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1 **First ⁴⁰Ar/³⁹Ar dating of intense Late Palaeogene lateritic weathering**
2 **in Peninsular India**

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Abstract

Lateritic surface processes have shaped large platform and cratons of the tropical belt. Constraining the timing of such processes is crucial to decipher their role in cratonic morphogenesis and their response to long-term climatic change and lithospheric deformation. Weathering histories have been documented for South America, Africa and Australia, but precise time constraints of the lateritic weathering processes in South India are still lacking. We present $^{40}\text{Ar}/^{39}\text{Ar}$ ages of supergene cryptomelane (K-Mn oxide) formed in the Sandur Mn ore deposits exposed on the highest lateritic paleolandsurface that once covered the Mysore plateau and the adjacent Deccan Traps. Significant $^{40}\text{Ar}/^{39}\text{Ar}$ ages are estimated between ~ 36 and ~ 26 Ma from well-defined plateaus in step heating ^{39}Ar release spectra and from best-fitted inverse isochrones. These ages constitute firm time constraints that document intense late Eocene-Oligocene lateritic weathering over Peninsular India under the influence of warm and wet climate comparable to that prevailing in tropical humid forests. These results imply that Southern India was weathered between ~ 36 and 26 Ma and may have been dissected mostly in the Neogene.

Keywords: $^{40}\text{Ar}/^{39}\text{Ar}$ radiometric dating; Cryptomelane; Supergene Mn-ore deposits; Lateritic paleolandsurface; Sandur Greenstone Belt; South India

1. Introduction

Exposed laterites and lateritic duricrusts on relics of paleolandsurfaces in cratonic domains result from successive long-duration chemical weathering and erosion processes, and as such represent morphoclimatic fingerprints of tropical regions (Ollier, 1988; Bárdossy and Aleva, 1990; Thomas, 1994; Tardy and Roquin, 1998; Beauvais and Chardon, 2013). Since the first definition and description of laterites in South India (Buchanan, 1807), many studies were conducted on Indian lateritic formations (Subramanian and Mani, 1978; Babu, 1981; Sahasrabudhe and Deshmukh, 1981; Krishna Rao et al., 1989b; Widdowson and Cox, 1996; Widdowson and Gunnell, 1999), including manganiferous supergene ore deposits (Sawkar, 1981; Krishna Rao et al., 1982, 1989a; Dessai, 1985; Roy, 1992; Mohapatra et al., 1996; Sethumadhav et al., 2010). Except few geological stratigraphic constraints, radiometric dating of surface processes involved in the formation and evolution of Indian lateritic paleolandsurfaces are still lacking. Laterites are interlayered with marine sequences of late Eocene and early Miocene age (McGowran et al., 1977; Prasad, 1983). Lateritisation processes have also been indirectly dated with Neogene palynofossils found in colluvial/alluvial sediments overlying in-situ formed Mn ore deposits ~ 100 km south of Belgaum (Fig. 1). These fossils constrain intense weathering that led to Mn ore concentration to be Neogene at the latest (Krishna Rao et al., 1989a). The age of lateritic weathering has also been roughly estimated on the basis of the paleomagnetic properties of iron oxides formed in laterites around 16°N (Schmidt et al., 1983), i.e., “Early Tertiary” (Paleogene?) for the high-level Belgaum laterites and “Late Tertiary” (Neogene?) for the low-level laterites found west of the Western Ghats escarpment (Fig.1).

Chemical rock weathering commonly resulted in the accumulation of metallic elements (Al, Fe, Mn) in residual laterites and depletion of more soluble elements (Na, K, Si, Ca). Potassium, Na and/or Ba may, however, be retained by supergene Mn oxides in large crystallographic lacuna existing between the MnO_6 octahedra chains that define the typical tunnel-structure of the hollandite manganese oxide group. Cryptomelane belongs to this group with the general stoichiometric formulae $\text{K}_x \text{Mn}^{\text{IV}}_{8-x} \text{Mn}^{\text{III}}_x \text{O}_{16}$, x ranging from 0.2 to 1 and K_x counterbalancing Mn^{III} substitution for Mn^{IV} (Burns and Burns, 1979; Turner and Buseck, 1979; Post and Burnham, 1986; Pasero, 2005). Cryptomelane is a major Mn oxide of many lateritic Mn-ore deposits in South India (Sivaprakash and Phil, 1980; Krishna Rao et al., 1982; Mohapatra et al., 1996; Sethumadhav et al., 2010). This mineral hosts radioactive ^{40}K atoms, which can decay into ^{40}Ar . The high radiogenic ^{40}Ar retentive power of cryptomelane makes this mineral a suitable chronometer for $^{40}\text{Ar}/^{39}\text{Ar}$ step heating dating of weathering processes in lateritic soils including duricrusts and manganiferous ore deposits (Vasconcelos et al., 1992; Vasconcelos et al., 1994a; Ruffet et al., 1996; Vasconcelos, 1999).

This study quantifies the radiometric ages of lateritic weathering processes that have affected the rocks of the Dharwar craton on the Mysore Plateau by $^{40}\text{Ar}/^{39}\text{Ar}$ dating of cryptomelane from supergene Mn-ore deposits. We focus on the Fe-Mn sedimentary formations belonging to the Sandur Greenstone Belt (SGB) (Figs. 1 and 2a), which hosts the biggest Indian manganese reserve (Kameshwar et al., 2000). Our results provide tangible time constraints (Late Eocene – Oligocene) on the paleoweathering processes and morphoclimatic evolution of the potentially oldest lateritic paleolandscape preserved on the Mysore plateau and Traps in southern India.

2. Geological and geomorphological setting

2.1. Sandur Greenstone Belt geology

The SGB is a synform-shape greenstone belt that consists of mafic/ultramafic volcanics and sediments (Fig. 2a) having undergone low-grade regional metamorphism during latest Archean orogeny that shaped the Dharwar craton (Roy and Biswas, 1979; Mukhopadhyay and Matin, 1993; Chadwick et al., 1996; Chardon et al., 2008). The sedimentary sequences contain elements of metavolcanic rock, which are relatively rich in Ni, Cr, Co and Sc. The upper part of the greenstone stratigraphic pile includes platform-like sediments bearing stromatolites, Mn formations and carbonaceous Mn phyllites (Fig. 2a).

In the eastern part of the belt, the upper part of the sequence includes banded iron formations (BIF), felsic volcanics and terrigenous sediments (Fig. 2a). SHRIMP U-Pb dating of zircon in two interlayered felsic volcanics indicate ages ranging from 2651 ± 18 Ma to 2659 ± 40 (Nutman et al., 1996). The BIF and Fe-Mn formations are related to marine transgression and regression cycles, respectively (Roy, 2006). These cycles implied alternating chemical and clastic sedimentation, which is also evidenced by specific iron formation facies and argillites within the BIF (Manikyamba et al., 1993). The lower part of the sequence is confined to the Deogiri region and is essentially composed of manganiferous stromatolitic limestones, dolomites and argillites (Fig. 2a) while the upper part is richer in Fe (Roy, 2000). Iron and Mn fractionation was controlled during eustatic sea-level variations by differentiated redox conditions of two stratified water masses on the shallow and deep shelf (Roy, 1992; Manikyamba and Naqvi, 1997). The Fe-Mn sediments have later been affected by low-grade metamorphism by 2475 ± 65 Ma (whole rock Pb/Pb isochron dating), which is the

maximal recrystallisation age of the limestones (Russell et al., 1996; Chadwick et al., 2000). Such a metamorphism led to Mn enrichment of the sediments (Roy, 2000), which are composed of Mn-carbonates, argillites, arenites and phyllites (Fig. 2a), and are the main protorees of the SGB Mn-ore deposits. These deposits grade upward into supergene weathering materials (Roy, 2006).

2.2. Geomorphology of southern India and Sandur Greenstone Belt

The geomorphology of southern India consists of high-elevation surfaces occurring as relics of a differentiated landscape developed on the Dharwar craton and Deccan traps (Fig. 1), and low-elevation surfaces along the western coast (Radhakrishna, 1993), both being overlain by laterites (Widdowson and Cox, 1996; Widdowson, 1997). The regional eastern dip of the high-elevation surfaces is interpreted to result from eastward tilting of Peninsular India due to the Seychelles rifting and Deccan Traps events (~ 65-63 Ma) (Radhakrishna, 1993). On the Deccan Traps, bauxitic lateritic weathering of the highest preserved landsurface must be younger than 62.3 ± 0.6 Ma, which is the age of the youngest basalts (Courtillet et al., 1986; Pande, 2002). On the Traps and the Mysore Plateau, relics of two younger surfaces step below the plateau remnants of the high bauxitic surface (900-1000 m). The three erosion surfaces remain as sparse geomorphic relics at the scale of the Mysore plateau, and have been numbered S2, S3 and S3_d, the latter being a denudation degradation landform of S3 (Gunnell, 1998). Spatially reduced relics of the highest massifs also overhang the Mysore plateau (Fig. 1).

Landsurfaces S2, S3 and S3_d are also identified on the SGB relief (Figs. 2b-c). The S2 surface levels the summits of the belt over an altitude range of 900 to 1100 m

and exposes deep in situ formed weathering mantles capped by Fe-Mn and Mn duricrusts at altitudes of ~ 1000-1016 m. Our study is mainly focused on the Mn ore open-pit mines carved in this surface (Figs. 2b-c). Relicts of paleo-land surface S3 extends below S2 over an altitude range of 750 to 850 m (Fig. 2c), and are covered by in-situ formed lateritic weathering profiles capped by iron duricrust instead of Mn ore concentrations. Relicts of land surface S3_d occur as benches below the S3 surface over an altitude range of 700-750 m above the modern river network (Figs. 2c). The S3_d surface is covered by reworked erosional materials including ferricrete and bedrock clasts originated from S3 lateritic profiles, sharing most characteristics with West African glacis/pediments (Beauvais and Chardon, 2013), but this surface is not affected by a late ferruginous or manganiferous lateritic duricrusting.

3. Material and methods

3.1. Ore bodies description and sampling

We have sampled across the main Mn ore bodies of the SGB deposits, which formed within weathered meta-sedimentary phyllites at different depths below the top of landsurface S2 at 1012-1015 m altitudes (Fig. 3). Description of samples precisely located on open-pit mine benches allowed the establishment of log sections (Fig. 3b and 3d). In Kappataswamy pit (N 14°59'59'' E 76°32'42'', Fig. 3a), the first ore body was sampled between 856 and 893 m of elevation along the E-W log section 1 on the western benches of the pit, and the second one between 904 to 935 m along the N-S log section 2 on the eastern benches (Fig. 3b). The third ore body is located in Channanghi KMK-E pit (N 14°59'50'' E 76°34'37'', Fig. 3c) at an elevation range of 926 to 994 m (log section 3, Figure 3d). The Mn-ore bodies are composed of different petrographic

facies. The float and platy Mn-ores constitute the stratiform ore type, which is conformably bedded within weathered mangan-phyllites preserving this meta-sedimentary protore structure (Mishra, 1978). The massive Mn ore is found between ~ 850 m and 892 m elevation in the first section, between 904 m and 912 m elevation in the second ore body of the Kappataswamy pit (Fig. 3b), and between 926 and 982 m elevation in Channanghi KMK-E pit (Fig. 3d). The massive ore is a podiform type with cavity fillings (geodes) and botryoidal structures, which formed by secondary Mn supergene enrichments (Mishra, 1978). Iron-Mn duricrust of ~ 10 m thickness caps the two Mn-ore deposits immediately above a segregated/tabular ore body (Figs. 3b-d).

The samples collected in the Mn-ore deposits (Fig. 3b-d) were cut with a circular saw to get the best angle for mineralogical observations on metallographic quality polished thin sections (200-300 μm -thick) made from the sawed fragments (Fig. 4a). Symmetrically to the thin sections, 500 μm -thick slabs are prepared for sub-sampling of the target cryptomelane grains suitable for $^{40}\text{Ar}/^{39}\text{Ar}$ dating (Hénocque et al., 1998).

3.2. Analytical techniques and cryptomelane characterisation

Reflected light microscopy and X-ray micro fluorescence ($\mu\text{-XRF}$) analyses allow precise characterisation of K-enriched Mn-oxide (cryptomelane) on the polished thin sections (Fig. 4b-c). Microscopic observations of thin sections were accomplished using a Leica DM-RXP microscope. The $\mu\text{-XRF}$ measurements were completed on slabs using a X-ray microscope (XGT7000, Horiba Jobin Yvon) equipped with a X-ray guide tube producing a finely focused and high-intensity beam with a 100 μm spot size, Rh X-ray tube, accelerating voltage of 30 kV and current of 1 mA. X-ray emission from the irradiated sample was detected via an energy dispersive X-ray (EDX) spectrometer

equipped with a liquid-nitrogen-cooled high purity Si detector. Micro mapping of the coupled Mn-K occurrences on slabs contributes to precise identification of cryptomelane, while Fe maps document the occurrence of Fe oxihydroxides (Fig. 4c). A Scanning electron microscope (SEM) has been also used for fine-scale observations of the cryptomelane crystals (Fig. 4d). Electron probe micro chemical analysis (EPMA) of minerals on polished thin sections was completed using a CAMECA SX-100 electron microprobe equipped with five wavelength-dispersive X-ray spectrometers (WDS).

Cryptomelane grains were separated from the slabs by hand picking. Cryptomelane separates were crushed to produce powders ($< 64\ \mu\text{m}$), which were analyzed by XRD using a Panalytical X'Pert Pro MPD with a Co $K\alpha$ X-ray source ($\lambda = 1.79\ \text{\AA}$) operating at 40 kV and 40 mA. The system is equipped with a linear X'Celerator detector and a flat monochromator. The analysis range was $10\text{-}85^\circ\ 2\theta$ with a step size of 0.033° and a counting time of 10 s per step.

After ultrasonically cleaning in absolute ethanol and conditioning in aluminium foil packets, other separated grains were irradiated for 50 hours in the TRIGA Mark-II reactor of Pavia University (Italia). The standard Taylor Creek Rhyolite sanidine 2 (TCRs-2) monitor, with an age of $28.05\ \text{Ma} \pm 0.08$ (Baksi et al., 1996), was analysed after every ten unknown samples to allow determination of the factor J. After a two-month “cooling” period, the irradiated cryptomelane grains were loaded in a double vacuum Staudacher-type furnace for step heating Ar isotopes measurements. The furnace temperature was calibrated by means of a classical thermocouple, and the gas purification was accomplished using a cold trap with liquid air and Al–Zr AP10 getters (one hot, one cold) for 8 minutes before the introduction into the VG3600 mass spectrometer. One minute was allowed for equilibration before analysis. ^{40}Ar and ^{39}Ar

were measured on a Faraday cup with a resistor of 10^{11} ohm, while ^{39}Ar , ^{38}Ar , ^{37}Ar , and ^{36}Ar were analysed using a scintillator and photomultiplier after interaction on a Daly plate. The analytical data are reported in data repositories (Tables DR1 and DR2), and the errors are quoted at the 1σ level. Ages were analysed in two ways: plateaus and inverse isochrones. Plateaus ages were calculated from at least three consecutive steps comprising up to 50% of total $^{39}\text{Ar}_K$ released and overlapping at the 2σ confidence level (Fleck et al., 1977; Maluski, 1985). Alternatively, plateau ages encompassing two steps releasing $\sim 30\text{-}40\%$ ^{39}Ar were also defined as “plateau-like” or “pseudo-plateau” (e.g., Feng and Vasconcelos, 2007; Vasconcelos et al., 2013). Ages derived from best-fit inverse isochrones in $^{36}\text{Ar}/^{40}\text{Ar}$ vs. $^{39}\text{Ar}/^{40}\text{Ar}$ correlation diagrams (Roddick et al., 1980) were accepted when mean square weighted deviation (MSWD) was less than 2.5 and the $^{40}\text{Ar}/^{36}\text{Ar}$ intercept within 2σ from the $(^{40}\text{Ar}/^{36}\text{Ar})_{\text{atm}}$ value of 298.56 ± 0.31 (Lee et al., 2006; Renne et al., 2009), since supergene cryptomelane crystallised at shallow depth, and thus incorporated excess argon, if any, in the atmospheric ratio. Age spectra also show the variations of $^{40}\text{Ar}^*$ and K/Ca (derived from $^{39}\text{Ar}/^{37}\text{Ar}$) to track the compositional effects on age.

4. Results and Discussion

Six samples (KPA-2.1, KPA-2.5, KPA-9, KPA-10, KPA-12a and KPA-12b) from the Kappataswamy pit (Fig. 3b) and one (KMK-3) from the Channanghi KMK-E pit (Fig. 3d) were analysed and dated. KPA-9 and KPA-12 show the platy preserved meta-sedimentary structure with Mn pods, while KPA-2, KPA-10 and KMK-3 are rather massive with cavity crystallisations (geode) and/or botryoidal pod concretions.

4.1. Mineralogical and geochemical characterisation of cryptomelane

Cryptomelane was identified on polished thin sections including μ -XRF and SEM imaging (Fig. 4), and characterized by microchemical (EPMA) and XRD analyses of separated grains (Figs. 5 and 6, Table 1). Samples KPA-9, KPA-12a and KPA-12b are the richest in cryptomelane, which occurs as cryptocrystalline plasmic domains formed within preserved inherited protore microstructures (Figs. 5a-c). Appropriately, sample KPA-9 exhibits a relict of Si-Mn sedimentary protore with residual smoothed small size quartz grains preserved in acicular cryptocrystalline cryptomelane (Fig. 4d). The two different generations of cryptomelane (Fig. 5a) have the same stoichiometric composition (Table 1). Separated grain of KPA-9 mainly contains cryptomelane, but also pyrolusite (βMnO_2) and quartz (Fig. 6). Sample KPA-12a is composed of two cryptomelane generations, the latest one filling the micro porosity developed across the earliest generation and/or the other mineral components such as pyrolusite and Fe oxihydroxides (Fig. 5b). The stoichiometric composition is very close to that of KPA-9 (Table. 1). Sample KPA-12b is very rich in cryptomelane (Fig. 5c) with higher K content than KPA-12a (Table. 1). The two separated grains are dominantly composed of cryptomelane but pyrolusite and goethite (γFeOOH) are also present (Fig. 6). Samples KPA-2, KPA-10 and KMK-3 are composed of successive cryptomelane crystallisations organized in banded overgrowth microstructures associated with other Mn oxides (Figs. 5d-f and 6). In the sample KPA-2, the first three bands are pure cryptomelane, the fourth is a cryptocrystalline mixture of nsutite and cryptomelane, and the fifth is dominantly cryptomelane, while pyrolusite crystallized in a cavity (Fig. 5). The cryptomelane crystals analysed in the different bands have different stoichiometric compositions with K varying from 0.45 to 0.57 atoms (Table 1). Sample KPA-10

consists of a ~ 5 mm size cryptomelane concretion with a transition zone of less than 1 mm separating it from pyrolusite (Fig. 5e). Separated grain of KPA-10 is mainly composed of cryptomelane with 0.5 K atoms (Table 1). Small quantities of pyrolusite, goethite and hematite (Fe_2O_3) are also present (Fig. 6). Sample KMK-3 (Fig. 5f) is a concretion of cryptomelane with less than 1.5 wt.% K (Table 1). The outer rims are composed of lithiophorite $(\text{Al,Li})\text{MnO}_2(\text{OH})_2$ and nsutite (γMnO_2) (Fig. 5f), which also occur as minor fractions in the XRD analysis of the separated grain (Fig. 6).

4.2. $^{40}\text{Ar}/^{39}\text{Ar}$ ages of cryptomelane

Two kinds of $^{40}\text{Ar}/^{39}\text{Ar}$ age spectra are distinguished from the step heating degassing depending on their shape and interpretation. The first group displays regular ^{39}Ar release spectra allowing calculation of plateau ages (Fig. 7a-e), which are commonly defined at the 2σ error and accounting at least for 40% of the total ^{39}Ar released from the sample (Vasconcelos and Conroy, 2003; Feng and Vasconcelos, 2007; Beauvais et al., 2008). The second group corresponds to disturbed ^{39}Ar release spectra (Fig. 8), which are more problematic to interpret. The analytical data are provided in the data repository tables DR1 and DR2.

4.2.1. Regular $^{40}\text{Ar}/^{39}\text{Ar}$ age spectra

The $^{40}\text{Ar}/^{39}\text{Ar}$ age spectra from the first group have been obtained from the first generation of cryptomelane crystallized in the meta-sedimentary protore (Figs. 5a and 5c, 7a and 7c), and for cryptomelane crystallisations in banded overgrowth microstructures of botryoids (Figs. 5f and 7d) or geodes (Figs. 5d-e, 7b and 7e). In each

case the calculated plateau and pseudo-plateau ages agree with statistically meaningful inverse isochron ages estimated in the correlation diagrams (Fig. 7).

However, these spectra present some anomalies at low and high temperatures steps. The age spectrum of KPA-9 (Fig. 7a) shows younger individual ages (steps 10 to 13), which may reflect the degassing of a late cryptomelane crystallisation (Figs. 4b and 5a). The sample KPA-10 displays an age spectrum with a clear ascending “stair case” pattern at low temperature (steps 1 to 7) with young ages affected by quite a large standard deviation (Fig. 7b). The isochron suggests a moderate excess of ^{40}Ar , which was interpreted as the result of primary mineral weathering (McDougall and Harrison, 1988; Vasconcelos et al., 1994a), which is unlikely in this sample (Figs. 5e and 7b, Table 1). The higher amounts of released $^{40}\text{Ar}^*$ and ^{39}Ar in intermediate temperatures (step 8 to 11), however, allow the best estimation of the cryptomelane crystallisation age (Vasconcelos, 1999; Hautmann and Lippolt, 2000; Li and Vasconcelos, 2002; Vasconcelos and Conroy, 2003; Feng and Vasconcelos, 2007). Sample KPA-12b is characterised by well-constrained plateau and isochron ages (Fig. 7c).

The age spectrum of KMK-3 exhibits an ascending “stair case” pattern at high temperature (steps 8, 9, 10, 13 and 14) with older ages than the plateau age (Fig. 7d). In such a case, it is usually admitted that radiogenic argon is released by two cryptomelane phases, the older one retaining radiogenic ^{40}Ar better than the younger one (Feng and Vasconcelos, 2007) resulting in a maximum plateau age estimate and an older integrated age (Li and Vasconcelos, 2002). The plateau age of KMK-3 has also quite a large error due to relatively low K content (Table 1 and Fig. 7d), and higher atmospheric correction, the isochron being less sensible and thus more precise.

The age spectrum of KPA-2.1 (Fig. 7e) is more disturbed and does not allow calculation of a standard plateau age (Fleck et al., 1977). Pseudo-plateaus ages have been, however, previously estimated from similar spectra by considering only two concordant steps including ~ 40% of ^{39}Ar released (e.g., Feng and Vasconcelos, 2007; Vasconcelos et al., 2013). Two pseudo-plateaus occur in intermediate temperature steps 4-5 and 7-8, allowing for a concordant age estimate of 35.86 ± 1.10 Ma by combining the four steps at the 2σ level (Fig. 7e). We note that the best-fitted isochron supports a similar age (Fig. 7e). The high temperature steps 13 to 16 suggest a hypogene contaminant (Vasconcelos et al., 1994a; Ruffet et al., 1996; Li and Vasconcelos, 2002), which could have aged the intermediate steps (e.g., the step 6). The low K content extraction in high temperature steps could have also suffered from imprecise blank corrections. Therefore, similarly to the sample KMK-3 (Fig. 7d), the best-fitted isochron might reasonably provide a more precise age (Fig. 7e).

4.2.2. *Disturbed $^{40}\text{Ar}/^{39}\text{Ar}$ age spectra*

The ^{39}Ar release spectra from the second group are characterized by an upward convex shape (Fig. 8) suggestive of a mixing of two or more supergene cryptomelane phases (Vasconcelos et al., 1995; Ruffet et al., 1996; Hautmann and Lippolt, 2000), which may result from the handpicking of single grains. Phase mixing is obvious in sample KPA-12a showing late cryptomelane crystallisation precipitated into the porosity ($\leq 500 \mu\text{m}$) generated by the dissolution of a first plasmic cryptomelane generation (Fig. 5b). Such a process may result in remobilization of low energy sites (Colin et al., 2005; Beauvais et al., 2008). Phase mixing often occurs during separation of grains from multiple overgrowth bands (Vasconcelos and Conroy, 2003). The grain

of KPA-2.5 was tentatively separated from a thin band (~ 500 μm) between a late pyrolusite crystallisation in a geodic void and a band composed of a mixing of cryptomelane and nsutite (Fig. 5d). The growth rate of cryptomelane crystals in banded overgrowth structures (botryoids or geodes) is generally very low (Vasconcelos et al., 1992; Dammer et al., 1996; Hénocque et al., 1998) and may result in very small size ($\leq 1 \mu\text{m}$) intimately intergrown cryptomelane phases in a same grain (Vasconcelos et al., 1994b; Vasconcelos, 1999).

The pronounced upward convex shape of ^{39}Ar release spectra precludes plateau age calculation, while correlation diagrams only allow estimation of minimum and maximum ages of the oldest and youngest mixing phases, respectively (Fig. 8a-b). Each step of the ^{39}Ar release spectra yields a mixing age of the intergrown cryptomelane generations (Vasconcelos, 1999; Vasconcelos and Conroy, 2003; Feng and Vasconcelos, 2007), implying that the ages in low and high temperature steps are overestimates of the youngest phase, while the ages in intermediate temperature steps are underestimates of the oldest phase (see also Ruffet et al., 1996; Beauvais et al., 2008). This interpretation is mainly based on previous experiments (Vasconcelos et al., 1995), which showed a maximal degassing of $^{40}\text{Ar}^*$ and ^{39}Ar between 700°C and 900°C with a peak at 800°C. In such a range of temperatures, the two isotopic components are released from the mineral structures (Vasconcelos et al., 1994b). Therefore, only maximum and minimum ages might be estimated from upward convex spectra in the temperature range specific of supergene cryptomelane degassing (Fig. 8). For KPA-2.5 (Fig. 8a), a minimum age of ~ 46 Ma (step 7) and a maximum age of ~ 30 Ma (step 10) could be at best estimated for the crystallisation of the earliest and latest cryptomelane phases, respectively. For KPA-12a the maximum and minimum ages of ~ 29 Ma (step

10) and ~ 34 Ma (step 8) are estimated for the youngest and oldest phases, respectively, in agreement with the estimated isochron age limits, although the MSWD value of the second isochron is quite large (Fig. 8b).

4.2.3. Cryptomelane $^{40}\text{Ar}/^{39}\text{Ar}$ ages summary

Despite mineral phase mixing of some samples, five significant plateau and isochron ages are estimated between ~ 26 and 36 Ma. The older ages (~ 33.5 and 36 Ma) are obtained either for the first cryptomelane generation in samples showing the preserved meta-sedimentary protore structure (Figs. 5a-c, 7a-c) or the first crystallisation band of an overgrowth geodic microstructure (Fig. 5d and 7e). The younger ages (~ 26-28 Ma) characterize the late cryptomelane crystallisations of banded overgrowth microstructures (Figs. 5e-f, 7b and 7d). The ages of the different mixing phases of KPA-2.5 and KPA-12a cannot be precisely determined. However, the two cryptomelane phase of KPA-12a (Fig. 5b) may have crystallised between ~ 29 Ma and ~ 34 Ma (Fig. 8c) that bracket the age of KPA-12b (Fig. 7c) sampled on the same open mine bench. The maximum age (~ 30 Ma) estimated for the youngest cryptomelane of KPA-2.5 is quite realistic compared to the older cryptomelane grain (KPA-2.1) separated from the same sample (Fig. 5d and 7e). Despite their limited number, our results usefully document Late Paleogene weathering history of southern India.

4.3. Potential implications for lateritic weathering and morphogenesis in southern India

The formation of cryptomelane in Mn-ore deposits is enhanced on landsurfaces covered by a humid tropical forest reflecting warm and/or at least wet climatic conditions (Vasconcelos et al., 1994a), like those prevailing for bauxite genesis in

tropical cratonic regions (Bárdossy and Aleva, 1990). Such hydroclimatic conditions are required for complete weathering of K-phyllosilicates into kaolinite with release of K^+ in Mn^{2+} -rich acidic solutions (pH ~ 4.5-5) that may ultimately lead to cryptomelane crystallisation under appropriate redox conditions (Parc et al., 1989). Therefore, massive formation of cryptomelane in Mn ore deposits and/or lateritic weathering profiles is a good indicator of humid/warm paleoclimatic conditions (Vasconcelos et al., 1994a; Vasconcelos and Conroy, 2003; Beauvais et al., 2008). Under such climatic conditions, wide areas were colonised by humid tropical forests over most of peninsular India (Kent and Muttoni, 2008; Chatterjee et al., 2013) enhancing the pervasive development of lateritic weathering profiles with differential sequestration of Al, Fe and Mn.

On the basis of global paleoclimatic reconstructions (e.g., Parrish et al., 1982), Bárdossy and Aleva (1990) argued for intermittent Cenozoic bauxitic weathering in South India. Such a bauxitic weathering intermittency was not observed in West Africa, where bauxites preserved in current landscapes have been dated from Late Palaeocene to Eocene at the latest (Prasad, 1983; Colin et al., 2005; Beauvais et al., 2008; Beauvais and Chardon, 2013). Nevertheless, two stages of lateritic weathering processes have been described in the bauxitic deposits of the Boknur-Navge plateau near Belgaum (Krishna Rao et al., 1989b) (Fig. 1). The first bauxitic stage characterizes the syngenetic lateritic weathering of the Deccan basalts under the action of slightly acidic meteoric waters, whereas the second stage is a late weathering epigenetic process of the early bauxite, under the effect of highly acidic meteoric Si-rich waters (Krishna Rao et al., 1989b). The Boknur-Navge bauxitic plateau is also a remnant of paleolandsurface S2 correlated to the S2 remnant of the SGB (Gunnell, 1998). Syngenetic bauxitic weathering on paleolandsurface S2 was likely active in the “Early Tertiary” (Paleogene

?) as suggested by paleomagnetic investigations on Fe-oxihydroxides in bauxites and laterites of the nearby Belgaum area (Schmidt et al., 1983; Kumar, 1986). Although no precise ^{40}Ar - ^{39}Ar age correlative of syngenetic bauxitic weathering of the Traps basalts was obtained in this study, lateritic weathering up to ~ 46 Ma may not be precluded (Fig. 8a). Epigenetic lateritic weathering of paleolandsurface S2 (Krishna Rao et al., 1989b) could however be recorded by our $^{40}\text{Ar}/^{39}\text{Ar}$ ages set.

The obtained ages (~26, ~ 28, ~ 33.5 and ~ 36 Ma) attest for intense Late Eocene to Mid-Oligocene lateritic weathering(s) of the highest paleolandsurface (S2) preserved as sparse relics on both the Mysore plateau and the Deccan Traps. These results suggest that the high-elevation landsurface of Peninsular India evolved under a wet and warm climate over a protracted period in the Late Paleogene, essentially during the Oligocene. Repeated lateritic weathering between 26 and 36 Ma apparently correlates with low clastic sediment fluxes and dominant carbonate sedimentation on the Arabian sea margin (Campanile et al., 2008) but the links between inland weathering, erosion and offshore sedimentation are poorly understood. Our $^{40}\text{Ar}/^{39}\text{Ar}$ results might also suggest that the dissection and rejuvenation of the high-elevation lateritic surface of Peninsular India took place in the Neogene as classically inferred (Radhakrishna, 1993). However, the relative contributions of lithospheric deformation and climate change to geomorphic rejuvenation of Peninsular India in the Neogene remain to be investigated.

5. Conclusion

We provide time constraints on lateritic weathering having affected Southern India by using $^{40}\text{Ar}/^{39}\text{Ar}$ radiometric dating of supergene cryptomelane in Mn ore deposits underlying a relict paleolandsurface (S2) that once covered the Mysore plateau

and Deccan Traps. Well-constrained plateau ages and best-fitted inverse isochrons allow defining $^{40}\text{Ar}/^{39}\text{Ar}$ ages at ~ 26, ~ 28, ~ 33.5 and ~ 36 Ma. These results document intense lateritic weathering in Southern India during the latest Eocene and the Oligocene under the effect of warm and wet climate similar to that prevailing in humid forest tropical environments. More specifically, the highest regional lateritic paleolandsurface (S2) of Peninsular India underwent chemical weathering at least between ~ 36 and ~ 26 Ma and was possibly dissected and rejuvenated afterwards, which suggests that most of the current relief of South Indian high-elevation terrains was acquired in the Neogene.

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666

667 **Figures and table caption**

668 Figure 1. Simplified map of Peninsular India showing first-order geomorphic and
669 geological features (adapted from Radhakrishna, 1993). B= Bababudan; N= Nilghiris;
670 S= Shevaroy.

671 Figure 2. Simplified geology of the Sandur Greenstone Belt adapted from Chadwick et
672 al. (1996). (b) Corresponding topography (ASTER GDEM30) and delineation of relicts
673 of paleosurfaces S2 and S3. (c) Topographic cross-sections (lines of section are located
674 on Figure 2b).

675 Figure 3. North-looking Google Earth views of (a) Kappataswamy and (b) Channanghi
676 KMK-E pits showing the location of the cross-sections through the Mn ore bodies. (c)
677 Logs of sections 1 and 2. (d) Log of section 3. Black numbered samples are those
678 analysed and dated in the current study.

679 Figure 4. Characterisation of procedure used for separating cryptomelane grains in
680 samples. (a) Polished thin section observed by (b) reflected light microscopy and (c) X-
681 ray micro fluorescence, which yields a composite picture with Mn in blue, K in green
682 and Fe in red. (d) SEM image showing typical cryptomelane acicular crystals in a
683 composite weathering matrix, which preserves altered quartz grains of the protore. P =
684 pyrolusite; C = cryptomelane; Pr = protore; Q = quartz.

685 Figure 5. Reflected-light photomicrographs of mineral assemblages of dated samples
686 showing EPMA numbered targets. (a) Two generations of cryptomelane (C1, C2) and
687 Pyrolusite (P) formed into the Mn protore (Pr). (b) Coexistence of two generations of
688 cryptomelane (C1, C2) with pyrolusite (P) and Fe-oxihydroxide (Fe). (c) Cryptomelane

(C) and pyrolusite with some Fe-oxihydroxides (Fe). (d) Cavity filling by banded overgrowth of a succession of cryptomelane (C) with pyrolusite (P) and a mixed overgrowth on nsutite+ cryptomelane (NC). (e) Massive cryptomelane and pyrolusite with some oxihydroxides (Fe). (f) Massive banded overgrowth of cryptomelane (C) with pyrolusite (P), lithiophorite (L) and nsutite. Sample names are reported on each photomicrograph.

Figure 6. X-ray diagrams of separated cryptomelane sub-samples. C = cryptomelane; L = lithiophorite; n = nsutite; Q = quartz; p = pyrolusite; G = goethite; h = hematite

Figure 7. ^{39}Ar - ^{40}Ar age spectra showing a well-defined plateau, with K/Ca (black) and Ar^* (grey) step curves (left), and inverse isochrone diagrams (right) of cryptomelane separates from Kappataswamy open pit samples (a-c and e), and Channanghi KMK-E open pit sample (d). MSWD = mean square weighted deviation.

Figure 8. Disturbed ^{39}Ar - ^{40}Ar age spectra with K/Ca (black) and Ar^* (grey) step curves (left) and inverse isochrone diagrams (right) of cryptomelane separates from Kappataswamy open pit samples. (a) Separated phase of KPA-2.5 (target 5 in Figure 5d). Samples KPA-2.5 and KPA-12a do not allow calculation of plateau age but only maximum and minimum ages of the mixed youngest and oldest phases, respectively (stippled line depicted from bold numbered steps in the spectra). MSWD = mean square weighted deviation.

Table 1. Electron Probe Micro Analyses (EPMA) of targeted cryptomelane phases (in bold) and other Mn oxides shown in the Figure 5. Each analyse is the average of three $5\mu\text{m}$ micro spots in a $50\mu\text{m}$ sized micro domain.

Highlights

- * First radiochronological dating of lateritic weathering in southern India
- * Late Palaeogene chemical weathering on the highest paleolandsurface preserved on the Mysore plateau and Deccan Traps
- * Potential rejuvenation and acquisition of the current relief of southern India in the Neogene

FIG. 1

Figure. 2
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FIG. 2

Figure. 3
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FIG. 3

Figure. 4
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FIG. 4

Figure. 5
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FIG. 5

FIG. 6

FIG. 7

Figure. 8

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FIG. 8

Table. 1
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Sample ID	KMK-3		KPA-2						KPA-9			KPA-10		KPA-12a		KPA-12b	
	1	2	1	2	3	4	5	6	1	2	3	1	2	1	2	1	2
	C	P	C	C	C	C	C	P	C	C	Pr	C	P	C	P	C	P
wt. %																	
Mn	60.15	60.97	59.72	60.01	59.38	61.07	59.13	62.95	56.62	57.52	12.38	59.11	60.67	59.10	61.86	59.82	60.37
Al	0.15	0.30	0.12	0.30	0.22	0.33	0.20	0.00	1.41	1.24	0.37	0.20	0.29	0.45	0.11	0.18	0.37
Fe	0.13	0.13	0.12	0.13	0.13	0.13	0.13	0.14	0.58	0.51	0.17	0.05	0.13	0.81	0.14	0.17	0.85
K	1.41	0.06	2.49	2.26	2.44	0.94	3.03	-	2.13	2.19	0.50	2.65	0.87	2.19	0.19	3.29	0.29
Na	0.03	0.01	0.09	0.14	0.11	0.03	0.08	-	0.06	0.08	0.01	0.10	0.13	0.08	0.02	0.07	0.01
Zn	-	-	0.01	0.02	0.02	0.01	0.02	-	0.01	0.02	0.01	0.01	0.01	0.01	0.01	0.01	0.01
Co	-	-	-	-	-	0.01	-	-	0.01	0.01	0.01	-	0.01	0.01	-	0.01	0.01
Ba	0.07	0.02	0.01	-	-	0.01	0.01	0.01	0.01	0.01	-	0.01	-	-	0.01	0.01	0.02
Ni	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
Ti	-	-	-	-	-	-	-	-	0.01	0.01	-	-	-	-	-	-	-
Mg	-	0.00	0.01	0.01	0.01	0.01	0.00	-	0.01	0.01	-	0.02	0.01	0.01	-	0.00	0.02
Si	0.06	0.33	0.07	0.08	0.06	0.09	0.07	0.08	0.38	0.20	37.58	0.06	0.10	0.08	0.17	0.07	0.12
Ca	0.02	0.04	0.12	0.10	0.11	0.06	0.11	0.00	0.11	0.12	0.02	0.18	0.10	0.11	0.04	0.13	0.15
O	35.59	36.23	35.60	35.92	35.49	36.24	35.44	36.80	35.35	35.52	50.53	35.36	36.02	35.68	36.44	35.89	36.02
Total	97.62	98.11	98.36	98.98	97.99	98.92	98.24	99.99	96.68	97.41	101.6	97.75	98.35	98.53	98.99	99.64	98.24
Stoichiometric composition of cryptomelane based on 16 oxygens																	
K	0.4		0.45	0.41	0.45	0.17	0.57		0.42	0.4		0.5		0.4		0.61	
Mn ^{IV}	7.6		7.55	7.59	7.55	7.83	7.43		7.58	7.6		7.5		7.6		7.39	
Mn ^{III}	0.4		0.45	0.41	0.45	0.17	0.57		0.42	0.4		0.5		0.4		0.61	

- : not detected ; C = Cryptomelane ; P = Pyrolusite ; Pr = Si-Mn protore

Table. 1