



## **Radiative Forcing from North American NO<sub>x</sub> Emissions: dependence upon location and season of emission**

Richard Damoah (1), David Stevenson (1) and Dick Derwent (2)

(1) Institute for Atmospheric and Environmental Science, The University of Edinburgh, UK  
(rdamoah@staffmail.ed.ac.uk), (2) Rdsscientific, Newbury, Berkshire, UK

The 3D tropospheric Chemistry Transport Model (CTM) STOCHEM has been used to study the changes in the distribution of methane CH<sub>4</sub> and ozone O<sub>3</sub> due to NO<sub>x</sub> emissions from a variety of sites in N. America. One month long NO<sub>x</sub> emission pulses resulted in: (1) CH<sub>4</sub> deficits that lead to negative radiative forcings; these decay with e-folding times of 10-15 years; (2) excess O<sub>3</sub> mixing ratio (positive radiative forcing) in the short-term that decays over a few months to produce deficits; and (3) long-term O<sub>3</sub> deficits (negative radiative forcing) which decay away in line with the CH<sub>4</sub> deficits. The total time-integrated net radiative forcing from negative CH<sub>4</sub> and long-term O<sub>3</sub> contributions, and positive short-term O<sub>3</sub> leave a small negative residual. However, the net radiative forcing from NO<sub>x</sub> emission pulses varies strongly with the season and location of the emissions.