures are generally uncommon except in the most aluminous rocks. Veining on a macroscopic scale is rare to absent and only occurs immediately adjacent to the granite. Bedding is only rarely observed because of the flat nature of the exposure and the apparent homogeneous nature of the sediments. Veins cutting the metasedimentary rocks include: (1) granitic dykes ca. 1 to 3 cm wide, (2) a few cassiterite-rich (i.e., 50-80%) veins ca. 0.5 to 1 cm wide, and (3) quartz-carbonate-sulphide veins (along the south end of the deposit only). Fractures mantled by narrow (1-3 mm) zones of bleaching are also present.

The metasedimentary rocks are composed predominantly of quartz, feldspars, biotite and muscovite with lesser carbonate, opaque minerals, tourmaline, apatite and zircon (Fig. 4a). The rocks are either dominated by detrital quartz and feldspar in a fine-grained, mica dominant matrix (psammites; Fig. 4a) or are composed of fine-grained phyllosilicates (pelites). The micas are of two distinct populations in terms of grain size. The coarser material may be of either regional metamorphic or contact metamorphic origin. The finer grained material probably reflects in part later hydrothermal alteration (see below). The rocks have at least one well-developed tectonic fabric (S); in some cases crenulations and C-S fabrics are observed. In a few samples chlorite-rich shear zones cross-cut the earlier, regional (?) S fabric. Porphyroblasts occur in only a few samples, are subrounded to elliptical, and are generally replaced by muscovite ± chlorite along the outer margins with a darker core area consisting of unidentified material.

It is rare for the metasedimentary rocks not to show signs of hydrothermal alteration, which includes the following features: (1) fine-grained white mica and orange-brown biotite which locally may replace most of the matrix (Fig. 4b,e); (2) locally abundant tourmaline which overgrew the main fabric (Fig. 4d); (3) irregular patches of chlorite replacing the matrix; (4) fractures or veinlets (see below) of different types with or without ore minerals and with bleached margins (Fig. 4e,f); and (5) local disseminations of one or more of cassiterite, sphalerite, fluorite and carbonate which have no apparent proximity to veins or fractures. The most common of the aforementioned features observed are the formation of fine-grained muscovite which may replace considerable amounts of the matrix, and the presence of orange-brown biotite with abundant opaque minerals and rutile inclusions. The fine-grained muscovite and biotite are quite distinct texturally from the coarser metamorphic micas of either regional or contact metamorphic origin and are, therefore, considered to be of hydrothermal origin.

Veinlets (1-3 mm) which cross-cut the metasedimentary rocks consist of several varieties in terms of varying proportions of minerals, but these can be broadly grouped into two main types: (1) muscovite-rich, either fine- or coarse-grained, with associated ±chlorite, cassiterite, ore minerals, tourmaline, carbonate (Fig. 4b,c,e,f); (2) chloride ± cassiterite, ore minerals, tourmaline (Fig. 4d). Where cross-cutting relationships could be observed (three cases), muscovite-bearing fractures were seen to offset and postdate chloride-bearing fractures. Both veinlets generally have bleached margins where fine-grained muscovite or chloride predominate. The veinlets are never abundant and in terms of modal mineralogy account for only trace amounts of the mineralogy of a typical thin section. Thus, where abundant hydrothermal mica has replaced the matrix, it must reflect pervasive passage of fluid along much finer channelways.

The combination of regional metamorphism, contact metamorphism, hydrothermal alteration and attendant mineralization and later superimposed shear deformation (Kon- tak and Cormier, 1991), makes it difficult to be more specific about the possibility of different generations of minerals without knowledge of mineral chemistry. Such work is in progress and will be reported elsewhere.

**Analytical Procedures**

Samples selected for chemistry were fine-grained vein-free rocks. All samples were first cleaned and weathered surfaces removed by sawing. Material was crushed to -200 mesh and all samples analyzed for major and trace element contents with eleven samples also analyzed for rare-earth elements (REE). Major elements were analyzed on fused disks and trace elements (Rb, Sr, Ba, V, Cr, Ni, Cu, Zn, Ga, Zr, Nb, Y) on pressed powdered pellets using an automated Phillips X-ray fluorescence spectrometer at St. Mary’s Uni-
Fig. 4. Photomicrographs (field of view ca. 2 mm wide; transmitted light) of metasedimentary rocks at East Kemptville. (a) Unaltered psammitic with coarse grains of quartz and feldspar in matrix of mica; note the weak foliation (horizontal) and sutured quartz grains. (b) Coarse cassiterite grain (cst) with coarse rosettes of muscovite growing towards the sericitized metasedimentary rock. Note that in the lower left corner some vestige of the former psammitic texture remains, but close to the cassiterite it has been replaced by fine muscovite. (c) Cassiterite (cst)-fluorite (fl)-quartz (clear)-muscovite (ms; intergrown with quartz) assemblage in psammitic rock. (d) Veinlet of chlorite with cassiterite and sulphides growing along it. Note the bleaching of the rock against the vein which has destroyed most of the pre-existing mica mineralogy. (e) Muscovite veinlet (ms) in psammite with symmetrically disposed border zone of bleached wallrock (bzbr). (f) Tourmaline-muscovite-rich area in psammite located adjacent to a granite dyke.

versity, Halifax, Nova Scotia. Additional trace elements (Sc, Hf, Ta, Co) were analyzed by instrumental neutron activation along with the REE (La, Ce, Nd, Sm, Eu, Gd, Tb, Yb, Lu); samples were irradiated at the McMaster University reactor
and counting done at St. Mary’s University. Complete details of the analytical procedures, including estimates of precision and accuracy are given in Dostal et al. (1986). The remaining trace elements (Li, Mo, W, As, Bi, Ag, Sb, U, Pb) were analyzed at Bondar-Clegg Company Limited, Ottawa, Ontario by inductively-coupled and DC plasma. We note that there appears to be a systematic enrichment in Gd for some unknown reason which is most apparent in the chondrite-normalized diagrams where a prominent inflection occurs at this point.

**GEOCHEMISTRY**

The data have been subdivided into three populations based on wt. % SiO₂: <60%, 60-70% and >70% which correspond to psammites, semi-pelites and pelites, respectively, in terms of their petrographic features. The average ±1 standard deviation for each of these populations is presented in Table 1, along with representative analyses of different shale suites compiled from the literature and the two greisenized leucogranite samples. The averages and ranges of some metals (Zn, Cu, As, Sb, W, Sn, Pb) are given in Table 2 for comparative purposes (see below). The complete list of geochemical data is available from the senior author upon request. The REE data are given in Table 3 along with the elemental ratios discussed in the text.

**Major Element Geochemistry**

Results of major element chemistry are summarized in Table 1 and graphically presented in Figures 5, 6 and 7. The three suites show consistent and systematic chemical variation with respect to the major elements such that there is a gradual change in elemental abundances from the pelites to the psammites. The data set is plotted in a series of binary element diagrams in Figure 5 using Al₂O₃ as the absissa since it shows the largest overall range. In Figure 5 the other major elements are seen to show well defined positive (TiO₂, MgO, Fe₂O₃, K₂O, P₂O₅, LOI) or negative (SiO₂, Na₂O) correlations; calcium appears to behave independently. The excellent co-variations observed in Figure 5 are typical of major element chemistry of sedimentary rocks (e.g., Feng and Kerrich, 1990; Pan et al., 1991) and reflect systematic changes in mineralogy. The two altered leucogranites plot away from the well-defined trends established for the metasedimentary rocks and there appears to be no tendency for the sedimentary rock data to trend toward that of the greisens as a result of alteration.

In Figure 6 the data are plotted in the binary discrimination diagrams of Bhatia (1983); these diagrams are based on a compilation of data for greywackes. The deviation of the pelite data from the identified fields is to be expected because the diagrams are designed for clastic rocks; however, the pelite data are included to illustrate the continuity of the chemistry from psammite to pelite regardless of the binary plot used. The psammitic data conform to the fields for continental island arc (B) and active continental margin (C) which are consistent with the generally inferred paleotectonic setting for these rocks. Thus, in general the data conform to major element chemistry for greywackes suggesting that their chemistry has not been modified despite their proximity to a mineralized centre. Again, there appears to be no tendency for the sedimentary chemistry to trend toward that of the greisen samples.

The major element chemistry of the pelites is compared to some average shale compositions in Table 1. As one might expect, the variation of provenance, depositional environment and diagenetic history have contributed to the differences in the chemistry of the suites. However, the Bendigo shale average is remarkable similar to the pelites, except for relative differences in Na₂O and LOI, suggesting again that the chemistry of the rocks may not have been modified significantly.

A final comparison of the chemistry for the East Kemptville data set is made with the average compositions of similar rock types in the greenschist facies from the eastern Meguma Terrane as compiled by Douma (in prep.). In Figure 7 the average compositions are normalized to Douma’s data and there appears to be a very close correspondence between the two data sets, with even slight relative enrichments and depletions observed being identical for both data sets.

**Trace Element Geochemistry**

The average trace element geochemistry for the three groupings are summarized in Table 1 and we note the general absolute enrichment of Rb, Zn, Cu and Sn and the high percent relative deviation associated with these elements. We also note that these elements and also Bi, Li and W are enriched relative to the average compositions of similar rocks from the eastern Meguma Terrane (Fig. 7).

**Rb, Sr, Ba, Li and Ga**

There is a strong positive correlation between wt. % K₂O and Rb, Ba, Li and Ga, while Sr shows a poorly defined negative correlation. These trends are substantiated by the averages shown in Table 1 and the correlations suggest that these elements essentially reside in micas. With respect to Rb, Sr, and Ba the data are plotted in Figure 8 and the following points noted. (1) There is no correlation between Rb and Sr (Fig. 8A) and whereas the Sr contents are comparable to other suites [NASC, AWS, PCS in Fig. 8A; see also Nathan (1976), Scratch et al. (1984), Feng and Kerrich (1990), Wintsch et al. (1991)], the absolute Rb contents and Rb/Sr ratios (up to 24) are anomalously high compared to the aforementioned suites. Note how the greisen samples are also characterized by elevated Rb/Sr ratios. In contrast, the Ba versus Sr plot (Fig. 8B) separates the data such that the pelites are characterized by Ba/Sr of about 3 to 10 while the psammites have Ba/Sr of mostly <3; note that this distinction is mainly based on the enrichment of the pelites in Ba (>500 ppm). Although the Ba values are elevated relative to the averages shown in Figure 8B, comparable levels of enrich-
Table 1. Whole rock geochemistry for Meguma Group metasediments, East Kemptville deposit.

<table>
<thead>
<tr>
<th>ROCK</th>
<th>Pelite (SiO$_2$ &lt;60%)</th>
<th>Semi-Pelite (SiO$_2$ 60-70%)</th>
<th>Psammitic (SiO$_2$ &gt;70%)</th>
<th>Greisen EK-86-1227</th>
<th>Greisen EK-86-1154</th>
<th>AWS</th>
<th>NASC</th>
<th>BSA</th>
<th>PCS</th>
<th>WSS</th>
</tr>
</thead>
<tbody>
<tr>
<td>SiO$_2$</td>
<td>56.25 (1.94)</td>
<td>66.37 (2.01)</td>
<td>74.39 (2.57)</td>
<td>72.35</td>
<td>58.9</td>
<td>64.82</td>
<td>57.42</td>
<td>62.6</td>
<td>54.31</td>
<td></td>
</tr>
<tr>
<td>TiO$_2$</td>
<td>0.87 (0.07)</td>
<td>0.71 (0.07)</td>
<td>0.55 (0.07)</td>
<td>0.03</td>
<td>0.04</td>
<td>0.79</td>
<td>0.74</td>
<td>0.73</td>
<td>0.67</td>
<td></td>
</tr>
<tr>
<td>AI$_2$O$_3$</td>
<td>3.39 (0.31)</td>
<td>2.53 (0.22)</td>
<td>1.69 (0.20)</td>
<td>0.89</td>
<td>1.18</td>
<td>2.86</td>
<td>3.46</td>
<td>2.42</td>
<td>8.08</td>
<td></td>
</tr>
<tr>
<td>FeO</td>
<td>0.90 (0.39)</td>
<td>0.41 (0.30)</td>
<td>0.73 (0.24)</td>
<td>1.21</td>
<td>2.27</td>
<td>2.2</td>
<td>3.68</td>
<td>0.20</td>
<td>0.61</td>
<td></td>
</tr>
<tr>
<td>MnO</td>
<td>2.20 (0.57)</td>
<td>2.63 (0.84)</td>
<td>3.35 (0.70)</td>
<td>3.71</td>
<td>2.26</td>
<td>1.6</td>
<td>1.12</td>
<td>0.44</td>
<td>1.4</td>
<td></td>
</tr>
<tr>
<td>MgO</td>
<td>5.05 (0.96)</td>
<td>3.33 (0.67)</td>
<td>2.15 (0.72)</td>
<td>2.75</td>
<td>4.04</td>
<td>3.6</td>
<td>3.87</td>
<td>5.00</td>
<td>4.65</td>
<td></td>
</tr>
<tr>
<td>CaO</td>
<td>2.82 (0.60)</td>
<td>1.07 (0.01)</td>
<td>1.16 (0.36)</td>
<td>1.80</td>
<td>2.3</td>
<td>6.3</td>
<td>NA</td>
<td>5.80</td>
<td>5.9</td>
<td></td>
</tr>
<tr>
<td>N</td>
<td>99.26 (0.47)</td>
<td>99.62 (0.49)</td>
<td>100.10 (0.57)</td>
<td>99.55</td>
<td>99.75</td>
<td>100.33</td>
<td>100.63</td>
<td>100.63</td>
<td>99.63</td>
<td></td>
</tr>
</tbody>
</table>

Trace Elements (ppm):

<table>
<thead>
<tr>
<th>Element</th>
<th>Pelite (X+$\leq$1σ)</th>
<th>Semi-Pelite (X+$\leq$1σ)</th>
<th>Psammitic (X+$\leq$1σ)</th>
<th>Greisen EK-86-1227</th>
<th>Greisen EK-86-1154</th>
<th>AWS</th>
<th>NASC</th>
<th>BSA</th>
<th>PCS</th>
<th>WSS</th>
</tr>
</thead>
<tbody>
<tr>
<td>Ba</td>
<td>940 (367)</td>
<td>560 (139)</td>
<td>332 (164)</td>
<td>53</td>
<td>219</td>
<td>580</td>
<td>63.6</td>
<td>-</td>
<td>492</td>
<td>444</td>
</tr>
<tr>
<td>Rb</td>
<td>571 (317)</td>
<td>510 (191)</td>
<td>375 (194)</td>
<td>666</td>
<td>753</td>
<td>140</td>
<td>125</td>
<td>73</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>Sr</td>
<td>159 (77)</td>
<td>101 (89)</td>
<td>177 (51)</td>
<td>123</td>
<td>42</td>
<td>300</td>
<td>142</td>
<td>69</td>
<td>8</td>
<td>-</td>
</tr>
<tr>
<td>Y</td>
<td>52.0 (15.1)</td>
<td>42.5 (12.8)</td>
<td>34 (11.7)</td>
<td>43</td>
<td>49</td>
<td>35</td>
<td>-</td>
<td>25</td>
<td>25</td>
<td>25</td>
</tr>
<tr>
<td>Zr</td>
<td>158 (15.3)</td>
<td>162 (17.8)</td>
<td>193 (43.2)</td>
<td>23</td>
<td>18</td>
<td>200</td>
<td>-</td>
<td>-</td>
<td>120</td>
<td></td>
</tr>
<tr>
<td>Nd</td>
<td>35.8 (2.6)</td>
<td>30.5 (2.6)</td>
<td>36 (18.1)</td>
<td>38</td>
<td>40</td>
<td>40</td>
<td>-</td>
<td>8</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>Pr</td>
<td>25.8 (2.1)</td>
<td>18.6 (18.1)</td>
<td>14.7 (27.7)</td>
<td>34</td>
<td>36</td>
<td>14</td>
<td>-</td>
<td>-</td>
<td>20</td>
<td>-</td>
</tr>
<tr>
<td>Ga</td>
<td>240 (1.53)</td>
<td>357 (195)</td>
<td>25.6 (211)</td>
<td>1085</td>
<td>945</td>
<td>130</td>
<td>-</td>
<td>-</td>
<td>114</td>
<td>-</td>
</tr>
<tr>
<td>Ce</td>
<td>54.5 (65)</td>
<td>86.5 (62.2)</td>
<td>73.1 (46)</td>
<td>336</td>
<td>279</td>
<td>75</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>75</td>
</tr>
<tr>
<td>Ba</td>
<td>3.3 (8.7)</td>
<td>32.6 (61.9)</td>
<td>19.5 (44.4)</td>
<td>7</td>
<td>8</td>
<td>58</td>
<td>-</td>
<td>-</td>
<td>57</td>
<td>520</td>
</tr>
<tr>
<td>V</td>
<td>148 (15.5)</td>
<td>94.3 (10.5)</td>
<td>69.2 (9.9)</td>
<td>5</td>
<td>5</td>
<td>130</td>
<td>-</td>
<td>-</td>
<td>188</td>
<td>183</td>
</tr>
<tr>
<td>Cr</td>
<td>131 (28.0)</td>
<td>101 (9.9)</td>
<td>83.5 (6.5)</td>
<td>35</td>
<td>28</td>
<td>90</td>
<td>125</td>
<td>105</td>
<td>1195</td>
<td></td>
</tr>
<tr>
<td>Li</td>
<td>298 (106)</td>
<td>265 (88)</td>
<td>266 (152)</td>
<td>224</td>
<td>181</td>
<td>16</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>Mo</td>
<td>3.2 (2.6)</td>
<td>2.1 (2.6)</td>
<td>3.4 (2.9)</td>
<td>9</td>
<td>13</td>
<td>13</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>W</td>
<td>10.5 (10.3)</td>
<td>10.3 (0.7)</td>
<td>12.3 (7.3)</td>
<td>24</td>
<td>24</td>
<td>28</td>
<td>-</td>
<td>23</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>As</td>
<td>17.3 (14.1)</td>
<td>27.3 (23.3)</td>
<td>40.5 (99)</td>
<td>5</td>
<td>5</td>
<td>28</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>Bi</td>
<td>3.9 (19.9)</td>
<td>3.6 (11.1)</td>
<td>4.5 (18.8)</td>
<td>3</td>
<td>3</td>
<td>28</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>Ag</td>
<td>0.5 (0.08)</td>
<td>0.5 (0.0)</td>
<td>0.5 (0.0)</td>
<td>0.5</td>
<td>0.5</td>
<td>0.5</td>
<td>-</td>
<td>0.41</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>Sb</td>
<td>7.3 (3.6)</td>
<td>5.6 (0.9)</td>
<td>5.2 (0.5)</td>
<td>5</td>
<td>8</td>
<td>2.0</td>
<td>-</td>
<td>-</td>
<td>1.3</td>
<td>-</td>
</tr>
<tr>
<td>Sn</td>
<td>10.0 (0.0)</td>
<td>10.0 (0.0)</td>
<td>10.0 (0.0)</td>
<td>15</td>
<td>24</td>
<td>3.7</td>
<td>2.4</td>
<td>5.9</td>
<td>1.7</td>
<td></td>
</tr>
<tr>
<td>As</td>
<td>8.5 (8.5)</td>
<td>10.9 (1.6)</td>
<td>6.2 (3.7)</td>
<td>1.5</td>
<td>2.2</td>
<td>13</td>
<td>14.6</td>
<td>16</td>
<td>27</td>
<td></td>
</tr>
<tr>
<td>Hf</td>
<td>3.3 (2.4)</td>
<td>4.7 (1.6)</td>
<td>4.9 (3.4)</td>
<td>2.1</td>
<td>2.1</td>
<td>2.1</td>
<td>6.3</td>
<td>-</td>
<td>4.7</td>
<td>-</td>
</tr>
<tr>
<td>Th</td>
<td>10.3 (0.9)</td>
<td>7.6 (28)</td>
<td>6.0 (28)</td>
<td>4.6</td>
<td>3.8</td>
<td>12</td>
<td>12.3</td>
<td>-</td>
<td>4.7</td>
<td>-</td>
</tr>
<tr>
<td>Ta</td>
<td>1.6 (0.2)</td>
<td>1.0 (1.0)</td>
<td>0.8 (0.4)</td>
<td>11.3</td>
<td>19.3</td>
<td>2.8</td>
<td>1.1</td>
<td>22</td>
<td>59</td>
<td></td>
</tr>
<tr>
<td>Co</td>
<td>15.0 (1.2)</td>
<td>6.9 (2.3)</td>
<td>5.0 (3.1)</td>
<td>0.4</td>
<td>2.6</td>
<td>19</td>
<td>27</td>
<td>-</td>
<td>-</td>
<td>-</td>
</tr>
</tbody>
</table>

Note that for some elements (Hf, Th, Ta, Co) the X+$\leq$1σ values are based on 2 pelites, 1 semipelite and 6 psammites only.

AWS = average world shale from Wedepohl (in Nathan, 1976).
BSA = argillite; from Bhatia (1985) for Bendigo metaturbidites, Australia.
PSC = average Aphebian shale (n=326) of the Canadian Shield from Cameron and Garrels (1980).
WSS = Witwatersrand Supergroup, Bogoysens shale (Wronkiewicz and Condie, 1987).
Table 2. Some ranges and averages for metals in sedimentary rocks.

<table>
<thead>
<tr>
<th>Element</th>
<th>Mo (ppm)</th>
<th>Zn (ppm)</th>
<th>Cu (ppm)</th>
<th>As (ppm)</th>
<th>Sb (ppm)</th>
<th>W (ppm)</th>
<th>Sn (ppm)</th>
<th>Pb (ppm)</th>
<th>Reference</th>
</tr>
</thead>
<tbody>
<tr>
<td>Mo</td>
<td>50-150</td>
<td>80-170</td>
<td>0.1-0.7</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>2.2-4</td>
<td>Feng and Kerrich (1990)</td>
</tr>
<tr>
<td>Zn</td>
<td>2-180</td>
<td>5-28</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>Maas and McCulloch (1991)</td>
</tr>
<tr>
<td>Cu</td>
<td>48-120</td>
<td>1-13</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>12-30</td>
<td>Floyd and Leveridge (1987)</td>
</tr>
<tr>
<td>As</td>
<td>74±9</td>
<td>11-2</td>
<td>1-170</td>
<td>0.1-0.7</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>15±1</td>
<td>Bhatia and Crook (1986)</td>
</tr>
<tr>
<td>Sb</td>
<td>52±8</td>
<td>8±3</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>24±1</td>
<td>-</td>
<td>Bhatia and Crook (1986)</td>
</tr>
<tr>
<td>W</td>
<td>-</td>
<td>28.4</td>
<td>2-180</td>
<td>1-70</td>
<td>-</td>
<td>-</td>
<td>36±6</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>Sn</td>
<td>-</td>
<td>2-24</td>
<td>Feng and Kerrich (1990)</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Pb</td>
<td>42.2</td>
<td>2-82</td>
<td>Maas and McCulloch (1991)</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>54.5</td>
<td>12-30</td>
<td>Floyd and Leveridge (1987)</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>62.6</td>
<td>15±1</td>
<td>Bhatia and Crook (1986)</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>7.37-77</td>
<td>11-2</td>
<td>-</td>
<td>4.6</td>
<td>3.3</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td></td>
</tr>
<tr>
<td></td>
<td>8.52-111</td>
<td>0-46</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td></td>
</tr>
<tr>
<td></td>
<td>9.95</td>
<td>45</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td></td>
</tr>
<tr>
<td></td>
<td>10.26</td>
<td>175±42</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td></td>
</tr>
</tbody>
</table>

1 only 2 analyses >50 ppm
2 average for continental island arc sediments
3 average for active continental margin
4 Archean weighted average (n=406)
5 Phanerobian weighted average (n=326)
6 average Phanerian shale (Cameron and Garrels, 1980)
7 greyscale (unmineralized)
8 argillites (unmineralized)
9 average shale
10 data for altered sediments (schists) 0-1 km from ore body (Panasqueira W-Sn deposit, Portugal)
11 only 3 analyses >44 ppm

Table 3. Rare earth element geochemistry for Meguma Group metasediments, East Kemptville deposit.

<table>
<thead>
<tr>
<th></th>
<th></th>
<th></th>
<th></th>
<th></th>
<th></th>
<th></th>
<th></th>
<th></th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td>La</td>
<td>30.00</td>
<td>37.10</td>
<td>29.10</td>
<td>36.40</td>
<td>17.70</td>
<td>3.04</td>
<td>21.30</td>
<td>25.00</td>
<td>21.85</td>
</tr>
<tr>
<td>Ce</td>
<td>66.30</td>
<td>74.60</td>
<td>57.10</td>
<td>73.70</td>
<td>36.70</td>
<td>10.70</td>
<td>44.50</td>
<td>52.70</td>
<td>45.70</td>
</tr>
<tr>
<td>Nd</td>
<td>30.40</td>
<td>33.80</td>
<td>26.80</td>
<td>34.30</td>
<td>16.90</td>
<td>7.26</td>
<td>19.30</td>
<td>23.20</td>
<td>21.70</td>
</tr>
<tr>
<td>Sm</td>
<td>5.73</td>
<td>6.61</td>
<td>5.12</td>
<td>6.10</td>
<td>3.13</td>
<td>1.53</td>
<td>3.77</td>
<td>4.16</td>
<td>3.80</td>
</tr>
<tr>
<td>Eu</td>
<td>1.35</td>
<td>1.50</td>
<td>1.13</td>
<td>1.18</td>
<td>0.67</td>
<td>0.14</td>
<td>0.68</td>
<td>0.86</td>
<td>0.77</td>
</tr>
<tr>
<td>Gd</td>
<td>6.71</td>
<td>6.22</td>
<td>4.44</td>
<td>6.69</td>
<td>4.19</td>
<td>2.66</td>
<td>3.87</td>
<td>4.51</td>
<td>3.95</td>
</tr>
<tr>
<td>Tb</td>
<td>0.92</td>
<td>0.90</td>
<td>0.75</td>
<td>0.88</td>
<td>0.53</td>
<td>0.24</td>
<td>0.57</td>
<td>0.58</td>
<td>0.52</td>
</tr>
<tr>
<td>Yb</td>
<td>2.77</td>
<td>2.59</td>
<td>2.18</td>
<td>2.55</td>
<td>1.50</td>
<td>0.27</td>
<td>1.76</td>
<td>1.84</td>
<td>1.58</td>
</tr>
<tr>
<td>Lu</td>
<td>0.43</td>
<td>0.37</td>
<td>0.30</td>
<td>0.36</td>
<td>0.24</td>
<td>-</td>
<td>0.27</td>
<td>0.28</td>
<td>0.28</td>
</tr>
<tr>
<td>ΣREE</td>
<td>144.61</td>
<td>163.69</td>
<td>126.92</td>
<td>162.16</td>
<td>81.16</td>
<td>25.84</td>
<td>96.02</td>
<td>113.13</td>
<td>99.84</td>
</tr>
<tr>
<td>(La/Lu)_N</td>
<td>7.15</td>
<td>10.28</td>
<td>9.94</td>
<td>10.36</td>
<td>7.56</td>
<td>8.08</td>
<td>9.15</td>
<td>8.29</td>
<td>8.59</td>
</tr>
<tr>
<td>(Eu/Eu*)_N</td>
<td>0.67</td>
<td>0.72</td>
<td>0.73</td>
<td>0.57</td>
<td>0.57</td>
<td>0.21</td>
<td>0.54</td>
<td>0.61</td>
<td>0.77</td>
</tr>
<tr>
<td>(Gd/Yb)_N</td>
<td>1.94</td>
<td>1.92</td>
<td>1.63</td>
<td>2.10</td>
<td>2.24</td>
<td>1.07</td>
<td>1.76</td>
<td>1.96</td>
<td>2.00</td>
</tr>
<tr>
<td>%SiO₂</td>
<td>56.46</td>
<td>58.66</td>
<td>69.09</td>
<td>70.61</td>
<td>76.42</td>
<td>72.35</td>
<td>71.00</td>
<td>77.69</td>
<td>70.70</td>
</tr>
</tbody>
</table>

Pelites: EK-86-1407, 1021; Semipelites: EK-86-1079; Psammites: EK-86-1406, 1240, 1255, 1410, 1077A, 1414; Greisens: EK-86-1154, 1227
Fig. 5. Binary element diagrams for major element geochemistry of the East Kemptville metasedimentary rocks (□) and greisens (*). Abscissa is wt. % Al₂O₃ in all the plots. In the top left plot the fields of psammites, semipelites and pelites are outlined (from left to right, respectively) and the two greisens are indicated by the asterisk. Note the generally good correlation of elements and lack of dispersion toward the two greisen samples.

Ferromagnesian elements (Cr, V, Ni, Co, Sc)

The ferromagnesian or transition elements show a progressive enrichment from psammitic through semipelitic to pelitic lithologies, as is typical of these elements in sedimentary rocks (Table 1 and below). There is a strong positive covariation in all the ferromagnesian trace elements with MgO, total Fe and TiO₂ (e.g., Fig. 9B) as expected. In terms of the absolute abundances of the ferromagnesian elements, levels of enrichment comparable to other Meguma Group samples (Fig. 7), the NASC (Gromet et al., 1984; Table 1) and other Lower Paleozoic sedimentary lithologies (Nathan, 1976; Scratch et al., 1984; Floyd and Leveridge, 1987; Wintsch et al., 1991). However, the samples are depleted in...
these elements with respect to what is found in Archean rocks (Taylor and McLellan, 1985; see also AWS in Table 1). In Figure 9A the samples are plotted in the Cr versus Ni diagram and, although not shown, the data fall in the field defined for Paleozoic sediments (Taylor and McLellan, 1985). Note too, that in Figure 9A and all other binary element diagrams (plots of TiO₂, Cr, Sc and Ni versus V) there are continuous trends. Compared to the NASC (Table 1) there are similar elemental abundances and also ratios for Cr/V, Ni/V, Sc/V and Co/V (0.5, 0.83, 0.21, 0.08 versus 0.96, 0.44, 0.10, 0.20 for the Meguma Group versus NASC, respectively).

As with other trace elements, the two greisen samples deviate from the above trends and averages for the metasedimentary rocks. In this case, there is considerable depletion of these samples with respect to the ferromagnesian elements, similar to that found in the fresh leucogranite (Kontak, 1990a).

**High field strength elements (Hf, Ta, Y, Nb, Zr), U and Th**

These elements are generally considered to be resistant to weathering and alteration processes (Taylor and McLellan, 1985; Bhatia and Crook, 1986), and thus can provide reliable estimates of original chemical variation. In the East Kemptville lithologies there is slight enrichment of Th, Nb, Y and Zr in the psammites over the pelite while Ta, Hf and U remain fairly constant (Table 1). The absolute levels of elemental enrichment, exclusive of U, are comparable to most post-Archean suites (e.g., Nathan, 1976; Scratch et al.,
Fig. 7. Normalized spidergram plots for average pelite and psammite compositions from East Kemptville deposit area. The data in Table 1 are normalized to the average compositions of Meguma Group lithologies using the data of Douma (in prep.). Note the coherent behaviour of the data sets with similar relative enrichment and depletion compared to Douma’s data set.

1984; Floyd and Leveridge, 1987) and some Archean suites for the HFSE only (e.g., Feng and Kerrich, 1990; Maas and McCulloch, 1991). Most notable is the elevated absolute enrichment of U at 10 ppm for all lithologies and consequently depressed Th/U and K/U ratios of 0.6-1.0 and <4100, respectively. These ratios compare to the normal Th/U and K/U ratios of 3-4 and 8000-12000, respectively, for the NASC, AWS (Table 1) and crustal averages (Taylor and McLellan, 1985; McLellan and Taylor, 1991).

In the two greisens there is considerable enrichment and depletion with respect to the HFSE, U and Th (Table 1), with average concentrations of U (18 ppm), Ta (15 ppm) and Nb (39 ppm) well above those seen in the metasedimentary rocks whereas Zr (21 ppm) and Th (4.2 ppm) are depleted relative to the metasedimentary rocks.

Sn, Cu, Zn, Pb, Ag, Mo, W, As, Bi, Sb

The ore forming and related elements (ORFE) at the East Kemptville deposit show highly variable degrees of enrichment between samples when compared to the average Meguma Group metasedimentary rock (Fig. 7). Table 2 lists some typical values of the ORFE for comparative purposes. Bismuth, Ag, Mo, and Sb are near or below their respective detection limits (note that Sb values are artificially inflated because of the high detection limit, i.e., 10 ppm) and show no enrichment. Lead is similar in its lack of enrichment in any lithology and shows typical background abundances compared to the data in Table 2. In contrast, Zn, Cu, W and Sn are variably enriched and show large variances in the data because of the erratic nature of mineralization; hence, elemental abundances vary from background levels to 100’s of ppm (e.g., Sn at about 500 ppm). Compared to the background levels tabulated in Table 2, the enrichment factors are approximately as follows: Sn = 20-50; W = 2; Zn = 2-5; Cu = 1-10.

In Figure 10 binary element diagrams illustrate the relationship between Sn, the base metals and Rb (representative of the LILE): (1) the best correlation is between Zn and Cu, and to a lesser degree Zn and Sn; (2) the most elevated Sn values are for the altered leucogranites and these are associated with depleted levels of Zn, Cu and Rb; (3) there is generally a trend of increased LILE content, as represented by Rb, and the ORFE; and (4) the greisen samples generally plot away from the extensions of the enrichment trends.
Fig. 9. Binary element diagrams for East Kemptville metasedimentary rocks (○) and greisens (*) (B,C,D,E) showing the covariation of ferromagnesian trace elements; note the excellent correlation in all plots with no dispersion of the data towards the greisen samples. Also note that in Figure 9A the data plot within the field (not outlined in diagram) for post-Archean fine-grained sedimentary rocks (from Taylor and McLellan, 1985).

defined by the metasedimentary data. These data would suggest that there is considerable enrichment of the OFRE in the metasedimentary rocks and that this enrichment is quite pervasive.

Arsenic shows no absolute enrichment compared to the background levels listed in Table 2. Only 3 of the 36 analyzed samples have >100 ppm As while the remaining are all <50 ppm. This is noted because the general enrichment of As in mineralized zones within the ore deposit occurs in close proximity to the metasediment-granite contact, especially in the north end of the deposit (C. Poulin, personal communication, 1990).
Fig. 10. Binary element diagrams for East Kemptville metasedimentary rocks (□) and greisens (*) showing the covariation of various ore forming elements (ORFE; Cu, Zn, Sn) and Rb (representative of LIL elements in general). Note that there is a general tendency for the ORFE to behave compatibly, while Rb first increases and then rapidly decreases as the content of ORFE increases (e.g., Zn, Sn).

**Rare-earth elements**

REE data have been obtained for 11 samples, including 2 pelites, 1 semipelite and 6 unaltered psammites and the 2 greisens. The data are presented in Table 3, including the significant ratios (e.g., (La/Lu)_N, Eu/Eu*), and the chondrite-normalized profiles are illustrated in Figure 11. As already noted above, the apparent inflections at Gd in the chondrite-normalized profiles for all the samples, especially the two greisen samples, are anomalous and attributed to analytical problems. However, since the Gd inflection is not great in the metasedimentary rocks (e.g., from extrapolating between...
Fig. 11. Chondrite-normalized rare-earth element diagrams for East Kemptville metasedimentary rocks and greisens (EK-86-1154, 1227). The inset diagram shows the chondrite-normalized profiles for unaltered Meguma Group metasedimentary rocks from the Beaver Dam area (unpublished data of Kontak). Note the extreme depletions of the REE in the greisen samples and the generally consistent patterns for the variably mineralized metasedimentary rocks.

Tb-Sm), it is not considered to influence significantly the ratios discussed [e.g., (Gd/Yb)_n].

The metasedimentary rocks have typical REE abundances and patterns for fine-grained, post-Archean sediments (e.g., Haskin et al., 1966; Nance and Taylor, 1976; Taylor and McLellan, 1985; Grauch, 1989; McLellan, 1989; McLellan and Taylor, 1991), including metaturbidites (Nathan, 1976; Cullers et al., 1974; Jenner et al., 1981; Floyd and Leveridge, 1987; Feng and Kerrich, 1990) with \( \Sigma_{REE} \) (ppm) of 154 ± 10, 127 and 109 ± 25 for pelites, semipelite and psammites (unaltered), respectively and \((La/Lu)_n = 8.7 ± 1.6\) for the pelites and 8.6 ± 0.8 for the psammites. The Eu/Eu* values are similar in all the metasedimentary rocks at 0.57 to 0.77 and \((Gd/Yb)_n < 2\) in the fresh sediments; both values are typical for post-Archean metaturbidites (Fig. 12; McLellan, 1989; McLellan and Taylor, 1991). The REE patterns for the East Kemptville metasedimentary rocks also resemble those of other Meguma Group metasedimentary rocks (see inset in Fig. 11) within the central and eastern Meguma Terrane (Liew, 1978; Muecke and Clarke, 1981; Kontak, unpublished data).

For the two greisens there are several differences compared to the metasedimentary REE data: (1) depletion of the \( \Sigma_{REE} \) of 25 ppm; (2) an increase in the Eu anomaly (0.21-0.32 for Eu/Eu*); and (3) a convex shape to the LREE pattern which is enhanced by depletion in La. However, compared to the REE data for the leucogranites (Kontak, 1990a), there is general consistency, except for the inflection at Gd, indicating that at least for the two samples analyzed there has been limited mobility of the REE during hydrothermal alteration.

Geochemical Profiles in the Metasedimentary Rocks

The samples located in Figure 3 have been grouped into three profiles referred to as A, B and C. The following comments are noted with respect to these profiles: (1) profile C includes all the samples along the granite-metasedimentary rock contact progressing from the south side (i.e., starts with sample 1021) and moving counter-clockwise around the granite-sediment contact (i.e., finishes with sample 1027). This profile also includes the two roof pendent samples (1021, 1079; see Fig. 3b); (2) profile A and C include the two greisen samples (1154 and 1227, respectively); (3) profile B includes samples 1239 and 1240 as the NW extension continuing from sample 1415 (see Fig. 3b); and (4) sample 1401 forms part of profile A (after samples 1154 and 1402).

The geochemical data are presented in these three profiles in Figure 13 in order to evaluate the potential relationship between elemental abundance and proximity to the granite. The SiO₂ profile illustrates the typical intercalated nature of the pelitic to psammitic lithologies and the variable thickness of units (e.g., section B). There is a decrease in absolute Rb content away from the granite when profiles A and B are compared to profile C. The relative enrichment of Rb in this profile is again noted with respect to the ORFE (Cu, Zn, Sn, As). For Cu and Zn, and to a lesser degree As, profile C is consistently enriched compared to the other profiles. Interestingly, the greatest enrichment in Sn is found for two samples from profile B, although on average profile C again has a greater relative enrichment compared to sample from profiles A and B.
The emplacement of intrusive rocks into upper crustal levels generally results in the development of hydrothermal convection cells that promote the circulation of heated waters during the cooling history of the intrusion. The result of this hydrothermal activity is the formation of certain hydrothermal mineral deposits, especially massive sulphides (Franklin et al., 1981), Cu-Mo porphyries (Titley and Beane, 1981; White et al., 1981) and skarns (Einaudi et al., 1981; Kwak, 1987). That such cells occur and involve the introduction of magmatic-derived fluids is well documented from the study of stable isotopes (Taylor, 1974, 1978), fluid inclusions (Roedder, 1984), petrological investigations (Ferry, 1978) and integrated studies (e.g., Gerstner et al., 1989; Novick and Labotka, 1990). By analogy to this aforementioned work and the well-established capacity for granites to exsolve fluids late in their crystallization history (e.g., Burnham, 1979), it is reasonable to expect that there has been movement of fluid from the leucogranite at East Kemptville through the surrounding metasedimentary rocks. There have, however, been few detailed studies that have documented the magnitude of fluid flow and elemental enrichment resulting from infiltration of magmatic-derived fluids (Cathles, 1981).

In the case of Sn deposits, two recent studies focused on the extent of primary dispersion about mineralized centres are particularly relevant to this investigation because of the similarity of geological setting and mineralization. Polya (1987, 1988) examined the dispersion of Sn, W, Zn, Cu and As around the Panasqueira ore body, Portugal, and found that the pelitic schists surrounding the deposit contain anomalous values of these elements (see Table 2) up to 8 km from the mineralized centre. Polya (1987) also estimated that >90% of the metals, except for W (~30%), are dispersed around the deposit. Meanwhile, Beer and Ball (1986) showed that of the five mineralized centres studied in southwest England, Sn is enriched in the surrounding metasedimentary rocks, and to a lesser extent W, up to at least 1 km from the known point source.

In the case of the East Kemptville deposit area, the lithogeochemical data indicate that there has been little modification of the major element chemistry and most of the trace element chemistry. For example, in terms of the major elements, comparison to either the average chemical signature of unmineralized Meguma Group rocks (Fig. 7) or binary discriminant diagrams (Fig. 6) indicates that the metasedimentary rocks appear unaltered. For the trace elements, use of discriminant diagrams [e.g., Th/Sc/Zr, La/Th/Sc (Bhatia and Crock, 1986) or La-Th, Sc-Th (McLellan, 1989); note that these diagrams are not shown] and comparison to the unmineralized data base for the Meguma Group (Fig. 7) indicate little mobility or enrichment for most elements. However, for Sn, U, Rb, Li and Zn there is variable enrichment compared to typical background values in metasedimentary rocks (Table 2) and also compared to the average values established for the Meguma Group (Fig. 7). Some other elements (Bi, W, Sb) may be slightly enriched but higher detection limits do not permit an unambiguous conclusion.

To further examine the spatial relationship of elemental enrichment profiles about the granite were constructed. These profiles (Fig. 13) indicate that generally speaking there is slight enrichment of some elements immediately adjacent to the granite. However, the general absence of veins in the metasedimentary rocks containing ore minerals (e.g., cassiterite, sphalerite) suggests that the elemental enrichment may reflect the presence of fine-grained, disseminated minerals and/or variable amounts of substitution within the secondary micaceous minerals (e.g., muscovite). In either case, the profiles indicate that both the amount and extent of enrichment within the metasedimentary rocks is exceptionally limited compared to the examples cited above.

The general positive correlation among and between the elevated trace elements suggest that their enrichment is related to a single hydrothermal event. Petrographically this enrichment is manifested by the highly variable amount and growth of new minerals. The development of white mica ± other minerals, both as disseminations and along veinlets or microfractures, is the most commonly observed phenomenon and can account for the elevated Rb and Li values. Additional observations include: (1) the growth of chlorite, tourmaline, fluorite, cassiterite and other ore minerals, and (2) the alteration of biotite. The presence of tourmaline in the metasedimentary rocks is noted because of its absence in the local granite.

The processes responsible for the elemental enrichment may have preceded the main mineralizing event at East Kemptville and, therefore, reflect expulsion of early-stage orthomagmatic fluids or, conversely, the elemental enrichment may be related to the same fluids which later were responsible for the main Sn and base metal mineralization. We have no way of distinguishing between these two scenarios, but consider the latter to be more plausible.
Fig. 13. Geochemical profiles of elements (Cu, As, Zn, Sn, Si, Rb) in the country rocks at the East Kemptville Sn deposit. See Figure 3 for location of the profiles. Note that profile C is for rocks from along the granite-sediment contact starting with samples along south side of the deposit area in Figure 3. The two greisen samples are included in the profiles and indicated by the appropriate symbol (*).
The lack of visible, bona fide veinlets in the metasedimentary rocks, combined with the generally fine-grained, disseminated nature of muscovite suggests that fluids were transported through the sediments along microfractures, grain boundaries or a structural anisotropy such as cleavage. This contrasts, for example, with the richly mineralized metasedimentary-hosted Sn-W-base metal veins of the Central Andean tin belt (e.g., Palma and Clark, 1982) where large fissure veins acted as fluid channelways for magmatically-derived, metal-bearing fluids. In any event, hydrofracturing was clearly not the preferred mechanism for dispersal of fluid at East Kemptville. Although there is clearly evidence of infiltration of a granite-derived fluid, the metasedimentary rocks may still be considered to have acted as a relatively impermeable trap for the mineralizing fluids as Richardson et al. (1982) originally suggested. This process would have the effect of promoting build up of a fluid reservoir and this would have enhanced the development of mineralization within the granite.

Evidence that a hydrothermal circulation cell was present at East Kemptville may be examined in light of isotopic studies done at the deposit (Kontak, 1988, 1990b). Neither S nor O isotope data indicate that an externally-derived fluid equilibrated with the metasedimentary rocks infiltrated the granite, while Sr isotopic data (Richardson et al., 1990b) proved inconclusive is discerning the role, if any, of the metasedimentary rocks played in terms of the deposit. In addition, although the presence of graphite of hydrothermal origin occurs in the mineralized greisens (Kontak, 1988), we have no idea how it entered the mineralizing system or at what structural level.

Finally, the lack of Mn-enrichment, either chemically and mineralogically, indicates that the manganiferous GHT horizon has not played a significant role in the genesis of the East Kemptville tin deposit, as suggested by Sangster (1990). The stratigraphy, bulk rock chemistry and mineralogy contrasts, for example, with that observed at the Duck Pond deposit (Hattie, 1989; Pitre and Richardson, 1989; Kontak et al., 1990), where the GHT horizon is present.

**Conclusions**

A lithogeochemical and petrographic study of 36 samples of metasedimentary rocks from the East Kemptville deposit area indicates that a primary dispersion halo exists. Although the rocks appear relatively unaltered and highly indurated, petrographic evidence suggestive of fluid passage includes the presence of fine-grained, disseminated muscovite and lesser amounts of coarser fracture-controlled muscovite, chlorite, tourmaline, cassiterite, ore minerals, fluorite, topaz and carbonate. Geochemically the rocks are characterized by normal major element chemistry when compared to the average for unaltered Meguma Group rocks of similar lithology. With regards to the trace elements, the REE, HFSE and most transition or ferromagnesian elements retain apparently primary elemental abundances. However, there is appreciable enrichment of some LILE (Rb, Li, U) and ore elements (Sn, Zn) which reflects infiltration of the ore fluid from the adjacent mineralized granite/greisens. Although fluid did infiltrate and interact with the metasediments, transport must have been restricted to microfractures, grain boundaries and planar fabrics such as cleavage, as no fissure veins are present. Consequently, fluid flow may have been retarded which would have favoured the build up of fluid within the granite carapace and promoted the mineralizing process as originally suggested by Richardson et al. (1982).

**Acknowledgements**

Field work at the East Kemptville deposit by D. Kontak has been funded through Canada-Nova Scotia Mineral Development Agreements (1984-1989, 1990-1992) and the cooperation and support of the personnel at Rio Kemptville Tin Corporation is sincerely acknowledged. Funding for the analyses for this project were provided by the CNSMDA and National Science and Engineering Research Council grant to J. Dostal. The capable assistance in manuscript preparation by personnel at the Nova Scotia Department of Natural Resources is once again appreciated. Comments on the manuscript by D. Mossman and R. Raeside contributed to improvements and are appreciated. This paper is published with permission of the Director of the Mineral Resources Division, Nova Scotia Department of Natural Resources.

**References**


— 1990c. A geological and geochemical study of the East Kemptville Sn-Zn-Cu-Ag deposit, southern Nova Scotia, Canada: mineralization hosted by a ca. 370 Ma topaz-muscovite leucogranite. In Program with Abstracts, 8th IAGOD Symposium, Ottawa, Ontario, p. 34.


In press. Geochemical, geochemical and isotopic studies of the East Kemptville (Sn-Zn-Cu-Ag) deposit, Yarmouth County, Nova Scotia, Canada. Proceeding of the Eighth Quadrennial IAGOD Symposium, E. Schweizerbart'sche Verlagshandlung, Stuttgart, Germany.


Sawyer, E.W. 1916. The influence of source rock type, chemical weathering and sorting on the geochemistry of clastic sediments from the Queticco metasedimentary belt, Superior Province, Canada. Chemical Geology, 55, pp. 77-95.


