1.006 Modelling urban δ 13C variations in the Greater Toronto Area.

Early Career Scientist

Presenting Author:

Stephanie Pugliese, Department of Chemistry, University of Toronto, Canada, stephanie.pugliese@utoronto.ca

Co-Authors:

Felix Vogel, Laboratoire des Sciences du Climat et de L'Environnement, CEA-CNRS-UVSQ, Université de Paris-Saclay, France

Jennifer G. Murphy, Department of Chemistry, University of Toronto, Canada Junhua Zhang, Environment Canada Toronto, Toronto, Ontario, Canada Qiong Zheng, Environment Canada Toronto, Toronto, Ontario, Canada Michael D. Moran, Environment Canada Toronto, Toronto, Ontario, Canada Elton Chan, Environment Canada Toronto, Toronto, Ontario, Canada Douglas Worthy, Environment Canada Toronto, Toronto, Ontario, Canada Craig A. Stroud, Environment Canada Toronto, Toronto, Ontario, Canada Shuzhan Ren, Environment Canada Toronto, Toronto, Ontario, Canada

Abstract:

Even in urbanized regions, carbon dioxide (CO₂) emissions are derived from a variety of biogenic and anthropogenic sources and are influenced by atmospheric transport across borders. As policies are introduced to reduce the emissions of CO₂, there is a need for independent verification of emissions reporting. In this work, we use carbon isotope (^{13}CO) $_{\rm 2}$ and $^{\rm 12}{\rm CO}_{\rm 2}{\rm)}$ simulations in combination with atmospheric measurements to distinguish between CO_2 sources in the Greater Toronto Area (GTA), Canada. This is being done by developing an urban $\delta^{13}C$ framework based on existing CO $_2$ emissions data and forward modelling. We first developed a CO_2 inventory for the GTA at a very fine spatial and temporal resolution $(0.02^{\circ} \times 0.02^{\circ})$ and hourly, respectively). We compared the outputs from the Lagrangian dispersion model FLEXPART (at $0.1^{\circ} \times 0.1^{\circ}$) and the chemistry transport model GEM-MACH (at $0.02^{\circ} \times 0.02^{\circ}$) to evaluate the impact of model resolution on the produced simulations. These model results are used in our framework in combination with region-specific δ^{13} C signatures of the dominant CO₂ sources; the product is compared against highly accurate 13 CO₂ and 12 CO₂ ambient data. Locally, anthropogenic CO-2 in urban areas is often derived from natural gas combustion (for heating) and gasoline/diesel combustion (for transportation); the isotopic signatures of these processes were measured to be significantly different (approximately $\delta 13C_{VPDB} = -$ 44 [] and -29 [] respectively) in the GTA and can be used to infer their relative contributions. Utilizing our δ^{13} C framework and differences in sectoral isotopic signatures, we quantify the relative contribution of CO₂ sources on the overall measured concentration and assess the ability of this framework as a tool for tracing the evolution of sector-specific emissions.