

Mass-transfer in solution: on the context and cause of spherulitic lacustrine carbonate deposits

Michael Rogerson (1), Ramon Mercedes-Martin (1), Alex Brasier (2), Hubert Vonhof (3), Tim Prior (1), Simon Fellows (1), John Reijmer (3), Martyn Pedley (1), and Rona McGill (4) (1) University of Hull, Hull, UK, (2) University of Aberdeen, Abderdeen, UK, (3) Vrije Universiteit Amsterdam,

Amsterdam, Netherlands, (4) SUERC, East Kilbride, UK

Understanding how laterally extensive spherulitic carbonate deposits originated in the 'Pre-Salt' Aptian volcanicinfluenced lacustrine basins of Brazil and Angola is challenging, especially as few good analog systems are known to compare the Pre-Salt to. Here, we report a particularly good analog system in the Carboniferous of the Scotland (the East Kirkton Limestone), and examine the geochemical and sedimentological context in which spherular radial calcite has developed. Using empirical and theoretical approaches, we analyse the link between metal mobilization from sub-surface volcaniclastic rocks, and the potential precipitation of carbonates, Mg-Si minerals and chalcedony in a lacustrine spherulitic carbonate setting. This suite of minerals at the surface can be explained by CO₂ ingassing to a springwater derived from reaction of alkali igneous rocks in the subsurface with meteoric groundwater. This forms a 'source-to-sink' system occurring entirely in solution in the subsurface, but demonstrably capable of forming significant depositional units at the surface. We hypothesise that analogous processes occurring on a larger scale are implicated in the development of the 'Pre-Salt' spherulitic carbonate deposits.