# A-TYPE MAGMATISM IN THE SIERRAS OF MAZ AND ESPINAL: A NEW RECORD OF RODINIA BREAK-UP IN THE WESTERN SIERRAS PAMPEANAS OF ARGENTINA

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#### Abstract

Two orthogneisses have been recognized in the sierras of Espinal and Maz (Western Sierras Pampeanas, NW Argentina) that were emplaced within a Grenvillian metasedimentary sequence. Microcline, plagioclase and quartz are the main rock-forming minerals, with accessory zircon, apatite-(CaF), magnetite, biotite (Fe/(Fe+Mg) = 0.88-0.91), ferropargasite (Fe<sub>total</sub>/(Fe<sub>total</sub> + Mg) = 0.88-0.89), titanite (with up to 1.61 wt.% Y<sub>2</sub>O<sub>3</sub>) and an REE-rich epidote. REE-poor epidote and zoned garnet (Ca and Fe<sup>3+</sup>-rich) are metamorphic minerals, while muscovite, carbonates and chlorite are secondary phases. Texture is mylonitic.

Two representative samples are classified as granite (from Sierra de Espinal) and granodiorite/tonalite (from Sierra de Maz) on the grounds of immobile trace elements. Some trace element contents are rather high (Zr: 603 and 891 ppm, Y: 44 and 76 ppm, 10000\*Ga/Al: 2.39-3.89) and indicate an affiliation with A-type granites (more specifically, the  $A_2$  group). Both samples plot in the field of within-plate granites according to their Y and Nb contents.

Concordant crystallization ages (zircon U-Pb SHRIMP) are  $842 \pm 5$  and  $846 \pm 6$  Ma respectively.  ${}^{87}\text{Sr}/{}^{86}\text{Sr}_{i\,(845)}$  ratios are 0.70681 and 0.70666;  $\epsilon \text{Nd}_{i\,(845)}$  values are -1.5 and +0.3 and depleted mantle Nd model ages ( $2T_{DM}*$ ) are 1.59 and 1.45 Ga respectively. These values indicate the involvement of an isotopically evolved source.  $2T_{DM}*$  values are compatible with the presence of inherited zircon crystals of up to 1480 Ma in one of the rocks, thus implying that magmas incorporated material from Mesoproterozoic continental source. This is also indicated by the relatively high contents of Y, Ga, Nb and Ce compared to magmas derived from sources similar to those of oceanic-island basalts.

These orthogneisses represent a period of extension at ca. 845 Ma affecting the Western Sierras Pampeanas continental crust that was already consolidated after the Grenvillian orogeny (1.2 - 1.0 Ga). They are thus a record of the early stages of Rodinia break-up. Metamorphic conditions during the subsequent Famatinian orogenic cycle (ca. 420 Ma, SHRIMP U-Pb on zircon) attained  $7.7 \pm 1.2$  kbar and  $664 \pm 70^{\circ}$ C.

<u>Keywords</u>: Rodinia; U-Pb SHRIMP dating; anorogenic magmatism; Mesoproterozoic; Western Sierras Pampeanas; Argentina.

#### Introduction

The Sierras Pampeanas in northwestern Argentina constitute large exposures of pre-Mesozoic crystalline basement in the foreland of the the Central Andes (Fig. 1A). They record a complex tectonomagmatic history from the Mesoproterozoic to the Late Paleozoic that has not yet been completely deciphered. In the Western Sierras Pampeanas evidence for a reworked Grenville-age basement was firmly demonstrated by McDonough et al. (1993). This basement has since been considered the counterpart of the Grenville orogen along the southern Appalachian margin of Laurentia, which drifted away in the late Neoproterozoic to early Paleozoic to finally dock against the proto-Andean margin of Gondwana in the Ordovician. Whether this process involved an allochthonous exotic terrane, i.e., the Precordillera terrane hypothesis (for a review see Thomas and Astini, 2003, Ramos, 2004), or para-authoctonous translation along the proto-Andean margin of Gondwana (e.g., Finney, 2007; Casquet et al. 2008a), remains a matter of dispute. However all authors agree that the Western Sierras Pampeanas Grenville-age terranes were contiguous to Laurentia by the end of Rodinia amalgamation at ca. 1.0 Ga. Moreover Casquet et al. (2008b) raised the hypothesis that, at the onset of the Grenvillian orogeny, the Western Sierras Pampeanas Grenvillian terranes were part of a larger continental mass that embraced other Meso- to Paleoproterozoic outcrops such as the Arequipa block in southern Peru and northern Chile. This view opens new ways to interpret the Proterozoic history of southern South America (Rapela et al., 2007).

Rodinia is the name given to a hypothetical supercontinent that comprised almost all continental masses on Earth during the late Mesoproterozoic. Geological evidence for the existence and evolution of such supercontinent has been growing since the early 1990's (e.g., Hoffman, 1991; Dalziel, 1991; Moores, 1991; see Li et al., 2008 for a review), but consensus is still lacking on issues such as the number of participating cratons, their relative positions and the chronology of the assembly and subsequent break-up of Rodinia. Rodinia break-up and dispersal began around 900 Ma ago or even earlier (e.g., Cordani et al., 2003), but evidence of widespread rifting associated with mantle plumes occurred much later, spanning the time interval between 825 and 740 Ma (Li et al., 2008).

Evidence for Rodinia break-up in the Western Sierras Pampeanas was first recognized in the Sierra de Pie de Palo (Fig. 1A). Here A-type orthogneisses hosted by reworked Grenvillian basement yielded a zircon U-Pb SHRIMP age of ca. 774 Ma. Protoliths were interpreted as resulting from an anorogenic magmatic event during early rifting of Rodinia (Baldo et al., 2006).

This contribution deals with a newly recognized anorogenic magmatic event at ca. 845 Ma in the sierras of Maz and Espinal (Fig. 1A) which records a still earlier event of Rodinia break-up. It thus adds to understanding of the Rodina break-up process in this part of Western Gondwana both in time and paleogeography.

# Geological setting

The sierras of Maz and Espinal (Fig.1) were first described by Kilmuray and Dalla Salda (1971). Basement outcrops in the area, including those of the adjacent Sierra de Ramaditas and Villa Castelli massif, consist of metamorphic rocks with sedimentary

and igneous protoliths intruded by a Lower Ordovician suite of metaluminous and peraluminous granites that occurs east of the sierras, near Villa Castelli (Dahlquist et al 2007) (Fig.1). The basement outcrops are covered by Late and post-Paleozoic sedimentary rocks in angular unconformity. Westward high-angle thrusting during the Andean orogeny led to superposition of the basement blocks over the sedimentary cover (Jordan and Almendinger, 1986).

In the basement of the sierras of Maz and Espinal three domains separated by first order shear zones and faults (Fig. 1-B) were distinguished on the basis of field, geochronology and isotope composition evidence (Casquet et al., 2006, 2008 b). The Eastern Domain, consisting for the most part of high grade rocks, i.e., garnet-sillimanite migmatitic gneisses with subordinate marbles and amphibolites, is younger than 1.0 Ga. Metamorphism took place during the Ordovician-Silurian Famatinian orogeny at ca. 440 Ma (see also Lucassen and Becchio, 2003). The Central Domain (also known as the Maz terrane) consists of medium-grade (kyanite-sillimanite-garnet-staurolite schists, quartzites, amphibolites and marbles) to high-grade intermediate-to-ultrabasic metaigneous rocks and metasedimentary rocks that underwent a Grenvillian-age orogeny starting at ca. 1.2 Ga (Porcher et al., 2004; Casquet et al., 2006). Massif-type anorthosites of ca. 1070 Ma are restricted to the Maz terrane. Anorthosites also show evidence for Famatinian metamorphic rejuvenation ( $431 \pm 40$  Ma) throughout the Maz terrane (Casquet et al., 2005). Moreover, metamorphic and geochronological discontinuities within the Maz terrane suggest that it is in fact composed of a number of slivers separated by shear zones of unknown age, probably Famatinian. The Western Domain consists again of metasedimentary rocks younger than ca. 1.0 Ga that underwent Famatinian metamorphism. One sequence of rocks composed of thick marble beds, calcic semipelitic schists and quartzites is probably equivalent to the late Neoproterozoic Difunta Correa metasedimentary sequence of the Sierra de Pie de Palo, and to isotopically equivalent rocks of the Sierra de Umango (Varela et al. 2001, Galindo et al. 2004, Rapela et al. 2005). Most rocks within this domain are low-grade but high-grade rocks are locally found. A recently described ca. 0.57 Ga alkaline syenite-carbonatite complex (Casquet et al., 2008 c) occurs in the eastern margin of the Sierra of Maz.

# Figure 1 should appear about here

Two orthogneisses have been found that have provided almost the same crystallization age of ca. 845 Ma (see below). Both are foliated sheeted bodies concordant with the external foliation and bedding. Field evidence suggests that the protoliths were intrusive and not tectonically emplaced. The first, in the Sierra of Maz (fig. 2-A), is a ca. 100m thick body of leucocratic mylonitic augen-gneiss with streaks of mafic minerals and concordant stretched pegmatites; the fabric is S=L. It is hosted by whitish quartzites, sillimanite-garnet gneisses and garnet amphibolites that are assigned on the basis of Nd model ages (Casquet et al., 2006) to the Western Domain referred to above. This domain underwent Famatinian metamorphism only. The second, in the Sierra del Espinal (fig. 2-B), is a mylonitic augen-gneiss hosted by a sequence of kyanite-staurolite-garnet schists, amphibolites, marbles and quartzites belonging to the Maz terrane (Casquet et al., 2006). The extent of this body is unknown, as only a section some 20 m thick is visible along one creek. This sequence underwent medium-grade Grenvillian metamorphism and a low-grade rejuvenation attributed here to the Famatinian orogeny (see below).

# Figure 2 should appear about here

#### Sampling and analytical methods

Two samples were collected that are representative of the two orthogneisses: MAZ-6037 was taken from an outcrop along the Maz Creek (29°11′20′′W - 68°28′48′′S). ESP-7066 was collected on the western slopes of the Sierra del Espinal, close to Puesto Villalba (29°05′23′′W-68°31′40′′S) (Fig. 1B).

Electron-microprobe analyses were performed on sample MAZ-6037 and on a host amphibolite (MAZ-12046), the latter for estimation of metamorphic conditions, at the Complutense University, Madrid (Supplementary data files obtainable from the Precambrian Research Data Repository).

Whole-rock powders of ESP-7066 and of a further sample of the MAZ orthogneiss (MAZ-12040) were analysed by ActLabs (Canada) for major elements (ICP) and trace elements (ICP-MS) (4-Lithoresearch code) (Table 1).

Rb–Sr and Sm–Nd isotope determinations were carried out at the Geochronology and Isotope Geochemistry Center of the Complutense University (Madrid, Spain) on an automated multicollector VG® SECTOR 54 mass spectrometer. Analytical uncertainties are estimated to be 0.01% for <sup>87</sup>Sr/<sup>86</sup>Sr, 0.006% for <sup>143</sup>Nd/<sup>144</sup>Nd, 1% for <sup>87</sup>Rb/<sup>86</sup>Sr, and 0.1% for <sup>147</sup>Sm/<sup>144</sup>Nd. Replicate analyses of the NBS-987 Sr-isotope standard yielded an average <sup>87</sup>Sr/<sup>86</sup>Sr ratio of 0.710227±0.00004 (n=10) and La Jolla Nd-isotope standard yielded an average <sup>143</sup>Nd/<sup>144</sup>Nd of 0.511844±0.00002 (n=10). Errors are quoted throughout as two standard deviations from measured or calculated values (Table 2).

U-Th-Pb zircon dating was performed on the two samples at the Research School of Earth Sciences, The Australian National University, Canberra, Australia, using SHRIMP RG (MAZ-6037) and SHRIMP II (ESP-7066). Separated zircons were mounted in epoxy resin together with chips of the FC1 Duluth Gabbro reference zircon. Reflected and transmitted light photomicrographs and cathodo-luminescence (CL) SEM images were used to decipher the internal structures and to target specific areas within the zircons. Analytical methods followed Williams (1998, and references therein). U/Pb ratios were normalized relative to a value of 0.01859 for the FC1 reference zircon, equivalent to an age of 1099 Ma (see Paces and Miller, 1993) and data were reduced using the SQUID Excel macro of Ludwig (2001) (Supplementary SHRIMP data files are obtainable from the Precambrian Research Data Repository). Uncertainties are quoted at the 1-sigma level. <sup>204</sup>Pb-corrected data are presented, but it should be noted that this is not optimal for some of the low-U areas analysed, giving rise to large uncertainties in the radiogenic  ${}^{207}$ Pb/ ${}^{206}$ Pb ratios and ages. All age calculations were carried out using Isoplot/Ex (Ludwig, 2003) and the resulting ages quoted in the text and figures are quoted with 95% confidence limits, including propagation of the uncertainties in the calibration of the U/Pb ratio of the reference zircons (0.40% for sample MAZ-6037 and 0.16% for ESP-7066).

# Petrography and mineral chemistry

The Sierra del Espinal orthogneiss consists of quartz and microcline, with lesser amounts of plagioclase, opaque phases and very abundant muscovite and chlorite as alteration products after biotite (Fig. 3-A and 3-B). Titanite and scarce zircon are the main accessory minerals. Because of the strong low-grade retrogression this rock was not further considered for electron microprobe analyses

The orthogneiss from the Sierra de Maz shows a mylonitic texture; microcline and plagioclase constitute rounded porphyroclasts, while quartz forms recrystallized ribbons (figure 3-C and 3-D). Microcline, plagioclase and quartz are the main rock-forming minerals and are inherited from the igneous paragenesis. Zircon, apatite, magnetite, biotite, amphibole (ferropargasite), titanite and an REE-rich epidote are accessory and except for some biotite and titanite are probably also inherited. However, garnet and an REE-poor epidote forming mantles around REE-rich epidote grains are metamorphic minerals, as are some biotite and titanite, and muscovite. The low Mn (7.87-1.74 wt% MnO) and high Ca (13.67-11.02 wt% CaO) contents are not characteristic of garnets of magmatic origin and are instead similar to metamorphic garnets from meta-granites (Le Goff and Ballevre, 1990, Kotopouli et al., 2000). Biotite and amphibole have high Fe/(Fe+Mg) ratios (0.88-0.91 and 0.88-0.89, respectively). Muscovite is of secondary origin (after biotite and feldspars) and is locally associated with opaque minerals. Irregular carbonate concentrations are locally observed.

# Figure 3 should appear about here

# Whole rock geochemistry

The two rocks are chemically different (Table 1). Unfortunately, identification of the gneiss protolith in terms of major element chemistry is subject to uncertainty because of the strong metamorphic overprint. A more realistic classification should be based on immobile elements such as HFS elements. On the Zr/TiO<sub>2</sub>\*0.0001 vs. Nb/Y plot (Winchester and Floyd, 1977), ESP-7066 is a rhyolite, i.e., granite, whereas MAZ-12040 plots in the field of rhyodacites/dacites, i.e. granodiorite to tonalite.

# Table 1 should appear about here

Relying on HFS elements only, contents of Zr and Y are high in both rocks (603, 891 ppm and 44, 76 ppm, respectively). However, Nb and Ce are not so notably enriched (24, 27 ppm; 180, 97 ppm, respectively), although their values exceed those considered usual for fractionated I- and S-type granites (as compiled by Whalen et al., 1987) and are more typical of A-type granitoids. In fact values of 10000\*Ga/Al are 2.39 and 3.89, respectively, i.e., close to or above the value of 2.6 recommended by Whalen et al. (1987) to distinguish A-type granitoids. On a Zr+Nb+Ce+Y diagram (Eby, 1990) the two rocks plot in the field of A-type granitoids (Fig. 4-A). On the other hand, contents of mobile elements such as Rb (low: 54-74 ppm), Sr (moderate: 97-232 ppm) and Ba (high: 1051-1787 ppm) are more typical of non-evolved granitoids. Normalized REE patterns are slightly fractionated (La<sub>N</sub>/Lu<sub>N</sub> = 13 and 4 respectively), with REE concentrations close to 100 times chondritic values (Fig. 4-B). The Eu anomaly is negative and moderate (0.64) in ESP-7066 and slightly positive (1.09) in MAZ-12040.

Both samples plot in the field of within-plate granites close to the field of volcanicarc granites on discriminant diagrams based on relative abundances of Ta, Y, Yb, Nb, Hf and Rb (Pearce et al., 1984; Harris et al., 1986). Moreover, the two samples belong to the  $A_2$  group as defined by Eby (1992), which represents magmas derived from continental crust that has been through a cycle of subduction-zone or continentcontinent collision magmatism (Figs 4-C and D).

# Figure 4 should appear about here

#### Isotope composition

 $^{87}$ Sr/ $^{86}$ Sr<sub>i</sub> ratios at 845 Ma – the probable crystallization age (see below) – are 0.70681 and 0.70666 for MAZ 12040 and ESP-7066 respectively (Table 2).  $\epsilon$ Nd<sub>i</sub> values at the age of 845 Ma are +0.32 to – 1.5. Depleted mantle 2-stage Nd model ages (2T<sub>DM</sub>) (De Paolo et al., 1992) are 1.45 and 1.59 Ga (Table 2). Sr and Nd isotope initial ratios are evidence that an isotopically moderately evolved source contributed to the magma composition. Moreover, Nd model ages suggest that this source might be sought in underlying Mesoproterozoic continental crust. This interpretation is reinforced by zircon data below. Whether a juvenile component was also involved in the magma composition cannot be confirmed with the available information.

# Table 2 should appear about here

### U-Pb SHRIMP geochronology

Zircons from MAZ-6037 are elongate to sub-equant, euhedral to sub-round grains or fragments that are generally between 200 to 300 µm in length. The CL images show a dominantly oscillatory-zoned internal structure, with many grains having a thin, <10µm bright CL rim (Fig. 5-A). Some grains show a more complex internal structure, with central areas of oscillatory zonation overgrown by a less clearly oscillatory zoned zircon, in turn overgrown by the very thin bright CL outermost rim for which no reliable analyses could be obtained. Thirty areas were analysed on 27 zircon grains (data obtainable from the Precambrian Research Data Repository). The Pb peaks were not correctly centred during the analyses of grains 5 and 6 and so no data is presented for these. The outer rim and core areas were analysed on grain 1 and both yield <sup>207</sup>Pb/<sup>206</sup>Pb ages that are within uncertainty. The 28 analyses presented in the Data Repository are a mixture of both the inner and outer areas of grains, both types yielding Th/U ratios that are in the usual range recorded by igneous zircon (0.3-0.7). Furthermore, on the Wetherill concordia plot (Fig. 5-B) it can be seen that the majority of data lie within uncertainty of concordia and there is no consistent difference between analyses of these two zircon types. Some dispersion is evident and this probably results from correction errors associated with very small amount of radiogenic Pb. Twenty-two of the 28 analyses yield a self-consistent Concordia age (as in Ludwig, 2001) of 846 ± 6 Ma (MSWD = 1.4) and this constrains the crystallisation age of the dominant zoned igneous zircon. The fact that both inner and outer zoned components are within uncertainty at ca. 846 Ma indicates that they are coeval to within the uncertainty of SHRIMP analysis.

The zircons from sample ESP-7066 constitute a more heterogeneous population than those described above. The grains are mostly ~200  $\mu$ m in length, but are more clearly sub-round, with a few subhedral, and under transmitted light they have clearly been more affected by a metamorphic event. The CL images show a range of complex structures. Whilst many show oscillatory zonation, there are older central cores to some grains, and more homogeneous embayments of low-luminescence metamorphic zircon in others (Fig. 5-C). This complexity is highlighted in the U-Pb data, with 17 analyses

on 14 zircon grains (see table in Data Repository). Two analyses of the presumed metamorphic embayments on grains 4 and 7 yield  ${}^{206}\text{Pb}/{}^{238}\text{U}$  ages of ca. 420 Ma, consistent with Paleozoic metamorphism. Older inherited cores and whole grains give ages of ~1480, ~1200 and ~1000 Ma (Fig 5-C). The dominant oscillatory-zoned zircons yield a concordant group of analyses with a Concordia age of 842 ± 5 Ma (MSWD =2.0, 9 analyses, Fig. 5-D). This ca. 842 Ma igneous zircon can be seen to enclose older components (both igneous and metamorphic), and in turn is itself rimmed and embayed by the ca. 420 Ma metamorphic zircon.

The two samples thus provide crystallization ages that are coincident within error at ca. 845 Ma, i.e., Early Cryogenian, according to the International Stratigraphic Chart (International Commission on Stratigraphy, 2008). Both have been affected by metamorphism which, at least in the case of ESP-7066, is shown to be Silurian.

## Figure 5 should appear about here

## Conditions of metamorphism

Peak metamorphic P-T conditions were assessed for the Maz site to contribute to a better knowledge of the orthogneisses petrology. Using of multivariate equilibria procedure was however hampered by the garnet composition, which is very rich in Ca and Fe, and outside the range of thermodynamic models. Therefore, the best estimate of metamorphic P-T conditions is obtained from the host rocks.

Calculations were made on a para-amphibolite close to the orthogneiss body (sample MAZ-12046) using THERMOCALC 3.1 (Powell and Holland, 1988. Garnet, amphibole, biotite, quartz, plagioclase and ilmenite constitute the dominant assemblage, while calcite, epidote and chlorite were formed on the retrograde path. Other minor phases include apatite, allanite and zircon. Syntectonic garnet displays a very slight zonation, with decreasing Mg (X<sub>Py</sub> from 0.12 to 0.117-0.105) and Fe (X<sub>Alm</sub> from 0.654 to 0.645-0.647) and increasing Ca (X<sub>Gro</sub> from 0.177 to 0.182-0.190) and Mn (X<sub>Sps</sub> from 0.048 to 0.053-0.059) from core to rim. Neither biotite nor amphibole is significantly zoned. Biotite has an average Mg# of 0.44. Amphibole (ferropargasite according to the classification of Leake et al., 1997) has Al<sub>Total</sub> between 2.668 and 2.875 apfu and #Mg = 0.43. The average Fe<sup>3+</sup>/(Fe<sup>2+</sup> + Fe<sup>3+</sup>) ratio is 0.10. Plagioclase ranges from An<sub>54</sub> to An<sub>67</sub> (mineral compositions are obtainable from the Data Repository). Peak metamorphic conditions were: P = 7.7 ± 1.2 kbar and T = 664 ± 70° C.

# Discussion

A-type magmatism is indicative of largely continental within-plate extensional settings (e.g., Eby, 1990, 1992; Bonin, 2007). Thus we infer that the A-type granitoids described here represent a period of extension at ca. 845 Ma (early Cryogenian) affecting the continental crust of the Western Sierras Pampeanas that was already consolidated after the Grenvillian orogeny. This magmatic event implies that rifting of the Western Sierras Pampeanas Grenvillian basement started earlier than previously established. The oldest crystallization age yet reported for anorogenic A-type granitoids in the Western Sierras Pampeanas is ca. 774 Ma (middle Cryogenian; Baldo et al., 2006). Compared to the juvenile isotope composition of the ca. 774 Ma Sierra de Pie de

Palo orthogneisses (Sr<sub>i</sub> = 0.7005 to 0.7030,  $\epsilon$ Nd = +4.1 to +4.9; Baldo et al., 2006), those of Maz and Espinal orthogneisses resulted from the involvement of an isotopically more evolved source. Moreover, depleted-mantle Nd model ages of up to 1.46 Ga and inherited zircon crystals of up to 1.48 Ga in sample ESP-7066 lead us to speculate that the ca. 845 Ma A-type magmatism at Maz, largely involved a Mesoproterozoic continental source. The latter is also indicated by the relatively high contents of Y, Ga, Nb and Ce compared to those of magmas directly derived from mantle sources (Eby, 1992).

Igneous rocks of uncertain chemical signature that might likewise correspond to the same anorogenic event referred to here have been recorded from other locations in the Western Sierras Pampeanas. Mulcahy et al. (2003) reported a U-Pb SHRIMP zircon age of  $839 \pm 10$  Ma for an orthogneiss from the Sierra de la Huerta, southeast of the Sierra de Pie de Palo (Fig. 1). Vujovich et al. (2005) reported depleted mantle Nd model ages between 782 and 806 Ma for ortho-amphibolites from the Sierra de Umango, west of the Sierras of Maz and Espinal (Fig.1). However, these rocks are more probably related to the second extensional event.

The Western Sierras Pampeanas Grenvillian basement has been correlated with other outcrops of Proterozoic basement in South America, i.e., the Arequipa-Antofalla block and Amazonia, on the basis of geochronology of detrital zircons and Nd and Pb isotope geochemistry (Casquet et al., 2006, 2008b). It is probable that these continental masses were accreted to Laurentia through the Grenville-Sunsás orogeny between 1.2 and 1.05 Ga and thus amalgamated to the Rodinia supercontinent, with Laurentia in a central position. The process involved a still highly conjectural history of collision and further protracted lateral displacement of Amazonia along the boundary between the two continents (e.g., Loewy et al., 2003, 2004; Tohver et al., 2002, 2004; Boger et al., 2005; Cordani and Teixeira, 2007, among others). In the Neoproterozoic and Early Paleozoic, the continental masses mentioned above, i.e., Laurentia, Amazonia, Western Sierras Pampeanas, Arequipa-Antofalla and other minor cratons such as Rio Apa (Cordani et al., 2008) remained attached, forming a large continent that was involved in the Pampean orogeny between 535 and 520 Ma (Casquet et al., 2008c). This orogeny resulted from collision with other Gondwanan cratons to the east (present coordinates), probably Kalahari, and led to closure of the intervening Clymene Ocean (Trindade et al., 2006) and the final amalgamation of SW Gondwana (Rapela et al., 2007).

A-type orthogneisses in the Western Sierras Pampeanas are thus a record of Rodinia break-up that took place through a sequence of events in the Neoproterozoic. Allegedly mantle-plume related break-up pulses that affected large areas of the Earth have been recognized in different continents at ca. 825, 780 and 750 Ma (Li et al., 2008). The age reported by Baldo et al. (2006) (ca. 774 Ma) matches well one of the pulses mentioned above, whereas the age of the metagranitoids reported in this contribution is the oldest yet. In the case of the Western Sierras Pampeanas Grenvillian basement these events resulted only in aborted rifts, inasmuch as no evidence for oceanic crust of Neoproterozoic age has been reported so far. Of relevance to our case is the opening of the Iapetus Ocean between Laurentia and Amazonia that took place near the Neoproterozoic-to-Cambrian transition after a long rifting initiated as early as 765 Ma (Tollo et al., 2004). Drifting apart from Laurentia led to development of passive margin sedimentary sequences on both sides of the Iapetus Ocean almost coeval with the Pampean collision (Casquet et al., 2008b). A record of this process is represented by

the Argentine Precordillera carbonate platform, located west of the Sierras Pampeanas (e.g., Astini et al., 1995). Regardless of whether this platform is an exotic terrane travelled from the western margin of the Iapetus and accreted to western Gondwana in the early Paleozoic (reviews in Thomas and Astini, 2003; Ramos, 2004) or a para-autochthonous terrane (e.g., Finney, 2007; Casquet et al., 2008c), it provides evidence that the Western Sierras Pampeanas Grenvillian basement was also part of the drifted conjugate margin of Iapetus.

From the above discussion we infer that break-up of Rodinia along the Grenvillian boundary between Amazonia (+ Western Sierras Pampeanas + Arequipa-Antofalla) and Laurentia was protracted, starting at least at ca. 845 Ma and ending through drifting and ocean opening in the early Cambrian.

A few examples of A-type granitoids from elsewhere in South America coeval with those described here were recognized by Basei et al. (2008) in the reworked basement of the Brasiliano – Panafricano Dom Feliciano Belt, southern Brazil. Ages of  $835 \pm 9$  MA (IDTIMS) and  $843 \pm 12$  Ma (SHRIMP) were obtained that are within error of those found here. Whether this rifting event in southern Brazil was spatially connected through the continental hinterland with that in the Western Sierras Pampeanas, or alternatively records coeval but independent extensional processes in a separate craton (such as the Rio de la Plata craton, Kröner and Cordani, 2003; Fig.3), remains conjectural. Recent geochronological, paleomagnetic and geological evidence for southern South American cratons in the Neoproterozoic (Rapela et al., 2007) seem to favour the second interpretation.

It is notable that the rifting event at ca. 845 Ma was almost coeval with consumption of the Brasiliano Ocean between the São Francisco/Congo and the Amazonia and Paraná cratons in the early Neoproterozoic (Kröner and Cordani, 2003). An intra-oceanic magmatic arc, the juvenile Goiás magmatic arc in Central Brazil, existed between ca. 890 and 800 Ma (Pimentel et al., 2000; Laux et al., 2005). Complete consumption of the Brasiliano Ocean took probably place at ca. 600 Ma (Laux et al., 2005). This evidence of subduction in the early Neoproterozoic has been taken as a proof that some large cratons such as São Francisco/Congo and others were not part of the Rodinia supercontinent (Cordani et al., 2003; Kröner and Cordani, 2003). We can only state that the rifting events at ca. 845 Ma and ca. 774 Ma were comparatively very short and took place while subduction of the Brasiliano Ocean was underway.

The age of the metamorphic rims in zircon grains from ESP-7066 (ca. 420 Ma, i.e., Silurian) confirms that a strong metamorphic overprint took place during the Ordovician–Silurian Famatinian orogeny. Although still poorly known in detail (time and P-T conditions) the Famatinian overprint in the sierras of Maz and Espinal was widespread, varying from greenschist to upper garnet-amphibolite facies conditions (Lucassen and Becchio, 2003; Casquet et al., 2008 b). Because no conclusive evidence exists in the sierras of Maz and Espinal of orogeny between ca. 840 Ma and the Famatinian orogeny (however, see Mulcahy et al., 2007 and Casquet et al., 2008c), high-grade metamorphism ( $664 \pm 70^{\circ}$ C) at the site of MAZ-6037 was probably Famatinian. The thin low-U zircon rims in sample MAZ-6037 were unfortunately undatable. Since sample ESP-7066 underwent metamorphism at much lower grade than MAZ-6037, it is suggested that zircon overgrowths in this case might be related to the pervasive influx of fluids leading to retrogression of the igneous association. Zircon

crystallization seems possible from aqueous fluids under low P and T conditions (< 500°C) and high water/rock ratios (Corfu et al., 2003, Dempster et al. 2004, Rasmussen 2005).

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Table	1	
	ESP-	MAZ-
	7066	12040
wt %		
SiO	77 50	68 81
ALO	7.00	12 60
$AI_2O_3$	7.90	13.00
$Fe_2O_3$	4.20	6.35
MnO	0.04	0.09
MgO	0.98	0.23
CaO	0.98	2.19
$Na_2O$	0.58	2.92
K <sub>2</sub> O	4.85	4.73
TiO <sub>2</sub>	1.17	0.56
$P_2 O_{\epsilon}$	0.09	0.17
	1.42	0.58
Total	00.68	100.30
Total	99.00	100.50
ррт	6	0
Sc	6	9
Ве	<1	2
V	43	7
Cr	48	40
Co	6	3
Ni	<20	<20
Cu	<10	20
Zn	49	120
Ga	10	28
Ga	10	1 4
Dh	1.2 54	1. <del>4</del> 74
KU C	J4 07	74
Sr	97	232
Y	43.7	75.9
Zr	603	891
Nb	23.6	27.5
Mo	<2	3
Cs	0.4	0.5
Ba	1051	1787
Hf	14.7	21.3
Та	2.26	1.56
W	13	<05
TI	0.33	0.33
Dh II	17	1/
D;	17 <01	1 <del>4</del> ∠0.1
DI Th	<0.1 15 5	<0.1 1 77
In	15.5	1.//
U	1.88	0.65
La	81.79	40.70
Ce	180.05	97.10
Pr	16.39	13.60
Nd	63.71	60.40
Sm	11.65	14.30
Eu	2.15	5.17
Gd	8.97	14.70
Th	1 45	2.52
Dv	8 <b>7</b> 1	1/ 80
Dy Uc	0.21	14.00
п0 Г	1.01	∠.ðð
Er	4.64	8.38
Tm	0.72	1.20
Yb	4.50	7.59
Lu	0.67	1.19

<x: below the detection limit (x)

Table 2

s	ample	Sm	Nd	Sm/Nd	<sup>147</sup> Sm/ <sup>144</sup> Nd	<sup>143</sup> Nd/ <sup>144</sup> Nd	<sup>143</sup> Nd/ <sup>144</sup> Nd <sub>i</sub>	8Ndi	TDM	2TDM	Rb	Sr	Rb/Sr	<sup>87</sup> Rb/ <sup>86</sup> Sr	<sup>87</sup> Sr/ <sup>86</sup> Sr	$^{87}Sr/^{86}Sr_i$	٤Sr <sub>i</sub>
N	MAZ-12040	14.3	60.4	0.2368	0.1431	0.512358	0.511564	+0.3	1459	1446	74	232	0.3190	0.9238	0.717962	0.706810	+47
F	ESP-7066	11.7	63.7	0.1837	0.1110	0.512084	0.511469	-1.5	1413	1595	54	97	0.5567	1.6136	0.726144	0.706666	+45









3\*Ga



#### Electron microprobe analyses

Analyses were performed with a JEOL JXA 8900M microprobe using with 15 kV, 20 nA (20 kV and 150 nA for epidote) and a beam diameter of 5  $\mu$ m. An element was considered to be present if its concentration exceeded twice the calculated detection limit.

Standards included albite (Si, Na), sillimanite (Al), almandine (Fe and Mn), kaersutite (Mg, Ca and Ti), potassium feldspar (K), fluorapatite (F, Cl, P), zircon (Hf, Zr), and synthetic compounds (U, Th, REE, Y, Ni, Cr).

Wt. %	Microcline	Plagioclase		Biotite	Titanite		Amphibole	Garnet		
	#6	#5	#29	#22	#47	#52	#41	#16	#11	
$SiO_2$	63.87	61.65	65.90	33.27	29.83	29.44	37.45	37.69	37.63	
TiO <sub>2</sub>	< 0.08	$<\!\!0.08$	< 0.08	2.68	36.51	35.51	1.45	0.06	0.21	
$Al_2O_3$	18.65	23.81	21.51	16.58	1.73	2.07	13.57	20.65	20.42	
$Fe_2O_3$	< 0.09	< 0.09	0.20		1.00	1.52	2.71	17.27	16.70	
FeO				31.21			25.67	9.21	11.09	
MnO	< 0.08	$<\!\!0.08$	< 0.08	0.35	0.09	0.05	0.27	6.09	4.73	
MgO	< 0.04	< 0.04	< 0.04	2.24	< 0.05	< 0.05	2.17	0.28	0.47	
CaO	< 0.04	5.28	3.06	< 0.04	29.78	28.88	11.33	11.57	11.02	
ZnO				0.26	< 0.16	< 0.16	0.00	< 0.14	< 0.14	
$Y_2O_3$					0.21	1.61	< 0.05			
Na <sub>2</sub> O	0.55	8.51	9.75	< 0.03	< 0.03	0.04	1.43	< 0.03	< 0.03	
$K_2O$	16.16	0.38	0.07	9.24			2.04			
F				< 0.14	0.21	0.26	0.00	< 0.14	< 0.14	
Cl				0.03	0.06	< 0.02	0.06			
$H_2O_{calc}$				3.69	0.31	0.41	1.86			
Total	99.23	99.63	100.49	99.53	99.72	99.79	99.99	102.82	102.27	
Formula	(apfu)									
Si	2.979	2.747	2.884	2.696	0.971	0.963	5.986	2.993	3.004	
Ti	0.000	0.000	0.000	0.164	0.894	0.874	0.174	0.003	0.013	
Al	1.025	1.251	1.109	1.584	0.066	0.080	2.557	1.933	1.921	
Fe <sup>3+</sup>	0.000	0.000	0.006		0.025	0.037	0.326	1.032	1.003	
Fe <sup>2+</sup>				2.115			3.431	0.612	0.740	
Mn	0.000	0.000	0.000	0.024	0.002	0.001	0.036	0.410	0.319	
Mg	0.000	0.000	0.000	0.271	0.000	0.000	0.516	0.034	0.056	
Ca	0.000	0.252	0.143	0.000	1.038	1.013	1.941	0.984	0.942	
Zn				0.015	0.000	0.000	0.000			
Y				0.000	0.004	0.028		0.000	0.000	
Na	0.049	0.735	0.828	0.000	0.000	0.003	0.443			
Κ	0.961	0.022	0.004	0.955			0.416			
F					0.021	0.027	0.000			
Cl				0.004	0.003	0.000	0.017			
$\mathrm{OH}_{\mathrm{calc}}$				1.996	0.066	0.090	1.983			

Selected electron microprobe analyses of minerals from sample MAZ-6037

Plagioclase: #5: core; #29: rim. Titanite: #47: core; #52: rim. Garnet: #16: core; #11: rim. Blank: not measured. <x: less than the stated detection limit (at  $2\sigma$ ). Sought but not found (detection limit in ppm,  $2\sigma$ ): Ni (660), Cr (450). H<sub>2</sub>O calculated by stoichiometry. Microcline, plagioclase and titanite: all Fe expressed as Fe<sub>2</sub>O<sub>3</sub>. Biotite: all Fe expressed as FeO. Amphibole: Fe<sup>2+</sup>/Fe<sup>3+</sup> calculated after Leake et al. (1997). Garnet: Fe<sup>2+</sup>/Fe<sup>3+</sup> calculated based on 12 O and 8 cations, after Droop (1987). Structural formulae

calculated on the basis of 8 O (microcline and plagioclase), 11 O eq. (biotite), 3 cations (titanite), and 23 O eq. (amphibole).

Wt. %	#6	#2
SiO <sub>2</sub>	33.79	37.40
TiO <sub>2</sub>	0.27	0.15
$Al_2O_3$	21.97	24.31
$Fe_2O_3$	8.10	11.09
FeO	4.25	1.26
MnO	0.16	0.14
MgO	0.09	0.00
CaO	18.40	24.24
$P_2O_5$	0.03	
$Y_2O_3$	1.01	< 0.06
$La_2O_3$	1.58	
$Ce_2O_3$	3.83	
$Pr_2O_3$	0.47	
$Nd_2O_3$	2.18	
$Sm_2O_3$	0.42	
$Gd_2O_3$	0.42	
$Dy_2O_3$	0.26	
$Er_2O_3$	0.14	
$Yb_2O_3$	0.11	
$Lu_2O_3$	0.04	
$ThO_2$	0.03	
$H_2O_{calc}$	1.74	1.89
Total	99.31	100.47
Formula	(apfu)	0.055
Formula Si	(apfu) 2.903	2.966
Formula Si Ti	(apfu) 2.903 0.018	2.966 0.009
Formula Si Ti Al	(apfu) 2.903 0.018 2.225	2.966 0.009 2.272
Formula Si Ti Al $Fe^{3+}$ $E^{-2+}$	(apfu) 2.903 0.018 2.225 0.524 0.205	2.966 0.009 2.272 0.662
Formula Si Ti Al Fe <sup>3+</sup> Fe <sup>2+</sup>	(apfu) 2.903 0.018 2.225 0.524 0.305 0.012	2.966 0.009 2.272 0.662 0.083
Formula Si Ti Al Fe <sup>3+</sup> Fe <sup>2+</sup> Mn	(apfu) 2.903 0.018 2.225 0.524 0.305 0.012	2.966 0.009 2.272 0.662 0.083 0.009
Formula Si Ti Al Fe <sup>3+</sup> Fe <sup>2+</sup> Mn Mg	(apfu) 2.903 0.018 2.225 0.524 0.305 0.012 0.012	2.966 0.009 2.272 0.662 0.083 0.009 0.000 2.050
Formula Si Ti Al $Fe^{3+}$ $Fe^{2+}$ Mn Mg Ca	(apfu) 2.903 0.018 2.225 0.524 0.305 0.012 0.012 1.694 0.002	2.966 0.009 2.272 0.662 0.083 0.009 0.000 2.059
Formula Si Ti Al Fe <sup>3+</sup> Fe <sup>2+</sup> Mn Mg Ca P V	(apfu) 2.903 0.018 2.225 0.524 0.305 0.012 0.012 1.694 0.002 0.046	2.966 0.009 2.272 0.662 0.083 0.009 0.000 2.059 0.000
Formula Si Ti Al Fe <sup>3+</sup> Fe <sup>2+</sup> Mn Mg Ca P Y Lo	(apfu) 2.903 0.018 2.225 0.524 0.305 0.012 0.012 1.694 0.002 0.046 0.050	2.966 0.009 2.272 0.662 0.083 0.009 0.000 2.059 0.000 0.000 0.000
Formula Si Ti Al Fe <sup>3+</sup> Fe <sup>2+</sup> Mn Mg Ca P Y La Ca	(apfu) 2.903 0.018 2.225 0.524 0.305 0.012 0.012 1.694 0.002 0.046 0.050 0.121	2.966 0.009 2.272 0.662 0.083 0.009 0.000 2.059 0.000 0.000 0.000 0.000
Formula Si Ti Al Fe <sup>3+</sup> Fe <sup>2+</sup> Mn Mg Ca P Y La Ce Pr	(apfu) 2.903 0.018 2.225 0.524 0.305 0.012 0.012 1.694 0.002 0.046 0.050 0.121 0.015	2.966 0.009 2.272 0.662 0.083 0.009 0.000 2.059 0.000 0.000 0.000 0.000 0.000
Formula Si Ti Al Fe <sup>3+</sup> Fe <sup>2+</sup> Mn Mg Ca P Y La Ce Pr Nd	(apfu) 2.903 0.018 2.225 0.524 0.305 0.012 0.012 1.694 0.002 0.046 0.050 0.121 0.015 0.067	2.966 0.009 2.272 0.662 0.083 0.009 0.000 2.059 0.000 0.000 0.000 0.000 0.000 0.000
Formula Si Ti Al Fe <sup>3+</sup> Fe <sup>2+</sup> Mn Mg Ca P Y La Ce Pr Nd Sm	(apfu) 2.903 0.018 2.225 0.524 0.305 0.012 0.012 1.694 0.002 0.046 0.050 0.121 0.015 0.067 0.013	2.966 0.009 2.272 0.662 0.083 0.009 0.000 2.059 0.000 0.000 0.000 0.000 0.000 0.000 0.000
Formula Si Ti Al Fe <sup>3+</sup> Fe <sup>2+</sup> Mn Mg Ca P Y La Ce Pr Nd Sm Cd	(apfu) 2.903 0.018 2.225 0.524 0.305 0.012 0.012 1.694 0.002 0.046 0.050 0.121 0.015 0.067 0.013 0.012	2.966 0.009 2.272 0.662 0.083 0.009 0.000 2.059 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000
Formula Si Ti Al Fe <sup>3+</sup> Fe <sup>2+</sup> Mn Mg Ca P Y La Ce Pr Nd Sm Gd Dy	(apfu) 2.903 0.018 2.225 0.524 0.305 0.012 0.012 1.694 0.002 0.046 0.050 0.121 0.015 0.067 0.013 0.012	2.966 0.009 2.272 0.662 0.083 0.009 0.000 2.059 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000
Formula Si Ti Al Fe <sup>3+</sup> Fe <sup>2+</sup> Mn Mg Ca P Y La Ce Pr Nd Sm Gd Dy Fr	(apfu) 2.903 0.018 2.225 0.524 0.305 0.012 0.012 1.694 0.002 0.046 0.050 0.121 0.015 0.067 0.013 0.012 0.007 0.004	2.966 0.009 2.272 0.662 0.083 0.009 0.000 2.059 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000
Formula Si Ti Al Fe <sup>3+</sup> Fe <sup>2+</sup> Mn Mg Ca P Y La Ce Pr Y La Ce Pr Nd Sm Gd Dy Er Yb	(apfu) 2.903 0.018 2.225 0.524 0.305 0.012 0.012 1.694 0.002 0.046 0.050 0.121 0.015 0.067 0.013 0.012 0.007 0.004 0.003	2.966 0.009 2.272 0.662 0.083 0.009 0.000 2.059 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000
Formula Si Ti Al Fe <sup>3+</sup> Fe <sup>2+</sup> Mn Mg Ca P Y La Ce Pr Nd Sm Gd Dy Er Yb Lu	(apfu) 2.903 0.018 2.225 0.524 0.305 0.012 0.012 1.694 0.002 0.046 0.050 0.121 0.015 0.067 0.013 0.012 0.007 0.004 0.003 0.001	2.966 0.009 2.272 0.662 0.083 0.009 0.000 2.059 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000
Formula Si Ti Al Fe <sup>3+</sup> Fe <sup>2+</sup> Mn Mg Ca P Y La Ce Pr Nd Sm Gd Dy Er Yb Lu Th	(apfu) 2.903 0.018 2.225 0.524 0.305 0.012 0.012 1.694 0.002 0.046 0.050 0.121 0.015 0.067 0.013 0.012 0.007 0.004 0.003 0.001 0.001	2.966 0.009 2.272 0.662 0.083 0.009 0.000 2.059 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000
Formula Si Ti Al Fe <sup>3+</sup> Fe <sup>2+</sup> Mn Mg Ca P Y La Ce Pr Nd Sm Gd Dy Er Yb Lu Th OH	(apfu) 2.903 0.018 2.225 0.524 0.305 0.012 0.012 1.694 0.002 0.046 0.050 0.121 0.015 0.067 0.013 0.012 0.007 0.004 0.003 0.001 0.001 1.000	2.966 0.009 2.272 0.662 0.083 0.009 0.000 2.059 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000

Electron microprobe analyses of epidote from sample MAZ-6037

#6: core, #2: rim. H<sub>2</sub>O calculated by stoichiometry to give 1 (OH)<sup>-</sup>. Fe<sup>2+</sup>/Fe<sup>3+</sup> calculated according to Ercit (2002). Blank: not measured. <x: less than the stated detection limit (at  $2\sigma$ ). Sought but not found (detection limit in ppm,  $2\sigma$ ):Tb (140), Ho (240), Tm (1200), U (100), Pb (80), Zr (120), Hf (100), F (1200), Cl (220), Cr (480), Na (260), K (260), Zn (1400).

*Electron microprobe analyses of minerals from an amphibolite (sample MAZ-12046) used for thermobarometric calculations.* 

Wt. %		Bio	otite					Ampł	ibole			
	#125	#130	#142	#151	#124	#126	#128	#132	#133	#137	#141	#154
$SiO_2$	35.55	35.44	34.95	36.20	40.24	41.52	40.84	40.83	40.80	40.47	40.95	40.44
$TiO_2$	1.95	2.42	2.16	2.26	0.84	0.68	0.63	0.56	0.74	0.68	0.66	0.59
$Al_2O_3$	17.18	16.67	17.54	17.44	15.50	15.28	15.26	15.64	15.05	15.52	15.47	16.02
$Fe_2O_3$					3.33	2.28	2.31	1.87	3.46	0.83	1.83	0.95
FeO	19.26	20.25	23.09	20.34	16.49	17.03	17.00	17.16	16.69	17.22	16.89	18.13
MnO	0.05	0.10	0.09	0.13	0.20	0.17	0.13	0.09	0.20	0.20	0.18	0.17
MgO	10.01	9.85	7.57	9.75	7.37	7.40	7.27	7.13	7.39	7.07	7.29	6.50
CaO	0.18	0.15	0.07	0.11	11.75	11.47	11.72	11.71	11.57	11.85	11.69	11.80
ZnO	0.20	< 0.12	< 0.12	< 0.12	< 0.12	< 0.12	< 0.12	< 0.12	< 0.12	0.17	< 0.12	0.14
Na <sub>2</sub> O	0.09	0.14	0.14	0.14	1.75	1.67	1.67	1.66	1.76	1.67	1.64	1.74
K <sub>2</sub> O	8.68	8.76	9.30	9.43	0.80	0.61	0.65	0.62	0.67	0.67	0.59	0.65
Cl	0.09	0.09	0.09	0.08	0.05	0.06	0.11	0.08	0.07	0.07	0.08	0.07
H2O calc.	3.83	3.84	3.82	3.92	1.98	1.98	1.95	1.96	1.97	1.94	1.96	1.95
Total	97.06	97.70	98.79	99.78	100.29	100.14	99.53	99.32	100.36	98.35	99.22	99.13
Formula	(apfu)											
Si	2.763	2.752	2.725	2.753	6.062	6.228	6.182	6.183	6.137	6.192	6.199	6.160
Ti	0.114	0.142	0.127	0.129	0.096	0.076	0.071	0.064	0.084	0.079	0.075	0.067
Al	1.574	1.526	1.612	1.563	2.752	2.700	2.723	2.792	2.668	2.799	2.760	2.875
Fe <sup>3+</sup>					0.378	0.258	0.263	0.213	0.392	0.096	0.208	0.109
Fe <sup>2+</sup>	1.252	1.315	1.506	1.294	2.078	2.136	2.152	2.173	2.100	2.204	2.139	2.310
Mn	0.003	0.007	0.006	0.008	0.025	0.021	0.017	0.011	0.026	0.026	0.022	0.022
Mg	1.160	1.140	0.880	1.106	1.656	1.655	1.639	1.610	1.657	1.612	1.645	1.476
Zn	0.011	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.019	0.000	0.016
Ca	0.015	0.013	0.006	0.009	1.897	1.842	1.901	1.900	1.864	1.943	1.896	1.926
Na	0.013	0.021	0.022	0.021	0.510	0.486	0.489	0.487	0.515	0.494	0.482	0.515
Κ	0.861	0.868	0.925	0.915	0.155	0.116	0.125	0.120	0.128	0.130	0.113	0.126
Cl	0.012	0.011	0.012	0.011	0.011	0.016	0.029	0.021	0.017	0.017	0.021	0.018
OH calc	1.988	1.989	1.988	1.989	1.989	1.984	1.971	1.979	1.983	1.983	1.979	1.982

<x: less than the stated detection limit (at  $2\sigma$ ). H<sub>2</sub>O calculated by stoichiometry. Biotite: all Fe expressed as FeO. Amphibole: Fe<sup>2+</sup>/Fe<sup>3+</sup> calculated after Leake et al. (1997). Structural formulae calculated on the basis of 11 O eq. (biotite) and 23 O eq. (amphibole).

Wt. %					Garnet				
	#121	#122	#127	#129	#135	#143	#144	#152	#153
$SiO_2$	38.07	37.76	38.14	38.14	37.84	37.81	37.59	38.04	38.21
$TiO_2$	0.13	0.14	0.05	0.00	0.00	0.07	0.11	0.14	0.00
$Al_2O_3\\$	21.46	21.19	21.62	21.25	21.63	21.72	21.48	21.81	21.39
$Fe_2O_3$	0.00	0.00	0.00	0.00	0.00	0.00	0.94	0.62	0.30
FeO	29.46	29.02	29.38	29.42	29.10	25.51	27.53	30.94	29.43
MnO	2.12	2.60	2.67	2.40	2.56	2.35	2.46	1.27	1.88
MgO	3.06	2.63	2.69	2.99	2.60	2.63	2.53	3.29	2.96
CaO	6.22	6.55	6.77	6.47	6.75	7.69	7.12	5.74	7.09
Na <sub>2</sub> O	< 0.03	< 0.03	< 0.03	< 0.03	< 0.03	< 0.03	0.30	0.03	< 0.03
Total	100.52	99.89	101.33	100.66	100.47	97.79	100.06	101.88	101.26
Formula	(apfu)								
Si	3.007	3.009	2.998	3.013	2.997	3.032	2.984	2.972	3.001
Ti	0.008	0.008	0.003	0.000	0.000	0.004	0.006	0.008	0.000
Al	1.998	1.990	2.003	1.979	2.019	2.054	2.010	2.008	1.980
Fe <sup>3+</sup>	0.000	0.000	0.000	0.000	0.000	0.000	0.056	0.036	0.018
Fe <sup>2+</sup>	1.946	1.934	1.931	1.944	1.928	1.711	1.827	2.022	1.933
Mn	0.142	0.175	0.178	0.160	0.172	0.160	0.165	0.084	0.125
Mg	0.360	0.312	0.316	0.352	0.307	0.315	0.299	0.383	0.347
Ca	0.526	0.559	0.570	0.548	0.572	0.661	0.606	0.480	0.596
Na	0.000	0.000	0.000	0.000	0.000	0.000	0.046	0.005	0.000

 $$<\!x$: less than the stated detection limit (at <math display="inline">2\sigma$ ).  $Fe^{2+}/Fe^{3+}$  calculated based on 12 O and 8 cations, after Droop (1987).

Wt. %		Plag	gioclase		Ilmenite
	#131	#136	#139	#140	#222
$SiO_2$	57.37	59.89	59.56	57.07	0.10
TiO <sub>2</sub>					46.76
$Al_2O_3$	26.24	24.72	25.11	27.07	1.55
Fe <sub>2</sub> O <sub>3</sub>	0.09	0.15	0.32	0.04	9.13
FeO					41.46
MnO					0.37
MgO	< 0.03	< 0.03	< 0.03	< 0.03	0.08
CaO	8.32	6.89	6.85	9.57	0.17
Na <sub>2</sub> O	7.01	7.90	8.02	6.31	
K <sub>2</sub> O	0.05	0.04	0.07	0.09	
Total	99.07	99.59	99.92	100.15	99.60
Formula	(apfu)				
Si	2.593	2.681	2.661	2.558	0.002
Ti					0.888
Al	1.398	1.304	1.322	1.430	0.046
Fe <sup>3+</sup>	0.003	0.005	0.011	0.001	0.173
Fe <sup>2+</sup>					0.875
Mn					0.008
Mg					0.003
Zn					0.004
Ca	0.403	0.331	0.328	0.460	
Na	0.614	0.685	0.695	0.548	
Κ	0.003	0.002	0.004	0.005	

Blank: not measured. <x: less than the stated detection limit (at  $2\sigma$ ). Plagioclase: all Fe expressed as Fe<sub>2</sub>O<sub>3</sub>. Structural formulae calculated on the basis of 8 O (plagioclase) and 2 cations (ilmenite).

										Radiogenic	Ratios							Age (Ma)		
Grain.	Zircon	U	Th	Th/U	Pb*	<sup>204</sup> Pb/	<b>f</b> <sub>206</sub>	<sup>206</sup> Pb/		<sup>207</sup> Pb/			<sup>207</sup> Pb/			<sup>206</sup> Pb/		<sup>207</sup> Pb/		%
spot	type	(ppm)	(ppm)		(ppm)	<sup>206</sup> Pb	%	<sup>238</sup> U	±	<sup>235</sup> U		±	<sup>206</sup> Pb	±	ρ	<sup>238</sup> U	±	<sup>206</sup> Pb	±	Disc
MAZ-6037																				
1,1	weakly zoned outer	100	45	0,45	12,5	0,000263	0,46	0,1447	0,0019		1,330	0,035	0,0667	0,0015	0,500	871	11	829	48	-5
1,2	core	85	60	0,71	10,3	0,000159	0,27	0,1397	0,0019		1,298	0,031	0,0674	0,0013	0,580	843	11	850	40	1
2,1	weakly zoned outer	128	47	0,37	15,8	0,000664	1,16	0,1421	0,0019		1,268	0,077	0,0647	0,0038	0,216	857	10	765	124	-11
3,1	weakly zoned outer	46	24	0,52	5,8	0,000840	1,48	0,1434	0,0023		1,241	0,076	0,0627	0,0037	0,260	864	13	700	126	-19
*4.1	weakly zoned	137	74	0,54	17,2	0,000028	0,05	0,1457	0,0018		1,360	0,028	0,0677	0,0011	0,590	877	10	859	35	-2
7,1	osczoned outer	159	90	0,57	18,9	0,000069	0,12	0,1379	0,0018		1,287	0,032	0,0677	0,0014	0,519	833	10	860	44	3
8,1	osczoned outer	139	50	0,36	16,4	0,000264	0,46	0,1366	0,0017		1,243	0,037	0,0660	0,0018	0,423	825	10	806	57	-2
*8.2	core	680	227	0,33	84,2	0,000033	0,06	0,1441	0,0015		1,340	0,017	0,0674	0,0004	0,857	868	9	851	13	-2
9,1	osczoned outer	181	85	0,47	21,8	0,000195	0,34	0,1396	0,0017		1,275	0,028	0,0663	0,0012	0,562	842	10	814	38	-3
10,1	osczoned outer	121	66	0,54	14,5	0,000162	0,28	0,1391	0,0018		1,303	0,031	0,0679	0,0013	0,555	840	10	866	41	3
11,1	core	90	45	0,50	11,0	0,000500	0,87	0,1402	0,0021		1,282	0,056	0,0663	0,0027	0,349	846	12	817	86	-3
12,1	osczoned outer	175	89	0,51	21,4	0,000157	0,27	0,1414	0,0018		1,347	0,031	0,0691	0,0013	0,548	853	10	901	39	6
13,1	core	122	71	0,58	14,5	0,000390	0,68	0,1379	0,0018		1,251	0,046	0,0658	0,0023	0,353	833	10	799	72	-4
14,1	core	551	210	0,38	67,7	0,000085	0,15	0,1428	0,0016		1,302	0,018	0,0661	0,0006	0,770	861	9	810	19	-6
*15.1	osczoned outer	91	46	0,51	10,5	0,000330	0,57	0,1340	0,0025		1,230	0,043	0,0666	0,0019	0,539	811	14	825	61	2
*16.1	osczoned outer	167	88	0,53	19,0	0,000544	0,95	0,1310	0,0016		1,165	0,044	0,0645	0,0023	0,330	794	9	758	76	-5
*17.1	core	1060	549	0,52	132,3	0,000025	0,04	0,1452	0,0016		1,345	0,017	0,0672	0,0004	0,880	874	9	843	13	-4
18,1	osczoned outer	173	75	0,43	20,7	0,000280	0,49	0,1382	0,0017		1,225	0,035	0,0643	0,0017	0,427	834	10	752	55	-10
18,2	core	910	470	0,52	109,2	0,000029	0,05	0,1396	0,0016		1,304	0,017	0,0678	0,0004	0,886	842	9	862	13	2
19,1	weakly zoned outer	113	44	0,39	14,0	0,000101	0,18	0,1442	0,0019		1,328	0,030	0,0668	0,0012	0,588	868	11	831	38	-4
20,1	osczoned outer	157	65	0,42	18,9	0,000278	0,48	0,1394	0,0020		1,285	0,038	0,0669	0,0017	0,484	841	11	833	54	-1
*21.1	osczoned outer	148	82	0,56	18,6	0,000078	0,13	0,1465	0,0021		1,389	0,032	0,0688	0,0012	0,621	881	12	891	37	1
22,1	osczoned outer	97	51	0,53	11,4	0,000114	0,20	0,1366	0,0019		1,265	0,033	0,0672	0,0015	0,523	825	11	843	46	2
23,1	osczoned outer	151	78	0,52	18,1	0,000168	0,29	0,1393	0,0019		1,266	0,030	0,0659	0,0013	0,586	841	11	803	40	-4
24,1	osczoned outer	137	78	0,57	16,7	0,000104	0,18	0,1418	0,0018		1,293	0,036	0,0661	0,0016	0,474	855	10	811	51	-5

# SHRIMP U-Pb data for samples MAZ-6037 and ESP-7066.

Grain.	Zircon	U	Th	Th/U	Pb*	<sup>204</sup> Pb/	f <sub>206</sub>	<sup>206</sup> Pb/		<sup>207</sup> Pb/		<sup>207</sup> Pb/			<sup>206</sup> Pb/		<sup>207</sup> Pb/		%
spot	type	(ppm)	(ppm)		(ppm)	<sup>206</sup> Pb	%	<sup>238</sup> U	±	<sup>235</sup> U	±	<sup>206</sup> Pb	±	ρ	<sup>238</sup> U	±	<sup>206</sup> Pb	±	Disc
MAZ-6037	(continued)																		
25,1	osczoned outer	163	80	0,49	20,1	0,00083	0,14	0,1432	0,0018	1,34	0 0,033	0,0679	0,0014	0,503	863	10	864	44	0
26,1	osczoned outer	135	79	0,58	16,7	0,000264	0,46	0,1438	0,0020	1,29	6 0,038	0,0654	0,0017	0,485	866	11	787	53	-9
27,1	osczoned outer	325	131	0,40	38,7	0,000147	0,26	0,1383	0,0016	1,28	8 0,021	0,0660	0,0008	0,685	835	9	807	25	-3
	Error in FC1 reference	zircon o	calibratio	n was	0.40%, r	not included	in above	errors but	required v	when comparing <sup>206</sup> Pb/ <sup>238</sup>	J data fro	m differer	nt analytic	cal sessio	ns.				
ESP-7066																			
*1.1	osczoned	279	18	0,07	42,3	0,000034	0,06	0,1768	0,0020	1,79	5 0,023	0,0736	0,0005	0,863	1049	11	1031	13	-2
*1.2	weakly zoned, late	305	129	0,42	53,3	0,000002	<0.01	0,2034	0,0022	2,25	5 0,027	0,0804	0,0004	0,896	1193	12	1207	10	1
2,1	weakly zoned	259	48	0,18	31,0	0,000125	0,22	0,1391	0,0015	1,20	9 0,021	0,0662	0,0008	0,663	840	9	812	26	-3
2,2	weakly zoned, late	111	56	0,51	13,5	0,000335	0,59	0,1421	0,0017	1,3 <sup>-</sup>	9 0,022	0,0673	0,0007	0,738	857	11	847	23	-1
3,1	weakly zoned, late	828	161	0,19	98,4	0,000014	0,02	0,1384	0,0014	1,28	0,014	0,0672	0,0003	0,918	836	8	843	9	1
*4.1	embayed overgrowth	530	8	0,01	30,5	0,000118	0,21	0,0669	0,0007	0,5 <sup>,</sup>	0 0,008	0,0553	0,0006	0,715	417	4	425	24	2
4,2	osczoned	421	209	0,50	50,5	0,000070	0,12	0,1395	0,0015	1,30	0,016	0,0677	0,0004	0,880	842	9	859	12	2
5,1	zoned tip	271	139	0,51	31,8	0,000196	0,34	0,1369	0,0015	1,28	4 0,017	0,0680	0,0005	0,841	827	9	870	15	5
6,1	weakly zoned outer	539	192	0,36	64,1	0,000000	<0.01	0,1385	0,0015	1,28	5 0,015	0,0673	0,0004	0,885	836	8	846	12	1
*7.1	embayed overgrowth	525	9	0,02	30,8	0,000060	0,11	0,0683	0,0008	0,52	3 0,010	0,0556	0,0009	0,567	426	5	434	36	2
8,1	zoned tip	181	113	0,62	21,9	0,000048	0,08	0,1409	0,0016	1,32	1 0,024	0,0680	0,0009	0,649	849	9	870	28	2
*3.2	zoned core	185	90	0,49	41,5	0,000013	0,02	0,2617	0,0030	3,34	7 0,044	0,0927	0,0006	0,864	1499	15	1483	13	-1
*10.1	weakly zoned, late	269	75	0,28	27,8	0,000030	0,05	0,1201	0,0013	1,09	3 0,019	0,0660	0,0009	0,642	731	8	807	28	9
11,1	weakly zoned, late	145	36	0,25	17,1	0,000008	0,01	0,1377	0,0016	1,27	5 0,020	0,0671	0,0007	0,739	832	9	842	22	1
*12.1	weakly zoned, late	672	57	0,08	93,7	0,00008	0,01	0,1623	0,0017	1,6 <sup>-</sup>	4 0,018	0,0721	0,0003	0,916	970	9	988	9	2
*13.1	patchy zoning, round	587	33	0,06	87,6	0,000013	0,02	0,1737	0,0018	1,75	3 0,021	0,0732	0,0004	0,892	1033	10	1019	11	-1
14,1	osczoned core	45	31	0,68	5,4	0,000445	0,78	0,1401	0,0022	1,27	9 0,034	0,0662	0,0014	0,598	845	14	813	44	-4

Error in FC1 reference zircon calibration was 0.16%, not included in above errors but required when comparing <sup>206</sup>Pb/<sup>238</sup>U data from different analytical sessions.

Notes : 1. Uncertainties given at the  $1\sigma$  level.

- 2.  $f_{206}$  % denotes the percentage of <sup>206</sup>Pb that is common Pb.
- 3. Correction for common Pb made using the measured <sup>204</sup>Pb/<sup>206</sup>Pb ratio.
- 4. For % Disc., 0% denotes a concordant analysis.
- 5. Asterisk indicates data not used in calculation Concordia age as in Ludwig (2001)