Relationships between magmatism and deformation in southeastern Proterozoic Australia

Thesis submitted in accordance with the requirements of the University of Adelaide for an Honours Degree in Geology/Geophysics

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CONSTRAINTS ON MESOPROTEROZOIC MAGMATISM AND DEFORMATION IN THE SOUTHERN GAWLER CRATON, SOUTH AUSTRALIA

MAGMATISM AND DEFORMATION IN YORKE PENINSULA

ABSTRACT

The ca. 1600–1580 Ma time slice is recognised as a significant period of magmatism and deformation throughout eastern Proterozoic Australia. Within the northern Yorke Peninsula, this period was associated with the emplacement of multiple phases of the Tickera Granite; an intensely foliated orange granite, a white leucogranite and a red granite. These granites belong to the broader Hiltaba Suite that was emplaced at shallow crustal levels, throughout the Gawler Craton. Geochemical and isotopic analysis suggests these granite phases were derived from a heterogeneous source region. The orange and red granites were derived from the Donington Suite and/or the Wallaroo Group metasediments with slight contamination from an Archean basement. The white leucogranite is sourced from a similar but slightly more mafic/lower crustal source. Phases of the Tickera Granite were emplaced synchronously with deformation that resulted in development of a prominent northeast trending structural grain throughout the Yorke Peninsula region. This fabric is a composite of two fold generations; early isoclinal folds that were refolded by later open upright folds. Isoclinal folding may have occurred during the ca. 1730–1690 Ma Kimban Orogeny, or just prior to emplacement of the Tickera Granite at ca. 1597–1577 Ma. The upright fold generation was contemporaneous with the emplacement of the Tickera Granite. The Yorke Peninsula shares a common geological history with the Curnamona Province, which was deformed during the ca. 1600-1585 Ma Olarian Orogeny, and resulted in development of early isoclinal (recumbent) folds overprinted by an upright fold generation, a dominant northeast-trending structural grain and spatially and temporally related intrusions. This suggests that an apparent correlation with the geological history of the Curnamona Province, and that the Olarian Orogeny may have also affected the southeastern Gawler Craton. Constraint on the timing of the earlier isoclinal fold generation in the Yorke Peninsula will allow further understanding of the similarities between the two regions.

KEYWORDS

Mesoproterozoic; Magmatism; Deformation; Yorke Peninsula; Tickera Granite; Hiltaba Suite; Olarian Orogeny; Gawler Craton; Curnamona Province

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INTRODUCTION

The ca. 1600–1580 Ma time slice is a recognised period of high grade metamorphism associated with intense crustal orogenesis within eastern Proterozoic Australia (e.g. Collins & Shaw 1995; Betts et al., 2002; Forbes et al., 2008; Stewart & Betts 2010; Forbes et al., 2012), which includes the Gawler Craton and Curnamona Province of South Australia (Fig. 1a,b). Within the Gawler Craton, this period was also associated with bimodal volcanism that resulted in outpouring of the ca. 1592 Ma Gawler Range Volcanics (Fanning et al., 1998) and emplacement of the ca. 1595–1575 Ma Hiltaba Suite Granites (e.g. Flint et al., 1993; Conor 1995; Creaser & Cooper 1993; Fanning et al., 2007) to shallow crustal levels (e.g. Daly et al., 1998; Stewart & Foden 2003). The Hiltaba Suite granites are widely distributed across the Gawler Craton and are suggested to comprise two phases emplaced at ca. 1590 Ma and ca. 1580 Ma (Stewart & Foden 2003; Hand et al., 2007). The granites vary across the A-, I- and S-type range, and have characteristic chemical signatures (such as high SiO₂: CaO, MnO, Sr ratios and Sm-Nd isotopes) depending on whether they are part of the older or younger phase of granites (Stewart & Foden 2003; Hand et al., 2007). Granites in the northern Yorke Peninsula in the southern Gawler Craton (Fig. 1c) are suggested to be part of the Hiltaba Suite, and include the deformed Tickera Granite (1598 \pm 7 Ma, 1575 \pm 7 Ma: Conor 1995) and undeformed Arthurton Granite (1583 \pm 7 Ma: Creaser & Cooper 1993). These granites intrude metasediments and metavolcanics of the ca. 1750 Ma Wallaroo Group (Cowley et al., 2003). Magmatic crystallization ages derived from the Tickera Granite range from ca. 1597–1577 Ma (Conor 1995), suggesting variable conditions and timing of emplacement. The timing of emplacement of the Tickera Granite is also synchronous with deformation, metamorphism and

metasomatism that is suggested to have affected the Wallaroo Group at ca. 1600–1575 Ma (Conor et al., 2010). However, details on emplacement relationships and associated events that affected the Yorke Peninsula region is lacking.

This study aims to constrain the relative timing of emplacement of multiple phases of the deformed Tickera Granite exposed at Point Riley in the northwestern Yorke Peninsula and place this in context of the southeastern Proterozoic Australia (Fig. 1c). This has been done through detailed mapping to assess relative intrusive relationships, structural fabrics and deformation styles within the granite phases and comparison with hosting Wallaroo Group metasediments. Whole rock geochemical and Sm–Nd isotopic analysis of the granite phases is used to assess the source region of the multiple phases identified, and place the Tickera Granite within the broader context of the ca. 1595–1575 Ma Hiltaba Suite (Creaser & Cooper 1993; Fanning et al., 2007). The magmatic and deformational history of the Yorke Peninsula is then discussed in relation to the ca. 1600–1585 Ma Olarian Orogeny (Page et al., 2005) that affected eastern Proterozoic Australia, particularly within the Gawler Craton and Curnamona Province.

REGIONAL GEOLOGY OF THE GAWLER CRATON

The Gawler Craton preserves a complex geological history spanning from the late Archean to the early Mesoproterozoic. Mesoarchean granitoids of the Cooyerdoo Granite were emplaced between ca. 3200–3150 Ma, and are recognised as the oldest known units within the Gawler Craton (Fraser et al., 2010a; Jagodzinski et al., 2011b; Reid & Hand 2012). Felsic magmatism associated with emplacement of the Coolanie Gneiss occurred at ca. 2823 Ma (Fraser and Neumann, 2010), and was followed by bimodal magmatism and sedimentation during the Neoarchean to early

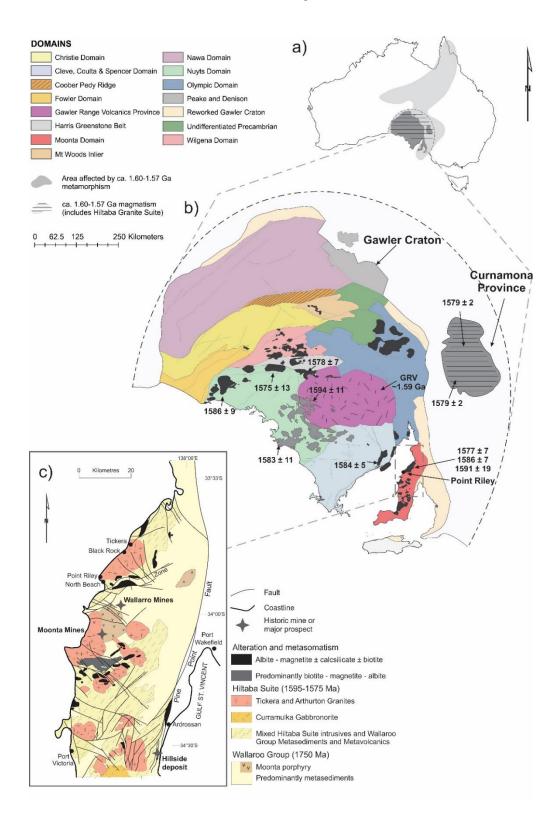


Figure 1. (a) Map of Australia showing the area of Proterozoic Australia affected by tectonothermal activity and magmatism during the ca. 1600–1580 Ma time slice; (b) Simplified geological map of the Gawler Craton showing the distribution of Hiltaba Suite plutons, the Gawler Range Volcanics and geological domains after Stewart & Foden (2003); (c) Solid geology map of the Yorke Peninsula displaying the distributions of the Arthurton and Tickera Granites, modified after Conor et al., (2010). Mapping and sample locations of Point Riley, and sample locations of Black Rock and Wallaroo North Beach are shown.

Paleoproterozoic, forming the Sleaford Complex and Mulgathing Complex in the in the southern and central western Gawler Craton respectively. The Gawler Craton was deformed and metamorphosed during the ca. 2480–2420 Ma Sleafordian Orogeny (Daly & Fanning 1993; Daly et al., 1998; Reid & Hand 2012). Recurrence of magmatism in the southeastern Gawler Craton at 2000 Ma lead to the development of the Miltalie Gneiss (Daly et al., 1998; Fanning et al., 1998, 2007).

Onset of basin formation resulted in deposition of the Hutchison Group at ca. 2000–1730 Ma (Parker et al. 1988; Hand et al., 2007). Basin development was interrupted at ca. 1850 Ma with emplacement of the Donington Suite along the eastern margin of the Gawler Craton during compressional deformation associated with the Cornian Orogeny (Reid et al., 2008). Subsequent emplacement of bimodal packages included the Broadview Schist and Myola Volcanics (ca. 1791 Ma: Parker et al., 1993; Fanning et al., 1988), and McGregor Volcanics (1740 Ma: Fanning et al., 1988). Sedimentary packages were also emplaced during this time interval and include the Price Metasediments (ca. 1767 Ma: Oliver and Fanning et al., 1997), Moonabie Formation (ca. 1756 Ma: Jagodzinski., 2005), the Wallaroo Group (1760–1740 Ma: Cowley et al., 2003), and unnamed sediments in the Nawa and Fowler Domains (Parker et al., 1993; Daly et al., 1998; Hand et al., 2007).

The Gawler Craton underwent a second tectonothermal event during the ca. 1730–1690 Ma Kimban Orogeny (Hoek & Schaefer 1998; Ferris et al. 2002; Hand et al. 2007). Intrusive rocks associated with the Kimban Orogeny include the ca. 1745–1700 Ma Peter Pan Supersuite (e.g. Wade & McAvaney 2016). Sedimentary packages include the ca. 1715 Ma Eba and Labrinyth Formations (Cowley and Martin 1991; Fanning et al., 2007). The Kimban Orogeny was followed by intrusion of the 1660–1640 Ma Tunkillia

Suite (Teasdale 1997; Daly et al. 1998; Ferris et al. 2002), and later eruption of the 1630 Ma Nuyts Volcanics (Rankin et al., 1990).

The St. Peter Suite intruded at ca. 1640–1608 Ma (Teasedale 1997; Daly et al. 1998; Swain et al., 2008; Symington et al., 2014). Bimodal volcanism that resulted in the emplacement of the Gawler Range Volcanics and Hiltaba Suite occurred between 1595–1575 Ma and was synchronous with regional deformation (e.g. Flint et al., 1993; Conor 1995; Creaser & Cooper 1993; Fanning et al., 2007). The Hiltaba Suite is suggested to have been generated from mafic underplating associated with lithospheric attenuation, and melting of the overlying continental crust (Flint et al., 1993; Zang et al., 2007). Geochemical variations within the Hiltaba Suite granites across the Gawler Craton is suggested to be due to their derivation from a depleted mantle source that mixed with various crustal components (Stewart & Foden 2003). Further deformation across the Gawler Craton occurred during the ca. 1570–1540 Ma Kararan Orogeny and ca. 1470–1450 Coorabie Orogeny (Hand et al., 2007).

REGIONAL GEOLOGY OF THE YORKE PENINSULA

The Yorke Peninsula is part of the Moonta Domain in the southeastern Gawler Craton, South Australia (Fig. 1b,c). The area comprises late Paleoproterozoic metasediments and metavolcanics of the ca. 1750 Ma Wallaroo Group (Cowley et al., 2003), which extends northward from Yorke Peninsula, below the Stuart Shelf and along much of the Olympic Domain (Fig. 1b; Conor et al., 2010). Fine-grained psammites, siltstones, calcsilicates, albitites, quartzites and iron rich sediments of the Wandearah Formation dominate the Wallaroo Group (Conor 1995; Cowley et al., 2003). These metasediments were contemporaneously deposited with bimodal A-type volcanics of the Weetula Formation and Matta Formation (Cowley et al., 2003).

The Wallaroo Group preserves evidence of metamorphism and multiple folding events (e.g. Conor et al., 2010). Metamorphic grade varies from upper greenschist in the northern Yorke Peninsula to amphibolite facies in south (Conor et al., 2010). Two phases of deformation are preserved in the Wallaroo Group metasediments. The first phase involved generation of isoclinal folds with limbs extending up to several kilometres in length. The timing of this deformation event is suggested to be related to the ca. 1730–1690 Ma Kimban Orogeny, however this is equivocal (e.g. Conor et al. 2010). The second deformation phase involved refolding of the isoclinal folds by open upright folds (Conor 1995; Conor et al., 2010). Late folding is suggested to be associated with deformation during ca. 1600–1570 Ma time period based on preservation of amphibolite-rich calcsilicate alteration within the axial plane of the open upright folds, and that is interpreted to be related to alteration associated with intrusion of the Hiltaba Suite granites (Conor et al., 2010).

Within the Yorke Peninsula, the mafic component of the Hiltaba Suite is recognised as the Curramulka Gabbronorite, which facilitated melting of the lower crust to form its

Within the Yorke Peninsula, the mafic component of the Hiltaba Suite is recognised as the Curramulka Gabbronorite, which facilitated melting of the lower crust to form its felsic component, the Arthurton and Tickera granites (Zang et al., 2007). The Tickera Granite, located in the northern Yorke Peninsula (Fig. 1c) preserves high strain deformation, which has been used to suggest that the granite was intruded into a tectonic regime in which shearing was prominent (Wurst 1994; Conor 1995). Conversely, the Arthurton Granite to the south is undeformed and suggested to postdate local deformation (Conor 1995; Conor et al., 2010).

METHODS

Point Riley on the western Yorke Peninsula preserves a well–exposed shore platform of multiple phases of the Tickera Granite. The platform was mapped in detail to show the distribution and intrusive relationships of granite phases and preserved structural fabrics (Fig. 2).

Twelve samples were collected from Point Riley and are representative of the four granite phases that were identified during mapping. These samples were spatial distributed across the mapping area, were *in situ* and as fresh as possible. Two additional samples representing the southern and northern extent of the Tickera Granite were collected; 2.87 km south-east of Point Riley at Wallaroo North Beach (sample 2149822) and 7.87 km northeast of Point Riley at Black Rock (sample 2149823). Sample locations are shown in Figure 2. The samples were used for a combination of petrography, whole rock geochemistry and Sm-Nd isotopic analysis (Table 1). Twelve samples were used for whole rock geochemical analysis (Table 1). All samples were prepared using a rock saw at the University of Adelaide. Crushing and whole rock geochemical analysis was undertaken at Australian Laboratory Services Pty. Ltd. using standard preparation methods (e.g. crushing and pulverising). Major and trace elements were analysed by inductively coupled plasma mass spectrometry (ICP–MS). Eight samples were analysed for Sm-Nd isotope geochemistry: six from Point Riley, one from North Beach and one from Black Rock (Table 1). Samples from Point Riley were selected to represent each of the granite phases observed and based on their spatial distribution and trace element and rare earth element geochemistry, particularly their Sm and Nd content. Samples collected from Wallaroo North Beach and Black Rock were analysed to provide a broader understanding of the Sm–Nd isotopic composition

of the Tickera Granite across the northern Yorke Peninsula. Sm–Nd analysis was undertaken following the method of Wade et al. (2006) using a Finnigan MAT262 and MAT261 thermal ionisation mass spectrometer. ¹⁴⁶Nd/¹⁴⁴Nd ratio was normalised to 0.721903, Nd blanks corrected for 200 pg and Sm for 150 pg.

Eight samples were selected for petrographic analysis (Table 1). Samples included the six samples used for Sm–Nd analysis. Samples were prepared using a rock saw at the University of Adelaide and prepared by Thin Section Australia. Petrographic analysis was undertaken using standard transmitted and reflected light microscopy to assess mineralogy and textural relationships.

Table 1. Analyses performed on samples collected from Point Riley, Black Rock and Wallaroo North Beach in this study.

Sample	Rock Type	Geochemistry	Petrography	Sm-Nd
2147273	Nodule	✓	✓	✓
2147279	Nodule	✓		
2147276	Orange	✓	✓	\checkmark
2147280	Orange	✓		
2147281	Orange	✓	✓	\checkmark
2147272	White	✓	✓	\checkmark
2147275	White	✓		
2147277	White	✓	✓	\checkmark
2147271	Red	✓		
2147274	Red	✓		
2147278	Red	√		
2147282	Red	√	✓	✓
2149823	Black Rock (Red)		\checkmark	✓
2149822	North Beach (Red)		✓	√

OBSERVATIONS AND RESULTS

Field Observations

The Point Riley area (Figs. 1b, c and 2) preserves well exposed shore platforms of the Tickera Granite. Four phases of granite were recognised and are divided based on grain size, mineralogy, composition and deformation style. The four phases were granite nodules, an intensely foliated orange granite, a white leucogranite and a red granite.

Granite nodules

Granite nodules are preserved throughout the intensely foliated orange granite (Figs. 2 and 3a). The nodules extrude from the surrounding foliated granite, are up to 30 cm length and oval in shape. The nodules comprise medium grained K–feldspar (60%; 1–5 mm), quartz (30%; 1–4 mm) and amphibole (10%; 2 mm) (Fig. 3a). Mineral grains vary from rounded (K–feldspar and quartz) to sub–rounded (amphibole) and elongated within a well–developed, northeast–trending foliation that dips steeply to the northwest. This foliation is also preserved in the surrounding foliated orange granite.

<u>Intensely foliated orange granite</u>

The dominant phase in the mapping area is an intensely foliated orange granite that is recessively weathered (Figs. 2 and 3a–c). The foliated orange granite shares similar mineralogy to the granite nodules. Mineral grains are sub–rounded to rounded, vary in size from medium– to coarse–grained and comprise K–feldspar (70%; 1–8 mm), quartz (~15%; 1–4 mm), biotite and amphibole (10% mafics; 1–2 mm), and small quantities of plagioclase (<5%; 1 mm). Hematite alteration is evident in this phase through its red/orange colour.

The orange granite preserves a well–developed, pervasive foliation (Figs. 2, 3a–c and 4a) defined by K–feldspar and quartz crystals. The foliation is northeast–trending and steeply northwest or southeast dipping. A locally well–developed cleavage is also preserved in the orange foliated granite, particularly in areas close to the contact boundaries with the red and white granite phases within which the cleavage is a dominant feature. The cleavage is discontinuous, northwest–trending and moderately to steeply northeast dipping.

Enclaves of undeformed orange granite are locally preserved within the intensely foliated orange granite in the northwestern part of the study area (Fig. 2). Metasedimentary xenoliths of the Wallaroo Group are also preserved within the same local area and range in size from 20cm–1m and comprise fine–grained metapelites (Fig. 2). Quartz and K–feldspar veins which are 5–30 cm thick cross–cut this phase throughout the study area.

White leucogranite

A white leucogranite forms prominent, blocky exposures throughout the study area (Figs. 2 and 3d–f). The leucogranite is medium– to coarse–grained and comprises plagioclase (60%; 1–3 mm), quartz (~35%; 1–2 mm) and minor mafics (<5%; 1 mm). All grains are sub–rounded.

Two distinct cleavages are evident in the white leucogranite (Figs. 2, 3d, f and 4b). The first is a locally well–developed, 5–30 cm spaced, northeast–trending and moderately to steeply northwest dipping cleavage. The second cleavage is a moderate to well–developed northwest–trending cleavage and dips steeply to the northeast. This cleavage is 10–100 cm spaced and is evident throughout the whole mapping area. A poorly

developed mineral lineation is locally preserved and is defined by the alignment of quartz and feldspar crystals.

Red granite

A deformed red granite intrudes through all phases as a sheeted dyke and outcrops at both high and low reliefs across the study area (Figs. 2 and 3b, e–g). The red granite comprises coarse, sub–rounded grains of K–feldspar (55%; 1–5 mm), quartz (35%; 1–3 mm) and minor amounts of amphibole (10%; 1–2 mm). Within the high relief outcrops of the red granite, cavities ranging in size from 3–5 mm are preserved (Figs. 2 and 3g). Hematite alteration is also evident by the red colouration of the granite.

A well–developed foliation is locally preserved along the margins of the red granite dyke within 50cm of the contact boundary with the intensely foliated orange granite (Figs. 2, 3b and 4c). The foliation is defined by elongate quartz and K–feldspar crystals, is northeast–trending and steeply dips to the northwest. A well–developed, 5–30 cm spaced, northwest–trending, moderately to steeply northeast or southwest dipping cleavage is locally preserved (Figs. 2 and 4c). This cleavage is continuous into the other granite phases.

Contact relationships

The contact boundaries between the foliated orange granite, white leucogranite and red granite are sharp in most cases (Figs. 3b, c, f). The white leucogranite locally intrudes the foliated orange granite as metre–scale fingers along the foliation plane of the enclosing orange granite phase. The fingers are 20–30 cm wide and up to 3m in length (Fig. 2). In the centre of the map area, small 5–10cm long apophyses of the white leucogranite observed intruding into the foliated orange granite (Figs. 2 and 3c).

Evidence of contact metamorphism between the red granite and white leucogranite is locally preserved (Figs. 2 and 3e). The contact metamorphic boundary ranges in thickness from 5–20 cm, and is manifest by a colour difference between the two phases, where the white leucogranite gradually becomes redder in colour. Approximately 2 m east of the contact metamorphic boundary, a raft of white leucogranite is preserved within the red granite dyke, which itself is surrounded by the white leucogranite (Figs. 2 and 3f). The white leucogranite raft is rounded, approximately 1x2 m size. In the northeastern section of the study area, evidence of magma mingling between the red granite dyke and white leucogranite is evident (Figs. 2 and 3h); contact boundaries are highly irregular and often unclear. Mixing appears to be incomplete. Iron staining on the surface of the white leucogranite is also a prominent feature in this part of the study area.

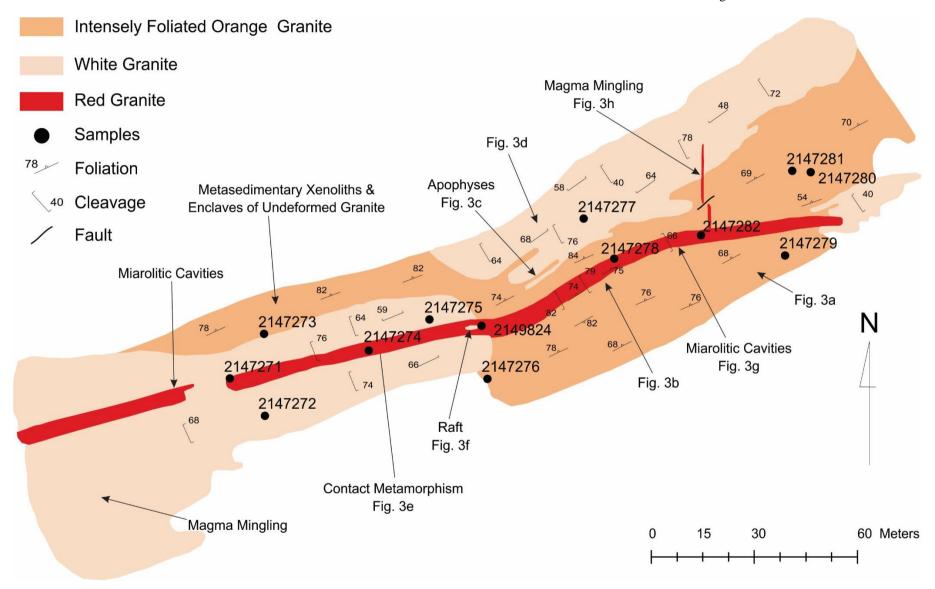


Figure 2. Geological surface map of Point Riley showing the mapped phases of the Tickera Granite, sample locations, structures observed and intrusive magmatic features. Map location shown in Figure 1c.

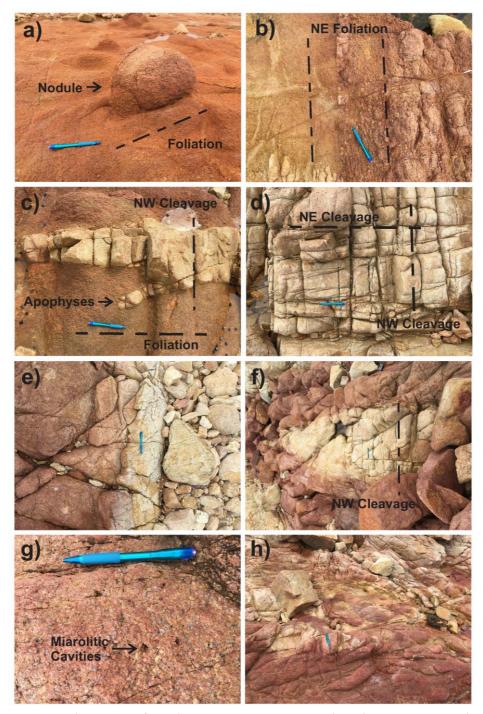


Figure 3. Representative photos of granite phases and contact relationships observed at Point Riley. (a) Granitic nodule within the intensely foliated orange granite. The recessive nature and the intense NE-trending foliation can be seen in the orange granite. Photo taken looking north; (b) Contact boundary between foliated orange granite (left) and red granite (right) showing the well-developed NE-trending foliation preserved within both granite phases. Photo taken looking southwest; (c) White leucogranite apophyses intruding into the surrounding foliated orange granite; (d) Two cleavages in the white leucogranite. Photo taken looking northwest; (e) Contact metamorphic boundary between the white and red granites. Northeast to the top of photo; (f) Raft of white leucogranite within the red granite dyke. Photo taken looking northwest; (g) Miarolitic cavities in the red granite. Photo taken looking southeast; (h) Magma mingling between the white and red granite phases. Photo taken looking east. Location of photos shown in Figure 2.

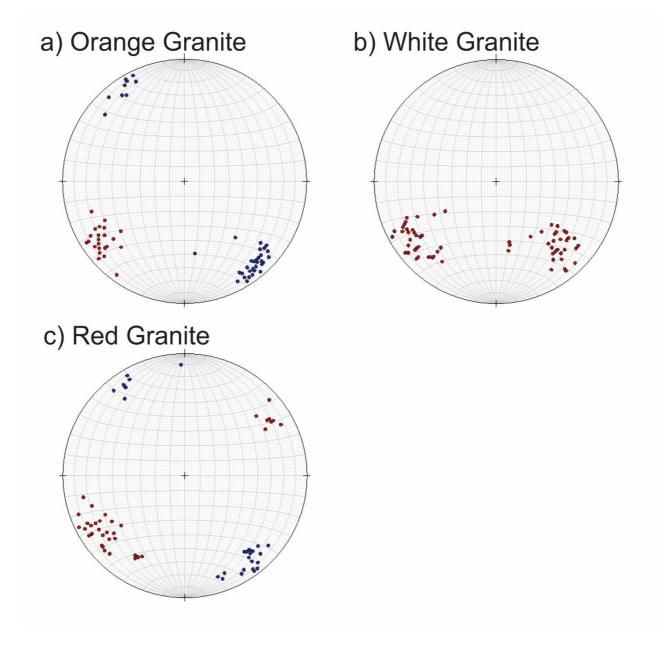


Figure 4. Stereonet plots showing foliation (blue) and cleavage (red) data plotted as poles to planes for the (a) orange, (b) white and (c) red granite phases.

Petrography

Granite nodules

The granite nodules primarily comprises microcrystalline K-feldspar, quartz, magnetite, biotite and plagioclase. Few K-Feldspars are microcline, and the majority show evidence of significant sericite alteration and hematite staining. Biotite is subhedral while magnetite is euhedral, and are both fine grained and randomly distributed throughout the granite. Hematite-filled veins also cross cut all minerals (Fig. 5a).

<u>Intensely foliated orange granite</u>

The orange granite comprises K–feldspar, quartz, biotite, magnetite, plagioclase and minor amphibole. Few K–Feldspars are microcline, and the majority show evidence of significant sericite alteration and hematite staining. Vermicelli type exsolution is commonly observed within feldspars. Biotite is euhedral and along with K-feldspar is aligned within the dominant foliation preserved in the granite. Localised brecciation shows a stockwork of hematite veining that is predominantly developed around altered feldspars and within quartz grains (Fig. 5b).

White leucogranite

The white leucogranite dominantly comprises plagioclase and quartz, with minor magnetite. Plagioclase is significantly sericite altered. Thin hematite veins occasionally cross cut quartz and plagioclase (Fig. 5c).

Red granite

The red granite generally comprises medium to coarse grained K-feldspar and quartz, with minor plagioclase, magnetite and biotite. Minor microcline is observed. Sericite and hematite alteration of feldspars is preserved (Fig. 5d). Minor perthitic exolutions are also observed. At North Beach, a prominent foliation defined by quartz and

significantly sericite altered and brecciated feldspars, is locally preserved within the red granite (Fig. 5e). At Black Rock, the red granite also contains amphibole, with higher modal proportion of biotite and more intense sericite alteration (Fig. 5f). Minor hematite is commonly preserved and is present in thin veins.

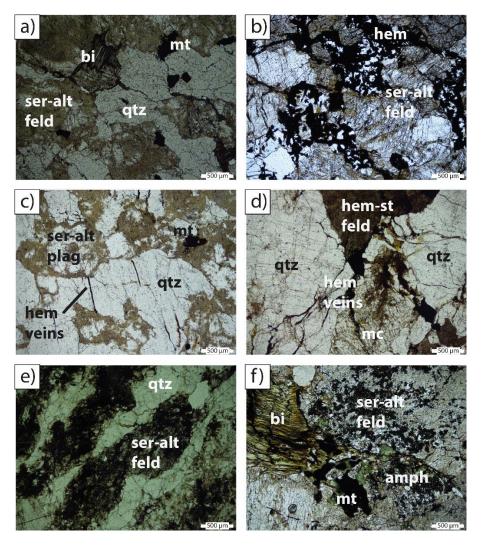


Figure 5. Representative plane polarised photomicrographs of Tickera Granite phases at Point Riley. (a) Granitic nodule illustrating sericitised feldspar, with biotite and magnetite localised around the edges; (b) Orange granite displaying a stockwork of hematite filled veins and altered feldspar. Sample 2147281; (c) White granite displaying sericitised plagioclase, quartz, minor magnetite and thin hematite filled veins. Sample 2147277; (d) Red granite displaying hematite stained feldspar, quartz, hematite veins and microcline; (e) North Beach red granite (2149822) displaying foliation defined by brecciated, altered feldspars and quartz; (f) Black Rock red granite (2149823) displaying abundance of biotite and amphibole, sericitised feldspars and minor magnetite. Mineral abbreviations: ser-alt feld: sericite-altered feldspar; bi: biotite; mt: magnetite; hem: hematite; ser-alt plag: sericite-altered plagioclase; qtz: quartz; mc: microcline; amph: amphibole

Geochemistry

Major Elements

Major element geochemical results for the nodules, orange, white and red granites are displayed in Figure 6 and Table 2. The SiO₂ content of the Tickera Granite ranges from 69.9–79.2 wt%. The orange foliated granite has SiO₂ content of 69.9–71.1 wt%, the nodules range from 71.1–73.4 wt%, whilst the white leucogranite clusters around 77 wt%. The red granite shows the widest variation in SiO₂ content of 70–79 wt% (Fig. 6). Al₂O₃, P₂O₅, MgO and TiO₂ display a negative trend with increasing SiO₂ content for all samples (Fig. 6a–d). Fe₂O₃ content has a steep negative correlation for SiO₂ content between 69.9–~72 wt%, which then becomes shallower with higher SiO₂ content (Fig. 6e). A negative trend is observed for CaO and Na₂O with increasing SiO₂ in the foliated orange granite, nodules and red granite phases (Fig. 6f-g). The white leucogranite however, has a relatively high CaO and Na₂O content compared to other phases (Fig. 6f–g). K₂O is scattered; the white leucogranite phase shows relatively low K₂O contents (Fig. 6h). MnO content for most samples are below detection limit except for two of the lower silica phase samples (2147271 and 2147280; Table 2, Fig. 6i). The white leucogranite is rich in CaO and Na₂O, and depleted in K₂O, reflecting the high albite content (Fig. 6j-k). The red granite is rich in K₂O and depleted in CaO and Na₂O, reflecting the high K-feldspar content (Fig. 6j-k). The orange granite and nodules are intermediate between the two and each have an outlier (Fig. 6j-k).

Table 2. Major and trace element geochemical data for samples from Point Riley. Sample location shown in Figure 2.

	Sample	Unit	2147276	2147280	2147281	2147273	2147279	2147272	2147275	2147277	2147271	2147274	2147278	2147282
Side	Description					Nodule	Nodule							
	SiO ₂	%				73.40	71.10							
	TiO_2	%	0.47	0.48	0.11	0.23	0.43	0.17	0.23	0.23	0.54	0.18	0.23	0.14
Mado % 0.01 0.02 0.01 0.01 0.01 0.01 0.02 0.01 0.02 0.01 0.01 0.01 0.02 0.01 0.07 0.03 0.03 0.22 0.07 0.03 0.04 0.04 0.04 0.04 0.04 0.05 0.04 0.05 0.02 0.01 0.05 0.03 0.02 0.01 0.03 0.02 0.03 0.03 0.03 0.03 0.04 0.03 0.02 0.03 0.02 0.03 0.03 0.03 0.03 0.03 0.04 0.04 0.03 0.03 0.03 0	Al_2O_3	%	14.00	14.35	14.95	13.60	14.20	13.50	14.00	14.00	11.95	12.90	11.70	12.50
MpO 6s 0.5s 0.5s 0.5l 0.13 0.12 0.15 0.05 0.05 0.16 0.05 0.10s 0.2cl 0.07 0.08 0.21 0.07 0.03 0.25 0.20 1.04 1.04 0.05 0.03 0.05 0.01 0.08 0.09 0.02 0.01 0.01 0.08 0.02 0.02 0.02 0.00 <t< th=""><th>Fe_2O_3</th><th>%</th><th>4.19</th><th>4.69</th><th>2.40</th><th>2.67</th><th>3.73</th><th>0.96</th><th>0.93</th><th>1.06</th><th>6.42</th><th>1.87</th><th>1.96</th><th>2.06</th></t<>	Fe_2O_3	%	4.19	4.69	2.40	2.67	3.73	0.96	0.93	1.06	6.42	1.87	1.96	2.06
Caco 6s 6.57 0.85 0.85 0.98 0.45 0.42 1.02 1.02 1.02 5.57 5.57 3.17 2.94 2.93 2.71 KSO % 4.26 3.30 6.83 4.30 1.33 1.22 1.21 1.41 5.31 3.86 4.87 6.06 6.06 0.09 0.02 0.01 0.01 0.08 0.03 0.05 0.04 0.04 0.05 0.03 0.04 0.04 0.05 0.05 0.04 0.04 0.05 0.05 0.05 0.04 0.04 0.05 0.05 0.03 0.04 0.04 0.05 0.05 0.04 0.04 0.05 0.03 0.04 0.04 0.04 0.05 0.03 0.04 0.04 0.05 0.05 0.05 0.03 0.04 0.04 0.05 0.04 0.00 0.04 0.05 0.03 0.03 0.03 0.00 0.00 0.03 0.03 0.03 0.03 </th <th>MnO</th> <th>%</th> <th>0.01</th> <th>0.02</th> <th>0.01</th> <th>0.01</th> <th>0.01</th> <th>0.01</th> <th>0.01</th> <th>0.01</th> <th>0.02</th> <th>0.01</th> <th>0.01</th> <th>0.01</th>	MnO	%	0.01	0.02	0.01	0.01	0.01	0.01	0.01	0.01	0.02	0.01	0.01	0.01
NacO	MgO	%	0.58	0.51	0.34	0.17	0.16	0.05	0.09	0.06	0.21	0.07	0.09	0.13
No. No.	CaO	%	0.57	0.85	0.08	0.55	0.93	1.62	1.08	1.04	0.36	0.28	0.33	0.26
Prob	Na ₂ O	%	4.03	4.11	3.97	4.04	5.52	5.20	5.57	5.57	3.17	2.94	2.93	2.71
No. No.	K_2O	%	4.26	3.80	6.83	4.30	1.31	1.22	1.24	1.43	5.01	5.86	4.87	6.06
Total % 99.21 100.91 107.66 99.64 98.46 99.98 100.65 100.65 98.81 98.66 101.75 103.00 F ppm 450.00 135.00 352.00 1835.00 222.00 105.00 172.00 173.00 355.00 353.00 352.00 450.00 350.00 353.00 352.00 350.00 353.00 352.00 350.00 353.00 352.00 350.00 350.00 350.00 350.00 350.00 350.00 350.00 350.00 350.00 350.00 350.00 350.00 350.00 350.00 350.0	P_2O_5	%	0.03	0.05	0.01	0.06	0.09	0.02	0.01	0.01	0.08	0.02	0.02	0.01
F ypm 590.00	LOI	%	1.17	1.25	0.96	0.61	0.98	0.43	0.59	0.54	0.65	0.53	0.41	0.46
Rb γpm 42000 32900 38300 22200 105.00 65.0 67.20 72.10 270.00 355.00 333.00 374.00 Sr ppm 1525.0 1815.0 3570 251.00 400.00 144.00 113.00 30.00 345.00 83.00 83.00 83.00 83.00 68.00 6.08 1.22 0.08 1.22 0.08 0.08 0.08 0.08 0.08 0.08 0.00 0.08 0.00 0.00 0.00 0.00 0.00 0.00 0.00 0.00 0.00 0.00 0.00 0.00 3.00 1.01 3.00 1.01 1.01 1.00 9.00 1.00 1.00 9.00 0.00 2.00 9.00 0.00 2.00 2.00 3.00 2.00 2.00 3.00 3.00 2.00 9.00 9.00 2.00 2.00 9.00 9.00 2.00 2.00 9.00 9.00 1.00 1.00 9.00 1.00	Total	%	99.21	100.91	100.76	99.64	98.46	99.98	100.65	100.65	98.81	98.66	101.75	100.34
Sr ppm 152,00 181,50 35.70 251,00 400,00 144,00 119,00 113,00 306,00 84,50 83,80 83,00 Cs ppm 658,00 722,00 177,00 1744 744,0 124,00 114,00 597,00 527,00 699,00 Th ppm 65,30 65,30 27,20 122,00 49,80 90,90 115,50 65,80 37,60 121,50 104,50 98,60 U ppm 631 8.49 1.81 1345 5.50 3.74 2.52 2.77 6.59 8.66 6.66 3.65 Pb ppm 6.68 2.50 3.30 141,00 227,00 3.80 3.50 2.20 3.80 3.50 3.20 3.40 4.10 6.53 3.30 Tr ppm 160.80 2.88 2.49 5.50 2.77 2.50 3.80 3.30 3.40 4.10 4.00 4.00 4.00	F	ppm	590.00				470.00		130.00				110.00	
Cs ppm 2.55 2.26 0.81 1.38 1.73 0.56 1.02 1.12 0.78 0.94 0.96 0.84 Ba ppm 668.00 683.00 788.00 712.00 177.00 174.00 75.40 122.00 114.00 597.00 507.00 68.00 68.00 68.00 68.00 68.00 68.00 68.00 68.00 68.00 2.70 122.00 104.00 98.00 90.00 15.00 68.00 6.00 5.00 3.00 17.00 38.00 12.00 16.60 5.00 2.00 2.00 9.00 18.00 2.20 13.40 4.00 </th <th>Rb</th> <th>ppm</th> <th>420.00</th> <th>329.00</th> <th>383.00</th> <th>222.00</th> <th>105.00</th> <th>60.50</th> <th>67.20</th> <th>72.10</th> <th>270.00</th> <th>355.00</th> <th>303.00</th> <th>374.00</th>	Rb	ppm	420.00	329.00	383.00	222.00	105.00	60.50	67.20	72.10	270.00	355.00	303.00	374.00
Ba ppm 668.00 683.00 758.00 712.00 177.00 74.40 75.40 122.00 1140.00 597.00 527.00 699.00 Th ppm 25.70 65.30 27.20 122.00 129.00 115.50 65.80 37.60 121.50 1601.30 98.00 Pb ppm 6.00 5.00 3.00 17.00 38.00 12.00 5.00 3.00 9.00 5.00 2.00 9.00 Nd ppm 16.80 2.40 2.40 59.00 44.10 2.00 14.60 4.50 2.55.00 24.20 2.22.3 13.40 Ta ppm 16.80 2.40 5.00 2.70 6.10 7.80 7.70 17.40 2.55.00 24.20 22.20 13.30 Zr ppm 3.00 8.00 4.50 7.90 7.70 6.10 7.80 7.70 17.40 2.90 1.40 2.00 Y ppm	Sr	ppm	152.00	181.50	35.70	251.00	400.00	144.00	119.00	113.00	306.00	84.50	83.80	83.90
Th ppm 25.70 65.30 27.20 122.00 49.80 99.90 115.50 65.80 37.60 121.50 104.50 98.60 U ppm 6.31 8.49 1.81 13.45 5.50 3.74 2.52 2.77 6.59 8.66 6.66 3.65 Pb ppm 6.00 5.00 3.00 12.00 8.60 4.69 2.20 2.90 2.00 2.00 4.16 4.50 2.55 24.20 22.20 13.40 Ta ppm 16.80 2.48 2.40 59.00 24.10 2.00 21.00 26.10 25.00 24.00 22.20 13.40 Ta ppm 2.80 8.70 8.00 8.00 4.90 27.70 6.10 7.80 7.70 17.40 2.90 1.40 2.00 Y ppm 12.00 14.20 6.70 49.20 21.60 9.80 12.70 8.70 17.40 12.90	Cs	ppm	2.55	2.26	0.81	1.38	1.73	0.56	1.02	1.12	0.78	0.94	0.96	0.84
U ppm 6.31 8.49 1.81 13.45 5.50 3.74 2.52 2.77 6.59 8.66 6.66 3.65 Pb ppm 6.00 5.00 3.00 12.00 5.00 3.00 9.00 5.00 2.00 9.00 Nd ppm 16.80 2.480 5.90 44.10 20.00 14.60 4.50 25.50 24.20 22.20 13.30 Ta ppm 116.80 2.70 1.80 5.00 2.70 2.90 3.80 3.90 3.30 4.10 6.50 3.30 Ta ppm 319.00 313.00 14.90 277.00 7.70 6.10 7.80 7.70 17.40 2.90 1.40 6.00 Y ppm 8.00 8.00 4.50 7.70 7.70 6.10 7.80 7.70 17.40 2.90 1.40 2.00 Y ppm 4.00 4.10 3.00 2.00	Ba	ppm	668.00	683.00	758.00	712.00	177.00	74.40	75.40	122.00	1140.00	597.00	527.00	609.00
Pb ppm 6.00 5.00 3.00 17.00 38.00 12.00 5.00 3.00 9.00 5.00 2.00 9.00 Nd ppm 16.80 24.80 2.40 59.00 44.10 20.00 14.60 2.50 24.20 22.20 13.30 Ta ppm 16.80 2.480 5.00 2.70 305.00 21.00 271.00 303.0 4.10 6.50 3.30 Tr ppm 319.00 313.00 149.00 277.00 305.00 212.00 271.00 261.00 708.00 103.00 42.00 61.00 Y ppm 8.00 8.00 4.90 5.00 1.00 1.00 1.00 1.00 2.00 1.00 1.00 1.00 2.00 5.00 1.00 1.00 1.00 1.00 7.00 5.00 3.00 4.00 7.00 5.00 4.00 7.00 5.00 4.00 7.00 4.00 7.00 5.00 </th <th>Th</th> <th>ppm</th> <th>25.70</th> <th>65.30</th> <th>27.20</th> <th>122.00</th> <th>49.80</th> <th>90.90</th> <th>115.50</th> <th>65.80</th> <th>37.60</th> <th>121.50</th> <th>104.50</th> <th>98.60</th>	Th	ppm	25.70	65.30	27.20	122.00	49.80	90.90	115.50	65.80	37.60	121.50	104.50	98.60
Nd ppm 16.80 24.80 2.40 59.00 44.10 20.00 14.60 4.50 25.50 24.20 22.20 13.40 Ta ppm 2.60 2.70 1.80 5.00 2.70 2.90 3.80 3.90 3.30 4.10 6.50 3.30 Hf ppm 319.00 313.00 149.00 277.00 305.00 212.00 271.00 261.00 7.70 17.40 2.90 1.40 200 Y ppm 12.00 14.20 6.70 49.20 21.60 9.80 12.70 8.70 18.00 62.90 1.63 11.40 Sc ppm 6.00 5.00 1.00 4.00 5.00 1.00 1.00 1.00 7.00 5.00 3.00 4.00 V ppm 6.00 5.00 1.00 1.00 1.00 1.00 1.00 1.00 1.00 1.00 1.00 1.00 1.00 1.00 <th< th=""><th>U</th><th>ppm</th><th>6.31</th><th>8.49</th><th>1.81</th><th>13.45</th><th>5.50</th><th>3.74</th><th>2.52</th><th>2.77</th><th>6.59</th><th>8.66</th><th>6.66</th><th>3.65</th></th<>	U	ppm	6.31	8.49	1.81	13.45	5.50	3.74	2.52	2.77	6.59	8.66	6.66	3.65
Ta ppm 2.60 2.70 1.80 5.00 2.70 2.90 3.80 3.90 3.30 4.10 6.50 3.30 Zr ppm 319.00 313.00 149.00 227.00 305.00 212.00 271.00 261.00 708.00 103.00 42.00 61.00 Hr ppm 8.00 8.00 4.50 7.70 7.70 61.0 7.80 7.70 11.40 2.90 1.40 2.00 Y ppm 6.00 5.00 1.00 4.00 5.00 1.00 1.00 1.00 7.70 5.00 3.00 4.00 V ppm 40.00 41.00 30.00 27.00 48.00 <5	Pb	ppm	6.00	5.00	3.00	17.00	38.00	12.00	5.00	3.00	9.00	5.00	2.00	9.00
Zr ppm 319.00 313.00 149.00 277.00 305.00 212.00 271.00 261.00 708.00 103.00 42.00 61.00 Hf ppm 8.00 8.00 4.50 7.90 7.70 6.10 7.80 7.70 17.40 2.90 1.40 2.00 Y ppm 6.00 5.00 1.00 4.90 1.20 8.70 18.00 62.90 16.30 11.40 Se ppm 6.00 5.00 1.00 4.00 5.00 1.00 1.00 1.00 5.00 3.00 4.00 5.00 4.00 1.00 1.00 2.00 1.00 1.00 1.00 2.00 1.00 1.00 1.00 1.00 1.00 2.00 1.00 1.00 2.00 1.00 1.00 1.00 1.00 1.00 1.00 1.00 1.00 1.00 1.00 1.00 1.00 2.00 1.00 1.00 2.00 3.00 4.00	Nd	ppm	16.80	24.80	2.40	59.00	44.10	20.00	14.60	4.50	25.50	24.20	22.20	13.40
Hr ppm 8.00 8.00 4.50 7.90 7.70 6.10 7.80 7.70 17.40 2.90 1.40 2.00 Y ppm 12.00 14.20 6.70 49.20 21.60 9.80 12.70 8.70 18.00 62.90 16.30 11.40 Se ppm 6.00 5.00 1.00 5.00 1.00 1.00 7.00 7.00 5.00 3.00 4.00 V ppm 40.00 40.00 30.00 27.00 48.00 < 5	Ta	ppm	2.60	2.70	1.80	5.00	2.70	2.90	3.80	3.90	3.30	4.10	6.50	3.30
Y ppm 12.00 14.20 6.70 49.20 21.60 9.80 12.70 8.70 18.00 62.90 16.30 11.40 Se ppm 6.00 5.00 1.00 4.00 5.00 1.00 1.00 1.00 7.00 5.00 3.00 4.00 V ppm 40.00 41.00 30.00 27.00 48.00 <5	Zr	ppm	319.00	313.00	149.00	277.00	305.00	212.00	271.00	261.00	708.00	103.00	42.00	61.00
Sc ppm 6.00 5.00 1.00 4.00 5.00 1.00 1.00 1.00 7.00 5.00 3.00 4.00 V ppm 40.00 41.00 30.00 27.00 48.00 ≤5 ≤5 ≤5 26.00 <5	Hf	ppm	8.00	8.00	4.50	7.90	7.70	6.10	7.80	7.70	17.40	2.90	1.40	2.00
V ppm 40.00 41.00 30.00 27.00 48.00 <5	Y	ppm	12.00	14.20	6.70	49.20	21.60	9.80	12.70	8.70	18.00	62.90	16.30	11.40
Cr ppm 20.00 20.00 10.00 10.00 10.00 <10	Sc	ppm	6.00	5.00	1.00	4.00	5.00	1.00	1.00	1.00	7.00	5.00	3.00	4.00
Co ppm 4.00 7.00 1.00 2.00 2.00 1.00 2.00 3.00 1.00 2.00 2.00 2.00 2.00 1.00 3.00 4.1 3.00 2.00 2.00 <1	v	ppm	40.00	41.00	30.00	27.00	48.00	<5	<5	<5	26.00	<5	6.00	13.00
Ni ppm 1.00 2.00 2.00 2.00 1.00 1.00 3.00 <1	Cr	ppm	20.00	20.00	10.00	10.00	10.00	<10	<10	10.00	10.00	10.00	<10	10.00
Cu ppm 8.00 27.00 2.00 5.00 4.00 3.00 5.00 2.00 4.00 6.00 3.00 4.00 Zn ppm 7.00 7.00 2.00 6.00 5.00 4.00 5.00 5.00 8.00 7.00 5.00 5.00 Ga ppm 7.00 7.00 22.00 19.80 23.80 19.50 19.90 20.30 18.90 18.10 17.40 17.20 Ag ppm 4.05 40.5 <0.5	Co	ppm	4.00	7.00	1.00	2.00	2.00	2.00	1.00	2.00	3.00	1.00	<1	2.00
Zn ppm 7.00 7.00 2.00 6.00 5.00 4.00 5.00 5.00 8.00 7.00 5.00 5.00 Ga ppm 22.50 21.70 22.00 19.80 23.80 19.50 19.90 20.30 18.90 18.10 17.40 17.20 Ag ppm <0.5	Ni	ppm	1.00	2.00	2.00	2.00	1.00	1.00	3.00	<1	3.00	2.00	2.00	<1
Ga ppm 22.50 21.70 22.00 19.80 23.80 19.50 19.90 20.30 18.90 18.10 17.40 17.20 Ag ppm <0.5	Cu	ppm	8.00	27.00	2.00	5.00	4.00	3.00	5.00	2.00	4.00	6.00	3.00	4.00
Ag ppm <0.5	Zn	ppm	7.00	7.00	2.00	6.00	5.00	4.00	5.00	5.00	8.00	7.00	5.00	5.00
Sn ppm 6.00 5.00 3.00 5.00 8.00 4.00 6.00 6.00 9.00 4.00 5.00 3.00 La ppm 16.70 39.00 4.00 47.50 65.80 10.10 6.20 3.40 51.40 29.70 31.30 18.50 Ce ppm 41.00 77.10 7.60 111.00 126.00 27.20 18.10 5.90 98.30 60.50 69.00 39.60 Pr ppm 4.83 8.29 0.70 15.65 13.85 5.08 3.52 1.12 8.65 7.08 7.15 4.30 Nd ppm 16.80 24.80 2.40 59.00 44.10 20.00 14.60 4.50 25.50 24.20 22.20 13.40 Sm ppm 3.48 4.37 0.52 13.35 8.20 4.32 3.28 1.16 4.13 5.73 3.74 2.34 Eu ppm 0.71	Ga	ppm	22.50	21.70	22.00	19.80	23.80	19.50	19.90	20.30	18.90	18.10	17.40	17.20
La ppm 16.70 39.00 4.00 47.50 65.80 10.10 6.20 3.40 51.40 29.70 31.30 18.50 Ce ppm 41.00 77.10 7.60 111.00 126.00 27.20 18.10 5.90 98.30 60.50 69.00 39.60 Pr ppm 4.83 8.29 0.70 15.65 13.85 5.08 3.52 1.12 8.65 7.08 7.15 4.30 Nd ppm 16.80 24.80 2.40 59.00 44.10 20.00 14.60 4.50 25.50 24.20 22.20 13.40 Sm ppm 3.48 4.37 0.52 13.35 8.20 4.32 3.28 1.16 4.13 5.73 3.74 2.34 Eu ppm 0.71 0.65 0.20 2.05 1.41 0.91 0.79 0.47 0.87 0.99 0.54 0.56 Gd ppm 0.3	Ag	ppm	< 0.5	< 0.5	< 0.5	< 0.5	< 0.5	< 0.5	< 0.5	< 0.5	< 0.5	< 0.5	< 0.5	< 0.5
Ce ppm 41.00 77.10 7.60 111.00 126.00 27.20 18.10 5.90 98.30 60.50 69.00 39.60 Pr ppm 4.83 8.29 0.70 15.65 13.85 5.08 3.52 1.12 8.65 7.08 7.15 4.30 Nd ppm 16.80 24.80 2.40 59.00 44.10 20.00 14.60 4.50 25.50 24.20 22.20 13.40 Sm ppm 3.48 4.37 0.52 13.35 8.20 4.32 3.28 1.16 4.13 5.73 3.74 2.34 Eu ppm 0.71 0.65 0.20 2.05 1.41 0.91 0.79 0.47 0.87 0.99 0.54 0.56 Gd ppm 2.47 2.74 0.62 10.10 5.13 2.78 2.41 1.08 2.77 6.45 2.50 1.59 Tb ppm 0.37	Sn	ppm	6.00	5.00	3.00	5.00	8.00	4.00	6.00	6.00	9.00	4.00	5.00	3.00
Pr ppm 4.83 8.29 0.70 15.65 13.85 5.08 3.52 1.12 8.65 7.08 7.15 4.30 Nd ppm 16.80 24.80 2.40 59.00 44.10 20.00 14.60 4.50 25.50 24.20 22.20 13.40 Sm ppm 3.48 4.37 0.52 13.35 8.20 4.32 3.28 1.16 4.13 5.73 3.74 2.34 Eu ppm 0.71 0.65 0.20 2.05 1.41 0.91 0.79 0.47 0.87 0.99 0.54 0.56 Gd ppm 2.47 2.74 0.62 10.10 5.13 2.78 2.41 1.08 2.77 6.45 2.50 1.59 Tb ppm 0.37 0.39 0.13 1.42 0.70 0.38 0.36 0.20 0.42 1.13 0.43 0.27 Dy ppm 0.43 0.	La	ppm	16.70	39.00	4.00	47.50	65.80	10.10	6.20	3.40	51.40	29.70	31.30	18.50
Nd ppm 16.80 24.80 2.40 59.00 44.10 20.00 14.60 4.50 25.50 24.20 22.20 13.40 Sm ppm 3.48 4.37 0.52 13.35 8.20 4.32 3.28 1.16 4.13 5.73 3.74 2.34 Eu ppm 0.71 0.65 0.20 2.05 1.41 0.91 0.79 0.47 0.87 0.99 0.54 0.56 Gd ppm 2.47 2.74 0.62 10.10 5.13 2.78 2.41 1.08 2.77 6.45 2.50 1.59 Tb ppm 0.37 0.39 0.13 1.42 0.70 0.38 0.36 0.20 0.42 1.13 0.43 0.27 Dy ppm 2.32 2.53 0.89 8.64 4.27 2.01 2.39 1.43 2.97 8.18 2.99 1.98 Ho ppm 0.43 0.5	Ce	ppm	41.00	77.10	7.60	111.00	126.00	27.20	18.10	5.90	98.30	60.50	69.00	39.60
Sm ppm 3.48 4.37 0.52 13.35 8.20 4.32 3.28 1.16 4.13 5.73 3.74 2.34 Eu ppm 0.71 0.65 0.20 2.05 1.41 0.91 0.79 0.47 0.87 0.99 0.54 0.56 Gd ppm 2.47 2.74 0.62 10.10 5.13 2.78 2.41 1.08 2.77 6.45 2.50 1.59 Tb ppm 0.37 0.39 0.13 1.42 0.70 0.38 0.36 0.20 0.42 1.13 0.43 0.27 Dy ppm 2.32 2.53 0.89 8.64 4.27 2.01 2.39 1.43 2.97 8.18 2.99 1.98 Ho ppm 0.43 0.50 0.22 1.65 0.79 0.36 0.48 0.32 0.64 1.86 0.61 0.39 Er ppm 1.47 1.62	Pr	ppm	4.83	8.29	0.70	15.65	13.85	5.08	3.52	1.12	8.65	7.08	7.15	4.30
Eu ppm 0.71 0.65 0.20 2.05 1.41 0.91 0.79 0.47 0.87 0.99 0.54 0.56 Gd ppm 2.47 2.74 0.62 10.10 5.13 2.78 2.41 1.08 2.77 6.45 2.50 1.59 Tb ppm 0.37 0.39 0.13 1.42 0.70 0.38 0.36 0.20 0.42 1.13 0.43 0.27 Dy ppm 2.32 2.53 0.89 8.64 4.27 2.01 2.39 1.43 2.97 8.18 2.99 1.98 Ho ppm 0.43 0.50 0.22 1.65 0.79 0.36 0.48 0.32 0.64 1.86 0.61 0.39 Er ppm 1.47 1.62 0.81 4.93 2.28 1.00 1.53 1.12 2.37 5.96 1.94 1.28 Tm ppm 0.24 0.28	Nd	ppm	16.80	24.80	2.40	59.00	44.10	20.00	14.60	4.50	25.50	24.20	22.20	13.40
Gd ppm 2.47 2.74 0.62 10.10 5.13 2.78 2.41 1.08 2.77 6.45 2.50 1.59 Tb ppm 0.37 0.39 0.13 1.42 0.70 0.38 0.36 0.20 0.42 1.13 0.43 0.27 Dy ppm 2.32 2.53 0.89 8.64 4.27 2.01 2.39 1.43 2.97 8.18 2.99 1.98 Ho ppm 0.43 0.50 0.22 1.65 0.79 0.36 0.48 0.32 0.64 1.86 0.61 0.39 Er ppm 1.47 1.62 0.81 4.93 2.28 1.00 1.53 1.12 2.37 5.96 1.94 1.28 Tm ppm 0.24 0.28 0.15 0.76 0.37 0.17 0.26 0.20 0.45 0.97 0.34 0.23 Yb ppm 1.77 1.97	Sm	ppm	3.48	4.37	0.52	13.35	8.20	4.32	3.28	1.16	4.13	5.73	3.74	2.34
Tb ppm 0.37 0.39 0.13 1.42 0.70 0.38 0.36 0.20 0.42 1.13 0.43 0.27 Dy ppm 2.32 2.53 0.89 8.64 4.27 2.01 2.39 1.43 2.97 8.18 2.99 1.98 Ho ppm 0.43 0.50 0.22 1.65 0.79 0.36 0.48 0.32 0.64 1.86 0.61 0.39 Er ppm 1.47 1.62 0.81 4.93 2.28 1.00 1.53 1.12 2.37 5.96 1.94 1.28 Tm ppm 0.24 0.28 0.15 0.76 0.37 0.17 0.26 0.20 0.45 0.97 0.34 0.23 Yb ppm 1.77 1.97 1.14 5.22 2.82 1.18 1.90 1.51 3.39 6.19 2.38 1.67 Lu ppm 0.30 0.33	Eu	ppm	0.71	0.65	0.20	2.05	1.41	0.91	0.79	0.47	0.87	0.99	0.54	0.56
Dy ppm 2.32 2.53 0.89 8.64 4.27 2.01 2.39 1.43 2.97 8.18 2.99 1.98 Ho ppm 0.43 0.50 0.22 1.65 0.79 0.36 0.48 0.32 0.64 1.86 0.61 0.39 Er ppm 1.47 1.62 0.81 4.93 2.28 1.00 1.53 1.12 2.37 5.96 1.94 1.28 Tm ppm 0.24 0.28 0.15 0.76 0.37 0.17 0.26 0.20 0.45 0.97 0.34 0.23 Yb ppm 1.77 1.97 1.14 5.22 2.82 1.18 1.90 1.51 3.39 6.19 2.38 1.67 Lu ppm 0.30 0.33 0.19 0.78 0.41 0.18 0.31 0.22 0.64 0.93 0.32 0.25 Eu/Eu* 0.74 0.57 1.08	Gd	ppm	2.47	2.74	0.62	10.10	5.13	2.78	2.41	1.08	2.77	6.45	2.50	1.59
Ho ppm 0.43 0.50 0.22 1.65 0.79 0.36 0.48 0.32 0.64 1.86 0.61 0.39 Er ppm 1.47 1.62 0.81 4.93 2.28 1.00 1.53 1.12 2.37 5.96 1.94 1.28 Tm ppm 0.24 0.28 0.15 0.76 0.37 0.17 0.26 0.20 0.45 0.97 0.34 0.23 Yb ppm 1.77 1.97 1.14 5.22 2.82 1.18 1.90 1.51 3.39 6.19 2.38 1.67 Lu ppm 0.30 0.33 0.19 0.78 0.41 0.18 0.31 0.22 0.64 0.93 0.32 0.25 Eu/Eu* 0.74 0.57 1.08 0.54 0.66 0.80 0.86 1.28 0.79 0.50 0.54 0.89	Tb	ppm	0.37	0.39	0.13	1.42	0.70	0.38	0.36	0.20	0.42	1.13	0.43	0.27
Er ppm 1.47 1.62 0.81 4.93 2.28 1.00 1.53 1.12 2.37 5.96 1.94 1.28 Tm ppm 0.24 0.28 0.15 0.76 0.37 0.17 0.26 0.20 0.45 0.97 0.34 0.23 Yb ppm 1.77 1.97 1.14 5.22 2.82 1.18 1.90 1.51 3.39 6.19 2.38 1.67 Lu ppm 0.30 0.33 0.19 0.78 0.41 0.18 0.31 0.22 0.64 0.93 0.32 0.25 Eu/Eu* 0.74 0.57 1.08 0.54 0.66 0.80 0.86 1.28 0.79 0.50 0.54 0.89	Dy	ppm	2.32	2.53	0.89	8.64	4.27	2.01	2.39	1.43	2.97	8.18	2.99	1.98
Tm ppm 0.24 0.28 0.15 0.76 0.37 0.17 0.26 0.20 0.45 0.97 0.34 0.23 Yb ppm 1.77 1.97 1.14 5.22 2.82 1.18 1.90 1.51 3.39 6.19 2.38 1.67 Lu ppm 0.30 0.33 0.19 0.78 0.41 0.18 0.31 0.22 0.64 0.93 0.32 0.25 Eu/Eu* 0.74 0.57 1.08 0.54 0.66 0.80 0.86 1.28 0.79 0.50 0.54 0.89	Но	ppm	0.43	0.50	0.22	1.65	0.79	0.36	0.48	0.32	0.64	1.86	0.61	0.39
Yb ppm 1.77 1.97 1.14 5.22 2.82 1.18 1.90 1.51 3.39 6.19 2.38 1.67 Lu ppm 0.30 0.33 0.19 0.78 0.41 0.18 0.31 0.22 0.64 0.93 0.32 0.25 Eu/Eu* 0.74 0.57 1.08 0.54 0.66 0.80 0.86 1.28 0.79 0.50 0.54 0.89	Er	ppm	1.47	1.62	0.81	4.93	2.28	1.00	1.53	1.12	2.37	5.96	1.94	1.28
Lu ppm 0.30 0.33 0.19 0.78 0.41 0.18 0.31 0.22 0.64 0.93 0.32 0.25 Eu/Eu* 0.74 0.57 1.08 0.54 0.66 0.80 0.86 1.28 0.79 0.50 0.54 0.89	Tm	ppm	0.24	0.28	0.15	0.76	0.37	0.17	0.26	0.20	0.45	0.97	0.34	0.23
Eu/Eu* 0.74 0.57 1.08 0.54 0.66 0.80 0.86 1.28 0.79 0.50 0.54 0.89	Yb	ppm	1.77	1.97	1.14	5.22	2.82	1.18	1.90	1.51	3.39	6.19	2.38	1.67
	Lu	ppm	0.30	0.33	0.19	0.78	0.41	0.18	0.31	0.22	0.64	0.93	0.32	0.25
(La/Yb) _{Nb} 6.38 13.38 2.37 6.15 15.77 5.78 2.21 1.52 10.25 3.24 8.89 7.49	Eu/Eu*		0.74	0.57	1.08	0.54	0.66	0.80	0.86	1.28	0.79	0.50	0.54	0.89
	$(La/Yb)_{Nb}$		6.38	13.38	2.37	6.15	15.77	5.78	2.21	1.52	10.25	3.24	8.89	7.49

 $Eu/Eu* = (Eu/0.087)/(sqrt((Sm/0.231)*(Gd/0.306))) \; (Taylor \; \& \; McLennan \; 1885)$

 $REE\ Frac = (La/Yb)_{Nb} = (La/0.367)/(Yb/0.248)\ (Taylor\ \&\ McLennan\ 1985)$

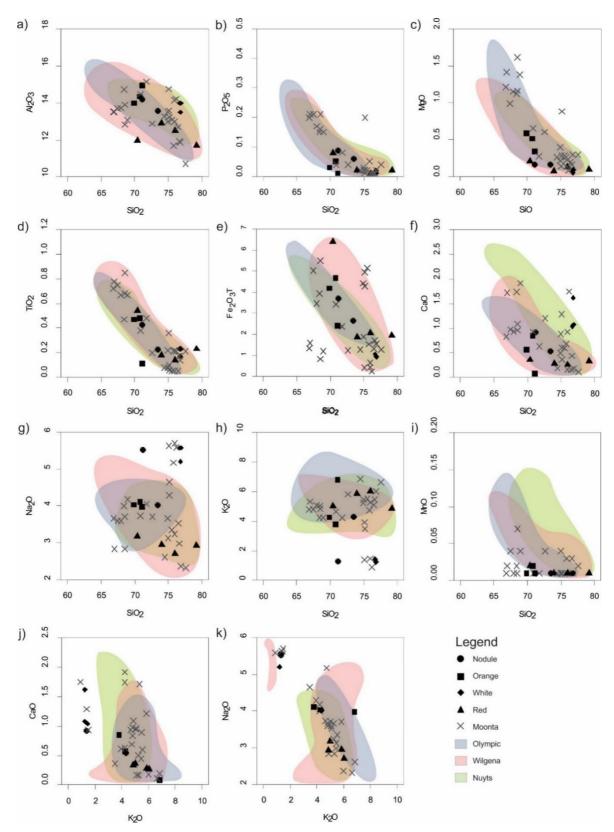


Figure 6. Major element variation diagrams of major elements for samples from Point Riley. Sample location shown in Figure 2. Data from the Moonta (Creaser 1989; Wurst 1994; Giles 1980), Olympic (Creaser 1989; OZCHEM; SARIG), Wilgena (Foden & Stewart 2005; SARIG) and Nuyts (Foden & Stewart 2005; SARIG) domains are also shown for comparison.

Trace and Rare Earth Elements

Trace and rare earth element geochemical results are displayed in Figure 7 and Table 2. A positive correlation is evident in Th with increasing SiO₂ content whilst a negative trend is observed for Sc, Zr, Ba, Ce and La (Fig 7a–f). Sr has an overall negative trend although there is a slight inflection at ~73 wt% SiO₂ for Sr (Fig 7g).

Primitive mantle normalised multi element trace element plots show well defined patterns for each of the granite phases, and are characterised by positive anomalies in Rb, Th and Ta, and negative anomalies in Ba, Nb, P and Ti (Fig. 8). There is a slight variance in the Zr content where all but three red granite samples have a positive anomaly. (Fig. 8). Similarly, Pb shows a positive anomaly in all samples except one from each of the orange and red granite phases that have slight negative anomalies.

Chondrite normalised rare earth element (REE) diagrams are presented in Figure 9. All samples show moderately steep REE patterns due to relative enrichment in light REE (LREE), with moderate negative Eu anomalies (Eu/Eu* = 0.49–0.88) (Fig. 9 and Table 2). The nodules display the highest REE enrichment, followed by the red and orange granite, then the white granite (Fig. 9). One sample from the orange and white granite phases are relatively depleted in REE compared with other samples, have an overall flatter REE pattern and positive Eu anomalies (Eu/Eu* = 1.08 and 1.28). REE fractionation (La/Yb)_{Nb} plotted against Eu/Eu* (Fig. 10) illustrates higher degrees of fractionation in the nodules, orange and red granite compared to the white leucogranite. The white leucogranite has consistently higher Eu values compared to the other granite phases, which preserve more variable Eu values.

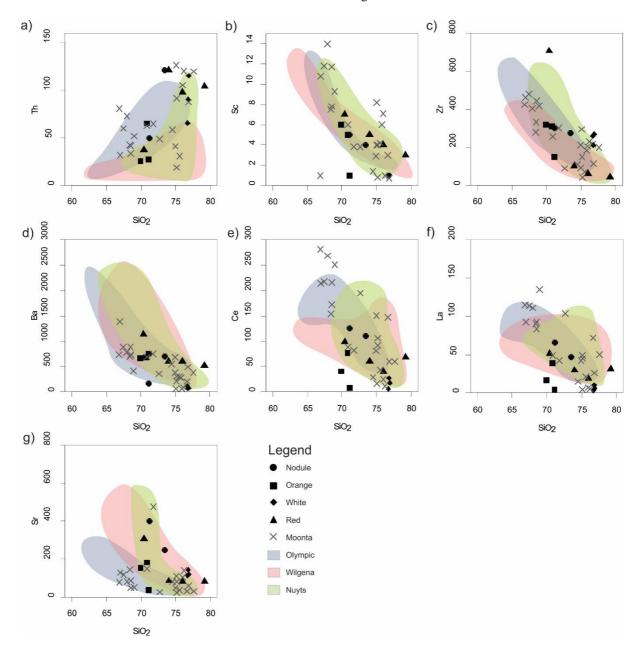


Figure 7. Trace element variation diagrams of trace elements for samples from Point Riley. Sample locations shown in Figure 2. Data from the Moonta (Creaser 1989; Wurst 1994; Giles 1980), Olympic (Creaser 1989; OZCHEM; SARIG), Wilgena (Foden & Stewart 2005; SARIG) and Nuyts (Foden & Stewart 2005; SARIG) domains are also shown for comparison.

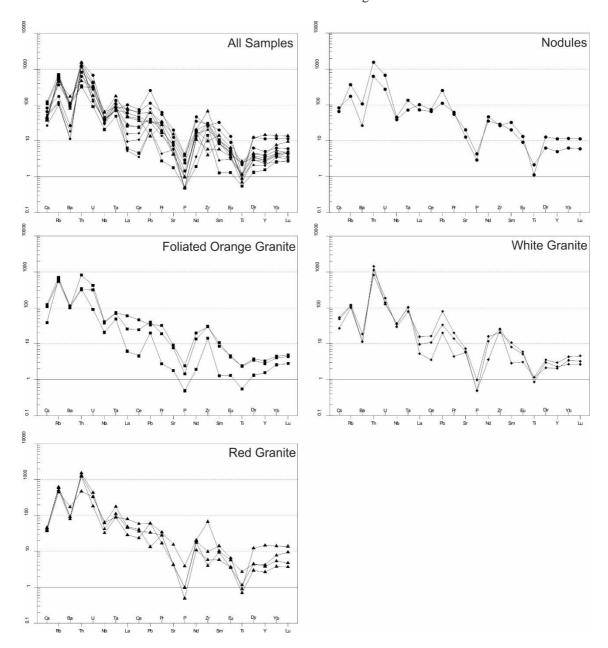


Figure 8. Primitive mantle normalised trace element spidergrams for samples from Point Riley. Data normalised to values of McDonough & Sun (1995). Sample locations shown in Figure 2.

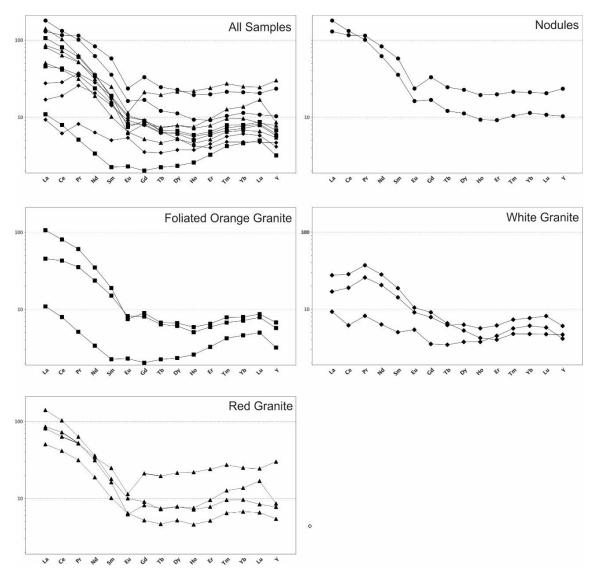


Figure 9. Chondrite normalised REE spidergrams for Point Riley samples. Data normalised to values of Taylor & McLennan (1985). Sample locations shown in Figure 2.

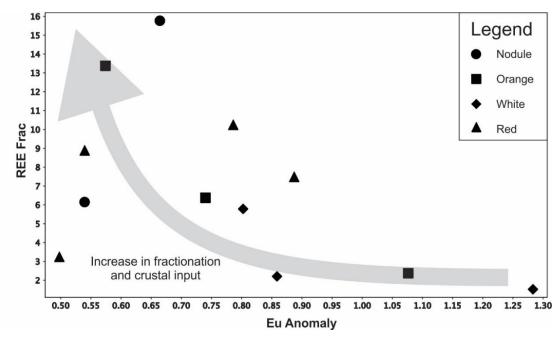


Figure 10. Rare earth element fractionation (REE Frac) versus Eu anomaly plot for the Point Riley samples. Data normalised to Taylor & McLennan (1985). Equations used for calculations are given in Table 3. Sample location shown in Figure 2.

Sm-Nd isotope analysis

Sm–Nd isotope results for two intensely foliated orange granites, one nodule, two white and three red granite samples are given in Table 3 and Figure 11. Initial εNd(i) values were calculated at 1590, 1585 and 1580 for the orange, white and red granite phases respectively based on the known age of the Tickera Granite (ca. 1597–1577 Ma; Conor 1995) and interpreted relative timing of emplacement of the granite phases. εNd(i) values for the orange granite and nodule are well grouped and range from –6.73 and –3.55. The white and red granite phases have broad εNd(i) values ranging from –9.78 and –2.42 and –13.81 and –5.13 respectively. ¹⁴⁷Sm/¹⁴⁴Nd ratios range between 0.1294 and 0.1390 for the orange granite and nodules, 0.1296 and 0.1561 for the white leucogranite, and 0.1098 and 0.1663 for the red granite. T_{DM} values for the orange granite and nodules range from 2601–2804 Ma, from 2383–2771 Ma for the white leucogranite and from 2433–3079 Ma for the red granites (Table 3).

Table 3. Sm–Nd isotopic data for selected samples from Point Riley, Black rock and Wallaroo North Beach. Sample location shown in Figures 1c and 2.

Sample No.	Granite Phase	Age (Ma)	Sm (ppm)	Nd (ppm)	¹⁴⁷ Sm/ ¹⁴⁴ Nd	¹⁴³ Nd/ ¹⁴⁴ Nd	2se	ε _{Nd} (0)	ε _{Nd} (i)	T _{DM} (Ma)
2147276	Orange	1590	3.7	16.9	.1315	.510239	0.000001	-19.98	-6.73	2804
2147281	Orange	1590	0.5	2.4	.1294	.510315	0.000011	-18.94	- 5.25	2641
2147273	Nodule	1590	14.3	62.1	.1390	.510402	0.000002	-15.27	-3.55	2601
2147272	White	1585	4.6	21.3	.1296	.510466	0.000001	-16.04	-2.42	2383
2147277	White	1585	1.1	4.3	.1561	.510090	0.000003	-17.98	- 9.78	2771
2147282	Red	1580	2.34	13.4	.1098	.510326	0.000002	-22.85	-5.29	2439
2149822	Red (North Beach)	1580	0.8	2.9	.1663	.509891	0.000003	-19.89	-13.81	3079
2149823	Red (Black Rock)	1580	15.5	84.7	.1105	.510334	0.000002	-22.56	-5.13	2433

 $^{^{147}}Sm/^{144}Nd\ CHUR\ T{=}0-0.1966\ (Goldstein\ et\ al.,\ 1984)$

 $^{^{147}}$ Sm/ 144 Nd DM T=0 – 0.2145 (Goldstein et al., 1984)

¹⁴³Nd/¹⁴⁴Nd CHUR T=0 – 0.512638 (Goldstein et al., 1984)

 $^{^{143}}Nd/^{144}Nd~DM~T=0-0.513150~(Goldstein~et~al.,~1984)$

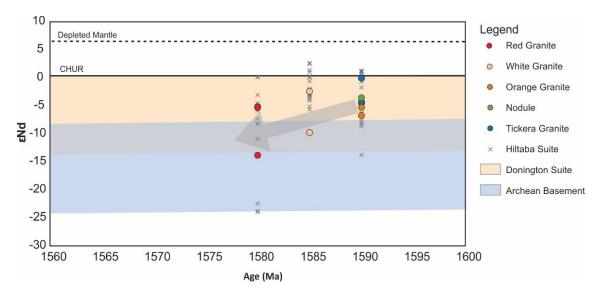


Figure 11. £Nd evolution diagram for granite phases at Point Riley, Black Rock and Wallaroo North Beach. Sample locations are shown in Figures 1c and 2. Data for the surrounding Tickera Granite (Giles 1980; Creaser 1989; Wurst 1994) and for Hiltaba Suite granites from other domains (Creaser 1989; Foden & Stewart 2005; Budd & Skirrow 2007; Skirrow et al., 2007; Rusak 2009) are also shown. Fields for the Donington Suite (Mortimer 1984; Creaser 1989; Turner et al., 1993; Bendall 1994; Schaefer 1998) and Archean Basement (Cooyerdroo Granite and Sleaford Complex; Turner et al., 1993; Swain et al., 2005; Fraser et al., 2010; OZCHEM; SARIG) are also displayed.

DISCUSSION

Relative timing of emplacement of granitic phases

The granitic nodules observed at Point Riley have similar mineralogy and chemistry to the intensely foliated orange granite that hosts them (Figs. 6, 7 and Table 2); however, they differ in degrees of relief due to weathering (Fig 3a). Granitic nodules similar in appearance to those observed at Point Riley have previously been described as granitic 'corestones', which are spherical masses surrounded by weathered rock of the same composition (Twidale & Vidal Romani 2005; Hirata et al., 2016). Corestones form as a result of spheroidal and chemical weathering (hydrolysis) in hard uniform rocks with well–developed joint patterns over an extended period of time (Ollier 1967; Hirata et al., 2016). The similarity of the Point Riley nodules in mineralogy, chemistry, and deformation style to the hosting orange granite, and the morphology of the nodules

suggests that the nodules are corestones, and are therefore the same phase as the orange granite.

Contact relationships are often used to infer relative timing of emplacement of granitic phases. Apophyses have previously been described as contact features between igneous phases, and have been used to demonstrate that the phase containing the apophyses is the relatively younger phase (e.g. De Wet et al., 1989). Apophyses observed at the contact boundary of the white leucogranite and the orange granite (Figs. 2 and 3c) indicate that the orange granite was emplaced before the white leucogranite. A contact metamorphic boundary preserved in the centre of the study area that shows thermal alteration of the white leucogranite by the red granite (Figs. 2 and 3e), suggesting the red granite is younger than the white leucogranite. However, in the northeast part of the study area, magma mingling between the red and white granite phases is observed (Figs. 2 and 3h). Magma mingling develops when two magmas partially mix and likely takes place both at and en route to the emplacement site (Snyder et al., 1997; Vernon et al., 1998), and has been interpreted to represent coeval magmatism (Medeiros et al., 2001). Evidence of magma mingling between the white leucogranite and red granite in the Point Riley area suggests that these two phases may locally be coeval.

A raft of the white leucogranite is preserved within the red granite dyke (Figs. 2 and 3f). This raft may have formed from melting of the white leucogranite as the red granite intruded, mingling of the two magmas (Vernon et al., 1998), or due to a 3–dimensional feature being exposed on a 2–dimensional rock platform. Evidence of separate and coeval timings of emplacement of the red and white granite phases indicates that the white leucogranite was both locally crystalline and partially molten when the red granite

intruded, suggesting the red granite intruded at the waning stages of white leucogranite magmatism. Overall, the intensely foliated orange granite and nodules are interpreted to be the same phase and are the oldest granites exposed in the Point Riley area. These were intruded by the white leucogranite, and then the youngest phase; the red granite.

Petrogenesis of Point Riley Granites

Geochemical data can be used to model the source and evolution of magmas (Winter 2010; Hertogen & Mareels, 2016). Granitic phases crystallising from the same magma chamber will cause the residual magma to become progressively more silica-rich through fractional crystallisation, therefore older granite phases will have the lowest SiO₂ content (Winter 2010; Hertogen & Mareels, 2016). As plagioclase has a higher melting temperature relative to K-feldspar and quartz, it is also expected that the older granite phases will have higher Ca and Na content relative to younger granite phases. At Point Riley, the orange and white granites show an increasing trend in SiO₂ where the white leucogranite is more SiO₂ rich, suggesting it represents the younger phase if both phases were from derived the same evolving source. Conversely, the red granite has the broadest SiO₂ content, which overlaps with the SiO₂ range of the orange and white granites (Figs. 6 and 7), suggesting the red granite could be from another source. However, no other geochemical data supports this notion. The orange to red granite phases show a decrease in Ca and Na content with decreasing SiO₂, however the white leucogranite shows the opposite trend (Fig. 6f, g), which is likely due to the high modal proportion of plagioclase in this phase. Additionally, the orange and red granites have higher K₂O content as they are predominantly K-feldspar rich. Additionally, the white leucogranite preserves relatively low Fe₂O₃ content and is a slight outlier in Al₂O₃

content (Fig. 6a, e). These geochemical variances are likely a result of distinctive differences in mineralogy and alteration as the white leucogranite contains negligible amounts of K-feldspar or hematite alteration, unlike the other granite phases. Based on these differences in mineralogy and geochemistry, it is inferred that the white leucogranite is from a different source region to the orange and red granite phases. Enriched light rare earth element (LREE) content in granites is considered indicative of higher degrees of crustal contamination as LREEs are more abundant in the crust while heavy rare earth elements (HREE) are more abundant in the mantle (Walters et al., 2013). Overall, the nodules, orange and red granite phases exhibit similar REE patterns showing enriched LREE contents and high LREE/HREE ratios (Table 2; Fig. 10). This supports the conclusion that these phases are from a similar crustal source region, although indication of locally variable sources are evident from the comparatively different REE patterns within the granite phases (Fig. 10). The white leucogranite preserves a significantly different REE pattern showing relatively lower LREE/HREE ratios compared to the other granite phases, suggesting a more mafic source. Positive Eu anomalies infer less fractionation of plagioclase and may have been inherited from a source that was characterised by positive Eu anomalies, while negative Eu anomalies suggests fractionation of plagioclase during formation (Winter 2010). One white and one orange granite sample (samples 2147277 and 2147281 respectively) display relatively flat to positive Eu anomalies and relatively low LREE/HREE ratios (Fig. 10), suggesting the incorporation of a more mafic material. Alternatively, the remainder of the samples are characterised by negative Eu anomalies, suggesting a crustal source. Two red granites and one nodule display high degrees of plagioclase fractionation but relatively low LREE/HREE ratios (Fig. 10), which also infers

incorporation of more mafic or lower crustal material into these phases. These interpretations suggest the granite phases do not solely have an evolved crustal source, rather a more variable source including amalgamation of a lower crustal and/or mafic material.

Sm–Nd isotopes can be used to determine the source region of magmas and the relative contribution of crustal versus mantle material (e.g. DePaolo et al., 1992). The εNd values for the samples analysed in this study cover a broad range from –2.44 to –13.81 (Fig. 11). Previously published εNd values for samples of the Tickera Granite which were taken from the sample pluton in the Moonta region (Giles 1980; Creaser 1989; Wurst 1994), show similar values despite having an older model age (Fig. 11). Overall, the phases of Tickera Granite in the Moonta region show increasingly negative εNd(i) values over time, which is indicative of progressive input of crustal material throughout magma evolution. This is likely due to the assimilation and fractional crystallisation of the surrounding wall rock as the magma ascends and evolves. One sample of the white leucogranite has a more juvenile εNd value (–2.42: 2147272) than the other granite phases, indicating some input from a more mafic source.

Overall, the mineralogy, geochemical and isotopic data for the Tickera Granite suggests that the nodules, orange and red granite phases are from a similar source, and the white leucogranite is from a slightly different, more mafic source region. Possible sources for the Tickera Granite in the Moonta Domain includes the underlying Wallaroo Group and Donington Suite. The ε Nd values for the Donington Suite partially overlap with the Point Riley samples (Fig. 11), and are general relatively juvenile, therefore is unlikely to be the sole source of the Tickera Granite. The Archean basement underlying the Wallaroo Group and Donington Suite is also considered as potential source region. The

ENd values of Archean basement (the Mesoarchean Cooyerdoo Granite and Neoarchean Sleaford Complex) ranges from approximately -8 to -24 (Turner et al., 1993; Swain et al., 2005; Fraser et al., 2010; Fig. 10), and are significantly more evolved than the majority of the samples analysed in this study (Fig. 11). ENd data indicates that the Donington Suite is a likely source for the Tickera Granite. However, it is difficult to assess the potential input from the Wallaroo Group as there is currently no ENd data available. Variable REE content of the granite phases suggests input from a crustal source of variable nature. The Wallaroo Group which includes psammites, siltstones, calcsilicates, albitites, quartzites, iron rich sediments and volcanics (e.g Cowley et al., 2003) has the potential to produce melt with variable REE content (and therefore εNd values), and cannot be discounted as a potential source region. Similarly, the Donington Suite comprises igneous rocks of mafic to felsic composition, which also has the potential to produce variable REE content and \(\varepsilon \) Nd values. Based on the εNd and REE data, it is suggested that the Tickera Granite may be derived from the Donington Suite and/or Wallaroo Group with some input from a more mafic or lower crustal source, which could potentially be the underlying Archean basement.

Comparison with the broader Hiltaba Suite

The Tickera Granite is generally inferred to be part of the Hiltaba Suite granites (e.g. Conor 1995; Conor et al., 2010). Comparison of whole rock geochemical data for samples of the Tickera Granite collected in this study with published data from Hiltabaaged (ca. 1595–1575 Ma) granites of the Moonta Domain (Fig. 1b) shows similar trends. Major and trace element data generally overlaps, with the exception of a cluster

of samples for the Moonta Domain that are elevated in P₂O₅, MgO and TiO₂ (Fig. 6b–d), and Sc, Th, Ce and La (Fig. 7b, c, e, f). These differences are likely due to samples being taken from the Arthurton Granite and slight differences between plutons in the Moonta region, but can generally be inferred to be of similar composition.

Comparison of the Tickera Granite samples from this study with the broader Hiltaba Suite granites from the Olympic, Wilgena and Nuyts Domains (Fig. 1b) shows that the data defines similar linear trends for most major and trace elements. Differences in chemistry of Hiltaba Suite granites between domains has previously been studied by Stewart & Foden (2003). The Moonta Domains considered to preserve elevated Th concentrations, and depleted CaO, Sr and MnO concentrations compared to the rest of the suite (Stewart and Foden 2005). The samples used in this study generally support this interpretation however, few of the samples show relatively high Sr values compared to the Moonta Domain and the rest of the Hiltaba Suite. However, the white leucogranite mapped at Point Riley has elevated Na₂O and depleted K₂O concentrations compared to other data from the Tickera Granite and broader Hiltaba Suite, with the exception of 3–4 data points from the Moonta Domain and three samples from the Wilgena Domain (Fig. 6k), which is attributed to this phase being plagioclase rich. Similarly, the high Ca/low K concentrations for samples of the white leucogranite at Point Riley is attributed to this phase being plagioclase rich and K-feldspar-poor (Fig. 6j). In general, the plagioclase-rich nature of the white leucogranite contrasts with the typically K-feldspar-rich nature of the Hiltaba Suite granites (e.g. Daly et al., 1993; Stewart & Foden 2003). Data in this study has also shown that the white leucogranite is from a different source to the other phases observed at Point Riley. Comparison of ENd

signatures of the Tickera Granite samples of this study with the broader Hiltaba Suite granites shows that the data are broadly comparable, (Fig. 11).

Similar major, trace element and isotopic data between the Tickera Granite and the majority of the Hiltaba Suite supports the notion that the Tickera Granite is part of the broader Hiltaba Suite. However, it is noted that there are localised phases within the Moonta Domain (including the Tickera Granite) and Wilgena Domain that suggests they were derived from a slightly different source region compared to the broader Hiltaba Suite. This is supported by Stewart and Foden (2005) who suggests that the Hiltaba Suite is a combination of mantle and crustal material, with the crustal component being less than 30%, and variations related to different amounts of compositions of mantle and crustal endmembers.

Structural fabrics and deformation style within Yorke Peninsula

The mechanisms by which deformation is accommodated in granites depends on a number of variables including crustal level of emplacement, granite rheology and morphology (e.g. Evans 1988; Tobisch & Paterson 1998). At higher metamorphic grades foliations develop, while at lower grades cleavages develop (Winter 2010). Structures can also manifest differently as a result of a composite of structures from progressive deformation events (Tobisch & Paterson 1998). A well–developed northeast–trending foliation is preserved in the orange granite and locally in the red granite (Figs. 3a, b and 4a, c). Conversely, the white leucogranite preserves a well–developed cleavage in the same northeast–trending orientation (Figs. 3d and Fig. 4b). The pervasive northeast-trending fabrics observed may have developed in the same deformation event, however, have manifested differently in the granite phases due to

differences in rheology, emplacement depths and/or due to a combination of structures from multiple deformation events.

Evidence of emplacement of the red granite at shallow crustal levels is demonstrated by the local preservation of cavities within this granite phase (Fig. 2, 3g). These cavities have been suggested to be miarolitic cavities (Conor et al., 2010), which develop during uplift as pockets of gas expand within a magma, and are therefore indicative of emplacement at shallow crustal levels (Petford 2003). The presence of these cavities in the red granite suggests this phase was emplaced at shallow crustal levels, supporting the suggestion that the degree of preservation of the dominant northeast—trending fabric in the Point Riley area is variable due to the variation of crustal levels where the granites were emplaced. However, it is noted that these features have not indisputably been demonstrated to be miarolitic cavities, but may have resulted from other processes such as weathering and the removal of K—feldspar phenocrysts. Similar cavities have been reported in the Arthurton Granite, and are interpreted to have resulted from exsolution of magmatic volatile phases (Zang et al. 2007).

The preservation of the northeast–trending foliation along the margins of the red granite, combined with the knowledge that magmas crystallise from the outer edges inwards (Winter 2010), suggests that the initial stages of emplacement of the red granite may have occurred during the waning stages of this deformation event. These northeast–trending structures are also subparallel to the orientation of the shoreline at Point Riley, and have been correlated with a major northeasterly trending structure just off–shore (Conor et al., 2010).

The northeast-trending fabric has also been observed in the Wallaroo Group metasediments in the Point Riley area (Wurst 1994; Arcaro 2000; Zang et al., 2006;

Conor et al. 2010), and is also seen as a strongly developed northeast-trending structural grain across the Yorke Peninsula (e.g. Wurst 1994; Arcaro 2000; Conor 2002; Conor et al. 2010; Conor 2016). This fabric may be associated with the later deformation event described by Conor et al., (2010) that resulted in development of northeast-trending upright folds. In the Wallaroo North Beach area, exposures of the upright folds are preserved within the Wallaroo Group metasediments, where they are observed to be refolding an earlier generation of isoclinal folds to produce a Type 3 fold interference pattern (See Plate 38 in Conor 2016; Wurst 1994; Arcaro 2000; Conor et al. 2010). Development of Type 3 fold interference patterns implies the axial trace of the earlier fold generation was subparallel to the axial trace of the overprinting fold generation, and that the axial plane of the folds were at a high angle (Ramsay 1962). This implies that the earlier isoclinal fold generation in the Point Riley area had a northeast trending axial trace and a shallowly inclined to recumbent axial plane, and that the northeast-trending fabric may be a composite of the earlier isoclinal folds and the later upright folds. The mechanism for development of recumbent folds with a northeast trending axial trace would likely be northwest–southeast directed shortening. Long-limbed, upright, tight to isoclinal folds with northeast-trending fold axial planes have been described in the eastern Yorke Peninsula from aeromagnetic imagery (Parker 1993; Wurst 1994; Conor 2002; Conor et al., 2010). The relationship between these isoclinal folds and the northeast-trending upright folds observed in the Wallaroo Group in northwestern Yorke Peninsula is unknown, however the northeast-trending orientation supports the notion that the two fold generations observed in the Yorke Peninsula may have produced a composite northeast-trending fabric that is seen throughout the region.

A second, poorly– to moderately–developed, northwest–trending cleavage is preserved within all granite phases at Point Riley at different degrees, and overprints the dominant foliation seen in the orange and red granites (Fig. 2, 3c, d, f). This cleavage is interpreted to have developed during a later deformation event, possibly when the granites in the Point Riley area were at shallower crustal levels where deformation is accommodated via cleavage rather than foliation development. Northwest-trending structures including conjugate faults, fractures and shear zones have been recognised in the western Yorke Peninsula (e.g. Arcaro 2000; Zang et al., 2006; Conor et al. 2010) and in aeromagnetic magnetic imagery (Conor 2002; Conor 2016). Formation of these structures are suggested to be a product of emplacement of the Tickera Granite, resulting in mineralisation in the region (e.g. Dickinson 1953; Wurst 1994; Zang et al., 2006; Conor et al., 2010). The similar orientation of cleavages at Point Riley and regional structures in the northwestern Yorke Peninsula suggests that later deformation may have been localised, and occurred subsequently after granite emplacement. . Few studies have been done to compare structural fabrics preserved within the Tickera Granite to the surrounding Wallaroo Group (e.g. Wurst 1994, Arcaro 2000), and to relate the timing of deformation within the granites and the hosting metasediments (Wurst 1994; Arcaro 2000; Conor et al. 2010). The timing of deformation events that effected the Tickera Granite has generally been constrained to be synchronous with the timing of intrusion of the Point Riley granite phases at ca. 1597–1577 Ma (Conor 1995). This implies that the later northeast–trending upright folds developed at this time. This correlation and interpreted timing of deformation is supported by Conor et al. (2010), who suggest that the later upright folds are attributed to deformation associated with intrusion of the Hiltaba Suite granites.

The timing of the earlier isoclinal fold event is equivocal, and has been inferred to be related to the ca. 1730–1690 Ma Kimban Orogeny (Conor et al. 2010). This conclusion is based on the absence of amphibolite—rich calc—silicate alteration in the isoclinal folds, which is observed within the upright fold generation and is interpreted to be associated with the Hiltaba Suite event (Conor et al., 2010). This suggests deformation of the two fold generations may have occurred during the Kimban Orogeny, but does not exclude the possibility that the isoclinal folds developed just prior to initial stages of granite intrusion at ca. 1595 Ma and generation of the upright folds. The latter may overlap with the timing of intense deformation and metamorphism across eastern Proterozoic Australia, including the Gawler Craton and Curnamona Province at ca. 1600–1580 Ma (e.g. Page et al. 2005; Hand et al., 2007; Forbes et al., 2008, 2012; Fig. 12).

Relationships with the northern Gawler Craton and Curnamona Province

The late Palaeo to early Mesoproterozoic marked a significant period of metamorphism, deformation, magmatism and mineralisation within eastern Proterozoic Australia (e.g. Collins & Shaw 1995; Betts et al., 2002; Giles & Nutman 2002; Hand et al. 2007; Forbes et al., 2008; Stewart & Betts 2010; Forbes et al., 2012). Within southern Proterozoic Australia, this event is unnamed, but is recognised within the Gawler Craton (e.g. Hand et al. 2007; Szpunar et al. 2007; Cutts et al. 2011; Forbes et al. 2011, 2012; Morrissey et al. 2014), and is identified as the ca. 1600–1585 Ma Olarian Orogeny within the Curnamona Province (e.g. Page et al. 2005; Forbes et al. 2008; Fig. 12.

Within the northern Gawler Craton, the metamorphic regime at ca. 1600–1580 Ma is well defined (e.g. Szpunar et al. 2007; Cutts et al. 2011; Forbes et al. 2011, 2012;

Morrissey et al. 2014), however the deformational regime at this time is not.

Deformation within the Mount Woods Inlier in the northeastern Gawler Craton (Fig. 1b) may have resulted in development of isoclinal folds during the Kimban Orogeny (Betts et al. 2003), or leading up to/synchronous with emplacement of the ca. 1584 Ma Balta Granite Suite (O'Sullivan 2010). Subsequent deformation involved the uplift and exhumation of the Mount Woods Inlier along the Southern Overthrust, which was initiated at ca. 1592–1582 Ma (Forbes et al., 2012).

Although the timing of the early isoclinal folding event in the Mount Woods Inlier is conjectural, it shares similarities to the early deformation history observed in the Yorke Peninsula. However, later generation of open upright folds similar to that observed in the Yorke Peninsula, has not been recognised in the Mount Woods Inlier. Overall, the lack of constraint on the Proterozoic deformational regime of the Gawler Craton, makes correlation with the deformation history of the Yorke Peninsula difficult.

The geological history of the Curnamona Province (Fig, 1b) at 1600–1580 Ma is better constrained, and shows a similar history to the Yorke Peninsula. Early isoclinal, recumbent (nappe) folds are interpreted to have developed during northwest–directed thrusting. The early folds were overprinted by northeast trending, open, upright folds during upper amphibolite to granulite–facies metamorphism as a result of northwest–southeast directed shortening (Marjoribanks et al., 1980; Hobbs et al., 1984; Forbes et al., 2004, 2005, 2007; Page et al., 2005). In the northern Broken Hill Inlier, overprinting of the fold generations lead to the development of Type 2 and Type 3 fold interference patterns (Forbes & Betts 2004; Forbes et al., 2004). The generation of open upright folds is also responsible for the dominant north–south structural grain throughout the Broken Hill Block (Forbes et al., 2004). Granites intruded the Curnamona Province

synchronously with and immediately following the deformation events, and include the ca. 1600–1570 Ma Ninnerie Supersuite (Page et al. 2005; Wade 2011), which was emplaced during the generation of the later upright folds (Gibson et al., 2004; Page et al. 2005). Granites of the Ninnerie Supersuite is interpreted to be equivalents of the 1595–1575 Ma Hiltaba Suite (Hand et al., 2008; Conor et al., 2010; Conor 2016).

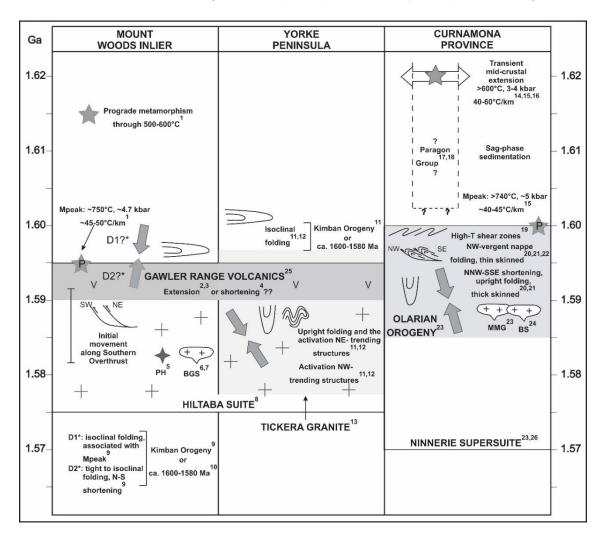


Figure 12. 1620–1570 Ma cladogram of selected Proterozoic terranes in South Australia, modified from Forbes et al. (2012). This includes the Mount Woods Inlier in the northern Gawler Craton, Yorke Peninsula in the southeastern Gawler Craton, and the Curnamona Province. Mineralisation: PH: Prominent Hill. Intrusives: BGS: Balta Granite Suite; MMG: Mundi Mundi Granites; BS: Bimbowrie Suite. References: Forbes et al. (2011); 2. Hitzman et al. (1992); 3. Betts et al. (2009); 4. Direen and Lyons (2007); 5. Belperio et al. (2007); 6. Fanning (1993); 7. Finlay (1993); 8. Creaser and Cooper (1993); 9. Betts et al. (2003); 10. O'Sullivan (2010); 11. Conor et al. (2010); 12. Hand et al. (2007); 13. Conor (1995); 14. Forbes et al. (2008); 15. Forbes et al. (2005); 16. Forbes et al. (2007); 17. Raetz et al. (2002); 18. Page et al. (2000); 19. Forbes et al. (2012); 20. Forbes et al. (2004); 21. Forbes and Betts (2004); 22. Gibson et al. (2004); 23. Page et al. (2005); 24. Ludwig and Cooper (1984); 25. Fanning et al. (1998); 26. Wade (2011).

Overall, the timing of at least one deformation event, the general orientation of structures and the timing of magmatism in the Yorke Peninsula and the Curnamona Province are similar. Both regions preserve evidence of two deformation events which have generally been constrained to similar times, and resulted in the development of a north–northeast trending structural grain within each region. In addition, if the generation of early isoclinal folds in the Yorke Peninsula occurred just prior to granite intrusion, these regions would share an even closer deformation history. Constraint on the timing of the earlier isoclinal folding in the Yorke Peninsula is needed to unequivocally determine the extent in similarity between the two regions.

CONCLUSIONS

The Tickera Granite is part of the broader Hiltaba Suite and comprises three phases; an early intensely foliated orange granite that was intruded by a white leucogranite and a later red granite. These localised phases differ compositionally, geochemically and isotopically, suggesting a heterogeneous source region. The white leucogranite originated from a similar but slightly more mafic source. Derivation of the Tickera Granite is likely from the Donington Suite and/or Wallaroo Group, with slight contamination from the underlying Archean basement.

The northeast–trending structures preserved at Point Riley are interpreted to be a composite fabric as a result of early isoclinal folds that were overprinted by open, upright folds. Early isoclinal folding is suggested to have occurred during the Kimban Orogeny or leading up to emplacement of the Tickera Granite at ca. 1597–1577 Ma. These northeast–trending structures are correlated with a regional northeast–trending fabric that defines the dominant structural grain across the Yorke Peninsula.

The Yorke Peninsula shares a similar deformational and magmatic history to the ca. 1600–1585 Ma Olarian Orogeny in the Curnamona Province, whereby each region preserves evidence of two deformation events, a dominant northeast trending structural grain and spatially and temporally related granite intrusions. Further constraint on the timing of early isoclinal folding in the Yorke Peninsula is required to determine the extent of similarities between the regions geological histories.

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