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**CONTROL ID:** 720661

**TITLE:** *In Situ*-produced vs. Meteoric  $^{10}\text{Be}$  in Hillslope Soils: One Isotope, Two Tracers, Different Stories

**PRESENTATION TYPE:** Assigned by Committee

**SECTION/FOCUS GROUP:** Earth and Planetary Surface Processes (EP)

**SESSION:** Sediment Supply, Storage, and Delivery as Controlled by Hillslope-Channel Coupling (EP02)

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**ABSTRACT BODY:** *In situ*-produced and meteoric  $^{10}\text{Be}$  are both powerful tools for tracing the production and transport of hillslope sediment. *In situ*-produced  $^{10}\text{Be}$  is used to infer sediment production rates as well as investigate sediment sources and transport. Meteoric  $^{10}\text{Be}$  may also be useful for inferring sediment production and transport rates in some landscapes, especially those that lack the target minerals for *in situ*-produced  $^{10}\text{Be}$ . Few studies have investigated the insights gained by a comparing *in situ*-produced and meteoric  $^{10}\text{Be}$  inventories. We present a series of paired  $^{10}\text{Be}$  inventories from different climatic and tectonic regimes to illustrate both the value and the potential pitfalls of coupling these geomorphic tracers.

The mean *in situ* and meteoric  $^{10}\text{Be}$  near surface (within a meter) inventories for our field areas are as follows: Great Smoky Mountains, NC, USA:  $3.6 \times 10^7$  atoms  $\text{cm}^{-2}$  and  $3.3 \times 10^{10}$  atoms  $\text{cm}^{-2}$ ; Laurely Fork, PA, USA:  $2.6 \times 10^6$  atoms  $\text{cm}^{-2}$  and  $3.0 \times 10^9$  atoms  $\text{cm}^{-2}$ ; Oregon Coast Range, OR, USA: no *in situ* data and  $3.87 \times 10^{10}$  atoms  $\text{cm}^{-2}$ ; North Island, New Zealand: no *in situ* data and  $1.8 \times 10^9$  atoms  $\text{cm}^{-2}$ ; and Amparafaravola, Madagascar:  $1.86 \times 10^7$  atoms  $\text{cm}^{-2}$  and  $8.0 \times 10^9$  atoms  $\text{cm}^{-2}$ . The associated inferred soil residence times, respectively, are: Great Smoky Mountains, NC, USA: 40.9 ky and 25.6 ky; Laurely Fork,

PA, USA: 2.9 ky and 2.3 ky; Oregon Coast Range, OR, USA: *n/a* and 30ky; North Island, New Zealand: *n/a* and 1.5 ky; and Amparafaravola, Madagascar: 21 ky and 6.2 ky. Soil residence times inferred from meteoric  $^{10}\text{Be}$  assume a global average delivery rate of  $1.3 \times 10^6 \text{ atoms cm}^{-2} \text{ yr}^{-1}$ . These soil residence times are minimum values that assume that all *in situ* and meteoric  $^{10}\text{Be}$  is accounted for. Discrepancies between inferred soil residence times most likely highlight some error in assumptions regarding meteoric  $^{10}\text{Be}$  retention in the soil mantles that we sampled. For example, if meteoric  $^{10}\text{Be}$  is not retained at the near surface where we collected our samples, then significant amounts of  $^{10}\text{Be}$  are not being accounted for in our inventory calculations.

If meteoric  $^{10}\text{Be}$  is fully retained by a given landscape, soil residence times inferred from each type of  $^{10}\text{Be}$  should agree. However depth profiles and downslope transects from each field area show differing degrees of meteoric  $^{10}\text{Be}$  mobility. We compare meteoric  $^{10}\text{Be}$  concentrations from each of our field sites to trends in CBD-extractable Al and Fe oxides, bulk soil pH, and mean grain size. Meteoric  $^{10}\text{Be}$  mobility correlates positively to trends in mobile Fe and Al oxides and negatively to soil pH. These data suggest that a meaningful comparison between a landscape's *in situ*-produced and meteoric  $^{10}\text{Be}$  inventories requires a thorough understanding of the geochemistry of the sampled soil mantle.

**INDEX TERMS:** [1826] HYDROLOGY / Geomorphology: hillslope, [1150] GEOCHRONOLOGY / Cosmogenic-nuclide exposure dating.

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