Redox-sensitive Element Uptake at the North-East Atlantic Benthic Boundary Layer Experiment Sites

John Thomson (jth@soc.soton.ac.uk)¹, Sharon Nixon¹, Ian Croudace¹, Tom Pedersen², Louise Brown³, Gordon Cook³ & Angus MacKenzie³

¹ Sothampton Oceanography Centre, Empress Dock, Southampton SO14 3ZH, UK

² Dept. Earth & Ocean Sciences, University of British Columbia, Vancouver V6T 1Z4, Canada

³ Scottish Universities Research & Reactor Centre, Scottish Enterprise Technology Park, East Kilbride G75 0QF, UK

At two of the three Benthic Boundary Layer Experiment sites in the north-east Atlantic, regularly-increasing radiocarbon age/depth profiles demonstrate that sediment accumulation has been relatively constant during the late Holocene, at rates of 4.4 cm.ky⁻¹ (3.1 g.cm⁻².ky⁻¹) at site B on Rockall Bank and 6.5 cm.ky⁻¹ (4.4 g.cm⁻².ky⁻¹) at site C on Feni Drift. The composition of the sediments is near-constant with depth at both sites, although they are quite distinct from each other (mean $CaCO_3 =$ 79.6% at site B and 54.0% at site C) as a result of contrasting particle size spectra. Under these two different but quasi steadystate depositional regimes, the geochemical behaviour of the redox-sensitive elements Cd, Mn, Mo, Re, Se and U in response to early diagenesis is readily discerned through down-core changes in elemental concentrations. Collectively, a colour change in the sediments, surficial MnOx enriched layers and ²¹⁰Pb excess profiles allow an estimate of the mean position of the oxic/post-oxic boundary in the sediments at both sites. This boundary is situated deeper than the ²¹⁰Pbexcess surface mixed layer and shallower than the ¹⁴C surface mixed layer which represent mixing on 102- and 103-year time scales, respectively. This implies that the well-defined surface MnOx enrichment and its associated ²²⁶Ra and Mo fractions must have been efficiently recycled by downwards bioturbative mixing into anoxic conditions, subsequent reduction and diffusive migration back up into oxic conditions. This process must have last occurred in <40 years at site C and in <103 years at site B. Authigenic enrichment of Se, Cd and U begins to increase immediately below the oxic/post-oxic boundary, although Re enrichment does not occur until a few cm deeper. The mean authigenic uptake enrichments of Se, Cd, U and Re are 0.15, >0.35, 0.4 and >0.0034 $\mu g.g^{g-1}$ at site B and 0.75, >0.27, 0.98 and >0.0051 $\mu g.g^{g-1}$ at site C.