Intrusion related gold deposits in the Tanami and Kurundi-Kurinelli goldfields, Northern Territory, Australia: Constraints from LA-ICPMS analysis of fluid inclusions

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ABSTRACT

Gold deposits in the Tanami and Kurundi-Kurinelli goldfields of the Northern Territory, Australia, have geological and geochemical characteristics that distinguish them from the typical vein-hosted, orogenic gold deposits that occur in other parts of Australia. This study used cathodoluminescence (CL) and laser ablation inductively coupled plasma mass spectrometry (LA-ICPMS) analyses of fluid inclusions to investigate the source of mineralising fluids from both the highly mineralised Tanami goldfield and weakly mineralised Kurundi-Kurinelli goldfields in northern Australia. Cathodoluminescence showed three generations of quartz in fractured and mineralised veins from the Groundrush deposit in the Tanami goldfield but the remaining deposits had uniform dark CL images.

The LA-ICPMS trace element data indicate an evolution from B-As–enriched, magmatic fluid to a highly Sr-Ba–enriched fluid which results from interaction with sedimentary rocks. The fluids from both regions have chlorinity normalised Zn and Pb concentrations at the magmatic end of the metamorphic–magmatic mixing trend for these elements. This is in agreement with previous studies showing that gold mineralisation is coeval with magmatism in the Tanami region. The relatively high salinity of the fluids in both the Tanami and Kurundi–Kurinelli goldfields, and their trace element contents indicates that the deposits in both regions are intrusion-related gold deposits. The higher gold endowment of the Tanami region may result from the multiple gold-mineralisation events in this region whereas fluid flow associated with mineralisation in the Kurundi–Kurinelli goldfields appears to be of limited extent.

1. Introduction

The distinction between intrusion-related gold deposits (IRGD) and orogenic gold deposits (OGD) remains a subject of debate (e.g. Groves et al., 2003; Hart and Goldfarb, 2005). Australia has a range of gold deposits that may be classified as either orogenic-gold or intrusion-related gold deposits (Mernagh et al., 2007). OGDs typically form along convergent margins during the late part of the deformational-metamorphic history of terrane accretion (Groves et al., 2003). These deposits have a strong structural control involving faults or shear zones, folds, and/or zones of competency contrast. Host rocks are extremely variable, although there is an overall trend from volcanic-or intrusion-hosted deposits in Archean provinces to sedimentary rock-hosted deposits in Phanerozoic provinces. Although the host rocks vary from subgeenschist to granulate facies, the majority of OGDs are hosted in greenschist facies rocks. Quartz ± carbonate veins are ubiquitous and the deposits may extend to depths of up to 1 km or more, with only subtle metal zoning, and strong, lateral zonation of wall-rock alteration. A distinctive metal association (Au-Ag ± As ± B ± Bi ± Sb ± Te ± W) is also characteristic of OGDs (McCuaig and Kerrich, 1998; Ridley and Diamond, 2000; Goldfarb et al., 2005).

IRGDs may be classified into two broad groups. The first group occur in copper poor or absent portions of typical porphyry copper systems, associated with oxidised magmatic complexes. The second group occur in base metal poor or absent portions of intermediate to reduced magmatic complexes and are typically associated with Au-Bi-telluride or Au-As-Mo-Sb systems. This second type bears many similarities to OGDs and is what is referred to as an IRGD in this paper.

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IRGDs typically occur in a continental tectonic setting well inboard of inferred or recognized convergent plate boundaries, in a landward or back-arc position, or within continental collisional settings (Thompson and Newberry, 2000). Intrusion-related mineralisation tends to be younger than the pervasive fabric of the host rocks, whereas, orogenic mineralisation is synchronous with, or postdates pervasive fabrics and structures. IRGDs also contain a metal suite that is very similar to OGDS which sometimes also includes distal Pb-Zn. However, Blevin (2005) has stated that the base metal contents of IRGDs are highly variable and may be almost absent. The Au-rich portion of granitoid related mineral deposits may be locally distant from the higher temperature parts of the hydrothermal system, and obvious genetic links between the Au-rich portions of IRGD deposits and their related granitoids may be difficult to establish (Blevin, 2004).

IRGD deposits are related genetically to felsic igneous rocks (granite/rhyolite) of intermediate oxidation state (magnetite contents may be low or absent). The deposits exhibit a range of characteristics that vary over a wide range of emplacement depths (< 1 km to > 7 km). Deposits in shallow crustal settings (< 5 km) are associated with stocks, sills, dykes and volcanic domes and include systems with epi-thermal-style veins to breccia and stockwork similar to porphyry-type settings. Deeper systems (> 5 km) have characteristics of mesothermal environments, and are hosted by plutons containing sheeted veins, greisen and disseminated gold.

Many researchers favour a metamorphic fluid source for OGDS (e.g. Phillips, 1993; Goldfarb et al., 2005; Ridley and Diamond, 2000; Goldfarb and Groves, 2015; Phillips and Powell, 2010; Beaudoin and Chariaud, 2016; Fusswinkel et al., 2017). These deposits are typically associated with aprotic-carbonic fluids having low to intermediate salinity (3–13 wt% NaCl equiv.), XCO2 ≈ 0.1–0.15, δ18O = 6–13‰, and P-temperatures (~250–350 °C) and porphyry-type settings. Deeper systems (> 5 km) have characteristics of mesothermal environments, and are hosted by plutons containing sheeted veins, greisen and disseminated gold.

Fortunately, the differences between magmatic and metamorphic fluids can be determined by laser ablation inductively coupled plasma mass spectrometry (LA-ICPMS) analysis. There have been several LA-ICPMS studies of magmatic fluid inclusions (e.g., Audétat et al., 1998, 2008; Ulrich et al., 1999, 2002; Rusk et al., 2004; Seo et al., 2012; Harlaux et al., 2017). At present most of the data for magmatic fluids are from porphyry-epithermal systems and a few are from Li-enriched, evolved granite intrusions. Detailed studies of fluid inclusions associated with magmatic fluids are now also being reported (e.g., Klemm et al., 2004; Marsala et al., 2013; Rauchenstein-Martinek et al., 2014; Fusswinkel et al., 2017).

Fusswinkel et al. (2017) reported LA-ICPMS data from fluid inclusions from the Pampalo OGD, located in the Neoarchean Hattu schist belt in Eastern Finland and demonstrated that it was possible to obtain reliable LA-ICPMS data from magmatic-igneous rocks. The Pampalo OGD, located in the Neoarchean Hattu schist belt in Eastern Finland, is a small deposit (<1 km to >7 km) that has been divided into two main groups (Ahmad et al., 2013): Ware Group and Tanami Group (Fig. 2). Voluminous granitoids, dominated by met-aluminous, I type biotite granodiorite and monzogranite, outcrop over more than half of the Tanami Region. These granites intruded at 1825–1790 Ma and have been subdivided on geological criteria into the Birthday, Frederick and Grimwade suites (Fig. 2). At present most of the data for magmatic fluids are from porphyry-epithermal systems and a few are from Li-enriched, evolved granite intrusions. Detailed studies of fluid inclusions associated with magmatic fluids are now also being reported (e.g., Klemm et al., 2004; Marsala et al., 2013; Rauchenstein-Martinek et al., 2014; Fusswinkel et al., 2017).

The Tanami goldfield is located 80 km northwest of The Granites goldfield, contains four significant deposits: Bullakitchie, Bunkers Hill, Shoe and the Callie mine. The Callie mine, planed at the Dead Bullock Soak (DBS) goldfield. Callie is the largest deposit in this goldfield with a resource of 7.6 Moz gold (Boggott et al., 2016). Other mines in this goldfield include Gahn, Villa, Triumph-Colliwobble and DBR. At the Callie mine, sheeted, planed at weakly boudinaged quartz veins up to 20 cm thick host free gold with variable carbonate, chloride, biotite, feldspar, and sulfide minerals. These veins dip 20° to 63° and define a series of E-W-trending corridors. Ore veins are generally surrounded by a narrow zone of wall rock alteration that varies with depth, with chloride being progressively replaced by biotite as the dominant phyllosilicate (Huston et al., 2007).

The Granites goldfield, located 40 km to the east of the DBS goldfield, contains four significant deposits: Bullahkutchie, Bunkers Hill, Shoe and Quorn. In the Granites goldfield, besides the quartz-only veins, gold also occurs in chloride bands within quartz–chlorite veins and as grains associated with sulfide and arsenide minerals disseminated in banded-iron formation (BIF). The highest gold grades are associated with sulfide-rich amphibole schist and with carbonate-rich facies BIF (Huston et al., 2007).

The Tanami goldfield is located 80 km northwest of The Granites goldfield. More than 40 deposits are located within a 15 km long, northeast-trending zone locally known as the Tanami Mine Corridor. Mineralisation in the Tanami goldfield is hosted by an alternating succession of basalt and mudstone belonging to the Mount Charles Formation (Fig. 2). Gold occurs in quartz-carbonate veins associated with sericite-pyrite alteration. The ore shoots range from 1 to 20 m thick, 20 to 300 m in length and 10 to 70 m vertical extent. Higher gold
grades occur adjacent to ore shoot boundaries. The main gangue minerals are calcite, siderite and quartz and pyrite is the main sulphide, comprising 5–20% of vein composition.

The Groundrush gold deposit is located about 40 km northeast of the Tanami goldfield and lies near the Frankenia granitic ‘dome’. The deposit sits within a 90–160 m thick dolerite sill intruding meta-greywacke of the Ferdies Member, Dead Bullock Formation (Fig. 2). Mineralised lodes are 10–30 m wide and consist of anastomosing zones of northeast-trending quartz veins dipping 70–80° and striking west. Gold occurs as free metal, or is associated with arsenopyrite and to a lesser extent, with pyrite and pyrrhotite.

Significant deposits in the Western Australian part of the Tanami Region include Coyote, and Kookaburra (Bagas et al., 2014; Ahmad et al., 2013). At least three sets of quartz veins are recognised at the Coyote deposit but only the first two sets are mineralised. Localised alteration assemblages accompanying these vein sets include 1) quartz–sericite–pyrite–chlorite, silica–dolomite spotting and veining, 2) uncommon quartz–K-feldspar, graphite, and 3) chlorite–biotite. Only the chlorite–biotite assemblage can be related to the auriferous veins, although some may be related to late carbonate-bearing veinlets (Bagas et al., 2009). At Kookaburra, gold is associated with quartz–carbonate ± pyrite ± arsenopyrite veins with variable amounts of sericite, K-feldspar and marcasite, and minor to trace quantities of biotite, amphibole, apatite, sphalerite, and pyrrhotite. The Buccaneer deposit was the first intrusion-hosted gold system discovered in the Tanami region (Bagas et al., 2009). It is hosted in the southern part of the ca 1795 Ma Twin Bonanza Porphyry, which also hosts the Cypress gold prospect in the north.

2.2. Warramunga Province

The Proterozoic Warramunga Province forms the central part of the Tennant Region, north and northwest of the Davenport Province and south of the Tomkinson Province (Donnellan, 2013; Fig. 3). To the east and west, the province extends subsurface beneath the Phanerozoic Georgina and Wiso basins, respectively. The oldest rocks of the Warramunga Province predate the ca 1850 Ma Tennant Event, and are divided into the Warramunga Formation, and the correlative Junalki Formation and Woodenjerrie beds. These rocks are overlain in the south by the Hatches Creek Group in the Davenport Province, but in the north, they are overlain by the Tomkinson Creek Group in the Tomkinson Province (not shown in Fig. 3).

Geochronological data indicate that there were four episodes of felsic volcanism in the Warramunga Province. These span the time interval between Junalki Formation volcanism (ca 1860 Ma) and the end of Ooradidgee Group time (ca 1810 Ma). Hoatson et al. (2007) assigned mafic magmatism in the Warramunga Province to three magmatic events. These are the ca 1850 Ma Mumbilla Event, the 1840–1820 Ma Edmirringe Event and the 1820–1800 Ma Mount Hay Event. The Mount Hay Event includes gabbro and dolerite sills that intrude the Kurinelli area of the southern Warramunga Province.

Compared to the Tanami region, the Kurundi – Kurinelli goldfields in the Warramunga Province are rather poorly endowed. Official production records indicate that only 25 kg of gold was produced from this region between 1926 and 1995 (Ahmad et al., 2009). Gold is present in quartz veins occupying bedding parallel faults or shears within sediment – dolerite contact zones, or within sedimentary and volcanic rocks of the Ooradidgee Subgroup. In the Kurundi goldfield (Fig. 3), gold is present in bedding-parallel quartz veins, which are 1–3 m wide and up to 200 m long. These veins are hosted in a variety of lithologies including quartz sandstone and shale of the Kurinelli Sandstone (Great Davenport, Power of Wealth, Aztec and Priesters); acid volcanics of the Epenarra Volcanics (Millars) and the Junalki Formation (Opengidgi); basalt of the Edmirringe Volcanics (Kurundi); sandstones of the Taragan Sandstone (Cairns); granophyre and quartz-feldspar porphyry (Davidsons). In the Kurinelli goldfield (Fig. 3) the gold is generally located in northeasterly-trending, quartz-filled shear zones within dolerite/gabbro and sandstone (Rooneys Formation), or at the dolerite – sediment contact. Quartz veins, emplaced subparallel to bedding, are 0.2 – 2 m wide and can be traced for up to 200 m along strike. Mineralisation is erratic and occurs as fine- to coarse-grained native gold, in places associated with chalcopyrite and pyrite.

![Fig. 1. Map of the geological regions in the centre of the Northern Territory, Australia. The yellow dots depict gold mines and prospects in the Tanami goldfield and the Kurundi-Kurinelli goldfields, respectively. The red lines show the major highways in this region. All data were obtained from the Spatial Territory Resource Information Kit for Exploration (STRIKE) web application provided by the Northern Territory Geological Survey. The light red boxes indicate the locations within the Northern Territory of Figs. 2 and 3. (For interpretation of the references to colour in this figure legend, the reader is referred to the web version of this article.)](image-url)
3. Previous fluid inclusion studies

3.1. Fluid inclusions in the Tanami region

Mernagh and Wygralak (2007) have carried out fluid inclusion studies on the major gold deposits and prospects in the Tanami Region. Six main types of inclusions, based on their phase proportions at room temperature, were identified in this study and are listed below:

Type A. Primary, CO₂-rich inclusions. They are very abundant and contain a CO₂-rich liquid phase or CO₂-rich liquid + vapour phases. They may also contain small amounts of CH₄ or N₂ and a minor aqueous phase.

Type B. Primary, two-phase, CH₄ (± CO₂) + H₂O inclusions. These inclusions are not as abundant and contain 10–30 vol% CH₄-rich vapour.

Type C. Primary, monophase, CH₄ ± N₂ ± CO₂ vapour inclusions. These inclusions are quite rare and can be distinguished from Type A inclusions by their CH₄- or N₂-rich nature.

Fig. 2. Bedrock geology and gold mines and gold occurrences of the Tanami region (from Ahmad et al., 2013). The yellow dots depict gold occurrences. (For interpretation of the references to colour in this figure legend, the reader is referred to the web version of this article.)
Type D. Two-phase, aqueous inclusions. They contain 5–15 vol% vapor, and are abundant in all stages of veining and may be primary or secondary.

Type E. Rare, secondary, aqueous inclusions with one or more daughter minerals. They typically contain halite and may also contain a vapour bubble. No gases were detected by Raman spectroscopy in the vapour bubble.

Type F. Secondary, two-phase brine inclusions. These are generally large, irregular, CaCl₂-bearing inclusions with 5–15 vol% vapour. Once again, no gases were detected by Raman spectroscopy in the vapour bubble.

Only the primary fluid inclusions associated with mineralisation will be considered and discussed in this paper.

The ore bearing quartz veins in the Tanami region typically contain coexisting Type A and D inclusions (Table 1). Microthermometric data indicate that the gold deposits formed over a range of depths and at homogenisation temperatures from 184 to 430 °C. No pressure corrections were applied to these temperatures. The Groundrush deposit formed at the greatest temperatures and depths (260–430 °C and ≤11 km), whereas deposits in the Tanami goldfield formed at the lowest temperatures (184–359 °C) and at the shallowest depths (1.5–5.6 km). Other deposits (e.g., The Granites, Callie, and Coyote) formed at depths between those of the Tanami goldfield and the Groundrush deposit and at temperatures ranging from 240 to 360 °C. Furthermore, the more deeply formed deposits were found to be enriched in CH₄ (with up to 85 mol.% CH₄ in the vapour phase) and higher level deposits were enriched in CO₂ (85–100 mol.% CO₂ in the vapour phase). Fluids from deposits hosted mainly by sedimentary rocks generally also contained appreciable quantities of N₂ (≤100 mol.% in the vapour phase). Calculated δ¹⁸O data for the ore fluids indicated a magmatic or mixed magmatic/metamorphic source for the ore fluids in the Tanami Region (Mernagh and Wygralak, 2007).

Interpretation of the fluid inclusion, alteration, and structural data suggests that mineralisation may have occurred via a number of processes (Mernagh and Wygralak, 2007). Gold occurs in veins associated with brittle fracturing and other dilational structures, but in the larger deposits, there is also an association with iron-rich rocks or carbonaceous sediments, suggesting that both structural and chemical controls are important. The major mineralisation process appears to be fluid immiscibility (i.e. boiling/effervescence) as indicated by the coexistence of vapour-rich (Type A) and liquid-rich (Type D) inclusions in fluid inclusion assemblages (FIAs) in the veins (Mernagh and Wygralak, 2007). However, the presence of pyritic alteration haloes around the quartz veins in some deposits may indicate that desulphidation by fluid–rock interaction may also be a significant ore deposition mechanism for these deposits. The pyritic alteration is generally more prevalent in the higher crustal-level deposits.

3.2. Fluid inclusions in the Warramunga Province

Wygralak and Mernagh (2008) conducted fluid inclusion studies on the major gold deposits and prospects in the Warramunga Province. They recognised two types of fluid inclusions, Types A and D, in this region (Table 1). The Type A, CO₂-rich fluid inclusions have salinities < 7 wt% NaCl eq. while the coexisting Type D, aqueous inclusions have salinities up to 10 wt% NaCl eq. These FIAs homogenised to liquid over the temperature range 400–440 °C and by analogy with other OGDs (Ridley and Diamond, 2000), these inclusions are thought to represent the main mineralising fluid in this region. The second type of FIAs contained only Type D, two-phase, aqueous inclusions which homogenised to liquid over a broad temperature range of 140–340 °C. This fluid has high salinity ranging from 18 to 22 wt% NaCl eq. and the lack of gas-rich inclusions in these FIAs indicate that they may have resulted from the trapping of a later basinal brine.

4. Analytical methods

4.1. Fluid inclusion petrography

This study uses the same samples from high-grade ore zones as previously reported in Mernagh and Wygralak (2007) and Wygralak.
terials used for analysis are listed in Table 2. In order to avoid
and Mernagh (2008). These previous studies have carefully determined
premature fracturing of the host quartz in this study, a stepwise
drilling procedure was used. Firstly the host quartz were pre-drilled at
lower repetition rate and small spot size, and then with relative high
replication rate and big spot size (depending on the fluid inclusion size)
to open and ablate the entire inclusion, followed by another ablation
with bigger spot size to obtain the trace element concentration in the
quartz matrix. Generally, a hole of 10 µm in diameter was pre-drilled
with a 4 Hz repetition rate in the quartz. Then a hole of 16 µm was
drilled in the same spot with a 4 Hz repetition rate, followed by hole of
24 µm with a 10 Hz repetition rate to open and ablate the whole
inclusions. Then to obtain the trace element concentration in the quartz
matrix, this was followed by another hole of 32 µm with a 10 Hz re-
petition rate. A comparison of this new method of determining the
composition of the quartz matrix with conventional methods shows
agreement for all elements is better than 20% (for Na, K, Ca and Mg
the differences are < 5%), which is acceptable for fluid inclusion
analyses. This method has the advantage of providing a more accurate
composition of the quartz matrix around the inclusion, whereas se-
parate ablations carried out before or after fluid inclusion analysis are
prone to contamination issues.

The adoption of this stepwise laser drilling procedure greatly in-
creased the number of successful ablations with a quantifiable ICPMS
signal. As reported by Fusswinkel et al. (2017), the most important
criterion was a stable and steady ablation of the host quartz for several
seconds before and after inclusion breach, without fracture formation
(leaseage of fluid) at the bottom of the ablation pit or explosive opening
of inclusions (spallation), because both processes may lead to in-
complete sampling of inclusion contents. Fracturing of quartz can result
in partial fluid leakage into the sample along newly formed crack sur-
faces extending out from the ablation pit, which results in very long,
drawn out signals with low intensity. Spallation usually results in un-
controlled opening of the inclusion cavity and explosive fluid release,
which may not ensure complete vaporisation of inclusion contents
(Table 5).

The SILLS software package (Guillong et al., 2008) was used for LA
ICPMS data reduction. All elements were quantified against NIST
SRM610 as the external standard. Absolute concentration values were
then calculated by internal standardisation against Na concentrations
determined by microthermometry. Most fluid types contained sig-
nificant amounts of other cations as well, so that the NaCl equivalent
salinity (NaCl eq.) is not identical to the true fluid salinity. However,
true Na concentration for internal standardisation can be reasonably

### Table 1


<table>
<thead>
<tr>
<th>Deposit</th>
<th>Inclusion Type</th>
<th>XCO2 (vap)</th>
<th>XN2 (vap)</th>
<th>XCH4 (vap)</th>
<th>Wt% NaCl equiv</th>
<th>T (°C)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Tanami Goldfield</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Callie A</td>
<td>D</td>
<td>0.0–1.0 (0.81)</td>
<td>0.0–0.76 (0.17)</td>
<td>0.0–0.12 (0.02)</td>
<td>3.8–7.3 (6.9)</td>
<td>254 to 326 (311)</td>
</tr>
<tr>
<td>Callie D</td>
<td>ND</td>
<td>ND</td>
<td>ND</td>
<td>ND</td>
<td>5.1–20.0 (14.6)</td>
<td>290 to 404 (274)</td>
</tr>
<tr>
<td>Groundrush B</td>
<td>0.15–0.87 (0.45)</td>
<td>0.0–0.27 (0.03)</td>
<td>0.13–0.85 (0.51)</td>
<td>1.2–13.9 (7.5)</td>
<td>213 to 490 (346)</td>
<td></td>
</tr>
<tr>
<td>Groundrush D</td>
<td>ND</td>
<td>ND</td>
<td>ND</td>
<td>ND</td>
<td>0.2–10.4 (6.9)</td>
<td>161 to 277 (207)</td>
</tr>
<tr>
<td>Coyote A</td>
<td>0.73–1.0 (0.79)</td>
<td>0.0–0.09 (0.06)</td>
<td>0.0–0.20 (0.07)</td>
<td>0.0–6.7 (2.1)</td>
<td>185 to 408 (308)</td>
<td></td>
</tr>
<tr>
<td>Coyote B</td>
<td>0.0–0.18 (0.02)</td>
<td>0.0–0.38 (0.08)</td>
<td>0.62–1.0 (0.90)</td>
<td>0.0–12.5 (2.7)</td>
<td>245 to 389 (325)</td>
<td></td>
</tr>
<tr>
<td>Coyote C</td>
<td>0.47–0.59 (0.5)</td>
<td>0.3–0.63 (0.39)</td>
<td>0.0–0.24 (0.11)</td>
<td>–</td>
<td>312 to 417 (359)</td>
<td></td>
</tr>
<tr>
<td>Coyote D</td>
<td>ND</td>
<td>ND</td>
<td>ND</td>
<td>1.1–11.8 (5.5)</td>
<td>183 to 434 (250)</td>
<td></td>
</tr>
<tr>
<td>Kurundi – Kurinelli goldfield</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Dempseys Choice A</td>
<td>ND</td>
<td>0.15–1.0 (0.67)</td>
<td>0.0–0.41 (0.10)</td>
<td>0.0–0.22 (0.02)</td>
<td>–</td>
<td>360 to 445 (399)</td>
</tr>
<tr>
<td>Dempseys Choice D</td>
<td>ND</td>
<td>ND</td>
<td>ND</td>
<td>2.0 to 22.1 (7.8)</td>
<td>180 to 313(243)</td>
<td></td>
</tr>
<tr>
<td>Kurinelli A</td>
<td>0.29–0.97 (0.86)</td>
<td>0.0–0.13 (0.08)</td>
<td>0.0–0.29 (0.06)</td>
<td>–</td>
<td>366 to 419(406)</td>
<td></td>
</tr>
<tr>
<td>Kurinelli D</td>
<td>ND</td>
<td>ND</td>
<td>ND</td>
<td>16.9 to 19.4 (18.0)</td>
<td>109 to 275(178)</td>
<td></td>
</tr>
<tr>
<td>Power of Wealth A</td>
<td>0.45–1.0 (0.91)</td>
<td>0.0–0.36 (0.08)</td>
<td>0.0–0.16 (0.03)</td>
<td>–</td>
<td>205 to 357(340)</td>
<td></td>
</tr>
<tr>
<td>Power of Wealth D</td>
<td>ND</td>
<td>ND</td>
<td>ND</td>
<td>1.7 to 11.5 (9.2)</td>
<td>130 to 428(323)</td>
<td></td>
</tr>
<tr>
<td>Kurundi A</td>
<td>0.64–0.9 (0.84)</td>
<td>0.0–0.08 (0.05)</td>
<td>0.08–0.22 (0.11)</td>
<td>–</td>
<td>154 to 418(335)</td>
<td></td>
</tr>
<tr>
<td>Kurundi D</td>
<td>ND</td>
<td>ND</td>
<td>ND</td>
<td>1.7 to 11.1 (8.8)</td>
<td>186 to 299 (232)</td>
<td></td>
</tr>
</tbody>
</table>

a See text for definitions.
b Mole fraction of CO2, N2 and CH4 in the vapour phase as determined by laser Raman microprobe analysis.
c The quoted figures represent the range of the data and the average value is given in parentheses.

### 4.2. Fluid inclusion microthermometry

As discussed above, extensive microthermometry has already been
done on these samples (Mernagh and Wygralak, 2007; Wygralak and
Mernagh, 2008). Additional fluid inclusion studies were carried out on
the samples selected for LA-ICPMS analysis to obtain accurate salinities of the selected FIAs for the standardisation of the LA-ICPMS data. The
new data is reported in the Electronic Supplementary Material. Mi-
crothermometry was carried out on a Linkam MDS600 heating/
cooling stage which was calibrated with synthetic H2O and H2O-CO2
fluid inclusion standards. A heating rate of 2 °C/min was used up to
30 °C and then a heating rate of 5 °C/min was used at higher tem-
peratures. After calibration the stage was accurate to ± 0.2 °C below
30 °C and ± 2.0 °C above 30 °C. All phase transitions were carefully
measured by taking photos at approximately one second intervals to
accurately determine the temperature of the phase transitions.

### 4.3. Cathodoluminescence studies

CL images were obtained from a JEOL JSM-7800F Field Emission
Scanning Electron Microprobe with the MonoCL4 detector from Gatan
Inc. and a 10 k eV beam.

### 4.4. LA-ICPMS analysis of the fluid inclusions

The analyses were conducted at the State Key Laboratory of Ore
Deposit Geochemistry, IGCAS by using the Agilent 7900 ICP-MS,
which is combined with a GeoLasPro 193 nm ArF excimer laser and
with synthetic fluid inclusions to monitor the accuracy (Lan et al.,
2017a,b). The instruments, analytical conditions and reference ma-
terials used for analysis are listed in Table 2. In order to avoid
well approximated by charge balancing using cation/chlorine ratios from LA-ICPMS signals, as described by Allan et al. (2005). Comparisons with synthetic inclusions (Allan et al., 2005; Zhou et al., 2016) show that this method produces acceptable results. Limits of detection were calculated using Eq. (6) in Pettke et al. (2012), which yields concentration of an element at the 95% confidence level. The method is implemented in the SILLS software package.

5. Results

5.1. Cathodoluminescence studies

In this study we have used cathodoluminescence (CL) to better understand the timing of fluid inclusion trapping in the samples that were selected for LA-ICPMS analysis. Fig. 6 shows the extra information.

Table 2
Summary of instruments, analytical conditions and reference materials used for the LA-ICPMS measurements.

<table>
<thead>
<tr>
<th>Laser ablation system</th>
<th>Instrument GeoLasPro 193 nm</th>
</tr>
</thead>
<tbody>
<tr>
<td>Energy Density (J/cm²)</td>
<td>10 10 10 10</td>
</tr>
<tr>
<td>Spot Size (µm)</td>
<td>10 16 24 32</td>
</tr>
<tr>
<td>Repetition rate (Hz)</td>
<td>4 4 24 32</td>
</tr>
<tr>
<td>Cell gas flow (L/min, He)</td>
<td>0.6</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>ICPMS system</th>
<th>Instrument Agilent 7900</th>
</tr>
</thead>
<tbody>
<tr>
<td>RF power (W)</td>
<td>1450</td>
</tr>
<tr>
<td>Cooling gas flow (L/min, Ar)</td>
<td>15</td>
</tr>
<tr>
<td>Auxiliary gas flow (ml/min, N₂)</td>
<td>3</td>
</tr>
<tr>
<td>Makeup gas flow (L/min, Ar)</td>
<td>1.0</td>
</tr>
<tr>
<td>Integration/Dwell time (ms)</td>
<td>10 for all elements except that 20 was used for ³⁵Cl, ³⁹K, ⁸¹Br, ¹⁰⁷Ag and 60 was used for ¹⁹⁷Au</td>
</tr>
</tbody>
</table>

Monitored isotopes

| ³⁵Cl, ³⁹K, ⁸¹Br, ¹⁰⁷Ag, ¹²³Th, ¹³³Cs, ¹³⁷Cs, ¹³⁷Ba, ¹⁸¹Ta, ¹⁸⁵W, ¹⁹⁷Au, ²⁰⁸pb, ²⁰⁹Bi, ²³⁵Th, ²⁵⁴U |

Reference material NIST SRM610
17.7 to 18.2 wt% NaCl eq. but as only a few inclusions from each zone slightly lower than inclusions in the later Q2 quartz, which range from the early Q1 quartz have salinities of 17.5–17.7 wt% NaCl eq. which is consistent. The data show that the inclusions studied by microthermometry in the Q1 and Q2 quartz in this sample prior to LA-ICPMS analyse. The pits shown in Fig. 6(b), the dark grey quartz (Q1) is the earliest phase which appears to have been fractured and infilled with the light grey quartz (Q2). This latter quartz also has alternating dark and light bands at approximately 45° which may represent growth banding. Finally, the whole sample is cut by a large number of thin, anastomosing veinlets of black quartz (Q3).

Microthermometry was carried out on selected inclusions from both the Q1 and Q2 quartz in this sample prior to LA-ICPMS analyse. The pits left after laser ablation are outlined with red circles in Fig. 6(b) and the salinity of these inclusions (in wt% NaCl eq.) is shown next to the circles. The data show that the inclusions studied by microthermometry in the early Q1 quartz have salinities of 17.5–17.7 wt% NaCl eq. which is slightly lower than inclusions in the later Q2 quartz, which range from 17.7 to 18.2 wt% NaCl eq. but as only a few inclusions from each zone were analysed, the result remains equivocal.

Unfortunately, it was not possible to distinguish different generations of quartz in the other deposits as they did not show any significant differences in the CL images. This is similar to quartz in orogenic Au deposits which typically generate a dull grey, homogenous CL response and mottled textures (Rusk, 2012).

### 5.2. Fluid inclusion LA-ICPMS

As discussed above we imposed strict quality principles to define successful ablations. Analyses were only accepted when we observed steady ablation of the host quartz for at least several seconds before and after opening the fluid inclusion (Fig. 7). The results from some fluid inclusions were rejected because of their small size or because they were located too close to the surface of the wafer, which makes them more susceptible to spallation. An example of a good LA-ICPMS analysis of a two-phase, aqueous inclusion is shown in Fig. 7(a). The stepwise opening of the laser hole is demonstrated by the alternating Si intensities. When the inclusion is opened, all element signals increase in synchronisation with the Na peak, which is used for internal standardisation. The composition of the host quartz is then determined by a subsequent ablation of the same hole with a 32 µm diameter laser beam. An example of a good LA-ICPMS analysis of a CO2-rich inclusion is shown in Fig. 7(b). Here, the same stepwise opening procedure is used but when the inclusion is breached only a small increase in the Na intensity is observed. All other elements are below detection limits. Therefore, only LA-ICPMS analyses of the two-phase aqueous inclusions are included in this study.

All the accepted LA-ICPMS data from the two-phase, aqueous inclusions show elevated concentrations of Na, K, As, B, Rb, Sr and Sb. The average element concentrations of successfully analysed FIAs for each ore deposit are plotted in Fig. 8. The full dataset of individual measurements is available as Electronic Supplementary Material. It can be seen from Fig. 8 that there are systematic differences between the different ore deposits, especially differences between the dolerite-hosted Groundrush deposit and the Callie and Coyote deposits in the Tanami goldfield. The Groundrush deposit has higher concentrations of Mg while the other deposits have a higher concentration of B, Ca, and Zn. In the Kurundi – Kurinelli goldfields, the Power of Wealth and Kurundi deposits have lower salinity, and hence, lower Na concentration than the other deposits. The Kurundi deposit also has high Fe, As and Sb concentrations. This may reflect the fact that the quartz veining is hosted by basalt of the Edmirringie Volcanics. The Kurinelli and Power of Wealth deposits have relative low concentrations of Mg compared to other deposits, which may indicate that the fluids have not interacted with mafic volcanics in these deposits.

The ratio of Na/K is of interest because of its potential as a geothermometer. If the fluids equilibrated with two feldspars, this ratio that can be obtained from the CL image of a quartz vein from the Groundrush deposit. In Fig. 6(a) the sample is viewed in ordinary transmitted light and contains a densely packed array of inclusions along with some trails of secondary inclusions but no other information on the relative timing of inclusion trapping is apparent in this image. However, the CL image is remarkably more detailed (Fig. 6(b)). As shown in Fig. 6(b), the dark grey quartz (Q1) is the earliest phase which appears to have been fractured and infilled with the light grey quartz (Q2). This latter quartz also has alternating dark and light bands at approximately 45° which may represent growth banding. Finally, the whole sample is cut by a large number of thin, anastomosing veinlets of black quartz (Q3).

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The ratio of Na/K is of interest because of its potential as a geothermometer. If the fluids equilibrated with two feldspars, this ratio...
should be a function of temperature, with lower ratios at higher temperature (Orville, 1963; Rusk et al., 2004). Most of the deposits have a fairly constant molar Na/K ratio, which ranges from 3.3 to 5.7. However, the Kurundi deposit has a much lower ratio of 1.6. The fluid inclusion data from the Kurundi deposit (Wygralak and Mernagh, 2008) indicate that this deposit actually formed at slightly lower temperatures (average \( T_h = 232 \) °C) than other deposits in this region (average \( T_h \) from 279 to 326 °C). Therefore, we conclude that the fluid in this deposit was not in equilibrium with the two feldspars. A comparison of the fluid composition across all the deposits shows, in general, that the fluids are richer in Na and Ca and have lower concentrations of K. Magnesium concentrations are generally much lower in the Kurundi – Kurinelli goldfields, perhaps reflecting a lower amount of mafic rocks in this region. Iron concentrations vary by several orders of magnitude and may reflect the different host rocks (i.e. mafic or arenaceous) of these deposits. Average Rb:Sr ratios for individual FIA vary considerably in each goldfield. In the Tanami goldfield the Rb:Sr ratios range from 0.073 to 0.500 while in the Kurundi – Kurinelli goldfields they range from 0.095 to 1.180.

Gold was consistently below the limits of detection (0.5 to 10.0 μg/g) for all fluid inclusions analysed in this study. Reliable data for Au concentrations in orogenic fluids are still very scarce. Garofalo et al. (2014) report concentrations of 0.5 μg/g to 5 μg/g for the Sigma gold deposit in the Abitibi Greenstone Belt, Canada, and Fusswinkel et al. (2017) report Au concentrations up to 0.6 μg/g for the Pampalo OGD in Finland.

Large et al. (2016) have shown that the sources of fluids can be inferred from their trace element composition and so we have compared the trace element ratios of the LA-ICPMS data from the two different goldfields in Fig. 9. Both goldfields show a similar spread in the range of Sr-B-Rb ratios which trend from the B-rich magmatic vapour field to the Sr-rich sedimentary influenced field (Fig. 9a and b). In comparison to the other deposits, samples from the Kurinelli deposit are enriched in Sr and plot mostly in the sedimentary influenced field. The
plot of Sr-As-Rb ratios for the Tamani goldfield (Fig. 9c) show a spread of values from B-rich, magmatic influenced fluid to the Sr-rich, sedimentary influenced fluid, with the Groundrush deposit appearing to span across both regions which may be due to the multiple phases of fluid flow. The data from the Coyote deposit, on the other hand, is mostly confined to the magmatic field. In the Kurundi – Kurinelli goldfields the trace element data from the Dempseys Choice and Power of Wealth deposits lie mostly within the magmatic field while the Kurinelli deposit, with its very high Sr ratios, has a strong sedimentary influence.

6. Discussion

6.1. Comparisons with other gold deposits

Large et al. (2016) have used data from a number of hydrothermal ore deposits to show that trace element ratios can be used to distinguish fluid end members. We have compared our data with their results in Fig. 9. Our results are similar to the study of the Carlin fluids (Large et al., 2016) and indicate a progression from B-As–enriched (magmatic) fluid to a highly Sr-Ba-enriched fluid as a result of interaction with sedimentary rocks. In the Tanami goldfield the Groundrush deposit appears to span across both regions which may be due to the multiple phases of fluid flow. The data from the Coyote deposit, on the other hand, is mostly confined to the magmatic field. In the Kurundi – Kurinelli goldfields the trace element data from the Dempseys Choice and Power of Wealth deposits lie mostly within the magmatic field while the Kurinelli deposit, with its very high Sr ratios, has a strong sedimentary influence.

Variations in the chemistry of the fluids at each deposit in this study (Fig. 8) may be due to either variations in the source fluids or fluid-rock interaction. The concentration of Mg appears to be sensitive to the presence of mafic rocks. The high Mg concentrations in fluids from the Groundrush deposit (2422 to 5227 µg/g) are in accord with the δ 18O data of the fluids (3.8 to 8.5‰) and Mg-rich biotite alteration in the Tanami region (Mernagh and Wygralak, 2007; Bagas et al., 2014) both of which suggest a magmatic or mixed magmatic/metamorphic source for the ore fluids (Mernagh and Wygralak, 2007). The Mg concentration in fluids from the Kurundi – Kurinelli goldfields are generally low with the exception of the Kurundi deposit which is hosted by basalt.

The concentrations of Fe, Zn and Cu in all deposits analysed in this study are considerably higher than those reported for the Pampalo orogenic gold deposit (Fusswinkel et al., 2017) and generally higher than metamorphic vein systems (Rauchenstein-Martinek et al., 2014) but the average value for Cu (80 ± 45 mg/g) is similar to that reported by Garofalo et al. (2014). This is probably due to the higher salinity of the fluid inclusions from the gold deposits in this study as these metals are dominantly complexed with chloride ions (Wood and Samson, 1998). Fig. 10 shows a compilation of the chlorinity normalised molar concentrations of Zn and Pb (i.e., mZn/mCl and mPb/mCl) for fluids from this study and also of magmatic-hydrothermal fluids and normal metamorphic fluids unrelated to orogenic gold deposits. It shows that the fluids in this study have Zn and Pb concentrations that lie towards the magmatic-hydrothermal end of the metamorphic – magmatic mixing trend. It is likely that the concentrations of Pb and Zn in the fluids was increased by the precipitation of sulfides such as pyrite, especially if the fluids initially have an excess of Fe over reduced S. The precipitation of sulphides results in a sulfur-deficient fluid, which can retain high concentrations of Pb and Zn down to temperatures as low as 200 °C (Heinrich et al., 2004). The concentration of Pb and Zn can be further enhanced by the increasing dissociation of the HCl0 species in response to cooling, which makes the fluid more acidic and capable of keeping Pb and Zn in solution (Tagirov et al., 1997).
few data are available at present, the generally higher Rb: Sr ratios obtained for deposits in this study indicate the involvement of magmatic fluids (Plimer and Elliott, 1979; Blevin and Chappell, 1995).

6.2. Intrusion related gold deposits in the Tanami and Kurundi-Kurinelli goldfields

Wall (2005) proposed that the gold mineralisation in the Tanami region was related to the thermal aureole around the granites. The author proposed that structurally-controlled plumbing systems channel the ore fluids into new or reactivated, pre-existing structures that formed at the time of granitoid emplacement. Tunks and Cooke (2007) carried out structural studies which indicated that gold veins in the Tanami district were part of an outer thermal aureole gold system that formed during the emplacement of the granitoids. Economic gold mineralisation was interpreted to occur late in the paragenetic history of the district. A summary of the geochronological data for the Tanami region is given in Table 3, which shows that the age of mineralisation is accepted as being approximately 1795 Ma (Bagas et al., 2014). The occurrence of intrusion related gold deposits (e.g. the Buccaneer and Cypress deposits) in the

Fig. 9. Comparison of the average trace element ratios from each deposit. (a) Sr-B-Rb ratios for the Tanami Goldfield, (b) Sr-B-Rb ratios for the Kurundi – Kurinelli goldfields, (c) Sr-As-Rb ratios for the Tanami Goldfield, (d) Sr-As-Rb ratios for the Kurundi – Kurinelli goldfields, (e) Sr-Cs-Rb ratios for the the Tanami Goldfield, and (f) Sr-Cs-Rb ratios for the Kurundi – Kurinelli goldfields. Hexagons show range for each fluid type (from Large et al. 2016) with intensity of shading emphasizing inferred end-member fluids, magmatic brine (blue), magmatic vapor (yellow), and sedimentary influenced (brown). (For interpretation of the references to colour in this figure legend, the reader is referred to the web version of this article.)
arsenic from Coyote’s quartz veins has a δ³⁴S value of –1.5‰ relative to the CDT and can be also interpreted in terms of pre-metamorphic fluids released from granitic intrusions coeval with orogenic gold deposition during late-stage compression in the Tanami region (Table 3). Bagas et al. (2008) have also suggested that mafic rocks in the Tanami region were deposited in a back-arc basin to island-arc setting, which is a favourable setting for IRGDs. The higher gold endowment of the Tanami deposits may be the result of multiple gold-mineralisation events in this region. Callie, the largest deposit in the Tanami region shows clear evidence of multiple gold-mineralisation events in this region (Wygralak and Mernagh, 2008) but the extent of fluid flow seems to be more limited as evidenced by the restricted extent of quartz vein ing (≤200 m). Therefore the lower endowment of the Kurundi – Kurinelli goldfields may be the result of a single mineralising event and/or the limited extent of fluid flow in this region.

7. Conclusions

This study has used cathodoluminescence (CL) and LA-ICPMS analyses of gold deposits in the Tanami and Kurundi-Kurinelli goldfields to
investigate the source of the fluids and the factors that may affect the gold endowment in the two regions. CL imaging has shown the presence of up to three generations of chemically distinct quartz in fractured and mineralised veins from the Groundrush deposit, whereas only vein quartz of uniform CL composition was observed in the other deposits.

The LA-ICPMS data from fluid inclusions from all the deposits show elevated concentrations of Na, K, As, B, Rb, Sr and Sb and indicate an evolution from B-As-enriched, magmatic fluid to a highly Sr-Ba-enriched fluid as a result of interaction with sedimentary rocks. Interaction with the sedimentary rocks appears to be more prevalent in the Kurundi – Kurinelli goldfields than in the Tanami goldfield. Rb: Sr ratios vary considerably but are generally higher than the ratios in orogenic gold deposits, and hence, indicate interaction with magmatic fluids.

The fluids from both regions in this study have chlorinity normalised Zn and Pb concentrations near the magmatic end of the metamorphic – magmatic mixing trend for these elements. Previous dating of sericite and xenotime associated with mineralisation (Cross et al., 2005; Fraser et al., 2012; Bagas et al., 2014) both indicate that mineralisation in the Tanami region was coeval with magmatism. This is supported by the occurrence of intrusion-hosted gold deposits in this region. Therefore, the timing of the mineralisation, the relatively high salinity of the fluids in both the Tanami and Kurundi – Kurinelli goldfields, and their trace element contents all indicate that the deposits in both regions are intrusion-related gold deposits. The higher gold endowment of the Tanami region may result from the multiple gold-mineralisation events in this region whereas fluid flow associated with mineralisation in the Kurundi – Kurinelli goldfields appears to be of limited extent.

Declaration of Competing Interest

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

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Appendix A. Supplementary data

Supplementary data to this article can be found online at https://doi.org/10.1016/j.oregeorev.2019.103189.

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