

## **Evaluation of surface ozone chemistry in regulatory air quality models through the use of OMI observations and aircraft data**

Tim Canty (1), Allison Ring (1), Margaret Marvin (2), Sandra Roberts (2), Linda Hembeck (1), Glenn Wolfe (3), Ross Salawitch (1), and Russell Dickerson (1)

(1) University of Maryland, College Park, Dept. of Atmospheric and Oceanic Science, College Park, United States, (2) University of Maryland, College Park, Dept. of Chemistry, College Park, United States, (3) NASA Goddard Space Flight Institute/ UMBC Joint Center for Earth Systems Technology, Greenbelt, United States

Determining the chemical conditions that lead to the formation of surface ozone, a pollutant often regulated by federal governments, is of great interest to scientists and policy makers. The lack of detailed observations of ozone precursors, such as  $NO_x$  and volatile organic compounds (VOCs), limits our ability to precisely identify the nonlinear chemistry that determines ozone production over large regions of the United States. Earlier studies have used the ratio of satellite observations of column HCHO, a VOC proxy, to column  $NO_2$  to estimate ozone sensitivity. Others use changes in average maximum 8hr surface ozone to determine ozone production regime. Recent work has highlighted the regional and temporal variations of the HCHO/ $NO_2$  ratio that must be accounted for. This analysis focuses on the ratio of column HCHO/ $NO_2$  as observed from space and the representation of surface ozone chemistry in regulatory air quality models using a number of assumptions regarding the model framework.