

# The Difunta Correa metasedimentary sequence (NW Argentina): relict of a Neoproterozoic platform? — elemental and Sr-Nd isotope evidence

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## ABSTRACT

The Sierra de Pie de Palo (Western Sierras Pampeanas, Argentina) in the Andean foreland is mainly formed by a Mesoproterozoic basement and an Ediacaran metasedimentary cover referred to as the Difunta Correa metasedimentary sequence. The latter is key to understanding the characteristics of this region prior to the early Cambrian assembly of SW Gondwana. It is composed of low- to medium grade metamorphic rocks (metasandstones, mica-schists, Ca-pelitic schists, metaconglomerates, marbles and less abundant amphibolites) that can be grouped into four informal lithostratigraphic units. The chemical composition of these rocks allows to classify the siliciclastic protoliths as shales, Fe-shales and immature sandstones (wackes, sub-litharenites, litharenites and Fe-sandstones). The sediments were derived from an evolved felsic to intermediate continental source and were deposited on a continental passive margin overlaying a Mesoproterozoic basement that crops out at several places of the Western Sierras Pampeanas. Thick marine carbonate beds with seawater isotope composition, phosphatic clasts and the lack of contemporaneous, arc related igneous rocks, also support a passive margin sedimentation. Phosphatic clasts within metaconglomerates are described for the first time in the Sierras Pampeanas and were probably formed after an important Neoproterozoic glaciation (Marinoan). We further suggest, based on our data and previous works, that the passive margin probably belonged to the Paleoproterozoic MARA (acronym of Maz, Arequipa, Río Apa) continental block. MARA, which remained juxtaposed to Laurentia since the middle to late Mesoproterozoic orogenies until its eventual drifting in the late Neoproterozoic, finally accreted to SW Gondwana in early Cambrian times during the Pampean orogeny.

Key words: geochemistry; metasedimentary rocks; neoproterozoic; Western Sierras Pampeanas; phosphatic clasts; Laurentia.

## RESUMEN

*La Sierra de Pie de Palo (Sierras Pampeanas Occidentales, Argentina), situada en el antepaís Andino, está formada principalmente por un basamento mesoproterozoico y una cubierta metasedimentaria ediacareense. Esta cubierta, denominada secuencia metasedimentaria Difunta Correa, es una unidad clave para entender la paleogeografía de*

*esta región previo al ensamble del suroeste de Gondwana en el Cámbrico inferior. Esta unidad está compuesta por rocas metamórficas de grado bajo a medio (meta-arenitas, esquistos micáceos, esquistos pelíticos cálcicos, metaconglomerados, mármoles y escasas anfibolitas), las cuales pueden agruparse en cuatro unidades litoestratigráficas informales. La composición química de las rocas permite clasificar a los protolitos siliciclásticos como pelitas, Fe-pelitas y areniscas inmaduras (grauvacas, sub-litoarenitas, litoarenitas y areniscas ferruginosas). Los sedimentos se depositaron en un margen continental pasivo y son el resultado de la erosión de una fuente cortical félsica a intermedia, isotópicamente evolucionada. La existencia de potentes capas de carbonatos con composición isotópica de Sr de origen marino, sumado a la presencia de clastos fosfáticos y la falta de rocas ígneas de arco contemporáneas, apoyan la hipótesis de una sedimentación en un margen pasivo sobre un basamento mesoproterozoico que aflora en diversos lugares de las Sierras Pampeanas Occidentales. Se describen aquí, por primera vez, clastos fosfáticos en las Sierras Pampeanas que podrían estar vinculados a una de las principales glaciaciones neoproterozoicas (Marinoense). Además sugerimos, basados en nuestros datos y en trabajos previos, que dicho margen pasivo probablemente formó parte del bloque continental paleoproterozoico MARA (acrónimo de Maz, Arequipa y Río Apa). Este bloque permaneció unido a Laurentia desde las orogénesis del Mesoproterozoico medio y superior, hasta su deriva a fines del Neoproterozoico y acreción al margen suroeste de Gondwana en el Cámbrico inferior (orogenia Pampeana).*

Palabras clave: geoquímica de rocas metasedimentarias; neoproterozoico; Sierras Pampeanas Occidentales; clastos fosfáticos; Laurentia.

## INTRODUCTION

The Sierra de Pie de Palo (SPP) is one of the Western Sierras Pampeanas (WSP) of Argentina which resulted from Cenozoic uplift in response to shortening of the Andean foreland (Jordan and Allmendinger, 1986; Isacks, 1988). The WSP mainly consists of middle to late Mesoproterozoic basement and a late Neoproterozoic (Ediacaran) sedimentary cover (McDonough *et al.*, 1993; Casquet *et al.*, 2001; Galindo *et al.*, 2004; Rapela *et al.*, 2005). Basement and cover were overprinted by metamorphism and penetrative deformation during the widespread early to middle Ordovician accretionary Famatinian

orogeny (Casquet *et al.*, 2001; Mulcahy *et al.*, 2011; van Staal *et al.*, 2011). The metasedimentary succession was recognized and named by Baldo *et al.* (1998) as the Difunta Correa metasedimentary sequence (DCMS) and represents a poorly known Neoproterozoic basin. Likely, the DCMS also crops out in other ranges of the WSP, namely Maz, Espinal and Umango (Figure 1a) where a mesoproterozoic basement has also been recognized (Casquet *et al.*, 2008; Varela *et al.*, 2011). However, the original relationships between the basement and the metasedimentary cover are always obscured by shearing and faulting.

The DCMS constitutes a key entity to the paleogeography of the WSP during the late Neoproterozoic, before its accretion to southwestern Gondwana in the early Cambrian (Casquet *et al.*, 2012; Rapela *et al.*, 2015). These authors suggest that the WSP Mesoproterozoic basement developed after the MARA Paleoproterozoic craton (MARA, acronym of Maz, Arequipa, Río Apa). This craton was accreted to Laurentia -and

consequently reworked- in the middle to late Mesoproterozoic orogenies between *ca.* 1.3 and 1.0 Ga. MARA drifted away from Laurentia in the late Neoproterozoic, resulting in the opening of the Iapetus Ocean. This basement was further involved in the early Cambrian Pampean orogeny of the Eastern Sierras Pampeanas by collision with other southwestern Gondwana cratons (such as Kalahari). This collision resulted in the closure of the hypothetical Clymene Ocean (Trindade *et al.*, 2006), and its southeastern extension (present position), *i.e.*, Puncoviscana Ocean.

Conventional petrological provenance studies of detrital rocks (sandstones; Dickinson and Suczek, 1979; Zuffa, 1985; Jonhsson, 1993; Arribas *et al.*, 2007) are hindered if the rocks underwent significant metamorphism and deformation. Primary components of the sedimentary rocks are deeply modified by processes such as recrystallization, formation of new minerals and development of a

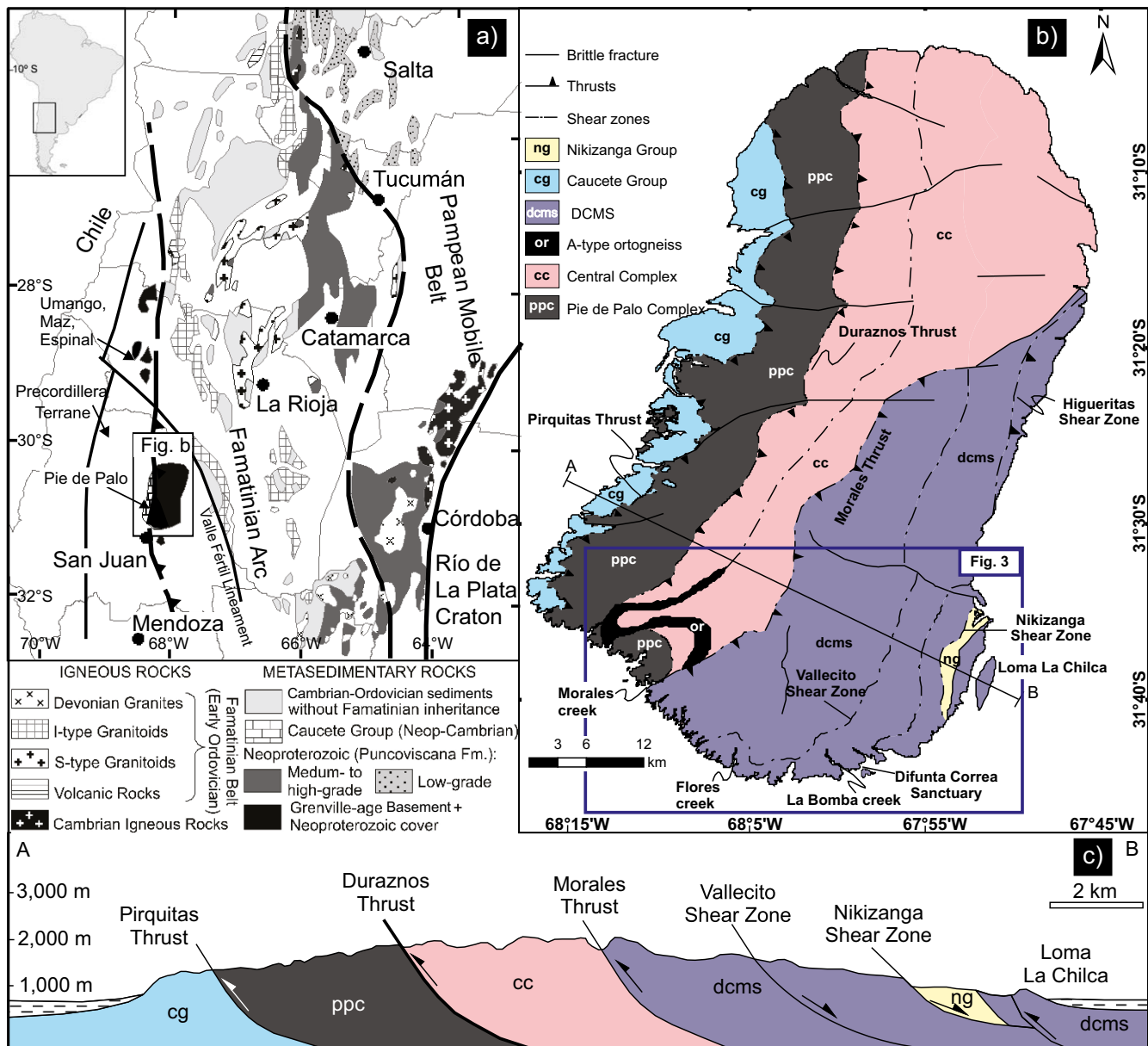


Figure 1. a) Geological setting of the Sierras Pampeanas showing the location of the Sierra de Pie de Palo (SPP); b) Geological map of the SPP; c) Schematic east-west cross-section of the SPP showing the main geological units and structural boundaries. DCMS: Difunta Correa metasedimentary sequence.

new tectonic fabric (foliation). In such cases, methods based on the chemical composition of the metasedimentary rocks help to define the source areas and to establish the tectonic setting of sedimentary basins (Bhatia and Crook, 1986; McLennan *et al.*, 1990, 2003; Zimmermann *et al.*, 2011). However, chemical data alone can lead to erroneous interpretation of the tectonic setting, especially in ancient basins (Armstrong-Altrin and Verma, 2005; Verdecchia and Baldo, 2010). Therefore, they are supported here by complementary evidence from regional geological knowledge and detrital zircon data (Rapela *et al.*, 2015).

Some major and trace elements (e.g., Si, Al, Fe, Mg, Mn, Ca, K, Ba, Sr, Rb and Cs) can be removed from sediments during weathering, diagenesis and low- to medium- grade metamorphism, and therefore they are not appropriate for constraining source areas and/or tectonic settings. On the other hand, elements such as Ti, La, Ce, Nd, Y, Th, Zr, Hf, Nb, and Sc within resistant detrital minerals preserve the initial inter-elemental ratios, and, in consequence, they are useful for provenance analyses (Bhatia and Crook, 1986; Floyd and Leveridge, 1987; McLennan, 1989; McLennan *et al.*, 1990; Zimmermann and Bahlburg, 2003).

The aim of this research is to infer source areas and the tectonic setting of the DCMS through regional, chemical, and isotope evidences of significant rock types, and to discuss the paleogeographic implications. Moreover, we present here the first mention of phosphatic clasts in the Sierras Pampeanas and evaluate their paleoclimatic and paleogeographic significance.

## GEOLOGICAL SETTING

The SPP consists of five lithological assemblages bounded by east-dipping shear zones and thrusts. From west to the east they are: the Caucete Group, the Pie de Palo Complex, the Central Complex, the DCMS, and the Nikizanga Group (Figure 1). All these units are overprinted by metamorphism and intruded by igneous rocks, both assigned to the Ordovician Famatinian orogeny (Pankhurst and Rapela, 1998; Casquet *et al.*, 2001; Mulcahy *et al.*, 2011; Baldo *et al.*, 2012).

The Caucete Group (Borrello, 1969; Vujovich, 2003; Galindo *et al.*, 2004; Naipauer *et al.*, 2010; van Staal *et al.*, 2011) is exposed along the western margin of the SPP, in the footwall of the major Piriquitas thrust (Figure 1b), and mainly consists of low- to medium-grade metamorphic rocks such as metasandstones and marbles, and to a lesser extent, metavolcaniclastic rocks. A late Neoproterozoic–early Cambrian depositional age was established for this group based on Sr-isotope composition of marbles (Galindo *et al.*, 2004) and U-Pb zircon ages from metasandstones (Naipauer *et al.*, 2010).

The Pie de Palo Complex (*sensu* Mulcahy *et al.*, 2011) lies between the Piriquitas and the Duraznos thrusts (Figure 1b and 1c) and is mainly composed of ultramafic and mafic rocks, largely metagabbros and garnet-amphibolites. Mesoproterozoic (ca. 1,067–1,204 Ma) U-Pb zircon crystallization ages were obtained for this complex (Vujovich *et al.*, 2004; Morata *et al.*, 2010; Rapela *et al.*, 2010; Mulcahy *et al.*, 2011). The Pie de Palo Complex is an igneous sequence formed in a back-arc setting (Vujovich and Kay, 1998).

The Caucete Group and the Pie de Palo Complex have been correlated with the cover and the basement, respectively, of the worldwide known Precordillera terrane. This correlation was based on Sr-isotope composition of marbles (Galindo *et al.*, 2004) and detrital zircon ages (Naipauer *et al.*, 2010) of Caucete Group, and on common Pb composition and geochronology of the ultramafic and mafic rocks of the Pie de Palo Complex (Abbruzzi *et al.*, 1993; Kay *et al.*, 1996). Many authors consider that the Precordillera is an exotic terrane of

Laurentian affinity that accreted to the proto-Andean margin during the Famatinian orogeny (Thomas and Astini, 2003; Ramos, 2004).

The Central Complex is defined here as a large block between the Duraznos and the Morales thrusts. The latter ductile thrust was recognized in the Morales creek (Figure 1b) and we were able to track it northward using satellite images. The boundary between the Central Complex and the DCMS is poorly known due to access difficulties. The Central Complex consists of schists, quartzites, marbles, migmatites, amphibolites, metavolcanic rocks and orthogneisses (Casquet *et al.*, 2001; Mulcahy *et al.*, 2011). One orthogneiss is an A-type granitoid of ca. 774 Ma attributed to the break-up of Rodinia (Baldo *et al.*, 2006). Other orthogneisses range from ca. 1,280 to ca. 1,027 Ma (McDonough *et al.*, 1993; Rapela *et al.*, 2010; Garber *et al.*, 2014), and therefore, the depositional age of the metasedimentary rocks of this complex has to be older than those of the igneous intrusions. In consequence, the Central Complex probably constitutes a mesoproterozoic basement.

The DCMS was described by Baldo *et al.* (1998) as a sequence made up of metapelites, Ca-pelitic schists, quartzites, quartz-feldspar metasandstones, marbles and para-amphibolites. The DCMS is widespread along the central and eastern parts of the SPP. A late Neoproterozoic depositional age for the DCMS was proposed by Galindo *et al.* (2004) on the basis that the Sr-isotope composition of the calcite marbles corresponds to sea-water composition at that time. The age of, at least, the lower and the middle part of the DCMS probably is ca. 600 Ma according to the more recent sea-water compilations and U-Pb detrital zircon ages (Rapela *et al.*, 2005, 2015; Murra *et al.*, 2014). We recognize four units in the DCMS; from bottom to top: La Loma, Flores, Vallecito, and La Bomba (Figure 2).

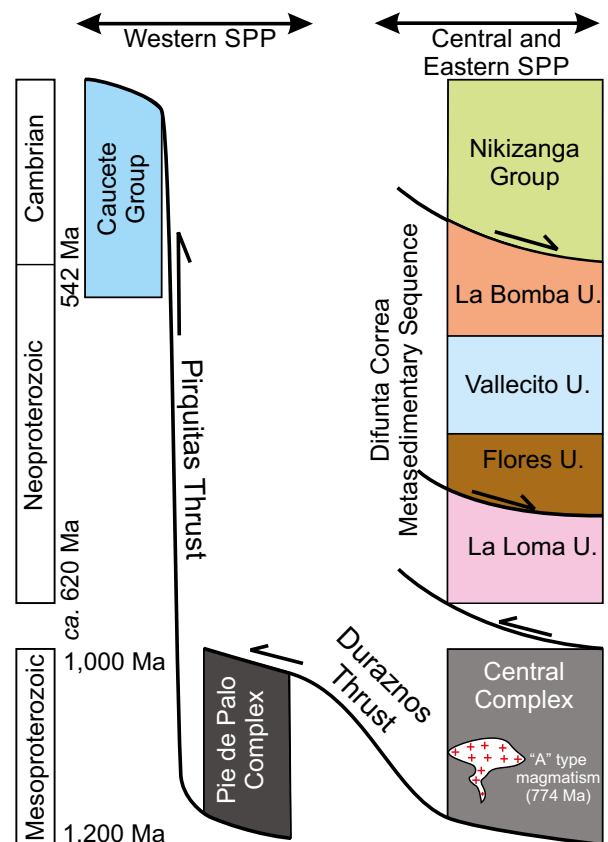


Figure 2. Main units of the Sierra de Pie de Palo (SPP) and structural relationships. Thicknesses are not to scale.

The Nikizanga Group crops out in the southeastern portion of the SPP and consists of low- to medium- grade metamorphic rocks (quartzites, marbles, graphitic schists and, to a lesser extent, amphibolites; Figure 1b). An early Cambrian depositional age was obtained for this group based on Sr-isotope composition of marbles (Filo del Grafito marbles; Galindo *et al.*, 2004), which was latter supported by U-Pb detrital zircon ages from a quartzite (Ramacciotti *et al.*, 2014).

## FIELD AND PETROGRAPHIC DESCRIPTIONS

The DCMS was affected by numerous shear zones, thrusts and brittle Andean faults that complicate the overall picture (Figure 3). Sedimentary structures and textures of the DCMS were largely obliterated and replaced by metamorphic textures as a consequence of strong deformation, recrystallization, and formation of new minerals during the Ordovician Famatinian metamorphism (Casquet *et al.*, 2001). Only the metaconglomerates preserve some primary structures (*e.g.*, graded-bedding and cross-bedding) and relict minerals in weakly deformed areas. Thus, the classification of protoliths was mainly based on geochemical data and not on mineral modes. The metasedimentary units are described below, from bottom to top (Figure 4):

La Loma unit consists almost exclusively of black quartz schists and widely outcrops in southeastern SPP (Figure 3). Schists are massive or banded and show a NE-SW striking, east-dipping, penetrative tectonic

foliation (Figure 5a). They were affected by numerous shear zones, and often show chevron folds (Figure 5b). The mineral assemblage mainly consists of quartz, with minor amounts of biotite, muscovite, plagioclase and zircon, with a fine-grained (< 1 mm) granoblastic texture (Figure 5c).

The Flores unit consists mainly of metaconglomerates, and less abundant Ca-pelitic schists. The former are matrix-supported with monocrystalline clasts of quartz and feldspar (2–15 mm), and locally, large (5–20 cm) phosphatic clasts which are described below. The metaconglomerates are lens-shaped and often grade in short distances into arenites (Figure 5d). The metamorphic mineral assemblage of the matrix is Qtz-Pl-Bt-Ms±Grt-Ep-Amph-Zrn-Ap-Op (abbreviations after Kretz, 1983; Figure 5e and 5f). Phosphatic clasts within the metaconglomerates are between 2 and 20 cm long and were flattened parallel to the main foliation. They consist of Ap-Qtz-Bt-Ilm (identified by electron microprobe) with microgranular texture (Figure 6a and 6b). Internally, the clasts show an alternation of phosphate-rich rock and thin, curved, quartz-rich veinlets (white arrows in Figure 6b). Sometimes an outer zone is found between the phosphatic clasts and the host conglomerate, containing abundant disseminated apatite, probably due to the exchange reaction during metamorphism (zone I in Figure 6a; see Discussion section). The mineral mode of phosphate rock was estimated using the software imageJ (Abramoff *et al.*, 2004). Sample SPP-22007D1 yielded: 45.5 % Ap, 41.6 % Qtz, 10.2 % Bt, 2.7 % Ilm (Figure 7). Ca-pelitic schists consist of Ms-Hbl-Grt-Bt-Ep-

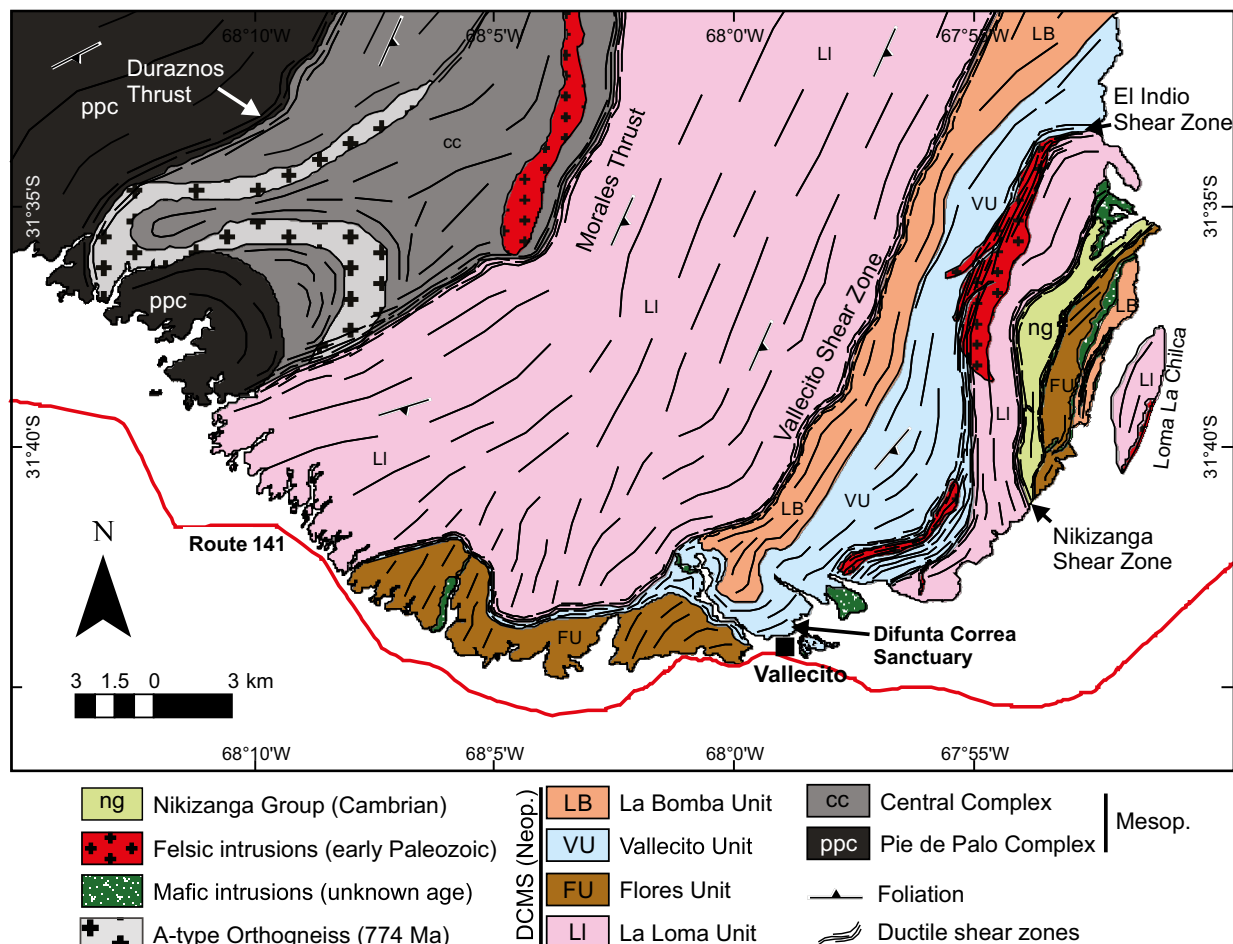


Figure 3. Geological map of the southern SPP showing the four units of the Difunta Correa metasedimentary sequence (DCMS).



Chl-Pl±Zrn-Op. They are characterized by the presence of Ca-minerals (amphibole and/or epidote) in association with typical metapelite minerals (*e.g.*, muscovite, biotite, garnet). Distinctive textural features consist of large, needle-shaped porphyroblasts of hornblende lying in the foliation plane with a radial arrangement (Figure 8a and 8b), and of garnet porphyroblasts discordant to the foliation (Figure 8b). Both textures suggest that minerals grew up late relative to foliation development. The textures and mineral composition of the DCMS Ca-pelitic schists are remarkably similar to *garben*-schists or *garbenschiefer* in the eastern Alps (Selverstone *et al.*, 1984).

The Vallecito unit, with a maximum apparent thickness of 3000 m, is mainly composed of calcite-rich marbles interbedded with calc-silicate rocks and quartzites. An abundant type of calc-silicate rock is para-amphibolite, *i.e.*, amphibolite formed after a sedimentary protolith (Rapela *et al.*, 2005). Marbles are often intruded by Ordovician granitoids (Baldo *et al.*, 2012) (Figure 8c) and exhibit a granoblastic texture with the minerals Cal±Dol-Ms-Bt-Chl-Pl-Qtz-Ttn-Zrn-Ap (Figure 8d).

La Bomba unit is composed of a succession of metapelites and metasandstones of variable composition (Figure 9a). The main rock types are: staurolite-garnet schists, garnet schists, hornblende-garnet metasandstones and calcite-rich metasandstones. The staurolite-garnet schists have a mineral assemblage of St-Grt-Bt-Ms-Pl-Qtz±Ap-Zrn and are characterized by the abundance of staurolite and garnet porphyroblasts set in a foliated mica-rich matrix (Figure 9b and c). The garnet schists consist of Ms-Grt-Bt-Qtz-Pl also with garnet as

porphyroblasts and a lepidoblastic to granoblastic matrix (Figure 9d). The hornblende-garnet metasandstones bear many similarities to Ca-pelitic schists (*garben*-schists) of the Flores unit, with the exception that they are more psammitic (Figure 10a-10c; compare with Figure 8a). The calcite-rich metasandstones are mainly composed of quartz and calcite, with minor amounts of muscovite, and show a granoblastic texture (Figure 10d).

## SAMPLING AND ANALYTICAL METHODS

Representative samples from the three meta-detrital units that form the DCMS were chosen for chemical and isotope analyses. Details of each sample are given in Table 1.

Chemical analyses were carried out on eleven samples (Table 2). Six samples were analyzed at Activation Laboratories Ltd. (Actlabs, Ontario, Canada) following the 4 Litho-research routine. Major elements were determined by ICP-AES, whereas minor and trace elements were determined by ICP-MS. The remaining five samples were analyzed by X-ray fluorescence (XRF) at the Laboratorio of the Instituto de Geología y Minería, Universidad Nacional de Jujuy, Argentina on a Rigaku FX2000 spectrometer with Rh tube (Table 2). Ground and homogenized samples were fused with lithium tetraborate for major element analyses. Trace element determinations were performed on rock powder pellets mixed with methyl methacrylate, and pressed at 20 t. Operating conditions were 50 kV and 45 mA.

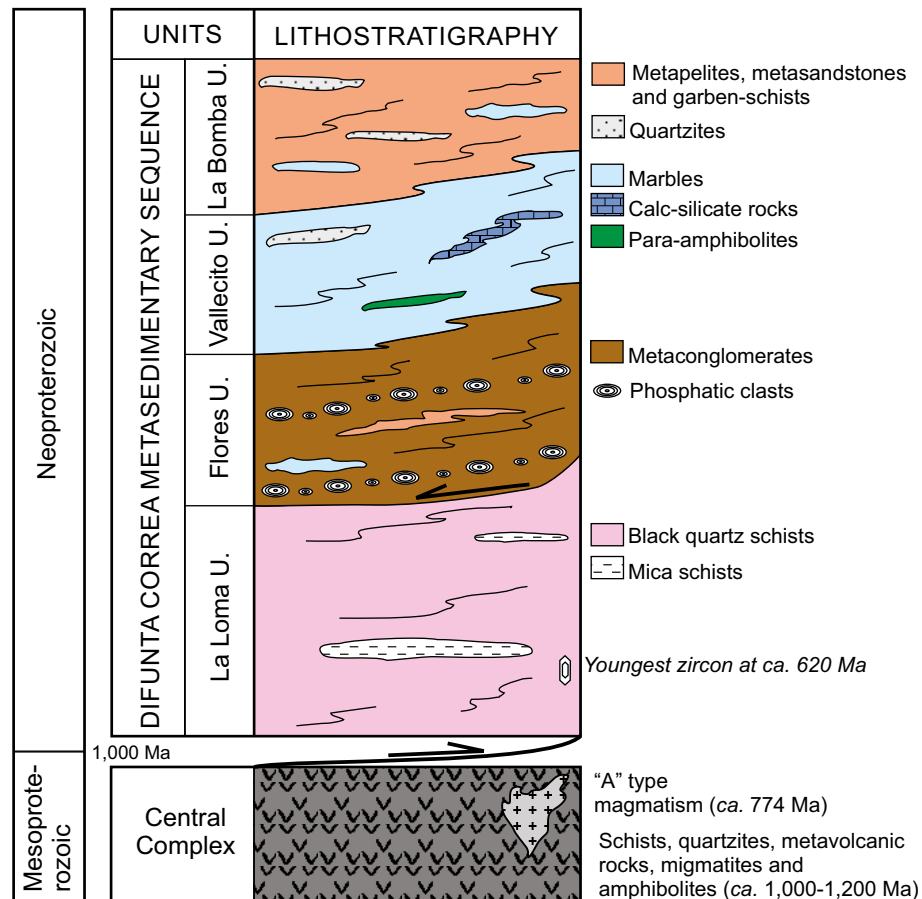


Figure 4. Schematic lithostratigraphic column of the Difunta Correa metasedimentary sequence. The youngest zircon age is from Rapela *et al.* (2005, 2015).



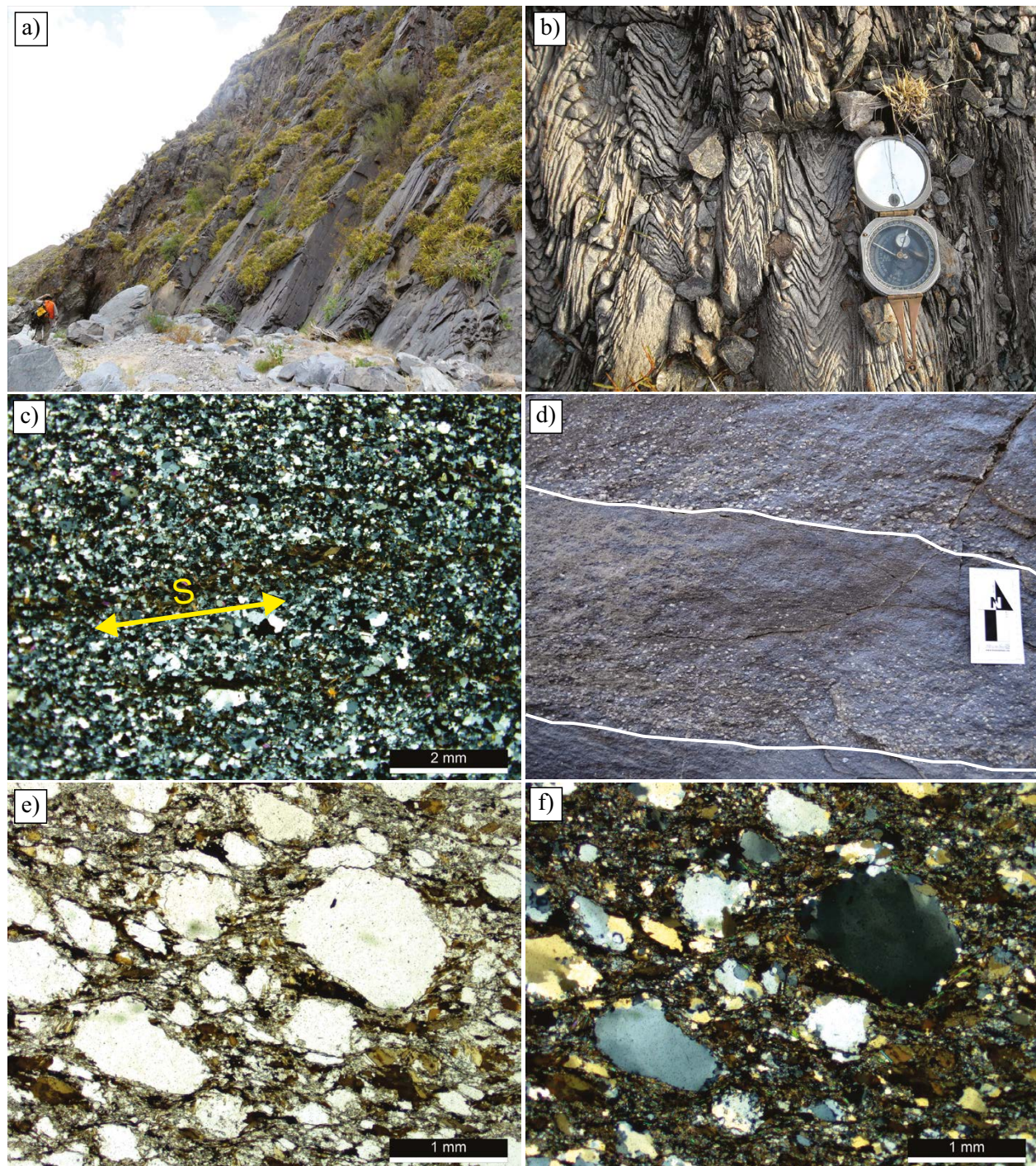


Figure 5. Field and petrographic characteristics of La Loma unit (a-c) and the Flores unit (d-f). a) Evenly foliated, black, quartz schists; b) Banded and folded (chevron folds) quartz schists; c) Fine-grained granoblastic texture mainly composed of quartz and, to a lesser extent, of oriented micas which define the foliation of the rock; d) Normal graded bedding in metaconglomerates; e) and f) PPL and crossed polars photomicrograph of metaconglomerates showing relict minerals (clasts) and recrystallized matrix.



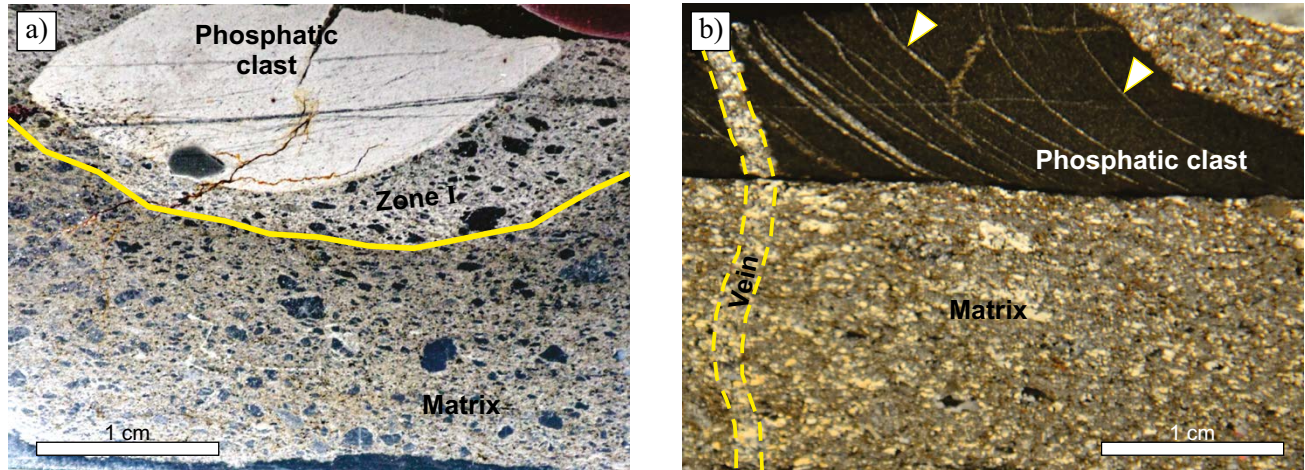


Figure 6. Phosphatic clasts within metaconglomerates of the Flores unit. a) Thin section of one phosphatic clast showing an outer zone of apatite-enriched host conglomerate (zone I); b) Crossed polars photomicrograph of a phosphatic clast with a truncated alternation of phosphate-rich rocks and thin veinlets of quartz (white arrow) and a cross-cutting quartz vein.

Rb-Sr and Sm-Nd isotope analyses were carried out on four detrital rocks. Additionally, Rb-Sr isotope data was collected from two phosphatic clasts within the metaconglomerates of the Flores unit (Table 3). Analytical work was carried out at the Centro de Geocronología y Geoquímica de Isótopos of the Universidad Complutense de Madrid on a PHOENIX<sup>®</sup> automated multicollector mass-spectrometer. Whole rocks (siliciclastic rocks) were first decomposed in 4 ml HF Merck-suprapure and 2 ml HNO<sub>3</sub> Merck-suprapure in Teflon digestion bombs for 48 hours at 120 °C, and finally in 6M HCl. Concentrations of Rb and Sr, as well as Rb/Sr atomic ratios, were determined by X-ray fluorescence spectrometry at the X-Ray Centro de Difracción de la Universidad Complutense following the methods of Pankhurst and O'Nions (1973). Sr and REE were separated using Bio-Rad AG50 x 12 cation exchange resin. The phosphatic clasts were decomposed in 6M HCl in Nalgene beakers and, after evaporation, dissolved with 3M HNO<sub>3</sub>; Sr was separated using an extraction chromatographic SrResinTM. Errors are quoted throughout as two standard deviations (2σ) from measured or calculated values. The decay constant used in the calculations is  $\lambda^{87}\text{Rb} = 1.42 \times 10^{-11} \text{ year}^{-1}$  recommended by the IUGS

Subcommission for Geochronology (Steiger and Jäger, 1977). Bulk Earth Sr-isotope composition was calculated from Allègre *et al.* (1983) and Taylor and McLennan (1985). Analytical uncertainties are estimated to be 0.01 % for  $^{87}\text{Sr}/^{86}\text{Sr}$  and 1 % for  $^{87}\text{Rb}/^{86}\text{Sr}$  ratios. Epsilon-Sr ( $\epsilon\text{Sr}$ ) values were calculated relative to a Bulk Earth present-day  $^{87}\text{Sr}/^{86}\text{Sr}$  value of 0.7045 and  $^{87}\text{Rb}/^{86}\text{Sr}$  of 0.0827 (DePaolo, 1988). Replicate analyses of the NBS-987 Sr-isotope standard yielded an average  $^{87}\text{Sr}/^{86}\text{Sr}$  ratio of  $0.710245 \pm 0.00003$  ( $n = 14$ ), (accepted value:  $0.71025 \pm 0.00005$ ; Faure, 2001)

Samarium and neodymium were determined by isotope dilution using spikes enriched in  $^{149}\text{Sm}$  and  $^{150}\text{Nd}$  and were separated from the REE group using Bio-beads coated with 10 % HDEHP. Errors are quoted throughout as two standard deviations (2σ) from measured or calculated values. The decay constants used in the calculations are the values  $\lambda^{147}\text{Sm} = 6.54 \times 10^{-12} \text{ year}^{-1}$  recommended by the IUGS Subcommission for Geochronology (Steiger and Jäger, 1977). Analytical uncertainties are estimated to be 0.006 % for  $^{143}\text{Nd}/^{144}\text{Nd}$  ratios and 0.1 % for  $^{147}\text{Sm}/^{144}\text{Nd}$  ratios. Epsilon-Nd ( $\epsilon\text{Nd}$ ) values were calculated relative to a present-day chondrite  $^{143}\text{Nd}/^{144}\text{Nd}$  value of 0.512638 and

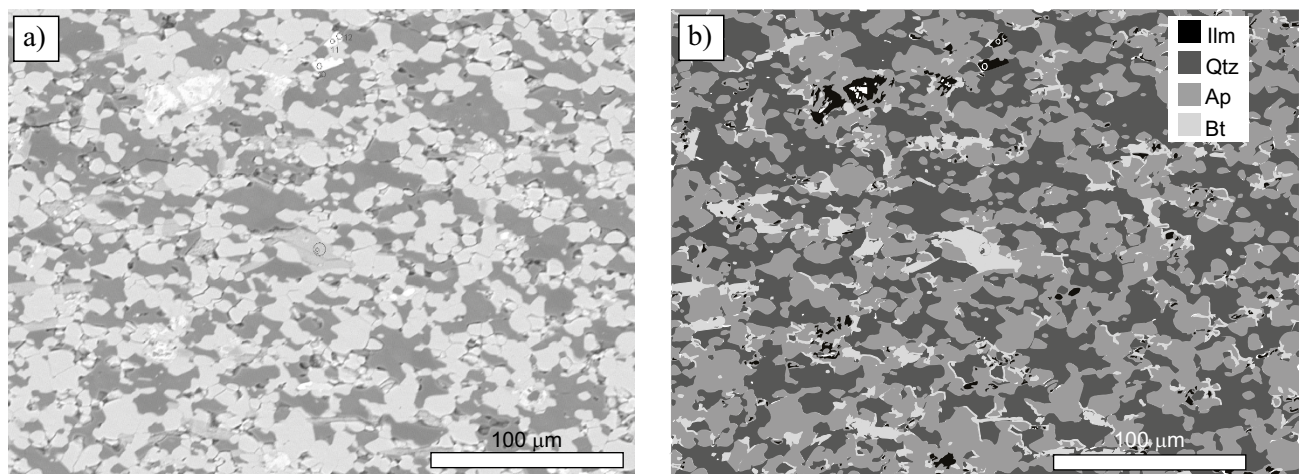


Figure 7. a) Secondary electron (SEM) image showing the slightly orientated microgranular texture of a phosphatic clast core (SPP-22007D1). The black areas are holes which were not considered for modal analysis, and are merged with quartz in b); b) Image a) processed with ImageJ software.



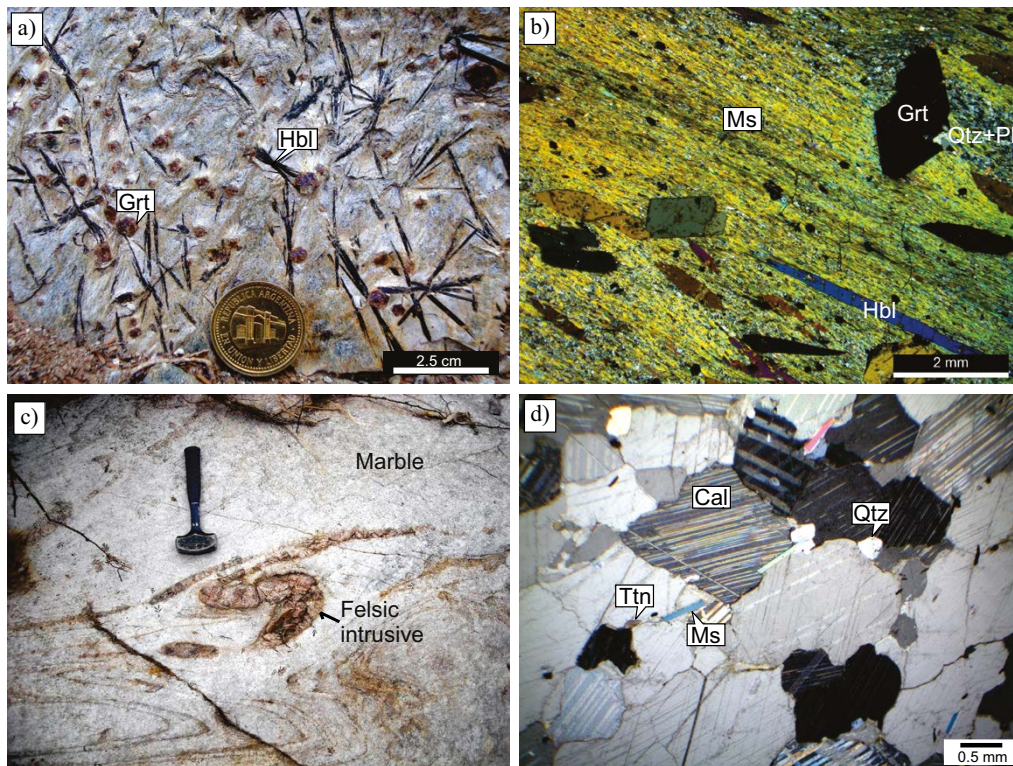


Figure 8. a) and b) Ca-pelitic schists (garben-schists) from the Flores unit. c) and d) Marbles from the Vallecito unit. c) Deformed marble with folded felsic intrusive, d) Crossed polars photomicrograph of marbles showing a granoblastic texture.

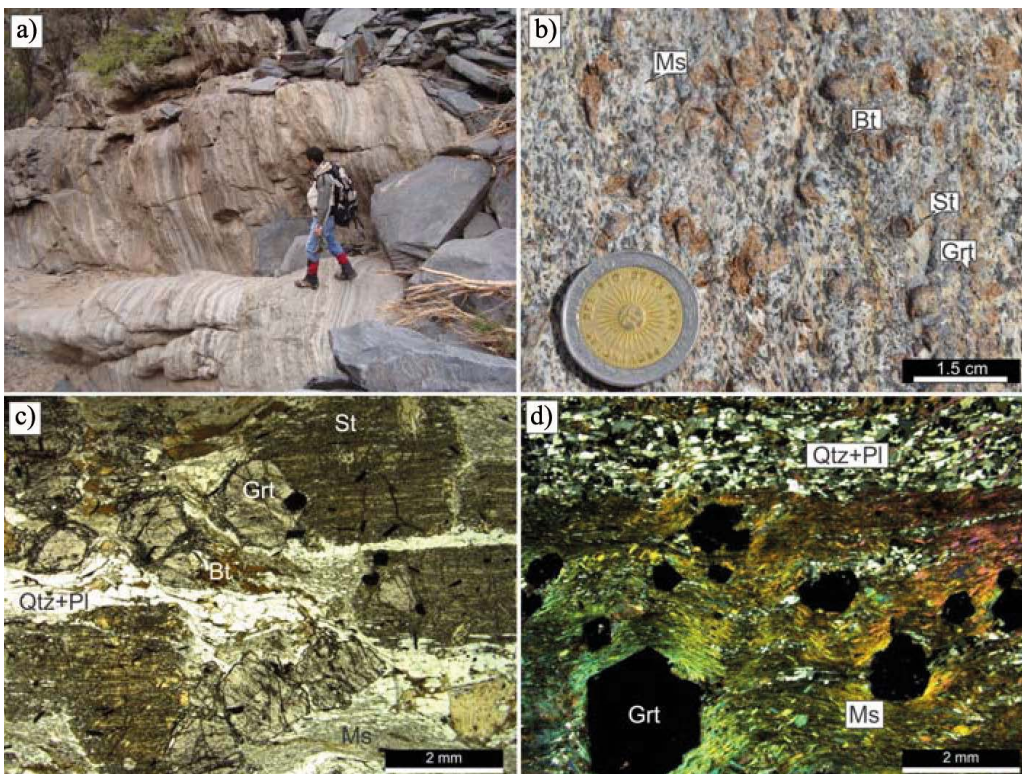


Figure 9. La Bomba unit. a) Succession of metapelites and metasandstones; b) Staurolite-garnet schist; c) PPL photomicrograph of a staurolite-garnet schist; d) Crossed polars photomicrograph of a garnet schist.



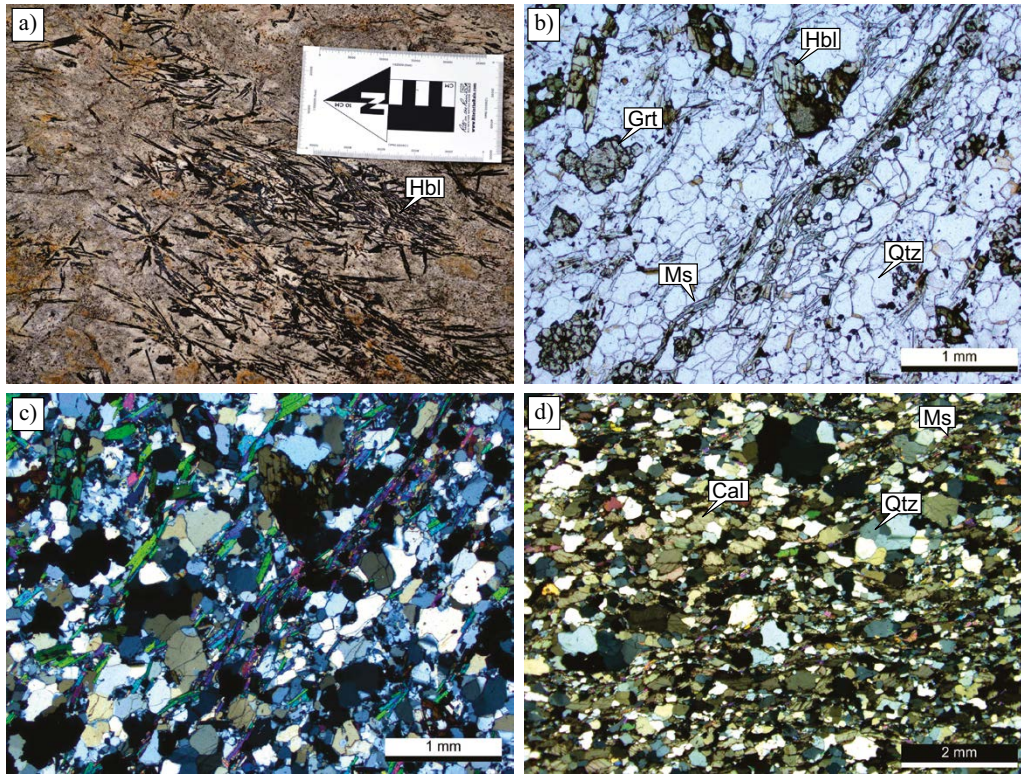


Figure 10. La Bomba unit. a) Hornblende-garnet metasandstone with non-oriented hornblende prisms, locally displaying a radial arrangement; b) and c) PPL and crossed polars photomicrograph of a hornblende-garnet metasandstone; d) Crossed polars photomicrograph of a calcite-rich metasandstone.

$^{147}\text{Sm}/^{144}\text{Nd}$  of 0.1967 (Jacobsen and Wasserburg, 1980; Goldstein *et al.*, 1984). Eight analyses of La Jolla Nd-standard measured during sample analysis gave a mean  $^{143}\text{Nd}/^{144}\text{Nd}$  ratio of  $0.511845 \pm 0.00001$  (accepted value:  $0.511858 \pm 0.00007$ ; Lugmair and Carlson, 1978). Single stage model ages ( $T_{\text{DM}}$ ) was calculated according to DePaolo (1988).

The composition of phosphate minerals was determined in one phosphatic clast (SPP-22007D1; Table 4) using a JEOL JXA8230 Superprobe at the Universidad Nacional de Córdoba, Argentina. Carbon-coated polished sections were analyzed at 15 kV using 20 nA beam and a range of natural and synthetic standards.

## RESULTS

### Detrital rocks geochemistry

Geochemical data from the DCMS siliciclastic rocks are shown in Table 2. According to Herron's (1988) plot (Figure 11) the samples are classified as pelites (shales and Fe-Shale) and sandstones (wackes, sub-litharenites, litharenites and Fe-sandstones). The Ca-pelitic schist from the Flores unit shows the lowest concentration of  $\text{SiO}_2$  (40.3–46.4 %) and the highest contents of  $\text{Al}_2\text{O}_3$  (26.2–28.9 %) and  $\text{K}_2\text{O}$  (6.6–7.2 %). Metapelites from La Bomba unit (staurolite-garnet

Table 1. Location and description of the analyzed samples. Mineral abbreviations after Kretz (1983).

Sample	Lat. (S)	Long. (W)	Unit	Rock type	Mineral assemblage
SPP-22013	31° 42' 24''	68° 01' 00''	La Bomba	Calcite-rich metasandstone	Qtz-Pl-Kfs-Cal-Ms
SPP-27000	31° 34' 46''	67° 52' 51''	La Bomba	Hornblende-garnet metasandstone	Grt-Hbl-Ms-Bt-Qtz-Pl
SPP-27001	31° 34' 46''	67° 52' 51''	La Bomba	Mica schist	Grt-Ms-Bt-Qtz-Pl
SPP-27002	31° 34' 46''	67° 52' 51''	La Bomba	Staurolite-garnet schist	St-Grt-Ms-Bt-Qtz-Pl
SPP-27003	31° 34' 46''	67° 52' 51''	La Bomba	Quartz schist	Qtz-Ms-Bt-Grt
SPP-424	31° 22' 55''	67° 58' 42''	La Bomba	Garnet gneiss	Grt-Bt-Ms-Qtz-Pl
SPP-22007D1	31° 43' 36''	68° 06' 00''	Flores	Phosphatic clast	Ap-Qtz-Bt-Ilm
SPP-22007C1	31° 43' 36''	68° 06' 00''	Flores	Phosphatic clast	Ap-Qtz-Bt-Ilm
SPP-22037	31° 39' 34''	67° 53' 04''	Flores	Metaconglomerate	Qtz-Pl-Kfs-Bt-Ms-Chl-Ep
SPP-27012	31° 35' 48''	67° 51' 36''	Flores	Metaconglomerate	Qtz-Pl-Kfs-Bt-Ms-Chl-Ep
SPP-6086	31° 43' 00''	68° 05' 48''	Flores	Metaconglomerate	Qtz-Pl-Kfs-Bt-Ms-Chl-Amph-Ep
PPL-15	31° 42' 22''	68° 05' 43''	Flores	Ca-pelitic schist	Grt-Hbl-Ms-Chl-Qtz-Pl-Ep
PPL-23	31° 42' 55''	68° 05' 42''	Flores	Ca-pelitic schist	Grt-Hbl-Ms-Chl-Qtz-Pl-Ep
SPP-22019	31° 40' 32''	67° 51' 55''	La Loma	Quartz schist	Qtz-Pl-Bt-Ms

Table 2. Chemical analyses. \*Samples analyzed at Actlabs. The remaining samples were analyzed at the Laboratory of the Instituto de Geología y Minería, Universidad Nacional de Jujuy, Argentina. Total Fe content is expressed as  $\text{Fe}_2\text{O}_3$ .

Difunta Correa Metasedimentary Sequence											
Unit	La Loma	Flores				La Bomba					
Rock type	Quartz schist	Metaconglomerates		Ca-pelitic schists		Metasandstones			Metapelites		
Sample	SPP-22019*	SPP-22037*	SPP-27012	PPL-23*	PPL-15*	SPP-22013*	SPP-27000	SPP-27003	SPP-27001	SPP-27002	SPP-424*
(weight %)											
SiO <sub>2</sub>	74.23	75.61	63.84	46.40	40.35	66.04	75.34	69.01	52.41	59.95	56.92
TiO <sub>2</sub>	0.64	0.68	1.07	1.45	1.60	0.27	1.09	0.85	1.05	1.00	1.59
Al <sub>2</sub> O <sub>3</sub>	11.69	9.72	14.62	26.22	28.88	4.23	11.20	13.80	23.26	18.79	17.84
Fe <sub>2</sub> O <sub>3</sub>	3.71	3.73	7.50	9.32	11.35	1.49	5.59	5.92	9.21	11.65	11.03
MnO	0.09	0.08	0.10	0.17	0.15	0.10	0.10	0.09	0.10	0.26	0.13
MgO	1.37	1.01	2.27	2.72	3.33	0.25	1.31	1.64	2.37	1.93	0.65
CaO	2.21	3.21	1.65	1.67	0.88	13.47	1.97	1.34	1.36	1.13	3.62
Na <sub>2</sub> O	3.32	2.55	2.87	2.21	0.95	0.67	2.72	2.35	1.27	0.65	6.64
K <sub>2</sub> O	1.84	0.94	3.39	6.64	7.23	0.94	0.72	3.09	5.84	3.07	1.36
P <sub>2</sub> O <sub>5</sub>	0.15	0.15	0.17	0.15	0.26	0.03	0.09	0.10	0.40	0.11	0.25
LOI	0.77	0.70	1.40	3.03	4.81	11.60	0.37	0.87	2.47	1.03	0.02
Total	100.02	98.38	98.87	99.98	99.80	99.09	100.49	99.04	99.75	99.55	100.05
(ppm)											
Cs	1.60	1.40	-	5.00	3.00	0.60	-	-	-	-	0.20
Rb	71.00	29.00	-	219.00	171.00	34.00	75.00	-	165.00	105.00	8.70
Ba	467.00	385.00	-	1,100.00	1,200.00	148.00	602.00	-	1,199.00	688.00	690.00
Th	10.10	6.22	-	10.40	13.40	6.20	12.00	-	12.00	13.00	0.90
U	2.46	3.24	-	2.30	3.90	2.49	4.00	-	4.00	3.00	0.70
Nb	11.90	6.40	-	-	-	4.10	18.00	-	21.00	21.00	40.00
Ta	1.11	0.70	-	<0.5	<0.5	0.56	-	-	-	-	1.60
K	15,276	7,804	-	55,127	60,026	7,804	5,953	-	48,452	25,447	11,291
La	30.50	28.30	-	39.20	65.60	12.10	-	-	-	-	56.10
Ce	67.70	62.50	-	79.00	127.00	27.20	-	-	-	-	121.00
Pb	8.00	9.00	-	26.00	27.00	-	-	-	-	-	-
Pr	8.42	7.28	-	-	-	3.12	-	-	-	-	-
Sr	231.00	256.00	-	143.00	91.00	179.00	202.00	-	194.00	124.00	340.00
Nd	35.20	28.40	-	35.00	60.00	11.70	-	-	-	-	67.00
P	654.85	654.85	-	654.85	1,135.07	130.97	371.08	-	1,759.37	458.40	1,091.42
Zr	598.00	176.00	-	176.00	126.00	283.00	254.00	-	190.00	138.00	790.00
Hf	13.80	4.30	-	7.00	10.00	6.40	6.00	-	5.00	3.00	16.00
Sm	7.78	5.69	-	7.80	14.20	2.36	-	-	-	-	13.50
Eu	1.57	1.27	-	1.50	2.20	0.45	-	-	-	-	4.78
Ti	3,844	4,054	-	8,695	9,618	1,595	6,548	-	6,290	5,967	9,534
Gd	6.97	4.86	-	-	-	2.16	-	-	-	-	-
Tb	1.15	0.78	-	0.80	1.50	0.37	-	-	-	-	1.70
Dy	7.03	4.54	-	-	-	2.19	-	-	-	-	-
Y	39.00	25.00	-	40.00	60.00	13.00	50.00	-	52.00	36.00	40.00
Ho	1.44	0.93	-	-	-	0.48	-	-	-	-	-
Er	4.28	2.75	-	-	-	1.49	-	-	-	-	-
Tm	0.66	0.43	-	-	-	0.24	-	-	-	-	-
Yb	4.38	2.80	-	6.50	8.20	1.64	-	-	-	-	2.70
Lu	0.70	0.45	-	0.95	1.20	0.27	-	-	-	-	0.38
Sc	7.54	7.49	-	26.50	31.00	2.85	-	-	-	-	25.00
Co	10.80	13.20	-	30.00	35.00	2.60	57.00	-	38.00	67.00	8.00
Cr	29.70	40.40	-	162.00	132.00	12.40	152.00	-	121.00	158.00	7.00
Cu	3.00	12.00	-	3.00	5.00	3.00	-	-	-	-	-
Ni	17.00	19.00	-	66.00	80.00	3.00	95.00	-	51.00	25.00	50.00
S	0.01	0.02	-	<0.01	<0.01	0.09	0.02	-	0.02	0.08	-
Sb	0.30	0.20	-	0.70	<0.10	<0.10	-	-	-	-	-
Sn	4.00	2.00	-	-	-	2.00	-	-	-	-	-
Tl	0.48	0.19	-	-	-	0.15	-	-	-	-	-
V	56.00	78.00	-	153.00	183.00	22.00	-	-	-	-	-
W	22.00	43.00	-	<1	<1	3.00	-	-	-	-	<1
Zn	65.00	47.00	-	129.00	75.00	19.00	-	-	-	-	-
Rb/Sr	0.31	0.11	-	1.53	1.88	0.19	0.37	-	0.85	0.85	0.03



Table 3. Isotope composition of samples from the Difunta Correa metasedimentary sequence.

Sample Rock type Unit	SPP-22013 Calcite-rich metasandstone La Bomba	SPP-22019 Quartz schist La Loma	SPP-22037 Meta-conglomerate Flores	SPP-6086 Meta-conglomerate Flores	SPP-22007C Phosphatic clast Flores	SPP-22007D Phosphatic clast Flores	Bulk Earth
Rb (ppm)	34	71	29	101	10.7	22.8	-
Sr (ppm)	179	231	256	160	158	408.2	-
Rb/Sr	0.18994	0.30736	0.11328	0.63125	0.06772	0.05585	-
<sup>87</sup> Rb/ <sup>86</sup> Sr	0.55042	0.89094	0.32796	1.83070	0.20170	0.16334	0.08920
<sup>87</sup> Sr/ <sup>86</sup> Sr	0.72377	0.72692	0.71431	0.73221	0.72163	0.72290	0.70470
<sup>87</sup> Sr/ <sup>86</sup> Sr <sub>465</sub>	0.72013	0.72102	0.71213	0.72008	0.72030	0.72182	0.70411
<sup>87</sup> Sr/ <sup>86</sup> Sr <sub>600</sub>	0.71906	0.71930	0.71150	0.71655	0.71991	0.72150	0.70394
εSr <sub>600</sub>	215	218	107	179	227	250	0
Sm (ppm)	2.36	7.78	5.69	6.65	-	-	-
Nd (ppm)	11.7	35.2	28.4	30.5	-	-	-
Sm/Nd	0.20171	0.22102	0.20035	0.21803	-	-	-
<sup>147</sup> Sm/ <sup>144</sup> Nd	0.12193	0.13361	0.12111	0.13180	-	-	-
<sup>143</sup> Nd/ <sup>144</sup> Nd	0.51203	0.51223	0.51223	0.51225	-	-	-
<sup>143</sup> Nd/ <sup>144</sup> Nd <sub>465</sub>	0.51166	0.51182	0.51186	0.51184	-	-	-
<sup>143</sup> Nd/ <sup>144</sup> Nd <sub>600</sub>	0.51156	0.51170	0.51176	0.51173	-	-	-
εNd <sub>600</sub>	-6.04	-3.20	-2.14	-2.69	-	-	-
TDM	1.64	1.53	1.33	1.47	-	-	-

schists) show similar values compared to the Ca-pelitic schist, whereas quartz schists (La Loma U.), metaconglomerates (Flores U.) and metasandstones (La Bomba U.) exhibit higher concentrations of SiO<sub>2</sub> (63.8–75.6 %) and the lowest contents of Al<sub>2</sub>O<sub>3</sub> and K<sub>2</sub>O (4.2–14.6 % and 0.7–3.4 % respectively; Table 2).

Geochemical variations among samples (Figure 12) reflect changes in their mineralogical composition. Modal variations of phyllosilicates (pelitic component) relative to quartz-feldspar (psammitic component) are supported by: 1) negative correlations of SiO<sub>2</sub> against TiO<sub>2</sub> ( $R^2 = 0.66$ ;  $p < 0.01$ ), Al<sub>2</sub>O<sub>3</sub> ( $R^2 = 0.98$ ;  $p < 0.001$ ), Fe<sub>2</sub>O<sub>3</sub> ( $R^2 = 0.71$ ;  $p < 0.01$ ), MgO ( $R^2 = 0.58$ ;  $p < 0.05$ ), and K<sub>2</sub>O ( $R^2 = 0.77$ ;  $p < 0.001$ ; Figure 12a–12e) and 2) positive correlations of TiO<sub>2</sub> against Fe<sub>2</sub>O<sub>3</sub> ( $R^2 = 0.62$ ;  $p < 0.01$ ), Al<sub>2</sub>O<sub>3</sub> ( $R^2 = 0.56$ ;  $p < 0.05$ ), and K<sub>2</sub>O ( $R^2 = 0.65$ ;  $p < 0.01$ ), and of Al<sub>2</sub>O<sub>3</sub> with K<sub>2</sub>O ( $R^2 = 0.85$ ;  $p < 0.001$ ; Figure 13). No clear patterns between CaO, Na<sub>2</sub>O, MnO and P<sub>2</sub>O<sub>5</sub> are recognized probably because they are mobile elements (Figure 12f–12i).

Relationships between major and trace elements are shown in Figure 14. The effect of the pelitic component is reflected by a positive correlation between K<sub>2</sub>O against Ba ( $R^2 = 0.90$ ;  $p < 0.001$ ) and Rb ( $R^2 = 0.92$ ;  $p < 0.001$ ) (Figure 14a and 14b). The highest concentrations of Zr and SiO<sub>2</sub> are found in metasandstones (La Bomba U.), quartz schist (La Loma U.) and metaconglomerates (Flores U.).

The REE concentration of metasedimentary rocks is strongly influenced by the grain size and mineral composition. In this sense, Cullers *et al.* (1987) found that the REE concentration is about 20 %

higher in shales than in sandstones. Heavy minerals, such as zircon and garnet, contribute significantly to the increase in the heavy rare earth element (HREE) content (Morton, 1991). The chondrite-normalized (Sun and McDonough, 1989) REE diagram of the DCMS samples is shown in Figure 15. The model average composition of PCS (Proterozoic Cratonic Sandstones; Condie, 1993) and PAAS (Post-Archean Australian Shales; Taylor and McLennan, 1985) are included for comparison. The REE content in metapelites from La Bomba unit and Ca-pelitic schist from the Flores unit ranges between 170.7 and 279.9 ppm, slightly enriched with respect to PAAS. Metasandstones (La Bomba U.), metaconglomerates (Flores U.) and quartz schist (La Loma U.) show higher REE values (65.7–177.7 ppm) than PCS. All samples are characterized by a light rare earth elements (LREE) enrichment ( $La_N/Eu_N = 2.87$ – $7.30$ ), and a flat array (except the sample SPP-424) to the HREE ( $Tb_N/Lu_N = 0.57$ – $1.17$ ). In general, all samples of the DCMS exhibit a negative Eu anomaly ( $Eu_N/Eu_N^* = 0.61$ – $0.74$ ), which is inherited from the sediment source.

#### Sr-Nd isotope composition of detrital rocks

The Sr-isotope composition of four siliciclastic samples is shown in Table 3. Siliciclastic rocks yielded <sup>87</sup>Sr/<sup>86</sup>Sr ratios recalculated to the time of sedimentation (*ca.* 600 Ma) between 0.71150 and 0.71930. The corresponding εSr<sub>600</sub> values range from 107 to 218, *i.e.*, well above Bulk Earth.

The <sup>143</sup>Nd/<sup>144</sup>Nd<sub>600</sub> values of the four detrital rocks range from 0.51156 to 0.51176. The εNd<sub>600</sub> values range from -6.04 to -2.14 and Nd depleted mantle model ages (T<sub>DM</sub>) are between 1.33 and 1.64 Ga.

#### Phosphatic clasts

Four grains of phosphate minerals were analyzed in one phosphatic clast from metaconglomerates of the Flores unit (Table 4). The composition (weight %) is very homogenous with 1.97 to 2.47 wt % F and minor contents of FeO, MnO and Cl. They are classified as apatite-(CaF), *i.e.*, fluor-apatite (Pasero *et al.*, 2010). The <sup>87</sup>Sr/<sup>86</sup>Sr ratio measured in two phosphatic clasts yielded values of 0.72163 and 0.72290 (Table 3).

Table 4. Mineral chemistry of phosphate minerals as weight% (sample SPP-22007D1). Total Fe content as FeO.

N° analysis	CaO	P <sub>2</sub> O <sub>5</sub>	F	FeO	MnO	Cl	Total
3	54.53	42.31	2.21	0.06	0.07	0.02	99.12
4	54.92	41.99	1.97	0.05	0.05	0.01	98.99
5	54.36	41.22	2.13	0.14	0.04	0.01	97.90
6	55.33	41.24	2.47	0.11	0.04	0.01	99.20
mean	54.79	41.69	2.20	0.09	0.05	0.01	98.82

## DISCUSSION

## Protoliths and sedimentary setting of the DCMS

Metamorphism is considered essentially an isochemical process, with restricted mobility of non-volatile elements (Fettes and Desmons, 2007; Vernon and Clarke, 2011). Hence, ratios between some major and trace elements and the REE contents in meta-detrital rocks should reflect the mineralogical composition of protoliths, sediment maturity, and possible source areas. High Field Strength (HFS) elements (*e.g.*, La, Th, Zr, Hf, Sc and Ti) can be used to infer the source area composition and the tectonic setting because they are immobile elements, and ratios between them do not significantly change during diagenesis and metamorphism (Bhatia and Crook, 1986). Elements like Th and La are useful for identifying igneous felsic sources, whereas Zr and La concentrations reflect the degree of either crustal recycling or input from siliceous sources. Sc and Ti are related to mafic sources (Bhatia and Crook, 1986).

Source areas composition and tectonic setting were estimated here using the plots proposed by Roser and Korsch (1988; Figure 16a and 16b), Floyd and Leveridge (1987; Figure 16c) and Bhatia and Crook (1986; Figure 16d-16f). The DCMS samples suggest predominantly felsic-to-intermediate source areas with scarce influence of mafic rocks. Furthermore, most data point to continental and passive margin sedimentation (Figure 16d-16f). As it was pointed out above, these tectonic setting discrimination diagrams have to be used with caution because they often show erroneous results (Armstrong-Altrin and Verma 2005; Verdecchia and Baldo, 2010). Thus, we use here these diagrams to better constrain the tectonic setting of the basin but regional geological evidence is also considered. Marbles with sea-water Sr-isotope composition (*i.e.*, Vallecito unit; Galindo *et al.*, 2004), phosphatic clasts in the Flores unit and the lack of contemporaneous arc related igneous rocks further support passive margin sedimentation. A similar interpretation was proposed by Rapela *et al.* (2015) based on detrital zircon geochronology of the DCMS.

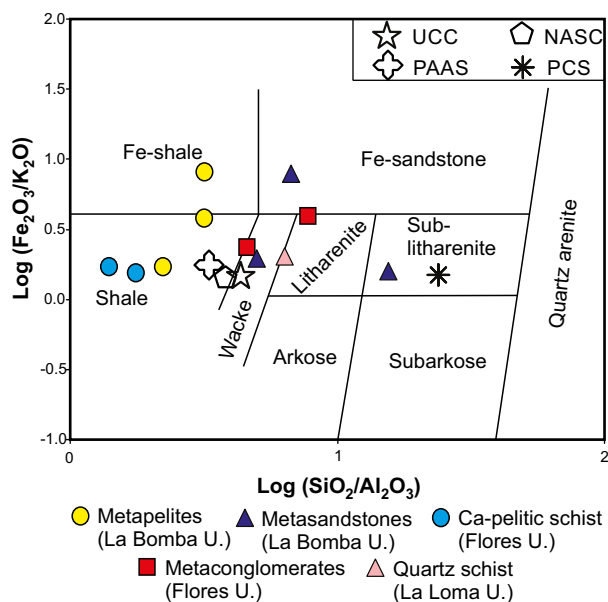


Figure 11. Herron (1988) classification diagram of sedimentary rocks. UCC: Upper Continental Crust (McLennan, 2001); PAAS: Post-Archean Australian Shales (Taylor and McLennan, 1985); NASC: North American Shale Composite (Gromet *et al.*, 1984); PCS: Proterozoic Cratonic Sandstones (Condie, 1993).

## Significance of Sr- and Nd-isotope composition of detrital rocks

As siliciclastic rocks are composed of detrital minerals and clasts, the whole-rock isotope signature may be a mixture of components coming from different sources with different isotope compositions and different ages. The variations in the  $^{87}\text{Sr}/^{86}\text{Sr}_{600}$  ratios (0.71150–0.71930) and the  $\epsilon\text{Sr}_{600}$  values (107–218) of the four siliciclastic samples (Table 3) can thus be attributed to a mixture of components from different sources. The high values of  $\epsilon\text{Sr}_{600}$  suggest that the siliciclastic rocks resulted from the erosion of an evolved continental crust.

Nd-isotope composition as shown by the  $\epsilon\text{Nd}_{600}$  values between -2.14 and -6.04 (Table 3) also suggests contribution mainly from an evolved continental source. Nd model ages ( $T_{\text{DM}}$ ) between 1.3 and 1.6 Ga further imply that the continental source was chiefly Mesoproterozoic (and probably older). Metasedimentary rocks of the Precambrian basement in the WSP (Sierra de Pie de Palo, Maz, Espinal and Umango; Figure 1a), yield Nd  $T_{\text{DM}}$  ages between 1.2 and 2.7 Ga (Figure 17; Varela *et al.*, 2003; Porcher *et al.*, 2004; Casquet *et al.*, 2008). A first group of ages (between 1.7 and 2.7 Ga) are restricted to the Maz suspect terrane in the sierras de Maz and Espinal. The latter consists of a strip of metasedimentary rocks that hosts a middle Mesoproterozoic juvenile magmatic arc and late Mesoproterozoic massif-type anorthosites with U-Pb ages between *ca.* 1.3 and 1.07 Ga (Casquet *et al.*, 2005a, b; Rapela *et al.*, 2010). On the other hand, the second group of model ages (between 1.2 and 1.6 Ga) involve a significant Nd contribution from Mesoproterozoic igneous sources and from reworking of older sedimentary and/or igneous rocks. This group corresponds not only to the DCMS of the SPP but also to metasedimentary successions with extensive marble beds in the sierras de Maz and Espinal (Casquet *et al.*, 2008) and in the westernmost Sierra de Umango, probably equivalent to the DCMS, where a Mesoproterozoic basement has also been recognized (Varela *et al.*, 2011). In all cases, as in the SPP, the original relationships between the second group of metasedimentary rocks and the Mesoproterozoic basement are overprinted by ductile shearing and/or faulting. However, the argument that the metasedimentary successions of the second group of Nd model ages were laid down on the Mesoproterozoic basement that outcrops in the WSP seems well-founded as they are commonly juxtaposed. Isotope geochemistry and detrital zircon evidence (*i.e.* the abundance of Mesoproterozoic ages) in this second group of metasedimentary rocks (Casquet *et al.*, 2008; Rapela *et al.*, 2015) further strengthen this interpretation.

We thus infer that the main source of the DCMS was the Mesoproterozoic basement that crops out in the sierras de Maz, Espinal, Umango and in the Central Complex of SPP. This basement underlaid the DCMS at the time of sedimentation. Moreover, rocks with Nd isotope composition similar to the DCMS are also exposed in southeastern Laurentia (present position) in the Grenville and Granite-Rhyolite provinces (Figures 17 and 18). These foreign sources have also been invoked - particularly the second- to explain some peculiarities of the detrital zircon patterns such as a population of grains between 1.3 and 1.5 Ga with no likely sources in the WSP Mesoproterozoic basement (Rapela *et al.*, 2015).

## Phosphatic clasts in the Flores unit

The  $^{87}\text{Sr}/^{86}\text{Sr}$  ratios in phosphorites (if not modified by later processes) should reflect the isotope composition of sea-water at the time of deposition (McArthur *et al.*, 1990; Compton *et al.*, 2002). Previous contributions (Galindo *et al.*, 2004) found  $^{87}\text{Sr}/^{86}\text{Sr}$  ratios between 0.70732 and 0.70742 in marbles of the DCMS which point to an early Ediacaran depositional age according to the Sr-isotope composition of sea-water in the middle to late Neoproterozoic (Halverson and Shields, 2011). The difference between the  $^{87}\text{Sr}/^{86}\text{Sr}$  ratios of the DCMS marbles and those of phosphatic clasts (0.71991 and 0.72150 at 600 Ma) sug-



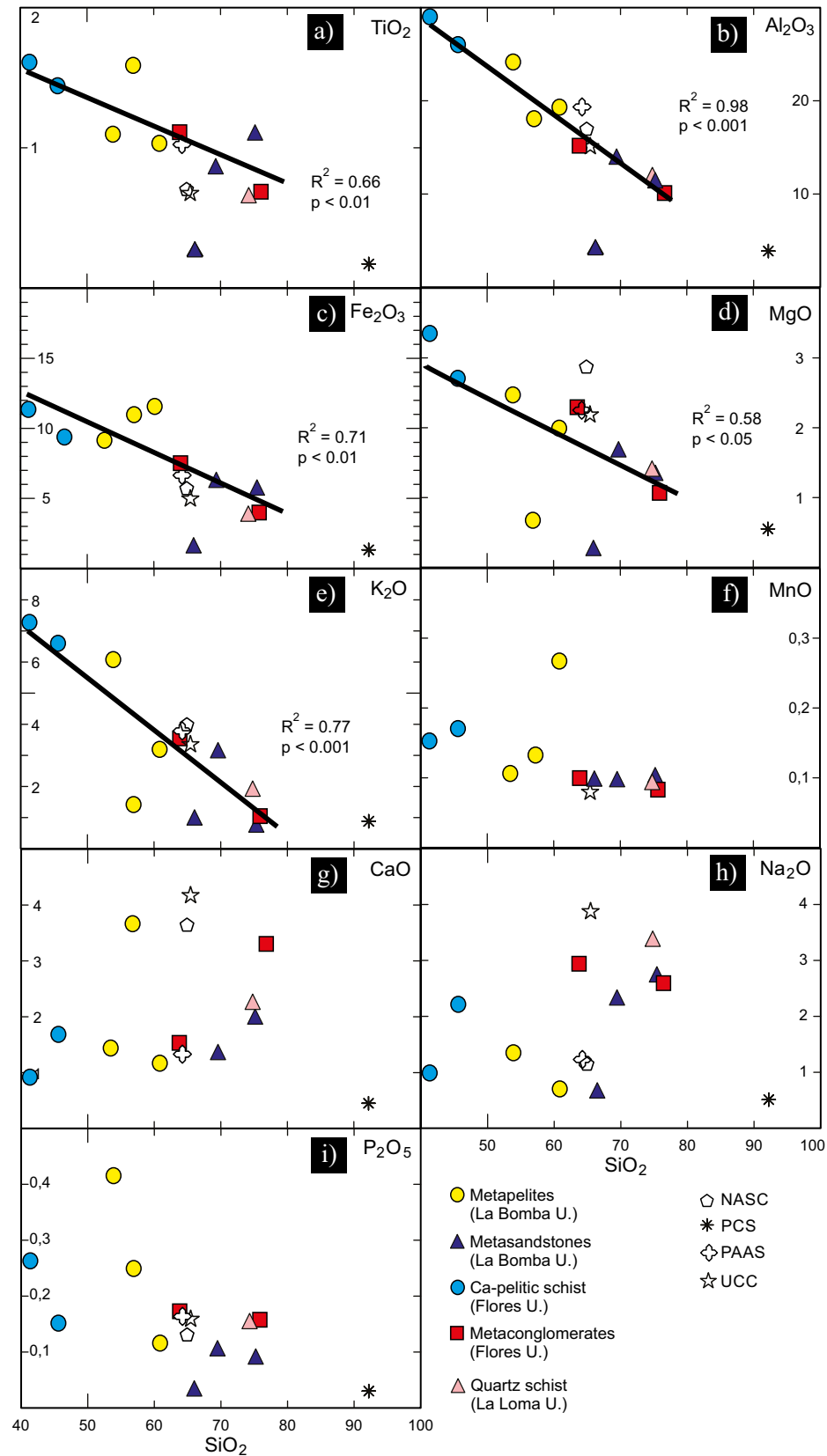


Figure 12. Chemical variation plots of metasedimentary rocks from the Difunta Correa metasedimentary sequence. Samples SPP-22013 and SPP-424 are not shown in the  $\text{CaO}$  and  $\text{Na}_2\text{O}$  plots (respectively) because contents of these components are off-scale. Values in weight percent. Sample SPP-22013 was not considered in correlation coefficients due to their carbonaceous cement.

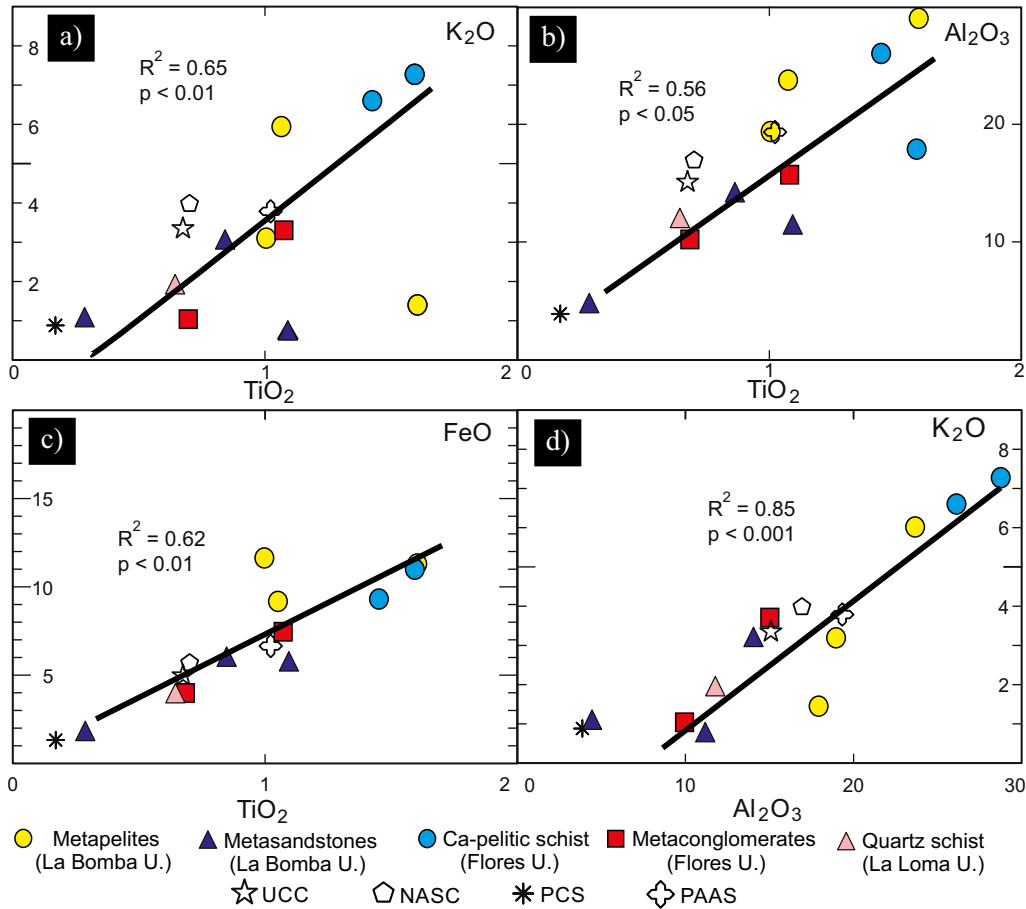


Figure 13. Variation plots of major elements against  $\text{TiO}_2$  and  $\text{Al}_2\text{O}_3$ . Values in weight percent. a) Sample PPL-424 was not considered in correlation coefficients. b) - d) Sample SPP-22013 was not considered in correlation coefficients due to their carbonaceous cement.

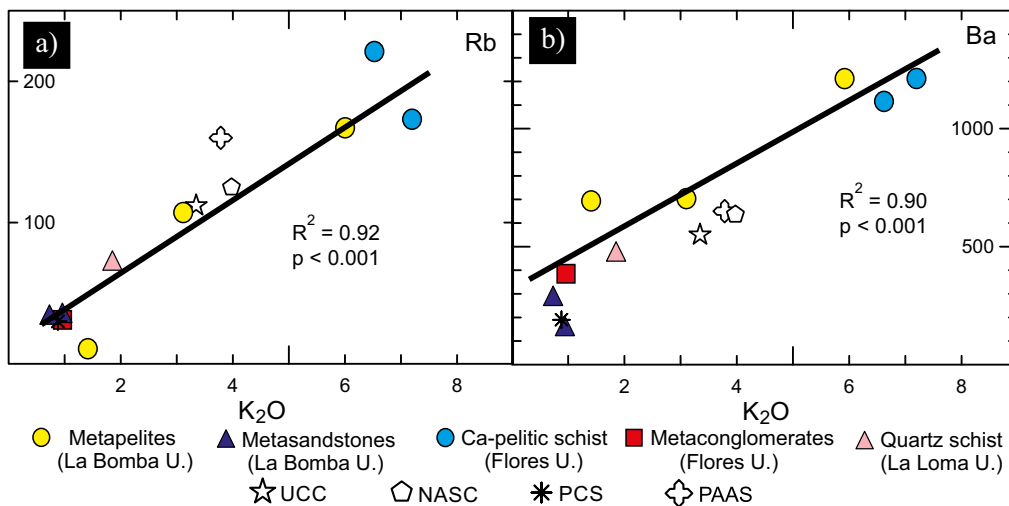


Figure 14. Variation plots of  $\text{Rb}$  and  $\text{Ba}$  of metasedimentary rocks from the Difunta Correa metasedimentary sequence. Values in weight percent for major elements and ppm for trace elements. Sample SPP-22013 was not considered in correlation coefficients due to their carbonaceous cement.



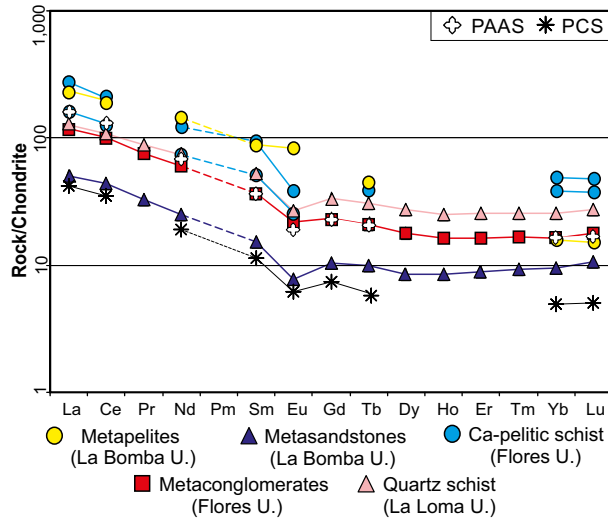


Figure 15. Chondrite-normalized (Sun and McDonough, 1989) REE diagram of samples from the Difunta Correa metasedimentary sequence.

gests that clast-forming phosphate was isotopically modified. In fact, radiogenic Sr is lost from clastic K-phyllosilicates (muscovite and K-clays minerals) during heating and can be taken up by apatite (Wasserburg *et al.*, 1964; Baadsgaard and Van Breemen, 1970), raising the  $^{87}\text{Sr}/^{86}\text{Sr}$  ratios of phosphatic rocks. Moreover, the  $^{87}\text{Sr}/^{86}\text{Sr}$  ratio of phosphatic clasts at the age of Famatinian metamorphism (ca. 465 Ma; Casquet *et al.*, 2001) is remarkably similar (0.72030 and 0.72182) to three of the siliciclastic samples from the DCMs (0.72009–0.72102; Table 3). Thus, phosphatic clasts probably underwent isotope exchange with the host siliciclastic matrix during metamorphism, favored by the relatively small size of the clasts and the strong compositional disequilibrium with the matrix. This exchange between phosphatic clasts and matrix can explain the outer zone (apatite rich) shown around some clasts (Figure 6a).

The phosphatic clasts were derived from the erosion of older phosphorite bodies. Phosphogenic events in the Neoproterozoic were episodic and of global extent (Cook and McElhinny, 1979; Bendor, 1980; Sheldon, 1980, 1981; Cook and Shergold, 2005; Xiao *et al.*, 2012) and time-related with glaciations (Cook and McElhinny, 1979; Sheldon, 1980, 1981; Cook, 1992; Bertrand-Sarfati *et al.*, 1997; Cook

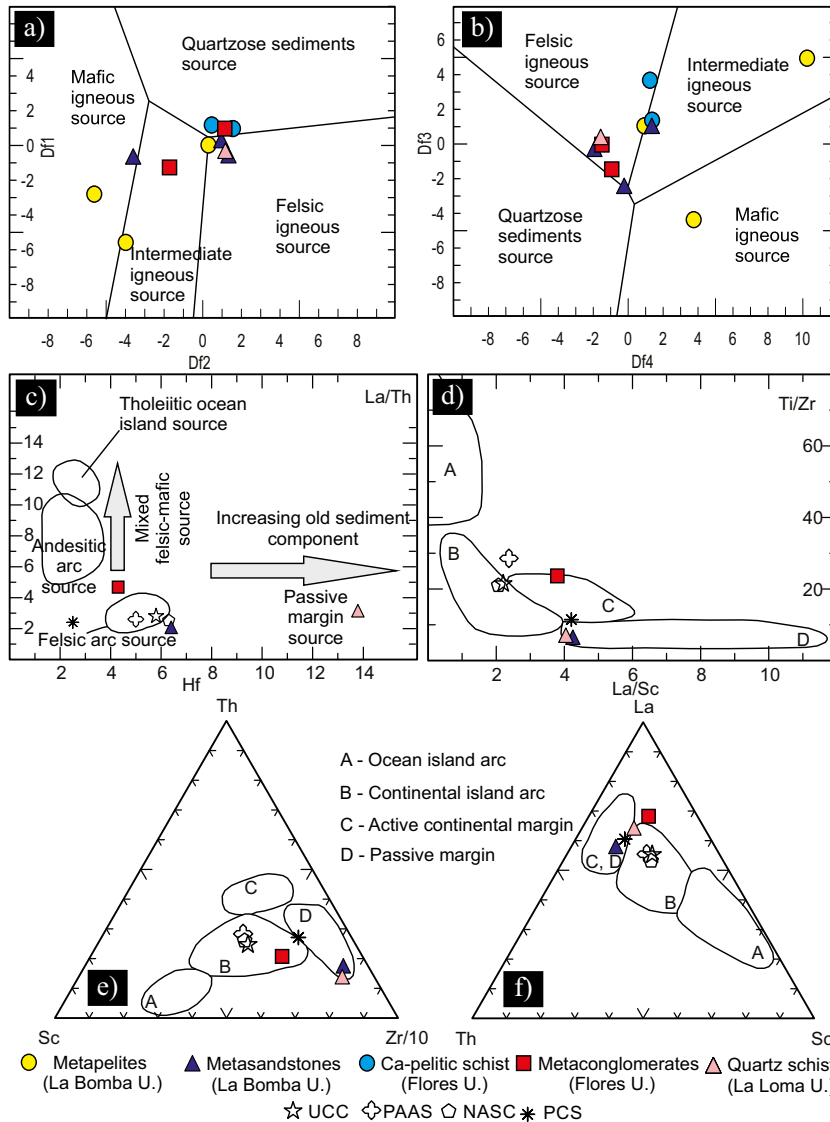


Figure 16. Source areas and tectonic setting diagrams. a) and b) Roser and Korsch, (1988). Discriminant function (Df) calculated as:  $Df1 = 56.500 \text{ TiO}_2 / \text{Al}_2\text{O}_3 - 10.879 \text{ Fe}_2\text{O}_3 / \text{Al}_2\text{O}_3 + 30.875 \text{ MgO} / \text{Al}_2\text{O}_3 - 5.404 \text{ Na}_2\text{O} / \text{Al}_2\text{O}_3 + 11.112 \text{ K}_2\text{O} / \text{Al}_2\text{O}_3 - 3.89$ ;  $Df2 = 30.638 \text{ TiO}_2 / \text{Al}_2\text{O}_3 - 12.541 \text{ Fe}_2\text{O}_3 / \text{Al}_2\text{O}_3 + 7.329 \text{ MgO} / \text{Al}_2\text{O}_3 + 12.031 \text{ Na}_2\text{O} / \text{Al}_2\text{O}_3 + 35.402 \text{ K}_2\text{O} / \text{Al}_2\text{O}_3 - 6.382$ ;  $Df3 = 0.445 \text{ TiO}_2 + 0.070 \text{ Al}_2\text{O}_3 - 0.250 \text{ Fe}_2\text{O}_3 - 1.142 \text{ MgO} + 0.438 \text{ CaO} + 1.475 \text{ Na}_2\text{O} + 1.426 \text{ K}_2\text{O} - 6.861$ ;  $Df4 = -1.773 \text{ TiO}_2 + 0.607 \text{ Al}_2\text{O}_3 + 0.760 \text{ Fe}_2\text{O}_3 - 1.500 \text{ MgO} + 0.616 \text{ CaO} + 0.509 \text{ Na}_2\text{O} - 1.224 \text{ K}_2\text{O} - 9.090$ . In all discriminant functions  $\text{Fe}_2\text{O}_3$  represents total Fe content. c) Floyd and Leveridge (1987). d) - f) Bhatia and Crook (1986).

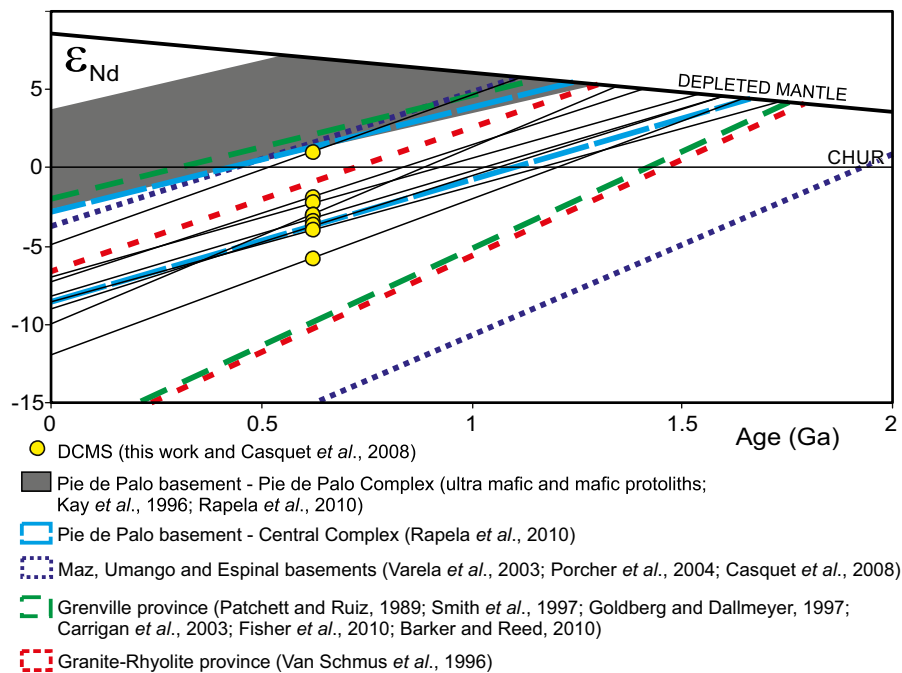


Figure 17. Comparison of isotope data of the Difunta Correa metasedimentary sequence (DCMS), with the probable source areas.

and Shergold, 2005). Considering that the marbles of the Vallecito unit probably formed in the early Ediacaran (ca. 600; Murra *et al.*, 2014; Rapela *et al.*, 2015) and that the Flores unit is stratigraphically older, we can speculate that phosphorite protoliths of the phosphatic clast probably formed near the Cryogenian-Ediacaran boundary, after the Marinoan (ca. 630 Ma) glaciation. However, unequivocal evidence for glaciogenic rocks in the lower DCMS has not been found yet.

The Neoproterozoic Sierras Bayas Group in the Tandilia System that overlies the Río de la Plata craton in southeastern Buenos Aires province, contains limestones with Sr-isotope composition similar to the DCMS marbles (Gómez Peral *et al.*, 2007, 2014). Two phosphogenic events were recognized in the Tandilia System (Gómez Peral *et al.*, 2014) that could thus be correlated with the protolith of the DCMS phosphatic clasts. In fact, Gaucher *et al.* (2008) suggested that the WSP was attached to the Río de la Plata craton in the Neoproterozoic and constituted a source area for the Sierras Bayas Group. However, detrital zircon age patterns from the Sierras Bayas Group (Rapela *et al.*, 2007; Gaucher *et al.*, 2008; Cingolani *et al.*, 2010) are significantly different from those of the DCMS (Rapela *et al.*, 2005; 2015). Although the Sierras Bayas Group contains late Mesoproterozoic zircons (between ca. 1.0 and 1.2 Ga) the dominant peaks are ca. 1.5 Ga, 2.0–2.2 Ga, 2.45 Ga and 2.7–2.8 Ga. However, the latter are absent or poorly represented in the DCMS. Therefore, detrital zircon data suggests that the source areas of the DCMS and the Sierras Bayas Group were different.

#### A paleogeographic scenario

Elemental and isotope geochemistry, as well as geological evidence suggest that the Ediacaran DCMS was probably deposited on a felsic to intermediate continental basement at a continental passive margin. The main source of the DCMS was probably the Mesoproterozoic basement that crops out in the sierras de Maz, Espinal, Umango and in the Central Complex of SPP. This basement underlaid the DCMS at the time of sedimentation. Furthermore, rocks with Nd isotope composition similar to the DCMS are also exposed in southeastern Laurentia (present position), in the Grenville and Granite-Rhyolite

provinces (Figures 17 and 18). These Laurentian sources have also been invoked –particularly the second– to explain the detrital zircon patterns with no likely sources in the WSP Mesoproterozoic basement (Rapela *et al.*, 2015).

The relationships between the DCMS and the mafic-ultramafic Pie de Palo Complex remain unknown because both are separated by the Duraznos thrust that underwent displacement during the Ordovician and the Silurian (Mulcahy *et al.*, 2011) (Figure 2). Moreover, the juvenile Pie de Palo Complex – and the Caucete Group – probably was part of the Precordillera terrane, exotic or simply allocthonous, which reached a position close to the present in the Middle to late Ordovician (Ramos *et al.* 1998; Thomas and Astini, 2003; Astini and Dávila, 2004).

Our data are compatible with the idea that the WSP basement was part of the MARA continental block (Casquet *et al.*, 2012). According to this hypothesis the southeastern passive margin of Laurentia in the Neoproterozoic was not the Appalachian margin but the eastern (present position) MARA margin, where the original sediments of the DCMS were laid down in the Puncoviscana/Clymene Ocean (Figure 18).

#### CONCLUSIONS

Metasedimentary rocks from the DCMS were classified as shales, Fe-shale and immature sandstones (wacke, sub-litharenite, litharenite and Fe-sandstone) based on chemical parameters. Modal variations of phyllosilicates (mainly pelitic component) and quartz-feldspar (psammitic component) are supported by negative correlations of  $TiO_2$ ,  $Al_2O_3$ ,  $Fe_2O_3$ ,  $MgO$  and  $K_2O$  with  $SiO_2$  and positive correlations of  $Fe_2O_3$ ,  $Al_2O_3$ , and  $K_2O$  against  $TiO_2$ , and of  $Al_2O_3$  with  $K_2O$ . Almost all samples exhibit a negative Eu anomaly ( $Eu_N/Eu_N^* = 0.61–0.74$ ), which indicates a typical continental crust composition.

At least part of the DCMS is of marine origin and the detrital sediments came from a felsic to intermediate continental source and were deposited on the continental passive margin of the MARA block

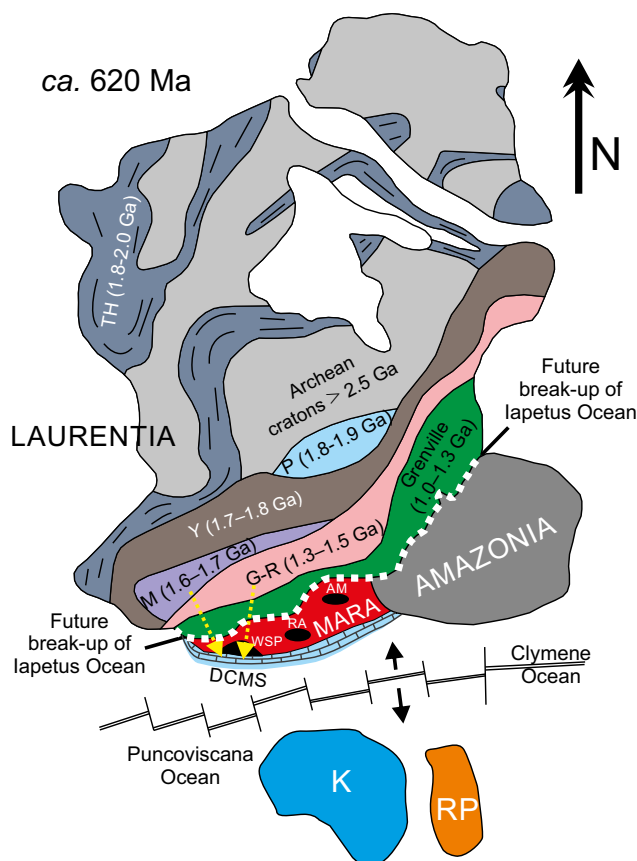


Figure 18. Paleogeographical reconstruction of Laurentia, Amazonia, Río de la Plata (RP) and Kalahari (K) showing the guessed position of MARA (acronym of Maz, Arequipa, Rio Apa) craton at ca. 620 Ma. Ages of Laurentian Precambrian orogenic belts and cratons according to Goodge *et al.* (2004) and Tohver *et al.* (2004) (Laurentia in its present position). Outcrops of basement with Mesoproterozoic ages in MARA have been included: AM, Arequipa Massif (Peru); RA, Rio Apa (Brazil, Paraguay); WSP, Western Sierras Pampeanas (Argentina). Laurentia: TH, Trans-Hudson and related mobile belts; P, Penokean orogen; Y, Yavapai orogen; M, Mazatzal orogen; G-R, Granite-Rhyolite province. Arrows show the hypothetical direction of sediment transport from the probable source areas of the Difunta Correa metasedimentary sequence (DCMS).

in the Puncoviscana/Clymene Ocean. This source probably laid in the Mesoproterozoic basement of the WSP and in the Grenville and Granite-Rhyolite provinces of southeastern Laurentia.

The phosphatic clasts from the Flores unit probably correspond to reworking of phosphorites formed after one of the worldwide Neoproterozoic glaciations. Because the age of the lower tract of the DCMS can be constrained to be younger than ca. 630 Ma we suggest that this glaciation was the Marinoan, although glaciogenic sediments have not been firmly recognized yet in this part of the Sierras Pampeanas.

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