



Effects of the 2006 El Niño on tropospheric composition as revealed by data from the Tropospheric Emission Spectrometer (TES)

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[1] The Tropospheric Emission Spectrometer (TES) is unique in providing multi-year coincident tropospheric profiles of CO, O₃ and H₂O. TES data show large differences in these gases over Indonesia and the eastern Indian Ocean in October–December 2006 relative to 2005. In 2006, O₃ was higher by 15–30 ppb (30–75%) while CO was higher by >80 ppb in October and November, and by ~25 ppb in December. These differences were caused by high fire emissions from Indonesia in 2006 associated with the lowest rainfall since 1997, reduced convection during the moderate El Niño, and reduced photochemical loss because of lower H₂O. The persistence of the O₃ difference into December is consistent with higher NO_x emissions from lightning in 2006. TES CO and O₃ enhancements in 2006 were larger than those observed during the weak El Niño of 2004. **Citation:** Logan, J. A., I. Megretskaya, R. Nassar, L. T. Murray, L. Zhang, K. W. Bowman, H. M. Worden, and M. Luo (2008), Effects of the 2006 El Niño on tropospheric composition as revealed by data from the Tropospheric Emission Spectrometer (TES), *Geophys. Res. Lett.*, 35, L03816, doi:10.1029/2007GL031698.

1. Introduction

[2] El Niño Southern Oscillation (ENSO) is the most important mode of interannual variability in the tropical atmosphere, and has a strong influence on the distribution of tropospheric O₃ in the tropics. During an El Niño event, the normally warm waters and associated convection over the western Pacific and maritime continent move towards the eastern Pacific. As a result of the changes in large scale circulation and convection, O₃ increases over the maritime continent and decreases over the central Pacific [e.g., Ziemke and Chandra, 2003]. The intense El Niño in late 1997 and the associated drought led to major forest fires in Indonesia, with increases in the tropospheric column of O₃ (TCO) of 40–75% in the region [Chandra *et al.*, 1998; Kita *et al.*, 2000; Thompson *et al.*, 2001]. Model studies showed that about half of the increase in O₃ was caused by changes in dynamics and the remainder by emissions of O₃ precursors

from the fires [Sudo and Takahashi, 2001; Chandra *et al.*, 2002].

[3] Chandra *et al.* [2007] reported on the effects of the weak El Niño in late 2004 on tropospheric O₃ and H₂O. They derived data for the TCO from the Ozone Monitoring Instrument (OMI) and the Microwave Limb Sounder (MLS) on board the Aura satellite, and used MLS data for H₂O at 215 hPa. They found that the TCO increased by 10–20% over the maritime continent and decreased by a similar amount over the eastern Pacific, while H₂O showed similar changes of opposite sign.

[4] There was a moderate El Niño in late 2006. Sea surface temperature anomalies in the Niño 3.4 region exceeding 0.5°C for 3-month running means during 5 consecutive seasons are considered warm events (El Niño conditions). Anomalies were 0.9°, 1.1°, and 1.1° for the last three months of 2006, compared to 0.9°, 0.9°, and 0.8° for these months in 2004 (www.cpc.noaa.gov/products/analysis_monitoring/ensostuff/ensoyears.shtml). Conditions were neutral in 2005.

[5] There were large fires in Indonesia between August and November of 2006 with estimated emissions of 82 Tg CO, much higher than emissions during the same months in 2004 and 2005, 24 Tg and 14 Tg CO respectively, but much lower than emissions from the fires in 1997, 193 Tg CO [van der Werf *et al.*, 2006] (<http://ess1.ess.uci.edu/~jranders>).

[6] We report here on the significant perturbations to O₃, CO, and H₂O from the 2006 Indonesian fires as seen by the Tropospheric Emission Spectrometer (TES), and compare to observations from 2005 as a neutral year; we refer to the differences for 2006–2005 as anomalies. TES provides coincident, vertically resolved, tropospheric profiles for these species. The O₃ profiles have vertical resolution of ~6 km, allowing a new perspective on the ENSO perturbation to O₃. We also compare the effects of the 2006 El Niño with those of the 2004 event.

2. Observations

[7] TES is a Fourier transform IR emission spectrometer [Beer *et al.*, 2001]. It was launched on the Aura satellite in July 2004 in a sun-synchronous polar orbit with an equator crossing time of ~13:45. We use data from global surveys (GS) which consist of 16 orbits over 26 h; a new GS starts every other day. The nadir vertical profiles are spaced 1.6° apart along the orbit track and have a footprint of 5 × 8 km².

[8] TES retrievals are described by Bowman *et al.* [2006], Clough *et al.* [2006], and Kulawik *et al.* [2006]. The O₃ and CO prior information is derived from a simulation with the MOZART model [Brasseur *et al.*, 1998]; the prior profiles

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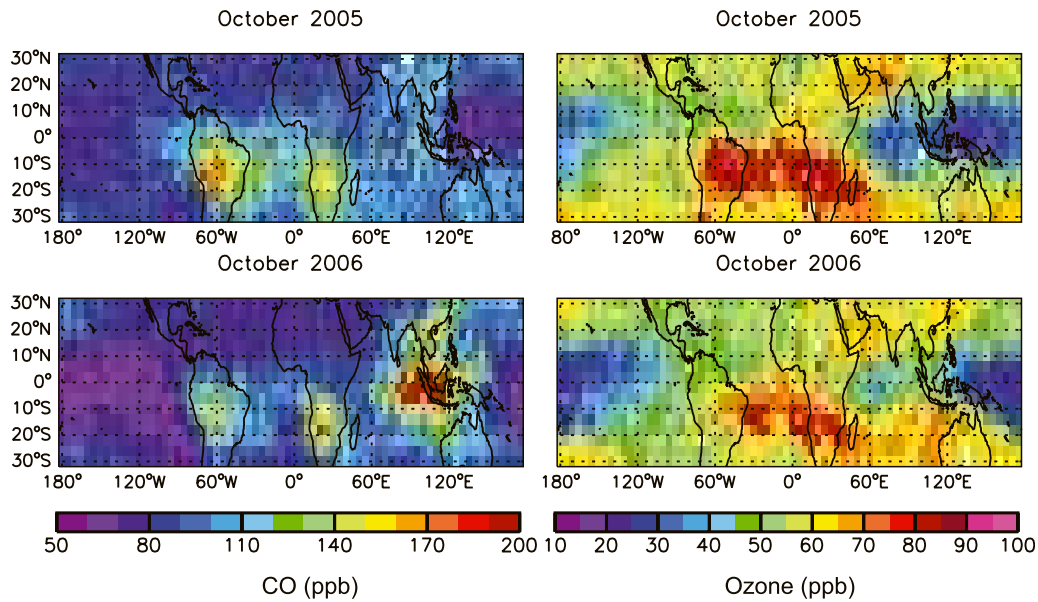


Figure 1. TES measurements of (left) CO and (right) O₃ in ppb for October of 2005 and 2006 at the 511 hPa retrieval level.

are monthly means in blocks of $10^\circ \times 60^\circ$ (latitude by longitude). Prior profiles for H₂O are derived from the GEOS data assimilation analysis fields. Typical averaging kernels (AKs) are shown in Worden *et al.* [2007] for O₃, Luo *et al.* [2007a] for CO, and Shephard *et al.* [2008] for H₂O. To ensure that the spatial structure we are analyzing is not caused by spatial patterns in the prior, we reprocess the TES profiles using a single prior (the average of those for 30°N – 30°S), following Zhang *et al.* [2006].

[9] The degrees of freedom for signal (DOFS) is a metric for the vertical information in the retrieved profile [Rodgers, 2000]. The DOFS for O₃ in the tropical troposphere is 1.5–1.7, showing that TES can distinguish between lower and upper tropospheric O₃ [Jourdain *et al.*, 2007]. The DOFS for CO in the tropics was 1.0–1.5 after launch, but degraded to 0.6–0.8 because of changes in the instrument optical alignment. Warm up of the optical bench in November 2005 improved the TES CO signal, and average values for the DOFS doubled, with many values >2 in the tropics [Rinsland *et al.*, 2006]. The typical DOFS for H₂O is 3–5, giving vertical resolution of ~ 3.5 km [Shephard *et al.*, 2008].

[10] We use V002 of TES data. Validation with ozone-sonde data shows that TES O₃ profiles are biased high by 3–10 ppb [Nassar *et al.*, 2008]. TES CO measurements are consistent with those of MOPITT [Luo *et al.*, 2007a] and are within 15% of aircraft data [Luo *et al.*, 2007b; J. Lopez *et al.*, TES carbon monoxide validation during two AVE campaigns using the Argus and ALIAS instruments on NASA's WB-57F, submitted to *Journal of Geophysical Research*, 2007]. TES H₂O profiles are within 5–15% of Vaisala radiosonde data, and within 5–40% of data from a cryogenic frostpoint hygrometer [Shephard *et al.*, 2008].

[11] Data quality and cloud filtering criteria used to select the TES profiles are given in the auxiliary material¹. Results

are shown as monthly means binned on a grid of $4^\circ \times 5^\circ$ at the 511 hPa retrieval level. As the AKs are relatively broad, the same pattern is seen over a range of several kilometers. We also show the zonal pattern for 0° – 12°S , with data binned every 15° of longitude.

3. Results

[12] Mixing ratios of CO in the mid-troposphere exceeded 200 ppb over Indonesia in October 2006, compared to ~ 110 ppb in October 2005, while those of O₃ were 45–55 ppb and 25–35 ppb respectively (see Figure 1). In September 2006, CO was generally less than 120 ppb in the regions with elevated CO in October. Over S. America, the south tropical Atlantic, and parts of southern Africa, both CO and O₃ were lower in October 2006 than in 2005.

[13] Figure 2 shows the difference in CO, O₃, and H₂O between 2006 and 2005 for October to December. The CO anomaly over Indonesia and the eastern Indian Ocean in October and November is >80 ppb in the center of the feature, with a maximum anomaly >125 ppb. CO is highest in the lower half of the troposphere in October and November 2006 (not shown). By December, the CO anomaly has decreased to less than ~ 30 ppb, and it is gone by January. We ascribe the high CO in 2006 to enhanced burning in Indonesia associated with the El Niño related drought, when fire emissions were almost six times those in 2005 in the GFED2 inventory [van der Werf *et al.*, 2006].

[14] Over the eastern Pacific, S. America, and the south Atlantic, CO is lower in October 2006 by 15–30 ppb, and over southern Africa it is lower by <15 ppb; differences are less than ± 15 ppb in the later months. The lower CO over S. America in October 2006 is likely caused by lower fire emissions; estimates in the GFED2 inventory for August–September are ~ 37 Tg CO from S. America in 2006, half the amount in 2005.

[15] The O₃ anomaly over Indonesia and the eastern Indian Ocean persists from October to December. Ozone

¹Auxiliary materials are available in the HTML. doi:10.1029/2007GL031698.

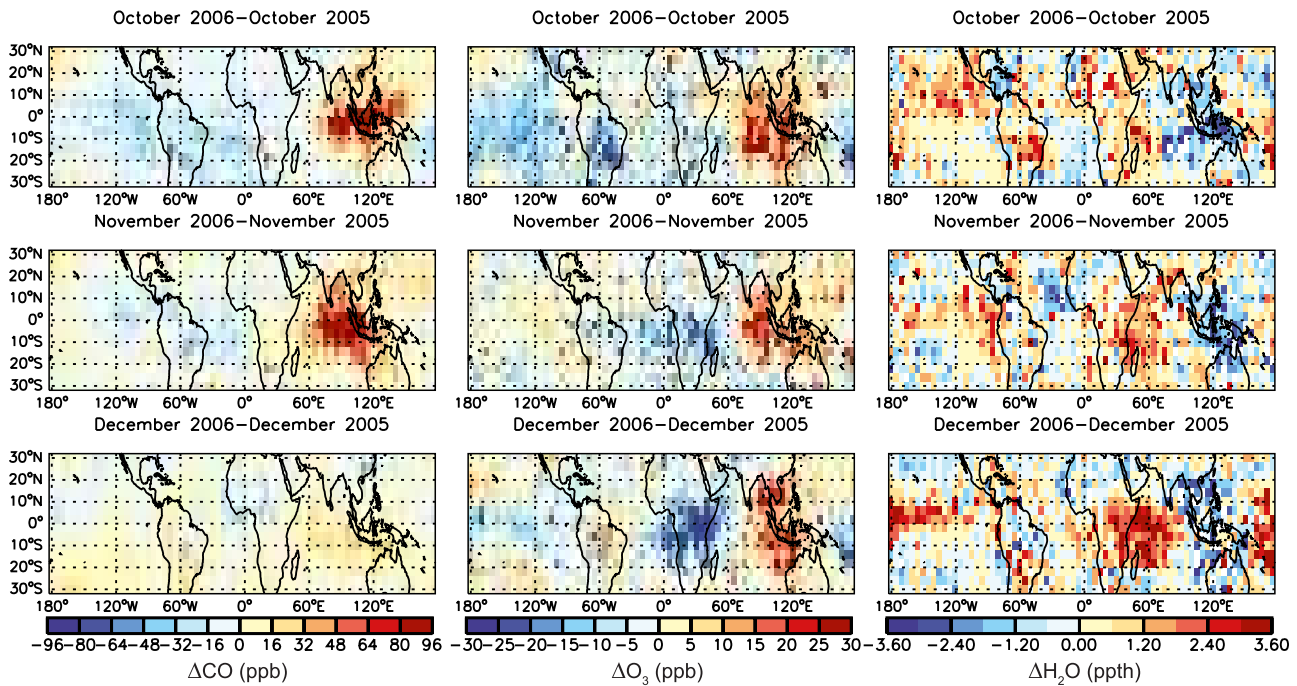


Figure 2. The difference between 2006 and 2005 for (left) CO, (middle) O₃, and (right) H₂O for October to December at the 511 hPa retrieval level; CO and O₃ are in ppb, H₂O in ppt. h.

differences are 10 ppb to >30 ppb from 15°N to 15°S for 3 months. As O₃ is typically very low here, this is an increase of 30–75%. The differences in Figure 2 are confirmed by ozonesonde profiles from Kuala Lumpur and Java (<http://croc.gsfc.nasa.gov/shadoz>). Figure 3 shows that the anomaly is slightly larger in the lower troposphere (LT) than the upper troposphere (UT) in October; the converse is true in November. The positive O₃ anomaly is associated with a negative H₂O anomaly (Figure 2). The high O₃ over Indonesia in 2006 is likely caused by photochemical production in the LT from the high fire emissions, reduced photochemical loss (lower H₂O), and reduced convection (El Niño conditions), the same mechanisms discussed in the context of the 1997 El Niño anomaly [Chandra *et al.*, 1998; Sudo and Takahashi, 2001]. Enhanced lightning in late 2006 compared to 2005 may also play a role as discussed below.

[16] October shows the dipole pattern found in TCO data for 1997 and in years with minor El Niños [Chandra *et al.*, 1998; Ziemke and Chandra, 2003], with a negative O₃ anomaly of 10–15 ppb (15–30%) over the central Pacific. The low O₃ in October is caused by enhanced convection over the western Pacific, which mixes higher UT O₃ into the LT, a region of net photochemical loss in the remote Pacific [e.g., Schultz *et al.*, 1999]. Maps of outgoing long-wave radiation (OLR) were used to determine regions of convection (<http://www.cdc.noaa.gov/HistData>). OLR maps show that there was a much smaller region of convection in the western Pacific in November than in October 2006, which may explain the lack of a negative O₃ anomaly there in November.

[17] Ozone is lower over Brazil and Bolivia by 10–30 ppb in October 2006 (likely related to lower fire emissions), and is higher in December by 5–15 ppb. Ozone is 15–30 ppb lower over equatorial Africa in December

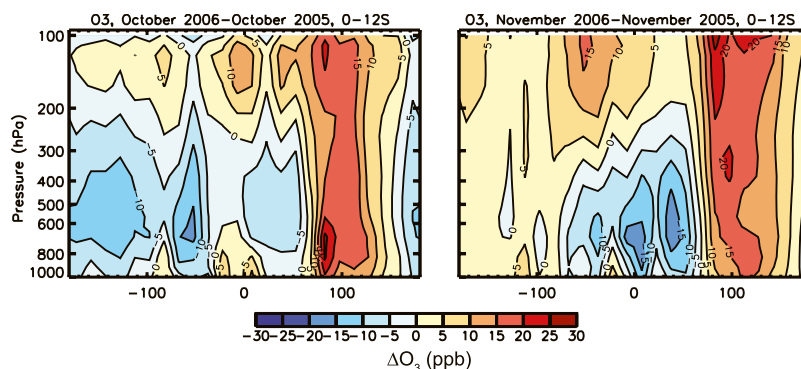


Figure 3. Difference between O₃ in 2006 and 2005, for 0–12°S, for (left) October and (right) November. The color scale for the O₃ difference (in ppb) is the same as that for Figure 2.

Table 1. Tropospheric O₃ Column Anomaly Over Indonesia in 1997, 2004, and 2006^a

| Instrument | | October | November | December |
|------------|-------------------|---------|----------|----------|
| EPTOMS | 1997 ^b | 20.3 | 14.4 | 11.4 |
| TES | 2004 ^c | | 6.6 | 5.1 |
| TES | 2006 ^c | 11.5 | 11.1 | 10.2 |

^aAnomalies measured in DU are given for the region 5°N–10°S, 90°–120°E. Data for 1996 and 1997 were obtained from http://code916.gsfc.nasa.gov/Data_services/cloud_slice/data/tropo.txt.

^bAnomaly with respect to 1996.

^cAnomaly with respect to 2005.

2006. These negative anomalies are confined to the LT (Figure 3).

[18] Water vapor is lower over Indonesia in October to December of 2006 than in 2005 by a few ppt (parts per thousand), and higher over the Pacific (Figure 2). There are not large absolute differences over the Pacific until December, although there are large relative differences (Figure S1). The changes in H₂O between 2005 and 2006 are caused by the eastward movement of convection during the El Niño year. There is much higher H₂O (>80%) over eastern Africa and the western Indian Ocean in December 2006, which likely contributes to the lower O₃ there, because of enhanced photochemical loss. The largest relative anomaly in H₂O is located from ~600 hPa to 250 hPa at all longitudes (not shown).

[19] The differences in values between 2004 and 2005 for CO and O₃ for November and December are smaller in magnitude and spatial extent than those between 2006 and 2005 (Figure S2). The CO anomaly in November 2004 is 25–45 ppb between Borneo and Sumatra, but is <25 ppb over the eastern Indian Ocean. Ozone is higher by 5–20 ppb in a small region over Indonesia in both months in 2004. We calculated TCO anomalies from the TES O₃ profiles over Indonesia (5°N–10°S, 90°–120°E) and found that the TES TCO anomalies for 2004–2005 are similar (within 1–2.5 DU) to those derived from OMI and MLS by Chandra *et al.* [2007].

4. Discussion

[20] We compare the TCO anomalies over Indonesia from TES for 2004 and 2006 to those derived for 1997 from total column O₃ data by Ziemke and Chandra [2003] in Table 1. The TCO anomaly in 2006 was smaller than that in 1997, particularly in October, but almost twice as large as that in 2004. We find that the magnitude of the TCO anomaly in October of 2004, 2006, and 1997 is related to the magnitude of CO emissions from fires in Indonesia derived by van der Werf *et al.* [2006], 24 Tg, 82 Tg, and 193 Tg respectively. The relatively high burning in 2006 is related to the strength of the drought, as discussed by G. R. van der Werf *et al.* (Climate controls on the variability of fires in the tropics and subtropics, submitted to *Global Biogeochemical Cycles*, 2007), who show that July to October of 2006 was the driest period over Indonesia since 1997.

[21] MOPITT data show the highest CO over Indonesia in late 2006 since observations began in 2000, consistent with the relative emissions estimates in the GFED2 inventory (G. R. van der Werf *et al.*, manuscript in preparation,

2007). Prior to 2006, CO over Indonesia was highest in 2002, followed by 2004, both El Niño years with droughts and relatively high fires [Edwards *et al.*, 2006].

[22] The O₃ anomaly was almost as large in December 2006 as in November, while that for CO was much less, consistent with the fires ending in early November. The O₃ anomaly may have persisted longer than that for CO because of NO_x production by lightning when convection moved over Indonesia bringing rain. Figure 4 shows difference plots derived from the Lightning Imaging Sensor (LIS) data [Mach *et al.*, 2007] (<http://thunder.msfc.nasa.gov>). There was more lightning over Indonesia in November and December 2006 than in 2005 by a factor of 2–3, although there was less in October.

[23] The tropospheric NO₂ column over Indonesia given by the SCIAMACHY and OMI instruments [Boersma *et al.*, 2007] was higher in October–November of 2006 than in 2005 in the vicinity of fires; NO₂ was slightly elevated in December 2006 in a broader area, consistent with a larger source of NO_x from lightning.

[24] Lower O₃ over equatorial Africa in December may also be caused by changes in convection between 2005 and 2006, although these are not necessarily ENSO related. There was a drought in eastern Africa in late 2005, and very high rainfall in 2006. More convection in 2006 would

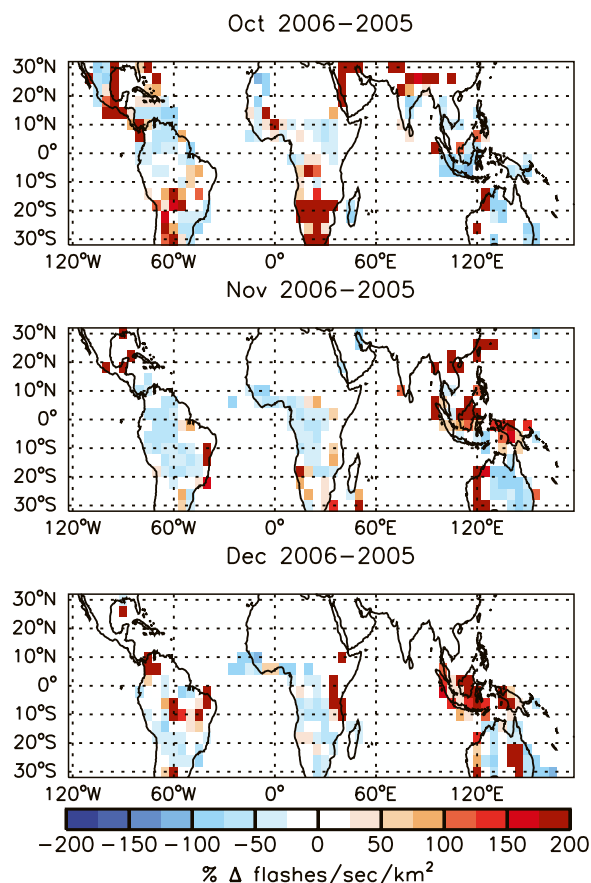


Figure 4. Percent difference in lightning flash density between 2006 and 2005, calculated from the Science Data product of LIS. The difference is shown for areas with a lightning flash density greater than 1×10^{-8} flashes $\text{sec}^{-1} \text{km}^{-2}$ in either year.

lead to more photochemical loss of O₃ in the LT, because of enhanced downward mixing of O₃ from the UT, and higher H₂O (Figure 2). There was also less lightning over much of southern Africa in late 2006 compared to 2005, except in easternmost Africa, so that NO_x emissions would be lower.

[25] We are using simulations with the GEOS-Chem model in conjunction with TES data to explore the mechanisms responsible for differences in tropical O₃ in 2005 and 2006, including the roles of dynamics, fire emissions, and lightning. The combination of vertically resolved, collocated data for CO, O₃, and H₂O from TES offers powerful constraints on model results.

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