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RESEARCH ARTICLE

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Key Points:

- First estimates of ECC ozonesonde uncertainties using SHADOZ data records
- Ozone uncertainties are generally within 15% and peak around the tropopause where ozone measurements approach the uncertainty estimates
- Uncertainties in background and sensor current dominate the troposphere, while conversion efficiency and flow rate dominate the stratosphere

Supporting Information:

- Supporting Information S1

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First Reprocessing of Southern Hemisphere Additional OZonesondes Profile Records: 3. Uncertainty in Ozone Profile and Total Column

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Abstract Reprocessed ozonesonde data from eight SHADOZ (Southern Hemisphere Additional OZonesondes) sites have been used to derive the first analysis of uncertainty estimates for both profile and total column ozone (TCO). The ozone uncertainty is a composite of the uncertainties of the individual terms in the ozone partial pressure (P_{O_3}) equation, those being the ozone sensor current, background current, internal pump temperature, pump efficiency factors, conversion efficiency, and flow rate. Overall, P_{O_3} uncertainties (ΔP_{O_3}) are within 15% and peak around the tropopause (15 ± 3 km) where ozone is a minimum and ΔP_{O_3} approaches the measured signal. The uncertainty in the background and sensor currents dominates the overall ΔP_{O_3} in the troposphere including the tropopause region, while the uncertainties in the conversion efficiency and flow rate dominate in the stratosphere. Seasonally, ΔP_{O_3} is generally a maximum in the March–May, with the exception of SHADOZ sites in Asia, for which the highest ΔP_{O_3} occurs in September–February. As a first approach, we calculate sonde TCO uncertainty (ΔTCO) by integrating the profile ΔP_{O_3} and adding the ozone residual uncertainty, derived from the McPeters and Labow (2012, doi:10.1029/2011JD017006) 1σ ozone mixing ratios. Overall, ΔTCO are within ± 15 Dobson units (DU), representing ~ 5 – 6% of the TCO. Total Ozone Mapping Spectrometer and Ozone Monitoring Instrument (TOMS and OMI) satellite overpasses are generally within the sonde ΔTCO . However, there is a discontinuity between TOMS v8.6 (1998 to September 2004) and OMI (October 2004–2016) TCO on the order of 10 DU that accounts for the significant 16 DU overall difference observed between sonde and TOMS. By comparison, the sonde–OMI absolute difference for the eight stations is only ~ 4 DU.

Plain Language Summary Electrochemical concentration cell ozonesondes are balloon-borne instruments that measure profiles of ozone from launch up to 35 km. The high vertical resolution allows researchers to capture features in measurement-deficient regions of the atmosphere: the boundary layer, tropopause region that marks the boundary between the troposphere and stratosphere, and the protective ozone shield layer around 20–25 km. The Southern Hemisphere Additional OZonesondes (SHADOZ) project has been archiving ozonesonde data from up to 17 stations since 1998 that are available publicly at <https://tropo.gsfc.nasa.gov/shadoz>. As with any instrument there are uncertainties in the measurements. For ozonesondes the uncertainty in the ozone measurement is associated with the mechanics of the electrochemical concentration cell sensor and the preparation procedures. For the first time we calculate the ozone profile uncertainties measured by the ozonesonde using the SHADOZ 1998–2016 database. The advantage of doing a detailed uncertainty analysis is that it reveals areas of the measurements where we can refine operational procedures to reduce the uncertainty and where additional research is needed to improve the basics of this measurement.

1. Introduction

1.1. Influence of ECC Ozonesonde Measurements

The electrochemical concentration cell (ECC) ozonesonde was developed by Walter Komhyr (Komhyr, 1969) and has been flown at ~ 80 stations worldwide since the late 1960s (Hassler et al., 2008; Logan, 1985; Oltmans et al., 2006; Stauffer et al., 2016; Tarasick et al., 2005; Tiao et al., 1986). The ECC instrument consists of a

gas-sampling pump connected to an ozone sensor and an electronic interface that connects the ozone sensor to a radiosonde for data telemetry (see Figure 1 in Komhyr et al., 1995). Measured parameters transmitted to the ground receiving station are ozone current, the ozonesonde's pump temperature, motor voltage and current, and ambient pressure, temperature, and relative humidity (RH) (*P-T-U*). In recent decades, winds and Global Positioning System-enabled measurements became available. In flight, the instrument is encased in a weatherproof box that is tethered to a balloon, capable of measuring ozone up to an altitude of ~35 km. The balloon ascent rate, typically around 5 m/s, and data transmission rate lead to a vertical resolution within 150 m.

Light-weight, compact, and relatively easy to prepare and launch, ozonesondes (also referred to here as sondes) fulfill an important role in providing high vertical resolution ozone (O_3) profiles from the surface to the middle stratosphere, capable of making measurements during polar night and in cloudy and rainy conditions. They are readily deployed from remote locations, such as over Antarctica, and high-latitude Europe, United States, and Canada. For example, during the Match campaigns over the Arctic (Rex et al., 1998, 2006; von der Gathen et al., 1995), trajectory pathways of ozone-depleted air parcels measured from one location can be forecast and sondes launched from other sites can intercept the low-ozone filaments. The Intercontinental Chemical Transport Experiment Ozonesonde Network Study (IONS) series of North American campaigns collected hundreds of soundings with daily launches from 8 to 20 sites over 3 to 6 week periods to augment aircraft in situ and lidar profiles (Thompson, Stone, et al., 2007; Thompson et al., 2008, 2011). Studies using tropical ozonesonde measurements have examined the tropical wave-one feature (Sauvage et al., 2006; Thompson et al., 2003, 2017), quasi-biennial oscillations and El Niño–Southern Oscillation features (Lee et al., 2010; Logan et al., 2003; Randel & Thompson, 2011; Thompson et al., 2001; Witte et al., 2008), and the tropical transition layer (Corti et al., 2006; Folkins et al., 1999; Fu et al., 2007; Gettelman & Forster, 2002; Randel et al., 2007; Thompson et al., 2012).

Ozonesonde data have become highly valued over the past 20 years as a large community is focused on O_3 trends in the upper troposphere and lower stratosphere (UT and LS), a region where satellite data do not provide the vertical resolution and sampling frequency that sondes typically do (Bodeker et al., 1998; Gebhardt et al., 2014; Kivi et al., 2007; Rao et al., 2003).

1.2. Reprocessing of ECC Ozonesonde Data

As popular as the sounding data have become, it is clear when long-term sonde-based O_3 records are examined in detail that changes in the ozonesonde instrument, calibration, and preparation techniques, and data processing methods lead to discontinuities and possibly artifact trends at individual sites (Witte et al., 2017). In addition, there are station-to-station variations in satellite O_3 biases versus sondes, biases among stations, and biases within the data record of an individual station that must be corrected for if sondes are used to assess measurement uncertainties and the reliability of O_3 profile trends (Thompson, Witte, et al., 2007; Thompson et al., 2003, 2014).

Accordingly, the ozonesonde community has worked for many years to establish quality assessment standards for ozonesondes. The first important step in this effort was the establishment in the mid-1990s of the World Calibration Center for Ozone Sondes (WCCOS) (Smit & Kley, 1998; Smit & Straeter, 2004a, 2004b). The periodic intercomparison experiments conducted in WCCOS, called Jülich Ozonesonde Intercomparison Experiments (JOSIE), operate with a standard O_3 reference ultraviolet photometer in a chamber. In the test chamber ozone is introduced under changing temperature and pressure conditions at a rate that simulates profiles that correspond to standard high-latitude, midlatitude, subtropical, and tropical conditions. The first JOSIE (JOSIE-1996; Smit & Kley, 1998) included non-ECC sondes that have largely been replaced at operational stations (De Backer et al., 1998; Fujimoto et al., 2004; Stübi et al., 2008).

From subsequent JOSIE campaigns (1998 and 2000) the focus of the WCCOS tests has been on ECC sondes manufactured by Science Pump Corporation (SPC) and Environmental Science (ENSCI) and characterizing how different SST (sensing solution type) perform under the various simulations (Smit et al., 2007). These campaigns revealed that differences in O_3 , as measured among different ECC sensors, are largely due to differences in preparation procedures and SST used by participating researchers. The outcome was that two combinations of instrument type and SST were recommended as standard operating procedures



Figure 1. Locations of the eight reprocessed Southern Hemisphere Additional OZonesondes sites for which uncertainty estimates are calculated. Table 1 lists the latitude/longitude per site.

(Smit et al., 2007; Smit & ASOPOS (Panel for the Assessment of Standard Operating Procedures for Ozonesondes), 2014). The same combination emerged as the WMO (World Meteorological Organization) preferred standard during the BESOS (Balloon Experiment on Standards for Ozonesondes) campaign in 2004 that evaluated the ozone response of commonly used ECC/SST pairings following JOSIE standard operating procedures guidelines under ambient conditions (Deshler et al., 2008).

After 2010, several Ozone Sonde Expert meetings and the SPARC-IO3C-IGACO-NDACC (Stratospheric Processes and their Relation to Climate, International Ozone Commission, International Gases and Aerosols Composition, Network for Detection of Atmospheric Chemical Composition Change) SI2N activities considered how to homogenize long-term data sets for trends because a number of stations had more than 30 years of ECC records, albeit with both ozonesonde and radiosonde changes. These activities and the results of JOSIE, BESOS, and further dual balloon soundings led to the creation of the O3S-DQA (Ozone Sonde Data Quality Assessment) panel report (Smit & O3S-DQA, 2012, hereafter referred to as Smit12) the goals of which are to (1) establish guidelines for reprocessing ozonesonde data records to remove inhomogeneities due to instrumental or procedural artifacts and (2) determine the contributions of the individual uncertainties of the different instrumental parameters to the O_3 measurement. The WMO/GAW (Global Atmospheric Watch) Report #201, referenced as Smit and ASOPOS (Panel for the Assessment of Standard Operating Procedures for Ozonesondes) (2014, hereafter referred to as the WMO/GAW Report), is a comprehensive summary of the Smit12 findings. The O3S-DQA panel report has also formalized the concept of transfer functions to compensate for instrument-SST changes. Deshler et al. (2017) have recently published the set of transfer functions based on the JOSIE and BESOS experiments and on unpublished field comparisons that incorporate a total of 197 tests with the SPC and ENSCI ECC sensors.

2. SHADOZ and Data Reprocessing

Since early 2016 we have been reprocessing the Southern Hemisphere Additional OZonesondes (SHADOZ) record (Witte et al., 2017; Thompson et al., 2017, hereafter referred to, respectively, as Witte17 and Thompson17) according to the O3S-DQA guidelines using customized software based on Skysonde (developed by Allen Jordan at National Oceanic and Atmospheric Administration (NOAA)/Earth Systems Research Laboratory/Global Monitoring Division (NOAA/ESRL/GMD)).

Figure 1 displays the map of the stations for which data are analyzed in detail in this study. Details of the first reprocessing of SHADOZ data from seven sites appear in Witte17. When added to data from six other SHADOZ stations, including four sets from stations reprocessed by NOAA/ESRL/GMD, 14 stations with continuous data of at least one decade have been evaluated in Thompson17. Those reprocessed SHADOZ data were compared to three backscatter ultraviolet-type satellite total column ozone (TCO) amounts spanning 1998–2016 and to colocated ground-based instruments at nine SHADOZ stations. Thompson17 showed that, compared to earlier evaluations (Thompson, Witte, et al., 2007; Thompson et al., 2003, 2012), offsets between ozonesonde and satellite TCO are reduced due to the homogeneity of the newly reprocessed ozonesonde data records. Most stations ended up with sonde TCO, satellite, and, where applicable, ground-based instruments, within 2% of one another.

In this paper we reprocess data for Réunion a second time and present the first reprocessed data for Costa Rica (various locations around San José) and Nairobi (refer to locations in Figure 1). However, the principal goal here is to report an uncertainty analysis of the ozonesonde measurement system, term by term, for the eight stations we have reprocessed at National Aeronautics and Space Administration/Goddard Space Flight Center (NASA/GSFC), as well as uncertainties in TCO amounts. In the latter case we compare the columns to Earth Probe/TOMS (Total Ozone Mapping Spectrometer) and OMI (Ozone Monitoring Instrument) satellite overpasses.

Following Witte17, this study continues the O3S-DQA goal of determining uncertainties in the ozonesonde measurement system. The treatment of uncertainty in this study closely follows the definitions described in Smit12. Section 3 describes details of the ozone measurement and the reprocessing of SHADOZ data to date. Section 4 analyzes the uncertainty relationships term by term. The profile uncertainties appear in section 5 and column uncertainties in section 6, followed by a summary (section 7).

3. Details of the Ozonesonde Measurement and Reprocessed SHADOZ Ozonesonde Data

3.1. The Ozonesonde Measurement

The ECC sensor measures O_3 using iodine/iodide electrode reactions. Two platinum electrodes are immersed in separate cathode and anode chambers of differing concentrations of potassium iodide (KI) solution. The anode cell contains a solution saturated with KI. Both cells contain an equal concentration of potassium bromide (KBr) and a phosphate buffer to maintain a neutral pH. An ion bridge connecting the two chambers allows ions to flow between the two cells but prevents mixing, thereby preserving their respective concentrations. Ambient air containing O_3 is pumped into the cathode cell and reacts with iodide (I^-) in solution to form iodine (I_2). To maintain electrochemical equilibrium, I_2 is converted back to I^- on the platinum electrode resulting in the release of two electrons. Thus, each O_3 molecule entering the sensor causes two electrons to flow through the ECC's external circuit, which it measures as a current. The resulting electrical current is proportional to the amount of O_3 in the sampled air. The electrochemical technique assumes no secondary reactions take place and a 1:1 stoichiometric relationship of the $O_3:I_2$ is maintained. The relationship between O_3 and the electrical current is defined by the following equation:

$$P_{O_3} = 4.307 \times 10^{-2} \frac{(I_M - I_B) T_P}{\Psi_P \Phi_P \eta_C}, \quad (1)$$

where

- P_{O_3} ozone partial pressure, mPa;
- I_M cell current, μA ;
- I_B cell background current, μA ;
- T_P ozonesonde pump temperature, K;
- Φ_P pump flow rate, ml/s;
- Ψ_P pump flow efficiency, unitless; and
- η_C conversion efficiency, which is generally assumed to be 1.

The constant, 4.307×10^{-2} , is the half ratio of the ideal gas constant ($8.314 \text{ J K}^{-1} \text{ mole}^{-1}$) to Faraday's constant ($9.6487 \times 10^4 \text{ C mole}^{-1}$). Equation (1) is similar to what is written in the WMO/GAW report.

3.2. SHADOZ Reprocessed Data

SHADOZ is the premier archive of tropical and subtropical ECC ozonesonde data. Since this National Aeronautics and Space Administration program started in 1998, SHADOZ has archived ozonesonde profiles from up to 17 tropical sites with support from NOAA/ESRL/GMD and international partners. Data are publicly available at <https://tropo.gsfc.nasa.gov/shadoz>. Information on the eight SHADOZ sites described in this study (Figure 1) appears in Table 1; each site's location is summarized. Launches are two to four times per month. We use 1998–2016 data from the six sites that have been reprocessed by Witte17. Costa Rica and Nairobi were later reprocessed based on Witte17 methods that closely follow Smit12 guidelines. Evaluation of these two data sets can be found in Thompson17.

Table 1
Southern Hemisphere Additional OZonesondes (SHADOZ) Sites Used in This Study

Site	Latitude, Longitude	Time period	Profile total
Ascension Is., UK	7.98°S, 14.42°W	1998 to August 2010 and 2016 ^a	632
Costa Rica ^b	9.94°N, 84.04°W	July 2005–2016	510
Hanoi, Vietnam	21.02°N, 105.80°E	September 2004–2016	245
Irene, S. Africa	25.90°S, 28.22°E	November 1998–2007 and September 2012–2016 ^a	337
Kuala Lumpur, Malaysia	2.73°N, 101.70°E	February 1998–2010 and March 2012–2016 ^a	388
La Réunion, France	21.10°S, 55.48°E	1998–2016	618
Nairobi, Kenya	1.27°S, 36.80°E	1998–2016	827
Natal, Brazil	5.42°S, 35.38°W	May 1998–2011 and September 2014–2016 ^a	595

^aNote that there are data gaps. ^bPrevious sites, Alajuela and Heredia, are within 0.5° of San Pedro (current site).

A transfer function based on Deshler et al. (2017) is applied to the Nairobi 1998 to October 2010 data to convert O₃ measured with a nonstandard ENSCI ECC/1% full buffer SST to the WMO and manufacturer recommended ENSCI/0.5% half buffer equivalent. A similar transfer function is used to homogenize August 2007–2016 Réunion data from an ENSCI/0.5% full buffer SST to the standard ENSCI/0.5% half buffer. Both applications reduce O₃ measurements by about 4%. Witte17 applied a transfer function to 6% of the profiles in the beginning of the Réunion record where SPC/0.5% half buffer SST was adjusted to the WMO recommended ENSCI/0.5% half buffer. An overview of transfer functions applied by Witte17 and in this study to the eight SHADOZ sites is found in Table 2. Transfer functions have been applied to over half the profiles at Réunion and Nairobi and 48% at Hanoi. Transfer functions applied at Ascension and Natal account for 10% and 17% of the data sets, respectively.

Witte17 concluded that O₃ trend assessments could not be made using original and initially reprocessed Réunion data due to the solution change after August 2007, for which transfer functions had not yet been applied. We examine the effect of applying this additional transfer function to the reprocessed Réunion data set by comparing time series of TCO. In this study, sonde TCO is calculated by integrating O₃ partial pressure up to 10 hPa and adding an O₃ climatology from balloon burst to the top of the atmosphere (TOA), taken from the McPeters and Labow (2012) look-up table (reported in Dobson units (DU)). From Figure 2a the reprocessed data used in Witte17 show a significant difference in mean TCO of 16.4 DU between the 1998–2006 and 2007–2016 time periods (black dashed lines). However, this difference is reduced to 6.5 DU (blue) after the additional transfer function has been applied. Thus, as of this study, the significant discontinuity found by Witte17 has almost disappeared, allowing for meaningful trend assessments. Later, we show the impact of the uncertainty of the transfer function to the total ozone uncertainty budget. Further evaluation of this newly homogenized data set has been carried out by Thompson17.

Table 2
Summary of Transfer Functions Applied for Each SHADOZ Site

Site	Transfer function	Dates applied	Conversion applied	Profiles
Ascension	Yes	1998–2001	ENSCI/1% to SPC/1%	10%
Costa Rica	No	—	—	—
Hanoi	Yes	2004 to May 2009	ENSCI/2% unbuffered to ENSCI/0.5%	48%
Irene	No	—	—	—
Kuala Lumpur	No	—	—	—
La Réunion	Yes	1998 to July 2000 8 records in 1998 and August 2007–2016	SPC/0.5% to ENSCI/0.5% ENSCI/0.5% full buffer to ENSCI/0.5%	7% 56%
Natal	Yes	March 1999 to July 2002	ENSCI/1% to SPC/1%	17%
Nairobi	Yes	1998 to June 2010	ENSCI/1% to ENSCI/0.5%	64%

Note. Conversions applied are to World Meteorological Organization recommended electrochemical concentration cell sensor/sensing solution type pairs: ENSCI/0.5% (half buffer) and Science Pump Corporation (SPC)/1% (full buffer).

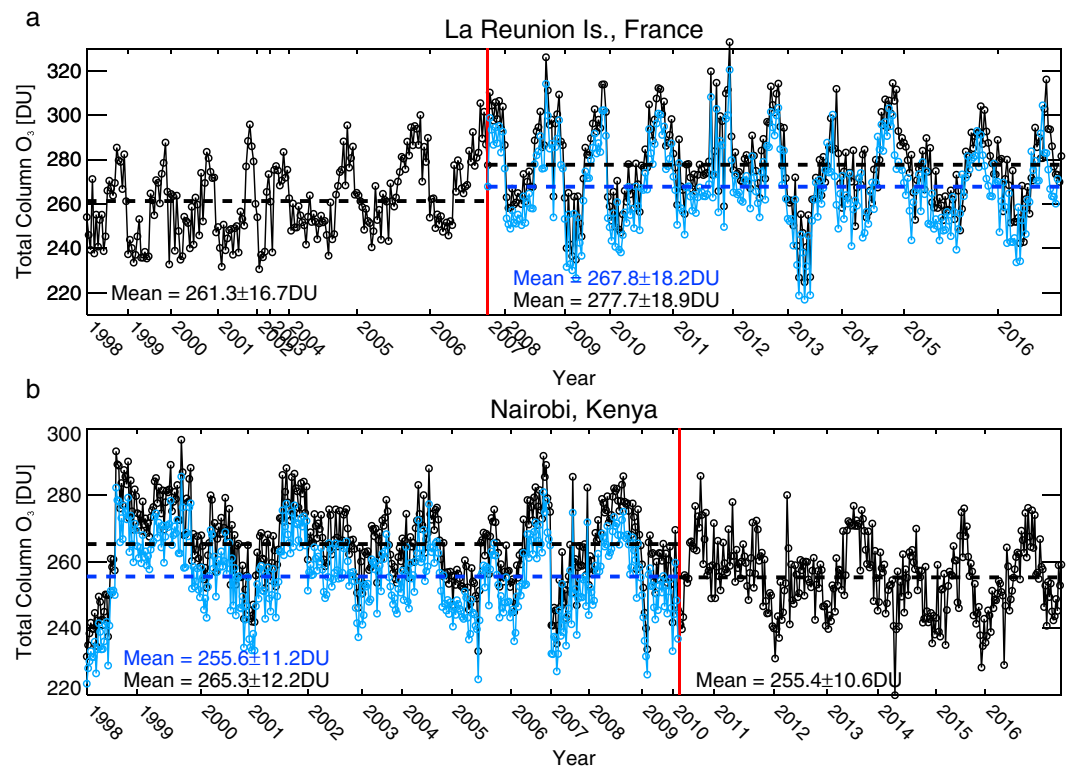


Figure 2. Time series of total column ozone (TCO) at (a) Réunion and (b) Nairobi. Years are shown on the x axis. Reprocessed data are in black, and reprocessed data + transfer function are in blue. Transfer functions have been applied in (a) since 2007 and in (b) before June 2010, marked with a vertical red line. Black dashed lines indicate the mean TCO for reprocessed data only (values included in black). The blue dashed lines shows the mean TCO for the period where transfer functions have been applied (values included in blue) DU = Dobson units.

Results of applying a transfer function to the early part of the Nairobi data set are shown in Figure 2b. As with Réunion, Nairobi retains a significant discrepancy of 9 DU prior to reprocessing between the periods 1998 to October 2010 and November 2010–2016 (black dashed lines). The offset disappears after a transfer function is applied (blue dashed line).

We examine more closely the impact of applying transfer functions to the Réunion and Nairobi data sets by comparing TCO values with OMI (Version 3) TCO overpass data (Levelt et al., 2006) as a reference. Results are shown in Figure 3 as histograms of the percentage difference with respect to sondes. Biases with respect to OMI at Réunion are significantly reduced after transfer functions have been applied (blue hashes, Figure 3a), where the Gaussian peak shifts from 4.3% to 0.7%. Most of the agreement falls within $\pm 5\%$. The homogenized Nairobi data, in Figure 3b, show a different impact. The Gaussian peak shifts from +2.2% to -1.5% , and the sonde bias with respect to OMI changes from a mostly positive (high-bias) regime to a negative (low-bias) regime. This low-bias relationship between the Nairobi sondes and satellite data is consistent with most of the SHADOZ network data sets (Hubert et al., 2016; Thompson17). Like Réunion, the agreement is $\pm 5\%$ making both these sites a stable reference for trends analysis and satellite validation.

4. The Ozone Uncertainty Equation

To improve the ozonesonde measurement system, Smit12 introduced the first instrumental uncertainty equation of the ECC-type ozonesonde based on the current best knowledge of the ECC performance under lab conditions. After reprocessing to remove all known inhomogeneities, the overall uncertainty in P_{O_3} , shown in equation (2), is the square root of the sum of the squares of the uncertainty in each term of the ozone partial pressure equation (equation (1)). The uncertainties are assumed to be random and Gaussian and therefore follow the Gaussian propagation of uncertainty. In equation (2), the assumption is that the uncertainties are not only random but also uncorrelated.

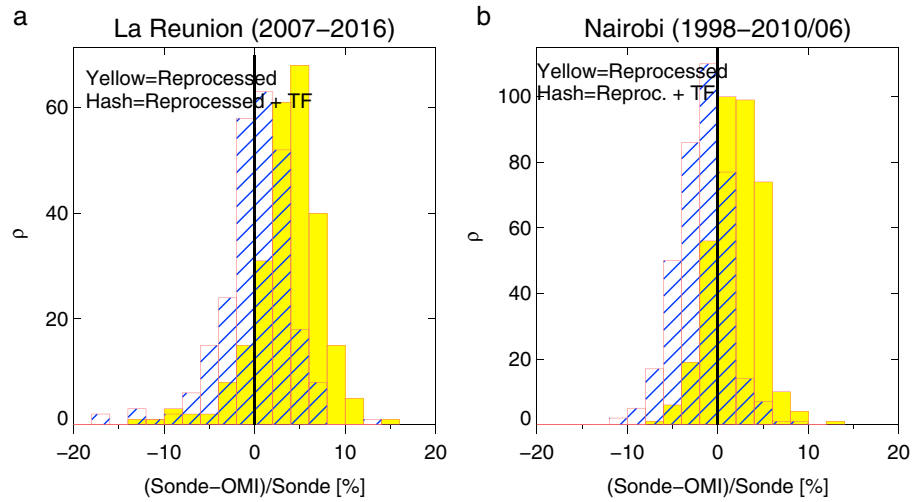


Figure 3. Histogram of the percentage difference in total column ozone between reprocessed sondes and Ozone Monitoring Instrument (OMI) overpasses with respect to sondes for (a) 2007–2016 Réunion and (b) 1998 to June 2010 Nairobi data. These are time periods for which transfer functions have been applied (blue hashed).

$$\frac{\Delta P_{O_3}}{P_{O_3}} = \sqrt{\frac{(\Delta I_M)^2 + (\Delta I_B)^2}{(I_M - I_B)^2} + \left(\frac{\Delta \eta_C}{\eta_C}\right)^2 + \left(\frac{\Delta \Phi_P}{\Phi_P}\right)^2 + \left(\frac{\Delta \Psi_P}{\Psi_P}\right)^2 + \left(\frac{\Delta T_P}{T_P}\right)^2} \quad (2)$$

where

- ΔP_{O_3} ozone partial pressure uncertainty,
- ΔI_M sensor current uncertainty,
- ΔI_B background current uncertainty,
- $\Delta \eta_C$ conversion efficiency uncertainty,
- $\Delta \Phi_P$ pump flow rate uncertainty at the ground,
- $\Delta \Psi_P$ pump flow efficiency uncertainty below 100 hPa, and
- ΔT_P pump temperature uncertainty.

The individual uncertainties are defined as a relative error $(\Delta x)/x$ and can be expressed as a percentage. The value of P_{O_3} and its error, or uncertainty in this case, can then be expressed as the interval $P_{O_3} \pm \Delta P_{O_3}$.

The first application of equation (2) was done for the McMurdo, Antarctic station data set (reference analysis can be downloaded at http://www.das.uwy.edu/~deshler/NDACC_O3Sondes/O3s_DQA/O3S-DQAGuideline_Summary_OzUncertainty_td.pdf). Van Malderen et al. (2016) also applied Smit12 uncertainties to the ECC ozonesonde measurements at Uccle (Belgium). Sterling et al. (2017) applied the formula to two SHADOZ stations (Hilo, Hawaii, and Pago Pago, American Samoa), Boulder, CO, and the South Pole, and Tarasick et al. (2016) applied their own uncertainty estimates to the Canadian ozonesonde records, taking into account the uncertainty terms in equation (2). We take advantage of these studies to compare with our methods and results.

Where applicable, based on Deshler et al. (2017) calculations, the overall uncertainty of the transfer function, ΔT_F , is 5%. We apply the Smit12 approach and add this uncertainty to the uncertainty in the conversion efficiency term.

Whereas in the report the $\Delta \Phi_P/\Phi_P$ term includes $\Delta \Psi_P/\Psi_P$, we separate the two terms for clarity. The rest of the instrumental uncertainty terms are defined in subsequent subsections.

Note that this study focuses only on the uncertainties of the ozonesonde instrument and does not take into account uncertainties due to radiosonde pressure offsets (offsets that lead to errors in the height registry of the computed ozone). Tarasick et al. (2016) do not correct for radiosonde errors but include a pressure offset uncertainty for the VIZ (± 1 hPa) and Vaisala RS-80 (± 0.5 hPa) manufacturers based on previous studies.

Conversely, Sterling et al. (2017) correct for pressure offsets but do not include the radiosonde pressure uncertainties in their ozone uncertainty calculations. The challenge of determining radiosonde errors, particularly in the non-Global Positioning System era, is still an ongoing debate.

The uncertainty in the ozone current, ΔI_M , is $\pm 0.01 \mu\text{A}$ for currents less than $1.0 \mu\text{A}$ and 1% elsewhere. This is taken as the overall resolution of the digital interface board (Smit12). V2 and V7 interface boards have similar piecewise uncertainties (Sterling et al., 2017). The Vaisala OIF11 and current generation OIF92 interface boards have an accuracy of $0.01 \mu\text{A}$ and $0.001 \mu\text{A}$, respectively (taken from the Vaisala manufacturer brochure). Uncertainty analysis done for the McMurdo station data set used $\Delta I_M = 0.1 \mu\text{A}$, while this uncertainty was not taken into account by Tarasick et al. (2016). Sterling et al. (2017) used similar ΔI_M during the digital era of the measurements that started in the 1990s.

SHADOZ sites encompass a wide range of radiosonde/ozonesonde systems that use a variety of interface boards with varying resolutions. For example, the Modem radiosonde used at Réunion, Lockheed-Martin-Sippican used at Ascension and Natal, and Vaisala employed at a number of SHADOZ sites use their own interfaces, which have different analog-to-digital converters and possibly different thermistors. This may impact the uncertainty estimate for the cell current and pump temperature measurement; however, further investigation is needed to determine the effect, if any. In this study, we simplify ΔI_M to $\pm 0.01 \mu\text{A}$ for currents less than $1.0 \mu\text{A}$ and 1% elsewhere, as recommended by Smit12.

Van Malderen et al. (2016) noted that equation (2) does not take into account the uncertainty due to the time lag of the response of the I_M , T_p , and I_B measurements. Tarasick et al. (2016) incorporated an e -folding response time to the ozone gradient to take into account slow ECC responses to changes in ozone due to variable ascent rates.

4.1. Background Current Uncertainty, ΔI_B

The background current (I_B) is the residual current measured by the sonde when sampling ozone-free air. Conventional processing of the sonde telemetry assumes that the background current remains constant during flight. There is no statistically robust method for estimating the uncertainty of the background current, ΔI_B . JOSIE studies used small sample sizes, fewer than 14 ECC sensors, to conduct the background current experiments published in Smit et al. (2007) and recommended in the WMO/GAW Report. During JOSIE-1996, significantly high backgrounds were recorded due, in part, to the slow decay in the ozone response and not allowing enough time for the background to drop lower (Johnson et al., 2002). Laboratory experiments by Vömel and Diaz (2010) tracked the decay of the cell current after exposure to ozone and showed that a much longer period of time (hours) can be required to approach initial values. There is also a dependence on the SST where experiments have shown a relationship between buffered KI SST and high ozone measurements due to a hysteresis effect (additional side reactions) that offset the ideal 1:1 stoichiometric ratio expected from the O_3 to I_2 reaction (Barnes et al., 1985; Johnson et al., 2002).

To further confound the issue, SHADOZ sites measure I_B in a number of different ways: some are based on an average value, the minimum value recorded during the conditioning process, prior to launch either in the lab or at the launch site, or set to an upper limit threshold for I_B values that exceed it. The many ways in which I_B is recorded are compounded by the quality of the ozone destruction filters used that does not guarantee uniformity and introduces a source of random uncertainty that cannot be easily quantified (Reid et al., 1996). This is particularly true for ozonesondes flown in the tropics where high humidity affects the ozone removal efficiency of the filter (Newton et al., 2016). Specific to our study that uses tropical-based sonde data, we use the 1σ uncertainty of $\pm 0.03 \mu\text{A}$ for sites that use the ENSCI ECC sensors and $\pm 0.02 \mu\text{A}$ for SPC sensors based on Witte17 calculations. The ΔI_B value is doubled, where I_B is missing or exceeds the threshold value of $0.05 \mu\text{A}$ (based on Witte17 reprocessing criteria). Sterling et al. (2017) adopted the same strategy using $\Delta I_B = \pm 0.02 \mu\text{A}$ for ENSCI ECCs. For the McMurdo measurements, ΔI_B was higher for SPC ($\pm 0.05 \mu\text{A}$), whereas Tarasick et al. (2016) took a different approach applying a pressure dependent correction. Table 3 summarizes the ΔI_B applied for the eight reprocessed sites based on the ECC sensor used. In general, one ECC type dominates a single site's data set. Included is the percent of profiles for which I_B is missing or exceeds the $0.05 \mu\text{A}$ threshold. The large spread of these percentages illustrates the difficulty in establishing low backgrounds in tropical environments as well as the variation in technique. For example, Costa Rica applies a constant

Table 3
ECC Sensor and Background Current Uncertainty (ΔI_B) Applied for Each SHADOZ Site

Site	ECC	ΔI_B (μA) (fraction of missing or $>0.05 \mu\text{A}$)
Ascension Is., UK	SPC	0.02 (7.2%)
	ENSCI (92 records)	0.03 (7.4%)
Costa Rica	ENSCI	0.03 (0%)
Hanoi, Vietnam	ENSCI	0.03 (16.0%)
	SPC (15 records)	0.02 (1.6%)
Irene, S. Africa	SPC	0.02 (44.7%)
Kuala Lumpur, Malaysia	SPC (1998 to November 2014)	0.02 (86.2%)
	ENSCI (December 2014–2016)	0.03 (14.0%)
La Réunion, France	ENSCI	0.03 (7.4%)
	SPC (40 records)	0.02 (0%)
Nairobi, Kenya	ENSCI	0.03 (74.2%)
Natal, Brazil	SPC	0.02 (0.5%)
	ENSCI (104 records)	0.03 (0.2%)

Note. ECC = electrochemical concentration cell; SHADOZ = Southern Hemisphere Additional OZonesondes; SPC = Science Pump Corporation.

0.02 μA to all profiles, and Natal recharges the cells with new solutions to determine the final background prior to launch.

In equation (2) I_B is subtracted from I_m in the denominator, and thus, its value has a significant impact on that term, particularly in the troposphere where I_m in the tropical troposphere is typically less than 1 μA and I_B can be as high as 0.05 μA . Notably, the tropopause is a region of very low ozone, usually <1 mPa in the tropics. Where P_{O_3} measures less than 1 mPa, I_m can approach I_B , thus increasing the dominance of the $(\Delta I_M)^2 + (\Delta I_B)^2 / (I_M - I_B)^2$ term (abbreviated to $\Delta I_B / \Delta I_M$ hereafter) in equation (2).

4.2. Conversion Efficiency Uncertainty, $\Delta \eta_C / \eta_C$

The conversion efficiency, η_C , comprises two parts: (1) the absorption efficiency (α_{O_3}) from the gas to liquid phase in the sensing solution and (2) the stoichiometry of the $\text{O}_3:\text{l}_2$ relationship ($S_{\text{O}_3:\text{l}_2}$) which is assumed to be 1:1. Interferences with this one-to-one relationship can arise from the buffering of the solution (Johnson et al., 2002; Vömel & Diaz, 2010). Setting the α_{O_3} equal to 1 applies to cases where the volume of the cathode solution is 3.0 cm^3 . For SHADOZ sites, such as Ascension, Natal, Irene, and Réunion, that use a 2.5 cm^3 cathode volume, we use the following equations in Smit12 to calculate α_{O_3} , as a function of pressure, P : $\alpha_{\text{O}_3}(P) = 1.0044 - (4.4 \times 10^{-5}) \times P$ for $100 \text{ hPa} < P < 1,050 \text{ hPa}$, and $\alpha_{\text{O}_3}(P) = 1.0$ for $P \leq 100 \text{ hPa}$. In the ozone partial pressure equation (equation (1)), the conversion efficiency is assumed to be unity and is typically excluded from the ozone equation. However, the uncertainty of this unity assumption does contribute to the overall ozone uncertainty. These are constant unitless values for a cathode solution volume of 3.0 cm^3 . Thus, $\Delta \eta_C / \eta_C$ can be expressed as follows:

$$\frac{\Delta \eta_C}{\eta_C} = \sqrt{\left(\frac{\Delta \alpha_{\text{O}_3}}{\alpha_{\text{O}_3}}\right)^2 + \left(\frac{\Delta S_{\text{O}_3:\text{l}_2}}{S_{\text{O}_3:\text{l}_2}}\right)^2}, \quad (3)$$

where $\alpha_{\text{O}_3} = 1.0$, $\Delta \alpha_{\text{O}_3} = \pm 0.01$, $S_{\text{O}_3:\text{l}_2} = 1.0$, and $\Delta S_{\text{O}_3:\text{l}_2} = \pm 0.03$. The $\Delta \alpha_{\text{O}_3}$ value is ± 0.01 for both cathode volumes (Gaussian 1σ value taken from Davies et al., 2003). This is a simplistic approach because $S_{\text{O}_3:\text{l}_2}$ will increase over the course of the sonde flight due to slow side reactions involving the phosphate pH buffers (Davies et al., 2000; Johnson et al., 2002). However, at present changes in $S_{\text{O}_3:\text{l}_2}$, while recognized, are poorly understood and require further research. Where a transfer function is applied, 0.05 is added to the $\Delta S_{\text{O}_3:\text{l}_2}$ term (Smit12).

4.3. Pump Flow Rate Uncertainty at the Ground, $\Delta \Phi_p / \Phi_p$

A common procedure in the ECC conditioning is the use of a soap bubble flow meter method to measure the volumetric flow rate of the pump, Φ_p (mL/s). However, calculating $\Delta \Phi_p / \Phi_p$ is not straightforward because not

Table 4
 Values of ΔC_{PH} and $\Delta\Phi_{corrected}/\Phi_{corrected}$ for Each SHADOZ Site

Site	ΔC_{PH}	$\Delta\Phi_{corrected}/\Phi_{corrected}$	
Ascension Is., UK	± 0.014	± 0.02	
Costa Rica	± 0.013	± 0.02 and $\pm 0.04^a$	
Hanoi, Vietnam	± 0.010	± 0.02 and $\pm 0.04^a$	
Irene, S. Africa	± 0.015	± 0.04	$T = 25 \pm 5^\circ\text{C}$, $\text{RH} = 50 \pm 25\%$, $P = 850 \text{ hPa}^b$
Kuala Lumpur, Malaysia	± 0.013	± 0.04	$T = 25 \pm 5^\circ\text{C}$, $\text{RH} = 50 \pm 25\%$, $P = 1,000 \text{ hPa}^b$
La Réunion, France	± 0.014	± 0.02	
Nairobi, Kenya	± 0.010	± 0.02 and $\pm 0.04^a$	
Natal, Brazil	± 0.014	± 0.02	

^aUsed for profiles where lab P - T - U is unknown. ^bBased on Witte17 tropical climatology.

all quantities are known. Witte17 applied correction formulae found in section 8.4 of Smit12 that compensate for the evaporation of the soap bubble solution and rely on the saturated water vapor pressure under ambient P - T - U conditions (C_{PH}). There is a second correction that takes into account the temperature difference between the internal pump base temperature and the ambient room temperature, $C_{PL} = (T_{\text{pump}} - T_{\text{lab}})/T_{\text{lab}}$. Our calculations reveal ΔC_{PL} values $\ll 1$, and thus, we exclude this term that has a negligible impact on the correction of Φ_p and its uncertainty. The final equation to calculate $\Delta\Phi_p/\Phi_p$ can then be expressed as follows:

$$\frac{\Delta\Phi_p}{\Phi_p} = \sqrt{\left(\frac{\Delta\Phi_{\text{reprocessed}}}{\Phi_{\text{reprocessed}}}\right)^2 + (\Delta C_{PH})^2}, \quad (4)$$

where $\Delta\Phi_{\text{reprocessed}}/\Phi_{\text{reprocessed}} = \pm 0.02$ and ΔC_{PH} is based on the minimum and maximum C_{PH} values in each site's data set, that is, $\pm(C_{PH,\text{High}} - C_{PH,\text{Low}})/2$ (Smit12). Here we assume that the uncertainties in the reprocessed flow rates are within the uncertainty of the original measured quantities. Table 4 summarizes the ΔC_{PH} and $\Delta\Phi_{\text{reprocessed}}/\Phi_{\text{reprocessed}}$ computed for the eight sites. At Irene and Kuala Lumpur, for which lab P - T - U are not documented, we double the $\Delta\Phi_{\text{reprocessed}}/\Phi_{\text{reprocessed}}$ term to ± 0.04 and calculate ΔC_{PH} using the Witte17 tropical climatology of $T_{\text{lab}} = 25 \pm 5^\circ\text{C}$, $\text{RH}_{\text{lab}} = 50 \pm 25\%$, and $P_{\text{sfc}} = \text{mean pressure surface at launch}$. In the case of Costa Rica, Hanoi, and Nairobi for which a fraction of their data sets are missing lab P - T - U , we use the mean C_{PH} for flights with known lab conditions. ΔC_{PH} remains the same since the mean does not affect the min/max range of C_{PH} values and $\Delta\Phi_{\text{reprocessed}}/\Phi_{\text{reprocessed}} = \pm 0.04$.

4.4. Pump Flow Efficiency Uncertainty, $\Delta\psi_p/\psi_p$

The pump flow rate, Φ_p , measured during the conditioning procedures is approximately constant up to 100 hPa and decreases steadily to the top of the atmosphere due to instrumental degradation at low pressures (Komhyr, 1986; Komhyr et al., 1995). From equation (1), the pump flow efficiency, Ψ_p , is based on empirically derived pump correction factors (PCF) that take into account the efficiency loss in Φ_p as a function of pressure. This study follows the WMO/GAW report recommendations and applies the Komhyr (1986) PCF for sondes launched with an SPC sensor and Komhyr et al. (1995) PCF for ENSCI sensors. These PCF compensate for the effect of the buffer that creates side reactions in the solution (Johnson et al., 2002). The exception is at Hanoi for which almost half the data set uses a NOAA sensing solution recipe of 2% unbuffered KI. The unique formula requires its own PCF due to the lack of the buffer in solution (Johnson et al., 2002). These three pump flow efficiencies and their $\pm 1\sigma$ uncertainties ($\pm\Delta\psi_p$) are listed in Table 5. We can interpret the effect of the buffer at low pressures as the difference in Ψ_p between the Johnson et al. (2002) and both Komhyr look-up tables: the difference ranges from 3% at 100 hPa to 15% at 5 hPa. The Deshler et al. (2017) transfer function that corrects for solution changes between the 0.5% half buffer and 1.0% full buffer solution intrinsically takes into account the effect of the buffer, and therefore, its overall uncertainty of $\pm 5\%$ carries within it the uncertainty in the buffering.

The O3S-DQA panel has recommended a revised table of PCF $\pm \Delta\psi_p$ based on the average of combined laboratory calibration experiments conducted by NOAA/ESRL/GMD, University of Wyoming, and Japan Meteorological Agency. This study uses the revised $\pm\Delta\psi_p$ values included in Table 5. Note that these

Table 5
Ozonesonde Pump Flow Efficiencies, Ψ_p , With 1σ Uncertainties ($\pm\Delta\Psi_p$)

Pressure (hPa)	Komhyr (1986) $\Psi_p \pm \Delta\Psi_p$	Komhyr et al. (1995) $\Psi_p \pm \Delta\Psi_p$	Johnson et al. (2002) ^a $\Psi_p \pm \Delta\Psi_p$	O3S-DQA panel $\pm \Delta\Psi_p$
Sfc-100	1.000	1.000	1.1.000	
100	0.993 \pm 0.005	0.993 \pm 0.005	0.967 \pm 0.011	\pm 0.010
50	0.982 \pm 0.006	0.982 \pm 0.005	0.950 \pm 0.012	\pm 0.012
30	0.978 \pm 0.008	0.972 \pm 0.008	0.935 \pm 0.012	\pm 0.014
20	0.969 \pm 0.009	0.961 \pm 0.012	0.919 \pm 0.014	\pm 0.017
10	0.948 \pm 0.010	0.938 \pm 0.023	0.876 \pm 0.020	\pm 0.022
7	0.935 \pm 0.012	0.920 \pm 0.024	0.842 \pm 0.025	\pm 0.028
5	0.916 \pm 0.014	0.890 \pm 0.025	0.803 \pm 0.032	\pm 0.037

Note. Ψ_p are taken from the World Meteorological Organization/Global Atmospheric Watch Report. Ozone Sonde Data Quality Assessment panel revised $\pm\Delta\Psi_p$ values are computed to replace Komhyr (1986) and Komhyr et al. (1995) $\pm\Delta\Psi_p$.
^aAverage of ENSCI-Z model pump efficiency calibrations.

uncertainties are similar to those of Johnson et al. (2002) for a 2% unbuffered SST; thus, we expect $\Delta\Psi_p/\Psi_p$ values to be similar across the eight study sites. The revised $\pm\Delta\Psi_p$ values are considered to be a more realistic representation of the PCF uncertainties and are better quantified, being based on hundreds of profiles (refer to Table 2 in Johnson et al., 2002). Refer to Table 3 for a summary of ECC sensor used at each site.

4.5. Pump Temperature Uncertainty, $\Delta T_p/T_p$

All SHADOZ sites use either an ENSCI-Z or SPC-6A model that measures the pump temperature internally. These measurements are considered to be a close approximation to the “true” pump temperature that is measured in the vicinity of the moving piston, T_{piston} (Smit12). Witte17 reprocessing includes a correction that accounts for the temperature difference between T_{piston} and the internal pump temperature. For SHADOZ data sets, the $\Delta T_p/T_p$ equation is expanded to

$$\frac{\Delta T_p}{T_p} = \sqrt{\left(\frac{\Delta T_{\text{reprocessed}}}{T_{\text{reprocessed}}}\right)^2 + \left(\frac{\Delta T_{\text{piston}}}{T_{\text{reprocessed}}}\right)^2}, \quad (5)$$

where both $\Delta T_{\text{reprocessed}}$ and ΔT_{piston} are $\pm 0.5^\circ\text{K}$ (Smit12). As with the reprocessed flow rates, we assume that the uncertainties in the reprocessed pump temperatures are within the uncertainties of the original measured values. Witte17 replaced missing pump temperature profile data at Kuala Lumpur (1998–2005) and Irene (1998–2006) with climatological values. For these data sets the 1σ values of the climatology (range between 3° and 4°C) are used to compute $\Delta T_{\text{reprocessed}}$ (see Figure A1 in Witte17).

5. Ozone Uncertainty Estimates in SHADOZ Profiles, ΔP_{O_3}

For the eight reprocessed SHADOZ sites, we compute ΔP_{O_3} and the individual uncertainty terms defined in equation (2) for each profile. The overall profile average of the uncertainty terms for each site is also shown in Figure 4. The vertical resolution is 50 m. We show the impact of applying a 5% transfer function uncertainty to the Ascension, Natal, Nairobi, Réunion, and Hanoi data sets (Figures 4a–4e, right panels, dashed black line). Note that the overall mean ΔP_{O_3} profile will lie between the solid black line, which is the data set for which no transfer function is applied, and the dashed line. From Figure 4 (right panels) we observe the following:

1. All sites show a peak in ΔP_{O_3} around the tropopause region (15 ± 3 km) due to I_B , as nicely illustrated in Figure 1 in Vömel and Diaz (2010).
2. The uncertainty in the background and O_3 current term ($\Delta I_B/\Delta I_M$, red) dominates the overall uncertainty of P_{O_3} in the troposphere. Where transfer functions have been applied (Figures 4a–4e), the conversion efficiency term (orange) becomes a significant contributor to the uncertainty.
3. The uncertainties in the conversion efficiency (orange) and flow rate (blue) terms dominate the overall uncertainty of P_{O_3} in the stratosphere.

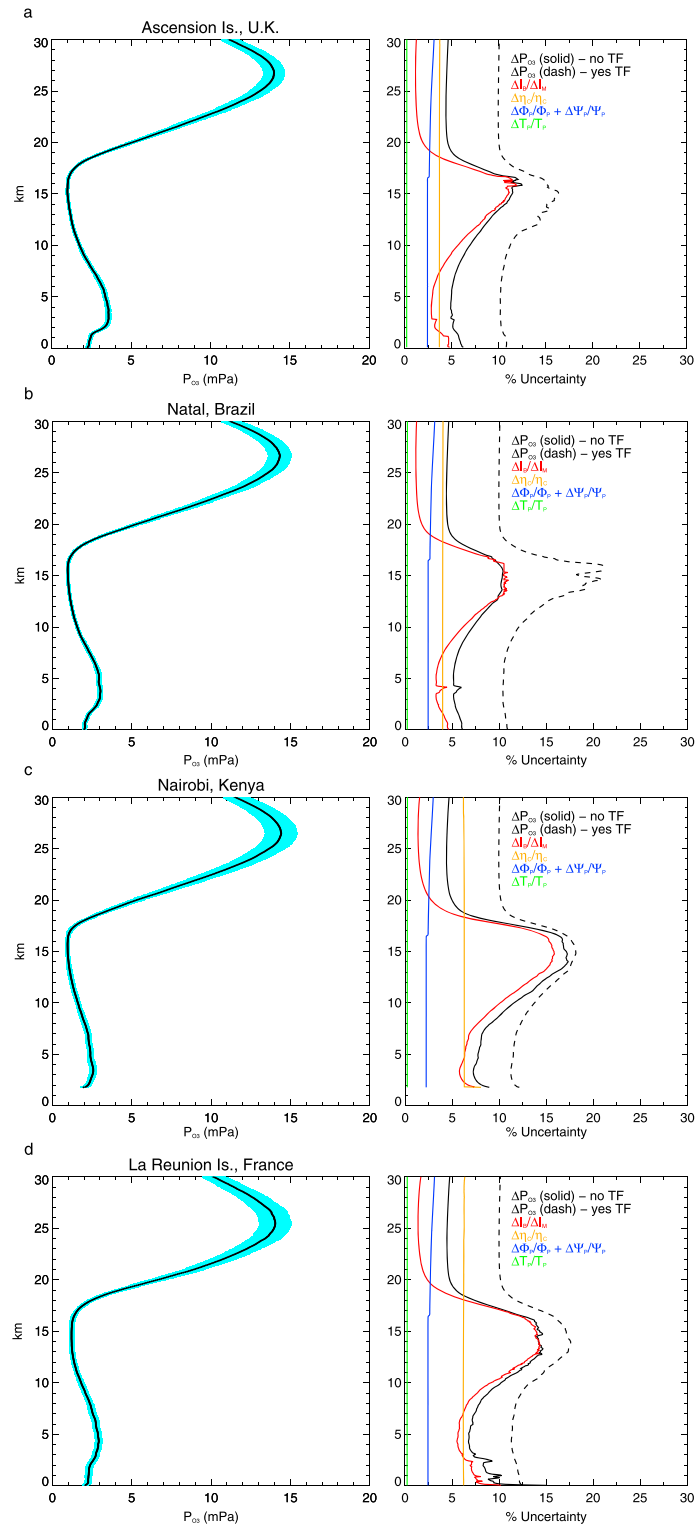


Figure 4. Left panels show the average profile of O_3 partial pressure (P_{O_3}) in mPa (black) and \pm uncertainty estimates (cyan) for the eight SHADOZ sites. Right panels are the individual uncertainty contributions to the total O_3 uncertainty (ΔP_{O_3}) (black) in percent. Solid black lines in the right-hand side panels for (a)–(d) are ΔP_{O_3} average profiles for which no transfer function was applied. Concurrently, dashed black lines are the average ΔP_{O_3} of profiles where a transfer function is applied. The $\Delta I_B/\Delta M$ term is an abbreviation of the $(\Delta I_M)^2 + (\Delta I_B)^2 / (I_M - I_B)^2$ term in equation (2). Refer to Table 1 for each site’s time period and total number of profiles used to calculate the averages. TF = transfer function

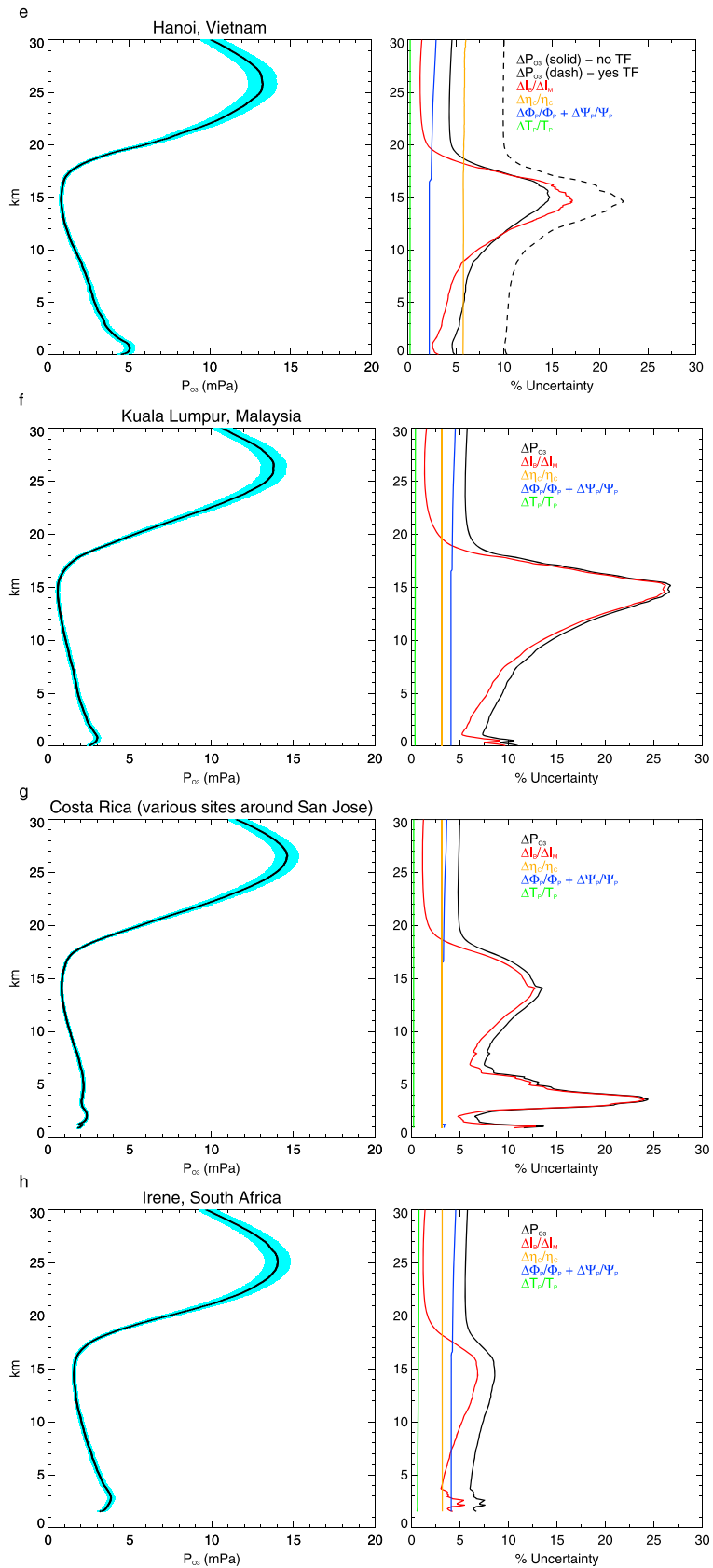


Figure 4. (continued)

4. The application of transfer functions (Figures 4a–4e, purple) has a significant impact on the overall uncertainty of P_{O_3} throughout the profile.
5. The contribution of the pump temperature uncertainty term ($\Delta T_p/T_p$, profiles in green) is minimal, accounting for only a few percent.
6. Overall uncertainties are within 15%. Notable exceptions are at Hanoi (Figure 4e, dashed line) and Kuala Lumpur (Figure 4f) around the tropopause region.
7. Costa Rica (Figure 4g) displays a unique maximum in ΔP_{O_3} and $\Delta I_B/\Delta I_M$ between 2 and 7 km.
8. Irene results show the smallest P_{O_3} uncertainties (less than 7%), relative to the other sites (Figure 5h), while Kuala Lumpur shows the largest values of ΔP_{O_3} .

In the vicinity of the tropopause (15 ± 3 km), P_{O_3} is a minimum (Figure 4, left panels). This is particularly true for Hanoi and Kuala Lumpur data sets that show the lowest mean P_{O_3} relative to the other sites. As the sensor current (I_M) approaches the measured background current value (I_B), the significant increase and dominance in the $\Delta I_B/\Delta I_M$ term indicates that the sensor measurement is approaching its detection limit. Here we define the detection limit where the total uncertainty is equal to the measured signal (~ 0.02 – $0.05 \mu\text{A}$ range). This is particularly true at Nairobi where the $\Delta I_B/\Delta I_M$ uncertainty term matches the transfer function uncertainty between 10 and 18 km (Figure 4c, right panel). Further investigation reveals a prevalence of $I_B > 0.05 \mu\text{A}$ in the Nairobi metadata, which contributes to a doubling of its uncertainty. In contrast, Irene (Figure 4h) shows the highest measurements of P_{O_3} in the same region concurrent with the lowest values in the $\Delta I_B/\Delta I_M$ and ΔP_{O_3} . Irene is located in the very edge of the subtropics and can exhibit midlatitude behavior (i.e., stratospheric fold events) as well as Southern Hemisphere pollution transport (Thompson et al., 2014). Hanoi and Kuala Lumpur are the only Asian tropical sites in the SHADOZ network and due to their geographic proximity to one another likely exhibit similar dynamics and transport features in the UT/LS region (Ogino et al., 2013). Thompson et al. (2012) show that Hanoi and Kuala Lumpur have a similar annual cycle with almost double the O_3 amount at the Hanoi site throughout the troposphere. In fact, values of O_3 at Kuala Lumpur and Hanoi are among the lowest relative to other SHADOZ sites, corroborating the Thompson et al. (2012) result. Both sites show a higher gradient in ΔP_{O_3} and a larger peak $>15\%$, relative to the other sites.

As noted above, there is a second peak in ΔP_{O_3} in the lower troposphere in the Costa Rican data set (Figure 4g). There is a notch in P_{O_3} between 2 and 7 km (left panel) that is due to sulfur dioxide (SO_2) interference from the nearby active Turrialba volcano (Diaz et al., 2012; Morris et al., 2010). Profiles that record near-zero I_M due to volcanic SO_2 plumes are not uncommon in a given year (Diaz et al., 2012). This is seen in the right panel of Figure 4g where the uncertainty in the I_B/I_M term (red) increases significantly as I_M approaches I_B and thus the detection limit of the sensor. The data set reveals that roughly 10% of all profiles have SO_2 interference, with years 2010–2012 being particularly volcanically active. This uncertainty peak is really due to the reduction in the conversion efficiency (η_C), whereby SO_2 interference reduces η_C to 0. This is not captured in the uncertainty discussion because chemical interference is not assumed in equations (1) and (2). The near-zero cell current measurements amplify the ΔP_{O_3} and $\Delta I_B/\Delta I_M$ terms generating values that are much larger than they should be.

Van Malderen et al. (2016) applied corrections to the Uccle data set for high- SO_2 interference in the Brewer-Mast ozonesondes using in situ SO_2 measurements from a nearby site. No corrections were necessary during the ECC sonde era (1996 to present) because SO_2 concentrations had diminished to levels that would not impact the ECC measurements. By filtering the obvious SO_2 layers in the Costa Rican data set, the peak would still be present but not as large or as dominant (Thompson et al., 2010).

Figure 5 presents the uncertainty terms arranged by season. P_{O_3} is shown in the background in silver. Similarly, we find that ΔP_{O_3} is a maximum around the tropopause and is dominated by the $\Delta I_B/\Delta I_M$ term. In particular, in the vicinity of the tropopause region we observe in Figure 5 the following:

1. The overall uncertainty of P_{O_3} is the highest in December–February (DJF) and March–May (MAM) (with or without transfer functions) for sites in the Atlantic and Africa: Ascension, Natal, Nairobi, Réunion, and Irene.
2. The overall uncertainty of P_{O_3} is highest in DJF and September–November (SON) at the two northern tropical Asian sites, Hanoi (Figure 5e) and Kuala Lumpur (Figure 5f), where ΔP_{O_3} peaks are over 20%. ΔP_{O_3} gradients are highest for these two sites.

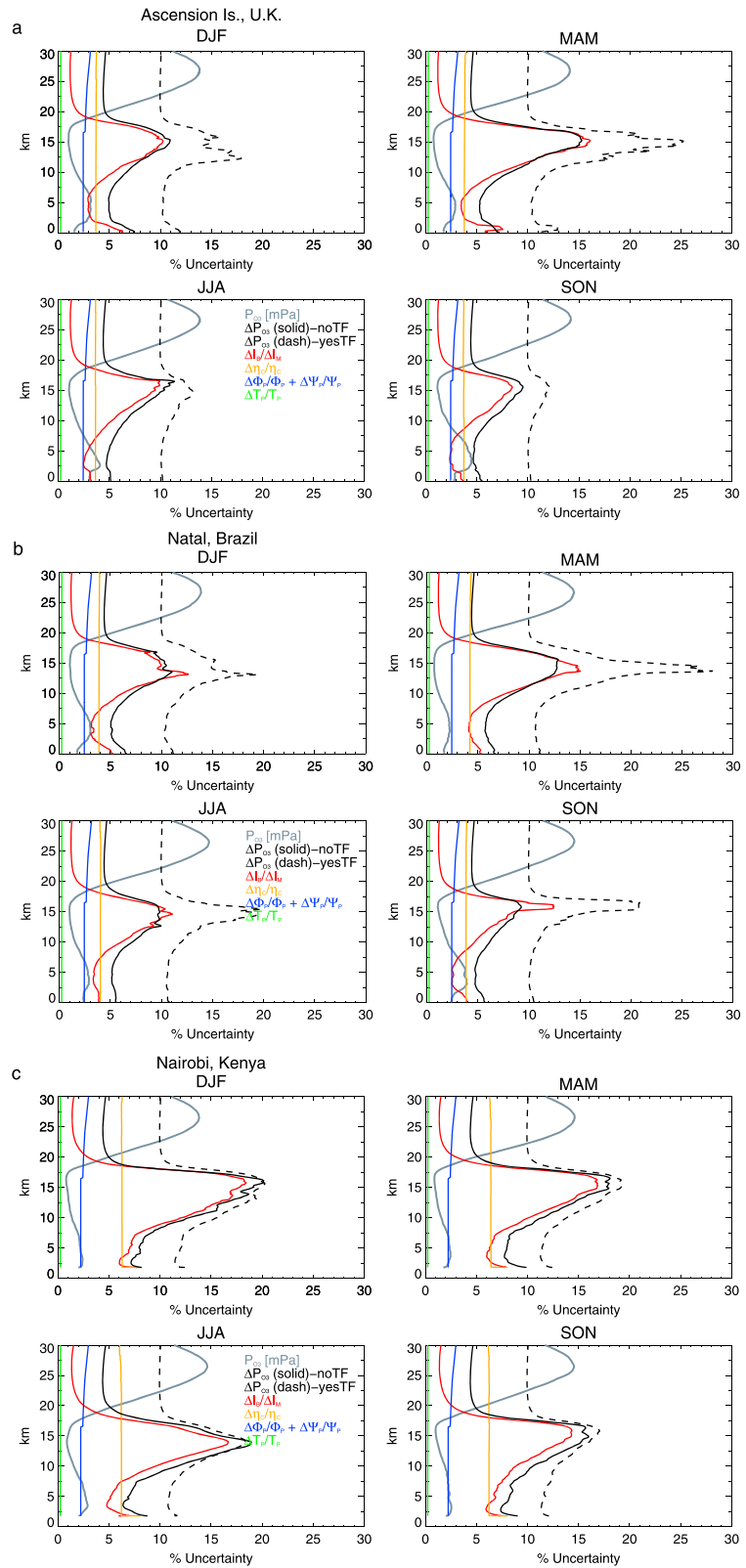


Figure 5. Similar to the right panels in Figure 4 but by season: DJF = December-January-February; MAM = March-April-May; JJA = June-July-August; SON = September-October-November. Average seasonal P_{O_3} is shown in silver in mPa units. Dashed black lines in (a)–(e) are the average ΔP_{O_3} profiles for which a transfer function was applied. The solid black line denotes the average ΔP_{O_3} for profiles that did not apply a transfer function. TF = transfer function.

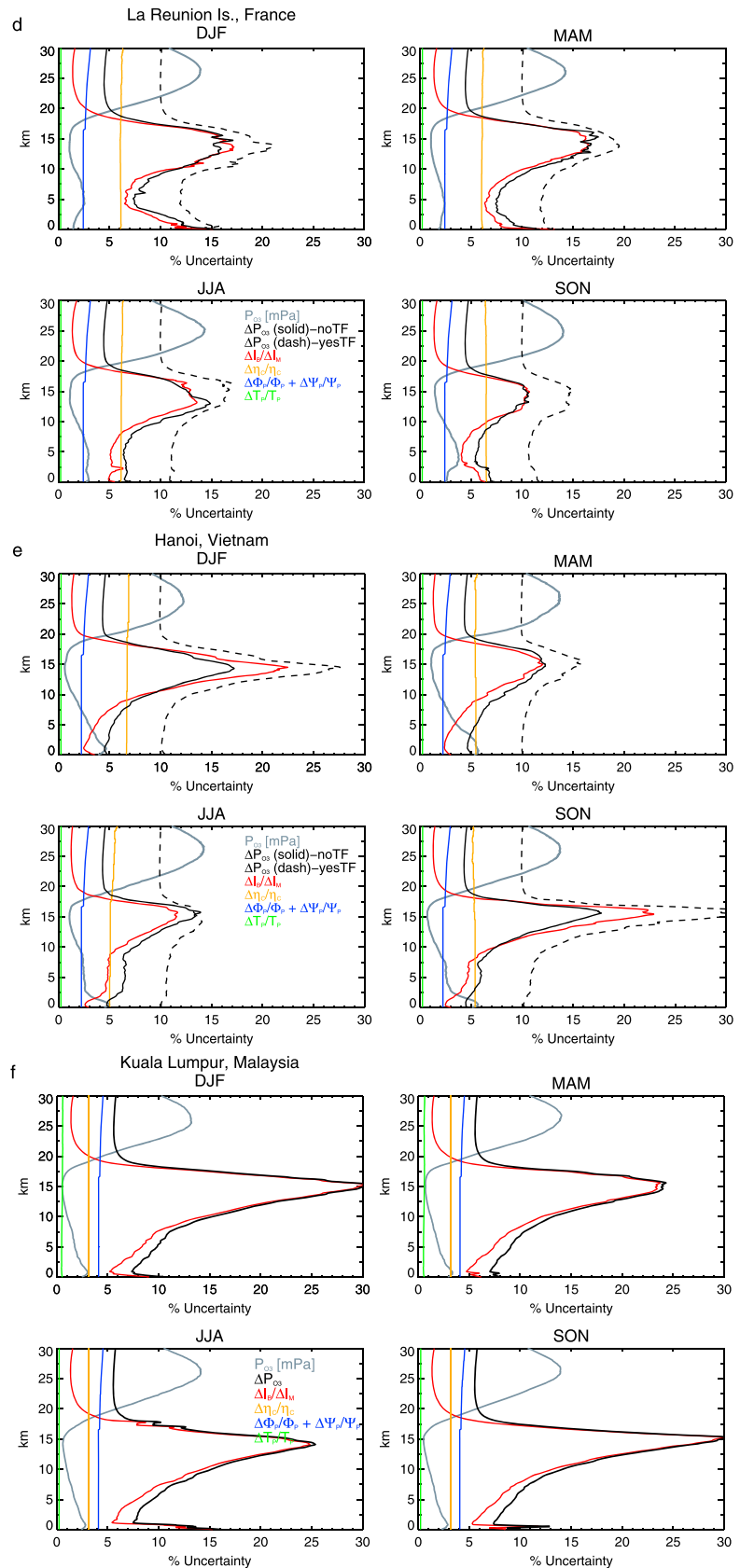


Figure 5. (continued)

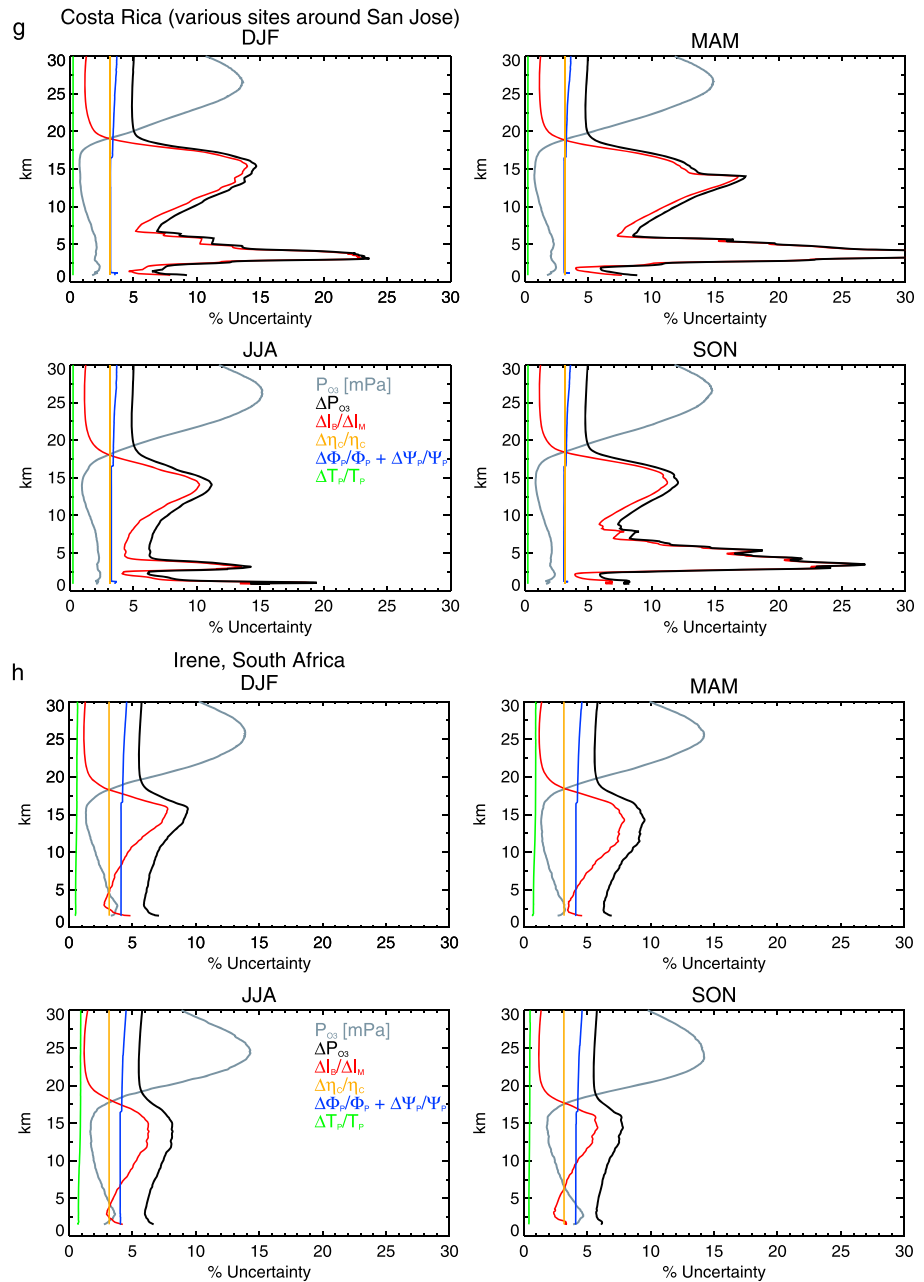


Figure 5. (continued)

3. High ΔP_{O_3} in the Costa Rican data set (Figure 5g) due to volcanic SO_2 interference between 2 and 7 km occurs throughout the year.
4. Relative to other sites, the overall uncertainty of P_{O_3} is smallest at Irene for all seasons (Figure 5h). SON is a minimum (ΔP_{O_3} is less than 8%).

Overall, the $\Delta I_B/\Delta I_M$ term dominates ΔP_{O_3} in the troposphere. The exception is where P_{O_3} measurements are high in the lowermost troposphere (silver profile), such as during the biomass burning season in SON over the Atlantic sites: Ascension (Figure 5a), Natal (Figure 5b) as well as Réunion (Figure 5d), and Irene (Figure 5h) and where the boundary layer is polluted such as at Hanoi (Figure 5e). P_{O_3} is elevated below 5 km for all seasons in the Irene data set (Figure 5h), accounting for the relative minima in $\Delta I_B/\Delta I_M$ (red). For all seasons, Kuala Lumpur (Figure 5f) stands out as having the highest ΔP_{O_3} in the tropopause region, particularly in DJF and

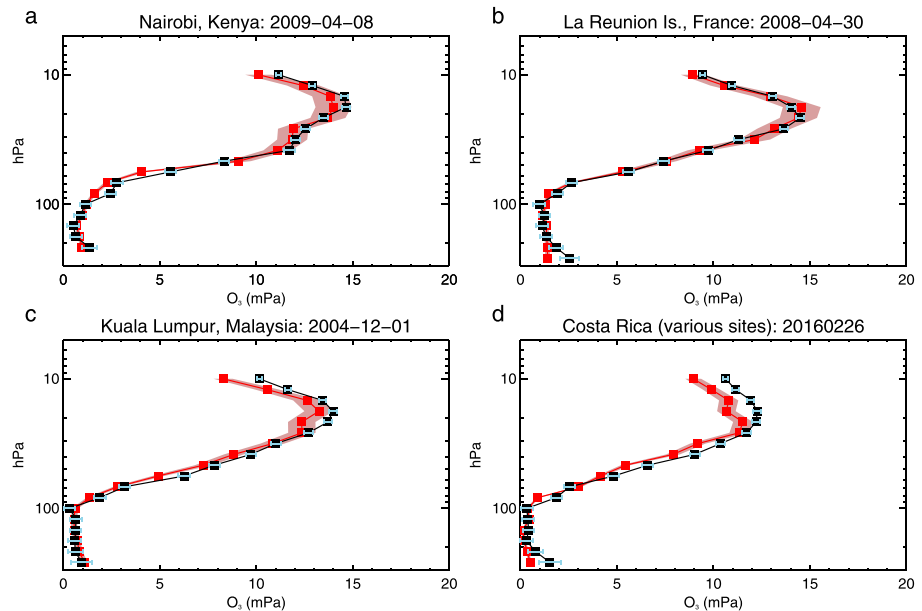


Figure 6. Individual profiles of sonde (red) and Microwave Limb Sounder (MLS) (black) for (a) Nairobi, (b) Réunion, (c) Kuala Lumpur, and (d) Costa Rica. Sonde profiles are matched to the MLS resolution. Red shading denotes the sonde $\pm\Delta P_{O_3}$. Blue bars show the MLS precision.

SON, where uncertainties peak sharply at around 30%, coincident with high $\Delta I_B/\Delta I_M$ (red). Interestingly, the Nairobi data set also shows a relative ΔP_{O_3} maximum in DJF (Figure 5c). Logan et al. (2003) found that ozone concentrations at Nairobi are lowest from December to April from the tropopause to 50 hPa (~20 km). This corroborates the uncertainty results in Figure 5c that show relatively higher ΔP_{O_3} during DJF and MAM where the $\Delta I_B/\Delta I_M$ uncertainty term peaks at 15 ± 3 km.

Examples of applying uncertainties to profile comparisons with satellite can be found in Figure 6. We use Microwave Limb Sounder (MLS) v4.2 overpass profiles (Waters et al., 2006) at pressure levels between 261 and 10 hPa matched to the sonde location (within 200 km and 18 h). The four profile examples represent a cross section of reprocessing techniques. Transfer functions were applied to Nairobi and Réunion profiles (Figures 6a and 6b), Kuala Lumpur profiles incorporated a pump temperature climatology (Witte17) (Figure 6c), and Costa Rica is a site that required very little reprocessing and therefore had a minor impacts to the overall data set, relative to the original v5 data set (Thompson17). One notes patterns of agreement/disagreement on a profile-by-profile basis. Applying sonde uncertainties will provide a robust metric of evaluating the accuracy of current and future generations of satellite O_3 profilers, particularly in the climate sensitive UT/LS region.

How do our uncertainty estimates compare with previous studies? The WCCOS simulation by Smit and ASOPOS (2014) of a tropical profile estimate a similar range of ΔP_{O_3} values that also maximize around the tropopause (up to ~17%) due to the dominance of the $\Delta I_B/\Delta I_M$ term. The main difference is that this term dominates ΔP_{O_3} values throughout the profile, whereas in our study, the flow rate and conversion efficiency uncertainty dominate in the stratosphere. Sterling et al. (2017) present ΔP_{O_3} mean profiles for April and October at Hilo, Hawaii, and American Samoa (two SHADOZ sites), which we interpret as similar to the seasonal plots shown in Figure 5. Here we observe that their range of ΔP_{O_3} values is similar to ours, that is, maxima at the tropopause (>10%) and minima elsewhere in the profile (~5%). Sterling et al. (2017) show a relative minimum (maximum) in MAM (SON), similar to that shown for Hanoi and Kuala Lumpur (Figures 5e and 5f). In comparison, midlatitude and high-latitude ozonesonde profiles from previous studies show that (1) the overall uncertainty is smaller, that is, less than ~10% throughout the profile, (2) the conversion efficiency uncertainty term is a significant contributor to ΔP_{O_3} throughout the profile, and for some sites like Uccle is the dominant uncertainty term, and (3) the impact of the $\Delta I_B/\Delta I_M$ term remains an important contributor to ΔP_{O_3} around the tropopause (*McMurdo uncertainty analysis*; T. Deshler,

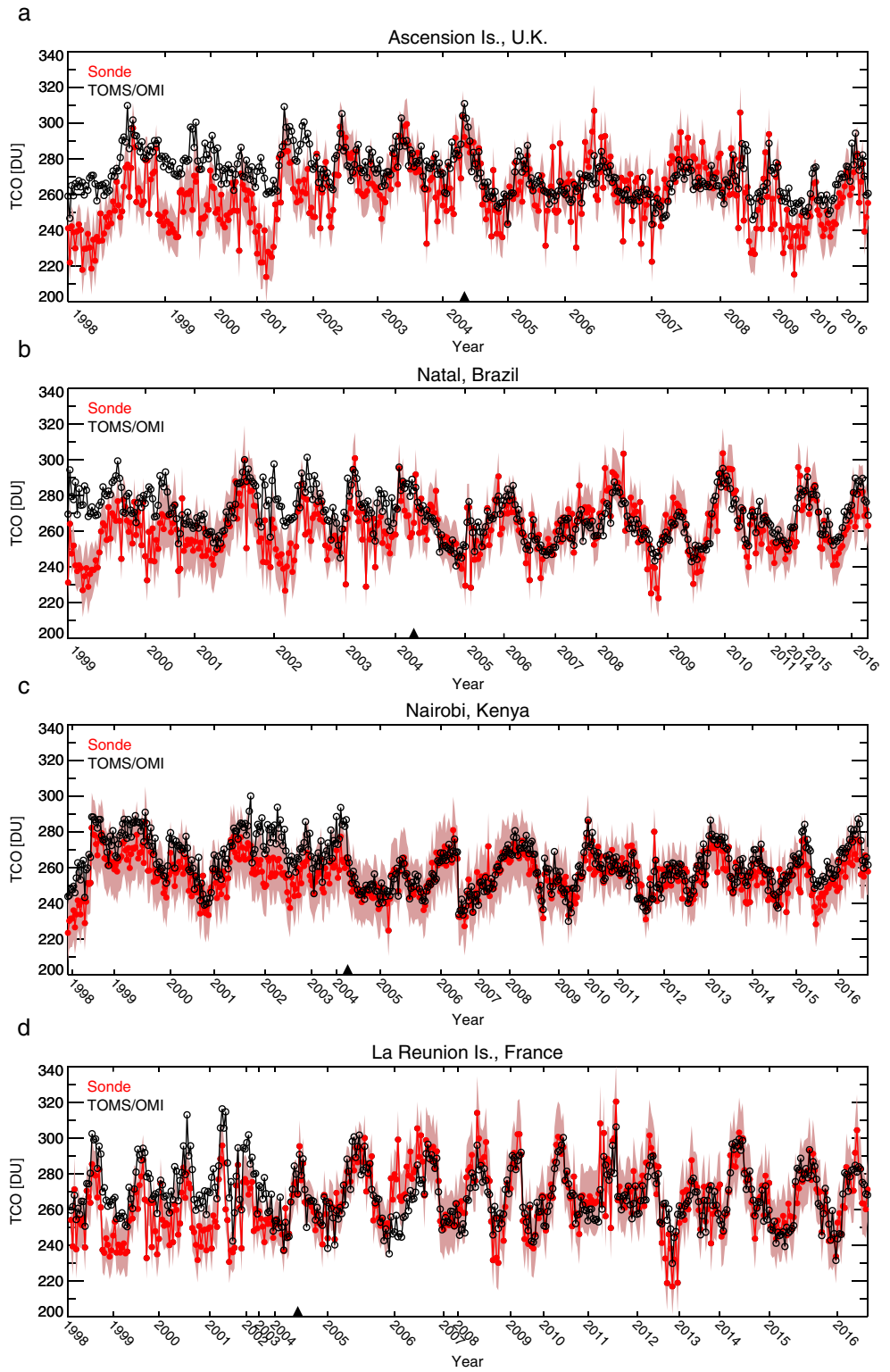


Figure 7. Time series of total column ozone (TCO) of every sonde launch (red) matched to a satellite overpass (Total Ozone Mapping Spectrometer (TOMS) and Ozone Monitoring Instrument (OMI), black). Red shading denotes the sonde $TCO \pm$ uncertainties (ΔTCO). Satellite data sets are filtered for clouds $>60\%$ and distance from the sites location >200 km, then compared to the date of each sonde launch at the given site. The transition from TOMS to OMI overpasses occur in October 2004 and are marked by the black triangle on the x axis. Results are plotted in sequence, accounting for the uneven spacing of the years on the x axis.

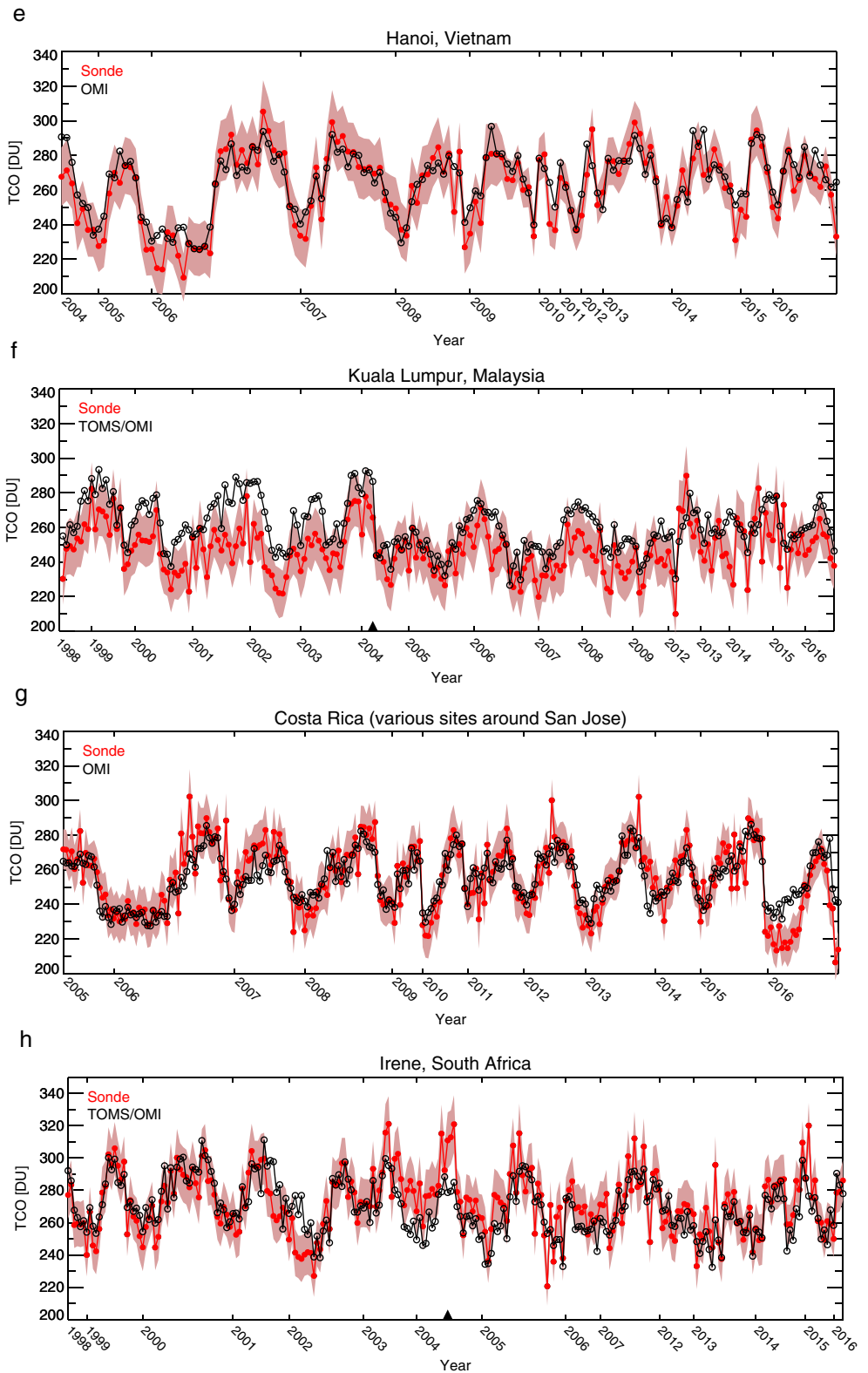


Figure 7. (continued)

Table 6
Mean Sonde TCO (TCO_{sonde}) and TOMS/OMI Satellite Overpasses for Each Site Computed From Data in Figure 7

Satellite period	Sample No.	$TCO_{sonde} \pm \Delta TCO_{sonde}$	Satellite $\pm 1 - \sigma$
Costa Rica (data since 2005)			
OMI	277	255.3 ± 12.6	255.4 ± 14.3
Hanoi (data since September 2004)			
OMI	147	261.7 ± 14.7	264.5 ± 17.6
Ascension			
TOMS/OMI	445	260.4 ± 13.1	271.6 ± 12.3
TOMS	220	259.1 ± 13.4	277.7 ± 10.9
OMI	225	261.6 ± 12.8	265.7 ± 10.6
Irene			
TOMS/OMI	250	272.4 ± 15.0	269.6 ± 16.4
TOMS	122	274.2 ± 15.4	274.7 ± 16.2
OMI	128	270.7 ± 14.7	265.8 ± 15.2
Kuala Lumpur			
TOMS/OMI	215	247.3 ± 14.9	260.6 ± 14.1
TOMS	86	249.7 ± 14.9	267.4 ± 14.5
OMI	129	245.8 ± 14.9	256.1 ± 11.9
La Réunion			
TOMS/OMI	454	264.9 ± 15.8	268.5 ± 15.7
TOMS	129	255.8 ± 13.1	272.4 ± 16.0
OMI	325	268.6 ± 16.8	266.9 ± 15.2
Nairobi			
TOMS/OMI	570	255.8 ± 15.9	262.1 ± 13.3
TOMS	199	257.7 ± 17.3	270.2 ± 12.9
OMI	371	254.7 ± 15.1	257.7 ± 11.3
Natal			
TOMS/OMI	436	261.1 ± 13.4	269.2 ± 12.7
TOMS	188	258.9 ± 14.4	275.3 ± 11.2
OMI	248	262.8 ± 12.7	264.5 ± 11.9

Note. Units are in Dobson units. ΔTCO_{sonde} is the mean uncertainty. The $\pm \sigma$ standard deviations are computed for the TOMS and OMI periods separately and the combined TOMS/OMI period. Note that matched satellite overpasses are filtered for clouds >60% and distance from the site location >200 km. OMI overpasses start in October 2004. TCO = total column ozone; OMI = Ozone Monitoring Instrument; TOMS = Total Ozone Mapping Spectrometer.

personal communication, 2015; Smit & ASOPOS, 2014; Sterling et al., 2017; Tarasick et al., 2016; Van Malderen et al., 2016).

6. Ozone Uncertainty Estimates in SHADOZ TCO

6.1. Calculating Total Column O₃ Uncertainty, ΔTCO

Profiles of O₃ partial pressure are integrated up to 10 hPa to generate integrated column amounts of O₃ in DU. To capture the O₃ maximum, we exclude profiles that burst at pressures less than 15 hPa. We limit our column integration to 10 hPa based on the Witte17 recommendation. They found that the accuracy of O₃ profile measurements below 10 hPa with respect to MLS did not improve even after reprocessing and remains highly variable and suspect. To compute the sonde TCO values, the McPeters and Labow (2012) O₃ climatology is used to extract the O₃ amount from the sonde's balloon burst to the TOA. This is termed the "O₃ residual" in this paper ($P_{O_3 \text{residual}}$). McPeters and Labow (2012) derived an O₃ climatology based on combining MLS (2004–2010) and ozonesonde (1988–2010) data sets. The climatology is a look-up table of monthly averaged ozone profiles from the surface to the TOA for 10° latitude zones. Previous studies have used the same sonde TCO formulation when analyzing SHADOZ data (Witte17; Thompson17, and references therein).

As a first approach, we compute sonde TCO uncertainties (ΔTCO) by integrating the O_3 partial pressure uncertainties (ΔP_{O_3}) generated from equation (2) and adding the uncertainty in the O_3 residual amount ($\Delta P_{\text{O}_3\text{residual}}$); that is, $\Delta\text{TCO} = \Delta P_{\text{O}_3} + \Delta P_{\text{O}_3\text{residual}}$. For example, if $\text{TCO} = \int_{\text{sfc}}^{\text{BurstLevel}} P_{\text{O}_3}(z) dz + P_{\text{O}_3\text{residual}}$, then $\Delta\text{TCO} = \int_{\text{sfc}}^{\text{BurstLevel}} \Delta P_{\text{O}_3}(z) dz + \int_{\text{BurstLevel}}^{\text{TOA}} \Delta P_{\text{O}_3\text{residual}}(z) dz$. To generate an equivalent look-up table of $\Delta P_{\text{O}_3\text{residual}}$ uncertainties, we use the table of monthly averaged 1σ O_3 mixing ratios taken from McPeters and Labow (2012) and convert to DU following Ziemke et al. (2001) formulation (refer to equation (1) in that study). Thus, we can extract the $\Delta P_{\text{O}_3\text{residual}}$ at the sonde's lower limit altitude (15–10 hPa) in the same way. Look-up tables of $\Delta P_{\text{O}_3\text{residual}}$ for the tropical latitude zones between 15 and 8 hPa are found in Table S1 in the supporting information. For example, for a 10 hPa pressure burst the $P_{\text{O}_3\text{residual}} \pm \Delta P_{\text{O}_3\text{residual}}$ range is (48–58 DU) \pm (1.5–3 DU). We recognize that this approach may be an oversimplified calculation and hope that this study motivates future evaluation and refinement of this method.

6.2. Comparisons With TOMS and OMI Overpasses

As a reference, sonde TCO $\pm \Delta\text{TCO}$ are compared with satellite overpass TCO from Earth Probe TOMS (V8.6 provided by the TOMS/OMI science team, McPeters et al., 1998) and the OMI V3, (Levelt et al., 2006). TOMS and OMI have a local equator crossing time of 11:16 and 13:30, respectively. Sonde launch times (local) vary between 10:00 and 1500. From Figure 7 we observe that, overall, the overpasses are within the uncertainty of the sondes (red shading). Table 6 gives a summary of each site's mean TCO $\pm \Delta\text{TCO}$ and includes the individual and combined TOMS and OMI mean overpass TCO $\pm 1\sigma$ standard deviation values. Mean $\pm \Delta\text{TCO}$ at the eight sites are comparable to $\pm 1\sigma$ from the satellite overpass data. The exception is at Kuala Lumpur where there are many cases in which TOMS and OMI are outside the uncertainty range (Figure 7f, red shading). This amounts to a 5% offset overall. Given that there is limited metadata information for this site (Witte17), we speculate that the offset seems to be consistent with the use of a nonstandard sensing solution formulae, similar to what was discovered at Réunion and Nairobi.

Ascension, Natal, Nairobi, and Réunion reprocessed data sets (Figures 7a–7d) show significantly larger O_3 offsets relative to the TOMS period (1998–2004/09) compared to the OMI time series. The cause for this persistent low bias is unknown and cannot be explained by reprocessing and system differences alone. Whereas Ascension and Natal used the Lockheed Martin Sippican radiosonde/ozonesonde system during that entire period, Réunion used a Vaisala RS80 system switching to a Modem system (2007 to present). The low bias relative to TOMS is not readily apparent in the Irene reprocessed data set (Figure 7h) for which a Vaisala system is used and is the only subtropical site in this study with a data record extending back to 1998. However, Kuala Lumpur, which also used a Vaisala system during the TOMS period, show a much larger sonde-TOMS difference compared to sonde-OMI in the time series (Figure 7f). Further investigation reveals a discontinuity between the TOMS and OMI time series at all sites, and in particular, at Kuala Lumpur (Figure 7f, TOMS overpasses in black). Table 6 includes the mean TCO for the individual TOMS and OMI periods at each site and reveals an almost 10 DU difference between averaged TOMS and OMI TCO with TOMS measuring higher than OMI (right column). For example, at Kuala Lumpur TOMS measures 11 DU higher TCO than OMI (TOMS = 267.3 ± 14.5 DU versus OMI = 256.1 ± 11.9 DU from Table 6). The Kuala Lumpur sonde data set does not show a similarly large discontinuity in its TCO time series between the TOMS and OMI periods (249.7 ± 14.9 DU during the TOMS period versus 245.8 ± 14.9 DU during the OMI period). Interestingly, the agreement between Irene sonde TCO and TOMS is excellent; however, the agreement between TOMS and OMI is offset by 9 DU (Table 6, right column). From Table 6, excluding Irene, the average difference between sonde and TOMS is 16 DU and is significantly higher than the 4 DU difference computed for the difference between sonde and OMI. This discontinuity is likely due to a change in the TOMS processing algorithm and is currently being investigated by the TOMS/OMI science team. This TOMS to OMI TCO discontinuity is not apparent in Witte17 and Thompson17 that used an older version 8 TOMS overpass data set taken from the Aura Validation Data Center (<https://avdc.gsfc.nasa.gov>).

The sonde low bias at Costa Rica (Figure 7g) observed at the end of 2015 points to a change in the ENSCI instrument. Other SHADOZ sites (Samoa, Fiji, and Hilo) that use ENSCI during that period also exhibit a low bias relative to satellite overpasses (Thompson17). The MLS/sonde profile comparison in Figure 6d highlights the significant underestimate in the sonde (red) observed after 2015 in Figure 7g and in

Thompson17. The sonde underestimate appears above 50 hPa and is typical of sonde/MLS profile comparisons after 2015.

7. Summary

For the first time, uncertainty estimates in profile and total column O₃ are computed using reprocessed data from eight SHADOZ sites. As a first approach, we also create a look-up table of monthly averaged O₃ residual uncertainties to compute the uncertainty in sonde TCO. Variations in reprocessing procedures at each site require a tailored approach when calculating uncertainties. For example, missing metadata such as background current and flow rate measurements require climatological values specific to each site and thus a doubling of the uncertainty; not all sites require a transfer function.

A significant fraction of the Réunion and Nairobi data sets have been homogenized with the use of transfer functions, 63% and 64%, respectively. We demonstrate the efficacy and success of applying the Deshler et al. (2017) formulae to generate consistent and stable reference data sets for trends analysis and satellite validation. Biases are reduced (Réunion = 0.7% and Nairobi = -1.5%) and preliminary agreement with OMI is within 5%.

Overall, O₃ profile uncertainties are less than 15% with a persistent maximum in the vicinity of the tropopause where O₃ values are low and approach the detection limits of the ECC sensor. Here the background and sensor current uncertainties dominate, as well as in the troposphere. Stations for which transfer functions have been applied show a significant contribution from the conversion efficiency uncertainty. In the stratosphere, the conversion efficiency and flow rate uncertainty terms dominate. We observe a unique second peak in the free troposphere in the Costa Rican data set due to SO₂ interference from volcanic plumes. Seasonally, uncertainties are a maximum in MAM when O₃ values are the lowest. Exceptions are at Hanoi and Kuala Lumpur, the only two northern tropical Asian sites in the SHADOZ network. Both display the highest uncertainties in DJF and SON, indicating unique meteorology relative to the other sites. The Irene data set displays the lowest O₃ uncertainties, among the eight sites.

Overall, TCO uncertainties are less than 15 DU and represent ~5–6% of the TCO, consistent with the stated uncertainties in Thomson17. TOMS/OMI overpass comparisons are within the sonde TCO uncertainties with the exception of Kuala Lumpur for which the sondes are low biased by about 5%. The persistent offset mimics that found in the Réunion and Nairobi data sets for which an error in the sensing solution formulae was found. However, due to the current lack of verification in the metadata, we can only speculate on the offset. In addition, there is a discontinuity between the latest TOMS v8.6 and OMI TCO overpasses at almost all sites. TOMS measures higher TCO than OMI on the order of 10 DU that is not readily apparent in the older version 8 overpasses used in Witte17 and Thompson17 studies.

The advantage of doing a detailed uncertainty analysis is that it reveals areas of the measurements where we can refine operational procedures to reduce the uncertainty and where additional research is needed to improve the basics of this measurement. (1) To improve O₃ measurements in the UT/LS in SHADOZ, we need to improve our understanding of the background current. This means that the actual measurement as done at present needs to be better quality controlled, that is, using a high-quality zero ozone air filter, lapsed time of measurement after reconditioning, and consistency of procedures. For example, the large spread of SHADOZ I_B observed in Witte17 (range is 0.01 μ A–0.12 μ A) is a result of varying procedures and not a property of the sonde instrument. It also means that the definition of the background current needs to be better agreed upon. (2) To reduce the uncertainty of the middle stratospheric measurements, that is, region of high O₃, we need to better characterize the pump efficiency and conversion efficiency at low pressures. This is also the reason that transfer functions become pressure dependent at lower pressures. (3) Careful and complete metadata collection of lab P - T - U during preparation will reduce the uncertainty of the RH correction (ΔC_{PH}), and careful metadata collection of the O₃ destruct filter methods being used during the background measurements will help in estimating the quality of that measurement.

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