

Measurements of Ambient Mercury and Related Species at the Mauna Loa Observatory 2002-2008

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EPA ORD and NOAA ESRL initiated measurements of gaseous elemental mercury (Hg^0), divalent reactive gaseous mercury (RGM), and particulate bound mercury ($\text{Hg}(\text{p})$) in 2002 at the Mauna Loa Observatory (MLO). Collocated elemental carbon, O_3 and SO_2 measurements were subsequently initiated in 2004. NOAA ESRL O_3 data was used to complete our data set from 2002 - 2004. Hg^0 concentrations ranged between 0.3 and 2.9 ng m^{-3} , and average \pm standard deviation was $1.6 \pm 0.5 \text{ ng m}^{-3}$; $\text{Hg}(\text{p})$ concentrations ranged between 1 - 1900 pg m^{-3} , and RGM ranged between 0.6 - 360 pg m^{-3} . Periods when $\text{Hg}(\text{p})$ is anti-correlated to Hg^0 are also periods when ozone is anti-correlated to Hg^0 (2002 - 2003), suggesting that air masses sourcing ozone differ from that of Hg^0 , possibly of stratospheric origin. The majority of SO_2 impacting Mauna Loa is expected to be of volcanic origin. SO_2 correlation with Hg^0 suggests that we observed significant Hg^0 during periods of volcanic activity, while anti-correlation periods indicate long distance transport influences for Hg^0 in contrast to the more regional volcanic influences for SO_2 . Elemental carbon is a good indicator of anthropogenic sources and is mostly correlated to Hg^0 , suggesting that most Hg^0 observed at MLO is of anthropogenic origin. This study is still in early stages of data analysis and validation, a complete presentation of the data and detailed analysis will be presented.

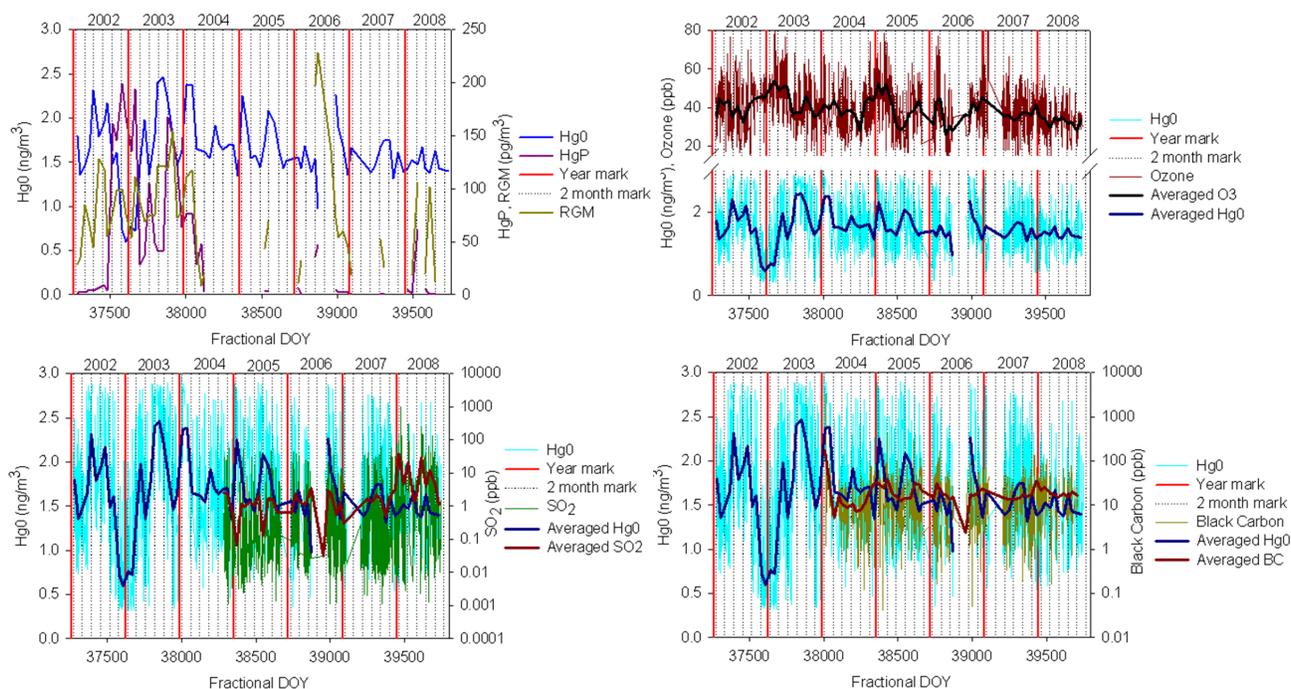


Figure 1. Hg^0 , HgP , RGM, ozone, SO_2 and black carbon concentration trends in Mauna Loa, Hawaii in 2002 – 2008. The solid thick lines represent the monthly average and the lighter solid line is the 3 hour resolution measurement for the chemical species.