

Production and Destruction of Novel Organic Species during simulated Hydrothermal Alteration of Seafloor Sediment

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Hydrothermal alteration of sedimentary organic matter is a fundamental process generating dissolved organic molecules in sediment-influenced hydrothermal ecosystems. Production of organic species other than simple hydrocarbons and some organic acids [1,2] remains largely unconstrained, however. To address this, we conducted two time-series experiments using a Au-TiO₂ apparatus to investigate NSO-production during hydrothermal seawater-sediment interactions at 250–360°C (35 MPa). The experiments heated two differing sediment types with similar wt.% organic carbon over several months - Arctic Mid-Ocean Ridge terrigenous sediment (KNIP) and Guaymas Basin diatomaceous ooze (GUAY). Both were progressively heated to 250°C and 330°C at similar seawater/sediment ratios, with GUAY heated further to 360°C while KNIP was cooled to 250°C, simulating conductive cooling of an organic-rich fluid.

Time-series measurements showed major element, gas and dissolved organic carbon (DOC) concentrations generally followed trends observed in previous studies at comparable conditions [1,2]. We observed remarkably similar trends for NSOs in both experiments. CH₃SH rapidly increased to mmolar concentrations at 250°C, vastly outpacing CH₄ production in the earliest stages. CH₃SH concentrations then decreased at 330–360°C, but increased again slightly in the cooled 250°C KNIP experiment. This implies CH₃SH production exceeds destruction at moderate temperatures over several months - a trend reversed at higher temperatures. C₂–C₅ carboxylic acid concentrations behaved similarly, rapidly increasing at 250°C and 330°C, then decreasing at 360°C (GUAY), and stabilizing in the cooled 250°C phase (KNIP). Formate abundance was temperature dependent, evidenced by lower, but identical, abundances in both 250°C phases in KNIP relative to the higher concentration 330°C phase. Despite bulk DOC and total dissolved nitrogen concentrations reaching 10s of mM at 250°C, solid-phase extractable DOC decreased rapidly with increased temperature in both experiments. Ongoing analysis of alcohols, ΣNH₄⁺, amino acids, structural characterization of DOC and mineral alteration products will also be presented. Our results demonstrate complex

but common trends for diverse NSO-organics during hydrothermal sediment alteration, with important implications for how these species respond to varying thermal maturation extents and timescales in natural systems.

[1] Seewald *et al.* (1990) *Applied Geochemistry*, 5(1-2), 193-209

[2] Kawagucci and Seewald (2019) *Geochemical Journal*, 53(4), 281-291