

Supplementary Information

Latitudinal distribution of reactive iodine in the Eastern Pacific and its link to open ocean sources

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IO mixing ratios for selected clear days were retrieved from the Multi-Axis Differential Optical Absorption Spectroscopy (MAX-DOAS) Differential Slant Column Densities (DSCDs) in two steps. First, O₄ DSCDs were forward modeled using the NIMO fully spherical Monte Carlo radiative transfer model (Hay et al., 2012) by prescribing aerosol profiles with varying aerosol optical depths and shapes. The surface albedo was set to 0.07 and the Henyey-Greenstein parameterization for aerosol scattering was used with an asymmetry parameter of 0.75, appropriate for sea salt aerosols, and a single scattering albedo of 0.97. Since the vertical distribution of O₄ in the atmosphere is known and decreases in proportion to the square of pressure, O₄ DSCDs are a good proxy for the average effective path lengths of observed photons in the boundary layer for the different viewing geometries (Wagner, 2004). Second, the aerosol profile that resulted in the best fit of the modeled O₄ DSCDs to the measurements was prescribed in the forward model calculation of weighting functions that characterize the sensitivity of the DSCDs to changes in the trace gas concentrations in different altitude layers. A linear maximum *a posteriori* (MAP) inversion (Rodgers, 2000) of the IO DSCDs was performed using the weighting functions to obtain vertical profile information. The MAP inversion is essentially a least squares fit of the model to the measurements using the weighting functions weighted by the measurement and *a priori* errors. An *a priori* profile is required as a constraint as the measurements alone do not contain sufficient information to arrive at

a single solution. A linearly decreasing *a priori* profile was used, based on previous measurements and chemical modeling, with a layer grid height of 50 meters from the surface to four kilometers. The *a priori* error for each layer was set to 80% of the peak *a priori* value in order to minimize the RMS of the model fit to the measurements while still providing some constraint on the profile shape. This percentage was chosen using the L-curve method described in Schofield (2003). In most cases, an aerosol profile could be found which resulted in a reasonable fit of the modeled O₄ DSCDs to the measurements. However, IO was often below the detection limit for higher elevation angles and the low information content in the DSCDs meant that the absolute errors in the retrieved IO mixing ratios were never below 0.3 pptv.

Therefore, MBL mixing ratios were also obtained using the O₄ DSCDs for the 1° elevation angle to derive effective path lengths. A similar technique has been used by other groups (Wagenr et al., 2004; Sinreich et al., 2010). The O₄ DSCDs were divided by the mean extinction coefficient of O₄ from the surface to 200 m a.s.l. to obtain the path lengths. This layer height was based on the average last scatter altitude calculated with NIMO using the different aerosol profiles determined by forward modeling. A wavelength correction, calculated with the RT model, was applied to the path length since the O₄ spectral analysis was performed on an absorption band at 360 nm whereas the IO analysis was centered on 427 nm. IO mixing ratios were then obtained by dividing the IO DSCDs by these corrected path lengths. This method is less sensitive than the inversion to the effect of clouds since the last scattering altitude for the 1° elevation angle is below the cloud base altitude (the spectra were filtered to remove much of the cloudy data) and most of the O₄ absorption can be assumed to occur on the extended path through this low layer. However, this technique relies on the assumption that the IO profile in the boundary layer is similar to the O₄ profile.

The IO mixing ratios calculated with this method were in agreement within the errors of the values calculated by the MAP optimal estimation technique [where IO was above the detection limit for all elevation angles \(Figure S2\)](#). The errors were derived from the DOAS fitting errors in the O₄ and IO DSCDs, combined with small errors in the mean O₄ extinction coefficient and the air density due to the uncertainty of the layer height. Further errors are likely to be introduced by the assumption that the IO layer has a constant mixing ratio up to the last scatter altitude and that the entire differential O₄ absorption, relative to the zenith sky viewing direction, occurs in the line of sight direction. Hence the errors on the O₄ method are most likely underestimated. The O₄ method retrieved mixing ratios, validated using the MAP inversion, were used for further analysis.

References:

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Wagner, T., B. Dix, C. v. Friedeburg, U. Friß, S. Sanghavi, R. Sinreich, and U. Platt (2004), MAX-DOAS O₄ measurements: A new technique to derive information on atmospheric aerosols—Principles and information content, *J. Geophys. Res.*, 109, D22205, doi:10.1029/2004JD004904.

Figures:

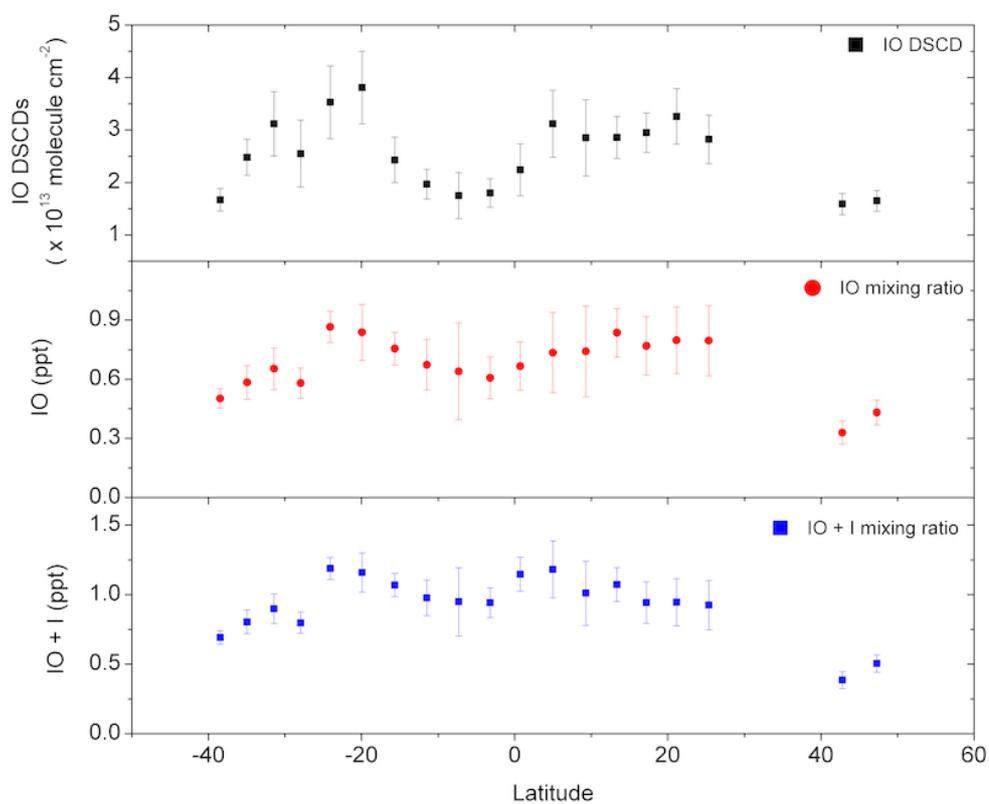


Figure S1: The latitudinal distribution of IO DSCDs (top panel), the calculated IO mixing ratios using the method described above (middle panel) and the total IO_x (= I + IO) mixing ratio estimated using the THAMO model (bottom panel).

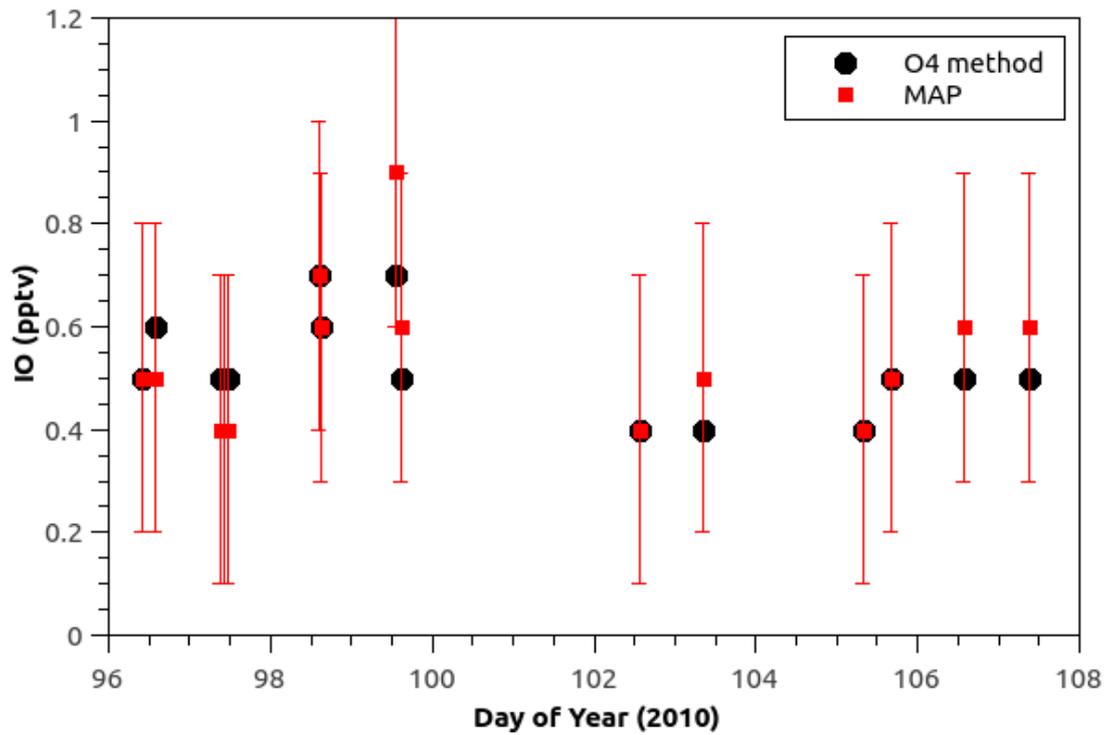


Figure S2: Comparison of retrieved IO mixing ratios using the O₄ method with those obtained using MAP optimal estimation. For clarity only the errors from the MAP method are shown. However, since the IO DSCDs were often below the detection limit for higher elevation angles, the low information content in the DSCDs meant that the absolute errors in the retrieved IO mixing ratios were the same as prescribed in the a priori (0.3 pptv), which is why the O₄ method is used for further analysis.