

Inversion of permafrost methane emissions using TM5-MP/4DVAR with TROPOMI measurements

SANTIAGO PARRAGUEZ¹, NIKOS DASKALAKIS¹,
MARIA KANAKIDOU^{1,2,3}, MIHALIS VREKOUSSIS^{1,4,5},
ARJO SEGERS⁶, OLIVER SCHNEISING¹ AND MICHAEL
BUCHWITZ¹

¹Institute of Environmental Physics (IUP), University of Bremen

²Environmental Chemical Processes Laboratory (ECPL),
University of Crete

³Center for Studies of Air Quality and Climate Change,
Foundation for Research and Technology Hellas

⁴Center of Marine Environmental Sciences (MARUM),
University of Bremen

⁵Climate and Atmosphere Research Center (CARE-C), The
Cyprus Institute

⁶Royal Netherlands Meteorological Institute

Presenting Author: sanparra@uni-bremen.de

Methane is a significant greenhouse gas (GHG) with a global warming potential (GWP) of 32 for the 100-year horizon. Its global concentration has increased by more than 2.6 times since the pre-industrial era, primarily due to anthropogenic activities, making it a major contributor to global warming. Since 2007, atmospheric CH₄ concentrations have been rising at a faster rate, reflecting a greater imbalance between its sources and sinks. As such, accurately assessing the sources of methane emissions is important to mitigate the consequences of climate change. Our current understanding of CH₄ sources is rather limited, resulting in uncertainties in both bottom-up and top-down estimation methods, along with inconsistencies between these results. Therefore, it is imperative to incorporate new data with high confidence levels into assimilation systems that estimate emission sources. The Arctic, which warms twice as fast as the global average and is highly sensitive to temperature changes, is a region of particular concern. There, a significant portion of the world's soil organic carbon is stored, mostly in permanently frozen peat. As temperatures increase, permafrost thaws faster, releasing stored carbon as CO₂ and CH₄ and creating a positive feedback loop on climate change.

Our study aims to assess the potential contribution of thawing permafrost to the recent increase in global CH₄ concentrations. To accomplish this, we estimate global CH₄ emissions at a spatial resolution of 1°×1° using the inverse modelling system TM5-MP/4DVAR. This framework is based on the TM5-MP atmospheric chemistry-transport model and its adjoint, the four-dimensional variational (4DVAR) data assimilation. The data assimilated in this work comprises tropospheric CH₄ column density measurements obtained from the TROPOMI instrument onboard the Sentinel 5-Precursor satellite, retrieved with the weighted function modified differential optical absorption spectroscopy (WFMD-IUP) algorithm, as well as near-surface CH₄ observations from the stations of NOAA network.