



Development and Application of Compound-Specific Chlorine Stable Isotope Analysis of 1,2-Dichloroethane

Orfan Shouakar-Stash (1), Ramon Aravena (2), Daniel Hunkeler (3), and Stephanie Fiorenza (4)

(1) Department of Earth and Environmental Sciences, University of Waterloo, Waterloo, Ontario, Canada, (orfan@uwaterloo.ca / Fax: 1-519-746-7484), (2) Department of Earth and Environmental Sciences, University of Waterloo, Waterloo, Ontario, Canada, (roaraven@uwaterloo.ca / Fax: 1-519-746-7484), (3) Center for Hydrogeology, University of Neuchâtel, Neuchâtel, Switzerland, (Daniel.Hunkeler@unine.ch), (4) Remediation Engineering & Technology, BP America, Houston, TX, USA

1,2-Dichloroethane (1,2-DCA) has been produced in larger quantities than any other chlorinated hydrocarbons, and is frequently detected in groundwater and has been classified as a priority pollutant by the USEPA. The dual isotope approach (^{13}C and ^{37}Cl) has shown a great potential for evaluating sources and processes that affect chlorinated ethenes in groundwater. In order to extend the application of the dual isotope approach in the case of 1,2-DCA, a new methodology was developed for determining compound-specific chlorine stable isotopes ($\delta^{37}\text{Cl}$) of 1,2-dichloroethane (1,2-DCA). The analytical methodology is carried out on a Continuous Flow - Isotope Ratio Mass Spectrometry (CF-IRMS). The technique depends on the measurement of fragments m/z 62 and m/z 64 of 1,2-DCA. The accuracy of the analytical technique was validated by conducting an off-line analysis technique. The analytical quantification limit of the methodology is better than $30 \mu\text{g/L}$ and the reproducibility is better than 0.15% . The extraction and separation of 1,2-DCA was conducted by a Solid Phase Micro Extraction (SPME) and Gas Chromatography (GC). The analytical methodology was tested in field samples collected at a site where 1,2-DCA is one of the main contaminants.