



Dating chert using in-situ produced ^{10}Be : Possible complications revealed on landslide scarps through a comparison with ^{36}Cl applied to coexisting limestone.

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This abstract and presentation highlights potential complications that may arise while using in situ produced ^{10}Be to date diagenetic silica (chert) exposure or burial event. The initiation and evolution of large gravitational collapses in sedimentary rocks were constrained using cosmic ray exposure dating. Because these collapses occurred in a stratigraphic level composed of chert (diagenetic silica) concretions interbedded in limestone layers, their development was studied by performing in situ-produced ^{36}Cl and ^{10}Be concentration measurements in both the limestone and coexisting diagenetic silica (chert), respectively. Following the routinely used decontamination and preparation protocols for ^{10}Be produced in diagenetic silica, large discrepancies were observed with exposure ages determined by ^{36}Cl within carbonate for samples originating from the same scarp. While ^{36}Cl exposure ages were clustered as expected for a unique single gravitational event, ^{10}Be exposure ages were scattered along the same studied scarps. To determine the origin of such a bias, petrological investigations were carried out for chert (diagenetic silica). Thin sections highlighted a complex mineralogical texture characterized by remnant silicified ooids showing calcitic cores, calcite inclusions and a dominant amorphous hydrated silica (grain $> 20 \mu\text{m}$). To decipher and characterize the potential origins of the excess measured ^{10}Be within diagenetic silica, all samples were first reprocessed following the routine decontamination protocol (HCL- H_2SiF_6 leachings and three partial HF dissolutions) but starting from three different grain size fractions (GS1: 1000-500, GS2: 500-250 and GS3: 250-50 μm). The resulting concentrations clearly showed a decreasing ^{10}Be content as a function of the grain size, but still yielded ^{10}Be exposure ages significantly higher than ^{36}Cl counterparts. Because potential adsorption of ^{10}Be at the surface of amorphous silica grains was suspected, partial dissolution steps following by a leaching step in hydroxylamine were investigated. Finally, it seems that an additional leaching in KOH allowed removal of the amorphous silica phase and the measured ^{10}Be concentrations to yield ^{10}Be exposure ages agreeing within uncertainties with the ^{36}Cl ones. This work suggests that measuring in situ produced ^{10}Be within chert (amongst other types of diagenetic silica, e.g. flint, hornstone, jasper, etc.) containing amorphous silica requires caution.