Rb-Sr isotopic characterization of pegmatite dikes from the Tuolumne Intrusive Suite, California, USA

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ABSTRACT

Pegmatite dikes are common in granitic plutons and reflect crystallization of the fluid-rich last increments of the system. I measured $^{87}\text{Sr}/^{86}\text{Sr}$ ratios for several pegmatites from the Tuolumne Intrusive Suite in the Sierra Nevada batholith, California to examine the possible role of mass transfer between wall rocks and intruding magma. Variation in isotopic composition between wall rocks and plutons provides a sensitive test of fluid-rock interaction.

Meter-scale pegmatite dikes along the margins of the intrusive suite cut both wall rocks and granitoids (including tonalite, granodiorite, and granite). Initial $^{87}\text{Sr}/^{86}\text{Sr}$ ratios in granitoids along an east-west traverse range from 0.7057 to 0.7066 (Kistler et al, 1986; Bateman et al., 1988; Gray et al., 2008). Wall rocks along the margins of the suite display variable $^{87}\text{Sr}/^{86}\text{Sr}$, ranging from 0.710 to 0.774 (Mills et al., 2009), significantly more radiogenic than the igneous materials. Age-corrected pegmatite $^{87}\text{Sr}/^{86}\text{Sr}$ ratios range from 0.706 to 0.709.

Pegmatites have higher $^{87}\text{Sr}/^{86}\text{Sr}$ ratios than granitoids, but lower than the metasedimentary wall rocks. I interpret these data as indicating interaction between hydrous fluids from late-stage granite crystallization and wall rocks. Pegmatites in the Tuolumne Intrusive Suite represent the final stages of pluton development, and hydrothermal circulation could drive fluid paths that traverse wall rocks and the crystallizing plutons, resulting in $^{87}\text{Sr}/^{86}\text{Sr}$ ratios between those of the two end members.
INTRODUCTION

Interaction between intruding magma and surrounding wall rock is broadly assumed, but geochemical data for this interaction are difficult to obtain. However, pegmatites may record mass exchange between intruding magmas and wall rocks. Although most studies of pegmatites concern their classification and mineralogy, pegmatites also represent the final stages of crystallization of plutonic systems. Processes governing pegmatite growth are still poorly understood.

A comparison of initial \(^{87}\text{Sr}/^{86}\text{Sr}\) ratios across lithologies may indicate whether pegmatites record mass exchange; initial \(^{87}\text{Sr}/^{86}\text{Sr}\) is a robust indicator of interaction because these isotopes do not fractionate during crystallization. Initial \(^{87}\text{Sr}/^{86}\text{Sr}\) ratios are far greater in wall rocks than in adjacent plutonic rocks (Kistler et al., 1986; Bateman et al., 1988; Gray et al., 2008; Mills et al., 2009; Gross, 2015). I sampled pegmatites at four locations throughout the Tuolumne Intrusive Suite in the Sierra Nevada batholith and calculated their initial \(^{87}\text{Sr}/^{86}\text{Sr}\) ratios. If the pegmatites record interaction between magma-derived hydrothermal fluids and wall rock, their isotopic ratios should be intermediate between the groups.

Pegmatite Petrogenesis

Jahns and Burnham (1969) described mechanisms for producing pegmatite from cooling granitic magma. As magma crystallizes anhydrous phases, the remaining liquid becomes enriched in water. Once the melt phase becomes water-saturated, fluid exsolves as a new phase, and pegmatites crystallize rapidly at low temperatures from such hydrothermal phases (Jahns and Burnham, 1969). This fractionation model for producing pegmatite was supported by Roda-Robles (2012) and Ballouard et al. (2015), which argued that pegmatites were derived from host granitoids by fractional crystallization.
Water reduces magma viscosity, nucleation rate, and polymerization, while increasing diffusivity and the ability for a melt to dissolve surrounding material (Simmons and Webber, 2007; Nabelek et al., 2010). Less viscous magma can more easily propagate as dikes, and increased capacity for magmas to dissolve may enable interaction with intruded materials. Elevated diffusivity and lower nucleation rates promote crystal growth, generating coarse textures characteristic of pegmatite (Jahns and Burnham, 1969, Davidson and Thomas, 2012).

Kontak et al. (2002) noted that devolatization of metasedimentary xenoliths may be a potential source of water for hydrothermal fluids. However, simultaneous release of CO₂ would reduce the activity of water and offset a metasedimentary water contribution. A granitic magma containing a few percent water by mass should concentrate enough water in the remaining melt during crystallization to exsolve a fluid phase by later stages of cooling.

Ballouard et al. (2015) argued that the Hercynian Guérande leucogranite of the Armorican Massif in France underwent alteration by hydrothermal fluids exsolved during cooling. Multiple pegmatite dikes and quartz veins through the leucogranite are thought to have crystallized from hydrothermal fluids. Therefore, Ballouard et al. (2015) concluded that fluids which caused hydrothermal alteration were involved in crystallizing pegmatites in the same plutonic body. Pegmatites have been documented as records of agents of hydrothermal alteration.

**GEOLOGIC SETTING**

The Sierra Nevada batholith (SNB) in eastern California (Fig. 1) makes up much of the Sierra Nevada range and spans about 600 km, trending NNW. The width of the batholith averages 100 to 120 km. The SNB lies between the Basin and Range to the east and the Great Valley to the west; the south end of the SNB is separated from the Mojave Desert by the Garlock fault, and the north end abuts the Cascade Range. Emplacement ages in the SNB are younger
from west to east, with Late Cretaceous intrusive bodies forming the east-central region. The Tuolumne Intrusive Suite (TIS) of Yosemite National Park, California, in the east-central portion of the SNB, is composed of several nested plutons.

The TIS is divided into several units based on composition and texture. The oldest, exterior units of the TIS are contemporaneous granodiorites and tonalites of Glen Aulin (Kga) and Kuna Crest (Kkc) on the west and east, respectively. Inward are the equigranular (Khde) and porphyritic (Khdp) Half Dome Granodiorite, Cathedral Peak Granodiorite (Kcp), and the Johnson Granite Porphyry (Kjp). Moving from margin to core, rocks are generally more felsic with coarser, porphyritic textures, with the exception of Kjp. Hornblende, biotite, and titanite phenocrysts are abundant in Khd, and megacrystic K-feldspar is present in Kcp. Tabular rafts of megacrystic K-feldspar are present in Kcp. Contacts between these units are generally gradational (Bateman and Chappell, 1979).

Metamorphic wall rocks along the margins of the TIS include metasedimentary and metavolcanic strata. The primary wall rocks of interest here are those near May Lake in the west and Saddlebag Lake in the East (Fig. 1). The Saddlebag Lake roof pendant consists of Devonian to Triassic metavolcanic and metasedimentary rocks (Brook, 1977; Lahren et al., 1990). Metamorphic rocks near May Lake consist of calc-silicate rocks, quartzite, pelitic quartzite, pelitic hornfels, and marble (Rose, 1957; Mills et al., 2009). Foliated metamorphic rocks at May Lake strike northeast and dip northwest.

Pegmatite dikes are widespread throughout the TIS. Pegmatites along the eastern margin near Saddlebag lake (Fig. 1) intrude both Kcp and surrounding metavolcanic rocks and strike oblique to foliation. In the west, near May Lake, thicker pegmatite dikes intrude granodiorite of Kga. Rose (1957) described pegmatites containing andalusite and corundum within the May
Lake screen. The modal assemblage of pegmatites described in Rose (1957) varies from that of the TIS pegmatites, which generally contain K-feldspar, quartz, albite, and biotite. Future work might investigate whether these dikes were chemically influenced by host metavolcanic rocks.

Figure 1: Geologic map of part of the Tuolumne Intrusive Suite (TIS). Unit boundaries were taken from Bateman et al. (1983). Inset shows the Sierra Nevada batholith in green with the TIS in gray.

PREVIOUS WORK

Tuolumne Intrusive Suite Rb-Sr Systematics

The TIS has long been seen as an archetype of a nested plutonic complex. Bateman and Chappell (1979) proposed that it was constructed of four separate intrusions (from oldest to youngest, Kga-Kkc, Khd, Khdp, and Kcp), each of which shouldered earlier pulses aside to form a nested, inward-younging complex. Frey et al. (1978) and Bateman and Chappell (1979)
proposed that compositional variation was controlled by crystal fractionation. However, these hypotheses are inconsistent with both thermal modeling and isotopic measurements.

Thermal modeling indicated the TIS would cool faster than permitted by geochronological evidence if it existed as amalgamated magma chambers (Glazner et al., 2004). U-Pb zircon geochronology showed that the whole suite took about 10 Ma to amalgamate (Coleman and Glazner, 1997; Coleman et al., 2004). Variation of initial $^{87}$Sr/$^{86}$Sr across the suite cannot be explained by simple fractionation of a few concentric magma chambers (Kistler et al., 1986; Bateman et al., 1988; Gray et al., 2008).

Kistler et al. (1986) found that initial $^{87}$Sr/$^{86}$Sr increases from 0.7057 to 0.7066 from the margin to the interior (Fig. 2). Most variation of initial $^{87}$Sr/$^{86}$Sr occurs in the outer units. Mixing of mantle and crustal derivatives may produce the range of initial $^{87}$Sr/$^{86}$Sr consistent with measured TIS isotopic compositions (Kistler et al., 1986). Monte Carlo simulations (Gray et al., 2008) showed that a combination of partial melting of a mantle source and mixing could produce observed compositional variation.

![Figure 2: Initial Sr for granodiorites from Kistler et al. (1986), Bateman et al. (1988), and Gray et al. (2008). Points lie along A - A' (Fig. 1), and span the suite from the western to eastern margin. The transect starts in Kga on the west and ends in Kkc in the east.](image)
Measured $^{87}\text{Sr}/^{86}\text{Sr}$ ratios of metasedimentary wall rocks from near May Lake age-corrected to 90 Ma ($^{87}\text{Sr}/^{86}\text{Sr}_{90}$) range from 0.710 to 0.774 (Mills et al., 2009). Mills et al. (2009) measured $^{87}\text{Sr}/^{86}\text{Sr}$ and $^{143}\text{Nd}/^{144}\text{Nd}$ in wall rocks and granitoids at May Lake to quantify interaction between the two. They found that measurable interaction was limited to within a meter or two of the contact. Spatially limited hybridization precludes geochemical influence from wall rocks on granitoids throughout the TIS. Metasedimentary rocks near Saddlebag lake have $^{87}\text{Sr}/^{86}\text{Sr}_{90}$ from 0.712 to 0.721 (Gross, 2015).

**SAMPLE DESCRIPTIONS**

Pegmatites, granitoids, and metasedimentary samples were collected at four locations in the TIS. Pegmatite dikes in Kga near May Lake are up to several meters thick and contain mainly K-feldspar, quartz, albite, biotite, and accessory phases including locally abundant tourmaline, Fe-Ti oxides, titanite, apatite, and zircon. K-feldspar is 10 to 20 cm along the longest axis (Fig. 3A). Quartz crystals are locally up to 3 cm in width but mostly occur as graphic intergrowths with K-feldspar and plagioclase. Zonation is not present on the outcrop scale, but some hand samples from May Lake exhibit zonation of quartz coarseness. Biotite books are about 1 x 4 cm. Books do not strike in a common orientation (Fig. 3C). Contacts are locally discolored by oxidation (Fig. 3C).
Three samples were collected from pegmatite dikes on the eastern margin of the TIS near Saddlebag Lake. Two of these pegmatites intrude Kcp and one intrudes a metavolcanic unit. These pegmatites contain K-feldspar (2-10 cm), megacrystic quartz, and interstitial albite. Graphic quartz textures are less common than at May Lake. Perthite lamellae are visible in hand sample as parallel veins. Pegmatite dikes are oblique to foliation in wall rocks (Fig. 3B).

Additional sample areas include Kcp near Twin Lakes, Khde at Olmsted Point, and north of Tenaya Lake (Fig. 1). Pegmatites from near Olmsted Point are narrow dikes about 5 to 10 cm thick and composed of quartz and K-feldspar crystals about 1 cm across. These bodies of pegmatite are thinner than those found along the margins of the TIS. Pegmatite dikes are locally found within aplite dikes. These samples have less complex mineral assemblages than dikes along the TIS margins. Pegmatite collected from the northern portion of the Kcp, near Twin Lakes, contains mostly K-feldspar along with quartz, albite, and biotite.
METHODS

A ceramic shatterbox was used to generate fine powders suitable for dissolution and XRF analysis. Samples larger than ~ 0.6 cm were crushed in a steel jaw crushe. A steel disc mill was used to reduce samples to sand size for heavy-liquid separation. Selected K-feldspar crystals from crushing and hand samples were either mounted or made into thick sections for analysis by scanning electron microscopy.

A hand magnet was used to separate iron filings introduced by the disc mill. Bromoform (specific gravity 2.85) was used to separate dense grains from the more abundant grains of quartz and feldspar. About 30 g of sample was placed in bromoform and stirred to aid in separation. Once settled, dense separates were flushed out of a separatory funnel. Separates were mounted in UV-curing epoxy and were polished with 1 μm α-alumina on a Struers Polisher to prepare for examination with a Tescan Vega TS5136 scanning electron microscope (SEM). Mounted K-feldspar and perthite thick sections were also examined by the SEM for textural observations and for obtaining EDS measurements on microperthite.

About 20 mg of each sample was dissolved for isotopic analysis. Each powder was placed in a Teflon insert with 2 mL of 13 M HNO₃ and 50 μL to 100 μL of 29 M HF, dependent on sample mass. The Teflon inserts were placed in steel jackets. Samples were removed after 48 hours at 180°C, and evaporated on a hot plate at 170°C for 2 hours. The residue was then dissolved in 550 μL of 3.5 M HNO₃.

Samples were then centrifuged and put through Sr-specific ion exchange columns using Eichrom Sr Spec resin (Eichrom Technologies). Each column was pre-rinsed with water and HNO₃. 500μL of sample solution was placed into each column. Once the samples completely drained, several cycles of 3.5 M HNO₃ were rinsed through the elution to remove all cations but
Sr. Distilled water flushed the column to evacuate remaining Sr into a collection beaker. About 30 μL (one drop) of H₃PO₄ was added to the eluted material to minimize evaporation. The solution was then evaporated until a small bead (~1 μL) remained.

Each Sr residue was mixed with 2 μL of 2M HCL and 2 μL of TaCl₅ solution, to aid Sr ionization. 2 μL load solution was loaded onto a rhenium filament. All analyses were conducted on a VG Sector 54 thermal ionization mass spectrometer, measuring \(^{87}\text{Sr}/^{86}\text{Sr}\), \(^{84}\text{Sr}/^{86}\text{Sr}\), and \(^{86}\text{Sr}/^{88}\text{Sr}\) ratios. Tripoli software was used for data reduction. A Rigaku Supermini X-ray fluorescence spectrometer was used to measure concentrations of Rb and Sr. XRF discs were prepared by mixing 10,000 g of finely crushed whole rock with 4,000 g of a mixture of 5% elvacite binder in acetone. After the material was mixed, it was allowed to sit overnight. After re-crushing in a ceramic mortar, the material was pressed into an aluminum cap at 50,000 psi. The XRF was calibrated against various USGS standards for Rb and Sr. Rb and Sr intensities were normalized to the intensity of the Pd Compton peak to correct for the mass absorption coefficient of each sample.

**DATA**

Rb-Sr isotopic ratios were measured for 11 samples from across the TIS, including eight pegmatite, one metasedimentary, and two granitoid samples. Sr isotopic ratios, Rb and Sr concentrations, and unit-wide age ranges from Coleman and Glazner (1997), Coleman et al. (2004), Matzel et al. (2005, 2006), and Memeti et al. (2011) were used to calculate initial \(^{87}\text{Sr}/^{86}\text{Sr}\). These initial ratios were then compared with those of granitoids and wall rocks from Kistler et al. (1986), Bateman et al. (1988), Gray et al. (2008), Mills et al. (2009), and Gross (2015).
Initial $^{87}\text{Sr}/^{86}\text{Sr}$ of Pegmatite

Previous U-Pb zircon geochronological data were used to estimate the pegmatites’ crystallization ages. Measured Sr isotopic ratios, Sr and Rb concentrations, and literature age ranges for corresponding units were applied to the $^{87}\text{Rb}$ decay equation to provide ranges of initial $^{87}\text{Sr}/^{86}\text{Sr}$. Ages for corresponding units and initial $^{87}\text{Sr}/^{86}\text{Sr}$ are all listed in Table 1. Literature ages consisted of the whole range of U-Pb zircon ages for particular units where pegmatites were located (Coleman and Glazner, 1997; Coleman et al., 2004; Matzel et al., 2005, 2006; Memeti, et al., 2011). The entire TIS was assembled from about 95 Ma to 85 Ma (Coleman et al., 2004), so this represents a maximum range for pegmatite crystallization ages. Age correction yielded two ranges of initial $^{87}\text{Sr}/^{86}\text{Sr}$ for each sample: one for a unit-wide age span and a larger uncertainty for the whole-suite age span. Age-related uncertainty is far greater than the analytical uncertainty of $^{87}\text{Sr}/^{86}\text{Sr}$ calculations. The variation in initial $^{87}\text{Sr}/^{86}\text{Sr}$ is a function of the age range and the Rb/Sr ratio (Fig. 4). Initial $^{87}\text{Sr}/^{86}\text{Sr}$ ranges are represented by the distance between y-intercepts of the isochrons shown.

Initial $^{87}\text{Sr}/^{86}\text{Sr}$ is spatially variable across the TIS. The May Lake and Saddlebag Lake areas have pegmatites with higher initial $^{87}\text{Sr}/^{86}\text{Sr}$ than those located in Khdp (Fig. 5). Average initial $^{87}\text{Sr}/^{86}\text{Sr}$ of pegmatite near May Lake is 0.7090 and near Saddlebag Lake is 0.7073. Initial $^{87}\text{Sr}/^{86}\text{Sr}$ for pegmatites on the interior portions of the suite is 0.7066. Ranges of $^{87}\text{Sr}/^{86}\text{Sr}$ for granitoids in corresponding units are shown as green bands in Figure 5. Nearly all of the pegmatites have elevated $^{87}\text{Sr}/^{86}\text{Sr}$ compared to the plutonic rocks of the TIS.
Table 1: Rb-Sr Isotopic Data

<table>
<thead>
<tr>
<th>Names</th>
<th>Type</th>
<th>Rb ppm</th>
<th>Sr ppm</th>
<th>(^{87}\text{Rb}/^{86}\text{Sr})</th>
<th>(^{87}\text{Sr}/^{86}\text{Sr})</th>
<th>(^{87}\text{Sr}/^{86}\text{Sr}_i)</th>
<th>Age est. (Ma)*</th>
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<tr>
<td>MLM1</td>
<td>Metasedimentary</td>
<td>25</td>
<td>155</td>
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<td>0.726575</td>
<td>0.72599</td>
<td>93.3 ± 0.9</td>
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<td>MLP1</td>
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<td>427</td>
<td>51</td>
<td>23.02729</td>
<td>0.739894</td>
<td>0.70930</td>
<td>93.3 ± 0.9</td>
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<tr>
<td>MLP2</td>
<td>Pegmatite</td>
<td>378</td>
<td>72</td>
<td>14.40508</td>
<td>0.727831</td>
<td>0.70869</td>
<td>93.3 ± 0.9</td>
</tr>
<tr>
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<td>0.70637</td>
<td>87.2 ± 1.1</td>
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<tr>
<td>NPGD</td>
<td>Granodiorite</td>
<td>149</td>
<td>539</td>
<td>0.75424</td>
<td>0.707347</td>
<td>0.70640</td>
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<td>OPP1</td>
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<td>223</td>
<td>46</td>
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<td>0.724164</td>
<td>0.70699</td>
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</tr>
<tr>
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<td>0.707167</td>
<td>0.70615</td>
<td>90.5 ± 1.5</td>
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</table>

*Ages are from geochronological data from Coleman et al. (2004), Coleman and Glazner (1997), Matzel et al. (2005, 2006), and Memeti et al. (2011). Margins of uncertainty represent extreme ranges from the sources listed above.*
Figure 4: Intercepts of age correction lines show the initial Sr ratios of OPP1 given the maximum age range of 90 ± 5 Ma (Coleman et al., 2004). The initial $^{87}\text{Sr}/^{86}\text{Sr}$ experiences a relatively small change even at this maximum age range.

Figure 5: Ranges of $^{87}\text{Sr}/^{86}\text{Sr}$ of igneous samples from across the Tuolumne Intrusive Suite. Blue lines show maximum extents of possible $^{87}\text{Sr}/^{86}\text{Sr}$ if the values are corrected to the age range of the entire TIS. Black lines represent $^{87}\text{Sr}/^{86}\text{Sr}$ corrected to the age ranges of units, with data from Coleman et al. (2004). Green bands represent $^{87}\text{Sr}/^{86}\text{Sr}$ for granitoids in the corresponding units.
**Electron Microscopy**

SEM imagery of feldspars form May Lake and Saddlebag Lake was used for both textural observations and semi-quantitative analyses. Perthite lamellae are among the most ubiquitous textural features. Cracks occur locally on-axis with the perthite lamellae. Parallel linear features that are oblique to the general orientation of perthite extend through the feldspar from MLP1. Narrow, tightly packed perthite lamellae truncate asymptotically against these features (Fig. 6E). Perthite lamellae are also present but less continuous throughout SLP2. Perthite lamellae compositions are around An$_{4}$Or$_{1}$Ab$_{95}$, and the K-feldspar is around An$_{0}$Or$_{90}$Ab$_{10}$ (Fig. 7). Not all albite appears to have formed by exsolution; granular, zoned albite in MLP1 contains graphic quartz and more abrupt margins with surrounding K-feldspar (Fig 6B).

Small speckled features are ubiquitous in both feldspar (Fig. 6A) and quartz (Fig. 6C), and in both localities (Fig. 6D). These speckled features appear in and out of focus in Figure 6A despite an even surface. These are likely not polishing artifacts due to their nonlinear shapes. The out-of-focus patches may represent features below the polished surface, such as fluid inclusions. K-feldspar and albite domains appear as patches in samples from both localities. This change in brightness signifies a slight difference in the mean atomic number and may reflect changes in the K/Na ratio or Ba content (Johnson and Glazner, 2010). Major element compositions did not change appreciably when X-ray point analyses were taken across these patches.
Figure 6: A, B, and C are pegmatites from May Lake, and D is a pegmatite from near Saddlebag Lake. E and F show polished sections of MLP1. A) Perthite lamellae (Ab) in a K-feldspar matrix (Or). Bright grains are Fe-Ti oxides (Il). B) Zoned albite containing graphic quartz. C) Quartz containing muscovite (ms) adjacent to albite and K-feldspar. D) K-feldspar with perthite oriented perpendicular to common axes. E) Parallel perthite-free bands in pegmatitic K-feldspar from near May Lake. F) Perthite from May Lake with indistinct margins between K-feldspar and albite domains.
DISCUSSION

Conditions

Major element oxide compositions of perthite domains were compared to assess the extent of exsolution in pegmatitic K-feldspar. MgO, TiO₂, and Fe₂O₃ comprise less than 0.3% of K-feldspars analyzed by XRF, suggesting the crystals were relatively free of inclusions. The whole-feldspar analyses of May Lake and Saddlebag Lake K-feldspars were closer to the orthoclase end, at about Or₇₅. This composition is expected, given proportions of perthite lamellae and K-feldspar observed in BSE images.

Compositions of K-feldspar and perthitic albite, plot on opposite ends of the feldspar ternary diagram (Fig. 7). The bulk compositions were roughly Or₇₅, the albite domains were about Ab₉₅, and the K-feldspar was about Or₉₀ (Fig. 7). These bimodal feldspar compositions indicate that pegmatites continued to develop at temperatures as low as 400°C (Glazner and Johnson, 2013). The implied cooling required for these feldspar compositions is consistent with the late stage nature of pegmatites (Jahns and Burnham, 1969; Glazner and Johnson, 2013). The
peristerite solvus may also have been involved in generating compositional variability in feldspars along the albite end of the ternary diagram (Johnson and Glazner, 2010).

Electron microscopy of K-feldspars, the most abundant phase in sampled pegmatites, supports low-temperature, late stage pegmatite development. The presence of possible fluid inclusions in quartz and K-feldspar (Fig. 6A) may signify high water contents required to crystallize pegmatite (Jahns and Burnham, 1969; Nabelek et al., 2010, Davison and Thomas, 2012). High water contents in pegmatite parental magmas reduce viscosity, enabling circulation through surrounding rock.

**Spatial Variation in $^{87}\text{Sr}/^{86}\text{Sr}_i$**

Initial $^{87}\text{Sr}/^{86}\text{Sr}$ calculated from whole rock Rb-Sr TIMS analyses were compared with surrounding granitoids and nearby metasedimentary wall rocks by projecting initial $^{87}\text{Sr}/^{86}\text{Sr}$ for pegmatites, granitoids (Kistler et al., 1986; Bateman et al., 1988; Gray et al., 2008), and wall rocks (Mills et al., 2009; Gross, 2015) onto transects A – A’ and B – B’ (Figs. 1, 8A, and 8B). Due to the limited spatial extent of wall rocks from near May Lake in Mills et al. (2009), I assumed that they are all located at the same point along the A – A’ transect. Plotting initial $^{87}\text{Sr}/^{86}\text{Sr}$ and $^{87}\text{Sr}/^{86}\text{Sr}_{90}$ versus distance to the nearest contact provides spatial context for isotopic variation (Fig. 9).

Generally, pegmatites near the margins of the TIS have initial $^{87}\text{Sr}/^{86}\text{Sr}$ between granitoids and adjacent wall rocks (figure 8A and 9). These pegmatites are significantly more radiogenic than the granitoids which they intrude (Fig. 5). Of all pegmatite samples along the margins, only one has an uncertainty in $^{87}\text{Sr}/^{86}\text{Sr}_i$ which overlaps the literature granitoid values.
Spatial variation is also apparent when comparing initial $^{87}\text{Sr}/^{86}\text{Sr}$ with distance from the nearest contact. If interaction between wall rocks and granitoids contributed to pegmatite compositions, initial $^{87}\text{Sr}/^{86}\text{Sr}$ of pegmatites along the exterior of the TIS should be greater than those on the interior. Although comparing the isotopic data to contact distance shows similar patterns to transect projections (Figs. 8A, 8B, and 9) this absolute spatial comparison more accurately depicts the distance from pegmatites to the contacts.
Models which invoke simple fractional crystallization (Jahns and Burnham, 1969; Roda-Robles, 2012) cannot solely account for differences in isotopic compositions between pegmatites and plutonic rocks of the TIS. Some external fluid or magmatic source must have contributed to the system prior to pegmatites’ rapid crystallization. The proximity between more radiogenic pegmatites and wall rock, along with the intermediate pegmatite $^{87}\text{Sr}/^{86}\text{Sr}$, suggest that mass transfer between metasedimentary wall rock and surrounding granitoids contributed to the whole-rock isotopic variation.

**Mechanisms**

Hypothetical mixing curves were calculated between several metasedimentary and granitoid samples to determine whether exposed compositions can account for pegmatite compositions. The bulk mixing of two magmas should produce compositions along a hyperbola between endmembers, when comparing initial $^{87}\text{Sr}/^{86}\text{Sr}$ and Sr concentration. Compositions of pegmatites are distinct from most of the calculated mixing curves between the wall rocks and granitoid samples (Fig. 10). Curves which overlap the pegmatite data imply that pegmatites are composed of about 90% wall rock. This ratio of wall rock to granitoid is unlikely, given pegmatites’ low-temperature development (Jahns and Burnham, 1969). If chemical interaction via mixing occurred between intruding magma and wall rocks, at least one of the two endmembers had a composition distinct from units currently exposed.

A more realistic endmember is depicted by the red curve in Figure 10. This curve represents mixing between one of the wall rocks with lower $^{87}\text{Sr}/^{86}\text{Sr}_{90}$ and a fluid derived from a granitic magma. Fluids exsolved from crystallizing magma would likely have lower Sr concentrations but identical initial $^{87}\text{Sr}/^{86}\text{Sr}$ as the granitic magma from which the fluid was
derived. This mixing curve passes through the pegmatite field (Fig. 10) and may represent a possible set of mixing compositions.

The spread of wall rock isotopic data limits estimates of wall rock-related endmembers. A partial melt depleted in Sr may be a more realistic wall rock endmember. However, mixing of partial melts and hydrothermal derivatives may account for pegmatite isotopic compositions in the TIS. The inability for samples to fall on a mixing curve suggests that more complex processes were involved, perhaps combining partial melting of wall rock and fractional crystallization of magmatic pulses. Additional, unobserved endmembers cannot be ruled out. Mass exchange by circulation of fluids through the surrounding wall rock is consistent with the variation of initial $^{87}\text{Sr}/^{86}\text{Sr}$ between pegmatites, wall rocks, and plutonic rocks throughout the TIS.

CONCLUSIONS

Pegmatites’ elevated initial $^{87}\text{Sr}/^{86}\text{Sr}$ and low Sr concentration, compared to plutonic rocks, suggest that pegmatites formed by the interaction of hydrothermal fluids with more
radiogenic wall rock on the margins of the TIS. Hydrothermal fluids circulated through surrounding wall rocks, and acquired a wall rock isotopic signature. This intermediate initial $^{87}\text{Sr}/^{86}\text{Sr}$ was subsequently preserved in pegmatites. Neither fractional crystallization of granitic magma, nor bulk mixing of exposed compositions can exclusively account for the intermediate initial $^{87}\text{Sr}/^{86}\text{Sr}$ of pegmatites along the margins and interior of the TIS. These isotopic data do not preclude the possibility that external hydrothermal fluids entered the system and contributed to pegmatite isotopic compositions.

Obtaining isotopic data from pegmatite dikes in Kjp and from additional locations along TIS-wall rock margins would further test the hypothesis that pegmatites record fluid-wall rock interaction. If pegmatites from the interior of the TIS have initial $^{87}\text{Sr}/^{86}\text{Sr}$ within the range of Kjp initial $^{87}\text{Sr}/^{86}\text{Sr}$, this would support the aforementioned hypothesis. Elevated initial $^{87}\text{Sr}/^{86}\text{Sr}$ for pegmatites on the TIS interior would suggest that another mechanism or crustal fluid contributed to isotopic variation.

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