INSTITUTO DE GEOCIÊNCIAS

# Metamorfismo e Tectônica da Região de Alfenas-Areado-Alterosa, MG: Contribuições Da Termobarometria e Datação da Unidade Granito-Migmatito Areado

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Dissertação apresentada ao Programa Mineralogia e Petrologia do IGC-USP para a obtenção do título de Mestre em Ciências

Área de Concentração: Petrologia Ígnea e Metamórfica

Orientador: Porf. Dr. Renato de Moraes

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# UNIVERSIDADE DE SÃO PAULO INSTITUTO DE GEOCIÊNCIAS

"Metamorfismo e Tectônica da Região de Alfenas-Areado-Alterosa, MG: Contribuições Da Termobarometria e Datação da Unidade Granito-Migmatito Areado"

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"The known is finite, the unknown infinite; intellectually we stand on an islet in the midst of an illimitable ocean of inexplicability. Our business in every generation is to reclaim a little more land, to add something to the extent and the solidity of our possessions."

T.H. Huxley, 1887

"The Cosmos is all that is or was or ever will be"

Carl Sagan, Cosmos, 1980

Dedicated to my uncle Narciso Benini Which was the grandfather that I've never had And unfortunately another victim of the Covid-19

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### ABSTRACT

Rocks from the Southern Brasilia Orogen basement occur between Alfenas and Areado, MG, Brazil, here called Areado Granite-Migmatite Unit. This unit is composed of garnet meta-syenogranite, hornblende-garnet metatexite, and biotite stromatic diatexite with different degrees of partial melting. Even with many geological maps, no detailed work has been done in this area to understand its geotectonic meaning or the complex pattern of metamorphism, partial melting, and deformation. This study characterizes the Areado Granite-Migmatite Unit and unravels the reason for a different pattern of deformation and partial melting of this unit and its geological and tectonic context. Petrography associated with modal analysis, structural data, quartz c-axis analysis, zircon dating in ICP-MC-MS, and trace elements chemical analysis in quartz and titanite to determine metamorphic, partial melting, and deformation evolution were used as tools in this investigation. The garnet meta-syenogranite has a crystallization age of 2069±5 Ma which indicates that this rock was generated in the last stages of the Mineiro Belt. This unit is the protolith of the migmatites, which is inferred for field observation and accessory minerals comparison. Lower intercepts from zircon indicate a metamorphic age between 671±58 and 657±40 Ma. Melt crystallization ages are inferred between 621 and 604Ma. Water influx and deformation during metamorphism are the cause of different rates of partial melt. The big variation in the proportion of alkali feldspar and plagioclase occurs as an effect of melting in the presence of water, together with segregation and melt loss processes. Quartz c-axis thermometer yielded deformation temperatures of 766-892 °C and another lowtemperature pulse between 613-714 °C, both between 11.5-15 kbar. Those values match the Zr-in-titanite thermobarometer data, which is 786-815°C for hightemperature crystals and 667-708°C for low-temperature titanite. Although these methods have given robust results, several issues must be assessed on these thermobarometers before their application. These data indicate a deformationalmetamorphic event of high temperature, associated with partial melting before the Três Pontas-Varginha thrust. Thus, the Areado Granite-migmatite Unit should be considered as part of the Southern Brasilia Orogen in later studies.

#### Resumo

Rochas do embasamento Orógeno Sul de Brasília ocorrem entre Alfenas e Areado, MG, Brasil, aqui denominadas Unidade Granito-Migmatito Areado. Esta unidade é composta por um meta-sienogranito, um granada-hornblenda metatexito e um biotita diatexito estromático com diferentes graus de fusão parcial. Mesmo com muitos mapas geológicos, nenhum trabalho detalhado foi feito nesta área para entender seu significado geotectônico ou o padrão complexo de metamorfismo, fusão parcial e deformação. Este estudo caracteriza a Unidade Granito-Migmatita Areado e desvenda a razão de um diferente padrão de deformação e fusão parcial desta unidade e seu contexto geológico e tectônico. Petrografia associada à análise modal, dados estruturais, análise do eixo c de quartzo, datação de zircão em ICP-MC-MS e análise química de elementos traço em quartzo e titanita foram utilizados para determinar evolução metamórfica, fusão parcial e deformação foram utilizadas como ferramentas nesta investigação. O granada meta-sienogranito possui idade de cristalização de 2069 ± 5 Ma, o que indica que essa rocha foi gerada nos últimos estágios do Cinturão Mineiro. Esta unidade é o protólito dos migmatitos, baseado nas observações de campo e comparação de minerais acessórios. Interceptos inferiores da Concordia e idades de borda de zircão indicam uma idade metamórfica entre 671 ± 58 e 657 ± 40 Ma. As idades de cristalização por fusão são inferidas entre 621 e 604Ma. O influxo de água associada à deformação durante o metamorfismo são as causas de diferentes taxas de fusão parcial. A grande variação na proporção de feldspato alcalino e plagioclásio ocorre como efeito da fusão na presença de água, juntamente com processos de segregação e perda de derretimento. O termômetro de eixo-c de quartzo gerou temperaturas de deformação de 766-892°C e outro pulso de baixa temperatura entre 613-714°C, ambos entre 11,5-15 kbar. Esses valores correspondem aos dados do termobarômetro de Zr-em-titanita, que é 786-815°C para cristais de alta temperatura e 667-708°C para titanita de baixa temperatura. Embora esses métodos tenham dado resultados robustos, existem várias questões que devem ser avaliadas nesses termobarômetros antes de sua aplicação. Esses dados indicam evento deformacional-metamórfico de alta temperatura, associado ao um derretimento parcial antes do empuxo Três Pontas-Varginha. Assim, a Unidade Granito-migmatito Areado deve ser considerada parte do Orógeno Brasília Sul em estudos posteriores.

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#### **Chapter I – Introduction**

The Southern Brasilia Orogen has been the subject of many academic researches (*e.g.* Fuck *et al.*, 1994; Brito-Neves *et al.*, 1999; Campos Neto, 2000; Trouw *et al.*, 2000), however, there are several regions inside this tectonic framework and still need a detailed approach to fully understand its evolution. It is composed of nappe systems which involve rocks from a magmatic arc (Socorro-Guaxupé Nappe), and rocks associated with the magmatic arc basins (Andrelândia Nappe system) and passive margins (Carrancas and Lima Duarte Nappe systems). All these systems were subjected to regional metamorphism from green schist to granulite facies (Campos Neto, 2000; Trouw *et al.*, 2000; Campos Neto *et al.* 2010, 2011).

At the northwest of this structure, the nappe thrusts a group of metagranitic rocks that went through partial melting, which here are denominated Areado Granite-migmatite unit. This area has been considered as part of the Campos Gerais Complex (Artur, 1988) which comprehends an intensely reworked unit during different tectonic cycles and has a huge lithological variety (Cavalcante *et al.* 1979) but no other detailed work has been made on the region to define the correct tectonic framework of the migmatites.

Despite the historical doubt, the Areado Granite-migmatite Unit presents a very complex petrological situation. In regional metamorphism, geologists observed the effect of the same pressure and temperature on rocks of a big area, however, the migmatites show different rates of partial melt and deformation inside a small area, which raises the question about the melting process that affected the area. Thus, this research shows data to unravel the geological history and petrological evolution that involved the Areado Granite-Migmatite Unit

#### I.1 Dissertation presentation

In the last few years, the Institute of Geosciences has suggested the students publish articles during the academic period and include them in their dissertations or thesis. This way of presentation presents a big advantage in the sense of making the dissertation more compact with the same academic quality, and the use of published articles shows an interesting approach since the articles have already been peer-assessed, which guarantees a high-quality text. With this idea, this dissertation follows the structure below:

**Chapter I – Introduction:** This section will introduce the subject of the research, along with the objectives, justification, and the location of the studied area.

**Chapter II – Geological Setting:** This chapter describes in bigger detail the geological context than presented in the article and the earlier studies of the Areado granite-migmatite unit.

**Chapter III – Analytical Methods:** This section presents the methodology used to achieve a good quality result, together with the scientific background of the methods, with a major focus on thermobarometry.

**Chapter IV – Results:** This chapter exhibits the published article "The role of deformation-assisted by water influx on partial melting of a granite protolith, and its role in the evolution of southwestern São Francisco Craton margin, Brazil" accepted on the Journal of South America Earth Sciences on October 17<sup>th</sup>, 2021. It shows the results and a major discussion about the data obtained from the Areado granite-migmatite unit.

**Chapter V – Concluding Remarks:** It presents a summary of the discussion presented in the article and a reflection about the use of the methods on the samples.

**Chapter VI – References:** A list of all citations used in this volume, with exception of chapter IV.

#### I.2 Objectives

The project covers structural, petrological, and geochronological aspects of the Areado Granite-migmatite Unit to define the relation of the unit with other rocks around it, the connection it has with the geological evolution of the area, and understand the metamorphic process that affected these rocks and caused a big difference in melting fraction. Thus, the objectives of this work are:

- Characterize petrologically the main lithotypes present in the Areado Granite-migmatite Unit and understand the relation between them.
- Analyze the structural variation, together with geological mapping to establish the connection of this unit with other units around it.
- Define the proper thermobarometers for these rocks and find the metamorphic peak of the rocks.
- Date both the crystallization age of the metagranite and the partial melting age to generate a geological model for the evolution of the unit.
- Integrate all data and clarify the tectonic framework that the Areado Granite-migmatite Unit is inserted and the metamorphic process that generated such different partial melting rates.

## I.3 Geographical location

The studied area is located between the cities of Alfenas and Areado, in the southern region of Minas Gerais. The main accesses to the area are the MG-491 that connects the cities to Varginha, the Fernão Dias highway (BR-381), and the roads MG-179 and MG-369, which pass between Pouso Alegre and Alfenas (Fig. 1)



Figure 1 – Geographical localization of the studied area and the main roads to access it

#### **Chapter II – Geological Settings**

The rocks investigated are located at the southernmost part of the Brasilia Orogen, a neoproterozoic fold belt, which borders the São Francisco Craton at its west and southwest margins. The São Francisco Craton is formed by greenstone belts, granites, gneisses, and migmatites of the archean and paleoproterozoic ages. At the craton eastern border, the so-called Mineiro Belt crops out, in which rocks formed or affected by a paleoproterozoic orogeny occur. As mentioned earlier, in a structural window, to west, between the Passos, to the north, and the Andrelândia Nappe Systems, to the south, there are old rocks, also of archean and paleoproterozoic ages, neither attributed to the São Francisco Craton nor to Mineiro Belt. As they were partially affected by this neoproterozoic orogeny, and so, they were related to the older Paramirim Craton.

#### II.1 Mineiro Belt

The Mineiro Belt is situated at the southeastern border of the São Francisco Craton, which is limited by the Jeceaba-Bom Sucesso and Congonhas-Itaverava lineaments (Campos and Carneiro, 2008; Ávila et al., 2010; Teixeira et al., 2015) and the Lenheiro shear zone (Teixeira and Figueiredo, 1991). The stratigraphy of the Mineiro Belt exhibits paleoproterozoic orthogneisses and igneous bodies associated with coeval supracrustal units (Alkmim and Teixeira, 2017). Orthogneisses and granites are of granite, granodiorite, and trondhjemite composition, with metaluminous to peraluminous and calc-alkaline affinity (e.g. Ávila et al., 2010, 2014; Teixeira et al., 2015), and present four different orogenic groups, based on isotopic and geochemical data, such as 2.47-2.41, 2.36-2.33, 2.23-2.20 and 2.17-2.09 Ga (Teixeira et al., 2017). The formation of these rocks is related to a long-term magmatic accretion, associated with oceanic subduction slab (Noce et al., 2000). Sm-Nd whole rock and Lu-Hf in zircon data attest the juvenile source of the Mineiro Belt magmatism (e.g. Ávila et al., 2010, 2014; Teixeira et al., 2015; Moreira et al., 2018). So far, the youngest recognized plutonic rocks that are attributed to the final amalgamation involved in the formation of Mineiro Belt date within the range of 2.17 and 2.10 Ga (Barbosa *et al.*, 2015) and are related to an accretionary orogeny (Teixeira *et al.*, 2015). These granites are separated into two groups, the first corresponds to rocks formed between 2.17-2.15 Ga and generated by partial melting of LREE enriched MORB and with crustal assimilation. Meanwhile, the second group presents ages between 2.14-2.10 Ga, they present geochemical similarities to metadiorites, depleted in Ti, Nb, and P, formed by fractional crystallization and contamination with crustal material (Barbosa *et al.*, 2015).

### II.2 Southern Brasilia Orogen

The Brasilia Orogen (Fuck *et al.*, 1994; Dardenne, 2000) is part of the Tocantins Province (Almeida, 1981) and it is interpreted as the result of the collision of the Central Goiás and Paranapanema blocks against the San Franciscan Plate (Brito Neves *et al.*, 1999; Campos Neto, 2000; Trouw *et al.*, 2000). This orogen is segmented into three domains (Fuck *et al.*, 2017): (i) the External domain, which comprises cratonic rocks and its undeformed cover; (ii) the Internal domain, which represents the metamorphic area of the orogen; and (iii) the Goias Magmatic Arc domain (see Fig 2, Chapter IV). The metamorphic grade increases from the External towards the Internal domain (Fuck *et al.*, 2017), reaching ultra-high temperature metamorphic conditions (*e.g.* Moraes *et al.*, 2002). The metamorphic event is neoproterozoic (*e.g.* Pimentel, 2016).

The Southernmost Brasilia Orogen denotes a thick-skinned nappe system, with a 300 km displacement (Brito-Neves *et al.*, 1999; Campos Neto, 2000). It was divided into a domain of passive margin related to the San Franciscan paleocontinent with basement rocks (*e.g.* Cioffi *et al.*, 2016; Peternel *et al.*, 2005), along with metasedimentary rocks, that corresponds to the São Vicente Complex, Carrancas and Lima Duarte Nappe Systems (*e.g.* Trouw *et al.*, 2008; Westin *et al.*, 2016; Westin and Campos Neto, 2013). The other domain recalls to the active margin of the Paranapanema with a metasedimentary unit from Andrelândia Nappe System (*e.g.* Campos Neto *et al.*, 2011, 2010), together with magmatic arc unit, the Socorro-Guaxupé Nappe (Campos Neto, 2000; Campos Neto and Caby, 2000; Janasi, 2002).

Dating zircon crystals from rocks of the Socorro-Guaxupé Nappe, with the U-Pb method, reveal the age of arc magmatism to be before the collision stage, ca. 730-640 Ma (Rocha et al., 2017, 2018). Metamorphic conditions are extreme, with 940 °C at 11 kbar, in the granulites at the base of the nappe (Moraes et al., 2015; Rocha et al., 2018), and diatexites at their top (Rocha et al., 2017). The age of this UHT metamorphism is ca. 630-625 Ma, proving to be syn-collisional (Campos Neto et al., 2004; Rocha et al., 2017, 2018), whereas zircon crystals from leucosome of granulites and diatexites crystallized in a considerable interval, from 612 to 600 Ma (Rocha et al., 2017, 2018; Tedeschi et al., 2018). Below these UHT granulites, the Três Pontas-Varginha Nappe occur, and it comprises high-pressure kyanite-garnet granulite, associated with subordinate garnet mafic granulite, and calc-silicate rocks (Trouw and Castro, 1996; Campos Neto, 2000; Campos Neto and Caby, 2000; Trouw et al., 2000; Garcia and Campos Neto, 2003; Reno et al., 2009; Motta and Moraes, 2017). Although several authors investigated the P-T conditions of metamorphism, the most recent calculations combine Zr-in-rutile and Ti-in-quartz data and produced 850 °C and minimum pressures of 15 kbar (Martinez, 2015).

#### II.3 Campos Gerais Complex

The Campos Gerais Complex is frequently reported as a "complex unit, intensively recycled and crustally reworked in successive tectonic cycles" (*e.g.* Cavalcante *et al.*, 1979; Artur, 1988). The Campos Gerais Complex is composed of a variety of rocks: gneiss and migmatites of granodiorite to tonalite compositions, with lenses of metamafic and metaultramafic rocks, schists, quartzite, and intrusive granite bodies (Cavalcante *et al.*, 1977; Artur, 1988; Szabó, 1989). This unit has been divided into North and South Domains (*e.g.* Morales *et al.*, 1983; Schrank *et al.*, 1990; Szabó, 1996). The first is composed of an autochthonous terrane with granite-greenstone belt characteristics like the Morro do Ferro Greenstone Belt sequence (Teixeira *et al.*, 1987; Szabó, 1996). Meanwhile, the second domain comprises paragneisses, and intercalated sequences of metasedimentary rocks with amphibolites and metaultramafic units (*e.g.* Morales *et al.*, 1983; Szabó, 1996). Recently, it was proposed that the Campos Gerais Complex should be restricted to the granites and orthogneisses

from the North Domain (Turbay *et al.*, 2008). It is also recognized that intrusive paleoproterozoic metagranites crosscut the rocks of the Campos Gerais Complex (Turbay and Valeriano, 2012).

The rock investigated in the present research has not yet been well mapped or properly separated in any regional map, in some of them it is mapped as part of the Andrelânida Nappe System (Campos Neto, 2000), but in some others, it is considered as part of the Campos Gerais Complex (*e.g.* Campos Neto *et al.*, 2004; Heilbron *et al.*, 2017). This data corroborates with observations made by Turbay *et al.* (2008). The age of metamorphism of the Campos Gerais Complex is 2.9 Ga (Wernick *et al.*, 1981; Teixeira *et al.*, 1989) . In recent work, the crystallization ages for the protoliths of some of these orthogneisses have been determined as 2.9 Ga, using U-Pb dating in zircon, with partial melting and metamorphism at 2.6 Ga, and possibly with the presence of some primitive crust, even older, of ages 3.2-3.6 Ga (Valeriano *et al.*, 2006).

### **Chapter III – Applied Methods**

### III.1 Geological Mapping

A regional geological reconnaissance has been made in order to define the structural relation between the migmatitic units of the area. Most outcrops of the area are presented as slabs for the diatexite unit, and blocks for the other units, with the exception of the Areado quarry where the meta-syenogranite lithotype is present. To achieve a better result, we made cross-sections with structural data measurements and generated a Geological Map using ArcMap 10.1 software.

### **III.2** Petrography

During the geological mapping stage, several samples were acquired to obtain more information about the mineralogical, structural, and textual characteristics of the rocks involved in this project. To do so, an Olympus BXP-40 microscope with an Infinity One camera coupled was used in the Optical Microscopic Laboratory from the Núcleo de Apoio à Pesquisa Geoanalítica (NAP) of IGc-USP. For the modal analysis, an Olympus U-FMP chariot was used to count minerals. Since most of the samples presented coarse crystals, we utilized a 1mm square grid for counting, with the minimum amount of 600 points counted in each thin section. In addition, for units with megacrysts, photographs were used to count the proportion of megacryst and groundmass. This was made with a 20x30 grid in Corel Draw X5 program.

#### **III.3 Structural Analysis**

A detailed structure description was made along with the collection of structural data using a Clar-type compass. These structural data were loaded on Stereonet software (v. 9.5.1) and projected on a south hemisphere Schmidt-Lambert stereogram to look for a correlation between the structures. The cross-sections were generated using the plotted structures on the map, no vertical exaggeration was used on them.

#### **III.4** Microstructural analysis

Mineral microstructures encountered in deformed rocks are usually associated with the preferential orientation of the main structure. Many mechanisms are responsible for the formation of oriented crystals, however, in the case of the same crystal shape, like quartz, the main process is the dislocation creep, which rotates the grains due to the change of habit and the interaction with neighbor minerals (Paschier & Trouw, 2005). The variation of sliding in relation to the crystalline structure of quartz is dependent on the temperature. In lower temperatures, the migration tends to follow the <a> crystallographic axis, while the crystals tend to dislocate towards the <c> crystallographic axis at higher temperatures (Kruhl, 1986, 1996). In the case of quartz grains, the increase of temperature also alters the recrystallization textures and the way that the grains dislocate, from bulging (BLG), to Sub-grain rotation (SGR) and grain boundary migration (GBM) respectively (Stipp *et al.*, 2002b).

The c-axis quartz thermometer was created by Kruhl (1998) that demonstrated the correlation of the spatial position of the quartz <c> axis with the deformation temperature. The quantification is made using the distribution of the <c> axis data on the stereogram and measuring the opening angle between the major concentrations. The author used both natural and experimental data to calibrate this thermometer and estimated a 50°C error margin.

Although it is a practical method, some factors can influence the temperature calculation. Both strain rate and water presence can cause a variation of the data. In addition to it, divergences in the way of the opening angle measurements can also interfere with the results (Law, 2014). Faleiros *et al.* (2016) raise the importance of the pressure as a factor of angle variation generated a calibration based on opening angle, pressure and temperature:

$$T(^{\circ}C) = 410.44 \ln OA + 14.22P(kbar) - 1272$$

OA stands for the c-axis opening angle, P for pressure, and T for temperature. This calibration also extended the thermometer interval to rocks from granulite facies and ultra-high temperatures, and it also presents an error margin of 50°C.

The Faleiros *et al.* (2016) calibration was used in this work to estimate the deformation temperature of the studied samples. A Zeiss Ortholux microscope with a four-axis Leinz Wetzlar universal-stage from the Laboratório de Óptico (IGc-USP) was used to obtain the c-axis measurements, and the data were treated on the Stereo 32 software to obtain the grindles and the opening angles. The calculation was made on a spreadsheet in Microsoft Excel 2013.

#### III.5 Thermobarometry

Two thin sections and one polished block were used to analyze the trace elements in titanite and quartz in a quadrupole ICP-MS iCAP Q – Thermo Scientific model coupled with a 213 A/F – New Wave Laser Ablation Sampler. During the procedure, Heraeus glasses were used as intern standard, for quality control and calibration of the equipment. Data was inserted in Glitter (v4.4.4) to

quantify the amount of trace elements; to do so, a stoichiometric proportion obtained from Deer *et al.* (1994) was used to calibrate the estimative. This data was used with different thermometers, which are explained below.

#### III.5.1 Titanium in quartz (TitaniQ)

Titanium is commonly found in igneous and metamorphic quartz crystals, controlled by a coupled substitution reaction between Si, Ti, and other cations (Wark and Watson, 2006). The diffusion of Ti in quartz is slower than other elements like aluminum and gallium, however, its valence facilitates the exchange. The excess of titanium inside the crystals can cause exsolution of TiO2 in the form of rutile needles, concentrating the element in a sector of the grain (Cherniak *et al.*, 2007). Wark and Watson (2006) calibrated a thermometer based on synthetic samples of quartz in the presence of TiO2 component, between 600 to 1000°C and 1GPa pressure, with a Ti detection limit of 15 ppm. The authors considered the pressure an irrelevant fact in the calibration of TitaniQ. However, Thomas *et al.* (2010) calibrated o TitaniQ as a thermobarometer with synthetic analysis of quartz and rutile, and silica-saturated fluids between 700 and 940°C and pressures of 5 and 20 kbar with a detection limit of 8 ppm. The calibration equation is:

 $RTlnX_{TiO_2}^{quartz} = -60952 + 1520T(K) - 1741P(kbar) + RTlna_{TiO_2}$ 

Where  $X_{TiO_2}^{quartz}$  represents the molar fraction of TiO2 in quartz grains, T is the temperature in Kelvin, P is pressure in kbar, and  $a_{TiO_2}$  is the activity of TiO2 on the rock. The experiments from Thomas *et al.* (2010) showed a great influence of the pressure on the concentration of Ti in quartz grains. This calibration was used in this research and the calculations were made on Microsoft Excel 2013.

#### III.5.2 Zirconium in titanite (Zr-in-Tit)

Titanite is a common accessory mineral of igneous and metamorphic rocks and has a huge potential to be applied as a thermobarometer, due to its capacity for element substitution in its crystalline structure. Zirconium (Zr) usually substitutes titanium in the mineral structure. Its diffusion in titanite is considered intermediate since it is slower than lead and oxygen, but faster than rare earth elements. It also tends to preserve the formation temperature better than other elements because of its slower diffusion gradient (Cherniak, 2006).

Hayden *et al.* (2008) calibrated a thermobarometer based on de diffusion of Zr in titanite in an interval of 600-1000°C and up to 10 kbar, with a detection limit of 75ppm. The formula of the calibration is:

$$logZr^{titanite} = 10.52 - \frac{7708}{T} - 960\frac{P}{T} - \log(a_{TiO_2}) - \log(a_{SiO_2})$$

Where  $Zr^{titanite}$  is the Zr concentration in the titanite crystal in ppm, T is temperature, P is pressure,  $a_{TiO_2}$  and  $a_{SiO_2}$  represent the TiO2 and SiO2 activity in the system, respectively. Due to the dependence of the activities, the thermometer is perfect for the appliance on rocks that has quartz and rutile, however, granites without rutile can be tricky to use it. The authors suggest using  $a_{TiO_2} = 0.6$  in that case since most granites present that activity. It is possible to use the Zr-in-titanite together with TitaniQ to constrain the pressure and temperature of a sample (Thomas *et al.* 2010).

#### III.6 U-Pb based geochronology

Dating the Areado migmatite-granite unit is essential to unravel the geological evolution of the area. As cited above, many geological maps of the region give divergent information about the geological history associated. To define the tectonic framework, associated with petrological studies, this research also abroad U-Pb dating of zircon and titanite looking for crystallization and metamorphism ages. Both zircons and titanite from the samples were separated

using the protocol described in Loios (2004). Then, the generated samples were used in three different sessions.

One of them was conducted on a Thermo Scientific Neptune LA-MC-ICP-MS and an Excimer 193nm laser from Photon Machine, at CPGeo, Institute of Geosciences, University of São Paulo to evaluate the sample ARE-04, which correspond to a non-active quarry, south of Areado, MG (23K 0379323/7631788). The second session involved another sample from the quarry (CGE-25) and other samples from the migmatites, north of Alfenas, MG (CGE-03Zr and CGE-05). This session was carried out at the Sensitive high-resolution ion microprobe (SHRIMP IIe/MC) from the Institute of Geosciences, University of São Paulo, Brazil. The last session happened on SHRIMP II at the John de Laeter Centre of Mass Spectrometry at Curtin University and evaluated U-Pb data from the sample TPR-40, which comes from an outcrop of the diatexite leucosome from Três Pontas, MG area.

#### **Chapter IV – Results**

#### Abstract

The Areado Granite-Migmatite Unit is part of the basement from the Southern Brasilia Orogen at southwestern São Francisco Craton margin, outcropping between Alfenas and Areado regions, Minas Gerais State, Brazil. This unit is composed of garnet meta-syenogranite, hornblende-garnet metatexite, and stromatic diatexite with different degrees of partial melting. This study characterizes the garnet meta-syenogranite, its migmatitic counterparts and investigates the different patterns of deformation and partial melting involved to produce the observed migmatites within the unit. The connection between the garnet meta-syenogranite, hornblende-garnet metatexite, and biotite diatexite is inferred from field observations and assemblage of accessory minerals. Petrography associated with modal analysis, structural data, guartz c-axis analysis, zircon U-Pb (LA-MC-ICP-MS and SHRIMP) dating, as well as trace elements chemical analysis in quartz and titanite to determine metamorphic, partial melting, and deformation evolution were used as tools in this investigation. The garnet meta-syenogranite has a crystallization age of 2069.8 ± 5 Ma. Lower intercepts from zircon indicate a metamorphic and partial melting age between  $671 \pm 58$  and  $657 \pm 40$  Ma. Melt crystallization ages are inferred between  $622 \pm$ 7 Ma and 604  $\pm$  6 Ma. Different rates of partial melt are structurally connected, being higher at the base of the unit, and are related with contrasting amounts of water influx and deformation. The deformation during metamorphism/partial melting was the main cause of the differential aspect of the produced rocks. Quartz c-axis thermometer yielded deformation temperatures of 765-935 °C and other pulses occurred between 590-715 °C, both between 11-15 kbar. Those values match the Zr-in-titanite thermobarometer data, which is 670-710 °C for low-temperature titanite and 785-815 °C for high-temperature crystals. These data indicate that the protolith was generated at a Rhyacian late magmatic stage of the Mineiro Belt and passed by a Neoproterozoic deformational-metamorphic event of high temperature.

*Keywords:* metamorphism, quartz c-axis, thermobarometry, water-influx melting.

### **IV.1** Introduction

Regional metamorphism is associated with orogeny, resulting in the formation of mountain ranges. It comprises an increase in pressure and temperature on active margins, leading to recrystallization and metamorphism, which has as a primary characteristic feature to produce similar *P*-*T* metamorphic conditions affecting a large crustal region. As the base of these mountain ranges cannot be accessed directly to be investigated, the rocks formed at these portions of the continental crust and exposed in ancient exhumated orogens are the targets to evaluate the processes acting at these sites, such as high-grade metamorphism and partial melting. They are accessed in the outcrops of metamorphic rocks, such as migmatites and granulites (*e.g.* Sawyer, 2008; Moraes *et al.*, 2015).

The studies of ancient high-grade metamorphic rocks can give us clues about the evolution of the lower crust in the past. Metamorphic studies of those rocks usually investigate mineral assemblages, their composition, ages, and textures to generate a pressure-temperature-time model and show how deep the rock was in the crust and how hot it was during the metamorphic peak and along its exhumation. The models can be made by thermodynamic modeling, analysis of metamorphic minerals pair for geothermometer generation, experimental studies, or analysis of trace elements in metamorphic minerals. In addition, metamorphic textures may reveal the processes that the migmatites passed through, such as partial melting, segregation, and crystallization. These data, associated with geochronological studies can show the geological behavior of the lower crust on large orogenic belts and give a broader view from the crustal evolution of the Earth and insights of the processes happening in new orogens, such as the Alps and the Himalayas.

In Brazil, the Southern Brasília Orogen is a thick-skinned structured nappe belt, where pieces of its basement were involved and worked during its formation in the Neoproterozoic (Fuck et al., 1997; Campos Neto, 2000; Valeriano, 2017). These slices are observed in several levels of crustal depth offering an opportunity to investigate the processes of transformation that are imposed in the relatively cold, old and dry crust when involved in much younger orogenic events. In southwestern Minas Gerais, Archean, and Paleoproterozoic rocks crop out in the infrastructure of the Southern Brasília Orogen, some related to the Mineiro Belt, a Paleoproterozoic orogen situated at the southern edge of the São Francisco Craton (Teixeira *et al.*, 2000).

Between the cities of Alfenas and Areado (Fig. 3), a garnet metasyenogranite, along with its metamorphic equivalents, a hornblende-garnet metatexite, and a biotite stromatic diatexite, here called the Areado Granite-Migmatite Unit, crops out as infrastructure of the uppermost portion of the Andrelândia Nappe System, one of the main metasedimentary units in the southernmost portion of the Brasília Orogen. In addition, the diatexite from this unit also appears as blocks close to Três Pontas city (see Fig. 3). Although the Areado Granite-Migmatite Unit occurs with a relatively small thickness, it spreads in a large area, a variable degree of deformation and partial melt is observed, which is not compatible with just simple variation of temperature of regional metamorphism. In addition, although several geological maps have been made in the region (*e.g.*, Cavalcante *et al.*, 1977; Artur, 1988; Ribeiro *et al.*, 2014; Campos, 2015), none of them has made detailed research on this unit, which difficult the definition of its tectonic framework, petrological and geological meaning.

In the present paper, it is investigated the relationship between metamorphism, heterogeneous deformation, and variation in the degree of partial melting within the Areado Granite-Migmatite Unit, to understand how the different metamorphic products were generated in a small geographic area. The investigation is based on petrography, deformation temperatures, which were determined using c-axis quartz thermometer (Faleiros *et al.*, 2016), metamorphic temperatures calculated using *in situ* analysis of trace element in quartz and titanite, using, respectively, the thermobarometers Ti-in-quartz (TitaniQ; Thomas *et al.*, 2010) and Zr-in-titanite (Hayden *et al.*, 2008). Also, the crystallization age of the garnet meta-syenogranite is determined using zircon crystals and LA-MC-ICP-MS, and the age of metamorphism and partial melting is determined with

SHRIMP using zircon and titanite grains, to constrain its metamorphic/partial melting and crystallization ages and tectonic history.

#### IV.2 Geological setting

The rocks investigated in this study are located in a key geotectonic position, where Archean to Paleoproterozoic rocks associated with the southwesternmost margin of the São Francisco Craton are surrounded by rocks of the southern part of the Neoproterozoic Brasília Orogen (Fig. 2). The São Francisco Craton, which was consolidated during the Neoproterozoic, is composed of TTG rocks and greenstone belt assemblages. It crops out in southern Minas Gerais and Bahia States, being mostly covered by sedimentary rocks of Bambui or São Francisco basins (Alkmim, 1993). Its eastern border is formed by the Mineiro Belt, a Paleoproterozoic orogen constituted of several magmatic arcs (Teixeira et al., 2000). To the west of the western limit from the São Francisco Craton, in a structural window, croup out a group of high-grade rocks of tonalitic to granitic protoliths along with minor occurrences of supracrustal rocks, including greenstone belts, some of them grouped in the Campos Gerais Complex (Cavalcante et al., 1977; Morales et al., 1983; Teixeira et al., 1987; Artur, 1988; Szabó, 1989; Schrank et al., 1990; Szabó, 1996). Available zircon U-Pb data for orthogneisses indicate Archean magmatic episodes between 3.0 and 2.7 Ga, metamorphic episodes at 2.2 and 2.0 Ga, and late granitic magmatism at 1.9 Ga (Valeriano et al., 2006; Turbay, 2010). The Campos Gerais Complex presents an E-W-trending elongated shape limited in north and south by the Passos, Andrelândia, and Três Pontas-Varginha Nappe Systems from the Brasília Orogen (Fig. 3). Although it displays the main structure parallel to these Neproterozoic nappes, no unequivocal geochronological data are supporting the interpretation that it was substantially involved in the Neoproterozoic deformation. Neoproterozoic dating in zircon grains from Campos Gerais Complex rocks is restricted to very imprecise lower intercept U-Pb ages between 650 and 510 Ma (Turbay, 2010).

The Brasilia Orogen is interpreted as the result of the collision of the Central Goiás Massif, Paranapanema and São Francisco-Congo Plates during the

Neoproterozoic (Fig. 3; Fuck *et al.*, 1994; Brito Neves *et al.*, 1999; Dardenne, 2000; Campos Neto, 2000; Trouw *et al.*, 2000; Valeriano *et al.*, 2004). The southern part of the orogen denotes a thick-skinned nappe system, with a 300 km displacement (Brito-Neves *et al.*, 1999; Campos Neto, 2000). It consists of schists and paragneiss, along with metabasic rocks associated to passive margin setting, called the Araxá Group, in Passos Nappe (Valeriano *et al.*, 2004; Valeriano, 2017) and Carrancas and Lima Duarte Nappe Systems (*e.g.*, Trouw *et al.*, 2008; Westin *et al.*, 2016; Westin and Campos Neto, 2013). Structurally above this



Figure 2 - Geological and tectonic map of the Southern Brasilia Orogen, associated with the southwestern margin of the Sao Francisco Craton. Dashed coloured lines separate the orogens in three domains: I- The External domain, II- The Internal Domain and III - Goiás Magmatic Arc. Keys:1—Ribeira belt; 2—São Francisco craton basement; 3—Paranoá Group; 4—Bambuí group cratonic cover; 5—Vazante Group; 6—Canastra Group; 7—Ibiá Group; 8—Araxá Group; 9—Araxá Group metamorphosed under granulite facies conditions; 10—Magmatic arc complexes; 11—Phanerozoic cover (Paraná Basin). In the southernmost part of the orogen, number 6, 8 and 9 are considered part of the Andrelândia Nappe system, while number 10 is called Socorro-Guaxupé Nappe (e.g. Campos Neto, 2000; Campos 2010, 2011; Cioffi et al., 2016). Modified from Valeriano (2017).



Figure 3 - Geological map of the southernmost area of the Brasilia orogen. This map shows with more detail the distribution of the units inserted in the Andrelândia Nappe System and Socorro-Guaxupé Nappe, along with the Areado Granite-Migmatite Unit, the Campos Gerais Complex and units related to Passos Nappe. The yellow rectangle marks the study area from this paper and the red star points the location of the sample TPR-40. This map was generated with data from Turbay and Valeriano (2004), and Cioffi et al. (2016).

this last sequence the Andrelândia Nappe System occurs, a metasedimentary sequence associated with an active margin deposition (Campos Neto *et al.* 2010, 2011). The uppermost rocks of this later nappe system are associated with the Três Pontas-Varginha Nappe, which comprises high-pressure kyanite-garnet-orthoclase granulite (Trouw and Castro, 1996; Campos Neto, 2000; Campos Neto and Caby, 2000; Trouw *et al.*, 2000; Garcia and Campos Neto, 2003; Reno *et al.*, 2009). Several studies of this unit produced results of high-pressure conditions for the metamorphic peak, between 800-900 °C and minimum pressures of 15 kbar (Trouw *et al.*, 1998; Campos Neto and Caby, 1999; 2000; Garcia and Campos Neto, 2003; Campos Neto *et al.*, 2010; Motta *et al.*, 2010; Martinez, 2015). U-Pb monazite ages from the Andrelândia Nappe system evidence a metamorphism age of 618-595 Ma for these rocks (Campos Neto *et al.*, 2011).
#### IV.3 Analytical methods

Modal QAP classification for three weakly deformed/metamorphosed garnet meta-syenogranite samples and four hornblende-garnet metatexite samples was performed by counting at least 500 points in thin sections. Proportions of megacrysts, when these are present, were estimated using outcrop photographs to establish megacrysts/matrix ratio, using a 20x30 squares grid of 43mm, to be integrated into modal proportion derived from thin sections.

Quartz c-axis orientations were measured using a four-axis Leitz Wetzlar universal-stage coupled to a Zeiss Ortholux microscope in thin sections cut perpendicular to the foliation and parallel to the stretching lineation. Quartz c-axis data were plotted in the Schmidt-Lambert lower hemisphere grids, using Stereo32 software.

Analyses of quartz and titanite were acquired using an iCAP-Q-ICP-MS, by Thermo Scientific with a solid-state New Wave 213 A/F laser coupled, using 100  $\mu$ m thick thin sections. In the investigated samples, ten quartz crystals were analyzed, in which a raster profile was made along with them, and fourteen titanite crystals were analyzed, but doing spots on each one (analytical uncertainties, 2 $\sigma$ , of 1.22 and 14.44, respectively). An internal calibration curve was built with Heraeus synthetic glasses as standard for quartz, and for titanite using standards from NIST, with data refinement and reduced using Glitter v. 4.4.4. Analytical conditions from LA-ICP-MS for these analyses are summarized in Table 1. All these laboratories are linked to the NAP-Geoanalítica Facilities of the Institute of Geosciences, University of São Paulo, Brazil.

Zircon and titanite grains were separated from four samples of all rocks that were collected in selected outcrops for U-Pb dating (ARE-04 and CGE-25 samples from the garnet meta-syenogranite; CGE-05 from the hornblendegarnet metatexite and CGE-03Zr from the biotite stromatic diatexite; see Fig. 4 for locations). One of the analyses was carried out using a Thermo Scientific Neptune LA-MC-ICP-MS and an Excimer 193nm laser from Photon Machine, at CPGeo Facility, Institute of Geosciences, University of São Paulo, Brazil. The other five data were obtained in Sensitive high-resolution ion microprobe (SHRIMP IIe/MC) at the Institute of Geosciences, University of São Paulo, Brazil. Concentrations were made following a protocol designed by Loios (2004) and analytical procedures, as described by Sato *et al.* (2008, 2014). For dataset treatment, Concordia plots were constructed with the ISOPLOT 4.11 spreadsheet (Ludwig, 2003). Analyses with common lead content over 5% and discordance over 5% were discarded. Once all the crystallization U-Pb ages obtained were older than 1.300 Ma, the <sup>207</sup>Pb/<sup>206</sup>Pb ages were used in a Probability Density Plot (PDP) to dispose of the most representative age graphically.

Table 1 - Analytical conditions for in situ analysis in quartz and titanite in LA-ICP-MS

Analyzed Material	Spot Size (µm)	Laser Fluency (J/cm <sup>2</sup> )	Repetition Rate (Hz)
Quartz (thin section)	80	3,765	20
Quartz (block)	80	14	20
Titanite	55	2	15

Additional zircon grains were extracted from biotite diatexite (TPR-40) via traditional methods of disaggregation, magnetic and heavy liquid separation. These were mounted in 25-mm diameter epoxy discs together with NBS-610 glass and primary and secondary zircon standard material, polished to approximately half-grain thickness, carbon-coated, and then imaged by cathodoluminescence (CL) using a TESCAN MIRA3 FESEM. The mounts were then cleaned and covered with a 40 µm gold coat prior to U-Pb isotopic analysis using the Sensitive High-Resolution Ion Microprobe (SHRIMP II). All this equipment is housed in the John de Laeter Centre of Mass Spectrometry at Curtin University.

SHRIMP II was operated at a mass resolution of 5,000, with a primary beam current of 2 nA and beam diameter of 20  $\mu$ m, and a 6-scan cycle was used during analysis (de Laeter and Kennedy, 1998; Kennedy and de Laeter, 1994). NBS-610 glass was used to calibrate the position of the <sup>204</sup>Pb peak. CUYZ (Curtin internal laboratory standard - 568.55 Ma) was used as the primary standard and OGC (Stern et al., 2009) as a secondary standard. Data were processed using SQUID II and Isoplot 3.75 (Ludwig, 2009, 2003). The <sup>207</sup>Pb/<sup>206</sup>Pb age of secondary standard determined during the analytical session was 3461.0 ± 8.4 Ma, which is within error of the published age (3465.4 ± 0.6 Ma; Stern et al., 2009). U–Pb data are presented in Table S1 (Supplementary Material).

## IV.4 The Areado Granite-Migmatite Unit

The northern and southern boundaries of the Areado Granite-Migmatite Unit (hereafter Areado Unit) are characterized by sheared tectonic contacts with units of very distinct nature and evolution. To the north, it overrides a unit of reddish muscovite quartzite that presents a schistosity marked by the preferred orientation of muscovite and opaque mineral-rich layers. In its southern boundary, the Areado Unit is overridden by (high-pressure) kyanite-garnetorthoclase granulite from the Três-Pontas Varginha Nappe. Very close to the contact, the granulite has a mylonitic foliation marked by stretched K-feldspar, garnet, kyanite, and quartz and all of them mark an intense mineral stretching lineation. The Areado Unit was divided into three subunits: garnet metasyenogranite, hornblende-garnet metatexite and biotite stromatic diatexite, as described below (Fig. 4A, B).

#### IV.4.1 Garnet meta-syenogranite

The garnet meta-syenogranite has a massive inequigranular blastoporphytic texture (Fig. 5A), with a weak foliation close to the edge of the body or within localized shear zones. Its color is grey with localized red garnet crystals. Euhedral phenocrysts of perthitic microcline, with size between 0.8 and 7 cm, are set in a groundmass composed of K-feldspar, plagioclase, quartz, biotite, garnet, titanite, ilmenite, zircon, apatite and allanite (Table 2). The groundmass has inequigranular seriate texture, varying from medium to coarse-grained. Rare leucosome veins with idioblastic hornblende occur (Fig. 5B, C).



Figure 4 - (A) - Geological map of the study area, marked in Fig 2. Here is shown the essential outcrops for this paper, the analysis applied in each rock, and the most representative structural data. For the full-size map, visit the Apendix section. The units described are 1- Garnet meta-syenogranite; 2- Hornblende-garnet metatexite; 3- Biotite stromatic diatexite; 4- Garnet Amphibolite; 5- Muscovite quartzite and 6- Kyanite-garnet granulite with rutile. Units are ordered by stratigraphic position. (B) - Cross-sections generated from the structural data present on the map. The vertical scale is equal to the horizontal scale. All cross-sections show that Areado Granite-Migmatite Unit structures are controlled by the thrust of the kyanite bearing granulite.

Quartz crystals have undulose extinction and rounded, lobated, and irregular contacts, suggesting that deformation was accommodated by grain boundary migration recrystallization mechanism (GBM). Plagioclase is tabular with optical compositional zoning and wedge-shaped albite twinning. Garnet exclusively occurs intergrown with K-feldspar and sometimes biotite in finegrained reaction-rim coronas surrounding partially corroded grains of coarsegrained plagioclase, K-feldspar and biotite (Fig. 6A). Films of plagioclase surrounding garnet and K-feldspar in these coronas are pseudomorphing a



Figure 5 - Photographs from: (A) - The main texture of the Garnet meta-syenogranite. It is possible to observe the phenocrysts of K-feldspar weakly oriented and a cross-cutting vein. (B) - Leucosome patch inside the Garnet meta-syenogranite with idioblastic hornblende inside. (C) - Central leucosome vein channelling smaller leucosome veins in contact with K-feldspar phenocrysts. The sketch highlights the leucosome patches forming in the contact of the phenocrysts and the groundmass of the rock, and its connection to the leucosome vein.

former melt phase (e.g., Sawyer, 1999, Fig. 5C). A second generation of biotite occurs as xenomorphic grains, mostly with partial replacement by chlorite. Ilmenite crystals are xenomorphic and present a corona of metamorphic titanite (Fig. 6B), when in contact with plagioclase. Idiomorphic to subdiomorphic grains of apatite present coronas of allanite.



Figure 6 - Photomicrographs: (A) - Igneous plagioclase crystal mantled by metamorphic garnet and metamorphic K-feldspar when in contact with biotite in the Garnet meta-syenogranite unit. This texture comprehends a reaction between igneous plagioclase and biotite, forming metamorphic garnet and k-feldspar. (crossed polarized light). (B) - Anhedral opaque crystal mantled by titanite rim on the Garnet meta-syenogranite, which represents the substitution of the opaque by titanite during metamorphism (uncrossed polarized light).

#### IV.4.2 Hornblende-garnet metatexite

The hornblende-garnet metatexite is an inequigranular rock with centimetric K-feldspar porphyroclasts, while minerals at the matrix are millimetric (Fig. 7A). The rock has a mylonitic foliation marked by K-feldspar porphyroclasts, recrystallized quartz lenses, and, by biotite, which has, locally, totally replaced by garnet. The leucosome appears as patches and in veins, which crosscut the foliation planes. Shear zones cut the foliation and some host leucosome, as well (Fig. 7A).

The leucosome is composed of K-feldspar, quartz, plagioclase (An<sub>28</sub>), and hornblende. Quartz grains present lobed boundaries and undulose extinction, which indicates it recrystallized by GBM. The K-feldspar porphyroclasts have inclusions of euhedral plagioclase, indicating its igneous origin, whereas the smaller crystals were originated by recrystallization and are sitting at porphyroclasts rims or matrix. Hornblende occurs in the leucosome, as idioblastic crystals with inclusions of quartz, plagioclase, K-feldspar, garnet, allanite, titanite and zircon.

The residuum is composed of porphyroclasts of K-feldspar, and a matrix of quartz, xenoblastic crystals of plagioclase and K-feldspar, garnet, titanite, allanite, zircon, and apatite. Garnet crystals are idioblastic, with titanite, allanite and zircon inclusions. The biotite is subdioblastic and occurs at the edge of garnet and hornblende crystals. Allanite exhibits medium to coarse-grained xenoblastic grains, with inclusions of subdioblastic titanite and apatite. Titanite occurs either as subdioblastic crystals, with straight borders against quartz and feldspar, or as xenoblastic agglomerates with garnet and allanite inclusions.

Despite the intense deformation, melt crystallization textures (e.g., Sawyer, 1999, 2001; Marchildon and Brown, 2002; Holness and Sawyer, 2008), such as small quartz or feldspar films surrounding garnet grains (Fig. 8A) and a second generation of K-feldspar filling porphyroclast fractures (Fig. 8B), both mimicking a trapped melt phase, are present.

Table 2 - Modal analysis from most representative samples. The mineral proportion already considers a matrix/phenocrystals proportion of 58/42 for the meta-syenogranite unit, and a matrix/porphyroclasts proportion of 74/26 for the hornblende-garnet metatexite unit. Both proportions were obtained by photography modal counting.

Unit	Sample type	Samples	Qtz	PI	Kfs	Bt	Grt	Tit	Ор	Hbl	Aln	Ар	Zr	Ms	Ep	Total
Meta- syenogranite	Whole Rock	ARE-4C	19%	21%	50%	6%	2%	1%	1%	Trace	Trace	Trace	Trace	Trace	Trace	100%
		ARE-4B	24%	16%	49%	7%	2%	Trace	1%	Trace	Trace	1%	Trace	Trace	Trace	100%
		CGE-25C	12%	24%	54%	7%	1%	1%	Trace	100%						
		Mean	18%	20%	51%	7%	2%	1%	1%	Trace	Trace	Trace	Trace	Trace	Trace	100%
Hornblende- garnet metatexite	Whole Rock	CGE-05A	21%	11%	53%	Trace	8%	2%	Trace	4%	1%	Trace	Trace	Trace	Trace	100%
		CGE-05B	19%	13%	54%	1%	7%	1%	Trace	4%	1%	Trace	Trace	Trace	Trace	100%
		CGE-06B	29%	7%	47%	1%	6%	1%	Trace	8%	1%	Trace	Trace	Trace	Trace	100%
		CGE-10B	23%	7%	55%	1%	7%	1%	Trace	5%	1%	Trace	Trace	Trace	Trace	100%
		Mean	23%	10%	52%	1%	7%	1%	Trace	5%	1%	Trace	Trace	Trace	Trace	100%
Biotite stromatic diatexite	Whole Rock	CGE-03C	27%	38%	12%	18%	Trace	2%	1%	100%						
		CGE-21	36%	37%	15%	7%	Trace	4%	Trace	100%						
		CGE-23B	36%	33%	10%	16%	Trace	3%	1%	100%						
		Mean	33%	36%	12%	14%	Trace	3%	1%	100%						
	Leucosome	CEG-03EL1	35%	26%	29%	4%	Trace	6%	Trace	100%						
		CGE-03EL6	34%	34%	27%	3%	Trace	1%	1%	100%						
		Mean	35%	30%	28%	4%	Trace	3%	Trace	100%						
	Residuum	CGE-03ER1	20%	35%	7%	30%	Trace	2%	6%	100%						
		CGE-03ER4	28%	36%	9%	21%	Trace	4%	2%	100%						
		Mean	24%	36%	8%	25%	Trace	3%	4%	100%						



Figure 7 - Photographs of (A) - Main texture from the hornblende-garnet metatexite. It is possible to see a shear zone, with deformed minerals at its edges, channelling the leucosome in a patch. (B) - Cross-section view from the leucosome and residuum bands from the biotite stromatic diatexite, emphasizing the stromatic character of this unit. (C) - E-W horizontal view from the biotite stromatic diatexite unit. This picture shows the behaviour of the mylonitic foliation of the rock, forming several duplex structures, as highlighted by the sketch. (D) - E-W horizontal view of the main structure of the biotite stromatic diatexite unit. The image focuses attention on the relation between the leucosome and residuum band, along with a rotated deformed porhyroclast.



Figure 8 - Photomicrographs of: (A) - Felsic film mimicking melt in garnet contacts at the hornblende-garnet metatexite residuum. This texture is associated with a melt crystallization (crossed polarized light). (B) - K-feldspar porphyroclast from the hornblende-garnet metatexite, surrounded by quartz, garnet and allanite. The crystallization of a metamorphic k-feldspar generation inside the porphyroclast is a melt crystallization texture and corroborates partial melting in the rock (crossed polarized light). (C) - Biotite stromatic diatexite melanosome highlighting a preserved igneous plagioclase crystal, together with a late-stage muscovite cross-cutting biotite (crossed polarized light).(D) - The overall texture from the biotite stromatic diatexite that presents the leucossome and residuum of the rock. The sketch focus on the different quartz textures, showing lobated crystals in the leucosome, in which prevailed the grain boundary migration (GBM) process, while the melanosome has microcrystals of quartz close to biotite, thus where bulging (BLG) process dominates in this unit (crossed polarized light). All samples were made perpendicular to the foliation of the rock and parallel to the lineation.

#### IV.4.3 Biotite stromatic diatexite

The biotite stromatic diatexite is the most voluminous subunit, sitting at the base of the Areado Unit (Fig. 4). The term diatexite is used here because the degree of partial melting was high enough to destroy all original igneous texture, and the syn-anatexis intense deformation was responsible to imprint a stromatic structure in the rock (Fig. 7B), marked by a white leucosome and a black or grey residuum, rich in biotite. The mylonitic foliation is best represented by intensely deformed biotite. Two different fold types are observed, intrafolial, and open harmonic types (Fig. 14A and B). In addition to the mylonitic foliation, boudins of biotite-rich layers, S-C foliation (Fig. 7C) and rotated porphyroclasts (Fig 7D) are recognized.

The residuum reaches between 70-60% in volume (Table 2), being composed of biotite, plagioclase, quartz, epidote, titanite, muscovite, allanite, and zircon. Biotite is subdioblastic, has microstructural features, such as mica fish and defines pairs of S-C foliations. Allanite occurs with a total or partial corona of epidote, indicating it was replaced by the last, in a similar way as described by Finger *et al.* (1998). Quartz occurs as grains with straight contact, usually forming bands. Most plagioclase crystals are rounded without twinning and present concentric extinction. Retrograde muscovite overgrowth biotite grains and form symplectites with quartz and K-feldspar (Fig. 8E), evidencing a replacement in an open system.

The leucosome is composed of plagioclase (An<sub>27</sub>), quartz, and K-feldspar. Although leucosome is easily recognized in the field, in thin sections, minerals are deformed, and pristine igneous textures are rare. Although with feldspar grains with irregular boundaries being more common, there are euhedral plagioclase crystals with igneous texture preserved (Fig. 8C). Rarely, some perthitic microcline porphyroclasts occur. Quartz in these layers present lobated boundaries.

#### IV.5 U-Pb zircon and titanite dating

In order to constrain the age of igneous crystallization and metamorphism, four zircon and two titanite mounts from studied rocks were prepared for U-Pb dating.

The selected zircon crystals generally present homogeneous typology varieties, with 100 and 400 µm size. The most common morphological types include elongate crystals, exhibiting rounded to oval-shaped terminations. Between the most common zircon internal features, it is possible to observe: (1) bright cores exhibiting tenuous oscillatory zones; (2) thin homogeneous rims, darker than the cores and possibly generated by recrystallization; (3) metamitic zones are developed preferably along the rim-core boundaries; (4) the absence of inherited cores; and (5) an intense fracturing in crystals of the biotite stromatic diatexite unit (See Apendix).

For U-Pb data acquirement in LA-MC-ICP-MS of the garnet metasyenogranite, thirty-eight spots were concentrated at the bright cores, and among those, fourteen analyses were used to calculate the Concordia date at 2069.8±5 Ma (Fig. 9A). SHRIMP IIe analyses used between twelve and fifteen spots to calculate the isocron intercepts, resulting in 2079±21 Ma date for the garnet metasyeno-granite (Fig. 9B) and 2083±12 Ma for the hornblende-garnet metatexite unit (Fig. 9C). Although zircon crystals from the biotite stromatic diatexite present a significant amount of fractures, six-spot analysis from metamorphic rims resulted in a Concordia date of 604.2±6.3 Ma (Fig. 9D). This date is younger than obtained by the lower intercept of the garnet metasyenogranite (657±40 Ma, Fig. 9B) and the hornblende-garnet metatexite (671±58 Ma, Fig. 9C).



Figure 9 - Isocron graphs showing U-Pb analysis results. (A) - Sample of the garnet meta-syenogranite zircons on LA-ICP-ME-MS with the calculated Concordia Age. (B) - Isocron from zircon analysis of the meta-syenogranite, showing upper and lower intercepts. (C) - Isocron from zircon analysis of the hornblende metatexite, with upper and lower intercepts. (D) - Zircon analysis results from the Biotite stromatic diatexite, along with the calculated Concordia age. (E) - Titanite isocron from the meta-syenogranite with the calculated Concordia age. (F) - Calculated Concordia Age for titanite data from the metatexite unit.(G) Concordia diagram for zircon analyzed via SHRIMP II. From (B) to (F), all analysis have been made on SHRIMP from IGc-USP.

In addition to the metamorphic ages obtained with zircon, titanite crystals from the garnet metasyeno-granite and hornblende-garnet metatexite have been dated. These crystals are granular, mostly with angular shapes, with 100 to 200  $\mu$ m. Cathodo luminescence (CL) images show no signs of zoning in the crystals, only a homogeneous dark area. U-Pb analysis in SHRIMP from the garnet metasyenogranite gave a calculated Concordia date of 621.8±7.0 Ma (Fig. 9E). Data from the hornblende-garnet metatexite have a high content of common Pb and should be carefully treated. Still, Concordia date calculated from those crystals is 557±19 Ma with high concordance probability (0.8, see Fig. 9F).

# IV.6 Quartz and titanite *in situ* analysis and thermobarometry

The quartz in situ analyses were made on thin sections of the sample CGE-05 and a polished rock block of the sample CGE-03D (Fig. 4). Titanite analyses were made on samples ALFE-24E and CGE-09A (Fig. 4). All LA-ICP-MS analyses were performed using the raster mode.

Fourteen large quartz crystals from samples CGE-05 and CGE-03D were analyzed, which yielded TiO2 concentration below 8 ppm (Tables A1 and A2 – see Apendix). The results vary between 2.8 and 7.8 ppm (CGE-05 thin sections), and from 1.2 to 6.3 ppm (CGE-03D polished block). The TiO2 content was calculated based on an intern calibration curve made with Heraeus standard glasses, and the collected data have very good quality (see Supplementary data). Taking an average result and using the calibration by Thomas et al. (2010), temperatures between 374-650 °C are calculated for the hornblende-garnet metatexite, and 355-623 °C for the biotite stromatic diatexite, between 1 and 15 kbar (Fig. 10A). These calculations took into account an aTiO2 of 0.5, which is considered a lower limit for crustal rocks (Hayden and Watson, 2007). Ashley et al. (2014) propose an aTiO2 value of 0.2 for deformed rocks since deformation can relocate Ti inside quartz structure. If it is applied to these analyses, the results change to 450-732 °C for the hornblende-garnet metatexite and 428-700 °C for the biotite stromatic diatexite (Fig. 10B).



Figure 10 - P-T diagram showing (A) - Calculated temperature with TitaniQ and Zr in titanite using titanium activity equal to 0.5 as suggested by Hayden & Ward (2007). (B) - Calculated temperature with TitaniQ and Zr in titanite using titanium activity equal to 0.2, as suggested by Ashley et al. (2014). The Zr in titanite was separated by textural group, indicated in Figure 13.

Thirteen titanite crystals from samples CGE-09 and ALFE-24 were analyzed, including twelve idioblastic and one xenoblastic grain (Fig. 11A, Tables A3 and A4). A stoichiometric composition from average titanite crystals was used for trace element calculation (Deer et al., 1992) for Glitter calibration. The titanite REE chondrite normalized patterns are different for each type of crystal (Fig. 11B). The idioblastic grains present a higher concentration of LREE, from La to Sm, and a discrete negative Eu anomaly. HREE have a slightly concave curvature between Gd and Dy, and a decrease from this element to Lu (Fig. 10B).

On the other hand, the xenoblastic titanite grain also has a positive slope for LREE, a pronounced negative Eu anomaly, and almost flat distribution of HREE, with a small decrease between Gd and Lu (Fig. 11B). These patterns indicate that xenoblastic titanite might have formed before or along with other HREE rich minerals, such as zircon or garnet (Bea, 1996), whereas idioblastic titanite must have formed after these phases, even after the xenoblastic grains. The negative Eu anomalies suggest that both types of titanite crystals formed at subsolidus conditions after all feldspars have crystallized from the melt and captured its Eu.

The Zr-in-titanite thermometer calibrated by Hayden et al. (2008) was applied considering aTiO2 as 0.5 and aSiO2 as 1.0, due to quartz presence in all

samples. Zirconium content of idioblastic titanite in CGE-09 has an average of 55 ppm, which yields temperatures of 559-702 °C for pressures between 1 and 15 kbar, respectively. Crystals from sample ALFE-24E have an average of 70 ppm of Zr, which yields temperatures of 569-713 °C, for the same pressure range. The xenoblastic grains show a zirconium content of 640 ppm that corresponds to a temperature interval of 665-827 °C (Fig. 10A). Using aTiO2 0.2, as suggested by Ashley et al. (2014), the results change to 550-692 °C for the CGE-09 sample, 560-703 °C for idioblastic crystals, and 654-814 °C for xenoblastic crystals in ALFE-24E (Fig. 10B). The uncertainty, based on the crystals analyses, is  $\pm$  30 °C for the TitaniQ and up to 20 °C for the Zr-in-titanite thermometer.



Figure 11 - (A) - Photomicrographs (uncrossed polarized light) illustrating different titanite textures on the metatexite unit residuum. The first one, correspond to a xenoblastic titanite, usually with allanite inclusions. The second are idioblastic titanites, that rarely presents any inclusions. (B) - RRE pattern from titanites normalised by the chondrite and separated by textural groups. This graph suggests that the formation of xenoblastic titatine absorbed most elements present in allanite during its formation. Idioblastic crystals' generation occurred afterwards when the system was already depleted in LREE.

## IV.7 Main structures and quartz c-axis based thermometry

A mylonitic foliation is the main structure of the study rocks. The mylonite character is represented by K-feldspar porphyroclasts and shear zones filled with leucosome in the hornblende-garnet metatexite (Fig. 7A), while S-C foliations and feldspar porphyroclasts are evident in the biotite stromatic diatexite (Fig. 7B, C). In both migmatitic units, drag folds occur as intrafolial tight folds and open folds. In the eastern and middle part of the map (Fig. 4A), the foliation has an average of N020/45SE and most structures indicate movement to NW (Fig. 12A, B). In this region, cross-sections (Fig. 4B) show that the Três Pontas-Varginha Nappe thrusts the Areado Unit with a similar shear sense, which is based on their asymmetric structures (Fig. 7C, 7D). Deformation created two different types of folds in the biotite stromatic diatexite, a tight intrafolial fold and a rounded hinge fold with an oblique axial plane (Fig. 13A). Field relationships indicate a three deformational stage process; the first one deformed a pre-existent metamorphic surface, developing a foliation S<sub>n</sub>. The second stage formed the tight folds and the main mylonitic foliation  $(S_{n+1})$ , and the third stage generated only the harmonic folds (Fig. 13B).



Figure 12 - Stereograms generated from the structural data obtained in the field. (A) - Stretching lineation from the biotite diatexite, mostly acquired at the eastern part of the map. (B) - Mylonitic foliation, axial plane and fold axis from the biotite diatexite. (C) - Data obtained at Areado are, and consists of mylonitic foliation from the hornblende diatexite along with foliation and stretching lineation from the kyanite granulite.

In the western part of the area, the average orientation of the foliation changes to N170/32NE, and the foliation poles match with the foliation measured in the kyanite-garnet-orthoclase granulite unit (Fig. 12C). Stretching lineation in the granulites exhibits a low-angle spread behavior (Fig. 12C).



Figure 13 - Photographs of folds found in the biotite stromatic diatexite. (A) - Intrafolial fold with an axial plane parallel to the mylonitic foliation. (B) - Photograph of the same unit illustrating the second generation of folds, with an oblique axial plane concerning the mylonitic foliation. Both sketches highlight the characteristics of the folds.

In order to evaluate if it is possible to identify each deformationtemperature pulse or just the last one, the quartz c-axis thermometer was applied to a set of samples (Kruhl, 1996; Morgan and Law, 2004; Law, 2014; Faleiros *et al.*, 2016). Quartz c-axis orientations were measured in eight thin sections (Fig. 4). All thin sections are cut parallel to stretching lineation and perpendicular to mylonitic foliation. Quartz c-axes were measured with a universal stage, and the opening angle was estimated based on the maximum concentrations.

The hornblende-garnet metatexite has quartz crystals with lobed contacts, typical of recrystallization due to GBM. The quartz c-axis pattern in stereogram forms a girdle correspondent to point c-maxima at a low angle to the stretching lineation (Fig. 14A). Most samples from the biotite stromatic diatexite present quartz recrystallization by BLG and these data present a girdle with point c-maxima in Y and an intermediate position between X and Z directions (Fig. 14B). However, sample CGE-23A presents lobated quartz well preserved, surrounded by small, recrystallized quartz crystals. In this case, the two different crystal types were measured separately. The small crystals data formed a girdle with a similar pattern described above, with a small rotation of the data in relation



Figure 14 - Quartz c-axis fabrics measurements on lower hemisphere equal-area projections. Foliation is oriented on the X-axis, and stretching lineation is parallel to the cut. (A) graphic represents a point c-maxima at a low angle to the stretching lineation, with a rotation concerning X-axis, just like graphs (D), (E) and (F). Figures 7(B) and (C) show a girdle with point c-maxima in Y and an intermediate position between X and Z pattern, but with a small rotation in (C). Figures 7(G) and (H) presents an asymmetric type I crossed girdle. The temperature estimation is based on 11-15kbar pressure, obtained from the other thermometers (Section 7). All images shows the sample measured (visit Fig 3 for their geographic position), which recrystallisation process (grain boundary migration (GBM) or bulging) prevails in the thin section and the unit it belongs.

to the X-axis (Fig. 14C). The lobated quartz data generated point cmaxima at low angles to the stretching lineation, however, with some imprint of low angle measurements (Fig. 14D). Other biotite stromatic diatexite samples have quartz grains recrystallized by GBM, and present quartz c-axis data of cmaxima at a low angle to the stretching lineation, with a c-axis rotation about Xaxis (Fig. 14E and 13F). In the samples from the kyanite-garnet-orthoclase granulite, from Três Pontas-Varginha Nappe, quartz grains present GBM recrystallization and, locally, ribbons. Both samples formed asymmetric type I crossed girdles (Law, 2014; Faleiros *et al.*, 2016; Fig. 14G, H).

The resultant deformation temperatures obtained using the calibration made by Faleiros *et al.* (2016) present two different clusters (Table 3 and Fig. 15). The pressure range used was that calculated with thermobarometry under *a*<sub>TiO2</sub> equal to 0.2, 11-15 kbar. The first group corresponds to high deformation temperature, varying from 766 to 934 °C, between upper amphibolite and granulite facies. This cluster matches the hornblende-garnet metatexite, the biotite stromatic diatexite from the Areado region, and quartz crystals with GBM texture of the biotite stromatic diatexite of the Alfenas region. The second cluster corresponds to an amphibolite facies deformation temperature, varying from 591 to 714 °C. Quartz c-axis measurements from the kyanite-garnet-orthoclase granulite give a range of 518-711 °C at the same pressure, which is a lower deformation temperature than the Areado Unit metamorphic peak.



Figure 15 - C-axis thermobaromether results plotted on a P-T diagram. The colours of the lines represent the mapping units from Fig 3.P-T estimative based on TitaniQ and Zr-in-Titanite calculations from Fig 14B is displayed, showing that minerals formed on that temperature corroborates with low-temperature deformation.

Unit	Sample	Opening Angle	Pressure (kbar)	Temperature (°C)	
Hornblende-garnet		4449	1	675	
metatexite	CGE-05A (GDM)	111	15	874	
		760	1	515	
Biotite stromatic diatexite	CGE-03C (BLG)	75	15	714	
		700	1	497	
	CGE-23D (DLG)	12	15	697	
		116°	1	693	
	CGE-23B (GBINI)	110	15	892	
		108°	1	664	
	CGE-35A (GDIVI)	100	15	863	
	$CGE_{41A}$ (GBM)	08°	1	624	
		90	15	823	
		61°	1	512	
Três Pontas-Varginha Nappe		04	15	646	
		75°	1	447	
		75	15	711	

Table 3 - Calculated temperatures for quartz c-axis deformation temperatures, based on Faleiros *et al.* (2016) calibration.

# **IV.8 Discussion**

#### IV.8.1 Physical controls of partial melting and metamorphism

From the garnet meta-syenogranite, increasing deformation, metamorphism and partial melting transformed the rock into the hornblendegarnet metatexite as described earlier. However, there are still some doubts about the precedence of the stromatic biotite diatexite. It shows biotite as the main mafic mineral, with garnet as an accessory mineral in some samples. This suggests that a high concentration of water in this unit stabilized primarily biotite. The unit also presents increased plagioclase vs K-feldspar ratio in relation to the protolith (A vs P 12:36 and 51:20, respectively). It is possible that this major difference of feldspar proportions indicates that the diatexite does not come from the same protolith, if melting occurred in equilibrium. Still, field evidences point to another direction.

Field structural relationships indicate progressive changes with the correlated increase of deformation intensity and melt fraction. Weakly deformed/metamorphosed garnet meta-syenogranite presents ~3 vol.% of random pockets of leucosome that turn into 12 vol.% of leucosome veins either

parallel to or that crosscut the mylonite foliation (Fig. 7A), culminating in a stromatic structure where the leucosome reaches 40 vol.% of the rock in biotite stromatic diatexite (Fig. 16). This relation indicates that deformation exerted a strong control on the partial melt rate. This is reinforced by the fact that dynamic recrystallization of feldspar is more intense in the migmatites dominated by the moderate melt fraction. The presence of the same igneous mineral association in all three rock groups, including K-feldspar large crystals and the accessory minerals, such as allanite, apatite, zircon, and ilmenite corroborate that all rocks have an equivalent protolith, the garnet meta-syenogranite.

In addition, experimental studies by Johannes and Holtz (1992) show that the melting reaction of large plagioclase crystals and a Qtz-Or composed glass produces an unchanged plagioclase core with a reaction zone composed of small crystals of An-rich plagioclase, glass, biotite, and K-feldspar intergrowth. This texture is similar to the one represented in Fig 6A. Experiments on synthetic granite composition, run by Whitney (1975), produced melt in presence of water, at a minimum temperature of 800 °C. The produced melt consumed more Kfeldspar and left a plagioclase-rich residuum, in a similar way to the observed rock here. It is also observable in phase relation for biotite-granite generated using natural samples from Dinkey Lakes, where the K-feldspar reacts and leaves the system before plagioclase (Piwinskii, 1968b, 1973; Boettcher and Wyllie, 1968; Stern and Wyllie, 1981), what in a natural system would favor to increase the compositional differences between the original garnet metasyeno-granite and its most extreme derivative, the biotite stromatic diatexite, where it is observed the major difference in bulk composition and feldspars ratio. These processes can change the modal proportion of feldspars during melting, segregation, meltloss, and crystallization history. The differential melting, followed by segregation and melt loss were probably the processes responsible to enhance the impoverishment in K-feldspar in the biotite stromatic diatexite. Thus, field relation between the rocks, accessories minerals, peritetic minerals (discussed below) and experimental data from many different studies corroborate with the hypothesis that the protolith of this unit is the hornblende-garnet metatexite, which started as the meta-syenogranite.

In this model, the bulk-rock composition cannot be considered as the main control of the partial melting variation. In addition, regional metamorphism usually

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Figure 16 - Sequence of photographs from the outcrop CGE-09 illustrating the transition between the hornblende-garnet metatexite unit (A) to the biotite stromatic diatexite unit (C) passing through a transition section (B). From images (A) to (B), it is possible to observe the comminution of K-feldspar porphyroclasts and the beginning of banding formation. In figure (C) the comminution process is complete, with only a few small porphyroclasts left.

affects large extensions of the continental crust, which means that temperature and pressure must be similar, or not vary too much, within the investigated region. This consideration allows excluding temperature and pressure as the variables responsible for the heterogeneous amount of melt present in the three subunits. Considering that water facilitates partial melting, as it lowers solidus temperature (e.g., Sawyer, 2010; Weinberg and Hasalová, 2015), and that deformation can facilitate water-influx from external sources (Carvalho et al., 2016), the possible relationship between deformation-assisted water influx and partial melting rates should be investigated. This can be done by considering the modal mineral proportions in the residuum and leucosome from garnet meta-syenogranite, hornblende-garnet metatexite, and biotite stromatic diatexite, in addition to the peritectic mineral phases present in each case.

The peritectic minerals in each rock type, along with modal analysis of the samples, show that water content has a key role in the melting, deformation, and crystallization processes of the Areado Granite-Migmatite Unit. The less deformed garnet meta-syenogranite presents minimal evidence for metamorphism in the form of garnet + K-feldspar ± plagioclase reaction-rim coronas surrounding rounded (corroded) grains of igneous plagioclase, K-feldspar, and biotite (Fig. 6A). Plagioclase in these coronas is present as films pseudomorphosing a former melt (Fig. 8A). The following melting reaction can explain these features:

$$Qtz + PI + Kfs1 + Bt1 = Grt + Kfs2 + Liq (1)$$
$$IIm + PI + Bt = Tit + Bt2 + Grt (2)$$

More intensely deformed garnet meta-syenogranite samples present biotite intergrown with garnet and K-feldspar in similar reaction-rim coronas, suggesting the following reaction, which has formed by limited water-influx:

$$Qtz + PI + Kfs1 + Bt1 \pm H2O = Grt + Bt2 + Kfs2 + Liq (3)$$

The deformed hornblende-garnet metatexite present peritectic hornblende and titanite in addition to garnet, biotite, and K-feldspar, suggesting the following reaction, which also needs limited water-influx:

Idioblastic hornblende can also be found in leucosome veins in the garnet meta-syenogranite unit. These veins collected melt from the edges of K-feldspar and are formed by the same reaction. Locally, partial melting consumed all water and biotite and residuum and leucosome got rid of all biotite and only garnet was produced by reaction 1.

The presence of hydrated peritectic phases, biotite or hornblende, is good evidence for melting induced by water-influx (e.g., Weinberg and Hasalová, 2015). However, it seems that water influx was limited, as garnet was produced as the only anhydrous phase, in some portions, consuming all biotite, and hornblende is also observed in some localized sites.

The modal analysis also shows evidence of water interference on melt crystallization (Table 2). Data obtained from sixteen samples shows that the hornblende-garnet metatexite presents more K-feldspar in its composition, while the biotite stromatic diatexite presents more plagioclase. This difference in proportion is caused by the amount of water in the system, where the higher content crystallizes more albite than orthoclase (e.g., Tuttle and Bowen, 1958; Luth et al., 1964; Huang and Wyllie, 1975; Holtz et al., 1992c, 1995; Johannes and Holtz, 1996). This implies that after partial melting took place, in the site where the biotite stromatic diatexite was formed, melt loss occurred, and the lost melt carried a composition enriched in K-feldspar components.

The diffusive contact between leucosome and host rock along with the absence of mafic selvedges are morphological characteristics of neosomes generated by water influx (Sawyer, 2010), features also observed in the field. The amount of water injected into the system caused the difference observed. There was more water injection on the east than on the west of the area, and from bottom to top, the degree of partial melting decreased towards the top. The water origin is still unknown, but its access was controlled by deformation, which is quite intense and synchronous with partial melting.

In summary, we believe that both migmatite units came from the garnet meta-syenogranite, but this occurred in steps, first with the formation of the hornblende-garnet metatexite and from it evolving to the biotite stromatic diatexite. Each roch shows different melting fractions and melt timing. The hornblende-garnet metatexite was formed by high grade metamorphism and dehydration melt with a small amount of water added to the system, while the biotite stromatic diatexite came later, with a more intense water-influx at the base of the unit, using the already modified hornblende-garnet metatexite as its precursor.

Even if, we cannot rule out the possibility that the biotite stromatic diatexite could be a generated from a different rock, such as the country rock of the garnet meta-syenogranite, as we could not find any zircon with a paleoproterozoic date for a direct connection, although the field relationships are straightforward.

#### IV.8.2 Structural evolution and thermobarometry

The structural analysis allowed us to infer that the contact between the Areado Granite-Migmatite Unit and Três Pontas-Varginha Nappe controls the main structures from Alfenas and Areado region. The mylonitic foliation is concordant with the thrust fault, and the kinematic indicators with top to the northwest are inverse to what is observed in the rest of Brasília Orogen, where top movement is primarily to ENE (Campos Neto, 2000; Dardenne, 2000; Valeriano et al., 2004).

Quartz microstructures and quartz c-axis data indicate two deformation mechanism predominance, one of high- and another of lower-temperature, the first mainly related to GBM, whereas the second is associated with BLG recrystallization mechanism. Low-temperature deformation is marked by spaced, millimeter-thick, deformation band cleavage composed of quartz and fine-grained muscovite and epidote. These deformation bands are parallel to the hightemperature mylonitic fabric and cannot be easily distinguished in the field from the high-temperature ones, so they are parallel and define a continuum in deformation processes.

The quartz c-axis thermometer indicates the presence of at least two deformation temperatures, one stage at ~ 835 °C, and another ~ 650 °C, both agree with observed quartz microstructures and, although empirical, quartz c-axis seems to produce good temperature intervals for rock deformation. Higher

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temperatures obtained from quartz c-axis analysis in the biotite stromatic diatexite, however, almost overcome the temperature limit of ultra-high-temperature. With 923 °C, it is an unexpected and unlikely result, since it would mean that there should be orthopyroxene in the rock and probably more than 80-90% of melt in rock with such composition (Johannes and Holtz, 1996), which is not what is observed in outcrops. In addition, with 80-90% melt, it is unlikely that grains could still be in contact to deform and no mylonitic foliation would be formed, as it is observed in outcrops. Although no correlation of water activity and quartz opening-angle has been proved experimentally (Law, 2014), some authors have reported that water-saturated systems tend to favor a prism [c] slip over a basal <a> slip, which would increase the opening-angle (e.g. Blumenfeld et al. 1986, Mainprice et al. 1986, Garbutt and Teyssier 1991, Okudaira et al. 1995). Thus, the calculated temperatures of lobated quartz grains in the biotite stromatic diatexite might have been affected by water presence and the calculated temperature for the stromatic diatexite might be overestimated.

When associated with the quartz deformation temperatures and the ages of metamorphism, the two deformational textures suggest that the deformation of this rock occurred in a single progressive phase. The crystallization of quartz crystals ruled by GBM processes occurred close to the metamorphic peak of the migmatites, along with the mylonitic foliation and leucosome that filled shear zones. The generation of the low-temperature quartz texture resulted from strain localization in less competent layers of the biotite stromatic diatexite during cooling. This process explains the preservation of lobated quartz crystals in the leucosome and the formation of BLG ruled quartz grains in the residuum of the rock. This conclusion is corroborated by quartz deformation temperatures calculated for the Três Pontas-Varginha Nappe granulite, which already indicate low deformation temperatures during thrusting.

The calculation of metamorphic peak conditions for quartz-feldspathic rocks is not a trivial task, and the application of new thermobarometers using trace elements in minerals seems to be a remarkable tool. However, these thermobarometers are dependent on the saturation of oxides on which they are based. Thus, the use of Ti-in-quartz and Zr-in-titanite on rocks is straightforward when saturated in TiO2 and SiO2, which is guaranteed by the free presence of,

respectively, rutile and quartz (e.g., Zack et al., 2004; Thomas et al., 2010). The garnet meta-syenogranite has abundant free quartz and titanite, but no rutile.

Although all analyses have excellent quality, the TitaniQ thermobarometer has not produced the expected results for *P*-*T* metamorphic calculations. The aim is to use this thermobarometer along with Zr-in-titanite to define the *P*-*T* conditions of the metamorphic peak, using the double calibration developed by Thomas *et al.* (2010). The formed intersection by the isopleths of Zr-in-titanite and Ti-in-quartz, using the concentration of the Zr in the idioblastic titanite crystals, occurs at pressures over 20 kbar, which is unrealistic in this geological scenario. One possibility for these results is if the activity of TiO<sub>2</sub> is lower than 1 or even lower, such as 0.5, as advised by authors when titanite is present. Since this rock has no rutile, the TiO<sub>2</sub> activity is smaller than 1, and the approximation of 0.6 given by Hayden *et al.* (2008) and Hayden and Watson (2007) is not compatible with this rock, and results at such high pressure are assumed to be without petrological meaning (~950 °C and 31 kbar).

The difference between the activity of the protolith to the migmatites can be explained by Ti resetting in quartz due to deformation (*e.g.,* Grujic *et al.*, 2011; Ashley *et al.*, 2014; Cavalcante *et al.*, 2014; Bestmann and Pennacchioni, 2015). Ashley *et al.* (2014) suggest using TiO<sub>2</sub> activity equal to 0.2 when the metamorphic rock is intensely deformed. If this value is applied, the results from TitaniQ and Zr-in-titanite of the idioblastic titanite yield an interval between 667-708 °C and 11.5-15 kbar for the metamorphic peak (Fig. 10B). This information correlates to the low-temperature deformation obtained by the quartz c-axis, and this has a good petrological and geological appeal. Xenoblastic titanite crystals were formed at 786-815 °C for the same pressure interval. This data corroborates with high-temperature acquired from GBM formed quartz grains.

Correlating the quartz c-axis thermometer and the other methods suggests that xenoblastic titanite generation crystallized during metamorphic peak (Fig. 10B). It also corroborates the hypothesis that idioblastic titanite occurred together with the low-temperature deformational event. These data are interpreted as two different timing of crystallization and deformation. The older one would reach about 850 °C, forming high-temperature deformation quartz crystal and xenoblastic titanite crystals, probably associated with partial melting/metamorphic peak and it is interpreted as a peritectic phase, as pointed

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earlier. Meanwhile, the younger would comprise the formation of idioblastic titanite and low-temperature deformation quartz grains, during melting crystallization or late-stage solid state deformation/metamorphism.

The Ti-in-quartz, Zr-in-titanite, and c-axis deformation thermometer analyses indicate that caution must be taken into account before using trace element thermometers. The two first methods are excellent tools when quartz and rutile are present in the rock that underwent metamorphism under temperatures at or higher than amphibolite facies. When one of the phases is absent, activity calculations for the thermometer application become a problem. Although some authors suggest a fixed TiO<sub>2</sub> activity value for certain rocks (*e.g.*, Hayden and Watson, 2007; Ashley et al. 2014), there is no recognized method to calculate the real activity, mainly when quartz deformation is associated (e.g., Grujic et al., 2011; Ashley et al., 2014; Cavalcante et al., 2014; Bestmann and Pennacchioni 2015). The activity variation has induced unusual thermobarometer results on these rocks, such as the high pressures obtained by TitaniQ and Zr-intitanite with 0.5 TiO<sub>2</sub> activity (Fig. 10A). Quartz c-axis deformation temperature exhibits interesting results for metamorphic rocks. However, the influence of strain rate and water activity on the rock might affect the results, and its influence has not been quantified (Law, 2014; Faleiros et al., 2016). As field and petrography data present evidence that anatexis occurred with water influx on these rocks, the water activity has a significant influence on too high deformation temperature on the biotite stromatic diatexite, probably respond to some of the very high-calculated temperatures, close to 900 °C (Fig. 15). Even if, as most calculated values are in the range of 830 °C, this value is compatible with the syn-anatectic deformation of this kind of rock and is considered of petrologic importance.

# *IV.83 Geological history and tectonic evolution of the Areado Unit magmatism and metamorphism*

The garnet meta-syenogranite unit presents preserved igneous textures and its transition to migmatites in the field. Its zircon Concordia dates yielded ages of 2069  $\pm$  5 Ma by LA-MC-ICP-MS and 2079  $\pm$  21 Ma by SHRIMP, while zircon U-Pb dates from the hornblende-garnet metatexite unit yielded 2083  $\pm$  12 Ma and 2075  $\pm$  23Ma for zircon extracted from leucosome from the sample TPR- 40. Based on the internal textures of the zircon crystals, it is interpreted as the crystallization ages of the igneous protolith of this unit. It shows that this unit protolith is older than the Socorro-Guaxupé Nappe magmatism, which occurred between 730 and 640 Ma (Rocha et al. 2017, 2018; Tedeschi et al., 2018). Also, the crystallization age at the very end of Rhyacian puts this granite outside of what is called the Campos Gerais Complex, which is composed mainly of Archean rocks (Szabó, 1989; Turbay et al., 2008), but it is also younger than most rocks that intrude it (Turbay et al., 2008). Our U-Pb data are closer to the Pouso Alegre Complex (weighted average of 2078.7  $\pm$  6.7 Ma, Cioffi et al., 2016), and indicate that the magmatic event that formed the protolith of the Areado Unit is related to the evolution of the Mineiro Belt.

The Mineiro Belt comprises a Paleoproterozoic orogenic belt with four arctype related orogenies identified (~2.47–2.41, 2.36–2.33, 2.23–2.20 and 2.17– 2.10 Ga; Seixas et al., 2012, 2013; Ávila et al., 2010, 2014; Barbosa et al., 2015). The protolith of the area shows that the last magmatic event from the accretion of the Mineiro Belt to older rocks from the São Francisco craton is 30 m.y. younger than described in the literature.

The metamorphic U-Pb dates obtained from the lower intercept of zircon rims, metamorphic zircon zones, and titanite also evidence that partial melt and metamorphic history of the Areado Unit occurred ca. 600 Ma and is connected to the metamorphic evolution of the Brasilia Orogen. Zircon grains from the garnet meta-syenogranite yielded a lower intercept of  $657 \pm 40$  Ma and the lower intercept for the grains of hornblende-garnet metatexite is  $671 \pm 58$  Ma. Zircon rims from the biotite stromatic diatexite present Concordia age of  $604 \pm 6$  Ma. Titanite was dated in two rocks, garnet meta-syenogranite and hornblende-garnet metatexite, yielding, respectively,  $622 \pm 7.0$  Ma and  $557 \pm 19$  Ma.

It is considered that the zircon lower intercept represents the closest results for the age of partial melting and metamorphic peak. This data corroborates with high-temperature ages obtained for the Três Pontas-Varginha Nappe (650 Ma for 800 °C, Reno et al., 2009). The younger ages, titanite with  $622 \pm 7.0$  Ma and  $557 \pm 19$  Ma would record colling ages, whereas zircon date of  $604 \pm 6$  Ma from the biotite diatexite records the age of leucosome crystallization.

Regarding the obtained titanite age, we envisage two possible scenarios for these results. The first model would represent a variation in the metamorphic behavior. The metamorphism from the samples of the protolith is more coherent to an anhydrous melting process, with the generation of water-free phases, such as garnet, while from the migmatites, the major process is a water-fluxed melting. Since zircon and titanite samples from the garnet meta-syenogranite gave an older age cluster (657-621Ma) and the migmatites present a younger cluster (604 Ma for the biotite diatexite zircons and 557Ma for hornblende-garnet metatexite unit), this age difference can represent the water influx timing. This would mean that the anhydrous metamorphism began earlier than the water entrance at the bottom of the rock package.

The second possibility is that those rocks passed by a long-term cooling process. Although titanite represents a mineral from the metamorphic peak, its age can be more representative of the cooling process, due to its lower closing temperature for the U-Pb system (600-650 °C, Mezger et al. 1991, Cherniak, 1993). It would be more likely that data from the garnet meta-syenogranite still marks the last stage of melt crystallization, meanwhile, the age obtained from the hornblende-garnet metatexite represents the age of closure of U-Pb system in titanite after its crystallization. These data agree with the cooling ages found in the literature (620 - 588 Ma, Reno et al., 2012). At this stage, melt from the Areado Unit has already crystallized, causing a strain localization on biotite-rich layers and forming the low-temperature quartz deformation fabric.

Since the main metamorphic event occurred during the Neoproterozoic, the Areado Granite-Migmatite Unit should be part of the Southern Brasilia Belt, as the infrastructure of the Andrelândia Nappe System, formed during the collision of the São Francisco and Paranapanema plates and the formation of the Gondwana supercontinent.

A possible tectonic scenario to generate the heat for the metamorphism of these rocks is envisaged here. The Areado Unit is immediately overridden by high-pressure granulites from the Três Pontas-Varginha Nappe, which reached peak metamorphic conditions of 850 °C and 14-16 kbar (Martinez, 2015), and represents that they might have the same heat source. Reno et al. (2009) report that these rocks were at 800 °C at 650 Ma in a post-peak

metamorphic stage. Both data match the metamorphic ages obtained from the zircon lower intercept from the Areado Unit and suggests that this unit was either on the metamorphic peak or on an early stage of cooling at 671-657 Ma. The compilation of these data evidence that the metamorphism of Três Pontas-Varginha Nappe and the Areado Unit is coeval, and both have the same heat source.

The Areado Unit presented a similar cooling history to the Três Pontas-Varginha Nappe, associated with melt crystallization. This hypothesis is supported by the zircon metamorphic rims age of 604 Ma that is similar to the ages of 588 Ma for these granulites (Reno et al., 2012).

The migmatites present many indications that they melted in the presence of a restricted amount of water, e.g., hydrated peritectic minerals, extrapolated quartz deformation temperatures, and the lack of selvedges between leucosome and residuum. The amount of water was restricted, as in several portions, the amount of produced melt is restricted, there is an anhydrous peritectic phase, such as garnet, and both features point to restriction to the amount of the fluxed water into the system. However, the source of the fluid that caused partial melt is unknown. The first hypothesis consists of the dehydration of lower metasedimentary units. The water would rise and promote partial melting, at a higher degree at the bottom of the pile, and a lower degree at an intermediate portion. The rocks that received less water, like the garnet meta-syenogranite and the hornblende-garnet metatexite, stabilized hornblende, garnet, and allanite, while the more hydrated rocks biotite and, later, epidote and muscovite. The rocks that are in structural contact with the Areado Unit at its bottom are quartzites, which do not have many hydrous phases and they seem not to be the water source. The shear zone that deforms the rocks was the path for the fluids to get access to the rocks.

#### IV.9 Concluding remarks

The Areado Granite-Migmatite Unit is a complex group of metamorphic rocks that represents the infrastructure of the Southern Brasilia Orogen. Its description and genesis are still in process of investigation, but even with several maps made between Alfenas and Areado, this unit was, until now, not well characterized. Some conclusions are summarized below:

The accessory mineralogy, the K-feldspar porphyroclast, and mainly field relationships suggest that the garnet meta-syenogranite is the protolith of the migmatites described here.

The presence of euhedral hornblende and the absence of selvedges at the border of the leucosome show that the main factor that controlled partial melting on these rocks is the water influx. The upper structural level minimum volume of partial melting is observed, and a major effect of metamorphism occurs on the lower unit instead. The source of the fluid is still unknown. A variation on the mineralogical composition of the biotite stromatic diatexite unit is controlled by the amount of water inserted in the system and deformation.

Structural data indicate the thrust faulting of Três Pontas-Varginha Nappe controls the mylonitic foliation of the area. Fold relationship from the biotite stromatic diatexite unit has at least two deformation stages. Tight folds with the main mylonitic foliation in the axial plane are older than harmonic open folds with the oblique axial plane concerning the main foliation. The correlation of this data with quartz fabric, which presents GBM and BLG recrystallization, suggests that both folds and main foliation were formed during a single deformational event associated with the high-pressure granulite thrust.

Applying quartz c-axis thermometer implied in two deformation temperature intervals, one that represents the metamorphic peak and a lower one. The lobated quartz localized in both migmatite units presents deformation temperatures between 766 and 892 °C, associated with the metamorphic peak in the hornblende-garnet metatexite and partial melt. Quartz grains formed by BLG have deformation temperatures between 613-714 °C, and this temperature range probably marks the stage at which the rock presents enough solid material to underwent solid-state deformation.

The application of TitaniQ, together with Zr-in-Titanite, did not produce consistent results for these rocks when 0.5 titanium activity was applied. The lack of saturation of titanium in the system, along with its resetting due to deformation, might be the issue. When a 0.2 TiO<sub>2</sub> activity is used, the metamorphic peak reaches 770-820 °C and 11.5-15 kbar for xenoblastic titanites, while idioblastic titanite indicates a cooling temperature of 667-708 °C.

Correlations between the TitaniQ, Zr-in-Titanite, and quartz c-axis thermometers suggest a progressive deformational-metamorphic pulse. High-temperature deformation on quartz and the formation of xenoblastic titanite crystals, are possibly close to the metamorphic peak and associated with partial melting of the rock. Generation of low-temperature quartz texture and idioblastic titanite crystals, along with mylonitization and folding represents a second event, with lower temperature. The metamorphic ages obtained for this are a low intercept from zircon (671-658 Ma) and titanite U-Pb ages of 621 Ma for the garnet meta-syenogranite unit, and zircon age of 604 Ma, and younger titanite age of557 Ma for the migmatites. These ages might represent the changes in the melting process, from an anhydrous to water-fluxed melting, or can resemble the cooling history of the unit. This metamorphic event is associated with the thrust of the Southern Brasilia Orogen over the Areado Granite-Migmatite Unit.

Crystallization age of 2.08 - 2.07 Ga from the garnet meta-syenogranite indicates that it is synchronous to the late stage of magmatic accretion from the Mineiro Belt, similar to what occurs in the Pouso Alegre Complex (weighted average of 2078.7 ± 6.7 Ma, Cioffi *et al.*, 2016). However, its high-grade metamorphism occurred during the Brasiliano Event. Thus, this unit cannot be associated with the Campos Gerais Complex, nor with the Andrelândia Nappe System, as it occurred in some older maps. It should be considered part of the Southern Brasilia Orogen, as part of its infrastructure.

Although most of the obtained data correlate with each other, caution must be taken prior to applying experimental thermometers. The titanium activity calculation is the central issue to use Ti-in-quartz and Zr-in-titanite on metamorphic rocks when rutile is not present. Although the literature suggests a small value for titanium activity, new ways to calculate it must be investigated.
Water activity seems to affect some results for GBM quartz analysis, causing an overestimation of high deformation temperature.

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## **Chapter V – Final Considerations**

The Areado Granite-Migmatite Unit is a complex block that had never had detailed research on it before this project to actually identify its geological framework and petrological evolution. In some maps, it was associated with the Campos Gerais complex (Cavalcante *et al.*, 1979; Artur, 1988). This work has set the correct classification of each rock inside the unit, along with the geological process that generated such a different partial melt rate inside the same region, and has contextualized the granite emplacement and the metamorphism in time.

Modal analysis shows that a big variation in the QAP proportion between the meta-syenogranite and the migmatites but the petrography and field relations clearly indicates that they are cogenetic. The answer of that matter was found in experimental petrology, where several experiments (*e.g.* Piwinskii, 1968b, 1973; Boettcher and Wyllie, 1968; Stern and Wyllie, 1981; Whitney, 1975) have got similar results when applying partial melting with high  $a_{H_2O}$  in the samples. That fact, associated with the hydrated peritectic minerals in all units and the diffusive contact between leucosome and the residuum points to the water-assisted melting as the main metamorphic process that affected these rocks. It is also interesting to point out that the presence of garnet as a peritectic mineral in both meta-syenogranite and metatexite and mostly absence in the diatexite suggests a variation in the amount of water that caused melting in each rock. The meta-syenogranite had low to no addition of water in the system, while the metatexite received a low amount and the diatexite was saturated on it.

The application of different thermobarometry methods was successful with some problems to consider. The quartz c-axis proved to be a useful tool to understand the thermal evolution of rocks, exhibiting the temperature difference on deformation zones. The main issue is the temperature calculation on rocks that have a high  $a_{H_2O}$ . As stated by Law (2014) and Faleiros *et al.* (2016), the increase of water in the system can cause an over estimative of the deformation temperatures, and this study also saw this event happening. It is necessary more research on the effect of water on quartz dislocation creep and a new calibration to avoid this issue. The use of TitaniQ and Zr-

in-titanite together to estimate the metamorphic peak of the Areado unit created some interesting results and can be indicated for this type of rock, especially when thermodynamic modeling failed, which was the case. However, a big pitfall can be the  $a_{TiO_2}$  estimative on metamorphic rocks since both thermobarometers depend on them. When the sample present rutile, it is safe to assume that the  $a_{TiO_2}$  is equivalent to one, but when it doesn't present the mineral, it becomes tricky to define the activity of the system. Hayden *et al.* (2008) suggested the use of 0.6 for rocks with granitic composition, but that assumption proved to be inefficient with the Areado unit. It is possible that the deformation and partial melt might change the Ti concentration on metamorphic quartz grains and change the activity as consequence.

Dating the migmatites permitted us to understand better the geological context of the Areado Unit. Crystallization ages of 2069.8  $\pm$  5 Ma proved that the protolith of the migmatites was generated at the last stages of the Mineiro Belt generation. Lower intercepts from zircon indicate a metamorphic and partial melting age between 671  $\pm$ 58 and 657  $\pm$  40 Ma and the melt crystallization ages of 622  $\pm$  7 Ma and 604  $\pm$  6 Ma shows that these rocks were reworked during the Brasilia Orogenic Event and from now on, this unit should be considered as part of the Southern Brasilia Orogen.

## **Chapter VI – References**

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Apendix

Sample AL	.FE – 24								
ррт	Crystal 1	Crystal 2	Crystal 3	Crystal 4	Crystal 5	Crystal 6	Crystal 7	Crystal 8	Crystal 9
Li6	2.21	3.13	<3.32	<1.69	2.68	<2.28	<1.08	<3.22	<1.19
Li7	1.45	2.35	<0.70	0.47	2.05	2.49	<0.21	<0.51	0.43
Na23	10.1	59.35	26.16	<6.73	<10.24	<10.54	<7.55	<18.91	<8.13
Al27	45.77	21.06	170	7.93	94.9	38.84	13.88	59.95	19.56
Si29	467434	467434	467434	467434	467434	467434	467434	467434	467434
Si30	392236	407811	503934	425029	384035	502665	595054	562864	373202
Ca42	321	<157	1720	<253	380	791	<255	<622	354
Ca43	<117	<70.27	<390	<108	<159	<165	<111	<290	<117
Ca44	1348	2148	3620	1106	1022	1629	2056	1093	1163
Ti49	2.78	3.04	4.27	4.73	2.63	2.38	3.54	3.49	2.63
Mn55	<0.23	<0.134	<0.78	<0.22	<0.31	<0.31	<0.23	<0.54	<0.229
Fe56	13.11	<2.07	289	8.61	11.11	11.06	3.92	<8.04	6.36
Fe57	<12.02	<7.13	227	<11.64	<17.25	<17.38	<11.39	<29.58	<12.14
Zn66	4.34	2.43	3.67	1.82	6.37	2.43	<0.64	3.11	2.68
Ga69	0.042	0.05	0.515	0.16	<0.075	<0.061	0.048	<0.130	<0.056
Ge70	0.44	0.63	0.97	0.55	0.5	0.27	1	0.76	0.58
Ga71	< 0.032	<0.019	0.391	0.4	<0.049	0.054	0.306	0.285	0.191
Ge72	0.56	0.63	3.26	1.04	<0.77	<0.73	0.86	2.02	0.73
Ge73	0.51	0.48	3.23	<0.312	0.66	0.59	1.33	2.38	1.01
Ge74	0.44	0.64	1.15	0.71	0.65	0.46	0.79	<0.59	0.4
Se77	<1.77	<1.21	<6.02	<1.89	<2.59	<2.90	<2.31	<3.46	<1.61
Rb85	0.043	0.03	0.097	<0.040	0.04	0.051	<0.034	<0.072	<0.044
Zr90	0.059	0.08	<0.063	<0.038	0.084	0.042	0.034	<0.064	<0.024
Ba138	0.174	0.366	0.84	<0.039	0.855	0.209	0.134	0.272	0.213
Pb208	0.111	0.056	0.207	0.06	0.105	0.149	0.101	0.241	0.079

Table A 1 - ICP-MS analysis on quartz grains in the hornblende-garnet metatexite unit

Sample CGE-03	D							
ррт	Crystal 1	Crystal 2a	Crystal 2b	Crystal 3	Crystal 4	Crystal 5	Crystal 6a	Crystal 6b
Li6	1.33	<0.37	<0.47	0.62	<0.53	<0.43	<0.62	<0.43
Li7	1.047	<0.088	<0.105	0.786	0.231	<0.089	0.22	0.144
Na23	35.05	11.17	24.04	40.34	9.45	18.8	40.27	22.35
Al27	48.81	46.54	38.55	62	7.23	26.65	51.71	50
Si29	467434	467434	467434	467434	467434	467434	467434	467434
Si30	479120	458431	531086	479427	476160	448245	438233	476920
Ca42	<20.63	<29.88	52.26	<28.69	<36.44	29.21	<39.23	<34.04
Ca43	18.29	82.33	26.54	43.25	<15.51	31.77	30.28	40.3
Ca44	1334	2149	1564	1223	1569	1594	1074	1489
Ti49	5.08	3.29	3.73	4.54	3.82	4.23	5.13	5.1
Mn55	0.451	0.447	0.215	0.391	<0.036	0.144	0.284	0.34
Fe56	38.04	17.93	12.83	50.62	9.62	28.44	49.98	42.01
Fe57	26.9	14.61	15.94	57.69	2.16	26.41	41.52	49.49
Zn66	1.74	1.83	0.67	1.47	0.3	0.669	1.11	1.53
Ga69	0.0524	0.042	0.219	0.0636	0.054	0.239	0.018	0.0602
Ge70	0.825	0.99	1.79	1.02	0.88	0.924	0.87	0.811
Ga71	0.0306	0.154	0.078	0.095	0.208	0.176	0.049	0.087
Ge72	0.827	1.22	1.035	0.854	1.48	1	1.01	1.14
Ge73	0.971	1.15	1.47	0.941	2.58	1.26	0.87	1.46
Ge74	0.854	0.762	0.977	0.996	1.016	0.985	0.914	0.956
Se77	<0.133	<0.21	0.54	<0.216	<0.26	0.16	<0.26	<0.21
Rb85	0.0605	0.05	0.0373	0.419	0.0206	0.0496	0.162	0.186
Zr90	0.0423	0.0258	410	0.265	0.0072	0.0139	0.0267	0.053
Ba138	0.574	1.165	0.234	0.605	0.0725	0.633	1.046	0.585
Pb208	0.1194	0.185	0.0625	0.413	0.0345	0.0966	0.154	0.174

Table A 2- ICP-MS analysis on quartz grains in the biotite stromatic diatexite unit

Sample ALFE-24								
ppm	Tit1	Tit2	Tit3	Tit4	Tit5	Tit6	Tit7	Tit5b
Li7	0.64	<0.46	<0.45	0.71	1.11	0.75	1.19	<0.56
Mg24	105	109	99.47	76.92	258	107	102	179
Al27	23430	26844	22877	22374	27004	23582	20831	22412
Si29	137283	151585	142098	135450	138056	134161	132170	133513
P31	132	176	132	113	88.42	125	123	59.46
Ca42	201150	212260	206025	201076	195883	206697	192695	204689
Ca44	192397	192397	192397	192397	192397	192397	192397	192397
Sc45	11.3	12.72	11.27	11.26	31.34	12.74	12.01	27.55
Ti46	157894	153169	160423	158903	150503	155139	169963	162284
Ti49	179707	175054	182191	185861	174690	177512	194157	190469
V51	5.23	4.19	5.63	4.09	10.85	5.37	5.08	13.05
Cr52	<0.33	0.78	0.42	0.37	1.97	0.45	0.51	2.25
Mn55	75.09	75.05	72.2	74.59	545	71.98	70.7	173
Fe56	6940	6639	7432	6450	15667	6820	7307	12317
Co59	<0.024	0.081	<0.021	0.048	0.091	0.048	<0.018	0.061
Sr88	50.83	63.33	59.06	75.15	23.04	56.04	56.5	34.9
Y89	2341	2409	1980	1321	13722	1733	2024	12640
Zr91	93.69	75.15	69.35	49.22	667	61.95	74.34	612
Nb93	3852	4159	3859	3831	3344	4032	4304	3761
Sn118	124	136	128	95.61	81.35	115	134	103
Sb121	<0.048	<0.050	<0.048	<0.049	<0.050	<0.056	<0.048	<0.059
Ba137	0.782	1.99	0.381	0.439	0.25	1.64	0.599	0.59
La139	40.52	112	23.95	17.84	595	38.6	32.38	232
Ce140	222	412	182	115	3658	158	177	1645
Pr141	65.31	98.85	57.27	34.61	881	49.91	56.06	471
Nd143	545	729	510	291	5920	418	476	3701
Sm147	419	502	410	246	2431	333	361	1886
Eu151	116	122	105	74.62	141	91.96	103	130
Gd157	687	786	650	418	2581	529	623	2247
Tb159	143	158	132	86.89	487	104	130	459
Dy161	754	843	710	436	2951	528	652	2816
Ho165	104	116	95.36	57.65	577	68.14	89.03	546
Er166	177	211	152	91.44	1421	112	144	1284
Tm169	16.64	21.69	13.36	8.53	189	10.47	12.82	175

Table A 3 - ICP-MS analysis on titanite grains at the hornblende-garnet metatexite unit

Yb173	78.72	107	57.75	39.77	1167	48.03	56.68	1137
Lu175	6.87	9.04	4.33	3.18	122	3.92	4.34	123
Hf178	7.32	7.93	6	4.52	47.42	5.6	6.36	50.91
Ta181	116	236	133	121	325	142	177	346
Pb206	12.58	12.53	10.1	8.18	97.97	9.3	10.84	41.33
Pb207	4.38	4.24	3.75	4.46	14.57	4.4	4.54	7.05
Pb208	3.98	4.13	3.53	4.12	19.28	4.37	4.58	7.22
Th232	2.38	3.6	2.08	1.257	130	1.67	2.33	37.27
U238	33.36	29.29	27.57	17.38	106	22.86	27.37	63.74

Sample CGE-09D								
Ppm	Crystal 1	Crystal 2	Crystal 3	Crystal 4	Crystal 5	Crystal 6	Crystal 6b	Crystal 7
Li7	<0.23	<0.27	<0.36	0.37	<0.23	8.31	12	0.59
Mg24	24.26	34.76	26.56	34.92	26.84	345	434	43.02
AI27	18141	15548	18233	17807	19666	171735	181193	16815
Si29	85647	92892	88410	92995	81112	217488	209644	113994
P31	32.15	71.58	37.5	56.14	46.34	47.29	40.08	63.61
Ca42	189881	183834	171540	169366	112647	117884	121888	157326
Ca44	192397	192397	192397	192397	192397	192397	192397	192397
Sc45	6.68	6.38	8.72	8.69	7.05	119	108	6.17
Ti46	145776	172651	157595	164787	169399	5519	7655	162844
Ti49	169049	196916	175065	175414	191860	6105	8296	192368
V51	1.416	1.53	1.62	2.23	1.62	4.64	8.71	2.18
Cr52	0.281	0.214	<0.21	0.64	0.804	1.32	1.19	0.47
Mn55	47.71	66.58	49.52	69.81	63.95	863	804	88.83
Fe56	6977	9089	7376	6972	8939	180043	214177	8222
Co59	0.0151	0.0303	<0.015	0.052	0.0079	2.53	3.69	0.045
Sr88	73.59	47.4	93.48	60.54	56.74	4009	2448	72.04
Y89	1887	1812	1612	2518	2155	3923	3821	2253
Zr91	25.37	73.23	33.19	68.67	54.14	226	114	77.59
Nb93	4751	3562	4745	4224	3858	3.95	2.19	3616
Sn118	38.59	37.91	48.94	43.3	30.15	9.64	9.18	41.7
Sb121	<0.023	<0.028	0.057	<0.036	<0.025	0.079	0.093	<0.046
Ba137	0.391	0.882	0.479	2.28	0.227	456	276	1.51
La139	14.1	64.81	20.53	66.82	36.21	62030	88288	53.8
Ce140	123	752	175	673	241	128617	172797	487
Pr141	27.66	104	40.4	107	66.37	15315	18841	112
Nd143	211	796	321	742	523	58789	65801	902
Sm147	156	459	200	413	324	6474	6071	542
Eu151	39.93	116	45.35	83.84	82.13	634	477	121
Gd157	243	541	284	529	475	3377	3092	693
Tb159	52.7	86.9	59.9	99.64	86.37	312	294	125
Dy161	297	383	326	510	413	1031	1007	618
Ho165	41.69	45.57	46.43	72.91	50.97	103	101	81.51
Er166	68.52	70.96	78.5	136	77.96	153	144	134
Tm169	6.05	6.17	7.07	13.84	6.62	10.48	8.71	12.43

Table A 4 - ICP-MS analysis on titanite grains at the hornblende-garnet metatexite unit (Sample CGE-09D)

Yb173	29.59	29.27	31.89	70.64	30.47	44.95	36.27	58.74
Lu175	2.27	2.23	2.34	5.94	2.51	4.99	3.8	4.62
Hf178	2.38	6.16	3.35	6.18	4.2	3.84	3.13	6.24
Ta181	87.48	69.41	140	116	60.76	0.199	0.172	131
Pb206	6.53	11.69	8.67	9.62	8.55	667	329	11.75
Pb207	4.94	4.12	6.66	5.33	4.03	593	277	6.03
Pb208	5.04	3.83	6.48	5.62	4.16	775	474	5.42
Th232	0.415	3.43	1.174	3.72	1.92	5055	5898	2.69
U238	6.6	26.35	8.08	15.52	14.72	255	181	21.52

grain/spot			Isotopic	ratios			Th	U	Th/U		AGES	(in Ma)		Concordance <sup>1</sup>
position	<sup>207</sup> Pb/ <sup>235</sup> U	σ	<sup>206</sup> Pb/ <sup>238</sup> U	σ	<sup>207</sup> Pb/ <sup>206</sup> Pb	σ	ppm	ppm		<sup>206</sup> Pb/ <sup>238</sup> U	σ	<sup>207</sup> Pb/ <sup>206</sup> Pb	Σ	%
23.1	6.7012	0.0593	0.3789	0.0043	0.1281	0.0002	188.4	579.0	0.325	2.071	0.020	2.070	0.003	100
19.2	6.6553	0.0636	0.3758	0.0044	0.1275	0.0004	58.8	65.9	0.893	2.056	0.021	2.061	0.006	100
15.2	6.6263	0.0378	0.3757	0.0026	0.1278	0.0007	72.3	50.5	1.431	2.056	0.012	2.066	0.010	100
13.2	6.6824	0.0434	0.3769	0.0028	0.1281	0.0009	25.2	20.9	1.201	2.062	0.013	2.069	0.012	100
13.1	6.7001	0.0273	0.3796	0.0023	0.1285	0.0007	58.9	37.0	1.593	2.075	0.011	2.075	0.009	100
12.1	6.7026	0.0337	0.3796	0.0028	0.1288	0.0006	68.9	41.6	1.655	2.074	0.013	2.079	0.009	100
24.1	6.8258	0.0621	0.3815	0.0044	0.1285	0.0004	72.0	45.5	1.582	2.083	0.021	2.075	0.006	100
19.1	6.8414	0.0676	0.3821	0.0045	0.1286	0.0005	94.9	57.0	1.6647	2.086	0.021	2.076	0.007	100
25.1	6.7500	0.0613	0.3774	0.0043	0.1278	0.0003	201.3	131.0	1.537	2.064	0.020	2.066	0.004	100
1.1	6.7687	0.2033	0.3832	0.0100	0.1293	0.0008	122.2	124.8	0.9795	2.091	0.047	2.086	0.010	100
2.1	6.7054	0.2010	0.3747	0.0098	0.1273	0.0008	15.2	26.6	0.569	2.052	0.046	2.059	0.010	100
3.1	6.7439	0.2084	0.3791	0.0103	0.1289	0.0007	185.8	517.3	0.359	2.072	0.048	2.080	0.009	100
4.1	6.6064	0.1978	0.3754	0.0098	0.1275	0.0007	169.3	229.5	0.738	2.055	0.046	2.062	0.009	100
6.1	6.4553	0.1945	0.3751	0.0098	0.1263	0.0008	66.8	47.4	1.408	2.053	0.046	2.045	0.011	100
7.1	6.5027	0.1941	0.3736	0.0097	0.1269	0.0008	46.2	47.1	0.981	2.046	0.046	2.054	0.010	100
9.1	6.7643	0.2016	0.3815	0.0099	0.1282	0.0007	105.2	137.8	0.763	2.083	0.046	2.071	0.010	101
11.1	6.4829	0.0298	0.3750	0.0023	0.1267	0.0007	41.3	32.8	1.2586	2.053	0.011	2.051	0.010	100
16.2	6.4674	0.0268	0.3722	0.0024	0.1261	0.0006	176.1	208.1	0.846	2.040	0.011	2.042	0.008	100
18.1	6.3536	0.0372	0.3690	0.0028	0.1270	0.0007	123.5	104.5	1.182	2.025	0.013	2.055	0.009	99
19.3	6.4567	0.0745	0.3724	0.0049	0.1264	0.0003	126.2	533.2	0.237	2.041	0.023	2.047	0.004	100
21.1	6.5165	0.0658	0.3733	0.0046	0.1269	0.0004	49.3	56.7	0.870	2.045	0.022	2.054	0.006	100
4.2	6.3345	0.1889	0.3571	0.0093	0.1247	0.0007	168.6	800.9	0.211	1.969	0.044	2.023	0.010	97
5.1	6.0534	0.1802	0.3456	0.0090	0.1259	0.0007	150.1	879.8	0.171	1.913	0.043	2.039	0.009	94
26.1	6.2686	0.0578	0.3663	0.0042	0.1241	0.0005	48.9	44.4	1.101	2.012	0.020	2.015	0.007	100
8.1	7.0235	0.2211	0.3903	0.0107	0.1325	0.0009	84.8	583.9	0.145	2.124	0.049	2.128	0.011	100
17.1	6.8375	0.0258	0.3815	0.0025	0.1292	0.0007	62.5	49.3	1.267	2.083	0.012	2.085	0.009	100
22.1	6.9573	0.0724	0.3851	0.0047	0.1301	0.0003	66.6	59.2	1.126	2.100	0.022	2.096	0.005	100
15.1	6.8399	0.0274	0.3806	0.0024	0.1294	0.0006	94.5	62.9	1.503	2.079	0.011	2.087	0.008	100
6.2	5.7253	0.1797	0.3360	0.0094	0.1243	0.0007	183.1	227.7	0.804	1.867	0.045	2.018	0.009	93
21.2	5.6802	0.0511	0.3288	0.0038	0.1244	0.0003	190.3	1158.5	0.164	1.833	0.018	2.018	0.004	91
10.1	5.7027	0.1707	0.3311	0.0086	0.1248	0.0007	84.7	103.2	0.821	1.844	0.042	2.025	0.010	91
16.1	7.1269	0.0344	0.3769	0.0026	0.1282	0.0007	113.0	73.0	1.547	2.062	0.012	2.072	0.009	100
14.1	6.9466	0.0281	0.3855	0.0023	0.1298	0.0006	152.2	99.5	1.529	2.102	0.011	2.092	0.008	100
20.1	7.3796	0.0808	0.4124	0.0051	0.1302	0.0005	39.3	35.8	1.099	2.226	0.023	2.098	0.007	106

Table A 5 - Summary of LA-ICP-MS U-Pb data on zircons

1 – Concordance calculated for <sup>206</sup>Pb/<sup>238</sup>U and <sup>207</sup>Pb/<sup>206</sup>Pb ages

Table /	46 -	Summary	of	SHRIMP	data	on	zircons
rabic /	- 01	Gunnary	01		uata	011	21100113

		(1)	(2)	(3)	(1)	(1)	(2)	(3)	% _			(1)				(	(2)		- 1	(*)	
206P 4	corr 4-cor	206Pb	206Pb	206Pb	207Pb	208Pb	208Pb	207Pb	Dis- 7cor	r	(1)	(1)	(1)		(1) ε	err (3)	(3)	(	3)	(3)	
204Pb 207Pb 208Pb b % ppm ppm F	om ppm	232Th /238U	/23811	/23811	/206Ph	/232Th	/232Th	/206Ph	cor- 208P	5 C	238U	207Pb	207Pb	2	06Pb c	or 238U	207Pb	207	'Pb <sup>'</sup>	206P	b err.
Spot /206Pb ±% /206Pb ±% /206Pb ±% /238U ±% 206Pb, U Th 20	SPb 208Pb	/238U ±% Age	Ace	Age	Age	Age	Age	Age	dant /2321	'h ±%	/206Pb	±% /206Pb	±% /235U	±% /	238U ±%	r /206Pb	±% /206Pb	±% /23	35U ±9	% /238	U ±% corr.
		Age	Aye	Aye	Aye	Age	Aye	Age													
CGE_03C-1.1 5.0E-4 13 0.113 0.4 0.082 1.2 0.47 1.2 0.84 444 86	72 4.7	0.20 0.34 1117 ±1	1 1080 ±11	1115 ±11	1728 ±17 119	99 ±46	-259 ±40	1697 ±11	+38 -0.01	3 -15.5	5.3	1.1 0.106	1.0 2.8	1.4	0.189 1.1 0	.8 5.3	1.1 0.1040	0.6 2	.7 1.	.2 0.18	9 1.1 0.9
CGE_03C-2.1 6.7E-4 11 0.109 0.4 0.079 2.4 0.40 1.5 1.15 478 152	67 3.7	0.33 0.27 976 ±1	0 944 ±10	995 ±10	1624 ±21 54	7 ±30	-211 ±29	1910 ±12	+43 -0.01	0 -13.9	6.1	1.1 0.100	1.1 2.3	1.6	0.163 1.1 0	.7 6.0	1.1 0.1169	0.7 2	.7 1.	.1 0.16	7 1.1 0.8
CGE_03C-3.1 2.7E-4 19 0.121 0.5 0.074 2.8 0.68 0.4 0.44 189 36	14 2.9	0.20 0.51 1539 ±1	6 1501 ±17	1535 ±17	1923 ±14 173	31 ±69	72.3 ±94	1878 ±14	+22 0.00	4 130.7	3.7	1.2 0.118	0.8 4.4	1.4	0.270 1.2 0	.8 3.7	1.2 0.1149	0.8 4	.3 1.	.3 0.26	.9 1.2 0.8
CGE_03C-4.1 9.5E-4 20 0.128 0.9 0.141 1.0 0.57 0.8 1.56 332 148	68 7.4	0.46 0.65 1373 ±1	4 1331 ±14	1389 ±15	1880 ±45 110	06 ±70	346 ±39	2038 ±18	+30 0.01	7 11.5	4.2	1.1 0.115	2.5 3.8	2.7	0.237 1.1 0	.4 4.2	1.2 0.1257	1.0 4	.2 1.	.5 0.24	1 1.2 0.7
CGE 03C-5.1 1.1E-3 24 0.075 6.3 0.031 29.9 0.25 1.6 1.90 88 0	7 -0.1	0.01 4.48 599 +8	3 600 +9	602 +8	587 +222 -35	45 -4395.09	-3158 +7428	728 +155	-2 -0.14	5 -217.3	10.3	1.4 0.060	10.2 0.8	10.3	0.097 1.4 0	1 10.2	1.4 0.0636	7.3 0	9 7	7 0.09	8 1.4 0.3
CGE 03C-61 3 0E-4 20 0122 0.8 0107 13 054 14 049 265 85	5 53	0.33 0.36 1301 +1	4 1345 +15	1301 +15	1025 +20 13	77 ±38	223 +51	1030 +17	+31 0.01	1 22.0	12	1.1 0.118	11 30	1.6	0.241 11 0	7 42	1.2 0.1183	0.0 3	0 1	4 0.24	1 12 07
	27 222	0.88 0.34 1004 +1	0 1092 122	1002 +22	2066 10 200	05 ±27	1900 +40	2057 +12	14 0.00	22.0	2.0	1.1 0.129	0.5 6.4	1.0	0.241 1.1 0	0 20	1.2 0.1100	0.0 6	2 1	2 0.24	2 12 0.7
	51 ZZ.Z	0.00 0.24 1994 ±1	5 1503 ±22	1333 122	2000 ±0 200	00 100	620 ±40	2037 ±13	++ 0.05	2.2	2.0	1.1 0.120	10.5 0.4	1.2	0.305 1.1 0	2.0	1.3 0.1270	10.0 0	0 1	E 0.30	2 1.3 0.0
	0.4	0.31 0.35 1716 ±1	/ 16// ±19	1716 ±16	2033 ±21 1/0	09 ±03	030 ±96	2035 ±21	+16 0.03	2 15.6	3.3	1.1 0.125	1.2 5.3	1.0	0.305 1.1 0	.7 3.3	1.2 0.1255	1.2 5		.5 0.30	5 1.2 0.6
CGE_03C-9.1 8.7E-4 21 0.076 4.6 0.036 21.1 0.22 0.6 1.55 151 1	13 0.0	0.01 2.94 604 ±/	602 ±8	604 ±8	/18 ±150 83	9 ±1731	-1373 ±3307	705 ±116	+17 -0.06	6 -232.7	10.2	1.3 0.063	7.1 0.9	7.2	0.098 1.3 0	.2 10.2	1.3 0.0629	5.5 0	.9 5.	.8 0.09	8 1.3 0.3
CGE_03C-10.1 1.5E-4 21 0.063 1.2 0.009 15.5 0.23 1.1 0.27 849 8	72 0.3	0.01 1.13 607 ±6	607 ±6	607 ±6	627 ±33 80	)8 ±280	500 ±500	614 ±30	+3 0.02	5 101.1	10.1	1.1 0.061	1.5 0.8	1.8	0.099 1.1 0	.6 10.1	1.1 0.0603	1.4 C	.8 1.	.7 0.09	9 1.1 0.6
CF_03C-10.2 6.5E-4 12 0.074 1.4 0.027 10.0 0.23 2.4 1.14 758 21	68 0.2	0.03 0.70 644 ±7	7 641 ±7	646 ±7	776 ±49 24	1 ±219	-546 ±329	843 ±33	+18 -0.02	7 -59.4	9.5	1.1 0.065	2.3 0.9	2.6	0.105 1.1 0	.4 9.5	1.1 0.0672	1.6 1	.0 1.	.8 0.10	5 1.1 0.5
CE_03C-11.1 7.6E-4 14 0.144 0.5 0.132 1.5 0.77 0.5 1.17 128 46	38 4.1	0.37 0.50 1910 ±2	1 1875 ±23	1910 ±22	2146 ±22 19'	18 ±76	1054 ±85	2144 ±14	+13 0.05	4 8.3	2.9	1.3 0.134	1.2 6.4	1.8	0.345 1.3 0	.7 2.9	1.3 0.1334	0.8 6	.3 1.	.4 0.34	5 1.3 0.8
GE 03C-12.1 7.3E-4 22 0.071 4.4 0.027 18.2 0.19 0.5 1.29 213 1	18 0.0	0.01 2.87 605 ±7	7 604 ±7	605 ±7	633 ±139 -7	0 -1864.49	-721 ±3183	663 ±107	+5 -0.03	5 -433.3	10.2	1.2 0.061	6.4 0.8	6.5	0.098 1.2 0	.2 10.2	1.2 0.0617	5.0 C	.8 5.	.2 0.09	8 1.2 0.3
GE 03C-13 1 5E-4 20 0.128 0.6 0.160 0.9 0.83 1.6 0.23 245 124	11 5	0.52 0.31 1927 +1	9 1911 +22	1924 +20	2040 +12 198	87 +31	1705 +59	2012 +14	+6 0.08	8 36	29	1.1 0.126	07 60	13	0.348 1.1 0	9 29	1.2 0.1238	0.8 5	9 1	3 0.34	8 12 08
GE 03C-14 ( 8.0E-4 18 0.072 3.8 0.025 28.9 0.25 11 1.42 192 1	6 -0.1	0.01 2.65 602 +7	7 602 ±7	604 +7	600 +126 -13	81 -1733.08	-13/1 +30/9	60/ ±00	-0 -0.06	1 -210 0	10.2	1.2 0.060	5.8 0.8	6.0	0.008 12 0	2 10 2	1.2 0.0626	42 0	8 /	5 0.00	8 12 0.0
	-0.1	0.01 2.03 002 11	4 4440 .04	1102 .00	4000 120 13	07 .05	-1041 10040	1700 .24	-0 -0.00	213.3 F 109.1	F.0	1.2 0.000	10 24	0.0	0.000 1.2 0	7 50	2.0 0.0020	4.0 0	0 7	C 0.00	1 20 07
JGE_03C-15. 3.5E-4 16 0.115 1.6 0.090 4.0 0.46 2.7 0.59 319 76	5 4.4	0.25 0.67 1164 ±2	1 1142 ±21	1103 ±22	1000 ±34 124	27 ±05	-99 ±106	1790 ±34	+30 -0.00	5 -106.1	5.0	1.9 0.111	1.9 3.1	2.1	0.202 1.9 0	.7 5.0	2.0 0.1095	1.9 3	.0 2.	.6 0.20	1 2.0 0.7
JGE_03C-16.1 3.3E-4 19 0.135 1.0 0.104 3.0 1.02 0.6 0.49 106 33	35 3.3	0.32 0.55 2081 ±2	3 2076 ±27	2080 ±24	2108 ±22 21	18 ±81	1986 ±158	2099 ±21	+2 0.10	3 8.4	2.6	1.3 0.131	1.2 6.9	1.8	0.381 1.3 0	.7 2.6	1.4 0.1301	1.2 6	.8 1.	.7 0.38	1 1.4 0.7
CGE_05Q-1.1 6.1E-4 19 0.134 0.6 0.174 1.5 0.65 2.8 0.92 111 49	36 5.7	0.45 0.52 2081 ±2	4 2089 ±28	2061 ±25	2038 ±25 246	63 ±78	2607 ±103	1886 ±19	-2 0.13	8 4.2	2.6	1.3 0.126	1.4 6.6	1.9	0.381 1.3 0	.7 2.7	1.4 0.1154	1.1 6	.0 1.	.5 0.37	7 1.4 0.7
CGE_05Q-2.1 8.2E-4 22 0.106 1.0 0.062 11.6 0.40 1.5 1.39 171 11	26 0.9	0.07 1.16 1064 ±1	3 1039 ±13	1058 ±13	1528 ±55 174	44 ±397	-1219 ±561	1424 ±29	+33 -0.05	9 -44.6	5.6	1.3 0.095	2.9 2.4	3.2	0.179 1.3 0	.4 5.6	1.4 0.0899	1.5 2	.2 1.	.7 0.17	8 1.4 0.5
CGE_05Q-3.1 1.5E-4 24 0.129 0.7 0.094 1.1 0.93 1.1 0.23 238 67	71 6.4	0.29 0.38 1922 ±1	9 1902 ±21	1918 ±20	2055 ±14 203	30 ±42	1430 ±102	2028 ±14	+8 0.07	3 7.4	2.9	1.1 0.127	0.8 6.1	1.4	0.347 1.1 0	.8 2.9	1.2 0.1249	0.8 6	.0 1.	.3 0.34	6 1.2 0.8
CGE_05Q-3.2 2.9E-4 18 0.131 0.4 0.135 1.9 0.90 1.4 0.44 192 79	59 7.5	0.43 0.37 1972 ±2	0 1959 ±23	1969 ±21	2058 ±12 203	33 ±52	1756 ±78	2036 ±14	+5 0.09	1 4.6	2.8	1.2 0.127	0.7 6.3	1.4	0.358 1.2 0	.9 2.8	1.2 0.1255	0.8 6	.2 1.	.2 0.35	7 1.2 0.8
CGE 05Q-4.1 3.1E-4 18 0.132 0.5 0.097 3.7 0.89 1.7 0.48 149 40	14 3.8	0.28 0.51 1899 ±2	0 1873 ±23	1893 ±21	2072 ±13 206	68 ±88	1258 ±115	2031 ±17	+10 0.06	4 9.5	2.9	1.2 0.128	0.8 6.0	1.4	0.343 1.2 0	.8 2.9	1.3 0.1251	0.9 5	.9 1.	.3 0.34	1 1.3 0.7
CGE 050-51 28E-4 19 0127 0.5 0.099 1.2 0.93 0.8 0.43 200 60	56 51	0.31 0.41 1820 +1	9 1796 +21	1820 +19	2004 +13 184	43 +47	1150 +76	1997 +11	+10 0.05	9 68	3.1	1.2 0.123	07 55	14	0.326 1.2 0	8 31	1.2 0.1228	0.6 5	5 1	2 0.32	6 12 09
GE 050-61 1 8E-4 20 0.130 0.4 0.103 1.2 0.85 0.6 0.28 216 71	6 61	0.34 0.39 1953 ±2	0 1036 ±22	105/ ±20	2065 ±10 10	50 ±10	1/07 ±85	2067 ±10	+6 0.07	7 5 9	2.8	1.2 0.128	0.6 6.2	13	0354 12 0	0 28	1.2 0.1227	0.6 6	2 1	2 0.35	1 12 0.0
	0 0.4	0.04 1.56 676 10	0 1330 122	672 +0	2005 110 130	00 +510	249 +947	770 +170	10 0.01	2 2426	2.0	1.4 0.060	0.0 0.2	0.0	0.111 1.4 0	1 01	1.6 0.0661	9.1 1	0 0	4 0.11	0 15 0.3
	0.2	0.04 1.50 070 ±	, 071 ±10	073 ±9	004 1200 110	09 ±319	240 1047	779 ±170	+23 0.01	2 343.0	9.0	1.4 0.009	9.7 1.0	5.0	0.111 1.4 0		1.5 0.0051	0.1 1	.0 0.	.4 0.11	0 1.0 0.0
JGE_05Q-7.1 1.2E-4 26 0.129 0.4 0.119 1.2 0.89 0.8 0.17 169 65	5 6.3	0.40 0.39 2060 ±2	1 2059 ±24	2058 ±22	2066 ±9 205	98 ±39	2076 ±92	2053 ±11	+0 0.10	5 4.7	2.7	1.2 0.128	0.5 6.6	1.3	0.376 1.2 0	.9 2.7	1.2 0.1267	0.6 6	.6 1.	.2 0.37	6 1.2 0.9
GE_05Q-8.1 1.6E-3 21 0.083 6.4 0.057 17.6 0.27 2.3 2.81 119 3	1 0.0	0.03 2.01 646 ±	9 646 ±9	649 ±9	619 ±261 -22	23 -1072.97	-56 ±1453	/5/ ±1/2	-4 -0.00	3 -2586.1	9.5	1.5 0.060	12.1 0.9	12.2	0.105 1.5 0	0.1 9.4	1.5 0.0645	8.2 0	.9 8.	.5 0.10	6 1.5 0.3
CGE_05Q-9.1 6.0E-4 20 0.118 1.5 0.126 3.6 0.63 0.6 0.99 102 25	20 2.1	0.25 1.22 1312 ±1	5 1275 ±16	1295 ±16	1789 ±41 18'	16 ±101	679 ±109	1575 ±37	+29 0.03	4 16.3	4.4	1.3 0.109	2.3 3.4	2.6	0.226 1.3 0	.5 4.5	1.4 0.0974	2.0 3	.0 2.	.3 0.22	.2 1.4 0.5
CF_05Q-10. 2.8E-4 19 0.125 1.0 0.077 1.8 0.67 0.9 0.44 175 35	14 3.0	0.21 0.51 1671 ±1	7 1637 ±19	1666 ±18	1969 ±21 185	58 ±62	449 ±127	1928 ±20	+17 0.02	2 28.5	3.4	1.2 0.121	1.2 4.9	1.7	0.296 1.2 0	.7 3.4	1.2 0.1181	1.1 4	.8 1.	.6 0.29	5 1.2 0.7
CE_05Q-11.' 4.3E-4 20 0.135 1.1 0.173 3.7 0.95 1.6 0.65 67 36	21 3.4	0.56 0.52 2032 ±2	5 2021 ±28	2032 ±27	2094 ±26 202	26 ±85	1859 ±110	2097 ±28	+3 0.09	6.2	2.7	1.4 0.130	1.5 6.6	2.0	0.370 1.4 0	.7 2.7	1.5 0.1299	1.6 6	.6 1.	.9 0.37	1 1.5 0.6
GE 05Q-12. 2.3E-4 20 0.132 1.1 0.161 2.3 0.94 1.2 0.35 122 61	38 5.8	0.52 0.39 1971 ±2	1 1953 ±24	1966 ±22	2084 ±21 203	39 ±55	1736 ±88	2054 ±23	+6 0.09	0 5.3	2.8	1.2 0.129	1.2 6.4	1.7	0.358 1.2 0	.7 2.8	1.3 0.1268	1.3 6	.2 1.	.7 0.35	7 1.3 0.6
GE 05Q-13, 2.5E-4 18 0.130 0.4 0.124 1.9 0.96 1.0 0.38 169 65	53 6.2	0.40 0.38 2017 +2	0 2011 +24	2016 +21	2057 +12 204	40 +52	1898 +89	2049 +14	+2 0.09	8 4.9	2.7	1.2 0.127	0.7 6.4	1.4	0.367 1.2 0	9 2.7	1.2 0.1264	0.8 6	4 1	2 0.36	7 1.2 0.8
CGE 25-11 1.0E-2 3 0.244 0.4 0.511 1.1 0.58 0.4 17.50 359 48	3 87	0.14 0.53 1030 ±1	3 005 ±11	976 +15	1670 +08 37	37 +306	2031 +158	365 +50	+41 0.10	6 82	5.8	1.3 0.103	53 24	5.5	0 173 13 0	2 61	1.6 0.0539	22 1	2 2	4 0.16	3 16 04
	26 10.1	1 01 0 25 1040 +2	4 1024 +25	1060 +25	2047 172 19	60 +01	1729 +62	2120 +42	16 0.00	0.2	2.0	1.0 0.100	41 61	4.4	0.252 1.4 0	2 20	1.6 0.0000	2.2 6	1 2	0 0.10	5 1.0 0.4
	10.1	1.01 0.33 1949 ±2	4 1934 ±23	1900 ±23	2047 173 100	05 151	1730 ±02	2120 142	+0 0.09	3 3.7	2.0	1.4 0.120	4.1 0.1	4.4	0.353 1.4 0	2.0	1.5 0.1317	2.4 0	4 2	5 0.33	3 1.5 0.5
CGE_25-3.1 4.2E-4 19 0.155 0.9 0.209 2.0 0.92 1.5 0.04 140 117	4 11.3	0.66 1.14 1993 ±2	2 1962 ±25	1965 ±25	2056 ±23 200	04 ±55	1957 ±00	2003 ±30	+4 0.10	2 3.0	2.0	1.3 0.127	1.3 0.3	1.0	0.362 1.3 0	./ 2.0	1.5 0.1232	2.0 6		.5 0.30	1 1.5 0.6
CGE_25-4.1 3.9E-3 14 0.151 7.2 0.253 11.4 0.25 1.4 6.87 366 45	39 4.5	0.13 0.56 756 ±1	2 728 ±14	730 ±15	1552 ±279 214	49 ±466	590 ±/64	822 ±353	+54 0.03	J 131.3	8.0	1.7 0.096	14.9 1.7	15.0	0.124 1.7 0	.1 8.3	2.2 0.0665	16.9 1	.1 18	3.1 0.12	0 2.2 0.6
CGE_25-9.1 3.7E-4 17 0.091 1.0 0.054 1.8 0.32 1.3 0.64 505 75	53 2.6	0.15 0.64 879 ±1	2 861 ±12	881 ±13	1326 ±29 76	64 ±47	-99 ±55	1379 ±21	+36 -0.00	5 -55.4	6.8	1.5 0.085	1.5 1.7	2.1	0.146 1.5 0	.7 6.8	1.5 0.0879	1.1 1	.8 1.	.8 0.14	7 1.5 0.8
CGE_25-10.1 8.1E-4 27 0.144 3.2 0.296 3.6 0.86 1.1 1.20 25 22	8 2.3	0.89 0.77 2117 ±3	6 2111 ±43	2100 ±41	2149 ±72 226	66 ±108	2207 ±160	2040 ±84	+2 0.11	5 7.7	2.6	2.0 0.134	4.1 7.2	4.6	0.389 2.0 0	.4 2.6	2.3 0.1258	4.8 6	.7 5.	.4 0.38	.5 2.3 0.5
CGE_25-11.1 6.6E-5 25 0.119 0.8 0.081 1.1 0.55 1.4 0.11 543 124	26 10.0	0.24 0.30 1542 ±1	5 1504 ±16	1536 ±15	1923 ±15 174	43 ±28	387 ±77	1866 ±16	+22 0.01	9 20.1	3.7	1.1 0.118	0.8 4.4	1.4	0.270 1.1 0	.8 3.7	1.1 0.1141	0.9 4	.2 1.	.4 0.26	9 1.1 0.8
CGE_25-12.1 6.2E-3 11 0.150 1.2 0.404 2.0 0.14 2.7 10.95 355 46	6.0	0.13 0.86 614 ±1	1 614 ±8	571 ±9	604 ±407 272	22 ±383	2732 ±116		-2 0.14	5 4.5	10.0	1.9 0.060	18.8 0.8	18.9	0.100 1.9 0	.1 10.8	1.7 -0.0048	46.3 -0	).1 -45	5.7 0.09	3 1.7 0.3
CGE 25-13.1 1.6E-4 21 0.128 1.0 0.154 2.6 0.78 0.7 0.24 227 103	57 10.1	0.47 0.33 1897 +3	1 1875 +35	1886 +33	2044 +20 208	82 +66	1685 +109	1964 +27	+8 0.08	7 6.7	2.9	1.9 0.126	1.1 5.9	2.2	0.342 1.9 0	9 2.9	2.0 0.1205	1.5 5	6 2	2 0.34	0 2.0 0.7
CGE 25-14.1 1.5E-4 18 0.129 1.0 0.186 0.7 0.70 2.2 0.22 400 226	19 21.8	0.58 0.92 1912 +1	8 1890 +21	1900 +20	2056 +18 206	67 +33	1747 +61	1972 +22	+8 0.09	0 3.7	2.9	1.1 0.127	1.0 6.0	1.5	0.345 1.1 0	7 2.9	1.2 0.1211	1.2 5	7 1	6 0.34	3 12 07
CGE 25-15.1 5.6E-4 24 0.131 1.1 0.280 1.2 0.78 0.6 0.97 99 90	26 69	0.04 0.40 1805 +2	2 1870 121	1002 ±25	2004 +33 191	38 ±45	1603 +50	2055	+6 0.00	7 31	2.0	1.4 0.122	10 59	23	0342 14 0	6 20	15 0 1260	16 6	0 2	1 0.24	3 15 06
CCE 25-10.1 0.0E-4 24 0.101 1.1 0.200 1.2 0.70 0.0 0.07 00 00	57 144	0.05 0.50 1000 ±2	∠ 10/3 ±23 0 1920 ±24	1962 123	2004 200 100	50 ±+0	1620 ±20	2000 ±20	+0 0.00	1 3.1	2.5	1.7 0.123	07 57	2.5	0.0+2 1.4 0	0 2.9	1.0 0.1209	0.0 5	0 4	4 0.34	5 1.0 0.0
002_2010.1 2.72 7 22 0.120 0.4 0.200 0.0 0.01 1.7 0.30 190 104	// 14.1	0.00 0.00 1002 ±1	5 1055 ±21	1002 121	2022 IIJ 10	57 ±30	1023 130	2020 ±17	+3 U.UO	τ 2.3	3.0	1.2 0.120	0.7 0.7	1.4	0.000 1.2 0	3.0	1.3 0.1240	0.9 5	1.	.+ 0.33	0 1.0 0.0

ate the common and radiogenic portions, respectively.

ale the common and adapting pointing, respectively, above errors but required when comparing data free mounts), corrected using measured 204Pb, ming 206Pb/238U-207Pb/235U age-concordance ming 206Pb/238U-208Pb/232Th age-concordance

Table A7 - Summary of SHRIMP data on titanite crystals

										4-corr	4-corr			(1) 206Pb	(2) 206Pb	(3)	<sup>th</sup>	(1) 207Pb	2	(1)	(2)	) Ph	(3) 207Pb	% Dis-	7corr		(1)		(1)		(1)	(1)			(3)		(3)		(3)		(3)	
	204Pb	207	Pb	208Pb	206P	b	%	ppm	ppm	ppm	ppm	232Th		/238U	/238U	/238	Ŭ	/206Pb	/2	32Th	/232	Th	/206Pb	cor-	208Pb		238U		207Pb <sup>*</sup>	:	207Pb	206P	, ,	err	238U		207Pb		207Pb		206Pb	err.
Spot	/206Pb ±%	/206	Pb ±%	/206Pb	±% /238l	J ±%	206Pb <sub>c</sub>	Ü	Th	206Pb <sup>°</sup>	208Pb <sup>*</sup>	/238U	±%	Age	Age	Age	1	Age		Age	Age	е	Age	dant	/232Th	±%	/206Pb	±%	/206Pb	±%	/235U	£% /238	J ±%	corr	/206Pb	±%	/206Pb	±%	/235U	±%	/238U	±% corr.
CGE_05Q-1.1	3.6E-2 5	0.5	80 1.6	1.39	1.2 5.91	4 3.4	64.00	18	3	1.3	0.1	0.17	3.42	512 ±49	509 ±28	498	±87 67	9 ±2584	1081	±1762	964 :	±781	460 ±169	90 +26	0.0489	82.9	12.10	10.0	0.0621	121	0.708	21 0.082	7 10.	0.08	12.45	18.2	0.038	64	0.42	68	0.0803	18.2 0.33
CGE_05Q-2.1	2.7E-2 5	0.4	56 2.6	1.04	2.4 4.30	3 2.6	48.68	29	3	2.0	0.1	0.10	3.46	501 ±44	497 ±39	501	±83 74	7 ±1529	499	±1666	208 ±	±1242	749 ±73	9 +34	0.0104	599.1	12.37	9.1	0.0642	72	0.715	73 0.080	8 9.1	0.12	12.37	17.3	0.064	35	0.72	37	0.0808	17.3 0.34
CGE_05Q-3.1	3.1E-2 5	0.5	05 1.6	1.16	1.2 4.46	5 2.6	55.00	28	3	2.0	0.0	0.11	3.31	512 ±35	512 ±22	521	±53 54	5 ±1914	10.4	±1841	-24 :	±854	976 ±48	7 +6	-0.0012	-3604.3	12.09	7.0	0.0584	88	0.666	88 0.082	7 7.0	0.08	11.89	10.7	0.072	24	0.83	25	0.0841	10.7 0.33
CGE_05Q-4.1	1.3E-2 6	0.2	50 4.6	0.52	5.0 2.712	2 3.0	23.33	86	12	6.9	0.2	0.14	1.89	573 ±17	573 ±17	576	±21 57	7 ±779	427	±520	424 :	±663	729 ±46	7 +1	0.0212	158.2	10.75	3.1	0.0593	36	0.760	36 0.093	0 3.1	0.09	10.69	3.8	0.064	22	0.82	23	0.0935	3.8 0.44
CGE_05Q-5.1	3.6E-2 6	0.5	77 2.0	1.34	1.4 4.99	7 3.4	63.52	24	3	1.8	0.0	0.15	3.81	554 ±58	553 ±33	565	±94 60	5 ±3067	51.1	±2428	7.4 ±	1152	1113 ±74	9 +9	0.0004	15649.3	11.15	10.9	0.0600	142	0.742	42 0.089	7 10.	9 0.08	10.92	17.3	0.077	38	0.97	40 /	0.0915	17.3 0.35
CGE_05Q-6.1	2.1E-2 6	0.3	73 3.3	0.85	4.0 3.778	3 2.6	38.11	36	8	2.9	0.2	0.22	2.18	580 ±28	577 ±24	578	±38 71	2 ±1214	637	±659	554 :	±620	618 ±65	6 +19	0.0278	113.5	10.62	5.0	0.0631	57	0.819	57 0.094	2 5.0	0.09	10.66	6.9	0.060	30	0.78	33 (	0.0938	6.9 0.52
CGE_25-1.1	5.6E-3 13	0.1	43 8.2	0.74	2.5 2.33	5 1.2	9.92	219	498	19.3	11.6	2.35	0.41	630 ±14	629 ±15	685	±22 66	1 ±602	518	±30	516	±34 2	2107 ±85	6 +5	0.0259	6.7	9.74	2.3	0.0616	28	0.872	28 0.102	7 2.3	0.08	8.92	3.3	0.131	49	2.02	50 (	0.1122	3.3 0.51
CGE_25-2.1	5.2E-3 7	0.1	35 8.1	0.65	2.6 2.44	7 2.6	9.17	210	408	18.7	9.6	2.01	0.42	638 ±12	639 ±15	683	±20 58	3 ±491	524	±23	528	±37 ·	1891 ±64	5 -10	0.0265	7.1	9.61	2.0	0.0594	23	0.853	23 0.104	1 2.0	0.09	8.95	3.0	0.116	36	1.78	37 (	0.1117	3.0 0.47
CGE_25-3.1	7.3E-3 7	0.1	69 7.1	0.85	1.0 2.40	2.8	12.94	196	498	17.6	11.8	2.63	0.77	641 ±13	640 ±16	709	±24 72	4 ±531	526	±20	521	±31 2	2287 ±47	9 +12	0.0261	6.0	9.56	2.2	0.0635	25	0.916	25 0.104	6 2.2	0.09	8.61	3.6	0.145	28	2.32	29 (	0.1162	3.6 0.34
CGE_25-4.1	5.9E-3 11	0.1	48 8.5	0.62	4.5 2.419	9 2.5	10.52	237	409	20.9	9.4	1.79	0.42	630 ±18	629 ±19	668	±27 67	6 ±616	514	±42	510	±51 ·	1825 ±83	8 +7	0.0256	10.2	9.75	2.9	0.0621	29	0.878	29 0.102	6 2.9	0.10	9.16	4.2	0.112	46	1.68	48 (	0.1092	4.2 0.55
CGE_25-5.1	5.6E-3 15	0.1	42 8.6	0.70	3.3 2.43	8 1.1	9.97	225	466	20.1	11.0	2.14	0.41	637 ±15	637 ±15	684	±21 61	5 ±695	526	±39	528	±40 ·	1954 ±95	9 -4	0.0265	7.7	9.63	2.5	0.0603	32	0.863	32 0.103	8 2.5	0.08	8.93	3.3	0.120	54	1.85	55 /	0.1120	3.3 0.57
CGE_25-6.1	4.4E-3 23	0.1	23 8.4	0.54	4.3 2.49	9 1.7	7.81	242	386	20.8	8.6	1.65	0.41	615 ±19	615 ±18	650	±24 57	5 ±706	497	±54	501	±44 ·	1732 ±60	8 -7	0.0251	8.9	10.00	3.2	0.0592	32	0.817	33 0.100	0 3.2	0.10	9.43	3.9	0.106	33	1.55	35 /	0.1061	3.9 0.51
CGE_25-7.1	5.3E-3 7	0.1	36 6.7	0.71	2.6 2.36	5 1.1	9.42	220	484	19.2	11.0	2.28	0.40	625 ±12	626 ±13	680	±20 55	7 ±430	506	±21	510	±29 2	2086 ±79	2 -13	0.0256	5.7	9.83	2.0	0.0587	20	0.824	20 0.10	8 2.0	0.10	8.98	3.2	0.129	45	1.98	47 /	0.1113	3.2 0.57
CGE_25-8.1	5.6E-3 14	0.1	41 7.4	0.80	2.1 2.35	3 2.1	9.96	203	507	17.1	11.3	2.57	0.41	602 ±14	603 ±14	662	±22 57	4 ±623	498	±28	499	±27 2	2181 ±90	7 -5	0.0250	5.5	10.21	2.4	0.0592	29	0.799	29 0.097	9 2.4	0.08	9.25	3.5	0.136	52	2.03	54 (	0.1081	3.5 0.52
CGE_25-9.1	5.0E-3 14	0.1	32 4.6	0.76	1.6 2.40	7 1.3	8.81	191	478	16.5	10.5	2.59	0.42	617 ±14	617 ±13	689	±22 60	0 ±478	491	±26	492	±19 2	2337 ±56	3 - 3	0.0246	3.8	9.96	2.4	0.0599	22	0.830	22 0.100	4 2.4	0.11	8.87	3.3	0.149	33	2.32	35 (	0.1127	3.3 0.55
CGE_25-10.1	4.2E-3 7	0.1	20 5.7	0.54	3.6 2.30	7 1.0	7.43	262	439	22.7	9.6	1.73	0.40	620 ±11	621 ±12	663	±17 55	9 ±325	487	±24	492	±30 ·	1866 ±49	6 -12	0.0247	6.2	9.90	1.8	0.0588	15	0.818	15 0.101	0 1.8	0.12	9.24	2.7	0.114	27	1.70	29	0.1083	2.7 0.60
CGE_25-11.1	5.5E-3 13	0.1	40 7.0	0.85	2.2 2.30	0 1.9	9.69	206	567	17.8	12.9	2.84	0.40	616 ±14	616 ±14	688	±24 62	4 ±561	507	±25	506	±25 2	2355 ±113	37 +1	0.0254	5.0	9.97	2.4	0.0606	26	0.837	26 0.100	3 2.4	0.09	8.87	3.7	0.151	67	2.34	69 (	0.1127	3.7 0.60
CGE_25-13.1	4.5E-3 7	0.1	24 3.9	0.74	0.9 2.31	5 1.1	7.92	207	523	18.5	11.6	2.61	0.38	639 ±12	640 ±12	720	±20 56	0 ±275	497	±15	501	±15 2	2417 ±30	3 -15	0.0251	3.0	9.60	1.9	0.0588	13	0.845	13 0.104	2 1.9	0.15	8.46	2.9	0.156	18	2.55	19 (	0.1182	2.9 0.52
CGE_25-14.1	3.9E-3 12	0.1	21 4.9	0.66	1.4 2.40	7 1.0	6.87	206	457	18.0	10.1	2.29	0.37	625 ±12	622 ±12	687	±18 76	0 ±312	495	±20	485	±18 2	2245 ±35	1 +19	0.0243	3.8	9.82	1.9	0.0646	15	0.907	15 0.101	9 1.9	0.13	8.89	2.7	0.141	20	2.19	21 (	0.1125	2.7 0.49
CGE_25-15.1	4.6E-3 7	0.1	27 4.8	0.70	2.3 2.27	3 1.1	8.21	209	466	17.0	9.9	2.31	0.39	584 ±11	584 ±11	638	±18 61	1 ±307	472	±18	470	±20 2	2135 ±65	5 +5	0.0235	4.3	10.54	1.9	0.0602	14	0.787	14 0.094	9 1.9	0.13	9.60	3.0	0.133	37	1.91	39	0.1041	3.0 0.59

Errors are 1-sigma; Pbc and Pb indicate the common and radiogenic portions, respectively.

Ernor is Stander albraich weit and Pointauare une common and radiugemic portons, respectively. Error in Stander calibration was 10.7% (not included in above errors but required when comparing data from different mounts). (1) Common Pb corrected using measured 204Pb. (2) Common Pb corrected by assuming 206Pb/238U-207Pb/238U age-concordance (3) Common Pb corrected by assuming 206Pb/238U-208Pb/238U age-concordance

Spot ID	CL response	<sup>204</sup> Pb	U (ppm)	Th (ppm)	Th/U	<sup>206</sup> Pb/ <sup>238</sup> U	2SE (%)	<sup>207</sup> Pb/ <sup>235</sup> U	2SE (%)	<sup>207</sup> Pb/ <sup>206</sup> Pb	2SE (%)	<sup>206</sup> Pb/ <sup>238</sup> U Age
TPR40-1.2	bright	0.162	167.6	75.0	0.447	0.3929	3.2	6.97	3.3	0.1287	0.9	2136.1
TPR40-7.1	bright	-0.115	55.3	50.8	0.919	0.3653	3.0	6.50	3.1	0.1290	0.5	2007.2
TPR40-8.1	bright	0.043	197.5	266.8	1.351	0.3994	3.1	7.04	3.2	0.1278	0.6	2166.1
TPR40-9.1	bright	0.098	99.0	107.5	1.086	0.3992	3.2	7.16	3.3	0.1302	0.8	2165.3
TPR40-10.1	bright	-0.056	57.8	78.5	1.357	0.3884	3.2	6.89	3.2	0.1287	0.4	2115.2
TPR40-13.1	bright	-0.016	225.4	207.6	0.921	0.3585	3.1	6.37	3.1	0.1289	0.5	1975.0
TPR40-14.1	bright	0.000	37.1	35.1	0.946	0.3751	3.4	6.49	3.5	0.1255	1.0	2053.2
TPR40-1.1	dark	0.106	1163.0	164.4	0.141	0.2501	3.2	4.04	3.2	0.1170	0.3	1439.1
TPR40-2.1	dark	0.124	1462.3	445.4	0.305	0.1765	3.0	2.67	3.0	0.1097	0.3	1047.7
TPR40-3.1	dark	0.067	1083.1	388.6	0.359	0.2062	3.1	3.17	3.1	0.1116	0.3	1208.6
TPR40-4.1	dark	0.062	1102.7	185.4	0.168	0.2488	3.0	4.03	3.0	0.1175	0.2	1432.4
TPR40-5.1	dark	0.112	1593.4	561.2	0.352	0.1741	3.0	2.38	3.0	0.0993	0.3	1034.8
TPR40-6.1	dark	0.042	1695.9	347.5	0.205	0.1880	3.1	2.63	3.1	0.1015	0.4	1110.3
TPR40-8.2	dark	0.101	2109.4	240.1	0.114	0.1695	3.0	2.41	3.0	0.1031	0.5	1009.6
TPR40-11.1	dark	0.007	1504.0	235.3	0.156	0.2635	3.0	4.23	3.0	0.1165	0.2	1507.5
TPR40-12.1	dark	0.056	1923.5	232.7	0.121	0.2345	3.0	3.55	3.0	0.1100	0.4	1357.8
TPR40-15.1	dark	0.105	789.5	395.8	0.501	0.2391	3.0	3.81	3.0	0.1155	0.3	1382.2

Table A8 - Summary of SHRIMPIle data on zircons crystals from sample TPR-40

1SE	<sup>207</sup> Pb/ <sup>206</sup> Pb Age	1SE	Discordance (%)
59	2080.2	26.0	6
53	2047.2	77.1	-3
57	2107.4	27.3	4
59	2130.0	32.9	-6
58	2064.6	19.5	-4
52	1971.0	15.3	-2
59	1866.4	41.0	-1
41	1855.6	8.3	28
29	1756.2	20.5	45
34	1658.1	11.2	37
39	1975.2	7.1	28
29	1416.8	18.4	39
31	1659.4	8.8	36
28	1612.1	11.0	43
41	1903.7	5.6	23
37	1767.0	8.4	27
37	1218.0	21.5	30



Cathodoluminescence (CL) image along with transmitted light image of U-Pb analysis in zircon at LA-MC-ICP-MS. Spot size of 30µm. Sample from the Meta-syenogranite unit (ARE-04). Ages 206/207Pb for each spot are reported.

12.1 - 2079 Ma 12.2 - 1931 Ma



CL image along with transmitted light image of U-Pb analysis in zircon at LA-MC-ICP-MS. Spot size of 30µm. Sample from the Meta-syenogranite unit (ARE-04). Ages 206/207Pb for each spot are reported.



26.1 - 2015 Ma

500µm


CL image along with transmitted light image of U-Pb analysis in zircon at SHRIMP. Spot size of 20µm. Sample from the Meta-syenogranite unit (CGE-25). Ages 206/207Pb for each spot are reported.

13.1 - 2044 Ma

## 14.1 - 2056 Ma

15.1 - 2004 Ma

16.1 - 2022 Ma



CL image along with transmitted light image of U-Pb analysis in zircon at SHRIMP. Spot size of 20µm. Sample from the Hornblende-Garnet Metatexite unit (CGE-05). Ages 206/207Pb for each spot are reported.

## 9.1 - 1789 Ma

## 10.1 -1969 Ma

## 11.1 - 2094 Ma

## 12.1 - 2084 Ma

## 13.1 - 2057 Ma





## 1.1 - 1117 Ma











2.1 - 976 Ma



6.1 - 1391 Ma







CL image along with transmitted light image of U-Pb analysis in zircon at SHRIMP. Spot size of 20µm. Sample from the Biotite diatexite unit (CGE-03Zr). Ages 206Pb/238U for each spot are reported.

9.1 - 604 Ma

10.1 - 607 Ma 10.2 - 644 Ma

300µm



11.1 - 1910 Ma









CL image along with transmitted light image of U-Pb analysis in zircon at SHRIMP. Spot size of 20µm. Sample from the Biotite diatexite unit (CGE-03Zr). Ages 206Pb/238U for each spot are reported.

## 14.1 - 602Ma

## 15.1 - 1184 Ma

## 16.1 - 2081 Ma



## **TPR40-1.1**

**TPR40-1.2** 



**TPR40-5.1** 





# **TPR40-2.1** 0

TPR40-6.1



TPR40-10.1



**TPR40-3.1** 



**TPR40-7.1** 



**TPR40-11.1** 



**TPR40-4.1** 



**TPR40-8.1** 

**TPR40-8.2** 



TPR40-12.1



CL image of U-Pb analysis in zircon at SHRIMP. Spot size of 20µm. Sample from the Hornblende Metatexite unit (TPR-40).





**TPR40-13.1** 



## TPR40-14.1

## TPR40-15.1

# **Geological Map - Alfenas and Areado regions**



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