



LOCATING THE FLOOD/POST-FLOOD BOUNDARY USING THE RELATIVE DATING OF THE WEATHERING OF ORE DEPOSITS

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ABSTRACT

Erosion at the end of and after the Flood exposed the tops of primary ore deposits to subsequent weathering and the formation of supergene minerals. The $^{40}\text{Ar}/^{39}\text{Ar}$ and (U-Th)/He radioisotope methods applied to these supergene minerals provide the dates for when these minerals formed. But given the documented problems with the radioisotope methods, they can only provide at best relative dates. When the Flood waters retreated, the ground surface was dry, but extra time was needed to allow the water table to drop, soil to form and plants to grow before Noah stepped off the Ark, which marked the end of the Flood event. The weathering front then progressed downwards during the early post-Flood decades for supergene minerals to start forming. Residual post-Flood catastrophism may have involved mountains still rising and ore deposits still forming, such as the porphyry copper deposits associated with granite intrusions as the Andes continued to rise. Erosion exposed those later-formed ore deposits to subsequent weathering to produce supergene minerals well into the early post-Flood era. Thus, the relative ages of the supergene iron oxides, and potassium-bearing sulfates and manganese oxides produced span the whole Cenozoic. It is proposed that relative dates for the first formation of supergene minerals can possibly be used as a criterion for determining the placement of the Flood/post-Flood boundary at the K-Pg boundary with a relative age of 66 Ma. The few slightly earlier relative ages likely resulted from weathering that commenced before the Flood event ended, though deep weathering would have required decades. The spread of relative ages through the Cenozoic thus represents the progressive formation of supergene minerals as primary ore deposits emplaced during, and maybe after, the Flood were subsequently exposed to weathering by residual catastrophism. Continuing investigation of this criterion for placement of the Flood/post-Flood boundary seems warranted.

KEY WORDS

Flood/post-Flood boundary, ore deposits, weathering, erosion, supergene minerals, $^{40}\text{Ar}/^{39}\text{Ar}$ dating, (U-Th)/He dating, relative dating, K-Pg boundary

INTRODUCTION

The location of the Flood/post-Flood boundary in the geologic record has long been a contentious issue debated in the creationist literature (Snelling 2014a, c). However, it is crucial for the coherency and advancement of the Creation-Flood model of earth history for this issue to be resolved definitively. There are primarily two contenders which have been proposed for this boundary's location.

The first proposed location is at the Cretaceous/Paleogene (K/Pg) boundary (formerly known as the Cretaceous/Tertiary or K/T boundary) (Austin et al 1994). Above this boundary are lineages of mammal fossils that link extant mammals to their fossilized within-kind ancestors, even within the same geographic regions (Ross 2012, 2014a, b; Snelling 2014c; Whitmore and Wise 2008). Opponents of this view have failed to explain in a consistent and coherent scenario why both the Cenozoic fossil and related extant mammals are found in the same geographic regions, unless all belong to the same post-Flood population lineages (Oard 2007, 2010c, 2013a; Walker 2014a, b). For example, if Miocene fossil kangaroos found only in Australia are pre-Flood kangaroos buried by the Flood, then it is highly fortuitous that the related living kangaroos are also only found in Australia. However, Clarey (2016) raised the question of how large mammals could disperse in a post-Flood world prior to the Ice Age land bridges.

The alternate proposed location is at or very near the Pliocene/Pleistocene boundary (or what used to be called the Tertiary/Quaternary boundary). Several arguments have been advanced. Some proponents argue that there is too large a volume of Cenozoic sedimentary and volcanic rocks, which would make the post-Flood world too catastrophic, when instead they maintain those sediments and volcanics required Flood rate and scale geologic processes well beyond local catastrophes (Clarey 2015a, c, 2017; Holt 1996; Oard 2010b, 2011, 2013a). Others point to examples of large-scale Cenozoic erosion and its products that they claim had to happen when the Flood waters were receding in order to explain the scale of erosion that is observed (Holt 1996; Oard 2004, 2007, 2013a, b).

Whitmore and Garner (2008) attempted to develop a list of criteria which can be used to delineate the location of this boundary in strata sequences. Included in their list was the category "true paleosols". They reasoned that since the Flood involved the rapid accumulation of sediments without the passage of long periods between the deposition of individual layers, we would not expect the development of true soil horizons during the Flood. On the other hand, after the Flood weathered horizons and paleosols would be expected to become more abundant due to the extended time for their development. Nevertheless, they ranked this as only a tertiary criterion, because true paleosols are difficult to diagnose and they

could also be present in pre-Flood strata sequences.

However, the supergene weathering of many ore deposits at today's earth surface may still offer an additional criterion for determining the location of the Flood/post-Flood boundary in the geologic record. This is because other criteria can be used to establish that the primary ore deposits were indeed formed during the Flood or even pre-Flood. For example, most porphyry copper deposits are hosted by granite plutons which intrude into fossil-bearing, Flood-deposited sedimentary layers. On the other hand, Precambrian banded iron formations (BIFs) were largely formed before the Flood and may even date back to the Creation Week (Snelling 2009) but were only exposed at the earth's surface as a result of the erosion at the end of the Flood as the waters retreated. Furthermore, lateritic iron and manganese deposits were deposited as sedimentary layers on top of or between fossil-bearing, Flood-deposited sedimentary layers. Of course, some of these porphyry copper and lateritic iron and manganese deposits may also have formed in the first decades after the Flood due to the declining residual catastrophism.

Some consideration needs to also be given to the chronology of the Flood, and especially the timing and actions of the Flood waters when they retreated. This is highly relevant to the question of when the primary ore deposits were exposed by erosion to weathering, which then initiated the formation of the supergene minerals. For example, the granite plutons hosting the porphyry copper deposits were intruded some 2-5 km below the ground surface at that time, so that 2-5 km of overlying host rocks had to first be eroded away to expose the primary copper ore to weathering and formation of the supergene minerals that have been radioisotope dated. The timing of this weathering relative to the end of the Flood is crucial to considering whether the radioisotope ages derived for the supergene minerals can pinpoint where the Flood/post-Flood boundary should be placed in the geologic record.

SUPERGENE WEATHERING OF ORE DEPOSITS

Supergene metallic ore deposits form when common rock types or deeply buried primary metallic ore bodies are exposed at or within 1000 m of the earth's surface. It is necessary to avoid confusion here by defining the term supergene. Supergene is "said of a mineral deposit or enrichment formed near the surface, commonly by descending solutions" (Neuendorf et al. 2005, p.645). Thus, supergene alteration of primary ore deposits, which is due to the downward passage of weathering and oxygenated groundwaters and occurs at in situ ground temperatures, should not be confused with hypogene or hydrothermal alteration of primary ore deposits. Hypogene is "said of a mineral deposit formed by ascending solutions" (Neuendorf et al. 2005, p.315). Hydrothermal alteration generally produces a different set of minerals from those produced by supergene alteration. And hydrothermal alteration usually occurs at depths >1 km, whereas supergene alteration occurs at <1 km depth and is related to groundwater fluctuations and oxygen in the atmosphere penetrating downwards from the land surface.

Rocks and ore deposits that were formed at high temperatures and high pressures when exposed at and near the earth's surface have their equilibrium disturbed. This causes their mineral constituents to react and undergo transformations so as to adjust to the new

lower temperatures, pressures, and higher oxygen concentrations and moisture conditions. They undergo chemical weathering, which promotes the oxidation, dissolution, remobilization, re-precipitation, and re-concentration of metals of economic interest.

Recurrent dissolution, transport, and re-deposition of metals through time has created a chemically stratified weathering profile (Reich and Vasconcelos 2015) that contains a comprehensive record of these chemical reactions which have occurred at the earth's surface. Today's rates of these reactions are invariably climate-dependent, reflecting ambient temperature, availability of liquid water (involving rainfall intensity and seasonality), evapotranspiration rates, and biological and microbiological activity (Vasconcelos 1999a).

The phenomenon known as supergene enrichment refers to the secondary, in situ, accumulation of metals (for example, Cu, Zn, Ag, Au, Ni, or U) as a result of three essential processes:

- 1) The electrochemical oxidation of primary sulfides, oxides or native metals (for example, native copper Cu^0 to Cu^{2+});
- 2) The transport of the released metals as soluble metal species (for example, CuSO_4^0 , AuCl_4^{1-}); and
- 3) The reprecipitation of the metals by reduction (for example, Cu^{2+} to native copper Cu^0), by supersaturation (for example, Mg^{2+} in magnesite deposits), or by cation-exchange (for example, Ni^{2+} exchange for Mg^{2+} in smectite- or serpentine-group minerals).

In particular, oxidation processes leading to mineral leaching are commonly catalyzed by specialized Fe- and S-oxidizing bacteria. Oxidation processes are also active in the surficial vadose zone and the capillary fringe above the water table. Leaching processes also respond to changes in physicochemical properties, such as the partial pressure of oxygen ($p\text{O}_2$), and its effect on the redox potential (Eh), as well as the activity of H^+ of descending aqueous solutions (Reich et al. 2009; Sillitoe 2005).

Weathering profiles hosting supergene ore deposits may extend down to 1000 m below the surface, but they are mostly inaccessible to scientific investigation. When these systems are drilled for their mineral potential and eventually exposed by open-pit mining operations – some open pits may be 5–6 km wide and more than 1 km deep – they provide access to paleoclimatic records that are otherwise unavailable.

Minerals found in supergene ore deposits record information about chemical reactions and geochemical (and paleoclimatic) conditions prevailing during the formation of the deposits. Detailed descriptions of these chemical reactions which form the main minerals produced by supergene weathering of ore deposits are provided in the Appendix.

DATING OF SUPERGENE MINERALS

Supergene metal deposits host a comprehensive record of climate-driven geochemical reactions that may span the entire Cenozoic. Products of these reactions can be dated by a variety of radiogenic isotopic methods, such as $^{40}\text{Ar}/^{39}\text{Ar}$, (U–Th)/He, U–Pb, and U-series. High spatial resolution methods have distinct advantages when dating minerals from soils and weathering profiles that contain complex assemblages of intimately intergrown minerals precipitated at distinctly different times. The most commonly

used geochronological methods for weathering geochronology are $^{40}\text{Ar}/^{39}\text{Ar}$ laser incremental heating analysis of K-bearing supergene minerals (particularly the hollandite-group K–Mn-oxides and alunite group sulfates) and (U–Th)/He analysis of supergene oxides and hydroxides (hematite and goethite).

Some of the minerals found in the supergene weathering zones of ore deposits can be dated using the $^{40}\text{Ar}/^{39}\text{Ar}$ radioisotope method (Vasconcelos 1999a, b). It could be argued that method is unreliable and so cannot provide absolute dates (Snelling 2016a, 2017a). However, when it is carefully used it can still provide relative dates, anchored to the biblical parameters for subdividing and dating the rock record (Vardiman, Snelling and Chaffin 2005). So supergene weathering minerals such as alunite $[\text{KAl}_3(\text{SO}_4)_2(\text{OH})_6]$, jarosite $[\text{KFe}^{3+}_3(\text{OH})_6(\text{SO}_4)_2]$, and hollandite $[\text{Ba}, \text{K}(\text{Mn}^{4+}_6\text{Mn}^{3+}_2)\text{O}_{16}]$ have proven to be good candidates for $^{40}\text{Ar}/^{39}\text{Ar}$ dating of the time they formed. That potentially becomes the relative times these minerals formed after the Flood waters receded and the land was exposed to weathering.

The $^{40}\text{Ar}/^{39}\text{Ar}$ method is most widely applied because many minerals precipitated by weathering reactions contain K (as detailed in Table 1 in the Appendix), and many of these minerals are relatively stable once formed. If these phases retain ^{40}K and ^{40}Ar quantitatively, then nuclide abundances can be used to determine when the mineral formed (details in Vasconcelos 1999b). The analysis of a representative suite of K-bearing supergene minerals from a vertical section through a weathering profile may be used to estimate weathering rates and to infer paleoclimatic conditions (de Oliveira Carmo and Vasconcelos 2006). A probability density distribution of mineral precipitation ages identifies times in the past when climatic conditions favored mineral dissolution and reprecipitation. Chemical reactions recorded by mineral precipitation require water as a reactant; therefore, the frequency distribution of ages through time permits one to identify periods in the geological past that were relatively wet (Vasconcelos 1999a, b).

As with K-bearing minerals, the Fe-bearing goethites and hematites generated by water–rock interactions during the formation of supergene ore bodies can be dated. The decay of trace amounts of U and Th in goethite and hematite (Lippolt et al. 1998) results in ^4He by-products that can be used for dating mineral precipitation, as long as the ^4He , U, and Th are retained. But properly quantifying ^4He retention in goethite and hematite was not possible until scientists combined the (U–Th)/He and $^4\text{He}/^3\text{He}$ methods (Shuster et al. 2005; Heim et al. 2006; Monteiro et al. 2014).

The interpretation of geochronological results for samples from weathering/supergene profiles of ore deposits requires the identification of the dated reactions (Vasconcelos 1999a). For this purpose the dated mineral samples must be characterized petrographically, and the reactions must be identified from paragenetic relationships.

METHODS

Results from applying this methodology are already available in the relevant literature (Vasconcelos et al. 2015). Geochronological data was compiled from a number of studies – South American supergene copper deposits (Alpers and Brimhall 1988; Arancibia et al. 2006; Mote et al. 2001; Sillitoe and McKee 1996; Vasconcelos

1999a); South American supergene manganese deposits (de Oliveira Carmo and Vasconcelos 2006; Spier et al. 2006; Vasconcelos 1999b); other supergene manganese deposits in Africa (Beauvais et al. 2008), Australia (Dammer et al. 1999; Feng and Vasconcelos 2007; Li and Vasconcelos 2002; Vasconcelos 2002; Vasconcelos et al. 2013), China (Deng et al. 2014; Li et al. 2007), India (Bonnet et al. 2014), and Europe (Hautmann and Lippolt 2000); and supergene iron deposits in Australia (Heim et al. 2006; Vasconcelos 1999b; Vasconcelos et al. 2013), and Brazil (Monteiro et al. 2014).

The frequency of mineral precipitation, determined by dating a representative number of samples of a particular mineral collected from distinct parts of the supergene orebody, reflects times in the geological past when weathering conditions were conducive to water–rock interaction. The frequency of mineral precipitation through time permits identifying periods in the geological past when climatic conditions were most conducive to chemical weathering and supergene ore genesis.

RESULTS

The available data were compiled in three histograms in Fig. 1 according to the supergene minerals dated (Mn oxides, goethite and hematite, or alunite-jarosite) and the dating methods used [$^{40}\text{Ar}/^{39}\text{Ar}$ or (U–Th)/He], showing the frequencies of conventional ages in the range 0–70 Ma in 1 Ma increments.

The distribution of these supergene minerals through time appears to help to identify periods in the geological past conducive to the dissolution and reprecipitation of ore elements in the weathering environment. Each mineral species records slightly different conditions. For example, Mn oxides record wetter conditions needed for the reduction–dissolution processes needed to dissolve and reprecipitate Mn oxides in the weathering environment. In contrast, the formation and preservation of supergene alunite and jarosite required relatively dry conditions after mineral precipitation, typically achieved by drawdown of the water table during a transition from humid or semi-arid to hyper-arid conditions.

DISCUSSION

There are several issues that must be carefully considered before conclusions can be drawn from these data in Fig. 1.

1. The Usefulness of Radioisotope Ages as Relative Ages

It is first necessary to establish that it is still potentially valid to use radioisotope ages in a relative sense, even though it is well established that there are significant problems with the radioisotope dating methods to render the resultant ages as not absolute (Faure and Mensing 2005; Snelling 2000).

All the radioisotope dating methods are ultimately calibrated against the U–Pb and Pb–Pb methods. However, Snelling (2017a) has documented from the literature the residual uncertainties in determinations of the ^{238}U and ^{235}U decay rates, especially the latter, which is somewhat dependent on the determinations of the ^{238}U decay rate. Recently, copious measurements of the $^{238}\text{U}/^{235}\text{U}$ ratio in a very wide variety of rocks, ores, minerals and meteorites has revealed that values vary widely. Indeed, the $^{238}\text{U}/^{235}\text{U}$ ratio value of the same mineral can be very different in different rocks, including zircons, which are so often used in geochronology.

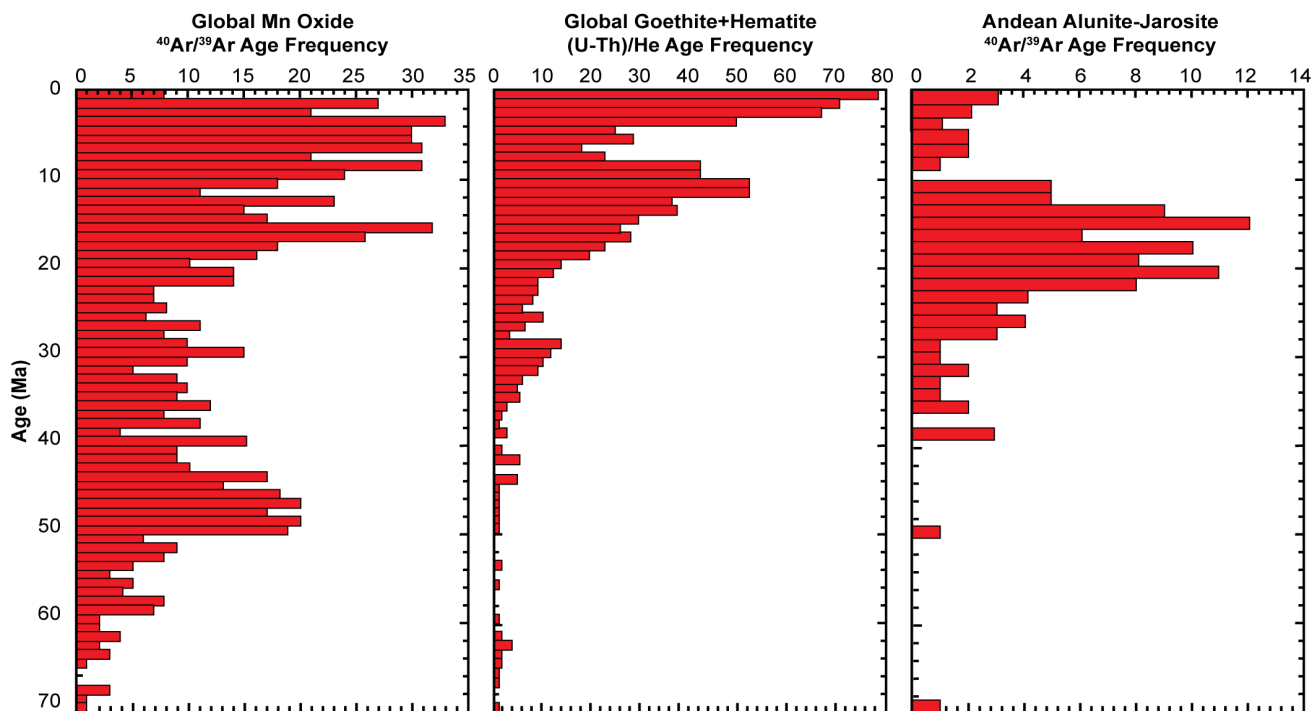


Figure 1. (a) Histogram illustrating the global distribution of $^{40}\text{Ar}/^{39}\text{Ar}$ ages for supergene Mn oxides. (b) Histogram illustrating the distribution of (U-Th)/He ages for the supergene iron oxides goethite + hematite. (c) Histogram illustrating the distribution of $^{40}\text{Ar}/^{39}\text{Ar}$ ages for supergene alunite-jarosite in the Andean region. (After Vasconcelos et al. 2015.)

In any case, it was established by Vardiman et al. (2005) from five lines of evidence (discordant radioisotope ages, helium diffusion, radiohalos, fission tracks, and radiocarbon) that radioisotope decay rates may have been grossly accelerated in a catastrophic event in the recent past (namely, the Flood). Furthermore, it appeared that this accelerated decay was systematic based on the atomic weight, mode of decay, and the present decay rate of the parent radioisotopes. This hinted at the cause being due to changes in the binding forces holding the particles in the nuclei together. The key here is that the acceleration factor was systematic, which means that during the Flood the rocks would have “aged” at consistent rates around the globe as the rocks accumulated.

Thus, in a given area a rock unit that has accumulated a lot of radioisotope decay (for example, a rock unit formed in the first month of the Flood) will yield a very much older radioisotope age than a rock unit that has suffered from much less radioisotope decay (because of forming in the last month of the Flood). Furthermore, if inheritance, the initial conditions and any contamination are roughly the same in all samples from those rock units, then because the different radioisotope parents have been grossly accelerated systematically according to their atomic weights and present half-lives, their apparent model and isochron ages calculated based on the present decay rates of the parent radioisotopes should essentially agree with one another.

Therefore, while the model and isochron ages will be grossly inflated and thus not be absolute ages, they will still be potentially useful as relative ages because they will be in the right order according stratigraphic relationships (Snelling 2010). This assists us in unraveling the time relationships between rock units in a given area when stratigraphic relationships are obscured by lack of

outcrops due to erosion and weathering. However, it can also assist us in correlating the rock units in one area with those in another, even on another continent. After all, the radioisotope contents of rock units are simply chemical characteristics of them in much the same way as their petrologic and mineralogical characteristics, and even their fossil contents. Furthermore, the parent radioisotopes in rock units will have all suffered the same amounts of grossly accelerated decay and thus potentially give the same grossly inflated apparent model and isochron ages which will be the same relative ages.

Baumgardner (2012) has already discussed in detail the usefulness of radioisotope dates as relative ages and similarly argued that accelerated radioisotope decay rates during the Flood would systematically grossly “age” rocks in the order in which they formed in the same way globally. He reviewed the results of the RATE project (Vardiman et al. 2005), emphasizing that the existence of helium in zircons, radiohalos and fission tracks are all evidence that hundreds of millions of years’ worth of radioisotope decay (as calculated from today’s measured decay rates) had occurred. This amount of decay thus needs to be accounted for by grossly accelerated decay during the Flood and perhaps also during the Creation Week. And because this accelerated decay was systematic as rock units were formed progressively during those biblical events of earth history, the resultant radioisotope dates obtained can be useful as relative ages.

In the context of this study it is thus argued that after the Flood waters receded at the same time in various places on different continents, weathering of primary ore deposits would have commenced. Both the receding Flood waters and residual post-Flood catastrophism would have resulted in erosion and exposure

of primary ore deposits to weathering and supergene mineral formation over subsequent decades. The supergene minerals produced by all that weathering should thus have similar or even identical radioisotope ages, which can thus be used as identical relative ages to correlate the progressive onset of weathering at the same time at various stages on different continents in the decades after the Flood ended. This methodology, depending on how the timing of the onset of weathering is interpreted in relation to the timing of the Flood/post-Flood boundary, could help us determine the placement of that boundary in the geologic record.

2. The Integrity and Usefulness of the Geologic Record

There has also been considerable debate about the integrity and usefulness of the so-called geological column, as if the diagram in textbooks does not match the realities of the observable rock record (Reed and Oard 2006). Resolving misunderstandings on this issue is fundamental to building a consistent viable Creation-Flood model that explains the observed details of field data of the literal physical rock record.

Creationist research has demonstrated conclusively that both individual fossil-bearing sedimentary units and megasequences of fossil-bearing sedimentary layers can be traced across continents and even between continents, precisely the evidence expected to be left behind by the global Flood cataclysm (Clarey 2015b, c, 2017; Morton 1984; Snelling 2008, 2009, 2015, 2016b, 2017b). Such transcontinental correlations of rock units underpin the construction of the geological column, ensuring its reality and integrity (Tyler and Coffin 2006). Even Oard (2010a) had to admit that “the geological column is a general Flood order”. Of course, there will always be a few minor exceptions because our knowledge is incomplete, and mistakes can be made by fallible humans, but ongoing field work and research continue to add to this “big picture”. Thus, in this study the integrity and usefulness of the geological column as the geologic record of earth history within the biblical chronology, especially the Flood, is accepted and utilized.

3. The Chronology and Actions of the Retreating Flood Waters

The biblical account suggests that the Flood ended when Noah stepped off the Ark where it had landed on “the mountains of Ararat” months earlier on Day 150 (Genesis 8:4). It is generally agreed that the Flood waters were at their peak on Day 150 of the Flood, even if they had first peaked by Day 40, and they then started subsiding from Day 150 onwards (Anderson 2014; Snelling 2014a). However, the rate at which the waters receded is unclear, though the water level likely dropped slowly at first because it was only some 74 days later on Day 224 that “the tops of the mountains were seen” (Genesis 8:5), though it should also be kept in mind those mountain peaks may have risen above the waters due to the mountain-building tectonic forces of catastrophic plate tectonics (Austin et al. 1994; Snelling 2009, 2014b). And subsequently the text describes the waters being “on the face of the whole earth” (Genesis 8:9). After that, it took another 90 days until Day 314 for the waters to be “dried off the face of the earth” and “the face of the ground” to be dry (Genesis 8:13) (Snelling 2009). Even then Noah waited another 57 days until Day 371 before “the earth had dried out” sufficiently for God to instruct Noah, his family and the animals to leave the Ark, thus ending the Flood event (Genesis

8:14-19) (Snelling 2009).

It should already be evident that these considerations have implications for modeling of the geology, geophysics and paleontology of the Flood event. For example, the peaking of the Flood waters so that all flesh died has implications for when the fossil-bearing strata were deposited globally (Snelling 2014c), while the rising of the mountains has implications for when catastrophic plate collisions and isostatic uplift occurred to form today’s mountain ranges (Snelling 2014b). And the timing and rate of subsiding water levels of the retreating Flood waters and their concurrent actions and motions would have implications for the erosion of the host rocks covering primary ore deposits. If erosion was too severe, not only would the cover rocks be eroded away, but the primary ore deposits as well. Radioisotope dating of the supergene minerals would provide relative dates of their formation.

Another significant stipulation should also be made here. When Noah stepped off the Ark to end the Flood event, even though the earth had dried out does not necessarily mean all residual geological and geophysical processes had ceased. Austin (1998) has documented that volcanic activity has decreased since the Flood, and Austin et al. (1994) postulated decelerating plate movements and collisions into the early post-Flood era. Thus, even ongoing volcanic and seismic activity today can be regarded as a consequence of the Flood. Furthermore, modeling post-Flood hypercanes generated by the warmer ocean waters at the end of the Flood (due to all the hot waters released from the fountains and via volcanism during the Flood), Vardiman (2003, 2012) demonstrated how intense storms would have dumped torrential rainfall on parts of the continents. Whitmore (2013) thus argued there was potential for resulting widespread post-Flood erosion and mass wasting processes. Therefore, while it was safe and dry for Noah to step off the Ark into the area in and near “the mountains of Ararat”, in other parts of the globe residual catastrophism could still have been eroding and depositing strata, and there could have been volcanoes erupting and mountains still rising, with ocean waters still covering continental areas that today are high and dry.

4. The Timing of Weathering and Formation of Supergene Minerals

Before the significance of the timing of weathering to form supergene minerals can be determined, it is crucial to discuss the timing of formation of the primary (hypogene) ore deposits, especially those that were subsequently exposed to chemical weathering.

It should be first noted that not all porphyry copper deposits are of Cenozoic age. Seedorff et al. (2005, Fig. 2, p.255) document that the formation of about a third of the porphyry copper deposits conventionally date back through the Mesozoic and Paleozoic to almost 500 Ma, including in the Andean and Rockies regions. Yet it is only the very latest Mesozoic and Cenozoic porphyry copper deposits that have been subject to supergene oxidation and enrichment (Sillitoe 2005), in spite of the fact that the older porphyry copper deposits are today located within 1000 m of the land surface. This is clearly due to the present arid climate conditions in the Andes and the American Southwest which are conducive to supergene processes, which thus links these supergene minerals to

the development of the current landscape.

The fact is that lateritic deposits formed by supergene weathering acting on primary ore deposits and suitable source rocks are all Cenozoic (Freyssinet et al. 2005). Even though bauxite deposits have formed from and thus sit on top of source rocks as old as the Precambrian (pre-Flood), all mined bauxite deposits are of Cenozoic age and are located in the present global subtropical and tropical belt where the climate was conducive to their formation (Freyssinet et al. 2005, Figs. 1 and 2). Similarly, all lateritic Ni deposits are Cenozoic and yet have formed from source rocks spanning from the Precambrian through the Phanerozoic (Freyssinet et al. 2005, Fig.11).

Other classes of primary ore deposits of all conventional ages also demonstrate that supergene weathering only affected them during the Cenozoic due to present landscape development. Volcanogenic massive sulfide deposits (Franklin et al. 2005), banded iron formations (Clout et al. 2005) and many sediment-hosted Pb-Zn deposits (Leach et al. 2005), of all conventional ages including Precambrian and early Phanerozoic, all formed at the sediments or volcanics surface/water interfaces. Yet even though many of these deposits have distinctive syngenetic or subsequent hydrothermal alteration features, none of these ore deposits have supergene alteration associated with them that is not Cenozoic and related to the current land surface and climate. Similarly, epithermal precious and base metal deposits (Simmons et al. 2005), iron oxide copper-gold deposits (Williams et al. 2005), gold deposits in metamorphic terranes (Goldfarb et al. 2005), and granite-related ore deposits (Černý et al. 2005), which are emplaced underground by hypogene processes, and whose ages span the Phanerozoic and some of the Precambrian; none of these ore deposits have any supergene alteration associated with them that is not Cenozoic and related to the current land surface and climate.

These observations are highly relevant to the objective of this study. In the conventional model it would be expected that supergene ore deposits formed at various times during the long past eons, rather than supergene ore deposits almost only being formed post-Mesozoic and related to Cenozoic to present landscape development and climate processes. Thus, while these observations are not consistent with a long-age model of earth history, they are totally consistent with only a single global Flood model of short earth history. Furthermore, these observations are relevant to their use as a criterion for determining the placement of the Flood/post-Flood boundary in the geologic record.

What is then next relevant to this study is the timing of the erosion that exposed the primary ore deposits to the weathering responsible for the formation of the supergene minerals which have been radioisotope dated. Clearly, when Noah stepped off the Ark marking the end of the Flood event, large-scale erosion would have ceased over large continental regions around the globe, so the corresponding Flood/post-Flood boundary in the geologic record should be marked by a major global-scale erosion surface, onto which residual local deposits would have accumulated during the early post-Flood era. That was the rationale behind the Austin et al. (1994) choice of the Cretaceous/Paleogene (K/Pg) boundary, because they assumed the Cenozoic then represented post-Flood

deposits from residual catastrophism. Localized erosion surfaces coinciding with faunal extinctions would thus have occurred during the post-Flood era. In any case, what is relevant here is that the continuing Cenozoic erosion, which Oard (2004, 2013a, b) agrees with, could have progressively exposed more ore deposits to weathering and supergene mineral formation. This is relevant to many porphyry copper deposits which formed during the Cenozoic mountain-building of the Andes, Rockies, and Alpides (Alpine-Himalayan orogenic belt) and were thus only exposed to weathering by subsequent erosion.

However, since the Flood account states the waters had “dried off the face of the earth” and “the face of the ground” was dry by Day 314 (Genesis 8:13), this could imply erosion had ceased and thus weathering of the primary ore deposits and formation of supergene minerals could have commenced in the 57 days prior to Noah stepping off the Ark and before the Flood event “officially” ended. In that scenario the radioisotope ages obtained for the formation of the supergene minerals would thus not help us date the Flood/post-Flood boundary. However, even if the “face” or surface of the ground was dry does not guarantee that the groundwater table was not still very high and close to the surface. That could be one reason why God left Noah and his cargo still onboard the Ark for those extra 57 days, to allow the water table to drop and soils to form, as well as for plants to grow. The weathering that produces supergene minerals only occurs in the oxidizing zone above the water table (Dill 2015; Reich and Vasconcelos 2015), though it would still be wet due to evaporation of groundwater and further locally-intense rainfall. So not only did the ground surface need to be dry before weathering and supergene mineral formation could commence, but the water table had to drop down below the tops of the primary ore deposits before their oxidization could commence. Thus, the radioisotope ages obtained for the formation of the supergene minerals could still represent the relative timing of the Flood/post-Flood boundary in the geologic record.

However, could weathering and supergene mineral formation commence before the Flood ended, during the drying out phase after the Flood waters retreated? Certainly, weathering would have commenced, but it would have taken subsequent decades rather than just the last 57 days of the Flood to form the supergene minerals. This is because it takes extended time periods for the chemical weathering reactions and the lowering of the water table to penetrate deeply enough to have formed the supergene minerals at the large scale necessary to produce these supergene ore deposits. Thus, the supergene minerals are much more likely to date from the early decades after the Flood ended.

Two other considerations must also be discussed. First, the Flood involved huge volumes of water being trapped in sediments as thick strata sequences were rapidly deposited subaqueously. This water would have been oxygenated from the intense global rainfall, and acidic and warm due to the volcanic water fountains of the “great deep” and depth of burial. Then as those sediments were subsequently compacted under their own weight and due to tectonic forces, it might be expected that these warm acidic waters would rise rapidly through the strata sequences and thus potentially generate look-alike weathering profiles and supergene minerals on their way to the earth’s surface. However, even though we do have

evidence that hot acidic waters rose to the earth's surface and even produced ore deposits (for example, Franklin et al. 2005; Simmons et al. 2005), in the process they produced hydrothermal alteration that has distinctly different minerals and patterns of minerals than found in supergene ore deposits.

Second, could oxygenated surface waters have been forced hydrodynamically downward in the receding waters phase of the Flood and accomplish the same supergene alteration? It is possible that some surface water would penetrate downwards, but as noted above, the chemical weathering that produces supergene minerals only occurs in the oxidizing zone above the water table where it is drier, and air reaches down from the surface (Dill 2015; Reich and Vasconcelos 2015). Besides, as noted above, the chemical reactions that produce the supergene minerals (see the Appendix) take decades to generate ore deposits from the repeated wetting and drying of the chemical weathering profile.

5. The Radioisotope Ages of the Supergene Minerals

In conventional terms, the earliest dates of formation of these supergene minerals is at 70 Ma (Fig. 1). That is just before the K-Pg boundary, currently determined at 66 Ma (Gradstein et al. 2012). And then the frequency in numbers of mineral dates of formation per absolute date increases steadily up through the conventional Cenozoic. This frequency pattern is the same for global Mn oxide $^{40}\text{Ar}/^{39}\text{Ar}$ ages (Fig. 1a), global goethite + hematite (U-Th)/He ages (Fig. 1b), and Andean alunite-jarosite $^{40}\text{Ar}/^{39}\text{Ar}$ ages (Fig. 1c).

However, it is interesting that the dates of formation of alunite and jarosite in the Andes really only start around 40 Ma, and then peak in frequency around 15-20 Ma (Sillitoe and McKee 1996). This is, in part, because many of the primary (hypogene) porphyry copper deposits only formed during the mid-Cenozoic, at the time when the Andes were being built by the active plutonism and volcanism associated with the subduction of the Pacific plate under the South American plate. So, the supergene minerals then only formed subsequently from those deposits after they formed in the mid-Cenozoic. That the chemical weathering responsible for the supergene minerals was connected to the present climate and landscape development was demonstrated conclusively by the U-series dating of supergene assemblages in Chilean porphyry copper deposits by Reich et al. (2009)

It is similarly relevant and significant that the most recent set of goethite (U-Th)/He ages from the Hamersley region of northwestern Australia closely parallel the global goethite + hematite (U-Th)/He ages in Fig. 1b (Miller et al. 2017). The Hamersley province is a passive continental margin with a moderately elevated (~800 m) interior composed of ridges and plateaus dominated by chemically resistant Archean to Paleoproterozoic banded iron formations (BIFs) and quartzites, and incised valleys consisting predominantly of shales and dolomites (Vasconcelos et al. 2013). The region has been structurally stable with little erosion for the entire Cenozoic. Much of the elevated interior is blanketed with ferruginous weathering profiles and related deposits, many of which are exposed and accessible in iron ore mines or prospects.

The prolonged process of iron mobilization and precipitation, along with later cycles of dissolution and reprecipitation of existing iron oxides, has produced a thick horizon rich in goethite with readily

visible and physically separable growth zones that supposedly span many millions of years (Heim et al. 2006; Vasconcelos et al. 2013). For such samples forming at earth-surface conditions, their goethite (U-Th)/He ages are established upon mineral crystallization (Shuster et al., 2005). Goethite acquires oxygen from the water in which it forms, and after that there is no open-system behavior of the oxygen or helium in the goethite (Yapp 1991). Thus, these (U-Th)/He ages date the formation of the goethite at the water table near the surface of this Precambrian terrain (Miller et al. 2017).

Most creation geologists would regard this Precambrian terrain as pre-Flood, and likely even formed during the Creation Week (Snelling 2009). For this terrain to now be exposed in the post-Flood world implies that whatever strata were deposited over it and covered it during the Flood were eroded away by the retreating Flood waters, which also eroded much of its present topography. Thus, the goethite in the present weathering profile had to form after the Flood waters retreated and the ground surface dried out, with weathering and goethite formation commencing as the Flood ended and the water table was subsequently lowered. The goethite (U-Th)/He ages span the Cenozoic from 70.5 Ma to 4.8 Ma, with intermediate ages of 68.0 Ma, 61.7 Ma, 47.3 Ma, 26.0 Ma and 8.5 Ma (Miller et al. 2017). These ages plot within the spread of ages in the histogram in Fig. 1b and are also consistent with the same pattern of $^{40}\text{Ar}/^{39}\text{Ar}$ ages for Mn oxides and alunite-jarosite in Fig. 1a and 1c respectively.

Other examples further substantiate this pattern for the timing of supergene mineral formation. The supergene iron and manganese ore deposits of Minas Gerais, Brazil have developed at the present land surface from early Precambrian banded iron formations and Mn-rich sedimentary units, so the $^{40}\text{Ar}/^{39}\text{Ar}$ ages for the supergene minerals are all Cenozoic (de Oliveira Carmo and Vasconcelos 2006; Spier et al. 2006). Similarly, in Africa supergene manganese deposits have developed at the present land surface on Proterozoic (Precambrian) sedimentary units in the Katanga (De Putter et al. 2015), Kalahari (Gutzmer et al. 2012) and Burkina Faso regions (Colin et al. 2005; Beauvais et al. 2008), and the $^{40}\text{Ar}/^{39}\text{Ar}$ ages for Mn oxides are all Cenozoic, some of them being plotted in Fig. 1a.

In China Mn oxides have developed at the surface on Devonian carbonates (Li et al. 2007), while in India Mn oxides have developed in laterites due to chemical weathering of the underlying Cretaceous Deccan basalts (Beauvais et al. 2016; Bonnet et al. 2014, 2016). In both instances the $^{40}\text{Ar}/^{39}\text{Ar}$ ages for Mn oxides are all younger Cenozoic, indicative of the chemical weathering at the current landscape surface.

In Australia the Permian Mt. Leyshon epithermal gold deposit and its surrounding hydrothermal alteration reach close to the present land surface from where chemical weathering has penetrated down to produce supergene alunite-jarosite group minerals in the host rocks and ore (Scott 1990). K-Ar ages of these supergene minerals are late Cenozoic, similar to $^{40}\text{Ar}/^{39}\text{Ar}$ ages for Mn oxides in the current weathering profile in a nearby part of Australia (Feng and Vasconcelos 2007). Similarly, in Spain the late Paleozoic Las Cruces volcanogenic massive sulfide deposit was exhumed and affected by subaerial oxidation connected to the current landscape (Tornos et al. 2017). $^{40}\text{Ar}/^{39}\text{Ar}$ dating of supergene alunite yielded

various Cenozoic ages, consistent with $^{40}\text{Ar}/^{39}\text{Ar}$ ages of supergene alunite and jarosite associated with the weathering of other similar ore deposits within the Iberian Pyrite Belt.

These results are exactly as anticipated if weathering started after the retreating Flood waters had exhumed much of the current landscape. It is to be expected there would be a range of ages for the formation of supergene minerals, as some would start forming where weathering commenced as the Flood ended, and then further supergene minerals would form later in the early post-Flood decades in areas that experienced residual catastrophism and erosion which delayed the onset of weathering. Thus, the retreating Flood waters would have rapidly eroded off the overburden covering many primary ore deposits, and then as the Flood ended the supergene weathering began. Subsequently, it would have required the extended time in the decades after the Flood for the chemical reactions to produce these supergene minerals from their primary precursors.

The relative ages of formation of all these supergene minerals thus date the beginning of weathering reaching the level where the precursor primary ore minerals occur, but erosion and weathering would have started well before that date. So, in the context of the Flood, as the waters retreated erosion occurred and then weathering could have begun while Noah was still on the Ark. However, there would have been insufficient time in the closing weeks of the Flood for chemical weathering to have generated large enough quantities of supergene minerals for dating. Thus, if the Flood/post-Flood boundary is placed at the K-Pg boundary in the geologic record (a relative date of 66 Ma) (Austin et al. 1994; Whitmore and Garner 2008), then it is not unexpected that only a few of these relative ages for supergene mineral formation would slightly pre-date that boundary (the >66 Ma dates in Fig. 1).

Much caution obviously needs to be exercised in interpreting all these data. Nevertheless, this is a line of investigation that could still potentially yield some demarcation of where the Flood/post-Flood boundary should be placed in the geologic record. Thus, these relative ages of formation of supergene minerals from end-Flood and post-Flood weathering appear to be consistent with a Flood/post-flood boundary low in the Cenozoic, likely as low as the K-Pg boundary many Flood geologists prefer based on many other criteria (Austin et al 1994; Snelling 2009; Whitmore and Garner 2008; Whitmore and Wise 2008).

CONCLUSIONS

The radioisotope dating methods cannot produce absolute ages because of the demonstrated violations of the three assumptions foundational to them. Nevertheless, because many sources of error are systematic, these methods can still produce relative ages, that may then be useful in determining where rocks and minerals fit in the geologic record. It has also been shown that the so-called geological column is likely a reasonable representation of the geologic record of earth history, especially as the global extent of some fossil-bearing sedimentary layers and the megasequences that contain them are powerful evidence of the global Flood cataclysm as described in the Genesis account. Those who dispute the integrity and usefulness of this geologic record use minor exceptions to disparage this “big picture” Flood evidence, and fail

to propose an alternative coherent Flood model consistent with the physical rock record.

When the Flood waters retreated after likely peaking on Day 150, major erosion occurred. By Day 314 the Genesis account says the face of the ground had dried, but Noah waited another 57 days before God instructed him and his cargo to leave the Ark. The ground surface may have been dry, but extra time was needed to allow the water table to drop, soil to form and plants to grow. Thus, the weathering front started to progress downwards, a process that would have been a major shaper of the land surface and topography in the early post-Flood era, in which it required decades for the chemical weathering reactions to produce the supergene minerals.

Primary ore deposits were formed at depth before and during the Flood, but the erosion by the retreating Flood waters stripped away enough of the cover rocks for weathering of their tops to commence after the Flood ended and for supergene minerals to start forming post-Flood. Subsequent residual post-Flood catastrophism may have involved mountains still rising and ore deposits still forming, such as the porphyry copper deposits associated with granite intrusions as the Andes continued to rise. The erosion to expose those later-formed ore deposits and the subsequent weathering to produce supergene minerals from them may thus have occurred well into the early post-Flood era.

The supergene iron oxides, and potassium-bearing sulfates and manganese oxides, produced by weathering of ore deposits can be radioisotope dated to yield the relative ages of their formation. The relative ages obtained span the whole Cenozoic, indicative of progressive weathering and initiation of supergene mineral formation through the Cenozoic. Thus, it is proposed that the relative dates for the first formation of supergene minerals can be used as a helpful criterion for determining the placement of the Flood/post-Flood boundary in the geologic record.

The relative ages of supergene minerals thus favor a placement at the K-Pg boundary with a relative age of 66 Ma, the Flood/post-Flood boundary favored by many Flood geologists. The few slightly earlier relative ages likely resulted from weathering that commenced before Noah stepped off the Ark to end the Flood event, while the spread of relative ages through the Cenozoic represents the progressive formation of supergene minerals as primary ore deposits emplaced during, and some maybe after, the Flood were subsequently exposed to weathering by residual catastrophism. Continuing investigation of this criterion for placement of the Flood/post-Flood boundary seems warranted.

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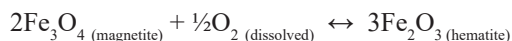
APPENDIX

Supergene Weathering of Ore Deposits

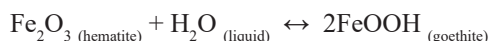
1. Iron Oxides

The oxidation of magnetite (Fe_3O_4) to hematite (Fe_2O_3) and the hydration of magnetite or hematite to goethite (FeOOH) are the product of chemical reactions (identifiable through ore microscopy) within supergene blankets overlying lateritic iron deposits. These chemical reactions are illustrated and labeled with blue numbered arrows in the Eh–pH diagram in Fig. 2 representing the thermodynamically favored pathways, and are:

Reaction (1)



Reaction (2)



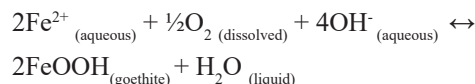
As reaction (1) indicates, the direct oxidation of magnetite to hematite is thermodynamically favored (ΔG^0 reaction = -49.375 kcal) when rocks, which formed under reducing conditions, enter the oxidation zone near the earth's surface. This process can be dated by the (U–Th)/He analysis of supergene hematite. On the other hand, the direct hydration of supergene hematite (martite) to goethite (reaction 2) is not thermodynamically favored (ΔG^0 reaction = $+0.567$ kcal). Yet, goethite formed after hematite is abundant in duricrusts overlying supergene iron deposits. It appears that the hydration of the primary ore oxides involves a two-stage process involving reactions thermodynamically favorable in the supergene zone of ore deposits. First, hematite is reductively dissolved to soluble Fe^{2+} by carboxylic acids (for example, acetic acid, a common organic acid in weathering profiles) (ΔG^0 reaction = -96.99 kcal):

Reaction (2a)



This is followed by the subsequent re-oxidation of Fe^{2+} by dissolved O_2 in weathering solutions and precipitation of goethite (ΔG^0 reaction = -104.286 kcal):

Reaction (2b)



These reaction pathways are shown as labeled blue arrows in Fig. 2. Microscopic evidence also suggests close links between these reactions and microorganisms (Monteiro et al. 2014).

2. Supergene Copper Minerals

Supergene metal deposits contribute significantly to the world's supply of selected base metals (Cu, Zn, Ni, Co) and structural metals (Al, Fe, Ni, V). Giant porphyry copper deposits contribute about 70% of the total global copper inventory and have median grades at ~ 0.4 wt% Cu (with hypogene and supergene Cu grades often being reported together). Yet supergene grades in most deposits are higher than hypogene grades.

In most porphyry copper orebodies, the stepwise hydrolysis and oxidation of primary pyrite-bearing assemblages leads to a decrease in the pH of descending groundwaters and the liberation of oxidized sulfur as SO_4^{2-} anions. There is a simultaneous breakdown of chalcopyrite (CuFeS_2), which produces soluble Cu^{2+} ions that are transported downwards, encountering progressively greater reducing conditions deep into the profile. This process is accompanied by “capping,” the precipitation of iron oxyhydroxides in the leached zone from which Cu is removed. The thickness of the leached cap is highly variable but can reach several hundred meters in porphyry copper deposits, particularly when the water table was deep enough during the supergene oxidation and enrichment phase (Reich and Vasconcelos 2015).

Copper is concentrated within the subjacent oxide zone, which forms laterally extensive deposits composed of assemblages called “green oxides” or “copper oxides”. This mineralogically and compositionally complex layer is composed of copper minerals including oxides, sulfates, hydroxy-chlorides, carbonates, silicates, and native copper. Among the copper minerals that may be encountered are cuprite (Cu_2O), tenorite (CuO), brochantite [$\text{Cu}_4\text{SO}_4(\text{OH})_6$], chalcantite [$\text{CuSO}_4 \cdot 5\text{H}_2\text{O}$], antlerite [$\text{Cu}_3\text{SO}_4(\text{OH})_4$], malachite [$\text{Cu}_2\text{CO}_3(\text{OH})_2$], azurite [$\text{Cu}_3(\text{CO}_3)_2(\text{OH})_2$], atacamite [$\text{Cu}_2\text{Cl}(\text{OH})_3$], turquoise [$\text{CuAl}_6(\text{PO}_4)_4(\text{OH})_8 \cdot 4\text{H}_2\text{O}$], native copper (Cu^0), and chrysocolla [$\text{Cu}_{2-x}\text{Al}_x\text{H}_{2-x}\text{Si}_2\text{O}_5(\text{OH})_4 \cdot n\text{H}_2\text{O}$, among many others.

The precipitation of “green oxide” mineral assemblages in the vadose zone is largely controlled by the enclosing rock type and pH (Fig. 3), forming thick (< 200 – 300 m) layers containing ore with > 1 wt% Cu grade. Furthermore, Cu and other metals dissolved in groundwater can migrate laterally when hydraulic conditions are favorable and form large “exotic-type” Cu-oxide deposits in gravel sequences that are far (distal) from the source.

Under more reducing conditions, the remaining Cu in the

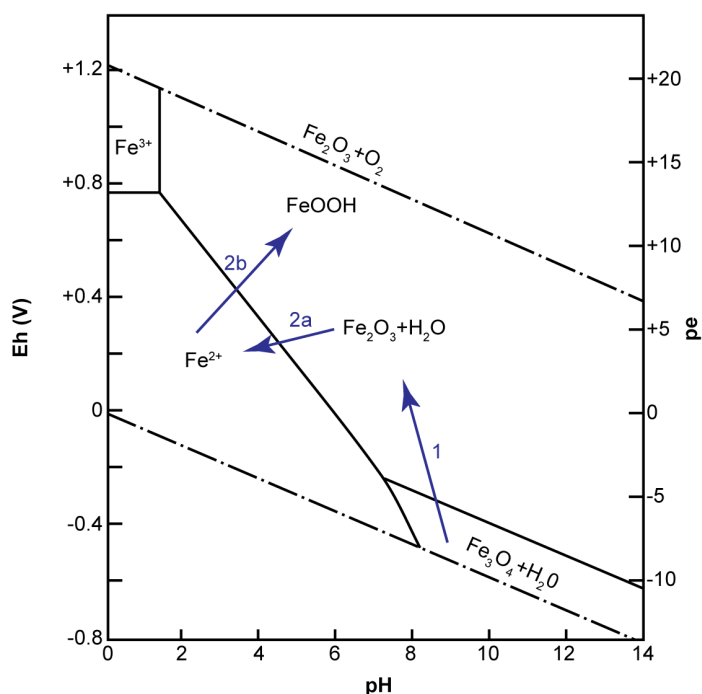


Figure 2. Eh–pH diagram for the Fe–O₂–H₂O system. Blue arrows illustrate the thermodynamically favored pathways for the oxidation (1), dissolution (2a), and precipitation (2b) reactions.

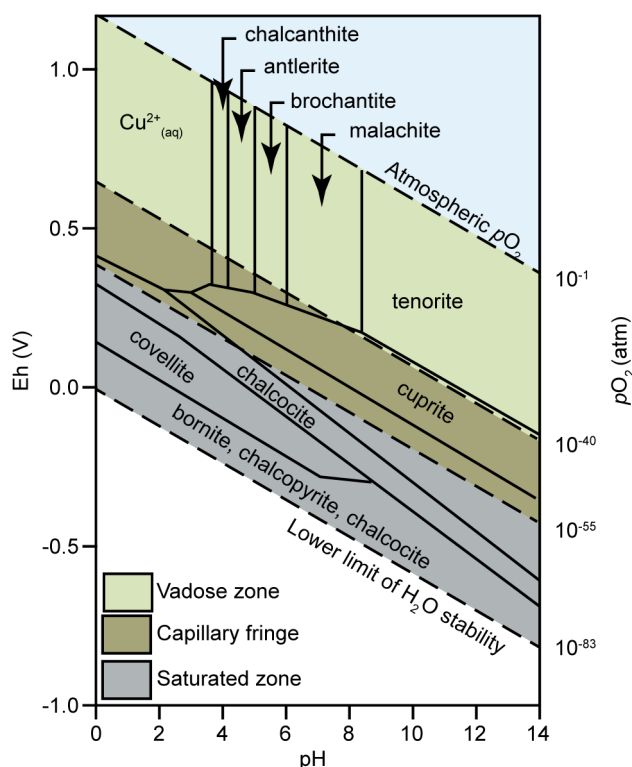


Figure 3. Graph of redox (Eh) and the partial oxygen pressure (pO_2) versus pH showing the stability of dissolved copper species and copper minerals in the supergene environment (after Sillitoe 2005). The diagram maps out possible occurrence of stable phases under particular redox and pH conditions along a supergene profile. Conditions shift vertically from the more reducing, saturated zone in gray at the bottom (below the water table), to the more oxidizing conditions towards the top of the profile (vadose zone, where the soil and rock pores contain air as well as water).

descending metal- and sulfate-rich solutions will form secondary sulfides in the saturated zone below the water table where free oxygen is almost absent ($pO_2 \sim$ below 10–40 atm.) (see Fig. 3). Formation of secondary sulfides occurs by replacement of Fe by Cu in the hypogene sulfides [pyrite (FeS_2), chalcocopyrite ($CuFeS_2$), and bornite (Cu_5FeS_4)]. The secondary chalcocite (Cu_2S) occurs on top, where Cu^{2+}/HS^- is high, while covellite (CuS) precipitates below, where Cu^{2+}/HS^- is lower. Enriched Cu sulfide zones in porphyry copper deposits are usually tens to hundreds of meters thick and can contain more than 1.5 gigatons of ore with 0.4–1.7 wt% Cu, invariably reaching higher Cu grades than those found in the primary (hypogene) orebodies (Reich and Vasconcelos 2015).

3. Supergene Potassium-Bearing Minerals

Alunite [$KAl_3(SO_4)_2(OH)_6$] and jarosite [$KFe_3(OH)_6(SO_4)_2$] are present in weathering profiles encompassing a large range of geological environments. They are common in the oxidation zones of ore deposits, paleosols, silcretes, cave deposits, and weathered river or marine terraces, or deltaic deposits (Vasconcelos 1999a). Supergene alunite and jarosite commonly occur intergrown with primary and other supergene minerals in the weathered zones of orebodies, and as precipitates in veins and cavities (Alpers and Brimhall 1998; Sillitoe and McKee 1996).

The most important parameters controlling the distribution of

alunite-jarosite in natural environments are pH, Eh, (SO_4^{2-}), (K^+), (Fe^{3+}), (Al^{3+}) [where the symbol () denotes activity of the aqueous species], H_2O fugacity, and temperature. In alunite-jarosite systems the chemical reactions which are dated by the K-Ar or $^{40}Ar/^{39}Ar$ methods are the simple precipitation of jarosite or alunite from aqueous solution (equations 1 and 4 in Table 1) or more complex mineral transformations (equations 2, 3, 5, 6, and 7 in Table 1).

During weathering processes, manganese is mobile as the cation Mn^{2+} , which predominates for most of the range of conditions characteristic of natural-water systems (Vasconcelos 1999a). At $pH > 10.5$, the complex $MnOH^+$ becomes the predominant form, and in solutions with high concentrations of HCO_3^- and SO_4^{2-} the complexes $MnHCO_3^+$ and $MnSO_4^{(aqueous)}$ may be important. Organic acids also control manganese solubility in the surficial environment. Mn^{3+} species may occur in strongly acid or organic-rich solutions. However, the tendency for the Mn^{3+} species to disproportionate indicates that it does not play a major role in the solution chemistry of manganese under surface conditions.

The chemical oxidation of Mn^{2+} and Mn^{3+} and the disproportionation of Mn^{3+} in aerated surface water lead to the precipitation of Mn^{4+} oxides. These oxides further catalyze the oxidation process. Thus, Mn oxide precipitation tends to occur on previously deposited oxides, forming accretionary growth bands. Supergene Mn oxides also catalyze the precipitation of other cations from solution (Bi, Ni, Co, Cu, K, Pb, and more), leading to the formation of complex supergene phases (Vasconcelos 1999a). Hollandite [$Ba, K(Mn^{4+}_6Mn^{3+}_2)O_{16}$], coronadite, cryptomelane, birnessite, romanéchite, todorokite, and vernalize are some of the complex Mn oxides commonly found in soils and weathering profiles.

Similar to jarosite-alunite systems, the K-Ar or $^{40}Ar/^{39}Ar$ dating of supergene Mn oxides may determine the direct precipitation of these oxides from aqueous solutions (equation 8 in Table 1) or determine the timing of more complex metasomatic reactions in the weathering environment (equations 9, 10, and 11 in Table 1). When multiple precipitation-dissolution-reprecipitation reactions occur, geochronology of supergene Mn oxides may date the influx of solutions, promoting the dissolution of the previously precipitated Mn oxides, the transport of aqueous Mn^{2+} elsewhere in the system, and the reprecipitation of new generations of Mn oxides in suitable sites. Because the most significant aqueous manganese species in the surficial environment is Mn^{2+} , the partial dissolution of previously precipitated Mn oxides implies the influx of acid and/or reducing solutions (equations 12, 13 and 14 in Table 1).

The aqueous Mn^{2+} species generated by reactions 12, 13 and 14 (Table 1) must be reprecipitated nearby within the weathering profile if these reactions are to be identified and dated. The kinetics of the oxidation of Mn^{2+} aqueous species and the precipitation of Mn^{4+} oxides is generally slow in neutral to slightly acidic oxygenated weathering solutions. In alkaline conditions ($pH > 8$) these reactions proceed more rapidly. The reactions are also catalyzed by bacteria and by mineral surfaces in the natural environment. The identification of the exact reaction controlling the precipitation of datable Mn oxides is desirable if the ages obtained are to be used to interpret paleoclimatic conditions.

Table 1. Representative reactions illustrating the precipitation of supergene K-bearing Mn sulfates and oxides in weathering profiles of ore deposits (after Vasconcelos 1999a, b).

(1)	Direct precipitation of jarosite from solution $\text{K}^+_{(\text{aqueous})} + 3\text{Fe}^{3+}_{(\text{aqueous})} + 2\text{SO}_4^{2-}_{(\text{aqueous})} + 6\text{OH}^-_{(\text{aqueous})} \leftrightarrow \text{KFe}_3(\text{SO}_4)_2(\text{OH})_6_{(\text{jarosite})}$
(2)	K ⁺ and SO ₄ ²⁻ metasomatism of iron oxides $3\text{Fe}(\text{OH})_3_{(\text{solid})} + \text{K}^+_{(\text{aqueous})} + 2\text{SO}_4^{2-}_{(\text{aqueous})} + 3\text{H}^+_{(\text{aqueous})} \leftrightarrow \text{KFe}_3(\text{SO}_4)_2(\text{OH})_6_{(\text{jarosite})} + 3\text{H}_2\text{O}_{(\text{liquid})}$
(3)	Oxidation and K ⁺ metasomatism of pyrite $12\text{FeS}_2_{(\text{solid})} + 4\text{K}^+_{(\text{aqueous})} + 30\text{H}_2\text{O}_{(\text{liquid})} + 45\text{O}_2_{(\text{aqueous})} \leftrightarrow \text{KFe}_3(\text{SO}_4)_2(\text{OH})_6_{(\text{jarosite})} + 16\text{SO}_4^{2-}_{(\text{aqueous})} + 36\text{H}^+_{(\text{aqueous})}$
(4)	Direct precipitation of alunite from solution $\text{K}^+_{(\text{aqueous})} + 3\text{Al}^{3+}_{(\text{aqueous})} + 2\text{SO}_4^{2-}_{(\text{aqueous})} + 6\text{OH}^-_{(\text{aqueous})} \leftrightarrow \text{KAl}_3(\text{SO}_4)_2(\text{OH})_6_{(\text{alunite})}$
(5)	K ⁺ and SO ₄ ²⁻ metasomatism of kaolinite $3\text{Al}_2\text{Si}_2\text{O}_5(\text{OH})_4_{(\text{kaolinite})} + 2\text{K}^+_{(\text{aqueous})} + 4\text{SO}_4^{2-}_{(\text{aqueous})} + 9\text{H}_2\text{O}_{(\text{liquid})} + 6\text{H}^+_{(\text{aqueous})} \leftrightarrow \text{KAl}_3(\text{SO}_4)_2(\text{OH})_6_{(\text{alunite})} + 6\text{H}_4\text{SiO}_4_{(\text{aqueous})}$
(6)	SO ₄ ²⁻ metasomatism of K-feldspar $3\text{KAlSi}_3\text{O}_8_{(\text{K-feldspar})} + 2\text{SO}_4^{2-}_{(\text{aqueous})} + 18\text{H}_2\text{O}_{(\text{liquid})} + 6\text{H}^+_{(\text{aqueous})} \leftrightarrow \text{KAl}_3(\text{SO}_4)_2(\text{OH})_6_{(\text{alunite})} + 9\text{H}_4\text{SiO}_4_{(\text{aqueous})} + 2\text{K}^+_{(\text{aqueous})}$
(7)	SO ₄ ²⁻ metasomatism of muscovite $\text{KAl}_3\text{Si}_3\text{O}_{10}(\text{OH})_2_{(\text{muscovite})} + 2\text{SO}_4^{2-}_{(\text{aqueous})} + 6\text{H}_2\text{O}_{(\text{liquid})} + 4\text{H}^+_{(\text{aqueous})} \leftrightarrow \text{KAl}_3(\text{SO}_4)_2(\text{OH})_6_{(\text{alunite})} + 3\text{H}_4\text{SiO}_4_{(\text{aqueous})}$
(8)	Direct precipitation of cryptomelane from solution $\text{K}^+_{(\text{aqueous})} + 8\text{Mn}^{2+}_{(\text{aqueous})} + 8\text{H}_2\text{O}_{(\text{liquid})} + 4\text{O}_2_{(\text{aqueous})} + \text{OH}^-_{(\text{aqueous})} \leftrightarrow \text{KMn}_8\text{O}_{16}(\text{OH})_{(\text{cryptomelane})}$
(9)	K ⁺ metasomatism of pyrolusite $8\text{MnO}_2_{(\text{pyrolusite})} + \text{K}^+_{(\text{aqueous})} + \text{OH}^-_{(\text{aqueous})} \leftrightarrow \text{KMn}_8\text{O}_{16}(\text{OH})_{(\text{cryptomelane})}$
(10)	Oxidative dissolution and K ⁺ metasomatism of rhodochrosite $8\text{MnCO}_3_{(\text{rhodochrosite})} + \text{K}^+_{(\text{aqueous})} + 4\text{O}_2_{(\text{aqueous})} + 8\text{H}_2\text{O}_{(\text{liquid})} + \text{OH}^-_{(\text{aqueous})} \leftrightarrow \text{KMn}_8\text{O}_{16}(\text{OH})_{(\text{cryptomelane})} + 8\text{H}_2\text{CO}_3_{(\text{aqueous})}$
(11)	Oxidative dissolution and K ⁺ metasomatism of Mn silicates $8\text{MnSiO}_3_{(\text{rhodonite})} + \text{K}^+_{(\text{aqueous})} + 4\text{O}_2_{(\text{aqueous})} + 16\text{H}_2\text{O}_{(\text{liquid})} + \text{OH}^-_{(\text{aqueous})} \leftrightarrow \text{KMn}_8\text{O}_{16}(\text{OH})_{(\text{cryptomelane})} + 8\text{H}_4\text{SiO}_4_{(\text{aqueous})}$
(12)	Reductive dissolution by acidification, followed by direct precipitation of cryptomelane from solution $\text{KMn}_8\text{O}_{16}(\text{OH})_{(\text{cryptomelane})} + 16\text{H}^+_{(\text{aqueous})} \leftrightarrow \text{K}^+_{(\text{aqueous})} + 8\text{Mn}^{2+}_{(\text{aqueous})} + 8\text{H}_2\text{O}_{(\text{liquid})} + 4\text{O}_2_{(\text{aqueous})} + \text{OH}^-_{(\text{aqueous})}$
(13)	Reductive dissolution by organic ligands, followed by direct precipitation of cryptomelane from solution $\text{KMn}_8\text{O}_{16}(\text{OH})_{(\text{cryptomelane})} + 8\text{CH}_2\text{O}_{(\text{aqueous})} + 4\text{O}_2_{(\text{aqueous})} \leftrightarrow \text{K}^+_{(\text{aqueous})} + 8\text{Mn}^{2+}_{(\text{aqueous})} + 8\text{HCO}_3^-_{(\text{aqueous})} + 9\text{OH}^-_{(\text{aqueous})}$
(14)	Reductive dissolution by soluble Fe ²⁺ , followed by direct precipitation of cryptomelane from solution $\text{KMn}_8\text{O}_{16}(\text{OH})_{(\text{cryptomelane})} + 31\text{H}_2\text{O}_{(\text{liquid})} + 16\text{Fe}^{2+}_{(\text{aqueous})} \leftrightarrow 16\text{Fe}(\text{OH})_3_{(\text{ferrhydrite})} + \text{K}^+_{(\text{aqueous})} + 8\text{Mn}^{2+}_{(\text{aqueous})} + 15\text{H}^+_{(\text{aqueous})}$
