

## **Reviewer #1:**

Review of Davis et al.

“This paper is comprehensive and well written. It provides a lot of information on the ozone and water vapour fields in various state-of-the-art reanalyses, including quantification of their accuracy, usefulness of the datasets, and possible improvements. As such, I expect this paper to be useful to the atmospheric sciences community, and likely to be highly cited. I recommend publication in ACP subject to the authors paying attention to the specific comments below.”

***We thank the reviewer for their comments, and indeed hope that this work will be a useful resource for the community.***

Specific comments

“P. 3,

L. 7: I suggest you indicate here what you will discuss in each section.”

***Done, at the end of section 1.***

“P. 4

L. 16: Do you need “notable”?”

***We removed this word.***

“L. 27: It would be helpful to the reader to identify the old and updated forecast model and data assimilation system.”

***We’ve added some information on the new and old systems to the sentence. Also, the details of the differences between CFSR and earlier NCEP reanalyses are discussed in Saha et al. 2010, and we’ve now made this more clear in the sentence.***

“P. 10

L. 71: was -> were. “

***Done***

“P. 27

L. 17: Maybe I am wrong, but I understood that there was a debate on the sign of trends in stratospheric water vapour during the late 1990s and early 2000s, with discrepancies between balloon and satellite measurements. Perhaps this has been resolved. Maybe the authors could mention this when they mention the work of Randel et al. (2006).”

***As the reviewer notes, there is a discrepancy between decadal trend estimates from balloon measurements and those from satellites. However, the Randel et al. 2006 paper and the text in question by the reviewer are referring to the drop in water vapor around the year 2000, and the measurements from both balloon and satellites are broadly in agreement on the existence of this drop. We have intentionally chosen to not include reanalysis trends in WV or O<sub>3</sub> in this paper.***

“P. 41

Table 1: If you are using US spelling, it should be “analyzed”. Same elsewhere.”

***Done. The entire document is now in US spelling.***

“P. 43

Fig. 1: What do the colours represent? Same for Fig. 2.”

***The colors denote the reanalyses. We have made this clearer by coloring the reanalysis text labels with the corresponding color.***

“P. 45

Fig. 3: It would be helpful if the authors could identify in the caption what the red/blue colours indicate, e.g., positive/negative values. Same for other figures.”

***Done***

“P. 47

Fig. 5: It would be helpful if the authors identified in the caption the colours referring to the reanalyses. Same for Fig. 6, 13, 14.”

***The captions are already quite verbose, and we don't think it is necessary to do this since in all of these figures a legend is given showing the reanalyses and their colors.***

## **Reviewer #2**

“This kind of paper is hard to review. It provides a summary of ozone and water vapour information in current reanalyses data sets, with an emphasis on the stratosphere. I guess the main conclusion of the paper was known before it was written: use the ozone and water vapour data with care and do not use for trend studies. However, it is nice to have some of the issues illustrated with figures in a comparative way. Therefore, there is no reason why the paper should not be published, however some explanations might require some clarifications. I will detail my questions below:”

“P3, 122: I do not know what this statement means. All systems try to model the micro- physics of water as good as they can, yet the results differ, because small differences in the treatment of water can have large effects?”

***This sentence was confusing and has been re-worded. The point of the sentence was to state that SWV is not well constrained by observations because no observations of SWV are assimilated, and that hence SWV is highly controlled by the differing physical representations of SWV. It now reads “Because stratospheric water vapour data are not directly assimilated, the concentration of water vapour in the stratosphere is highly variable amongst the reanalyses and is strongly affected by their representation of processes controlling it.”***

“P2, l30: This sentence is confusing. It tries to make two points in one sentence: Heating rate calculations and photochemistry. Which ozone is used when and where?”

***(We assume the reviewer means P3 here, not P2) The point of this sentence was to note the large variety of ways ozone is represented in reanalyses. The information pertaining to the reviewers question is all summarized in Table 1. We realized that Table 1 was not referenced until later in the manuscript, so we've added reference to Table 1 here, and have also changed the wording to make our point more clear.***

“P6, l3: This relates to the comment regarding P2, l30. I guess a clear discussion in the beginning would be fine. Alternatively, a corresponding sentence for each system. Which ozone and water vapour is used in the radiation (heating rates) and what is done for the chemistry (actinic fluxes, if required).

P7, l9: See above.”

***As noted above, this information is contained in Table 1. It is also discussed separately for each reanalysis.***

“P7, l34: was should read has.”

***Fixed***

“P9, l6: I am not sure why the stratospheric temperature bias changes the humidity product. Ice clouds in Antarctica.”

***In general, temperatures in the TTL affect how much water vapor enters the stratosphere. We have changed the wording here to make this point clear.***

“P11, l2: This sentence is not very clear. Which mean? What tendency? (For people in the know it will be clear, but . . .)”

***We agree with the reviewer that this sentence wasn't very clear. We've changed the sentence to be clearer about the potential impact of changes made in MERRA-2 on water vapor. The sentence now reads:***

***“The main innovation in MERRA-2 that could impact water vapor is the introduction of additional global constraints that ensure continuity of water mass in the atmosphere (Takacs et al., 2016).”***

“P11, l28: If I understand correctly, profiles (ozone and water vapour) are processed as described a few lines above on pressure levels and a common grid. However, TCO is calculated from model level data. Why not use the “ready made” products? Do some systems not provide their columns? How do you deal with orography?”

***We misspoke in this paragraph. We computed the monthly means ourselves “from the 6-hourly TCO fields”, not “from the 6 hourly model level data” as stated in the paper. We’ve re-worded this paragraph to make this clear. Orography is taken into account in using model level data because the lowest level is the surface, regardless of orographic height. As long as pressure is properly registered on the model levels (which it is in our JRA-25 calculations), using model level data is the most accurate way to compute total column ozone.***

“P12/13: I appreciate that the authors would like to compare the reanalyses systems with another data source. However the data used is neither independent nor in a fundamental form. Instead, merged data sets are utilised. Nothing wrong with this, but presumably other equally valid products exist and I am not sure why the data sets mentioned have been chosen . . . (I am also not very happy with the use of multi-instrument means without an appreciate of the spread, when comparing to the reanalyses systems.)”

***We appreciate the reviewers concern that some of the comparisons in this paper are to assimilated (non-independent) observations. However, we note that this is only true for ozone, as all of our comparisons to water vapor data are to independent measurements. Also, for ozone we have carefully noted instances where the comparisons are not to independent data sources. We chose SPARC DI and SWOOSH data because we are familiar with these data sets. Other merged data sets (e.g., GOZCARDS) could be used, but based on outside work assessing the merged data sets, we have no reason to believe that using a different data set would alter the conclusions here (see, e.g., Tummon et al., ACP, 2015; Hubert et al., AMT, 2016; Harris et al., ACP, 2015). Regarding the spread around the multi instrument mean, the spread is shown in all of the line plots containing the MIM (Figs. 5,6,13,14).***

“Instead of focusing on the text, I will now briefly comment on a small number of figures (my assumption being, that readers will be most interested in the graphical presentation of system differences). Given the large range in water vapour products and the small number of systems that provide it, I will not comment on the water vapour related figures (the figures are a health warning in themselves):

Figure 3 and 4: I struggle to combine the information in both figures. For example JRA- 25: In Figure 3 JRA-25 has a low bias with respect to SBUV everywhere. In Figure 4 JRA-25 has a positive bias from around 100 hPa to just below 10 hPa. Assuming that the largest column contribution stems from this region, I do not understand the consistency of the results. (Maybe I have over-read the explanation in the text . . .)”

***The reviewer has uncovered what turned out to be a major bug in the processing of the JRA-25 data. As noted in the manuscript, the JRA-25 total column ozone data were processed directly from the 6-hourly model level data set. The vertically resolved ozone came ultimately from the pressure level data supplied by JMA and NCAR via CREATE-IP. It turns out that the pressure level***

***data used the incorrect hybrid level model coefficients when converting from model levels to pressure levels, resulting in a downward “shift” of the entire ozone profile, as seen in the old version of Figure 4. This was caused by the use of the model “interface” hybrid coefficients, rather than the correct model “full level” coefficients when converting from the model levels to pressure levels.***

***We have gone back to the original model level data to properly create the monthly mean pressure level data set, and have updated all of the JRA-25 plots in this paper that contain vertically resolved data. The affected figures that have been updated are Figs. 4-6, 8, and 11 (the plots of TCO are correct). We have also corrected the relevant discussion in the manuscript and added text explaining our data processing for JRA-25 ozone.***

“Figure 7: I find this figure hard to understand. Presumably, by using equivalent latitudes, the differences in more than one variable are highlighted. PV will have been derived from very different dynamical cores and afterwards ozone has been mapped to it. Therefore, differences will arise from more than one change in the assimilation system and how PV has been derived (e.g. treatment of temperature, dynamical variables and ozone itself, etc.). Therefore, I am not entirely sure what the message on a global scale is . . . apart from they all look different.”

***The reviewer is correct that the plots showing ozone on isentropic-equivalent latitude coordinates could be affected by differences in both the dynamical and chemical (i.e., ozone) representation among the reanalyses. However, we believe Figure 7 is valuable because it illustrates the sort of analysis that people do in “research” studies, so we should know how derived quantities (such as ozone on EqL-theta coordinates) compare among the reanalyses. Also, the figure quite clearly illustrates some of the problems with ozone in the polar regions. The similarity of the PV contours among the different reanalyses in this region suggests that the problem lies more in the representation of ozone than in the representation of vortex dynamics. We have added text in Section 4.5 to clarify these points and the importance of this figure.***

# Assessment of upper tropospheric and stratospheric water ~~vapour~~vapor and ozone in reanalyses as part of S-RIP

Sean M. Davis<sup>1,2</sup>, Michaela I. Hegglin<sup>3</sup>, Masatomo Fujiwara<sup>4</sup>, Rossana Dragani<sup>5</sup>, Yayoi Harada<sup>6</sup>, Chiaki Kobayashi<sup>6,7</sup>, Craig Long<sup>8</sup>, Gloria L. Manney<sup>9,10</sup>, Eric R. Nash<sup>11</sup>, Gerald L. Potter<sup>12</sup>, Susann Tegtmeier<sup>13</sup>, Tao Wang<sup>14</sup>, Krzysztof Wargan<sup>11,15</sup>,  
5 Jonathon S. Wright<sup>16</sup>

<sup>1</sup>Earth System Research Laboratory, National Oceanic and Atmospheric Administration, Boulder, CO 80305, USA

<sup>2</sup>Cooperative Institute for Research in Environmental Sciences (CIRES), University of Colorado at Boulder, Boulder, CO 80309, USA

<sup>3</sup>Department of Meteorology, University of Reading, Reading, RG6 6BX, UK

10 <sup>4</sup>Faculty of Environmental Earth Science, Hokkaido University, Sapporo 060-0810, Japan

<sup>5</sup>European Centre for Medium-Range Weather Forecasts, Reading, RG2 9AX, UK

<sup>6</sup>Japan Meteorological Agency, Tokyo, 100-8122, Japan

<sup>7</sup>Climate Research Department, Meteorological Research Institute, JMA, Tsukuba, 305-0052, Japan

15 <sup>8</sup>Climate Prediction Center, National Centers for Environmental Prediction, National Oceanic and Atmospheric Administration, College Park, MD 20740, USA

<sup>9</sup>NorthWest Research Associates, Socorro, NM 87801, USA

<sup>10</sup>Department of Physics, New Mexico Institute of Mining and Technology, Socorro, NM 87801, USA

<sup>11</sup>Science Systems and Applications, Inc., Lanham, Maryland 20706, USA

<sup>12</sup>NASA Center for Climate Simulation, Code 606.2, NASA Goddard Space Flight Center, Greenbelt MD 20771, USA

20 <sup>13</sup>GEOMAR Helmholtz Centre for Ocean Research Kiel, Kiel, 24105, Germany

<sup>14</sup>NASA Jet Propulsion Laboratory/California Institute of Technology, Pasadena, CA 91109, USA

<sup>15</sup>Global Modeling and Assimilation Office, Code 610.1, NASA Goddard Space Flight Center, Greenbelt, MD 20771, USA

<sup>16</sup>Department of Earth System Science, Tsinghua University, Beijing, 100084, China

25

*Correspondence to:* Sean M. Davis (sean.m.davis@noaa.gov)

**Abstract.** Reanalysis data sets are widely used to understand atmospheric processes and past variability, and are often used to stand in as “observations” for comparisons with climate model output. Because of the central role of water ~~vapour~~vapor (WV) and ozone (O<sub>3</sub>) in climate change, it is important to understand how accurately and consistently these species are represented in existing global reanalyses. In this paper, we present the results of WV and O<sub>3</sub> intercomparisons that have been performed as part of the SPARC (Stratosphere-troposphere Processes and their Role in Climate) Reanalysis Intercomparison Project (S-RIP). The comparisons cover a range of timescales and evaluate both inter-reanalysis and observation-reanalysis differences. We also provide a systematic documentation of the treatment of WV and O<sub>3</sub> in current reanalyses to aid future research and guide the interpretation of differences amongst reanalysis fields.

The assimilation of total column ozone (TCO) observations in newer reanalyses results in realistic representations of TCO in reanalyses except when data coverage is lacking, such as during polar night. The vertical distribution of ozone is also relatively well represented in the stratosphere in reanalyses, particularly given the relatively weak constraints on ozone vertical structure provided by most assimilated observations and the simplistic representations of ozone photochemical processes in most of the reanalysis forecast models. However, significant biases in the vertical distribution of ozone are found in the upper troposphere and lower stratosphere in all reanalyses.

In contrast to O<sub>3</sub>, reanalysis estimates of stratospheric WV are not directly constrained by assimilated data. Observations of atmospheric humidity are typically used only in the troposphere, below a specified vertical level at or near the tropopause. The fidelity of reanalysis stratospheric WV products is therefore mainly dependent on the reanalyses’ representation of the physical drivers that influence stratospheric WV, such as temperatures in the tropical tropopause layer, methane oxidation, and the stratospheric overturning circulation. The lack of assimilated observations and known deficiencies in the representation of stratospheric transport in reanalyses result in much poorer agreement amongst observational and reanalysis estimates of stratospheric WV. Hence, stratospheric WV products from the current generation of reanalyses should generally not be used in scientific studies.

## 1 Introduction

Ozone and water ~~vapour~~vapor are trace gases of fundamental importance to the radiative budget of the stratosphere. Because of their impact on stratospheric temperatures, winds, and the circulation (e.g., Dee et al., 2011), ozone and water ~~vapour~~vapor are represented as prognostic variables in almost all current reanalysis systems. However, the degree of sophistication to which ozone and water ~~vapour~~vapor fields and their variability are represented depends on the reanalysis system, which observations it assimilates, which microphysical and chemical parameterizations it includes, and how those parameterizations affect the trace gas distributions. The accuracy and consistency of analysis and reanalysis ozone and water ~~vapour~~vapor fields in the upper troposphere and stratosphere has only been addressed for a limited subset of diagnostics and analysis/reanalysis systems by a few studies (e.g., Dessler and Davis, 2010; Jiang et al., 2015; Geer et al., 2006; Thornton et al., 2009).

As part of the SPARC (Stratosphere-troposphere Processes and their Role in Climate) Reanalysis Intercomparison Project (S-RIP), we conducted the first comprehensive assessment of how realistically and consistently reanalyses represent water ~~vapour~~vapor and ozone in the upper troposphere and stratosphere. In particular, the goals of this paper are to (1) provide a comprehensive overview of how ozone and water ~~vapour~~vapor are treated in reanalyses, (2) evaluate the accuracy of ozone and water ~~vapour~~vapor in reanalyses against both assimilated and independent (non-assimilated) observations, and (3) provide guidance to the community regarding the proper usage and limitations of reanalysis ozone and water ~~vapour~~vapor fields in the upper troposphere and stratosphere.

Towards this end, in the next section, we provide a description of how ozone and water vapor are treated by the various reanalyses to provide context for the comparisons presented in the rest of the paper. We then provide an overview of the observational data sets used for comparison to reanalyses in Section 3. Sections 4 and 5 contain the evaluations of reanalysis ozone and water vapor, respectively. In the final section, we conclude with a summary of the salient findings and guidance regarding the overall utility and limitations of reanalysis ozone and water vapor.

## 2 Description of ozone and water ~~vapour~~vapor in reanalyses

In this section, we provide information on how ozone and water ~~vapour~~vapor are represented in reanalyses. The information compiled here expands on that provided by Fujiwara et al. (2017), who presented a comprehensive overview of the reanalysis systems and their assimilated observations, including a basic discussion of the treatment of ozone and water ~~vapour~~vapor.

In most reanalyses, ozone and water ~~vapour~~vapor are prognostic variables that are affected by the assimilated observations (see Tables 1 and 2 for an overview of key aspects of these fields). The assimilated observations affecting the water ~~vapour~~vapor fields in reanalyses include some combination of radiosonde humidity profiles, GNSS-RO bending angles, and either radiances or retrievals from satellite microwave and infrared sounders such as TOVS, ATOVS, and SSM/I (see Appendix A for a list of all abbreviations). These observational data affect the reanalysis water ~~vapour~~vapor fields in the lower atmosphere, but radiosonde humidity data are not assimilated above a specified level in the upper troposphere (typically between 300 hPa and 100 hPa, see Table 2). Even though radiosonde humidity data may not be assimilated above a certain level, analysis increments are possible at higher levels unless the vertical correlations of the background errors are set to zero. Where relevant, this cutoff level above which analysis increments are disallowed has been noted in Table 2.

Because stratospheric water ~~vapour~~vapor data are not directly assimilated, the treatment of water ~~vapour~~vapor in the stratosphere is highly variable amongst the reanalyses. For the modern reanalyses, the concentration of water ~~vapour~~vapor entering the stratosphere is typically controlled by transport and dehydration processes occurring in the forecast model, primarily in the tropical tropopause layer (TTL). Higher in the stratosphere, chemical production of water ~~vapour~~vapor through methane oxidation is parameterized in some reanalyses, while others use a simple relaxation of the simulated water ~~vapour~~vapor field to an observed climatology.



As with water ~~vapour~~vapor, the treatment of ozone is quite different from reanalysis to reanalysis. The ozone treatment in reanalyses (see Table 1) ranges from ~~emitting prognostic~~using prescribed ozone and ~~using~~a climatology in the radiation calculations (NCEP R1/R2), to using a fully prognostic field with parameterized photochemistry that interacts with the radiation calculation (CFSR, ERA-40, ERA-I, MERRA, MERRA-2), to assimilating ozone with an offline chemical transport model for use in the forecast model radiation calculation (JRA-25, JRA-55).

The primary ozone observations assimilated by reanalyses are satellite nadir UV backscatter-based retrievals of vertically integrated total column ozone (TCO) or broad vertically weighted averages (e.g., SBUV data). These data come from a variety of satellites that have flown since the late 1970's, and reanalyses vary widely in what subset of the available data they assimilate (Figs 1-2). Some further differences exist amongst the reanalyses in their usage of different data versions from the same satellite instrument, and from different applications of data quality control and filtering. These differences in usage of input data may affect the reanalysis ozone fields.

Additional observation types using spectral ranges outside of the UV (namely microwave and IR) and exploiting different viewing geometries (such as limb-sounding) have been used, particularly by the newest reanalyses (ERA-I, MERRA-2). The assimilation of additional data, particularly vertically resolved data, should improve the quality of the ozone in reanalyses. However, the assimilation of new data sets could introduce sudden changes in the reanalysis ozone fields, and these transition times should be considered carefully when deriving or ~~analysing~~analyzing long-term trends.

## 2.1 NCEP-NCAR (R1) and NCEP-DOE (R2)

Neither NCEP-NCAR (R1) nor NCEP-DOE (R2) assimilates ozone data (Kalnay et al., 1996; Kanamitsu et al., 2002; Kistler et al., 2001). A climatology of ozone was used for radiation calculations.

Humidity information from satellites is not assimilated in R1 and R2 (Ebisuzaki and Zhang, 2011). In general, the treatment of water ~~vapour~~vapor is similar in R1 and R2, with only a few differences. One major difference is that humidity is not output above 300 hPa in R1, whereas it is output up to 10 hPa in R2. Another difference is that only relative humidity is output in R2, whereas in R1 both specific humidity and relative humidity are output. It is worth noting that in R1, specific humidity is a diagnostics variable, computed from relative humidity and temperature. Several fixes and changes were made in the treatment of clouds in R2, and these result in R2 being ~20% drier than R1 in the tropics at 300 hPa (Kanamitsu et al., 2002). As the focus here is on upper levels, we do not assess humidity fields from R1 or R2. It is worth noting that R1 shows negative long-term humidity trends between 500 and 300 hPa (Paltridge et al., 2009); however, these negative trends appear to reflect suspect radiosonde measurements at these levels and are not found in other reanalyses or satellite data (Dessler and Davis, 2010).

## 2.2 CFSR

The Climate Forecast System Reanalysis (CFSR) is a newer NCEP product following the NCEP R1 and R2 reanalyses but with numerous improvements (see Saha et al., 2010 for details), including an updated forecast model and data assimilation

system. CFSR was originally provided through the end of 2009, but output from the same analysis system was extended through the end of 2010 before transitioning to the CFSv2 analysis system starting in January 2011 (Saha et al., 2014). Because CFSv2 was intended as a continuation of CFSR, in this paper we refer to both CFSR (i.e., CFSRv1) and CFSv2 as CFSR. However, the system changeover did result in a discontinuity in the water ~~vapour~~vapor fields that is addressed later in this paper.

CFSR treats ozone as a prognostic variable that is analyzed and transported by the forecast model. The CFSR forecast model uses analyzed ozone data for radiation calculations. In the forecast model, ozone chemistry is parameterized using production and loss terms generated by the NRL CHEM2D-OPP (McCormack et al., 2006). These production and loss rates are provided as monthly mean zonal means, and are a function of local ozone concentration. The rates do not include the coefficients for temperature and overhead ozone column provided by McCormack et al. (2006), nor heterogeneous chemistry, although late 20<sup>th</sup> century levels of CFCs are used indirectly because CHEM2D-OPP is based on the CHEM2D middle atmospheric photochemical transport model, which includes ODS levels representative of the late twentieth century.

CFSR assimilates version-8 SBUV profile and TCO retrievals (Flynn et al., 2009) from *Nimbus-7* and SBUV/2 profiles and TCO retrievals from *NOAA-9*, *-11*, *-14*, *-16*, *-17*, *-18*, and eventually *NOAA-19* (Figs. 1-2). The ozone layer and TCO values assimilated by CFSR have not been adjusted to account for biases from one satellite to the next, although the use of SBUV version 8 is expected to minimize satellite-to-satellite differences. Despite the fact that CFSR assimilates TCO retrievals and SBUV ozone profiles, differences have been found between CFSR and SBUV(/2) ozone profile data (Saha et al., 2010). Most of these differences are located above 10 hPa, and appear to result from observational background errors that were set too high in the CFSR upper stratosphere by between a factor of 2 (at 10 hPa) and a factor of 60 (at 0.2 hPa). Because of this, assimilated SBUV(/2) ozone layer observations do not alter the CFSR first guess for pressures less than 10 hPa, and the model first guess is used instead. The observational background errors were fixed for CFSv2, starting in 2011.

Water ~~vapour~~vapor is treated prognostically in CFSR. There are several assimilated observation types that influence the analysis humidity fields in the troposphere, including GNSS-RO bending angles, radiosondes, and satellite radiances. However, as radiosonde humidity data is only assimilated at 250 hPa and greater pressures, there are no specific observations that constrain humidity in the stratosphere. Stratospheric humidity in CFSR is hence primarily governed by physical processes and parameterizations in the model, including dehydration within the TTL. The treatment of water ~~vapour~~vapor in the model can lead to negative water ~~vapour~~vapor values around and above the tropopause. These negative values are replaced by small positive values of 0.1 parts per million by volume (ppmv) for the radiation calculations, but are retained in the analysis products. CFSR does not include a parameterization of methane oxidation.

### 2.3 ERA-40

The ERA-40 forecast model included prognostic ozone and a parameterization of photochemical sources and sinks of ozone, as described by Dethof and Hólm (2004). This parameterization of ozone production/loss rates is an updated version

of the one proposed by Cariolle and Deque (1986, hereinafter CD86). In CD86, the net ozone production rate is parameterized as a function of the perturbation (relative to climatology) of the local ozone concentration, the local temperature, and the column ozone overhead. Compared to the CD86 formulation, the ozone parameterization in ERA-40 includes an additional term representing heterogeneous chemistry. This loss term scales with the product of the local ozone concentration and the square of the equivalent chlorine concentration, and is only turned on at temperatures below 195 K. The climatologies and coefficients used in the parameterization are derived from a photochemical model and vary by latitude, pressure, and month. The prescribed chlorine loading varies from year to year, from ~700 parts per trillion (ppt) in 1950 to ~3400 ppt in the 1990s. Instead of the CD86 ozone photochemical equilibrium values, ERA-40 made use of the Fortuin and Langematz (1995) ozone climatology.

The prognostic ozone was not used in the radiation calculations, which instead assumed the climatological ozone distribution reported by Fortuin and Langematz (1995). This choice was motivated by concerns that ozone–temperature feedbacks would degrade the temperature analysis if the assimilated ozone observations were of poorer quality than the temperature observations (Dethof and Hólm, 2004).

ERA-40 assimilated TOMS TCO and SBUV layer ozone retrievals from the end of 1978 onward (Figs. 1-2; See also Table 1, Dethof and Hólm, 2004; Poli, 2010). No ozonesonde measurements were assimilated, and no ozone data at all were assimilated before 1978. Ozone data prior to 1978 are thus primarily products of the photochemical parameterization. In addition, no ozone data were assimilated during 1989 and 1990 because the execution of the first ERA-40 stream (1989–2002) was started before the ozone assimilation scheme was implemented. Ozone background errors were also changed, such that the period January 1991 to October 1996 used different background errors than the rest of ERA-40 (Dethof and Hólm, 2004).

ERA-40 water ~~vapour~~vapor products below the diagnosed tropopause are substantially affected by assimilated observations. Three main periods can be identified (Uppala et al., 2005): until 1973, ERA-40 used only conventional in situ surface and radiosonde measurements; from 1973, satellite radiances from VTPR (1973–1978) and the TOVS instruments MSU, SSU, and HIRS (1978–onwards) were used in addition to these conventional data sources; from 1987, 1D-Var retrievals of TCWV from SSM/I radiances were added to the assimilation. Radiosonde humidity measurements were generally used at pressures greater than 300 hPa. No adjustments to the humidity field due to data assimilation were made in ERA-40 above the diagnosed tropopause. Thus, stratospheric water ~~vapour~~vapor in ERA-40 reflects TTL dehydration, transport, and methane oxidation. The latter was included via a simple stratospheric parameterization, in which WV was gradually relaxed to 6 ppmv at the stratopause (Untch et al., 1998). This relaxation was later found to produce too low WV concentrations at the stratopause as it was based on earlier studies when atmospheric methane levels were lower (Uppala et al., 2005). ERA-40 stratospheric humidity has also been shown to be too low overall, due primarily to a cold bias in TTL temperatures caused by an excessively strong Brewer–Dobson circulation (Oikonomou and O'Neill, 2006).

## 2.4 ERA-Interim

The treatment of ozone and water ~~vapour~~vapor in ERA-Interim is very similar to that in ERA-40. Notable differences include additional assimilated datasets and an improved treatment of water ~~vapour~~vapor in the upper troposphere and lower stratosphere (UTLS). Descriptions of the ozone system and assessments of its quality have been provided by Dee et al. (2011) and Dee et al. (2011).

As with ERA-40, total ozone from TOMS (Jan 1979–Nov 1989; Jun 1990–Dec 1994; Jun 1996–Dec 2001) and ozone layer averages from SBUV (1979–present) are assimilated (Figs. 1–2). ERA-Interim also assimilates TCO from OMI (Jun 2008–Jan 2009, Mar 2009–present) and SCIAMACHY (Jan 2003–Dec 2008), and ozone profiles from GOME (Jan 1996–Dec 2002), MIPAS (Jan 2003–Mar 2004), and MLS (Jan–Nov 2008, Jun 2009–present). A change in the assimilation of SBUV ozone profiles was implemented in January 2008. Before January 2008, assimilated SBUV profiles were low vertical resolution products derived over six vertical layers (0.1–1 hPa, 1–2 hPa, 2–4 hPa, 4–8 hPa, 8–16 hPa and 16 hPa–surface) from NOAA version 6 (v6) retrievals. These data were replaced by native 21-vertical-level SBUV profiles from v8 retrievals. The assimilation of ozone profile retrievals from Aura MLS started in 2008 (Fig. 2) using the reprocessed v2.2 MLS retrievals and carried on with the near-real-time v3 product from June 2009 onwards.

The ozone forecast model used in ERA-Interim has the same basic formulation as that used in ERA-40 but some aspects of the parameterization have been upgraded substantially, especially the regression coefficients. An account of the changes is provided by Cariolle and Teyssédre (2007). As in ERA-40, the radiation scheme in ERA-Interim does not use the prognostic ozone field.

A preliminary assessment of the temperature and wind fields revealed unrealistic temperature and horizontal wind increments generated near the stratopause by the 4D-Var assimilation scheme in an attempt to accommodate large local adjustments in ozone concentrations (Dee, 2008; Dee et al., 2011). As an ozone bias correction was not available in ERA-Interim to limit the detrimental effect of ozone assimilation on temperature and wind fields, the sensitivity of the latter to ozone changes was switched off in ERA-Interim. This change affected the period from 1 February 1996 onwards and the ten years from 1979 through 1988 that were run at a later stage.

Through December 1995, ERA-Interim ozone analyses perform better than their ERA-40 counterparts with respect to independent ozone observations in the upper troposphere and lower stratosphere, but perform slightly worse on average in the middle stratosphere (Dee et al., 2011). The assimilation of GOME ozone profiles (Jan 1996–Dec 2002) improves the agreement between ERA-Interim analyses and independent data, such that ERA-Interim outperforms ERA-40 throughout the atmosphere (including the middle stratosphere) from January 1996 through the end of ERA-40 in September 2002 (Dee et al., 2011).

The ERA-Interim humidity analysis is substantially modified from that in ERA-40 due to changes in both model physics and assimilated observations. A non-linear transformation of the humidity control variable was introduced to make humidity background errors more Gaussian (Uppala et al., 2005; Hólm, 2003; Hólm et al., 2002). This transformation normalizes

relative humidity increments by a factor that depends on background estimates of relative humidity and vertical level. A 1D-Var assimilation of rain-affected radiances over oceans was also added as part of the 4D-Var outer loop (Dee et al., 2011), which helps to constrain the spatial distribution of total column water ~~vapour~~vapor (TCWV). The ERA-Interim humidity analysis also benefits from several changes in the model physics, including changes in the convection scheme that lead to increased convective precipitation (particularly at night), reduced tropical wind errors, and a better representation of the diurnal phasing of precipitation events (Bechtold et al., 2004). The non-convective cloud scheme ~~wh~~as also been updated.

Perhaps of most relevance for humidity in the UTLS, the revised cloud scheme contains a new parameterization that allows supersaturation with respect to ice in the cloud-free portions of grid cells with temperatures less than 250 K (Tompkins et al., 2007). The inclusion of this parameterization results in substantial increases in relative humidity in the upper troposphere and in the stratospheric polar cap relative to ERA-40 (Dee et al., 2011). Methane oxidation in the stratosphere is included via a parameterization like the one used in ERA-40 but with relaxation to 6.8 ppmv at the stratopause (rather than 6 ppmv as in ERA-40), based on an analysis of UARS data by Randel et al. (1998).

As with ERA-40, no adjustments due to data assimilation are applied in the stratosphere (above the diagnosed tropopause). ERA-interim tropospheric humidity is affected by the assimilation of radiosonde humidity measurements, radiances from the TOVS (through 5 Sep 2006) and ATOVS (from Aug 1998) instrument suites, and TCWV retrievals based on rain-affected radiances from SSM/I (from Aug 1987). Recent ERA-Interim humidity analyses may also be affected by the assimilation of GNSS-RO bending angles (from May 2001) and/or AIRS all-sky radiances (from April 2004).

## 2.5 JRA-25 and JRA-55

Ozone observations were not assimilated directly in the JRA-25 and JRA-55 systems (Kobayashi et al., 2015; Onogi et al., 2007). Instead, daily three-dimensional ozone fields were produced separately and provided to the JRA forecast model (i.e. to the radiation scheme). Daily ozone fields in JRA-55 for 1978 and earlier are interpolated in time from a monthly mean climatology for 1980–1984. Daily ozone fields in both systems for 1979 and later are produced using an offline chemistry climate model (MRI-CCM1, Shibata et al., 2005) that assimilated satellite observations of TCO using a nudging scheme. Assimilated TCO retrievals are taken from TOMS on Nimbus-7 and other satellites for the period 1979–2004 and from Aura OMI after the beginning of 2005. Different versions of MRI-CCM1 and different preparations of the ozone fields have been used for JRA-25 and JRA-55. For JRA-25, MRI-CCM1 output were also nudged to climatological ozone vertical profiles to account for a known bias in tropospheric ozone that produces a bias in stratospheric ozone after nudging to observations of total ozone. This procedure produced reasonable peak ozone-layer values in the final ozone product. This vertical-profile nudging was not necessary for JRA-55, which used an updated version of MRI-CCM1. JRA-55 produces improved peak values in vertical ozone profiles relative to JRA-25, as well as a clear ozone ~~quasi~~biennial oscillation (QBO) signature.

As with other modern reanalyses, JRA-25 and JRA-55 humidity fields are affected by the assimilation of radiosonde humidity measurements and satellite radiances. The JRA-25 assimilation ~~analysed~~analyzed the logarithm of specific humidity (Onogi et al., 2007). Stratospheric humidity was dry-biased and generally decreased with time in JRA-25, in part

due to the lack of parameterized methane oxidation. The JRA-25 forecast model radiation calculations assumed a constant value of 2.5 ppmv in the stratosphere. Water ~~vapour~~vapor in the UTLS shows evidence of discontinuities at the start of 1991, which corresponds to the transition between the two major processing streams of JRA-25. Onogi et al. (2007) reported sudden jumps of +0.7 ppmv at 150 hPa and +0.9 ppmv at 100 hPa associated with this transition.

5 The treatment of water vapor in JRA-55 is similar in most respects to that in JRA-25. JRA-55 does not contain a parameterization of methane oxidation. Differences include a change in the upper boundary above which the vertical correlations of humidity background errors are set to zero, preventing spurious analysis increments at higher levels. This boundary is set at 5 hPa in JRA-55, relative to 50 hPa in JRA-25. Forecast model radiation calculations in JRA-55 use an  
10 annual mean climatology of stratospheric water ~~vapour~~vapor derived from UARS HALOE and UARS MLS measurements made during 1991–1997 in the stratosphere, rather than the constant 2.5 ppmv used in JRA-25. The introduction of an improved radiation scheme in JRA-55 greatly reduced lower stratospheric negative temperature biases that were present in JRA-25 during the TOVS period before 1998 (Kobayashi et al., 2015; Fujiwara et al., 2017), which may have beneficial  
15 impacts on JRA-55 stratospheric humidity products by impacting dehydration in the TTL. However, wWater ~~vapour~~vapor concentrations at pressures less than 100 hPa are not provided in the standard pressure-level products of these two reanalyses (although these concentrations are provided in model-level products), and are therefore not evaluated in this paper.

## 2.6 MERRA

Ozone is a prognostic variable in MERRA, and is subjected to assimilation, transport by assimilated winds (more precisely, the odd-oxygen family is the transported species), and parameterized chemistry. The MERRA general circulation model (GCM) uses a simple chemistry scheme that applies monthly zonal mean ozone production and loss rates derived  
20 from a 2-dimensional chemistry model (Stajner et al., 2008). Ozone data assimilated in the reanalysis include partial columns and total ozone (defined as the sum of layer values in a profile) from a series of SBUV instruments (Flynn et al., 2009) on various NOAA platforms (Figs. 1–2). Version 8 of the SBUV retrievals (Flynn, 2007) is used but the native 21 vertical layers are combined into 12 layers (each 5 km deep) prior to assimilation. All other assimilated data, including radiance observations, are explicitly prevented from impacting the ozone analysis directly. Since SBUV sensors measure backscatter  
25 solar ultraviolet radiation, only daytime observations are available; wintertime ozone in polar regions is thus poorly constrained by observations. Early NOAA satellites experienced orbital drifts that resulted in reduced daylight coverage over time. For example, the equatorial crossing time for NOAA-11 drifted from ~2PM in 1989 to ~5PM five years later, leading to limited SBUV coverage in 1994 (ozone observations were entirely unavailable south of 30°S during that austral winter). A similar orbital drift in the NOAA-17 satellite impacted the quality of the MERRA ozone products in 2012 before the  
30 introduction of observations from NOAA-19 SBUV in 2013. Outside of the exceptions described above and occasional short temporal gaps, SBUV provides good coverage of the sunlit atmosphere.

Background error standard deviations for ozone are specified as ~4% of the global mean ozone on a given model level. Horizontal background error correlation lengths vary from ~400 km in the troposphere to ~800 km at the model top.

Assimilated ozone fields are fed into the forecast model radiation scheme and are used in the radiative transfer model for radiance assimilation.

Water ~~vapour~~vapor is also a prognostic assimilated variable in MERRA; however, unlike ozone, moisture fields in the stratosphere are relaxed to a 2-D monthly climatology with a relaxation time of 3 days. This climatology is derived from water ~~vapour~~vapor observations made by the UARS HALOE and Aura MLS instruments (e.g., Rienecker et al., 2011 and references therein). This climatological constraint is introduced gradually over the layer between the model tropopause and 50 hPa, where pressure-dependent blending between the climatology and the GCM water ~~vapour~~vapor is applied. Water ~~vapour~~vapor above the tropopause does not undergo physically meaningful variations on timescales longer than the 3-day relaxation timescale except in the lowermost stratosphere where the climatology is given a smaller weight. No attempt was made to account for methane oxidation or trends in stratospheric methane concentrations.

MERRA assimilates specific humidity measurements from radiosondes at pressures above 300 hPa and marine surface observations. Moisture fields are affected by microwave radiance data from SSM/I and AMSU-B/MHS, infrared radiances from HIRS, the GOES Sounder, and AIRS, and rain rates derived from TMI and SSM/I. Background error statistics for water ~~vapour~~vapor were derived using the National Meteorological Center method and applied using a recursive filters methodology (Wu et al., 2002). The moisture control variable is pseudo-relative humidity (Dee and Da Silva, 2003).

## 2.7 MERRA-2

The key differences between the treatment of ozone in MERRA-2 (Gelaro et al., 2017) and that in MERRA are in the observing system and background error covariances. From January 1980 to September 2004, MERRA-2 assimilates v8.6 SBUV retrievals of partial columns on a 21-layer vertical grid (Bhartia et al., 2013) and total ozone computed as the sum of individual layer values. Compared to the v8 retrievals used in MERRA, the v8.6 algorithm uses upgraded ozone cross-sections and an improved cloud height climatology. These updates result in better agreement with independent ozone data and make SBUV more suitable for long-term climatologies (Frith et al., 2014; McPeters et al., 2013). Starting in October 2004, SBUV data ~~wasere~~ replaced by a combination of TCO from Aura OMI (Levelt et al., 2006) and stratospheric profiles from Aura MLS (Waters et al., 2006). The OMI data consist of TCO retrievals from collection 3 and are based on the v8.5 retrieval algorithm, which is an improvement of the v8.0 algorithm extensively evaluated by McPeters et al. (2008). The assimilation algorithm makes use of the OMI averaging kernels to account for the sensitivity of these measurements to clouds in the lower troposphere (Wargan et al., 2015). MLS data are from v2.2 between October 2004 and May 2015 and v4.2 (Livesey et al., 2017) afterwards. Users of the MERRA-2 ozone product should therefore be aware that the reanalysis record may show a discontinuity in 2004 with two distinct periods as follows: the SBUV period (1980–September 2004) and the EOS Aura period (from October 2004 onward). The analysis is expected to be of higher quality during the latter period due to the higher vertical resolution of Aura MLS profiles relative to SBUV profiles and the availability of MLS observations during night.

Ozone background error variance in the MERRA-2 model follows Wargan et al. (2015). The background error standard deviation at each grid point is proportional to the background ozone at that point and time. This approach introduces a flow dependence into the assumed background errors and allows a more accurate representation of shallow structures in the ozone fields, especially in the UTLS. As in MERRA, the ozone analyses are radiatively active tracers in both the forecast model and the radiative transfer model used for assimilation of satellite radiances. Bosilovich et al. (2015) provided a preliminary evaluation of the MERRA-2 ozone product. A more comprehensive description and validation, including comparisons with MERRA, is given in Wargan et al. (2017)

The treatment of stratospheric water ~~vapour~~vapor in MERRA-2 is similar to that in MERRA, with a 3-day relaxation to the same climatological annual cycle. The main innovation in MERRA-2 that could impact water vapor is the introduction of additional global constraints that ensure continuity of water mass in the atmosphere. ~~The main innovation is the introduction of additional global constraints that ensure the conservation of the dry mass of the atmosphere and rescale the water vapour tendency to remove the globally integrated mean from the analysis increment~~ (Takacs et al., 2016).

In addition to the moisture data assimilated in MERRA, MERRA-2 assimilates GNSS-RO data and radiances from the recently introduced infrared sensors IASI, CrIS, and SEVIRI. Radiances from these recent IR instruments are not highly sensitive to stratospheric water ~~vapour~~vapor. Stratospheric water ~~vapour~~vapor is therefore not intentionally adjusted by the assimilation of these observations but may be affected in small ways. Changes in the MERRA-2 observing system relative to MERRA are described in more detail by Bosilovich et al. (2015) and McCarty et al. (2016). The moisture control variable in the MERRA-2 assimilation scheme is pseudo-relative humidity normalized by the background error standard deviation. Background error covariances used in MERRA-2 have been significantly retuned relative to those used in MERRA (Bosilovich et al., 2015).

### 3 Data

In this section, we describe the approach we use to process the reanalysis ozone and water ~~vapour~~vapor fields, and the observations used to evaluate them. We note that some of these observational data are assimilated by the reanalyses. While comparisons between reanalyses and observations would ideally be based on independent observations, this is not always possible given the paucity of water ~~vapour~~vapor and ozone data in parts of the atmosphere. However, comparison to assimilated observations can serve a useful purpose by providing an internal consistency check on the ability of reanalysis data assimilation systems to exploit the data they assimilate.

#### 3.1 Reanalysis data processing

Most of the comparisons presented in this paper are based on monthly mean reanalysis fields calculated from the “pressure level” data sets provided by each reanalysis ~~centre~~center, and processed into a standardized format as part of the CREATE project (<https://esgf.nccs.nasa.gov/projects/create-ip/>). The one exception to this is JRA-25 ozone data, which we have



processed ourselves. This was done because the pressure level data product provided by JMA (“fcst\_phy3m25”) used incorrect hybrid model level coefficients when converting from model levels to pressure levels. The JRA-25 ozone data used here were computed directly from the 6-hourly model level data product (“fcst\_phy3m”). To facilitate intercomparison amongst reanalyses, the pressure level-based datasets have been re-gridded to a common horizontal grid (2.5° lon x 2.5° lat) and a common set of 26 pressure levels (1000, 925, 850, 700, 600, 500, 400, 300, 250, 200, 150, 100, 70, 50, 30, 20, 10, 7, 5, 3, 2, 1, .7, .5, .3, .1 hPa). Unless otherwise noted, climatological comparisons follow the WMO convention in using the 30-year 1981–2010 climatological norm (Arguez and Vose, 2011).

Reanalysis TCO data are monthly means computed from the 6-hourly ~~model-level~~TCO ~~-datafields~~. All of the ~~models~~ reanalyses provided 6-hourly TCO ~~on various native horizontal grids~~, except for JRA-25. For JRA-25, 6-hourly ozone mass mixing ratios were provided on model levels. The mixing ratios were integrated for each horizontal grid point to get TCO, and then monthly means were computed. For each reanalysis, the climatologies and departures from climatology were calculated and are presented on each data set's native horizontal grid. For comparisons to the SBUV and TOMS/OMI data, each ~~model-reanalysis~~ was interpolated to the native horizontal grid of each of the observational data sets. Reanalysis data were excluded for days containing no observational data, in order to make the most valid comparison with the data sets.

### 15 3.2 SBUV and TOMS/OMI total column ozone

Two datasets are used to evaluate the total column ozone in the reanalyses. The first is the SBUV Merged Ozone Data Set (Frith et al., 2014). The second is a combination of TOMS and Aura OMI OMTO3d total ozone observations (Bhartia and Wellemeyer, 2002). These two data sets provide a long, coherent span of observations for evaluation. TOMS data were processed using the TOMS V8 algorithm, while the OMI and SBUV data were processed using the TOMS V8.6 algorithm. Because data from SBUV and TOMS (and in many cases OMI) are assimilated by most of the reanalyses, these comparisons are not independent.

### 3.3 SPARC Data Initiative limb satellite observations

The SPARC Data Initiative (Fueglistaler et al., 2009; Gettelman et al., 2011) offers monthly mean zonal mean climatologies of ozone (Neu et al., 2014; Tegtmeier et al., 2013) and water ~~vapour~~vapor (Hegglin et al., 2013) from an international suite of satellite limb sounders. The zonal monthly mean climatologies have undergone a comprehensive quality assessment and are suitable for climatological comparisons of the vertical distribution and interannual variability of these constituents in reanalyses on monthly to multi-annual timescales. We use a subset of the instrumental records available, as specified below.

The observational multi-instrument mean (MIM) for ozone averaged over 2005–2010 is derived using the SPARC Data Initiative (in the following abbreviated as SDI) zonal monthly mean climatologies from ACE-FTS, Aura MLS, MIPAS, and OSIRIS. These instruments provide data for the full 6 years considered and show inter-instrument differences with respect to the MIM that are generally smaller than  $\pm 5\%$  throughout most of the stratosphere. Hence, temporal inhomogeneities that

could affect the MIM are avoided and the standard deviation in the MIM is relatively small. Differences from the MIM in the lower mesosphere and tropical lower stratosphere are somewhat higher ( $\pm 10\%$ ) (Tegtmeier et al., 2013). The evaluation of the ozone QBO signal for 2005–2010 is based on the instruments OSIRIS, GOMOS, and Aura MLS, which produce the most consistent QBO signals (Tegtmeier et al., 2013).

5 | The observational MIM for water ~~vapour~~vapor averaged over 2005–2010 is derived using the SDI zonal monthly mean climatologies from Aura MLS, MIPAS, ACE-FTS, and SCIAMACHY. These instruments show inter-instrument differences that are generally within  $\pm 5\%$  of the MIM throughout most of the stratosphere (Hegglin et al., 2013). Differences from the MIM in the tropical upper troposphere increase to  $\pm 20\%$ .

### 3.4 Aura MLS satellite data

10 The evolution of ozone in the reanalyses is compared with that observed by Aura MLS. This instrument measures millimeter- and submillimeter-wavelength thermal emission from Earth's atmosphere using a limb viewing geometry. Waters et al. (2006) provide detailed information on the measurement technique and the Aura MLS instrument. Vertical profiles are measured every 165 km along the suborbital track with an along-track horizontal resolution of 200–500 km and a cross-track footprint of 3–9 km. Here we use version 4.2 (hereafter v4) MLS ozone measurements from September 2004 through  
15 December 2013. The quality of the MLS v4 data has been described by Livesey et al. (2017). The vertical resolution of MLS ozone is about 3 km and the single-profile precision varies with height from approximately 0.03 ppmv at 100 hPa to 0.2 ppmv at 1 hPa. The v4 MLS data are quality-screened as recommended by Livesey et al. (2017). V4 stratospheric (pressures less than 100 hPa) ozone values are within  $\sim 2\%$  of those in version 2.2 (v2), which is the version assimilated in MERRA-2  
20 shows high and low biases with respect to v2 at alternating levels, indicating improvement of vertical oscillations seen in v2 (Livesey et al., 2017) and v3 (Yan et al., 2016).

### 3.5 SWOOSH merged limb satellite data record

The Stratospheric Water and Ozone Satellite Homogenized (SWOOSH) database is a monthly-mean record of vertically  
25 resolved ozone and water ~~vapour~~vapor data from a subset of limb profiling satellite instruments operating since the 1980s (Davis et al., 2016). SWOOSH includes individual satellite source data from SAGE-II (v7), SAGE-III (v4), UARS MLS (v5/6), UARS HALOE (v19), and Aura MLS (v4.2), as well as a merged data product. A key aspect of the merged product is that the source records are homogenized to account for inter-satellite biases and to minimize artificial jumps in the record. The homogenization process involves adjusting the satellite data records to a “reference” satellite using coincident observations during time periods of instrument overlap. SWOOSH uses SAGE-II as the reference for ozone and Aura MLS  
30 as the reference for water ~~vapour~~vapor. SWOOSH merged product data are used for timeseries evaluations that start before 2004, prior to the availability of Aura MLS. After August 2004, the SWOOSH merged product is essentially the same as the v4.2 Aura MLS data.

## 4. Evaluation of reanalysis ozone products

### 4.1 Total column ozone seasonal cycle

In this section, we compare SBUV TCO data to reanalysis products over the 1981–2010 climatology period. Figure 3 shows the seasonal cycle in total column ozone from SBUV as a function of latitude and month. Also shown are the differences between TOMS/OMI and SBUV, and between the different reanalyses and SBUV. The climatological TCO fields of the TOMS/OMI and the reanalyses are given as line contours in the difference plots. Supplementary Figure S1 shows the equivalent comparison for TOMS/OMI data. The reanalyses all reproduce the major features of the seasonal cycle and latitudinal distribution of TCO. This agreement is not surprising given that all of reanalyses shown in Fig. 3 assimilate TCO data from one of the two satellites (Fig. 1). As such, the comparisons here do not represent independent validation of ozone in reanalyses but rather represent a test of the internal consistency of the ozone data assimilation system. Hence it is not surprising that MERRA and MERRA-2 generally perform better against SBUV than against TOMS/OMI, while ERA-Interim and JRA-55 generally perform better against TOMS/OMI than against SBUV, since MERRA and MERRA-2 assimilate SBUV (but not TOMS/OMI), while ERA-Interim and JRA-55 primarily assimilate TOMS/OMI (but not SBUV).

Although the reanalysis TCO fields look quite similar, a handful of widespread biases are revealed by considering the differences between reanalyses and observations. The agreement between the two observational TCO datasets is within approximately  $\pm 6$  DU (2~3%), with SBUV generally having smaller values in the tropics and larger values at high latitudes relative to TOMS/OMI. Differences between the reanalyses and the TCO observations are generally slightly larger than the difference between the two observational datasets. ERA-40 produces substantially larger TCO values than observed, particularly at higher latitudes. JRA-25 contains significantly smaller TCO values than observed (~10 DU less), except during the springtime at high southern latitudes.

For reanalyses that only (or mainly) assimilate UV-based retrievals, the winter hemisphere high latitudes remain largely unconstrained by data assimilation. The impact of the TCO observations may also be limited by filtering choices. For example, assimilated observations are filtered to exclude low solar elevation angles (less than  $10^\circ$  for TOMS and less than  $6^\circ$  for SBUV) in both ERA-40 and ERA-Interim. This filtering further limits observational impacts on the ozone analyses at higher latitudes. Hence, for ERA-Interim, before the start of the Aura MLS assimilation in 2008, high latitude ozone fields essentially reflect the effects of transport and the ozone parameterization used. For ERA-40, Dethof and H6lm (2004) showed that the ozone model produces high biases in ozone concentrations at high latitudes ranging from ~20 DU in the summer hemisphere to ~50 DU in the winter hemisphere, which is broadly consistent with the comparison shown in Fig. 3.

### 4.2 Zonal mean ozone cross-sections

In this section, we compare zonal mean multi-annual mean cross sections of ozone between the different reanalyses and the SDI MIM. We perform the comparison for 2005–2010 using the subset of instruments described in Sect. 3.3. This shorter period has been chosen to avoid sampling issues that could be introduced by changes in instrument availability, which could

alter sampling patterns, or trends in the constituents, such as the increase in ozone depletion from the 1970s to the mid 1990s. ERA-40 is excluded from this and all other comparisons with the SDI MIM because it ended in 2002.

Figure 4 shows multi-annual zonal mean ozone from the SDI MIM and the relative differences between each reanalysis and the SDI MIM (calculated as  $100 \cdot (R_i - \text{MIM}) / \text{MIM}$ , where  $R_i$  is the reanalysis field). Also indicated using contours are the climatological ozone distributions of the reanalyses. The reanalyses all capture the general zonal mean distribution of ozone, including the global maximum in ozone volume mixing ratio in the tropical middle stratosphere and the tropopause-following isopleths immediately above the tropopause. Among the reanalyses, MERRA-2 best reproduces this overall structure, with relative differences within  $\pm 5\%$  throughout the middle and upper stratosphere. MERRA, CFSR, and ERA-Interim also perform generally well, but with MERRA overestimating concentrations in the ozone maximum ( $\sim 10$  hPa) relative to the SDI MIM. ERA-Interim shows relatively good agreement in the middle stratosphere with biases smaller than  $\pm 5\%$  but includes a low bias with magnitudes greater than 10% in the upper stratosphere. All reanalyses show biases exceeding  $\pm 10\%$  in the lowermost stratosphere, at pressures greater than 100 hPa. JRA-55 ~~shows evident is an~~ improvement relative to JRA-25, particularly in the polar regions. Negative biases in JRA-55 have approximately halved in the middle and upper stratosphere, compared to JRA-25. However, JRA-55 also shows somewhat higher positive biases around the tropical upper troposphere and lower stratosphere than JRA-25. The latter included a strong positive bias (greater than 20% relative to the MIM) in the lower and middle stratosphere and a strong negative bias ( $>20\%$ ) in the upper stratosphere. JRA-55 still shows a negative bias in the upper stratosphere, but the magnitude is approximately halved relative to that in JRA-25. It is worth noting that the diurnal cycle in ozone has not been explicitly accounted for in the observational MIM. Neglecting the diurnal cycle potentially contributes to differences between the reanalyses and observations in the upper stratosphere and lower mesosphere.

All reanalyses ~~(except the JRA products-55)~~ produce a positive bias in ozone in the Southern Hemisphere (SH) lower stratosphere. This indicates an inability to simulate Antarctic ozone depletion accurately due to a combined effect of limited data coverage, data filtering, and limitations of the reanalyses' chemistry schemes at high latitudes (Sect. 4.1). A dipole is apparent in the CFSR and ERA-Interim biases, with a high bias near  $\sim 100$  hPa located below a low bias near  $\sim 10$  hPa. This dipole may reflect a lack of information about the vertical location of the ozone hole in the TCO and SBUV observations assimilated by these systems. In contrast, MERRA includes a significant high bias ( $>10\%$ ) at Southern high latitudes that extends throughout the stratosphere.

### 4.3 Ozone monthly mean vertical profiles and seasonal cycles

Figures 5a and b show vertical profiles of ozone for January (2005–2010 average) for the reanalyses and the SDI MIM at two different latitudes,  $40^\circ\text{N}$  and  $70^\circ\text{S}$ , respectively, along with the relative differences for each reanalysis with respect to the MIM. In addition, Figures 5c and d show the seasonal cycles of ozone for three different pressure levels at  $40^\circ\text{N}$  and  $70^\circ\text{S}$ , respectively. The vertical profiles and the seasonal cycles reveal seasonal information on reanalyses-observation

differences that expands upon the annual zonal mean evaluation presented in Sect. 4.2. In general, the results shown reinforce the conclusions of the previous section.

5 Most reanalyses resolve the vertical distribution in January reasonably well at both latitudes, in particular in the middle stratosphere between around 50 and 5 hPa. MERRA-2, MERRA, and CFSR perform particularly well. JRA-25, on the other hand, is a clear outlier that ~~places the ozone maximum too low in the profile and~~ produces too little ozone ~~above in the vicinity of~~ the maximum. JRA-55 and ERA-Interim also underestimate ozone concentrations in the upper stratosphere by between 10 and 20% but are not as strongly biased as JRA-25 (which produces differences of more than 230%). All reanalyses show larger percentage differences from the MIM in the lower part of the profile at pressures greater than 100 hPa. The reanalyses seem to overestimate ozone at around 150 hPa by 20% in the Southern high latitudes, possibly related to not capturing accurately enough the extent of ozone depletion during spring. Below 200 hPa at both latitudes, all reanalyses underestimate observed ozone values.

15 The agreement between the reanalyses and observations varies by month, as can be seen in Figures 5c and d, which show the annual cycle for selected pressure levels (150, 50, and 10 hPa) and somewhat extended latitude bands of 30°N-50°N and 60°S-80°S, respectively. The agreement in the ozone seasonal cycle between the SDI observations and the reanalyses is better in the Northern Hemisphere (NH) mid-latitudes (where the seasonal cycles have a simple sinusoidal structure) than in the SH high latitudes. In the NH at 50 and 150 hPa, ozone reaches its annual maximum during boreal spring and its annual minimum during autumn, attributable to the strong seasonality in the Brewer-Dobson circulation. The seasonal cycle is shifted at 10 hPa, with a maximum in summer and a minimum in winter, attributable mostly to ozone photochemistry. Most of the reanalyses produce a fairly accurate ozone evolution at these levels with exceptions as follows: At 150 hPa, JRA-55 shows a strong low bias when compared to both observations and the other reanalyses during the NH winter/spring months. All the other reanalyses tend to overestimate the absolute ozone values, but agree rather well with the seasonal cycle in the observations in terms of amplitude and phase. At 50 hPa, the seasonal cycle produced by JRA-55 shows a more gradual decline in ozone concentrations into autumn relative to both observations and other reanalyses, ~~and JRA-25 shows too high ozone mean values throughout the year. At 10 hPa, JRA-25 produces a seasonal cycle that has a bi-modal structure more consistent with that observed in the tropics (not shown).~~ ERA-Interim, MERRA, and CFSR at 10 hPa tend to overestimate ozone during spring and early summer, while JRA-55 (JRA-25) tends to underestimate (overestimate) ozone during fall and winter.

30 Seasonal cycles in SH high latitudes have a more complex structure than those in the NH mid-latitudes due to generally weaker downwelling in the Brewer-Dobson circulation and the influence of Antarctic ozone depletion. As a consequence, the reanalyses have more difficulty in capturing the seasonal cycle. At 10 hPa, MERRA-2 shows the best agreement with the observations. CFSR also follows the observations relatively well, but overestimates the amplitude of the seasonal cycle, primarily because of values that are too low during May through July. MERRA and JRA-25 are outliers in that they do not contain the strong annual minimum observed during late boreal-austral autumn and early winter. At 50 hPa, MERRA and JRA-25 agree better with observations than at 10 hPa, but still underestimate austral springtime ozone depletion. Finally, at

150 hPa, the seasonality in the reanalyses varies widely and is inconsistent with that in the observations, with the exception of MERRA, which produces the most realistic seasonal cycle amplitude. MERRA-2 shows the closest agreement with observations at all levels except for 150 hPa, which is the next to lowest valid level of the MLS v2.2 ozone retrievals that it assimilates.

#### 5 4.4 Ozone interannual variability

Figure 6 shows time series of interannual variability of ozone and its anomalies in the SDI MIM and reanalyses during 2005–2010. The anomalies, which are calculated for each reanalysis by subtracting multi-year monthly means averaged over 2005–2010 from the monthly mean timeseries, are a good indicator of how well physical processes (such as transport) are represented in reanalyses. Time series are shown for the SH high latitudes (averaged over 60°S-80°S) at 50 hPa, and for the  
10 NH mid-latitudes (40°N-60°N) at 150 and 10 hPa. In all cases, MERRA-2 produces the closest match with the SDI MIM in terms of both the absolute values and the structure of its interannual variability. This agreement highlights the benefit of assimilating vertical profile observations from a limb-viewing satellite instrument. Although it has to be noted that the comparison is not done against truly independent observations in this case, since Aura MLS is included in the SDI MIM. MERRA-2 is an evident improvement over MERRA, which tends to disagree with the absolute ozone values of the  
15 observations at 150 hPa and to overestimate them at 10 hPa, and to underestimate interannual variability at both levels in the NH mid-latitudes. JRA-55 also shows clear improvement relative to JRA-25 with respect to the amplitude and structure of interannual variability, at least at 10 hPa in the NH mid-latitudes. Large excursions seen in JRA-25, such as the sudden drop in ozone at the beginning of 2008, are not present in JRA-55 or in the observations.

Although ERA-Interim ozone mean values mostly agree well with observations, the amplitude of its interannual variability  
20 is larger than observed. In particular, ERA-Interim overestimates the negative anomaly in NH midlatitudes at 10 hPa, and the positive anomaly in SH high latitudes at 50 hPa during 2008. The largest differences appear to affect ERA-Interim from mid-2009 when the assimilation of Aura MLS data restarted with the (v3) NRT product after months of data unavailability. CSFR also produces large interannual excursions during certain years (e.g., during spring 2006 and 2007 at 50 hPa in SH high latitudes). This issue may be related to SBUV only offering measurements between September to March, so that the  
25 assimilation system is not well constrained during the remainder of the year.

#### 4.5 Ozone time series in equivalent latitude coordinates

Equivalent latitude (EqL) is a common vortex-~~centred~~centered coordinate used in studies of the stratosphere (e.g., Butchart and Remsberg, 1986; Manney et al., 1999; and references therein). This coordinate is also useful as a geophysically-based coordinate in the UTLS (e.g., Santee et al., 2011), although interpretation becomes more complicated in this context (e.g.,  
30 Manney et al., 2011; Pan et al., 2012). The equivalent latitude of a potential vorticity (PV) contour is defined as the latitude of a circle ~~centred~~centered about the pole enclosing the same area as the PV contour (see Hegglin et al., 2006 for a visual illustration). Figure 7 shows the time series of v4 MLS ozone (Sect. 3.4) for late 2004 through 2013 in the lower stratosphere

(520 K), along with differences between MERRA, MERRA-2, ERA-Interim, CFSR, and JRA-55 and MLS ozone at the same level. MLS ozone is interpolated to isentropic surfaces using temperatures from MERRA. The EqL ozone time series are then produced using a weighted average of MLS data in EqL and time, with data also weighted by measurement precision (e.g., Manney et al., 2007; Manney et al., 1999). Figures S2-S3 in the Supplement show the equivalent evaluation for the 350K and 850 K potential temperature levels.

Figure 7 reveals that MERRA-2 matches MLS more closely over the full period than do the other reanalyses. This is expected because the stratospheric ozone reanalyses in MERRA-2 are largely constrained by the MLS stratospheric ozone profiles (v2 for the period shown here). This agreement is especially apparent during Antarctic winter and spring, when other assimilated ozone products (e.g., SBUV/2 and TOMS) cannot provide measurements due to darkness and simplified chemical parameterizations cannot adequately represent heterogeneous loss processes. The improved vertical resolution of MLS relative to SBUV/2 also better constrains the structure of the ozone hole, which is vertically limited. ERA-Interim also shows close agreement with MLS during the periods when it assimilates MLS ozone products (2008 and mid-2009 through present). The change in behavior in ERA-Interim between these time periods, and the general similarity of PV contours among the different reanalyses suggests that the poor representation of ozone in these regions is due more to the lack of assimilated ozone data than to the representation of polar dynamical processes in reanalyses.

Biases in the reanalyses that do not assimilate MLS and OMI ozone vary in magnitude and sign, not only among the reanalyses but also with altitude and latitude (see also Figs. S2-S3). High biases in MERRA and CFSR ozone during Arctic winter may be partially related to inadequate representations of ozone chemistry and an overall lack of measurements. We speculate that the latter is dominant due to the appearance of these biases even during years with minimal observed chemical ozone loss. JRA-55 biases increase strongly with altitude (cf., Figs. S2-S3), becoming even larger in the upper stratosphere. These large biases suggest that column ozone alone is insufficient to properly constrain the vertical distribution of the ozone analyses. but that assimilation of vertically resolved observations during polar night can provide a much better constraint on ozone in these regions.

#### 4.6 Ozone quasi-biennial oscillation signals

Variations in transport and chemistry associated with the quasi-biennial oscillation (QBO) in tropical zonal wind are among the largest influences on interannual variability in equatorial ozone. The QBO signal in tropical ozone has a double-peaked structure with maxima in the lower (50–20 hPa) and the middle-to-upper (10–2 hPa) stratosphere (Hasebe, 1994; Zawodny and McCormick, 1991). Ozone is mainly under dynamical control below 15 hPa, where the QBO signal results primarily from changes in ozone transport due to the QBO-induced residual circulation. In contrast, ozone is under photochemical control above 15 hPa. The QBO signal in these upper levels is understood to arise from a combination of QBO-induced temperature variations (Ling and London, 1986; Zawodny and McCormick, 1991) and QBO-induced variability in the transport of  $\text{NO}_y$  (Chipperfield et al., 1994). A realistic characterization of the time–altitude QBO structure is an important aspect of physical consistency in ozone data sets.

Figure 8 shows time–altitude cross-sections of deseasonalized ozone anomalies from 2005 to 2010 from the SDI MIM, along with the differences between the ozone anomaly fields from the reanalyses and the SDI MIM. The climatological QBO anomaly fields of the reanalyses are given as contours in the difference plots. Combined ozone measurements from the limb-viewing satellite instruments show a downward propagating QBO ozone signal with a shift in the phase around 15 hPa. All reanalyses exhibit some degree of quasi-biennial variability; however, differences are evident in the phase, amplitude, vertical extent, and downward propagation of these signals. The largest deviations from observations are in JRA-25, which displays positive anomalies from 2005 to mid-2007 followed by negative anomalies from mid-2007 through 2010 in place of the QBO signal above 15 hPa. In contrast, ERA-Interim shows predominantly negative anomalies in the 100–10 hPa pressure range before 2008 and positive anomalies afterwards. The changes in ERA-Interim coincide with the beginning of the assimilation of Aura MLS profiles beginning in 2008, which caused a shift to positive anomalies. Negative anomalies are present during the first half of 2009 when no MLS data were assimilated, followed by positive anomalies after the reintroduction of MLS data in June 2009 (Sect. 2.5).

CFSR and MERRA produce anomalies that are roughly consistent in amplitude and frequency with the QBO ozone signal in the satellite data. However, no clear downward propagation is apparent in ~~this signal~~~~these reanalyses.~~ ~~despite the existence of a downward propagating signal in SBUV data (McLinden et al., 2009), the only vertically resolved ozone measurements assimilated by CFSR and MERRA.~~ The vertical structure of the anomalies is also shifted. Instead of a pair of peaks in the lower stratosphere (50–20 hPa) and middle-to-upper stratosphere (10–2 hPa), a single peak emerges near 15 hPa. ~~This finding may be at least partially explained by the fact that the only vertically resolved ozone measurements assimilated by CFSR and MERRA come from SBUV. SBUV shows only a weak oscillatory behaviour, with a much smaller amplitude and without a properly downward propagating signal, attributable to the instrument’s vertically limited and rather low vertical resolution (McLinden et al., 2009; Kramarova et al., 2013).~~ JRA-55 and MERRA-2 produce a phase and amplitude of QBO variability like those observed in the satellite data. Overall, the features of the QBO (including the downward propagation) are much improved in MERRA-2 relative to MERRA (Coy et al., 2016), and in JRA-55 relative to JRA-25. Nearly all reanalysis data sets extend the QBO ozone signal to altitudes below 100 hPa; this upper tropospheric signal is not present (or not captured) in the satellite observations.

#### 4.7 Ozone hole area

The Antarctic “ozone hole” is a region of severe ozone depletion that starts in late August or early September and lasts until November or early December. The ozone hole is commonly defined as the area within the 220 DU TCO contour. Figure 9 shows average ozone hole areas based on TOMS/OMI observations and six reanalyses during 1981–2010. The average is computed over 21 September–20 October of each year. This period is chosen to avoid the partial coverage of the SH high latitudes that occurs in TOMS/OMI data during the early part of September. Observationally based ozone hole areas are larger than those produced by the reanalyses in almost all years between 1981 and 2002. The systematic negative bias in reanalysis-based ozone hole areas is consistent with reanalyses generally underestimating ozone loss. Most of the models



track the observations well starting in 2003. This is not a truly independent comparison (all reanalyses except for MERRA assimilate TOMS and/or OMI observations); however, it does show the general consistency among most reanalyses in reproducing realistic interannual and decadal changes in the size of the Antarctic ozone hole, except for a few outliers discussed below.

5 The newer reanalyses (MERRA-2, ERA-Interim, JRA-55, and CSFR) are all within 1 million km<sup>2</sup> (5.2%) of the observations, and generally produce root-mean-square (RMS) differences relative to TOMS/OMI of less than 0.9 million km<sup>2</sup> (14.6%). A notable exception to the latter is MERRA-2 with an RMS of 2.8 million km<sup>2</sup> (44.5%). This large RMS is attributable to an outlier year in 1994, when MERRA-2 had a very small ozone hole (Fig. 9). JRA-55 produces the smallest RMS difference relative to TOMS/OMI, while MERRA-2 model produces the smallest mean difference relative to these  
10 observations.

MERRA did not produce an ozone hole in 1994, and produced very small ozone holes in 1993, 1997, 2009, and 2010. For related reasons, MERRA-2 did not produce an ozone hole in 1994, and produced a relatively small ozone hole in 1993. The elimination or reduction of the ozone hole during those years was caused by a lack of ozone observations for constraining the ozone field, as the processes that contribute to the development of the ozone hole are not represented in the parameterized  
15 ozone chemistry used in MERRA and MERRA-2. In 1994, orbital drift of the *NOAA-11* satellite that provided the SBUV/2 TCO data assimilated by both MERRA and MERRA-2 led to a lack of ozone observations south of ~30°S during early Austral spring. *NOAA-11* SBUV/2 coverage was also limited in 1993. While both MERRA and MERRA-2 use *NOAA-11* SBUV, the version 8.6 data assimilated in the latter allowed less stringent quality screening criteria. Specifically, MERRA-2 uses observations made at solar zenith angles greater than 84°, excluded in MERRA, if they are otherwise marked as “good”.  
20 This results in a slightly better coverage of *NOAA-11* SBUV in MERRA-2, explaining its better performance in 1993 and even 1994. The MERRA ozone hole was only weakly constrained by observations in late September 1997 because *NOAA-11* data only extended to 60°S–75°S between 21 September and 20 October. MERRA-2 does not have a low bias in ozone hole size during 1997 because it used data from *NOAA-14* rather than data from *NOAA-11*. The MERRA ozone hole was also affected by orbital drift in the *NOAA-17* satellite and the concomitant loss of SBUV/2 observations at high southern latitudes  
25 during the austral springs in 2009 and 2010. MERRA-2 is unaffected during these years because of its assimilation of ozone observations from Aura OMI and MLS.

ERA-40 did not assimilate ozone data in 1989 and 1990. This resulted in a high bias in ozone concentrations and a very small ozone hole. The ERA-40 model also severely underestimated ozone hole area in 1997, most likely due to a gap in assimilated TCO from the Earthprobe TOMS instrument between August and December that year (Fig. 1; note that *NOAA-9*  
30 SBUV/2 profiles were assimilated during this timeframe as shown in Fig. 2). By contrast, the area of the ERA-Interim ozone hole was too large in 1995. This may be due to a lack of assimilated TCO observations in ERA-Interim during 1995 (Fig. 1).

## 4.8 Long-term evolution of ozone

Figure 10 shows the evolution of deseasonalized TCO anomalies from the reanalyses and assimilated observations from SBUV and TOMS/OMI. Also shown are the differences between the reanalyses and the primary TCO observations they assimilate. Both observational data sets show similar features, including a general trend toward decreasing ozone in the SH high latitudes, consistent with the Antarctic ozone hole depletion discussed in the previous section. However, in Fig. 10, comparison to the data set assimilated by a given reanalysis is done because differences between the TOMS/OMI and SBUV data sets show an apparent step change at the beginning of 2004. A comprehensive set of plots showing this step change, as well as reanalysis/observation differences separately for each data source, is provided in the supplementary material (Figs. S4-S5).

As expected, reanalyses agree more closely with TCO data that they assimilate than with data that they do not assimilate. For example, MERRA, MERRA-2, and CFSR assimilate SBUV data. The influence of SBUV on these reanalyses can be seen in the QBO-related anomalies in the tropics (particularly after ~1998) that are present in both the SBUV data and in the reanalyses that assimilate it. Differences between these reanalyses and SBUV are smaller in magnitude and more homogeneous in space and time than differences between these reanalyses and TOMS/OMI. The discontinuity in 2004 is particularly pronounced when MERRA and CFSR are compared against TOMS/OMI (Fig. S5). Similarly, differences between the ECMWF reanalyses and TOMS/OMI are generally more homogeneous and smaller in magnitude than differences between the ECMWF reanalyses and SBUV (Fig. S4). The period during which ERA-40 did not assimilate any ozone data (1989–1990) is also evident in Fig. 10. The stark contrast between this period and the surrounding years indicates the importance of data assimilation in constraining reanalysis ozone fields.

Figure 11 shows differences between reanalysis ozone fields and SWOOSH satellite limb profiler merged ozone data on two pressure levels (10 hPa and 70 hPa). This plot helps to evaluate disruptions in the temporal homogeneity of reanalysis ozone fields caused by changes in the assimilated observational data, and also provides a partially independent dataset for comparison with the reanalyses. The SWOOSH record is based primarily on v4.2 Aura MLS ozone starting in August 2004, so comparisons with reanalyses that assimilate MLS (i.e., MERRA-2 and ERA-Interim) after that time are not independent. However, none of the observations used to construct the SWOOSH record prior to August 2004 were assimilated by these reanalyses.

At 10 hPa, CFSR, MERRA, and MERRA-2 show the best agreement with observations. At this level, ERA-Interim and JRA-25 have positive biases in both SH and NH midlatitudes, while JRA-55 has a negative bias relative to SWOOSH in the tropics.

Overall, reanalysis ozone products do not exhibit large discontinuities at 10 hPa. As expected, both MERRA-2 and ERA-Interim show extremely good agreement with SWOOSH during the period in which they assimilate Aura MLS ozone data. Biases in these reanalyses undergo a step change when they start assimilating ozone profiles from Aura MLS ozone. For example, MERRA-2 assimilates Aura MLS data from August 2004 (Fig. 2), and at that time biases in 10 hPa ozone relative

to SWOOSH drop suddenly to less than 5% at all latitudes. This reduction is also apparent in ERA-Interim, which assimilates Aura MLS ozone data during 2008 and then from June 2009 through the present. Similar sudden reductions in ozone biases relative to SWOOSH are seen in ERA-Interim in both early 2008 and the latter half of 2009.

5 Differences between reanalysis ozone fields and SWOOSH are larger at 70 hPa. A strong discontinuity in the MERRA-2 time series occurs in mid-2004 when it begins to assimilate Aura MLS ozone data. To a lesser extent there is also a discontinuity (in 2008 and again in mid-2009) when ERA-Interim begins assimilating Aura MLS ozone data. The large positive bias in MERRA-2 that starts in mid-2004 is also seen in comparisons to (non-assimilated) ozonesondes (Wargan et al., 2017). This positive bias is related to vertical averaging of the MLS data before assimilation by MERRA-2 (Wargan et al., 2017).

10 For the other reanalyses that don't assimilate MLS, there are generally not strong discontinuities that can be tied to observing system changes. There does seem to be a change in the ERA-Interim differences at the beginning of 2003 when it begins to assimilate vertically resolved data from MIPAS and TCO from SCIAMACHY. Beyond the discontinuities discussed above, at 70 hPa differences between the reanalysis ozone fields and SWOOSH are relatively consistent in time, with negative biases prevailing in [JRA-25](#), CSFR, MERRA, and MERRA-2 (pre-Aura MLS), patchy biases in ERA-Interim, and mostly positive biases in [JRA-25](#) and [JRA-55](#) (especially in the tropics).

## 5 Evaluation of reanalysis water ~~vapour~~vapor

In this section, we evaluate reanalysis estimates of water ~~vapour~~vapor in and above the tropopause layer against available observations. In keeping with the S-RIP remit, this section focuses exclusively on evaluations of reanalysis water ~~vapour~~vapor products in the upper troposphere and stratosphere.

### 20 5.1 Zonal mean water ~~vapour~~vapor cross-sections

Figure 12 shows multi-annual zonal mean water ~~vapour~~vapor for 2005-2010 from the SDI MIM along with relative differences between each reanalysis and the MIM (calculated as  $100 \cdot (R_i - \text{MIM}) / \text{MIM}$ , where  $R_i$  is a reanalysis field). In contrast to ozone, the reanalyses do not consistently capture the zonal mean vertical distribution of water ~~vapour~~vapor. The pressure-level products provided by [JRA-25](#) and [JRA-55](#) do not include ~~analysed~~analyzed stratospheric water ~~vapour~~vapor fields, while CSFR produces a stratosphere that is much too dry (low biases exceeding 60%). ERA-Interim, MERRA, and MERRA-2 show water ~~vapour~~vapor fields that are close to observations. These three systems resolve the distinct minimum in water ~~vapour~~vapor mixing ratios just above the tropical tropopause, the second minimum in the lower stratosphere at SH high latitudes, and the increase in water ~~vapour~~vapor with increasing altitude. In contrast to other reanalyses, MERRA and MERRA-2 also extend up to the lower mesosphere (not shown), and, albeit with some limitations, they both capture the water ~~vapour~~vapor maximum found in the upper stratosphere (e.g., Hegglin et al., 2013), although slightly underestimated

compared to observations, consistent with the simple parameterization as a 3-day relaxation to a climatology (Sects. 2.6 and 2.7).

CFRSR is much too dry throughout the stratosphere and does not capture the typical structure of water ~~vapour~~vapor isopleths. This bias is due in part to the lack of assimilated observations to constrain the water ~~vapour~~vapor reanalyses at these altitudes and in part to the absence of a methane oxidation parameterization in the forecast model (Sect. 2.3). All reanalyses contain high biases relative to the SDI MIM at pressures greater than 100 hPa (see also Jiang et al., 2015), although this may be in part explained by the increase in measurement uncertainty of satellite limb sounders with decreasing altitude in the upper troposphere (Hegglin et al., 2013). Several studies have shown that Aura MLS contains a dry bias in the upper troposphere/lower stratosphere around 200 hPa (e.g., Davis et al., 2016; Vömel et al., 2007), and similarly a dry bias has been found in the upper troposphere for ACE-FTS (Hegglin et al., 2008).

## 5.2 Water ~~vapour~~vapor monthly mean vertical profiles and seasonal cycles

Figures 13 a and b show vertical profiles of water ~~vapour~~vapor for January (2005–2010 average) for the reanalyses and the SDI MIM at two different latitudes 40°N and 70°S, respectively, along with the relative differences for each reanalysis with respect to the MIM. Figures 13c and d show the seasonal cycles of ozone for three different pressure levels at 40°N and 70°S, respectively. In general, the results shown reinforce the conclusions of the previous section.

The comparisons in Figs. 13 a and b reveal very good agreement (within  $\pm 10\%$ ) between ERA-Interim, MERRA, MERRA-2, and the observations at altitudes above 100 hPa. As mentioned in the previous section, water ~~vapour~~vapor from CFRSR is unrealistic in the stratosphere, with values much lower than those observed. The reanalyses show large inconsistencies between their absolute values at altitudes below 100 hPa, leading to sharp increases in their relative differences with respect to the MIM of  $>100\%$ . These relative differences are systematically positive except for in CFRSR and JRA-25, pointing towards potential negative biases in the water ~~vapour~~vapor observations at these altitudes (e.g., Hegglin et al., 2013). The results may also indicate that the reanalyses produce an excessively moist tropical upper troposphere and/or excessive mixing of moist tropospheric air into the extratropical lowermost stratosphere. The 100 hPa level is one of the most important levels for stratospheric water ~~vapour~~vapor studies, because it is near the level where stratospheric water ~~vapour~~vapor entry mixing ratios are set in the tropics (Fueglistaler et al., 2009) and because it is near the peak region of the radiative kernel for water ~~vapour~~vapor in the extratropics (Gettelman et al., 2011).

The agreement between the reanalyses and observations varies by month, as shown in Figs. 13 c and d for selected pressure levels (250, 100, and 50 hPa) and latitude bands (30°N-50°N and 60°S-80°S). At NH mid-latitudes (30°N-50°N; Fig. 13c) at 250 hPa, all reanalyses are biased high relative to the observations by more than 100%, lending further support to the results by Jiang et al. (2015), who compared the reanalyses to Aura MLS alone, which is known to have a low bias around this altitude (Davis et al., 2016; Hegglin et al., 2013; Vömel et al., 2007). JRA-25 and JRA-55 have the smallest high biases relative to observations at 250 hPa. At 100 hPa and 50 hPa, ERA-Interim, MERRA, and MERRA-2 perform best, with approximately correct mean values, but somewhat underestimated seasonal cycle amplitudes. As noted earlier, a

significant portion of the agreement in MERRA and MERRA-2 results from the relaxation of stratospheric water ~~vapour~~vapor towards a climatology that is based in part on Aura MLS data (which are also included in the SDI MIM). JRA-55 (JRA-25) has mean values that are much too large (small) at 100 hPa. In addition to being too dry at 100 and 50 hPa, CSFR also has incorrect amplitude and phase of the seasonal cycle at these levels.

5 At SH high latitudes (60°S-80°S; Fig. 13d), all reanalyses show approximately the right phase, but overestimate mean values and amplitudes at 250 hPa, similar to the results at NH mid-latitudes. At 100 and 50 hPa, ERA-Interim captures the phase and amplitude of the observed seasonal cycle best when compared to the other reanalyses, but exhibits a slight low bias at 50 hPa. MERRA and MERRA-2 show also quite good agreement in terms of mean value, amplitude, and phase at 100 hPa, but overestimate mean values at 50 hPa, and also show a somewhat early minimum followed by an increase in  
10 September that occurs about a month earlier than observations. JRA-25 somewhat underestimates the mean value, but shows a similar phase and amplitude as the observations at 100 hPa. JRA-55 on the other hand, strongly overestimates the amplitude of the seasonal cycle at this level with mean values that are much too high. CSFR shows too low values at both 100 and 50 hPa, but captures the seasonality somewhat better than it does in the NH mid-latitudes.

### 5.3 Interannual variability in water ~~vapour~~vapor

15 Figure 14 shows time series of interannual variability in water ~~vapour~~vapor and its anomalies based on observations and reanalysis products during 2005–2010. At 250 hPa in NH midlatitudes (40°N-60°N), the reanalyses generally follow the observed interannual variability extremely well, especially JRA-25, JRA-55, and MERRA. CSFR seems to exhibit an underlying positive trend in its timeseries that is stronger than that observed. And as noted previously, all reanalyses are wetter than observations at this level by approximately a factor of two.

20 At 100 hPa in the tropics (a level that is often used to estimate stratospheric water ~~vapour~~vapor entry mixing ratios), all reanalyses except CSFR compare reasonably well with the observed anomalies. Perhaps surprisingly, JRA-25 captures the interannual anomalies quite well despite being biased in its seasonal cycle. CSFR shows no clear interannual variability and produces water ~~vapour~~vapor mean values as low as 0 ppmv. CSFR begins to produce more realistic water ~~vapour~~vapor concentrations at these levels in 2010 with values that are larger and in better agreement with observations than those in the  
25 other reanalyses. This change is discussed further in Sect. 5.4 Note that the SDI MIM for this level only includes Aura-MLS and ACE-FTS due to known problems in SCIAMACHY and MIPAS data in this region (Hegglin et al., 2013).

At 50 hPa in the SH high latitudes (60°S-80°S), MERRA and MERRA-2 have roughly correct water ~~vapour~~vapor mean values, whereas CFSR and ERA-Interim are too low. MERRA and MERRA-2 both place the minimum during austral winter (from dehydration processes in the cold polar vortex) about one month too early. Except for CFSR, the other reanalyses  
30 capture the correct structure in the interannual variability, including the prominent positive anomaly in 2010. MERRA and MERRA-2 show less variability than observed, which is unsurprising given their strong relaxation to the climatology.

#### 5.4 Tropical tape recorder in water ~~vapour~~vapor

Representations of the tropical tape recorder (Mote et al., 1996) provide an additional illustration of problems in reanalysis stratospheric water ~~vapour~~vapor products. Figure 15 shows the time–height evolution of water ~~vapour~~vapor in reanalyses and the merged SWOOSH observations averaged over the 15°S–15°N tropical band. Anomalies are calculated separately for each data set, relative to the mean seasonal cycle at each level for the period 1992-2014 (except ERA-40, which is 1992-2002), when all reanalyses (except ERA-40) overlap. Variations in these fields reflect changes in the mixing ratio of water ~~vapour~~vapor entering the tropical lower stratosphere, as driven by variations in tropical tropopause temperatures and the subsequent vertical propagation in the ascending branch of the stratospheric overturning circulation. Interannual variability in both water ~~vapour~~vapor entry mixing ratios and ascent rate (the vertical slope of the signal) is superimposed on this mean seasonal cycle. Although reanalyses do not reproduce observed water ~~vapour~~vapor concentrations in the stratosphere, most reanalyses do produce a tropical tape recorder signal.

As previously discussed, CFSR (Fig. 15a) produces water ~~vapour~~vapor concentrations near zero in the stratosphere for most of the record, although unrealistically wet values appear above 20 hPa at certain times (e.g. 1995 and 1999). These upper stratospheric wet anomalies (and several others that occurred before 1992) all correspond to transitions in the main CFSR production stream (see Fig. 2, Fujiwara et al., 2017). We hypothesize that these wet anomalies are a remnant of a wet bias in the model ~~initialisation~~initialization that remains after the ~1-year spinup. Additional step changes in water ~~vapour~~vapor are evident at the beginning of 2010 and at the beginning of 2011. The latter step change corresponds to the transition from CFSR (CDAS-T382) to CFSv2 (CDAS-T574) at the beginning of 2011. As discussed in Sect. 2.2, CFSv2 is intended as a continuation of CFSR but has differences in model resolution and physics relative to the original system. Although the reasons for the step change at the beginning of 2010 are not known definitively, we note that CFSR was extended for the year 2010 following its original completion over the 1979-2009 time period. This extension used the original CDAS-T382 system but with some slight changes to the forecast model. It is likely that the CFSR 2010 run was performed without a sufficiently long spin-up period, or that a change to the model configuration resulted in the observed water ~~vapour~~vapor discontinuity beginning in 2010.

ERA-40 and ERA-Interim (Fig. 15 c, e) are generally drier than the SWOOSH observations (Fig. 15 k), although the ERA-Interim represents an evident improvement over ERA-40 in this respect. Both MERRA and MERRA-2 (Fig. 15 g, i) are close in magnitude to SWOOSH, but this agreement is expected given that both systems relax stratospheric water ~~vapour~~vapor to a climatology based on Aura MLS and HALOE (Sect. 2.6, 2.7).

The reanalyses all produce tape recorder slopes that are more vertical than suggested by the observations, indicating that vertical upwelling in the tropical stratosphere is too strong in reanalyses. Although biases and differences in tropical stratospheric upwelling have been addressed quantitatively for a subset of reanalyses elsewhere (Abalos et al., 2015; Jiang et al., 2015), the SWOOSH data shown in Fig. 15 enable a comparison that extends beyond the Aura MLS record. This

extension allows for comparison to ERA-40, and shows that ERA-Interim benefits from a much-improved representation of stratospheric water ~~vapour~~vapor and its variability relative to its predecessor.

Figure 15 also shows interannual variability in tropical stratospheric water ~~vapour~~vapor as represented by the anomaly from the mean seasonal cycle at each level. Interannual variability in the tape recorder signal is related to interannual variability in cold-point tropopause temperatures (Fig. 15 m), with warm anomalies at the tropopause corresponding to wet anomalies in the tape recorder and vice versa. Although the reanalyses produce almost identical interannual variations in tropical tropopause temperatures over the period considered here, their interannual variations in stratospheric water ~~vapour~~vapor differ substantially. The strong relaxation to climatology applied in MERRA and MERRA-2 results in very little interannual variability above 60 hPa because of the short nudging timescale for WV (3 days). ERA-40 produces a very large wet anomaly during the 1997–1998 El Niño that coherently propagates upwards. This anomaly is wetter than that suggested by SWOOSH and the other reanalyses. SWOOSH and the reanalyses all show a wet anomaly near 100 hPa in the tropics during the 1997–1998 El Niño, but this anomaly does not correspond to a strong warm excursion in cold-point temperature.

Randel et al. (2006) reported the occurrence of a sudden drop in stratospheric water ~~vapour~~vapor that persisted for ~5 years during the early 2000s. This drop is evident in the cold-point temperature and SWOOSH water ~~vapour~~vapor anomalies (Fig. 15 l, m). The reanalyses generally capture the drop in stratospheric WV around 2000, with the caveat that the relaxation to a monthly mean climatology in MERRA and MERRA-2 damps the associated signals above the lowermost stratosphere.

## 6 Conclusions

In this paper, we described the basic treatment of ozone and water ~~vapour~~vapor in reanalyses, and presented comparisons both among reanalyses and between reanalyses and observations (both assimilated and independent). Here we briefly summarize the most influential characteristics and differences in the treatment of ozone and water ~~vapour~~vapor in reanalyses along with the key results of the intercomparisons.

The treatment of ozone and water ~~vapour~~vapor varies substantially among reanalyses. Some reanalyses prescribe ozone climatologies and do not treat ozone prognostically (R1, R2), some reanalyses specify ozone as a boundary condition generated by an offline chemical transport model (JRA-25, JRA-55), and some reanalyses treat ozone as a prognostic variable with parameterized photochemical production and loss (CFSR, ERA-40, ERA-Interim, MERRA, MERRA-2). Only ERA-40 and ERA-Interim contain a parameterization of heterogeneous ozone loss processes.

The reanalyses also assimilate different sets of ozone observations, with generally similar observation usage for reanalyses produced by the same reanalysis ~~entrecenter~~. All reanalyses that assimilate ozone observations rely heavily on total column ozone observations from some combination of satellites carrying the TOMS and SBUV sensors. Several recent reanalyses (including MERRA-2 and ERA-Interim) use the newest generation of vertically resolved ozone measurements (e.g., Aura MLS).

Reanalyses all assimilate tropospheric humidity information via some combination of radiosondes, satellite radiances, GNSS-RO bending angles, and retrievals of atmospheric hydrological quantities (e.g., total column water ~~vapour~~vapor or rain rate). None of the reanalyses assimilate WV observations in the stratosphere, although information from tropospheric observations may propagate upward in some systems. Beyond these similarities, the treatment of stratospheric water ~~vapour~~vapor varies substantially among the reanalyses. For example, the specific cut-off altitude up to which radiosonde humidity data are assimilated varies from one reanalysis to another, using either a fixed pressure level or the diagnosed tropopause. ERA-40 and ERA-Interim are the only reanalyses that include a water ~~vapour~~vapor source from methane oxidation. MERRA and MERRA-2 relax their fields to a water ~~vapour~~vapor climatology based on satellite observations (e.g., including Aura MLS), while other reanalyses simply do not provide valid data in the stratosphere (e.g., CSFR, JRA-25, JRA-55, R1, R2). These latter reanalyses prescribe a climatology or constant value for stratospheric water ~~vapour~~vapor as input to the forecast model radiative transfer code.

Given these differences amongst reanalysis treatments of ozone and WV, it is perhaps unsurprising that comparisons between reanalyses and observations also vary widely. Comparisons against assimilated observations of total column ozone (TCO) show that reanalyses generally reproduce TCO well, within ~10 DU (~3%). Key limitations that result in larger errors and uncertainties include a general lack of TCO data during polar night and the absence of heterogeneous chemistry from most reanalysis ozone schemes (except in ERA-40 and ERA-interim where it is introduced as a simple parameterization activated when the local temperature falls below 195K). The vertical distributions of stratospheric ozone and WV in reanalyses are unconstrained by observations through most of the record, owing to vertically-resolved data generally not being used in the assimilation systems. The situation for ozone is slightly better than that for WV, because stratospheric ozone observations are assimilated and because the ozone parameterizations are more advanced.

From the middle to upper stratosphere, reanalysis ozone profiles are within  $\pm 20\%$  of observations from the SPARC Data Initiative, although the comparisons are not truly independent for MERRA-2 or ERA-Interim because they assimilate data from Aura MLS, one of the instruments that contribute to the SPARC Data Initiative dataset. In the upper troposphere and lower stratosphere, biases increase to  $\pm 50\%$  for ozone.

MERRA-2 performs particularly well for ozone through much of the stratosphere. This is mainly due to the assimilation of the vertically resolved Aura MLS observations, which have helped to address difficulties in reproducing vertical distributions of ozone, particularly during polar night; however, these data are only available since late 2004 and are only assimilated by a few reanalyses. The use of reanalysis ozone for Antarctic ozone hole studies is therefore problematic. The reanalyses produce reasonable ozone holes when observations are available, but the timing and area of reanalysis ozone holes is highly biased when observations are unavailable. Also, apart from JRA-55, most reanalyses seem to exhibit a drift in the extent of the ozone hole area when compared to TOMS/OMI observations.

None of the reanalyses assimilate observations of stratospheric water ~~vapour~~vapor, resulting in large differences between reanalyses and independent observations. CFSR has an extreme dry bias in the stratosphere through 2009, with monthly mean values often approaching 0 ppmv. Although MERRA and MERRA-2 produce reasonable values for stratospheric



water ~~vapour~~vapor, these values represent a strong relaxation to a fixed annual climatology at pressures less than 50 hPa. Hence, mid- and upper-stratospheric water ~~vapour~~vapor does not undergo physically meaningful variations in MERRA or MERRA-2. ERA-40 and ERA-Interim produce a true “prognostic” water ~~vapour~~vapor field in the stratosphere. ERA-Interim produces surprisingly reasonable values given that its field is predominantly controlled by dehydration in the TTL and a very simple parameterization of methane oxidation. In the upper troposphere and lower stratosphere, reanalyses are around a factor of two wetter than the SPARC Data Initiative instruments used here, although the observations also have relatively large disagreements in this region.

Because of the lack of assimilated observations and the deficiencies in representation of the relevant physical processes, we recommend that reanalysis stratospheric water ~~vapour~~vapor fields should generally not be used for scientific data analysis, and stress that any examination of these fields must account for their inherent limitations and uncertainties. Future efforts toward the collection and assimilation of observational data with sensitivity to stratospheric water ~~vapour~~vapor, the reduction of reanalysis temperature biases in the TTL, and improvements in the representation of processes that control the entry mixing ratios or subsequent evolution of water ~~vapour~~vapor in the stratosphere could facilitate more reliable stratospheric water ~~vapour~~vapor fields in reanalyses.

## 15 **Code availability**

Code for creating the common-grid data files and plots are available from the corresponding author upon request.

## **Data availability**

The reanalysis data files necessary to create the “common grid” data files used here are available through the CREATE project website (<https://cds.nccs.nasa.gov/tools-services/create/>). Reanalysis total column ozone data was downloaded from the NCAR RDA (<https://rda.ucar.edu/>). SBUV data are available at [https://acd-ext.gsfc.nasa.gov/Data\\_services/merged/](https://acd-ext.gsfc.nasa.gov/Data_services/merged/). TOMS/OMI data are available at [https://disc.gsfc.nasa.gov/Aura/data-holdings/OMI/omto3d\\_v003.shtml](https://disc.gsfc.nasa.gov/Aura/data-holdings/OMI/omto3d_v003.shtml). SPARC DI data are available at <http://www.sparc-climate.org/data-center/data-access/sparc-data-initiative/>. Aura MLS satellite data are available at <https://disc.sci.gsfc.nasa.gov/Aura/data-holdings/MLS>. SWOOSH data are available at <https://www.esrl.noaa.gov/csd/swoosh/>.

25

## Appendix A

Major abbreviations and terms are defined below.

- ID-Var: 1-dimensional variational data assimilation scheme
- MIM Multi-Instrument Mean
- 5 20CR: 20th Century Reanalysis of NOAA and CIRES
- AIRS: Atmospheric Infrared Sounder
- Aqua: a satellite in the EOS A-Train satellite constellation
- ATMS: Advanced Technology Microwave Sounder
- ATOVS: Advanced TIROS Operational Vertical Sounder
- 10 Aura: a satellite in the EOS A-Train satellite constellation
- CDAS: Climate Data Assimilation System
- CFC: chlorofluorocarbon
- CFSR: Climate Forecast System Reanalysis of NCEP
- CFSv2: Climate Forecast System, version 2
- 15 CHEM2D: The NRL 2-Dimensional photochemical model
- CHEM2D-OPP: CHEM2D Ozone Photochemistry Parameterization
- CIRES: Cooperative Institute for Research in Environmental Sciences (NOAA and University of Colorado Boulder)
- CREATE: Collaborative REAnalysis Technical Environment
- CrIS: Cross-track Infrared Sounder
- 20 CTM: chemical transport model
- ECMWF: European Centre for Medium-Range Weather Forecasts
- EOS: NASA's Earth Observing System
- ERA-15: ECMWF 15-year reanalysis
- ERA-20C: ECMWF 20th century reanalysis
- 25 ERA-40: ECMWF 40-year reanalysis
- ERA5: A forthcoming reanalysis developed by ECMWF
- ERA-Interim: ECMWF interim reanalysis
- GFS: Global Forecast System of the NCEP
- GNSS-RO: Global Navigation Satellite System Radio Occultation (see also GPS-RO)
- 30 GPS-RO: Global Positioning System Radio Occultation (see also GNSS-RO)
- [GSI: Gridpoint Statistical Interpolation](#)
- HIRS: High-resolution Infrared Radiation Sounder
- IASI: Infrared Atmospheric Sounding Interferometer

IFS: Integrated Forecast System of the ECMWF  
IR: Infrared  
JCDAS: JMA Climate Data Assimilation System  
JMA: Japan Meteorological Agency  
5 JRA-25: Japanese 25-year Reanalysis  
JRA-55: Japanese 55-year Reanalysis  
JRA-55AMIP: Japanese 55-year Reanalysis based on AMIP-type simulations  
JRA-55C: Japanese 55-year Reanalysis assimilating Conventional observations only  
MERRA: Modern Era Retrospective-Analysis for Research [and Applications](#)  
10 MIM: Multi Instrument Mean  
MIPAS: Michelson Interferometer for Passive Atmospheric Sounding  
MLS: Microwave Limb Sounder  
MRI-CCM1: Meteorological Research Institute (JMA) Chemistry Climate Model, version 1  
MSU: Microwave Sounding Unit  
15 NASA: National Aeronautics and Space Administration  
NCAR: National Center for Atmospheric Research  
NCEP: National Centers for Environmental Prediction of the NOAA  
NMC: National Meteorological Center (now NCEP)  
NOAA: National Oceanic and Atmospheric Administration  
20 NRL: Naval Research Laboratory  
ODS: Ozone Depleting Substance  
OMI: Ozone Monitoring Instrument  
QBO: quasi-biennial oscillation  
R1: NCEP-NCAR Reanalysis 1  
25 R2: NCEP-DOE Reanalysis 2  
RDA: Research Data Archive  
RH: Relative Humidity  
RTTOV: Radiative Transfer for TOVS  
SEVIRI: Spinning Enhanced Visible and InfraRed Imager  
30 SBUV & SBUV/2: Solar Backscatter Ultraviolet Radiometer  
SCIAMACHY: Scanning Imaging Absorption Spectrometer for Atmospheric Chartography  
SDI: SPARC Data Initiative  
SPARC: Stratosphere-troposphere Processes And their Role in Climate  
S-RIP: SPARC Reanalysis Intercomparison Project

SSM/I or SSMI: Special Sensor Microwave Imager

SSI: Spectral Statistical Interpolation

SSU: Stratospheric Sounding Unit

TCWV: Total Column Water Vapor

5 TCO: Total Column Ozone

TIROS: Television Infrared Observation Satellite

TMI: Tropical Rainfall Measuring Mission (TRMM) Microwave Imager

TOA: top of atmosphere

TOMS: Total Ozone Mapping Spectrometer

10 TOVS: TIROS Operational Vertical Sounder

TTL: Tropical tropopause layer

UV: Ultraviolet

VTPR: Vertical Temperature Profile Radiometer

WV: Water Vapor

15

*Acknowledgements.* We thank the World Climate Research Programme and SPARC for supporting the S-RIP project. The authors are grateful to both past and present SPARC co-chairs Ted Shepherd, Greg Bodeker, Joan Alexander, and Neil Harris for their support and encouragement of S-RIP. The Information Initiative Center of Hokkaido University, Japan has hosted the S-RIP web server since 2014. We thank the reanalysis [centres](#) for providing their support and data products. The British Atmospheric Data Centre (BADC) of the UK Centre for Environmental Data Analysis (CEDA) has provided a virtual machine for data processing and a group workspace for data storage. We thank Peter Haynes, Gabriele Stiller, and William Lahoz for serving as the editors for the special issue “The SPARC Reanalysis Intercomparison Project (S-RIP)” in this journal. We thank Karen Rosenlof for valuable comments and suggestions on this manuscript. MF's contribution was financially supported in part by the Japan Society for the Promotion of Science (JSPS) through Grants-in-Aid for Scientific Research (26287117 and 16K05548). Work at the Jet Propulsion Laboratory, California Institute of Technology, was carried out under a contract with the National Aeronautics and Space Administration.

## References

- Abalos, M., Legras, B., Ploeger, F., and Randel, W. J.: Evaluating the advective Brewer-Dobson circulation in three reanalyses for the period 1979-2012, *J Geophys Res-Atmos*, 120, 7534-7554, 10.1002/2015JD023182, 2015.
- Arguez, A., and Vose, R. S.: The Definition of the Standard WMO Climate Normal The Key to Deriving Alternative  
5 Climate Normals, *B Am Meteorol Soc*, 92, 699-U345, 10.1175/2010BAMS2955.1, 2011.
- Bechtold, P., Chaboureaud, J. P., Beljaars, A., Betts, A. K., Kohler, M., Miller, M., and Redelsperger, J. L.: The simulation of the diurnal cycle of convective precipitation over land in a global model, *Q J Roy Meteor Soc*, 130, 3119-3137, 10.1256/qj.03.103, 2004.
- Bhartia, P. K., and Wellemeyer, C. W.: TOMS-V8 total O3 algorithm, in: *OMI Algorithm Theoretical Basis Document:*  
10 *OMI Ozone Products*, edited by: Bhartia, P. K., ATBD-OMI-02, 15–32, 2002.
- Bhartia, P. K., McPeters, R. D., Flynn, L. E., Taylor, S., Kramarova, N. A., Frith, S., Fisher, B., and DeLand, M.: Solar Backscatter UV (SBUV) total ozone and profile algorithm, *Atmos Meas Tech*, 6, 2533-2548, 10.5194/amt-6-2533-2013, 2013.
- Bosilovich, M., Akella, S., Coy, L., Cullather, R., Draper, C., Gelaro, R., Kovach, R., Liu, Q., Molod, A., Norris, P.,  
15 Wargan, K., Chao, W., Reichle, R., Takacs, L., Vikhliayev, Y., Bloom, S., Collow, A., Firth, S., Labow, G., Partyka, G., Pawson, S., Reale, O., Schubert, S. D., and Suarez, M.: MERRA-2: Initial Evaluation of the Climate, in: *Technical Report Series on Global Modeling and Data Assimilation*, edited by: Koster, R. D., NASA/TM–2015-104606, 43 2015.
- Butchart, N., and Remsberg, E. E.: The Area of the Stratospheric Polar Vortex as a Diagnostic for Tracer Transport on an Isentropic Surface, *J Atmos Sci*, 43, 1319-1339, Doi 10.1175/1520-0469(1986)043<1319:Taotsp>2.0.Co;2, 1986.
- 20 Cariolle, D., and Deque, M.: Southern-Hemisphere Medium-Scale Waves and Total Ozone Disturbances in a Spectral General-Circulation Model, *J Geophys Res-Atmos*, 91, 825-846, Doi 10.1029/Jd091id10p10825, 1986.
- Cariolle, D., and Teysse re, H.: A revised linear ozone photochemistry parameterization for use in transport and general circulation models: multi-annual simulations, *Atmos Chem Phys*, 7, 2183-2196, 2007.
- Chipperfield, M. P., Gray, L. J., Kinnersley, J. S., and Zawodny, J.: A 2-Dimensional Model Study of the Qbo Signal in  
25 Sage-Ii No2 and O3, *Geophys Res Lett*, 21, 589-592, Doi 10.1029/94gl00211, 1994.

- Coy, L., Wargan, K., Molod, A. M., McCarty, W. R., and Pawson, S.: Structure and Dynamics of the Quasi-Biennial Oscillation in MERRA-2, *J Climate*, 29, 5339-5354, 10.1175/JCLI-D-15-0809.1, 2016.
- Davis, S. M., Rosenlof, K. H., Hassler, B., Hurst, D. F., Read, W. G., Vömel, H., Selkirk, H., Fujiwara, M., and Damadeo, R.: The Stratospheric Water and Ozone Satellite Homogenized (SWOOSH) database: A long-term database for climate studies, *Earth Syst. Sci. Data*, 8, 461-490, doi:10.5194/essd-8-461-2016, 2016.
- Dee, D.: Importance of satellites for stratospheric data assimilation, Seminar on Recent developments in the use of satellite observations in numerical weather prediction, ECMWF, Reading, UK, 3-7 September 2007, 2008.
- Dee, D. P., and Da Silva, A. M.: The choice of variable for atmospheric moisture analysis, *Mon Weather Rev*, 131, 155-171, Doi 10.1175/1520-0493(2003), 2003.
- 10 Dee, D. P., Uppala, S. M., Simmons, A. J., Berrisford, P., Poli, P., Kobayashi, S., Andrae, U., Balmaseda, M. A., Balsamo, G., Bauer, P., Bechtold, P., Beljaars, A. C. M., van de Berg, L., Bidlot, J., Bormann, N., Delsol, C., Dragani, R., Fuentes, M., Geer, A. J., Haimberger, L., Healy, S. B., Hersbach, H., Holm, E. V., Isaksen, L., Kallberg, P., Kohler, M., Matricardi, M., McNally, A. P., Monge-Sanz, B. M., Morcrette, J. J., Park, B. K., Peubey, C., de Rosnay, P., Tavolato, C., Thepaut, J. N., and Vitart, F.: The ERA-Interim reanalysis: configuration and performance of the data assimilation system, *Q J Roy Meteor Soc*, 137, 553-597, 2011.
- 15
- Dessler, A. E., and Davis, S. M.: Trends in tropospheric humidity from reanalysis systems, *J Geophys Res-Atmos*, 115, 10.1029/2010jd014192, 2010.
- Dethof, A., and Hólm, E. V.: Ozone assimilation in the ERA-40 reanalysis project, *Q J Roy Meteor Soc*, 130, 2851-2872, 10.1256/qj.03.196, 2004.
- 20 Ebisuzaki, W., and Zhang, L.: Assessing the performance of the CFSR by an ensemble of analyses, *Clim Dynam*, 37, 2541-2550, 2011.
- Flynn, L.: Solar Backscatter Ultraviolet Instrument (SBUV/2) Version 8 Ozone Retrieval Algorithm Theoretical Basis Document (V8 ATBD), [http://ozoneaq.gsfc.nasa.gov/media/docs/SBUV2\\_V8\\_ATBD\\_020207.pdf](http://ozoneaq.gsfc.nasa.gov/media/docs/SBUV2_V8_ATBD_020207.pdf), 2007.
- Flynn, L. E., McNamara, D., Beck, C. T., Petropavlovskikh, I., Beach, E., Pachevsky, Y., Li, Y. P., Deland, M., Huang, L.
- 25 K., Long, C. S., Tiruchirapalli, R., and Taylor, S.: Measurements and products from the Solar Backscatter Ultraviolet (SBUV/2) and Ozone Mapping and Profiler Suite (OMPS) instruments, *Int J Remote Sens*, 30, 4259-4272, 2009.

- Fortuin, J. P. F., and Langematz, U.: An Update on the Global Ozone Climatology and on Concurrent Ozone and Temperature Trends, *P Soc Photo-Opt Ins*, 2311, 207-216, 1995.
- Frith, S. M., Kramarova, N. A., Stolarski, R. S., McPeters, R. D., Bhartia, P. K., and Labow, G. J.: Recent changes in total column ozone based on the SBUV Version 8.6 Merged Ozone Data Set, *J Geophys Res-Atmos*, 119, 9735-9751, 10.1002/2014JD021889, 2014.
- 5
- Fueglistaler, S., Dessler, A. E., Dunkerton, T. J., Folkins, I., Fu, Q., and Mote, P. W.: Tropical Tropopause Layer, *Rev Geophys*, 47, 2009.
- Fujiwara, M., Wright, J. S., Manney, G. L., Gray, L. J., Anstey, J., Birner, T., Davis, S., Gerber, E. P., Harvey, V. L., Hegglin, M. I., Homeyer, C. R., Knox, J. A., Krüger, K., Lambert, A., Long, C. S., Martineau, P., Molod, A., Monge-Sanz, B. M., Santee, M. L., Tegtmeier, S., Chabrilat, S., Tan, D. G. H., Jackson, D. R., Polavarapu, S., Compo, G. P., Dragani, R., Ebisuzaki, W., Harada, Y., Kobayashi, C., McCarty, W., Onogi, K., Pawson, S., Simmons, A., Wargan, K., Whitaker, J. S., and Zou, C. Z.: Introduction to the SPARC Reanalysis Intercomparison Project (S-RIP) and overview of the reanalysis systems, *Atmos. Chem. Phys.*, 17, 1417-1452, 10.5194/acp-17-1417-2017, 2017.
- 10
- Geer, A. J., Lahoz, W. A., Bekki, S., Bormann, N., Errera, Q., Eskes, H. J., Fonteyn, D., Jackson, D. R., Juckes, M. N., Massart, S., Peuch, V. H., Rharmili, S., and Segers, A.: The ASSET intercomparison of ozone analyses: method and first results, *Atmos. Chem. Phys.*, 6, 5445-5474, 10.5194/acp-6-5445-2006, 2006.
- 15
- Gelaro, R., McCarty, W., Suárez, M. J., Todling, R., Molod, A., Takacs, L., Randles, C. A., Darmenov, A., Bosilovich, M. G., Reichle, R., Wargan, K., Coy, L., Cullather, R., Draper, C., Akella, S., Buchard, V., Conaty, A., Silva, A. M. d., Gu, W., Kim, G.-K., Koster, R., Lucchesi, R., Merkova, D., Nielsen, J. E., Partyka, G., Pawson, S., Putman, W., Rienecker, M., Schubert, S. D., Sienkiewicz, M., and Zhao, B.: The Modern-Era Retrospective Analysis for Research and Applications, Version 2 (MERRA-2), *J Climate*, 30, 5419-5454, 10.1175/jcli-d-16-0758.1, 2017.
- 20
- Gottelman, A., Hoor, P., Pan, L. L., Randel, W. J., Hegglin, M. I., and Birner, T.: The extratropical upper troposphere and lower stratosphere *Rev Geophys*, 49, n/a-n/a, 10.1029/2011RG000355, 2011.
- Hasebe, F.: Quasi-Biennial Oscillations of Ozone and Diabatic Circulation in the Equatorial Stratosphere, *J Atmos Sci*, 51, 729-745, Doi 10.1175/1520-0469(1994)051<0729:Qboooa>2.0.Co;2, 1994.
- 25
- Hegglin, M. I., Boone, C. D., Manney, G. L., Shepherd, T. G., Walker, K. A., Bernath, P. F., Daffer, W. H., Hoor, P., and Schiller, C.: Validation of ACE-FTS satellite data in the upper troposphere/lower stratosphere (UTLS) using non-coincident measurements, *Atmos Chem Phys*, 8, 1483-1499, 2008.



- Hegglin, M. I., Tegtmeier, S., Anderson, J., Froidevaux, L., Fuller, R., Funke, B., Jones, A., Lingenfelter, G., Lumpe, J., Pendlebury, D., Remsberg, E., Rozanov, A., Toohey, M., Urban, J., von Clarmann, T., Walker, K. A., Wang, R., and Weigel, K.: SPARC Data Initiative: Comparison of water vapor climatologies from international satellite limb sounders, *J Geophys Res-Atmos*, 118, 10.1002/jgrd.50752, 2013.
- 5 Hólm, E., Andersson, E., Beljaars, A., Lopez, P., Mahfouf, J.-F., Simmons, A. J., and Thépaut, J.-N.: Assimilation and modelling of the hydrological cycle: ECMWF's status and plans, *ECMWF Tech. Memo*, 383, 2002.
- Hólm, E.: Revision of the ECMWF humidity analysis: Construction of a Gaussian control variable, *Proceedings of the ECMWF/GEWEX Workshop on Humidity Analysis*, Reading, UK, 8 - 11 July, 2003.
- Jiang, J. H., Su, H., Zhai, C. X., Wu, L. T., Minschwaner, K., Molod, A. M., and Tompkins, A. M.: An assessment of upper  
10 troposphere and lower stratosphere water vapor in MERRA, MERRA2, and ECMWF reanalyses using Aura MLS observations, *J Geophys Res-Atmos*, 120, 2015.
- Kalnay, E., Kanamitsu, M., Kistler, R., Collins, W., Deaven, D., Gandin, L., Iredell, M., Saha, S., White, G., Woollen, J., Zhu, Y., Chelliah, M., Ebisuzaki, W., Higgins, W., Janowiak, J., Mo, K. C., Ropelewski, C., Wang, J., Leetmaa, A., Reynolds, R., Jenne, R., and Joseph, D.: The NCEP/NCAR 40-year reanalysis project, *B Am Meteorol Soc*, 77, 437-471,  
15 1996.
- Kanamitsu, M., Ebisuzaki, W., Woollen, J., Yang, S. K., Hnilo, J. J., Fiorino, M., and Potter, G. L.: Ncep-Doe Amip-Ii Reanalysis (R-2), *B Am Meteorol Soc*, 83, 1631-1643, 10.1175/BAMS-83-11-1631, 2002.
- Kistler, R., Kalnay, E., Collins, W., Saha, S., White, G., Woollen, J., Chelliah, M., Ebisuzaki, W., Kanamitsu, M., Kousky, V., van den Dool, H., Jenne, R., and Fiorino, M.: The NCEP-NCAR 50-year reanalysis: Monthly means CD-ROM and  
20 documentation, *B Am Meteorol Soc*, 82, 247-267, Doi 10.1175/1520-0477(2001)082<0247:Tnnyrm>2.3.Co;2, 2001.
- Kobayashi, S., Ota, Y., Harada, Y., Ebata, A., Moriya, M., Onoda, H., Onogi, K., Kamahori, H., Kobayashi, C., Endo, H., Miyaoka, K., and Takahashi, K.: The JRA-55 Reanalysis: General Specifications and Basic Characteristics, *J Meteorol Soc Jpn*, 93, 5-48, 10.2151/jmsj.2015-001, 2015.
- Kramarova, N. A., Frith, S. M., Bhartia, P. K., McPeters, R. D., Taylor, S. L., Fisher, B. L., Labow, G. J., and DeLand, M.  
25 T.: Validation of ozone monthly zonal mean profiles obtained from the version 8.6 Solar Backscatter Ultraviolet algorithm, *Atmospheric Chemistry and Physics*, 13, 6887-6905, 2013.

- Levelt, P. F., Van den Oord, G. H. J., Dobber, M. R., Malkki, A., Visser, H., de Vries, J., Stammes, P., Lundell, J. O. V., and Saari, H.: The Ozone Monitoring Instrument, *Ieee T Geosci Remote*, 44, 1093-1101, 10.1109/TGRS.2006.872333, 2006.
- Ling, X. D., and London, J.: The Quasi-Biennial Oscillation of Ozone in the Tropical Middle Stratosphere - a One-Dimensional Model, *J Atmos Sci*, 43, 3122-3137, Doi 10.1175/1520-0469(1986)043<3122:Tqboo>2.0.Co;2, 1986.
- 5 Livesey, N. J., Read, W. G., Wagner, P. A., Froidevaux, L., Lambert, A., Manney, G., Valle, L. F. M., Pumphrey, H. C., Santee, M. L., Schwartz, M. J., Wang, S., Fuller, R. A., Jarnot, R. F., Knosp, B. W., and Martinez, E.: Version 4.2x Level 2 Data Quality and Description Document, JPL D-33509 Rev. B, [https://mls.jpl.nasa.gov/data/v4-2\\_data\\_quality\\_document.pdf](https://mls.jpl.nasa.gov/data/v4-2_data_quality_document.pdf), 2017.
- Manney, G. L., Zurek, R. W., Oneill, A., and Swinbank, R.: On the Motion of Air through the Stratospheric Polar Vortex, *J Atmos Sci*, 51, 2973-2994, 1994.
- 10
- Manney, G. L., Michelsen, H. A., Santee, M. L., Gunson, M. R., Irion, F. W., Roche, A. E., and Livesey, N. J.: Polar vortex dynamics during spring and fall diagnosed using trace gas observations from the Atmospheric Trace Molecule Spectroscopy instrument, *J Geophys Res-Atmos*, 104, 18841-18866, Doi 10.1029/1999jd900317, 1999.
- Manney, G. L., Daffer, W. H., Zawodny, J. M., Bernath, P. F., Hoppel, K. W., Walker, K. A., Knosp, B. W., Boone, C., Remsberg, E. E., Santee, M. L., Harvey, V. L., Pawson, S., Jackson, D. R., Deaver, L., McElroy, C. T., McLinden, C. A., Drummond, J. R., Pumphrey, H. C., Lambert, A., Schwartz, M. J., Froidevaux, L., McLeod, S., Takacs, L. L., Suarez, M. J., Trepte, C. R., Cuddy, D. C., Livesey, N. J., Harwood, R. S., and Waters, J. W.: Solar occultation satellite data and derived meteorological products: Sampling issues and comparisons with Aura Microwave Limb Sounder, *J Geophys Res-Atmos*, 112, Artn D24s50
- 15
- 20 10.1029/2007jd008709, 2007.
- Manney, G. L., Hegglin, M. I., Daffer, W. H., Santee, M. L., Ray, E. A., Pawson, S., Schwartz, M. J., Boone, C. D., Froidevaux, L., Livesey, N. J., Read, W. G., and Walker, K. A.: Jet characterization in the upper troposphere/lower stratosphere (UTLS): applications to climatology and transport studies, *Atmos Chem Phys*, 11, 6115-6137, 10.5194/acp-11-6115-2011, 2011.
- 25 McCarty, W., Coy, L., Gelaro, R., Merkova, D., Smith, E. B., Sienkiewicz, M., and Wargan, K.: MERRA-2 Input Observations: Summary and Initial Assessment, NASA Technical Report Series on Global Modeling and Data Assimilation, NASA/TM-2016-104606, 46, 61, 2016.

- McCormack, J. P., Eckermann, S. D., Siskind, D. E., and McGee, T. J.: CHEM2D-OPP: A new linearized gas-phase ozone photochemistry parameterization for high-altitude NWP and climate models, *Atmos Chem Phys*, 6, 4943-4972, 2006.
- McLinden, C. A., Tegtmeier, S., and Fioletov, V.: Technical Note: A SAGE-corrected SBUV zonal-mean ozone data set, *Atmos Chem Phys*, 9, 7963-7972, 2009.
- 5 McPeters, R., Kroon, M., Labow, G., Brinksma, E., Balis, D., Petropavlovskikh, I., Veefkind, J. P., Bhartia, P. K., and Levelt, P. F.: Validation of the Aura Ozone Monitoring Instrument total column ozone product, *J Geophys Res-Atmos*, 113, Artn D15s14  
10.1029/2007jd008802, 2008.
- McPeters, R. D., Bhartia, P. K., Haffner, D., Labow, G. J., and Flynn, L.: The version 8.6 SBUV ozone data record: An  
10 overview, *J Geophys Res-Atmos*, 118, 8032-8039, 2013.
- Mote, P. W., Rosenlof, K. H., McIntyre, M. E., Carr, E. S., Gille, J. C., Holton, J. R., Kinnarsley, J. S., Pumphrey, H. C., Russell, J. M., and Waters, J. W.: An atmospheric tape recorder: The imprint of tropical tropopause temperatures on stratospheric water vapor, *J Geophys Res-Atmos*, 101, 3989-4006, Doi 10.1029/95jd03422, 1996.
- Neu, J. L., Hegglin, M. I., Tegtmeier, S., Bourassa, A., Degenstein, D., Froidevaux, L., Fuller, R., Funke, B., Gille, J., Jones,  
15 A., Rozanov, A., Toohey, M., von Clarmann, T., Walker, K. A., and Worden, J. R.: The SPARC Data Initiative: Comparison of upper troposphere/lower stratosphere ozone climatologies from limb-viewing instruments and the nadir-viewing Tropospheric Emission Spectrometer, *J Geophys Res-Atmos*, 119, 6971-6990, 2014.
- Oikonomou, E. K., and O'Neill, A.: Evaluation of ozone and water vapor fields from the ECMWF reanalysis ERA-40 during 1991,Äì1999 in comparison with UARS satellite and MOZAIC aircraft observations, *J Geophys Res*, 111, D14109,  
20 papers2://publication/doi/10.1029/2004JD005341, 2006.
- Onogi, K., Tsltsui, J., Koide, H., Sakamoto, M., Kobayashi, S., Hatsushika, H., Matsumoto, T., Yamazaki, N., Kaalhari, H., Takahashi, K., Kadokura, S., Wada, K., Kato, K., Oyama, R., Ose, T., Mannoji, N., and Taira, R.: The JRA-25 reanalysis, *J Meteorol Soc Jpn*, 85, 369-432, Doi 10.2151/Jmsj.85.369, 2007.
- Paltridge, G., Arking, A., and Pook, M.: Trends in middle- and upper-level tropospheric humidity from NCEP reanalysis  
25 data, *Theor Appl Climatol*, 98, 351-359, 10.1007/s00704-009-0117-x, 2009.

- Pan, L. L., Kunz, A., Homeyer, C. R., Munchak, L. A., Kinnison, D. E., and Tilmes, S.: Commentary on using equivalent latitude in the upper troposphere and lower stratosphere, *Atmos Chem Phys*, 12, 9187-9199, 10.5194/acp-12-9187-2012, 2012.
- Poli, P.: List of observations assimilated in ERA-40 and ERA-Interim (v1.0),  
5 <http://www.ecmwf.int/sites/default/files/elibrary/2010/11692-list-observations-assimilated-era-40-and-era-interim-v10.pdf>, 2010.
- Randel, W. J., Wu, F., Russell, J. M., Roche, A., and Waters, J. W.: Seasonal cycles and QBO variations in stratospheric CH<sub>4</sub> and H<sub>2</sub>O observed in UARS HALOE data, *J Atmos Sci*, 55, 163-185, Doi 10.1175/1520-0469(1998)055<0163:Scaqvi>2.0.Co;2, 1998.
- 10 Randel, W. J., Wu, F., Vomel, H., Nedoluha, G. E., and Forster, P.: Decreases in stratospheric water vapor after 2001: Links to changes in the tropical tropopause and the Brewer-Dobson circulation, *J Geophys Res-Atmos*, 111, Artn D12312 10.1029/2005jd006744, 2006.
- Rienecker, M. M., Suarez, M. J., Gelaro, R., Todling, R., Bacmeister, J., Liu, E., Bosilovich, M. G., Schubert, S. D., Takacs, L., Kim, G. K., Bloom, S., Chen, J. Y., Collins, D., Conaty, A., Da Silva, A., Gu, W., Joiner, J., Koster, R. D., Lucchesi, R.,  
15 Molod, A., Owens, T., Pawson, S., Pegion, P., Redder, C. R., Reichle, R., Robertson, F. R., Ruddick, A. G., Sienkiewicz, M., and Woollen, J.: MERRA: NASA's Modern-Era Retrospective Analysis for Research and Applications, *J Climate*, 24, 3624-3648, 2011.
- Saha, S., Moorthi, S., Pan, H. L., Wu, X. R., Wang, J. D., Nadiga, S., Tripp, P., Kistler, R., Woollen, J., Behringer, D., Liu, H. X., Stokes, D., Grumbine, R., Gayno, G., Wang, J., Hou, Y. T., Chuang, H. Y., Juang, H. M. H., Sela, J., Iredell, M.,  
20 Treadon, R., Kleist, D., Van Delst, P., Keyser, D., Derber, J., Ek, M., Meng, J., Wei, H. L., Yang, R. Q., Lord, S., Van den Dool, H., Kumar, A., Wang, W. Q., Long, C., Chelliah, M., Xue, Y., Huang, B. Y., Schemm, J. K., Ebisuzaki, W., Lin, R., Xie, P. P., Chen, M. Y., Zhou, S. T., Higgins, W., Zou, C. Z., Liu, Q. H., Chen, Y., Han, Y., Cucurull, L., Reynolds, R. W., Rutledge, G., and Goldberg, M.: The Ncep Climate Forecast System Reanalysis, *B Am Meteorol Soc*, 91, 1015-1057, 2010.
- Saha, S., Moorthi, S., Wu, X. R., Wang, J., Nadiga, S., Tripp, P., Behringer, D., Hou, Y. T., Chuang, H. Y., Iredell, M., Ek,  
25 M., Meng, J., Yang, R. Q., Mendez, M. P., Van Den Dool, H., Zhang, Q., Wang, W. Q., Chen, M. Y., and Becker, E.: The NCEP Climate Forecast System Version 2, *J Climate*, 27, 2185-2208, 10.1175/JCLI-D-12-00823.1, 2014.
- Santee, M. L., Manney, G. L., Livesey, N. J., Froidevaux, L., Schwartz, M. J., and Read, W. G.: Trace gas evolution in the lowermost stratosphere from Aura Microwave Limb Sounder measurements, *J Geophys Res-Atmos*, 116, 2011.

- Shibata, K., Deushi, M., Sekiyama, T. T., and Yoshimura, H.: Development of an MRI Chemical Transport Model for the Study of Stratospheric Chemistry, *Papers in Meteorology and Geophysics*, 55, 75-119, 10.2467/mripapers.55.75, 2005.
- Stajner, I., Wargan, K., Pawson, S., Hayashi, H., Chang, L. P., Hudman, R. C., Froidevaux, L., Livesey, N., Levelt, P. F., Thompson, A. M., Tarasick, D. W., Stubi, R., Andersen, S. B., Yela, M., Konig-Langlo, G., Schmidlin, F. J., and Witte, J. C.: Assimilated ozone from EOS-Aura: Evaluation of the tropopause region and tropospheric columns, *J Geophys Res-Atmos*, 113, 2008.
- Takacs, L. L., Suárez, M. J., and Todling, R.: Maintaining atmospheric mass and water balance in reanalyses, *Q J Roy Meteor Soc*, 142, 1565-1573, 10.1002/qj.2763, 2016.
- Tegtmeier, S., Hegglin, M. I., Anderson, J., Bourassa, A., Brohede, S., Degenstein, D., Froidevaux, L., Fuller, R., Funke, B., Gille, J., Jones, A., Kasai, Y., Kruger, K., Kyrola, E., Lingenfelter, G., Lumpe, J., Nardi, B., Neu, J., Pendlebury, D., Remsberg, E., Rozanov, A., Smith, L., Toohey, M., Urban, J., von Clarmann, T., Walker, K. A., and Wang, R. H. J.: SPARC Data Initiative: A comparison of ozone climatologies from international satellite limb sounders, *J Geophys Res-Atmos*, 118, 12229-12247, 10.1002/2013JD019877, 2013.
- Thornton, H. E., Jackson, D. R., Bekki, S., Bormann, N., Errera, Q., Geer, A. J., Lahoz, W. A., and Rharmili, S.: The ASSET intercomparison of stratosphere and lower mesosphere humidity analyses, *Atmos. Chem. Phys.*, 9, 995-1016, 10.5194/acp-9-995-2009, 2009.
- Tompkins, A. M., Gierens, K., and Radel, G.: Ice supersaturation in the ECMWF integrated forecast system, *Q J Roy Meteor Soc*, 133, 53-63, 10.1002/qj.14, 2007.
- Untch, A., Simmons, A., Hortal, M., and Jakob, C.: Increased stratospheric resolution in the ECMWF forecasting system, *Proceedings of the workshop on chemical data assimilation, KNMI de Bilt, Holland*, 1998.
- Uppala, S. M., Kallberg, P. W., Simmons, A. J., Andrae, U., Bechtold, V. D., Fiorino, M., Gibson, J. K., Haseler, J., Hernandez, A., Kelly, G. A., Li, X., Onogi, K., Saarinen, S., Sokka, N., Allan, R. P., Andersson, E., Arpe, K., Balmaseda, M. A., Beljaars, A. C. M., Van De Berg, L., Bidlot, J., Bormann, N., Caires, S., Chevallier, F., Dethof, A., Dragosavac, M., Fisher, M., Fuentes, M., Hagemann, S., Holm, E., Hoskins, B. J., Isaksen, I., Janssen, P. A. E. M., Jenne, R., McNally, A. P., Mahfouf, J. F., Morcrette, J. J., Rayner, N. A., Saunders, R. W., Simon, P., Sterl, A., Trenberth, K. E., Untch, A., Vasiljevic, D., Viterbo, P., and Woollen, J.: The ERA-40 re-analysis, *Q J Roy Meteor Soc*, 131, 2961-3012, 10.1256/qj.04.176, 2005.

- Vömel, H., Barnes, J. E., Forno, R. N., Fujiwara, M., Hasebe, F., Iwasaki, S., Kivi, R., Komala, N., Kyro, E., Leblanc, T., Morel, B., Ogino, S. Y., Read, W. G., Ryan, S. C., Saraspriya, S., Selkirk, H., Shiotani, M., Canossa, J. V., and Whiteman, D. N.: Validation of Aura Microwave Limb Sounder water vapor by balloon-borne Cryogenic Frost point Hygrometer measurements, *J Geophys Res-Atmos*, 112, ArtD24s37
- 5 Doi 10.1029/2007jd008698, 2007.
- Wargan, K., Pawson, S., Olsen, M. A., Witte, J. C., Douglass, A. R., Ziemke, J. R., Strahan, S. E., and Nielsen, J. E.: The global structure of upper troposphere-lower stratosphere ozone in GEOS-5: A multiyear assimilation of EOS Aura data, *J Geophys Res-Atmos*, 120, 2013-2036, 10.1002/2014JD022493, 2015.
- Wargan, K., Labow, G., Frith, S., Pawson, S., Livesey, N., and Partyka, G.: Evaluation of the Ozone Fields in NASA's
- 10 MERRA-2 Reanalysis, *J Climate*, 30, 2961-2988, 10.1175/jcli-d-16-0699.1, 2017.
- Waters, J. W., Froidevaux, L., Harwood, R. S., Jarnot, R. F., Pickett, H. M., Read, W. G., Siegel, P. H., Cofield, R. E., Filipiak, M. J., Flower, D. A., Holden, J. R., Lau, G. K. K., Livesey, N. J., Manney, G. L., Pumphrey, H. C., Santee, M. L., Wu, D. L., Cuddy, D. T., Lay, R. R., Loo, M. S., Perun, V. S., Schwartz, M. J., Stek, P. C., Thurstans, R. P., Boyles, M. A., Chandra, K. M., Chavez, M. C., Chen, G. S., Chudasama, B. V., Dodge, R., Fuller, R. A., Girard, M. A., Jiang, J. H., Jiang,
- 15 Y. B., Knosp, B. W., LaBelle, R. C., Lam, J. C., Lee, K. A., Miller, D., Oswald, J. E., Patel, N. C., Pukala, D. M., Quintero, O., Scaff, D. M., Van Snyder, W., Tope, M. C., Wagner, P. A., and Walch, M. J.: The Earth Observing System Microwave Limb Sounder (EOS MLS) on the Aura satellite, *Ieee T Geosci Remote*, 44, 1075-1092, 10.1109/TGRS.2006.873771, 2006.
- Wu, W. S., Purser, R. J., and Parrish, D. F.: Three-dimensional variational analysis with spatially inhomogeneous covariances, *Mon Weather Rev*, 130, 2905-2916, Doi 10.1175/1520-0493(2002)130<2905:Tdvaws>2.0.Co;2, 2002.
- 20 Yan, X. L., Wright, J. S., Zheng, X. D., Livesey, N. J., Vomel, H., and Zhou, X. J.: Validation of Aura MLS retrievals of temperature, water vapour and ozone in the upper troposphere and lower-middle stratosphere over the Tibetan Plateau during boreal summer, *Atmos Meas Tech*, 9, 3547-3566, 10.5194/amt-9-3547-2016, 2016.
- Zawodny, J. M., and McCormick, M. P.: Stratospheric Aerosol and Gas Experiment-Ii Measurements of the Quasi-Biennial Oscillations in Ozone and Nitrogen-Dioxide, *J Geophys Res-Atmos*, 96, 9371-9377, Doi 10.1029/91jd00517, 1991.

**Table 1.** Key characteristics of ozone treatment in reanalyses

Reanalysis	Primary TCO data sources	Vertical profile data sources	Stratospheric O <sub>3</sub> used in radiative transfer	Stratospheric O <sub>3</sub> treatment	Photochemical parameterization
NCEP R1	None	None	Climatology	None	None
NCEP R2	None	None	Climatology	None	None
CFSR	SBUV	SBUV	Analy <del>s</del> zed	Prognostic	CHEM2D-OPP
ERA-40	TOMS	SBUV	Climatology	Prognostic	CD86
ERA-I	Same as ERA-40	SBUV, GOME, MLS, MIPAS	Same as ERA-40	Same as ERA-40	Same as ERA-40
JRA-25	TOMS (1979–2004) <sup>a</sup> OMI (2004–)	Nudging to climatological profile	Daily values from offline CTM	Daily values from offline CTM	Shibata et al. (2005)
JRA-55	Same as JRA-25	None	Daily values from updated offline CTM	Daily values from updated offline CTM	Shibata et al. (2005)
MERRA	SBUV	SBUV	Analy <del>s</del> zed	Prognostic	Stajner et al. (2008)
MERRA-2	SBUV (1980–9/2004) OMI (9/2004–)	SBUV, MLS	Same as MERRA	Same as MERRA	Same as MERRA

<sup>a</sup> Offline CCM nudged to TOMS/OMI data.

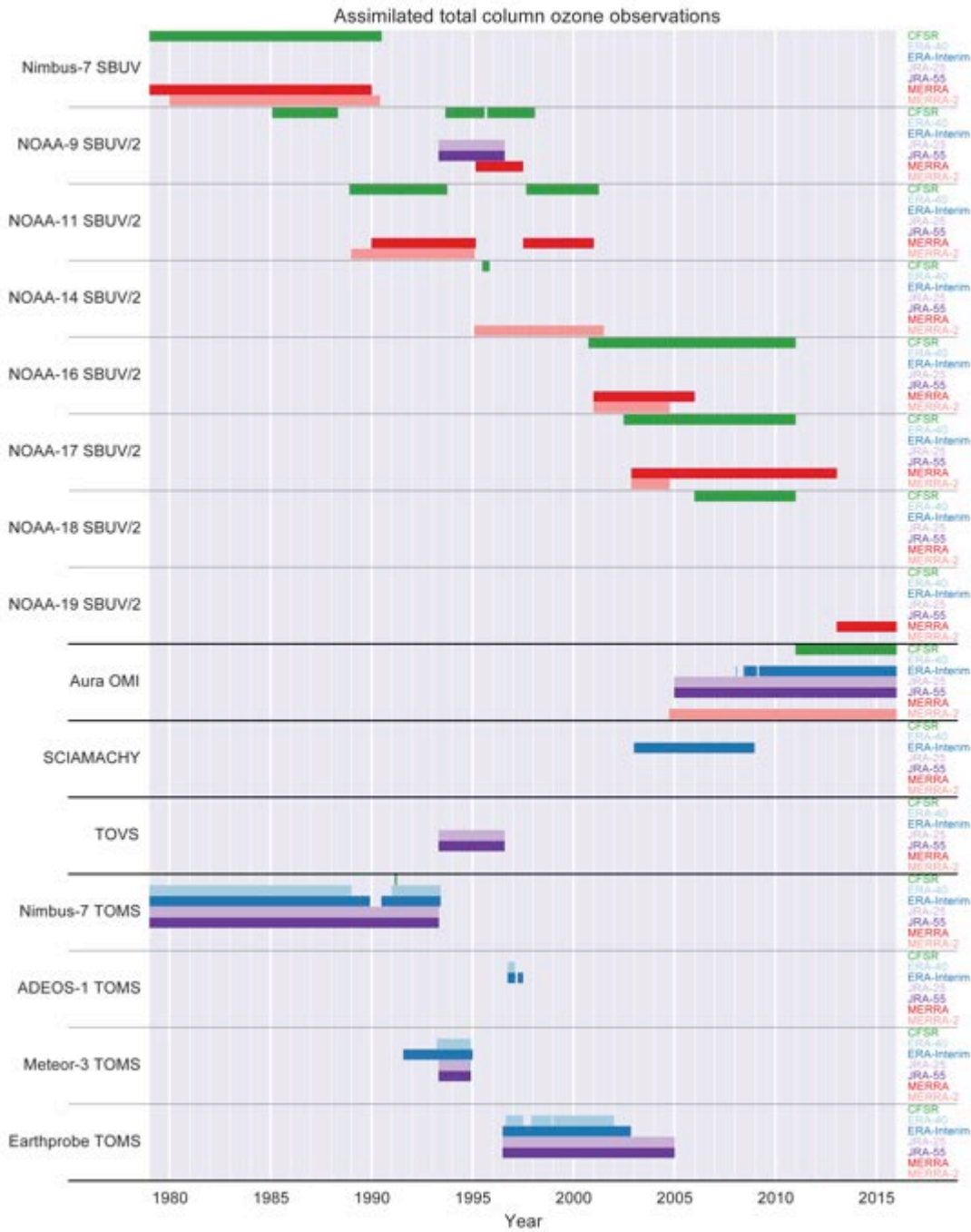
**Table 2.** Key characteristics of water ~~vapour~~vapor treatment in reanalyses

Reanalysis	Assimilation of satellite humidity radiances?	Highest level of assimilated WV observations	Highest level of analyzed WV <sup>1</sup>	Stratospheric WV used in radiative transfer	Stratospheric WV treatment	Stratospheric methane oxidation parameterization?
NCEP R1	No	300 hPa	300 hPa	Climatology	None	No
NCEP R2	No	300 hPa	10 hPa (RH only)	Climatology	None	No
CFSR	Yes	250 hPa	None	Analy <del>s</del> zed; negative values set to 0.1 ppmv	Prognostic	No
ERA-40	Yes	Diagnosed tropopause. Radiosonde humidity generally used to 300 hPa	Diagnosed tropopause	Analy <del>s</del> zed	Prognostic	Yes. Relaxation to 6 ppmv WV at stratopause
ERA-I	Yes	Same as ERA-40	Diagnosed tropopause	Analy <del>s</del> zed	Prognostic	Yes. Relaxation to 6.8 ppmv WV at stratopause
JRA-25	Yes	100 hPa	50 hPa	Constant 2.5 ppmv	Prognostic <sup>2</sup>	No
JRA-55	Yes	100 hPa	5 hPa	Climatological annual mean from HALOE and UARS MLS during 1991–1997	Prognostic <sup>2</sup>	No
MERRA	Yes	300 hPa	None	Analy <del>s</del> zed	3-day relaxation to zonal-mean monthly-mean satellite-based climatology	No
MERRA-2	Yes	300 hPa	None	Same as MERRA	Same as MERRA	No

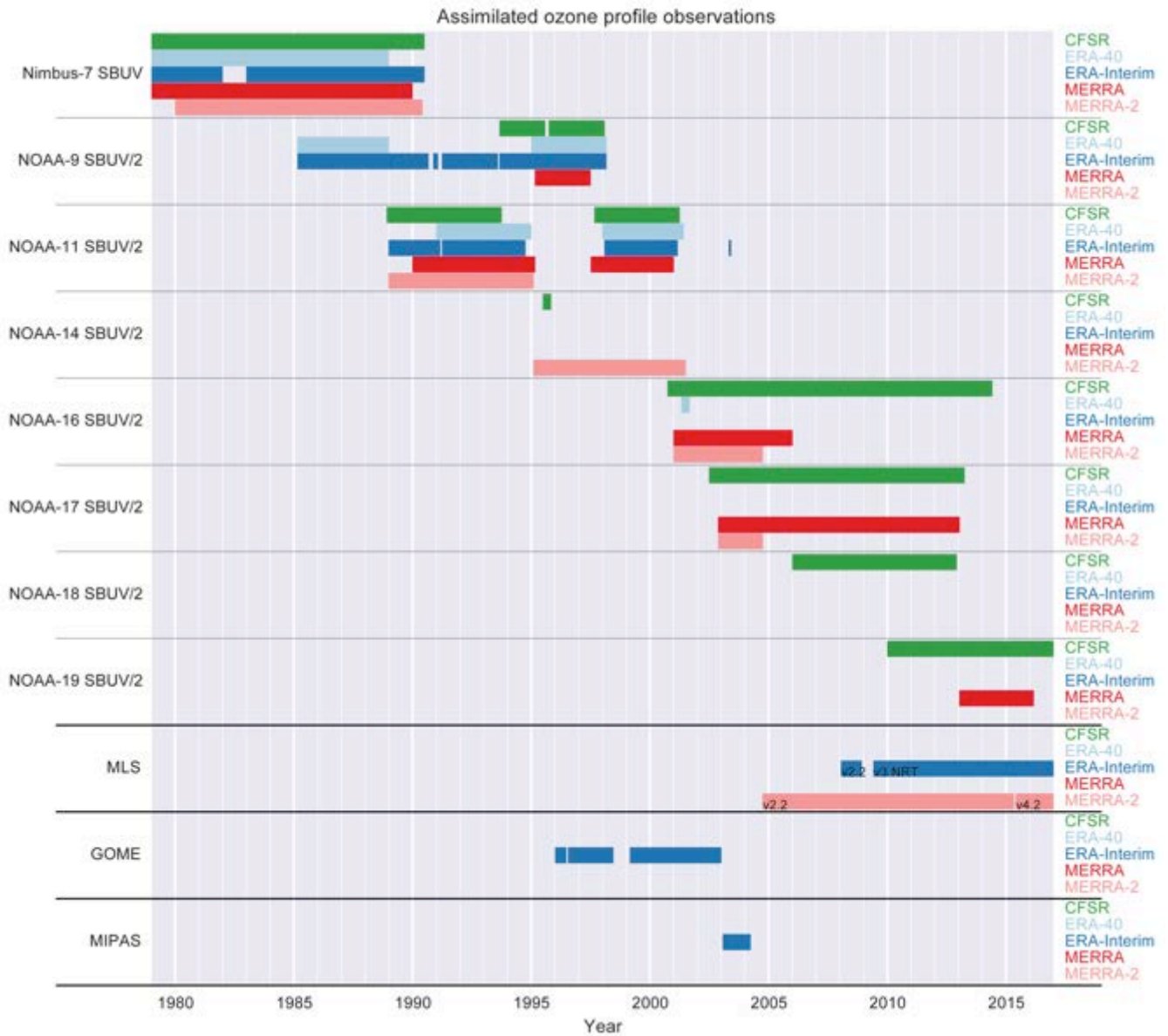
<sup>1</sup> Level above which assimilation-related increments are not allowed.

<sup>2</sup> Water ~~vapour~~vapor not provided above 100 hPa in pressure level analysis products.

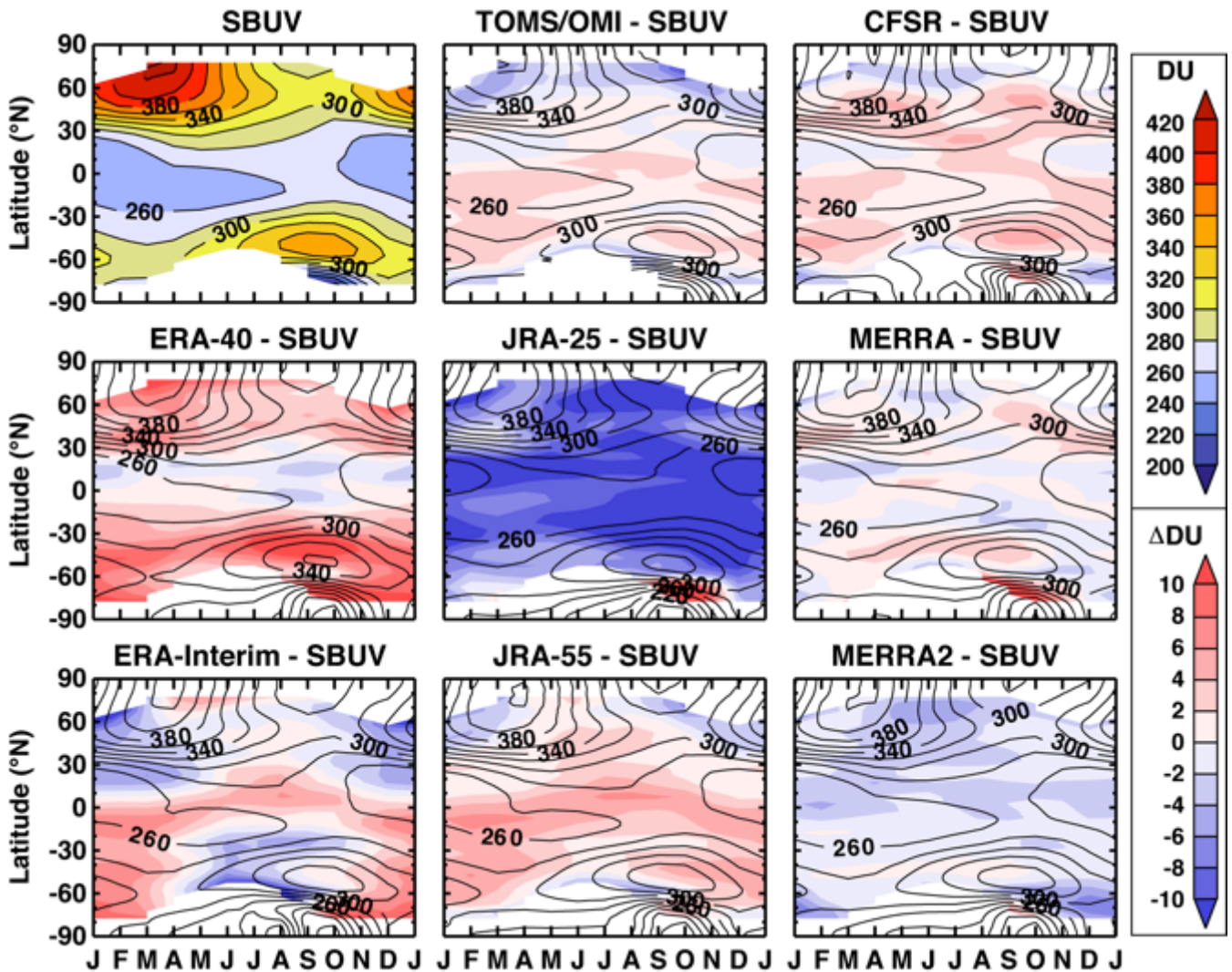




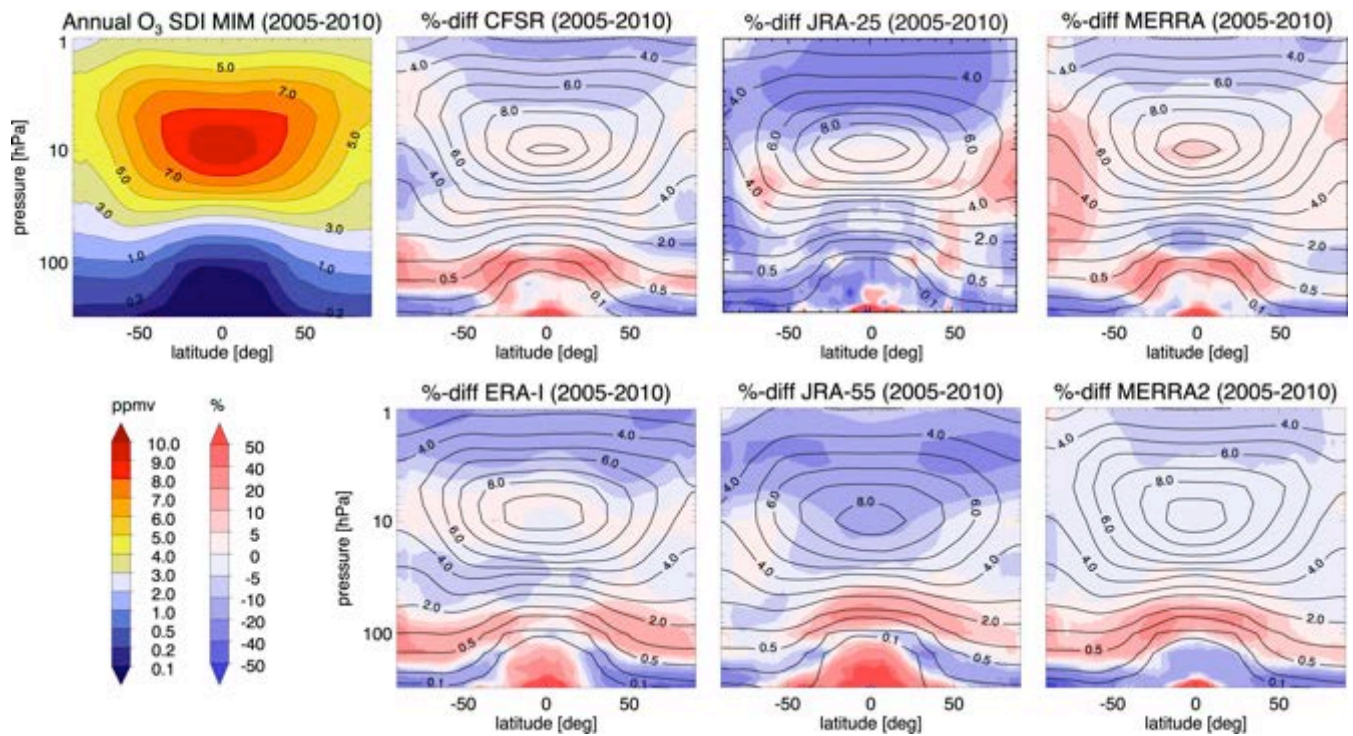
**Figure 1:** Total column ozone data by instrument as assimilated by the different reanalyses.



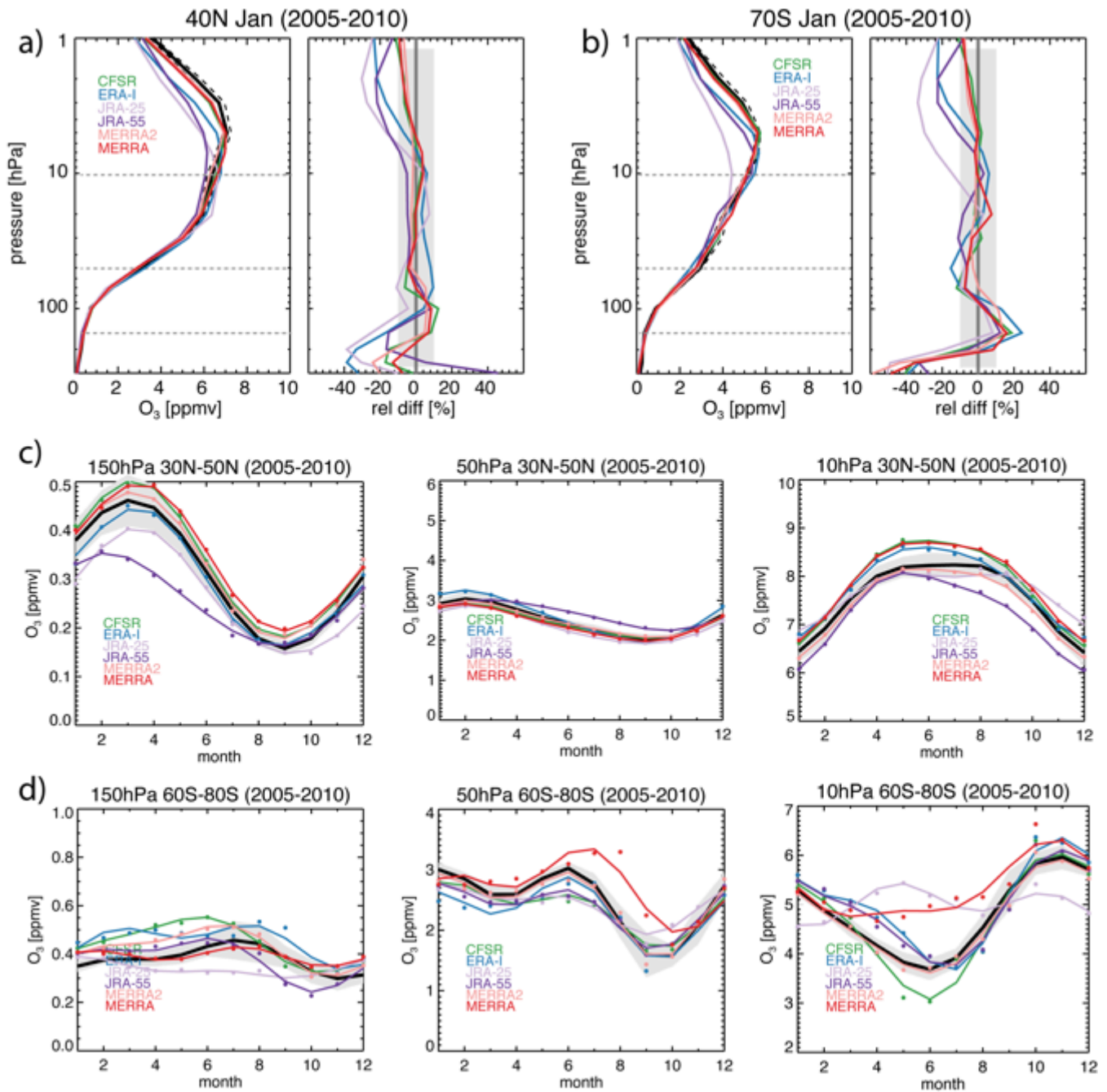
**Figure 2:** Ozone vertical profile observations by instrument as assimilated by the different reanalyses.



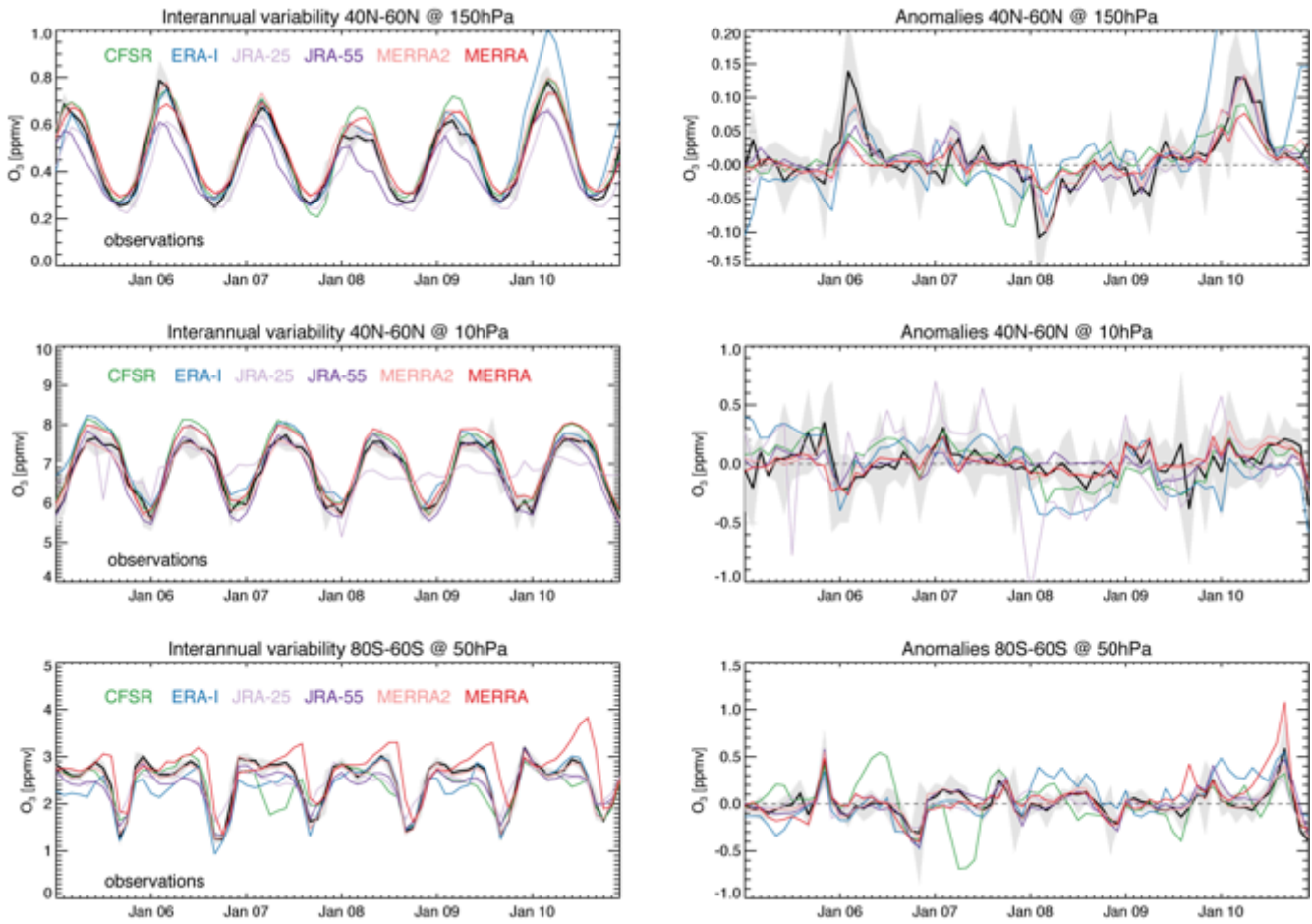
5 | **Figure 3:** Zonal- and monthly-mean total column ozone climatology over 1981–2010 from SBUV observations (uppermost left panel), along with the absolute differences between each reanalysis and SBUV. The difference between TOMS/OMI and SBUV is also shown (uppermost middle panel). (Other panels) Line contours show each reanalysis' respective climatology, and the shading shows differences from SBUV, with cool (blue) colors representing negative values and warm (red) colors representing positive values. Both climatology and observational reference to calculate differences for ERA-40 are for the time period Jan 1981–Aug 2002 in order to avoid sampling issues.



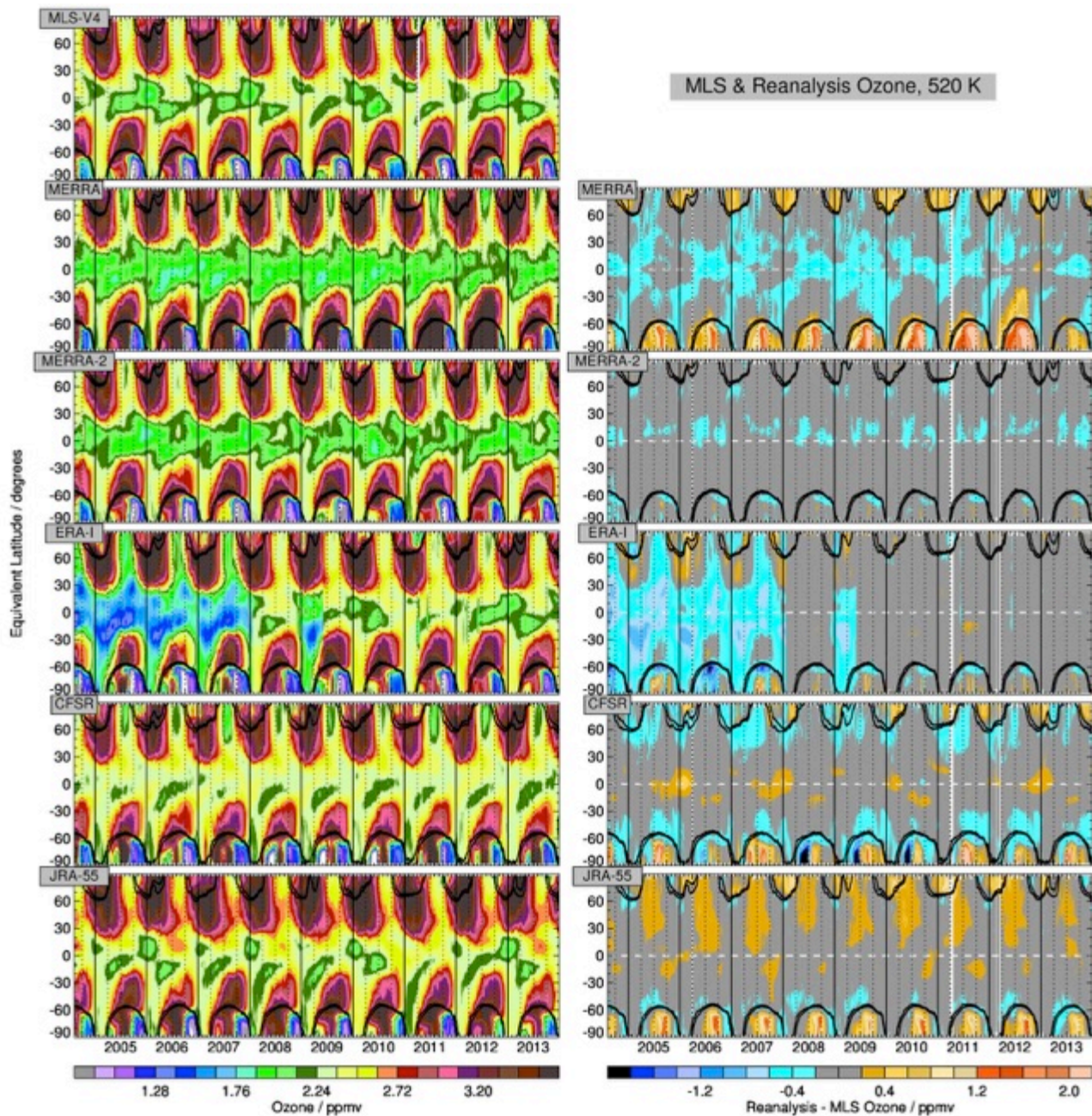
5 | **Figure 4:** Multi-annual zonal mean ozone cross sections averaged over 2005–2010 for the SPARC Data Initiative multi-instrument mean (SDI MIM) (upper left), along with the relative differences between reanalyses and observations as  $(R_i - \text{MIM})/\text{MIM} \times 100$ , where  $R_i$  is a reanalysis field (shading, other panels). Also shown in line contours are the respective zonal mean climatologies for the different reanalyses.



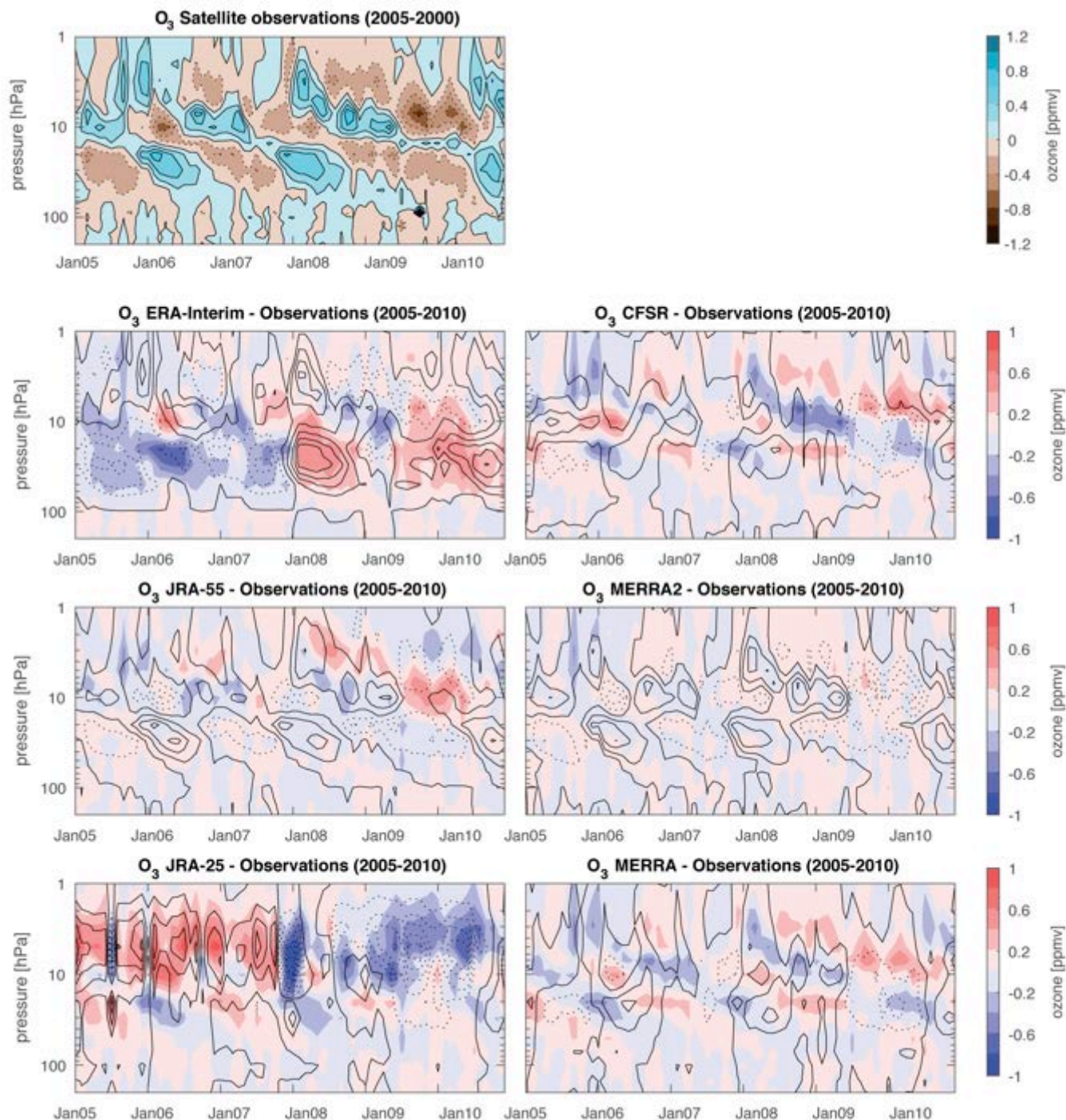
**Figure 5:** Multi-annual mean vertical ozone profiles over 2005–2010 for January at (a) 40N and (b) 70S from the SPARC Data Initiative multi-instrument mean (SDI MIM) (black) and the six reanalyses (colored). Absolute values are shown in the left and relative differences in the right panels for each comparison. Relative differences are calculated as  $(R_i - \text{MIM}) / \text{MIM} \times 100$ , where  $R_i$  is a reanalysis profile. Black dashed lines provide the  $\pm 1$ -sigma uncertainty (as calculated by the standard deviation over all instruments and years available) in the observational mean. Horizontal dashed lines in grey indicate the pressure levels (150, 50, and 10 hPa) for which seasonal cycles are shown in panels (c) and (d) for the two latitude ranges 30-50N and 60-80S, respectively. In the lower panels, the SDI MIM uncertainty is shown using grey shading.



5 **Figure 6:** Interannual variability (left column) and deseasonalized anomalies (right column) for ozone during 2005–2010 for the SPARC Data Initiative multi-instrument mean (SDI MIM, black) and the six reanalyses (colored). Results are shown for three different pressure levels and latitude ranges (top to bottom: 50 hPa at 60–80°S, 10 hPa at 20°S–20°N, and 100 hPa at 40–60°N). Grey shading indicates observational uncertainty ( $\pm 1$ -sigma) calculated as the standard deviation over all instruments and years available.



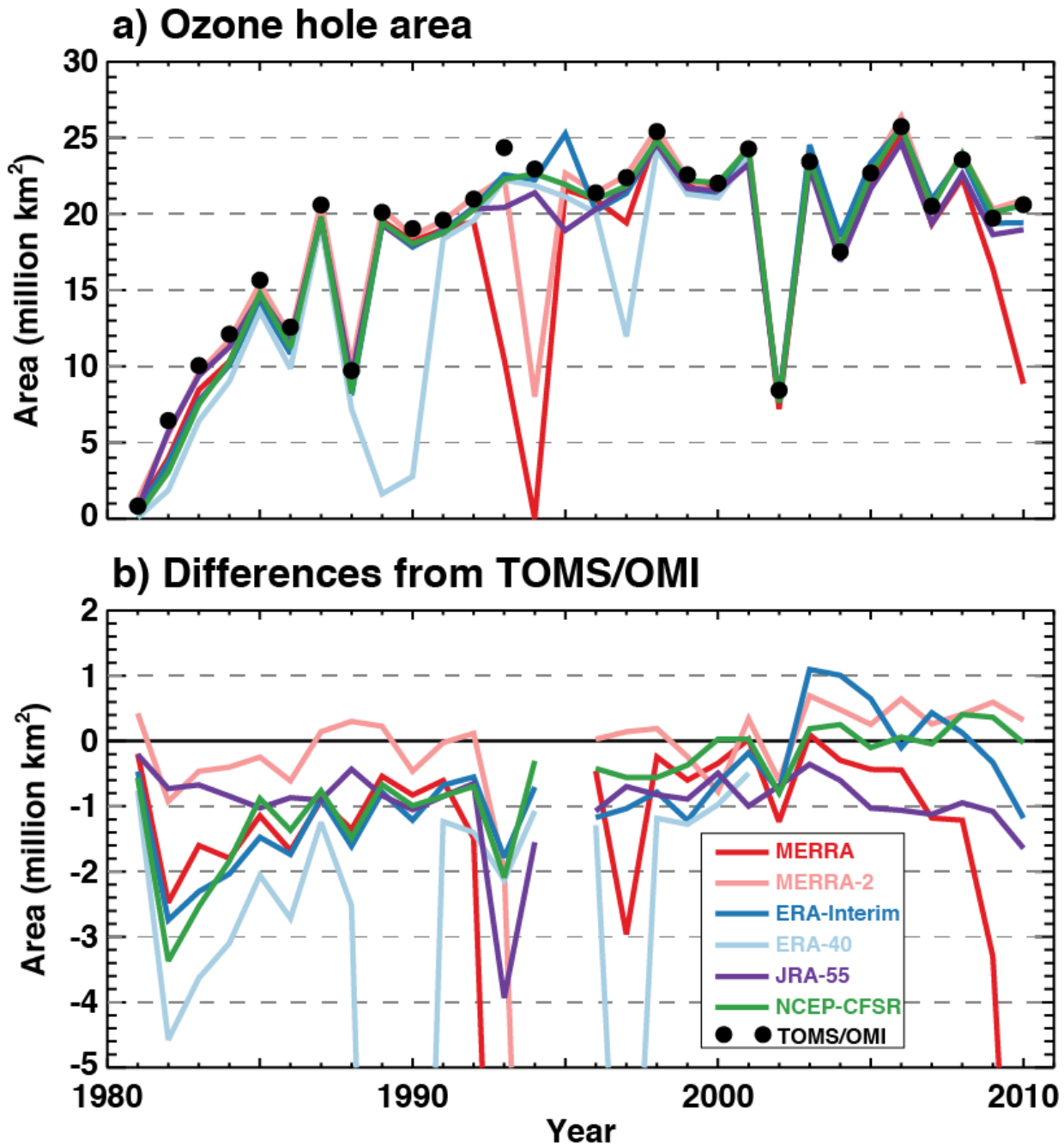
5 **Figure 7:** Comparison of the equivalent latitude–time evolution of each reanalysis ozone field and MLS on the 520 K isentropic surface ( $\sim 50$  hPa;  $\sim 20$  km altitude) during the Aura mission September 2004 – December 2013. (Left) Mixing ratios (ppmv) for MLS and the reanalyses MERRA, MERRA-2, ERA-Interim, CFSR, and JRA-55 (top to bottom). (Right) differences (ppmv) between each reanalysis and MLS ( $R_i - \text{MLS}$ ). Overlays are scaled potential vorticity (Manney et al., 1994) contours of  $1.4$  and  $1.6 \times 10^{-4} \text{ s}^{-1}$  from the corresponding reanalysis, which are intended to represent the wintertime polar vortex edge. Dynamical fields for the MLS panel are from MERRA.



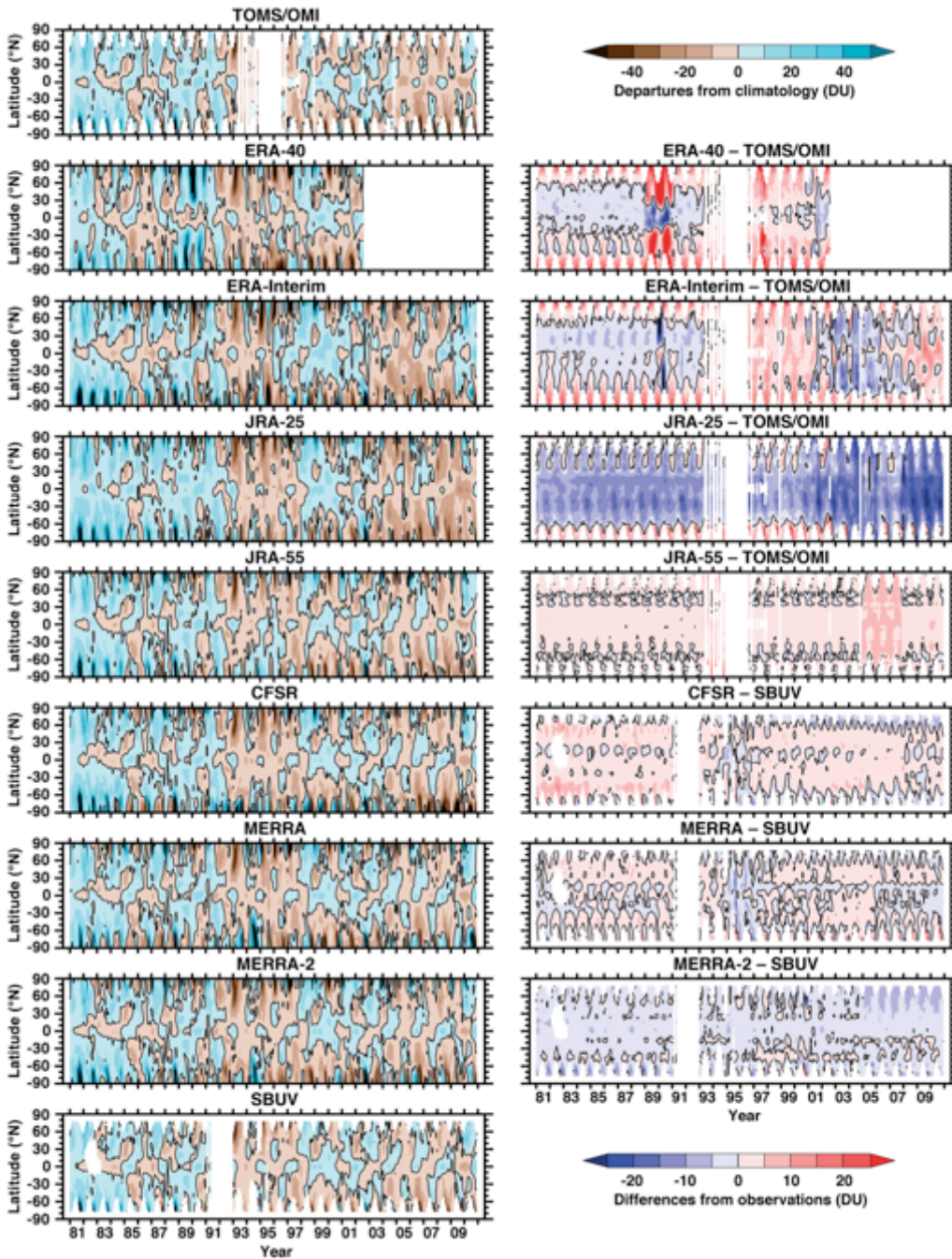
**Figure 8:** QBO ozone signal from the SPARC Data Initiative observations (upper left) during 2005–2010, defined as altitude–time cross-sections of deseasonalized ozone anomalies averaged over the 10°S–10°N tropical band. Observations are based on three satellite data sets. The other panels show the differences in QBO ozone signals between each reanalysis and the observations ( $R_i$ -MIM, shaded contours) with the black line contours showing the QBO ozone signal generated by each corresponding reanalysis.

5

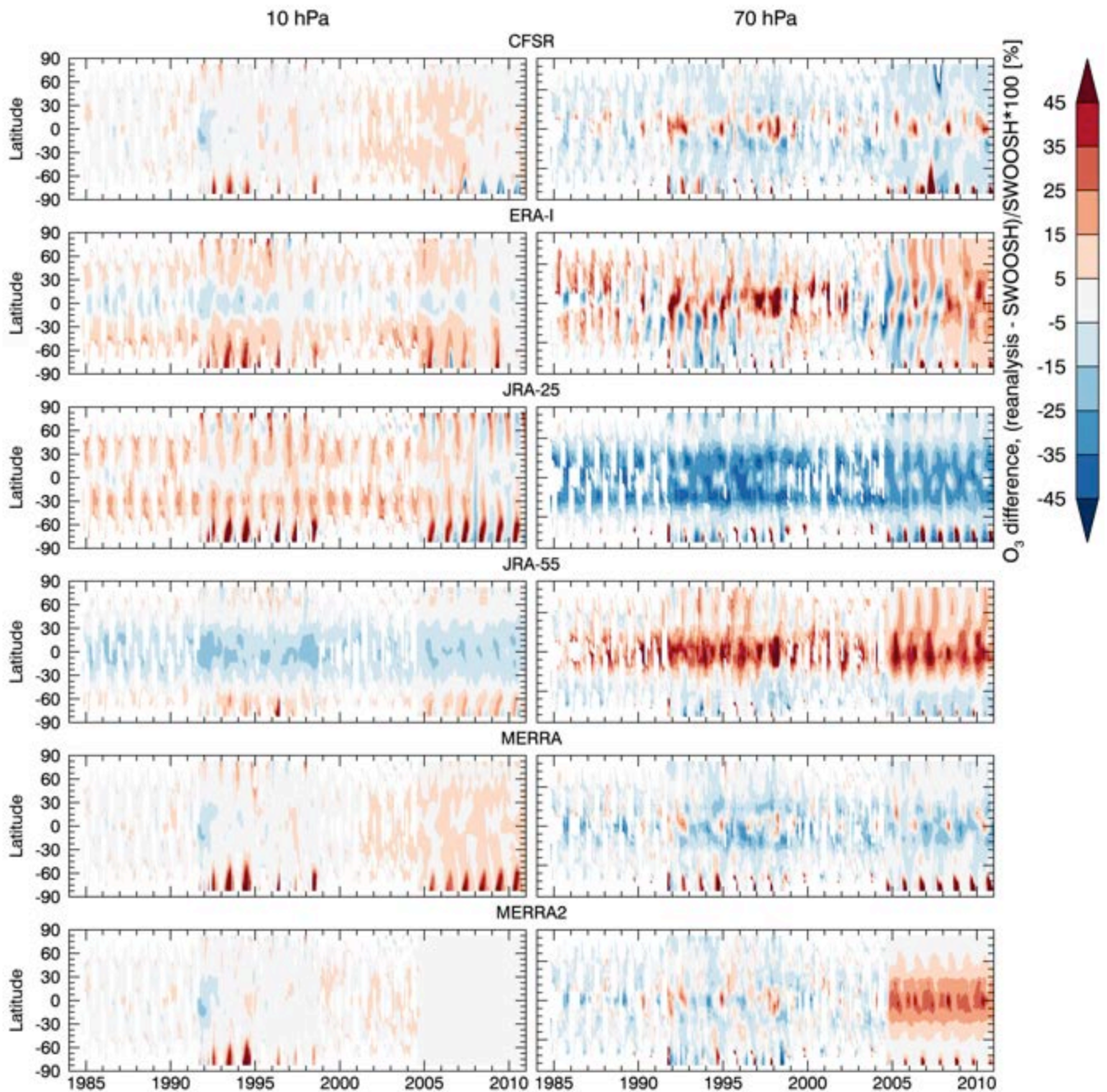




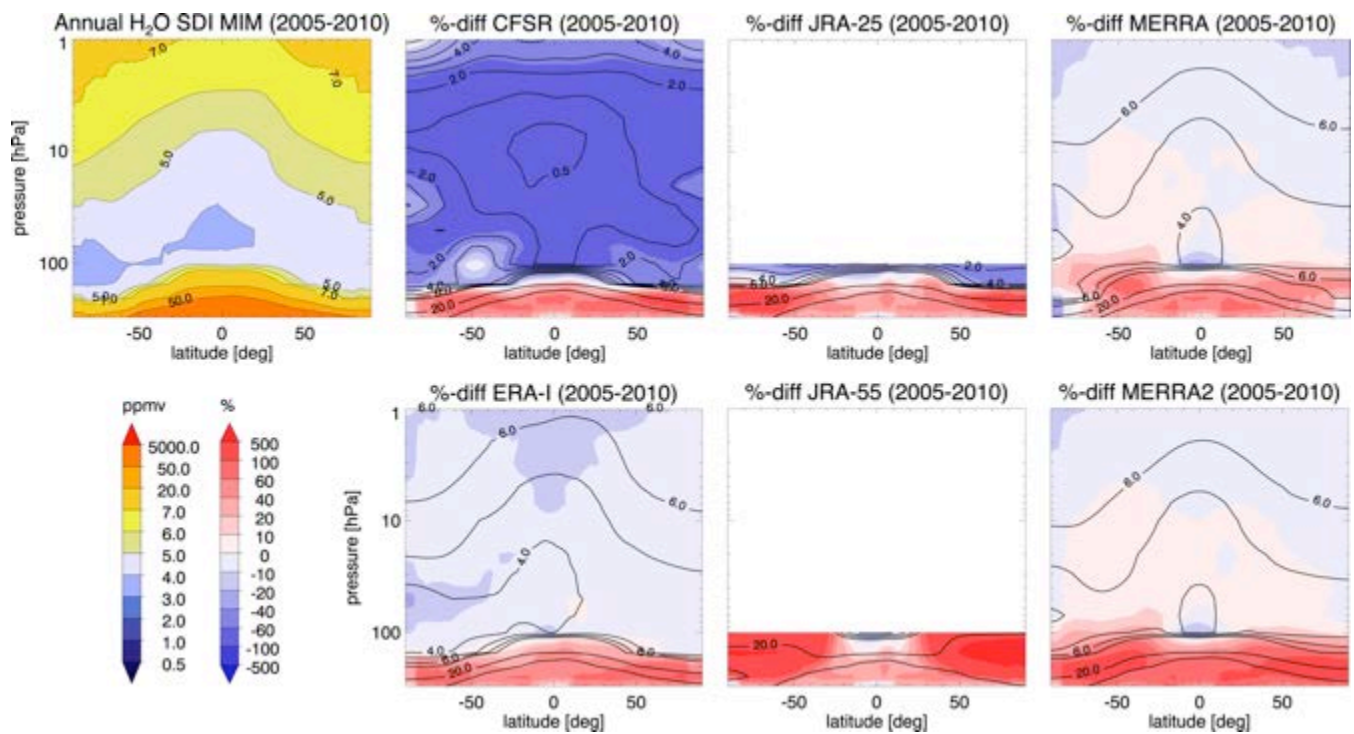
**Figure 9:** (a) Ozone hole mean area calculated from TOMS/OMI observations and the reanalyses for 21 September through 20 October of 1981–2010. (b) Differences between ozone hole mean areas from reanalyses and TOMS/OMI observations ( $R_i - \text{observed}$ ).



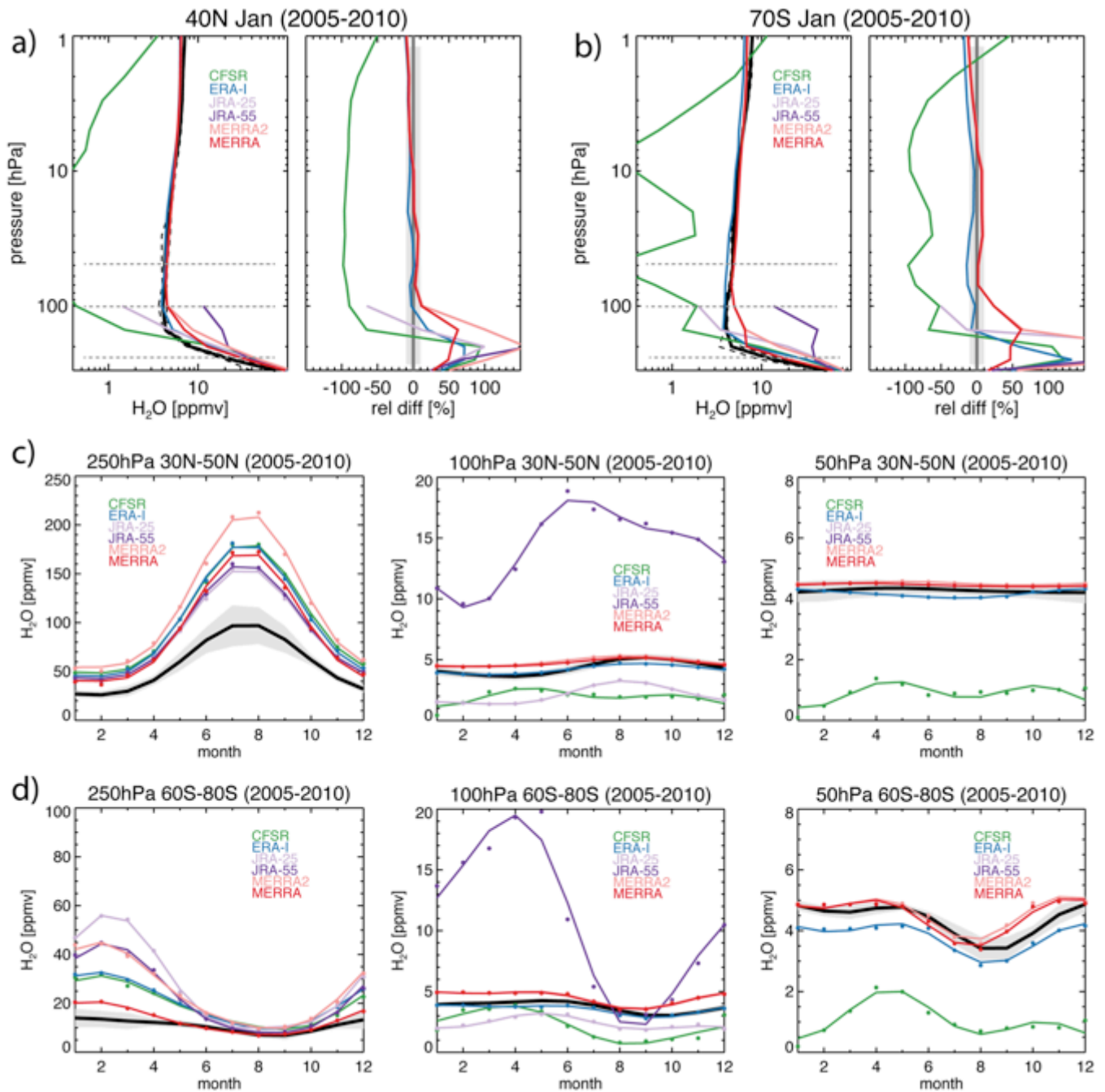
5 Figure 10: Departures of TCO from the zonal- and monthly-mean 1981-2010 climatology for TOMS/OMI (left column, top row), SBUV (left column, bottom row), and reanalyses (left column, other rows). (Right column) Differences between reanalyses zonal- and monthly-mean TCO and the primary TCO observations that they assimilate. The black contour is at 0 DU.



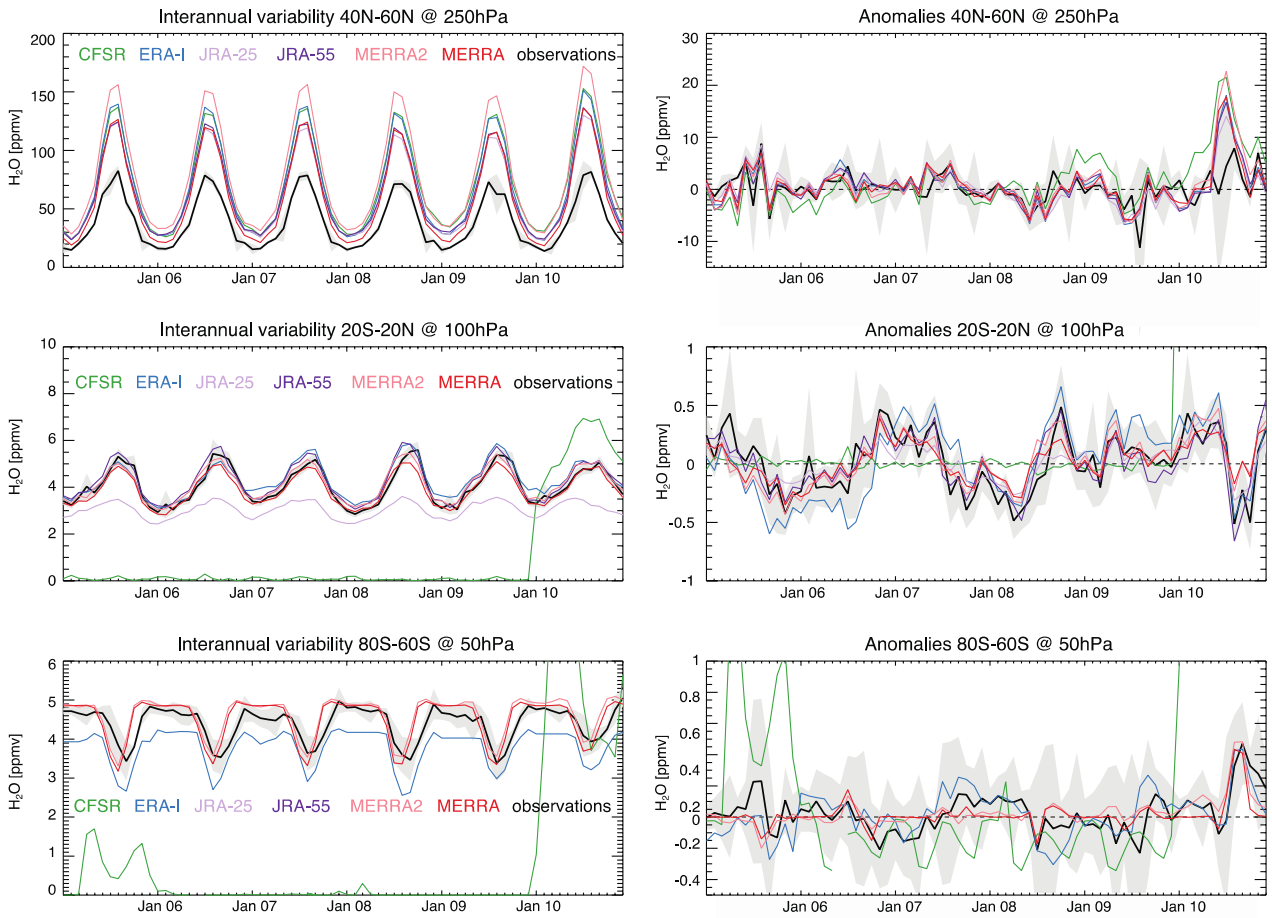
**Figure 11:** Latitude–time evolution of relative differences between ozone reanalyses and the merged SWOOSH ozone record at 10 hPa and 70 hPa. White indicates missing data, and light grey indicates near-zero differences (e.g., between MERRA2 and SWOOSH after mid-2004).



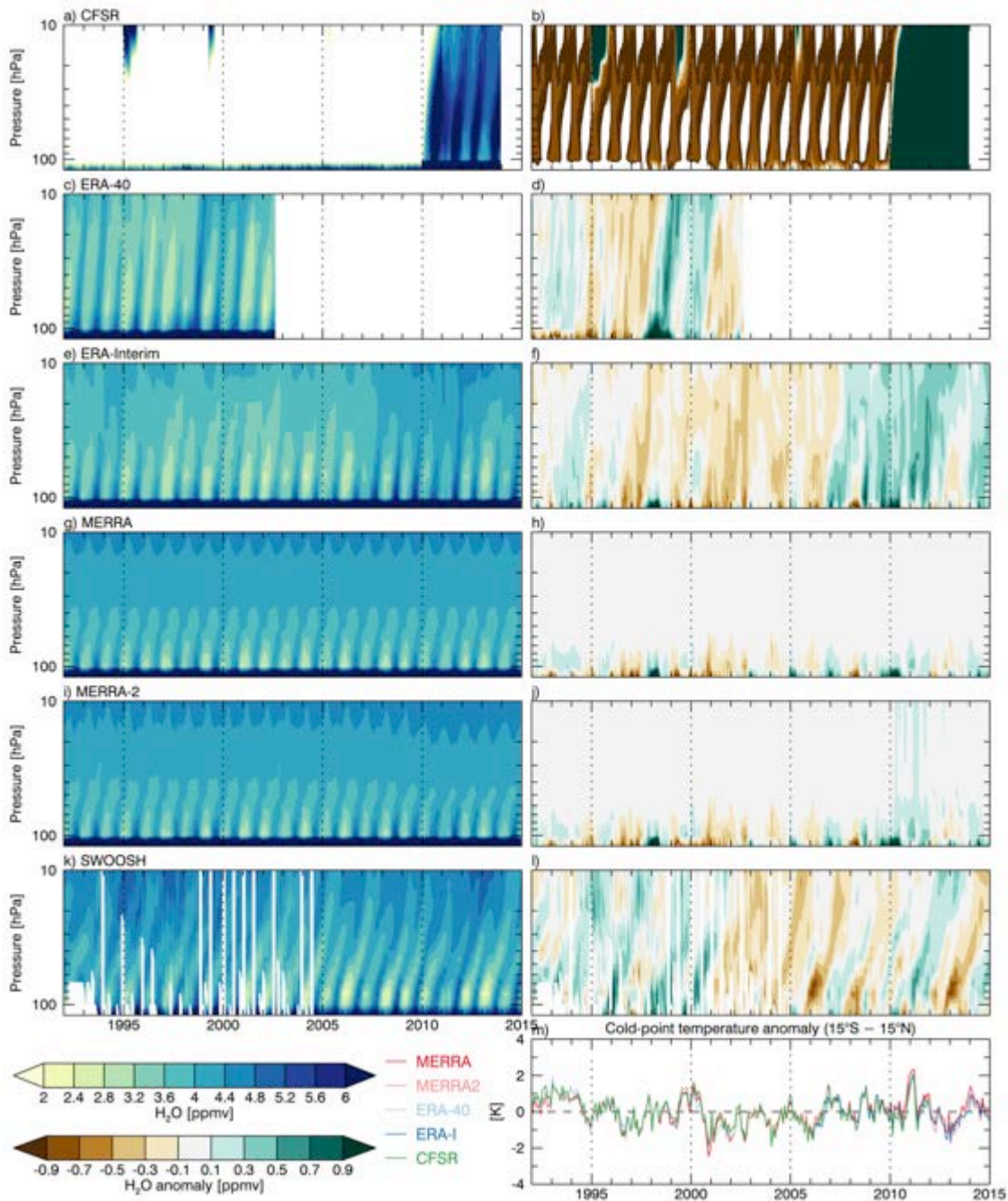
**Figure 12:** Multi-annual zonal mean water vapor cross sections averaged over 2005–2010 for the SPARC Data Initiative multi-instrument mean (SDI MIM) (upper left), along with the relative differences between reanalyses and observations as  $(R_i - \text{MIM})/\text{MIM} \times 100$ , where  $R_i$  is a reanalysis field. Also shown in contours are the respective zonal mean climatologies for the different reanalyses.



**Figure 13:** Multi-annual mean vertical water vapor profiles over 2005–2010 for January at (a) 40N and (b) 70S from the SPARC Data Initiative multi-instrument mean (SDI MIM) (black) and the six reanalyses (colored). Absolute values are shown in the left and relative differences in the right panels for each comparison. Relative differences are calculated as  $(R_i - \text{MIM}) / \text{MIM} * 100$ , where  $R_i$  is a reanalysis profile. Black dashed lines provide the  $\pm 1$ -sigma uncertainty (as calculated by the standard deviation over all instruments and years available) in the observational mean. Horizontal dashed lines in grey indicate the pressure levels (250, 100, and 50 hPa) for which seasonal cycles are shown in panels (c) and (d) for the two latitude ranges 30–50N and 60–80S, respectively.



5 **Figure 14:** Interannual variability (left column) and deseasonalized anomalies (right column) for water vapor during 2005–2010 for the SPARC Data Initiative multi-instrument mean (SDI MIM, black) and the six reanalyses (colored). Results are shown for three different pressure levels and latitude ranges (top to bottom: 50 hPa at 60–80°S, 100 hPa at 20°S–20°N, and 250 hPa at 40–60°N). Grey shading indicates observational uncertainty ( $\pm 1$ -sigma) calculated as the standard deviation over all instruments and years available.



**Figure 15:** The tropical tape recorder signal as represented in reanalyses and the SWOOSH merged satellite product, defined as the height–time evolution of water vapor averaged over the 15°S–15°N tropical band. Both absolute values (left column) and anomalies relative to the mean water vapor seasonal cycle at each level (right column) are shown. Anomalies are computed separately for each data set. Monthly mean anomalies in tropical (15°S–15°N) cold-point tropopause temperatures calculated from 6-h data on the native vertical resolution of each reanalysis model are shown for context (m).