

Modulation of Midtropospheric CO₂ by the South Atlantic Walker Circulation*

XUN JIANG

Department of Earth and Atmospheric Sciences, University of Houston, Houston, Texas

EDWARD T. OLSEN, THOMAS S. PAGANO, AND HUI SU

Science Division, Jet Propulsion Laboratory, California Institute of Technology, Pasadena, California

YUK L. YUNG

Division of Geological and Planetary Sciences, California Institute of Technology, Pasadena, California

(Manuscript received 19 November 2014, in final form 19 February 2015)

ABSTRACT

Midtropospheric CO₂ data from the Atmospheric Infrared Sounder (AIRS) are used in this study to explore the variability of CO₂ over the South Atlantic Ocean. It was found that the area-averaged CO₂ over the South Atlantic Ocean is less than that over South America by about 1 ppm during December–March. This CO₂ contrast is due to the large-scale vertical circulation over this region. During December–March, there is sinking motion over the South Atlantic Ocean. The sinking motion brings high-altitude air with a slightly lower concentration of CO₂ to the midtroposphere. Meanwhile, air rising over South America brings near-surface air with a higher concentration of CO₂ to the midtroposphere. As a result, the AIRS midtropospheric CO₂ concentration is lower over the South Atlantic Ocean than over South America during December–March. The detrended AIRS midtropospheric CO₂ difference correlates well with the inverted and detrended 400-hPa vertical pressure velocity difference between the South Atlantic and South America. Results obtained from this study demonstrate the strong impact of large-scale circulation on the vertical distribution of CO₂ in the free troposphere and suggest that midtropospheric CO₂ measurements can be used as an innovative observational constraint on the simulation of large-scale circulations in climate models.

1. Introduction

Recently retrieved CO₂ concentrations from satellites (Crisp et al. 2004; Miller et al. 2007; Chahine et al. 2008; Boesch et al. 2011; Crisp et al. 2012) offer a unique opportunity to explore CO₂ variations in the global domain. In addition to the trend and annual cycle (Keeling et al. 1995; Pearman and Hyson 1980; Cleveland et al. 1983; Bacastow et al. 1985; Keeling et al. 1996; Buermann et al. 2007; Jiang et al. 2008; Pagano et al. 2011, 2014), CO₂ concentrations also exhibit variability from intraseasonal

to interannual time scales (Bacastow 1976; Enting 1987; Feely et al. 1987; Keeling and Revelle 1985; Keeling et al. 1995; Dargaville et al. 2000; Dettinger and Ghil 1998; Keppel-Aleks et al. 2011; Jiang et al. 2012). Some recent studies further suggest that the large-scale circulations have significant influences on the temporal variability of midtropospheric CO₂ (Li et al. 2010; Jiang et al. 2010; Wang et al. 2011; Jiang et al. 2013a,b).

Using midtropospheric CO₂ retrieval from the Atmospheric Infrared Sounder (AIRS), it was found that the Madden–Julian oscillation (MJO) could influence the midtropospheric CO₂ concentrations in the tropical region (Li et al. 2010). The amplitude of the MJO signal in midtropospheric CO₂ is about 1 ppm (Li et al. 2010). In addition to the MJO signal, it was found that El Niño–Southern Oscillation (ENSO) could influence the midtropospheric CO₂ concentrations over the Pacific Ocean (Jiang et al. 2010, 2013a). There is more (less) midtropospheric CO₂ over the central Pacific Ocean and less

* Supplemental information related to this paper is available at the Journals Online website: <http://dx.doi.org/10.1175/JAS-D-14-0340.s1>.

Corresponding author address: Xun Jiang, Department of Earth and Atmospheric Sciences, University of Houston, 4800 Calhoun Rd., Houston, TX 77004.
E-mail: xjiang7@uh.edu

(more) midtropospheric CO₂ over the western Pacific Ocean during El Niño (La Niña). In the high latitudes, the midtropospheric CO₂ concentrations can be influenced by the large-scale dynamics—for example, the northern annular mode (Jiang et al. 2010) and stratospheric sudden warming (Jiang et al. 2013b). During strong (weak) polar vortex years, there is less (more) midtropospheric CO₂ in the polar region (Jiang et al. 2010). It was also found that the polar midtropospheric CO₂ concentrations could increase by 2–3 ppm within a few days after the stratospheric sudden warming events (Jiang et al. 2013b). In this paper, we investigate the influence of the South Atlantic Walker circulation (Wang 2002) on the AIRS midtropospheric CO₂.

2. Data

AIRS, version 5 (V5), midtropospheric CO₂ mixing ratios retrieved by the vanishing partial derivative (VPD) method (Chahine et al. 2005; Chahine et al. 2008) were used in this study. VPD CO₂ retrievals are obtained in a postprocessing stage ingesting the AIRS V5 level-2 products. The VPD algorithm applies the AIRS quality control (Suskind et al. 2011) to select retrievals and then minimizes the difference between level-2 cloud-cleared radiances and radiances computed for the level-2 atmospheric state. The solution is obtained when the partial derivatives of radiances in select channel sets with respect to temperature, water vapor, ozone, and CO₂ are individually equal to zero (Chahine et al. 2005; Chahine et al. 2008; Olsen and Licata 2015). The same constant value of CO₂ is used throughout the atmosphere across the globe as the initial guess in the original AIRS level-2 CO₂ retrieval. No additional assumptions for the distribution of atmospheric CO₂ are made (i.e., seasonal variations or vertical profiles from a model). The contribution function of channels employed to retrieve AIRS midtropospheric CO₂, which indicates the relative fraction of the radiance observed at the top of the atmosphere arising from the various layers in the atmosphere, is maximum between 500 and 300 hPa (Chahine et al. 2008; Olsen and Licata 2015). AIRS V5 level-3 midtropospheric CO₂ retrieval products gridded at 2° × 2.5° (latitude by longitude) from 60°S to 90°N are available from September 2002 to February 2011. Monthly mean AIRS level-3 midtropospheric CO₂ are available at <http://disc.sci.gsfc.nasa.gov/AIRS/data-holdings/by-data-product-v5/AIRX3C2M> (Olsen and Licata 2015). Validation by comparison to in situ aircraft measurements and retrievals by land-based upward-looking Fourier transform interferometers demonstrated that AIRS CO₂ is accurate to about 2 ppm between latitudes 40°S and 40°N (Chahine et al. 2005, 2008; Olsen and Licata 2015). The midtropospheric CO₂ retrieved via

the VPD method captures the correct seasonal cycle and trend compared with those from Comprehensive Observation Network for Trace Gases by Airliner (CONTRAIL) (Chahine et al. 2005).

Vertical pressure velocity data were used in this study to represent the South Atlantic Walker circulation. Vertical pressure velocity data are taken from National Centers for Environmental Prediction reanalysis 2 (NCEP2) data (Kistler et al. 2001), European Centre for Medium-Range Weather Forecasts interim (ECMWF-Interim) reanalysis data (Dee et al. 2011), Modern-Era Retrospective Analysis for Research and Applications (MERRA) reanalysis data (Rienecker et al. 2011), and atmosphere–ocean Coupled Model Intercomparison Project phase 5 (CMIP5) (Taylor et al. 2012).

3. Results

To investigate the influence of the large-scale circulation on the midtropospheric CO₂, we examined the AIRS midtropospheric CO₂ distributions from December to March (DJFM) averaged over 2003–10 in Fig. 1. December–March was chosen because this is when the CO₂ anomaly over the South Atlantic Ocean is greatest. NCEP2 400-hPa vertical pressure velocity (dP/dt) calculated for the same time period was overlain on the AIRS midtropospheric CO₂ in Fig. 1. There is more midtropospheric CO₂ over the western Pacific Ocean and less midtropospheric CO₂ over the eastern Pacific Ocean, which is related to the Walker circulation in this region as suggested by Jiang et al. (2010). Furthermore, midtropospheric CO₂ concentrations are the lowest over the South Atlantic Ocean, coincident with sinking air as shown by the white solid contours in Fig. 1. Sinking air can bring lower concentrations of CO₂ from high altitudes to the midtroposphere, resulting in AIRS measuring lower concentrations of midtropospheric CO₂ over the South Atlantic Ocean. The rising air over South America brings high concentrations of CO₂ from the surface to the midtroposphere, resulting in AIRS measuring higher concentrations of midtropospheric CO₂ over South America. As shown in Fig. 1, AIRS midtropospheric CO₂ concentrations over the South Atlantic Ocean are about 1 ppm lower than that over South America from December to March. There are 3000–4000 AIRS retrievals over this region. Chahine et al. (2005) demonstrated that the distribution of individual retrieval errors exhibits Gaussian properties. Thus the error for the area-averaged CO₂ is equal to CO₂ standard error (~2 ppm) divided by the square root of number of data. As a result, the error for the area-averaged mean CO₂ is below 0.1 ppm, which is smaller than the 1-ppm difference between the South Atlantic Ocean and South

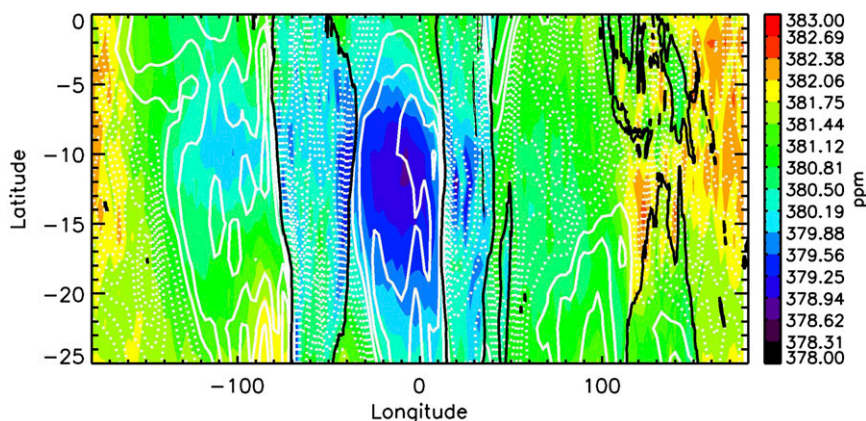


FIG. 1. AIRS midtropospheric CO_2 (ppm) averaged from December to March during 2003–10. Color represents AIRS midtropospheric CO_2 . White contours are the NCEP2 400-hPa dP/dt . Solid white contours refer to the sinking air. Dashed white contours refer to the rising air.

America. Our analysis also finds the NCEP2 400-hPa vertical pressure velocity distributions are well correlated with divergence fields at 850 and 200 hPa (see Fig. S1 in the supplemental material), with low-level divergence (convergence) and high-level convergence (divergence) for the South Atlantic Ocean (South America).

In addition to exploring the spatial distributions of vertical pressure velocity and midtropospheric CO_2 , we investigated the vertical cross section of the vertical pressure velocity across 70°W – 10°E averaged over 20° – 5°S from December to March during 2003–10. As shown in Fig. 2a, it is clear that air sinks over 35°W – 10°E and rises over 70° – 35°W . Consistent with expectations based on vertical motions, AIRS midtropospheric CO_2 concentrations are relatively low over 35°W – 10°E and relatively high over 70° – 35°W (Fig. 2b). The difference of midtropospheric CO_2 between the South Atlantic Ocean and South America areas is about 1 ppm from December to March.

Next, we investigated the temporal correlation between the South Atlantic Walker circulation and the midtropospheric CO_2 difference between the South Atlantic Ocean (20° – 5°S , 30°W – 10°E) and South America (20° – 5°S , 70° – 40°W). The detrended AIRS midtropospheric CO_2 difference between the South Atlantic Ocean and South America is shown by the black solid line in Fig. 3a. The linear trend was calculated using the first-order polynomial by the least squares method (Bevington and Robinson 2003) and was removed from the AIRS midtropospheric CO_2 difference. To indicate the South Atlantic Walker circulation, we also calculated the detrended 400-hPa vertical pressure velocity differences between the South Atlantic Ocean and South America from NCEP2 data (Kistler et al. 2001), ECMWF-Interim

reanalysis data (Dee et al. 2011), MERRA reanalysis data (Rienecker et al. 2011), and AMIP-type CMIP5 model simulations (Taylor et al. 2012), in which observed sea surface temperature were used to drive the models. A positive (negative) value of the vertical pressure velocity difference indicates that there is more (less) sinking motion over the South Atlantic Ocean than over South America. Detrended and inverted 400-hPa vertical pressure velocity differences from reanalysis data and different models are shown in Fig. 3a by the various color dashed lines. CO_2 and vertical pressure velocity differences from 2003 to 2008 are shown, since most model simulations ended in 2008.

As shown in Fig. 3a, the CO_2 difference (South Atlantic Ocean minus South America) is related to the vertical pressure velocity difference and there is a reoccurring seasonal variation. The CO_2 difference is negative (positive) when the vertical pressure velocity difference is positive (negative). It suggests that the concentration of midtropospheric CO_2 over the South Atlantic Ocean compared to that over South America is impacted by the vertical transport due to the South Atlantic Walker circulation. The relative concentration of midtropospheric CO_2 over the South Atlantic is diminished by sinking air and enhanced by rising air. Air converges at the surface and diverges in the upper levels over South America in DJFM (see Fig. S1 in the supplemental material), leading to rising air over South America, which in turn can bring high concentrations of CO_2 from the surface to the midtroposphere. Meanwhile, air diverges at the surface and converges in the upper levels over South Atlantic Ocean in DJFM, leading to sinking air over the South Atlantic Ocean, which in turn can bring low concentrations CO_2 from high altitudes to the midtroposphere. CO_2 difference

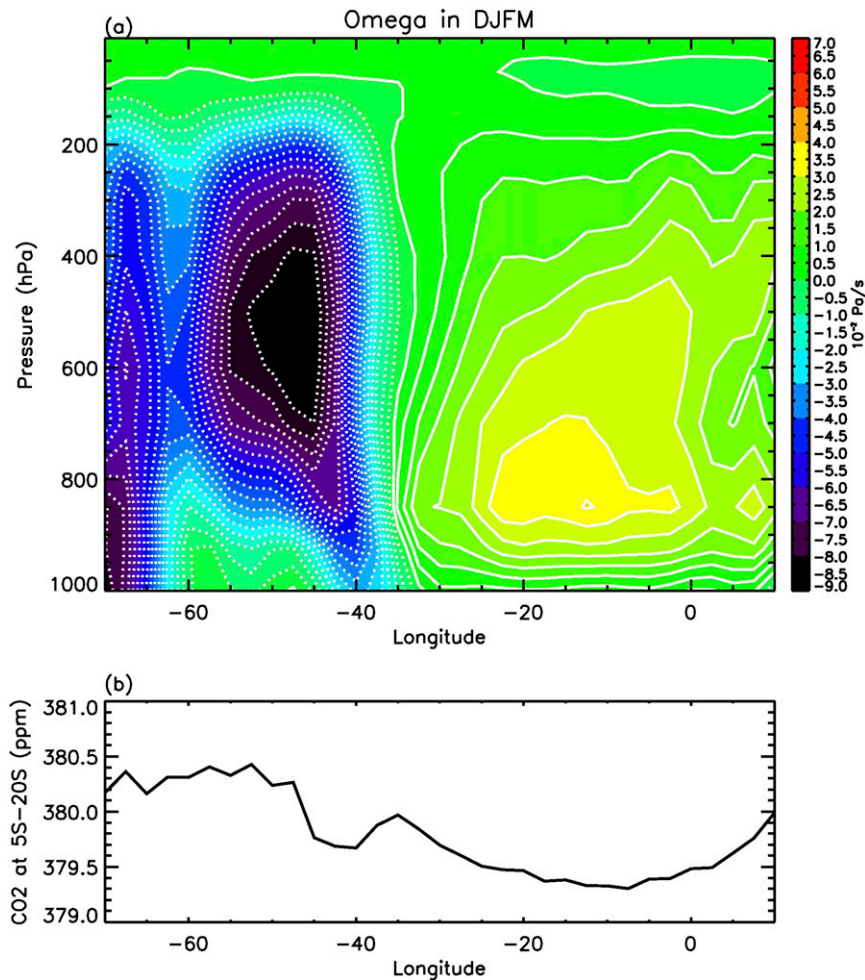


FIG. 2. (a) Vertical pressure velocity ($10^{-2} \text{ Pa s}^{-1}$) averaged over 20° – 5° S from December to March during 2003–10. Solid white contours refer to the sinking air. Dashed white contours refer to the rising air. (b) AIRS CO_2 (ppm) averaged over 20° – 5° S from December to March during 2003–10.

amplitudes are smaller in 2007 and 2008 than in earlier years, which might be related to CO_2 surface-emission contrasts between South America and the South Atlantic Ocean. The interannual variability of the CO_2 difference amplitudes will be explored in the future when CO_2 surface emission data are available.

As shown in Fig. 3a, the detrended AIRS CO_2 difference correlates well with the inverted and detrended vertical pressure velocity differences. The correlation coefficient between the detrended AIRS CO_2 difference (black solid line) and the inverted and detrended mean vertical pressure velocity difference (red dashed line) is 0.66. The corresponding significance level is 2%. The significance statistics for the correlation were generated by a Monte Carlo method (Press et al. 1992; Jiang et al. 2004). The correlation coefficients between the detrended AIRS CO_2 difference and the inverted and detrended vertical pressure velocity differences derived from reanalysis

datasets and CMIP5 model simulations range between 0.55 and 0.72 (Fig. 3b). The correlation coefficients are 0.67 for NCEP2, 0.64 for ERA-Interim, 0.55 for MERRA, and between 0.57 and 0.72 for CMIP5 simulations. Given the importance of large-scale circulation in driving global energy and water cycles, improving model simulations of large-scale circulation is critical to reducing the model spread in climate sensitivity estimates (Su et al. 2014). Since there are limited direct observations of vertical velocity, the midtropospheric CO_2 can be utilized as an indirect constraint on model representation of large-scale circulation—for example, the vertical velocity of the South Atlantic Walker cell.

4. Conclusions

AIRS midtropospheric CO_2 data were used in this study to investigate the impact of large-scale circulation

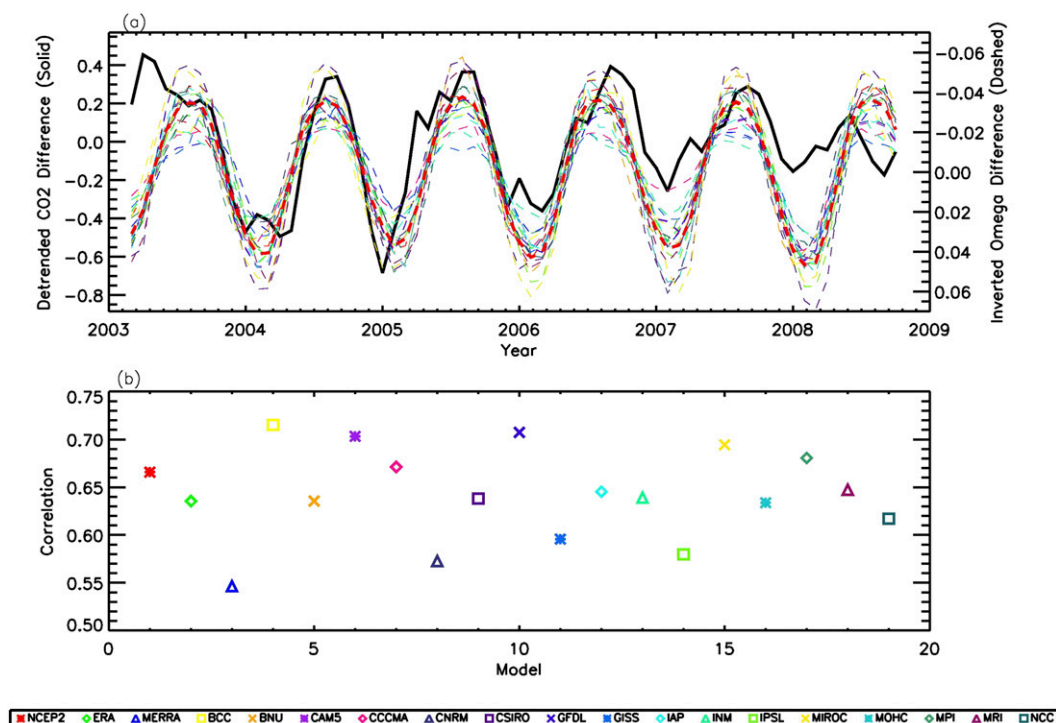


FIG. 3. (a) Difference of the detrended AIRS midtropospheric CO₂ between the South Atlantic Ocean (20°–5°S, 30°W–10°E) and South America (20°–5°S, 70°–40°W) (black solid line) and difference of the inverted and detrended 400-hPa dP/dt between the South Atlantic Ocean and South America from reanalysis data and various CMIP5 models (thin dashed lines). Thick red dashed line is the averaged dP/dt difference from all reanalysis data and model simulations. (b) Correlation coefficients between detrended CO₂ difference and detrended and inverted dP/dt differences from reanalysis data and CMIP5 models. A 3-month running mean has been applied to all time series to remove the high-frequency signals.

on midtropospheric CO₂ concentrations over the South Atlantic Ocean for the first time. It is found that concentrations of midtropospheric CO₂ over the South Atlantic Ocean are lower than over the surrounding region during December–March. As illustrated in Fig. 4, the sinking air over the South Atlantic Ocean brings lower-CO₂-concentration air from higher altitudes to the midtroposphere. Meanwhile, the rising air over South America brings near-surface air containing higher concentrations of CO₂ to the midtroposphere during December–March. As a response to the vertical motions, the AIRS retrieved midtropospheric CO₂ concentrations are about 1 ppm lower over the South Atlantic Ocean than over South America. We also find that the detrended AIRS midtropospheric CO₂ difference (South Atlantic Ocean minus South America) correlates well with the South Atlantic Walker circulation indicated by the inverted and detrended vertical pressure velocity differences from three reanalysis data and CMIP5 simulations. The vertical pressure velocity associated with the South Atlantic Walker circulation can lead to a difference as great as 1 ppm in the

midtropospheric CO₂. While all reanalysis datasets and CMIP5 simulations show a statistically significant correlation between the midtropospheric CO₂ difference and the South Atlantic Walker cell, the correlation coefficients range between 0.55 and 0.72. We conclude that

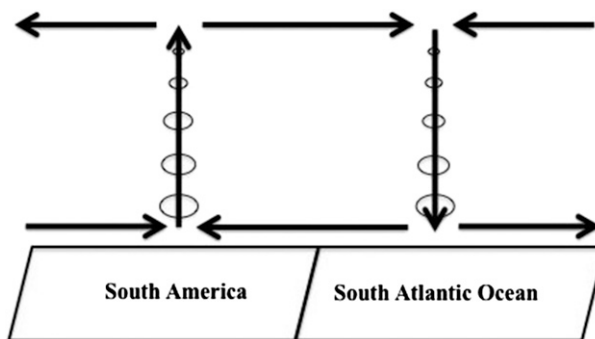


FIG. 4. Diagram to illustrate the influence of South Atlantic Walker circulation on the midtropospheric CO₂ concentrations. Air movements are highlighted by the arrows. Ovals represent the CO₂ concentrations, which are high near the surface and low in the high altitudes.

the clear modulation of large-scale circulation on the midtropospheric CO₂ enables an innovative observational constraint on the model representation of large-scale circulation (i.e., through the use of AIRS midtropospheric CO₂ measurements). Chemical transport models and CO₂ flux inversion projects are highly dependent on the accurate simulations of large-scale circulation to identify the impact of surface emissions on tracer distributions and evolutions. CO₂ measurements from AIRS and the newly available *Orbiting Carbon Observatory 2* (OCO-2) would likely enable a leap of progress in climate model improvements.

Acknowledgments. We thank two anonymous reviewers for helpful comments. XJ and YLY were supported by the AIRS project, OCO-2 project, and NASA Grant NNX13AC04G to UH and Caltech. Part of this research was carried out at the Jet Propulsion Laboratory, California Institute of Technology, under a contract with the National Aeronautics and Space Administration.

REFERENCES

- Bacastow, R. B., 1976: Modulation of atmospheric carbon dioxide by the Southern Oscillation. *Nature*, **261**, 116–118, doi:[10.1038/261116a0](https://doi.org/10.1038/261116a0).
- , C. D. Keeling, and T. P. Whorf, 1985: Seasonal amplitude increase in atmospheric CO₂ concentration at Mauna Loa, Hawaii, 1959–1982. *J. Geophys. Res.*, **90**, 10 529–10 540, doi:[10.1029/JD090iD06p10529](https://doi.org/10.1029/JD090iD06p10529).
- Bevington, P. R., and D. K. Robinson, 2003: *Data Reduction and Error Analysis for the Physical Science*. 3rd ed. McGraw-Hill, 336 pp.
- Boesch, H., D. Baker, B. Connor, D. Crisp, and C. Miller, 2011: Global characterization of CO₂ column retrievals from shortwave-infrared satellite observations of the Orbiting Carbon Observatory-2 Mission. *Remote Sens.*, **3**, 270–304, doi:[10.3390/rs3020270](https://doi.org/10.3390/rs3020270).
- Buermann, W., B. Lintner, C. Koven, A. Angert, J. E. Pinzon, C. J. Tucker, and I. Fung, 2007: The changing carbon cycle at the Mauna Loa Observatory. *Proc. Natl. Acad. Sci. USA*, **104**, 4249–4254, doi:[10.1073/pnas.0611224104](https://doi.org/10.1073/pnas.0611224104).
- Chahine, M., C. Barnett, E. T. Olsen, L. Chen, and E. Maddy, 2005: On the determination of atmospheric minor gases by the method of vanishing partial derivatives with application to CO₂. *Geophys. Res. Lett.*, **32**, L22803, doi:[10.1029/2005GL024165](https://doi.org/10.1029/2005GL024165).
- , and Coauthors, 2008: Satellite remote sounding of mid-tropospheric CO₂. *Geophys. Res. Lett.*, **35**, L17807, doi:[10.1029/2008GL035022](https://doi.org/10.1029/2008GL035022).
- Cleveland, M. S., A. E. Freeny, and T. E. Graedel, 1983: The seasonal component of atmospheric CO₂ Information from new approaches to the decomposition of seasonal time series. *J. Geophys. Res.*, **88**, 10 934–10 946, doi:[10.1029/JC088iC15p10934](https://doi.org/10.1029/JC088iC15p10934).
- Crisp, D., and Coauthors, 2004: The Orbiting Carbon Observatory (OCO) mission. *Adv. Space Res.*, **34**, 700–709, doi:[10.1016/j.asr.2003.08.062](https://doi.org/10.1016/j.asr.2003.08.062).
- , and Coauthors, 2012: The ACOS CO₂ retrieval algorithm—Part II: Global X_{CO2} data characterization. *Atmos. Meas. Tech.*, **5**, 687–707, doi:[10.5194/amt-5-687-2012](https://doi.org/10.5194/amt-5-687-2012).
- Dargaville, R. J., R. M. Law, and F. Pribac, 2000: Implications of interannual variability in atmospheric circulation on modeled CO₂ concentrations and source estimates. *Global Biogeochem. Cycles*, **14**, 931–943, doi:[10.1029/1999GB001166](https://doi.org/10.1029/1999GB001166).
- Dee, D. P., and Coauthors, 2011: The ERA-Interim reanalysis: Configuration and performance of the data assimilation system. *Quart. J. Roy. Meteor. Soc.*, **137**, 553–597, doi:[10.1002/qj.828](https://doi.org/10.1002/qj.828).
- Dettinger, M. D., and M. Ghil, 1998: Seasonal and interannual variations of atmospheric CO₂ and climate. *Tellus*, **50B**, 1–24, doi:[10.1034/j.1600-0889.1998.00001.x](https://doi.org/10.1034/j.1600-0889.1998.00001.x).
- Enting, I. G., 1987: The interannual variation in the seasonal cycle of carbon dioxide concentration at Mauna Loa. *J. Geophys. Res.*, **92**, 5497–5504, doi:[10.1029/JD092iD05p05497](https://doi.org/10.1029/JD092iD05p05497).
- Feely, R. A., R. H. Gammon, B. A. Taft, P. E. Pullen, L. S. Waterman, T. J. Conway, J. F. Gendron, and D. P. Wisegarver, 1987: Distribution of chemical tracers in the eastern equatorial Pacific during and after the 1982/1983 ENSO event. *J. Geophys. Res.*, **92**, 6545–6558, doi:[10.1029/JC092iC06p06545](https://doi.org/10.1029/JC092iC06p06545).
- Jiang, X., C. D. Camp, R. Shia, D. Noone, C. Walker, and Y. Yung, 2004: Quasi-biennial oscillation and quasi-biennial oscillation-beat in the tropical total column ozone: A two-dimensional model simulation. *J. Geophys. Res.*, **109**, D16305, doi:[10.1029/2003JD004377](https://doi.org/10.1029/2003JD004377).
- , Q. Li, M. Liang, R. Shia, M. T. Chahine, E. T. Olsen, L. Chen, and Y. L. Yung, 2008: Simulation of upper tropospheric CO₂ from chemistry and transport models. *Global Biogeochem. Cycles*, **22**, GB4025, doi:[10.1029/2007GB003049](https://doi.org/10.1029/2007GB003049).
- , M. T. Chahine, E. T. Olsen, L. L. Chen, and Y. L. Yung, 2010: Interannual variability of mid-tropospheric CO₂ from Atmospheric Infrared Sounder. *Geophys. Res. Lett.*, **37**, L13801, doi:[10.1029/2010GL042823](https://doi.org/10.1029/2010GL042823).
- , —, Q. Li, M. Liang, E. T. Olsen, L. Chen, J. Wang, and Y. L. Yung, 2012: CO₂ semiannual oscillation in the middle troposphere and at the surface. *Global Biogeochem. Cycles*, **26**, doi:[10.1029/2011GB004118](https://doi.org/10.1029/2011GB004118).
- , J. Wang, E. T. Olsen, M. Liang, T. S. Pagano, L. Chen, S. J. Licata, and Y. L. Yung, 2013a: Influence of El Niño on mid-tropospheric CO₂ from Atmospheric Infrared Sounder and Model. *J. Atmos. Sci.*, **70**, 223–230, doi:[10.1175/JAS-D-11-0282.1](https://doi.org/10.1175/JAS-D-11-0282.1).
- , —, —, T. S. Pagano, L. Chen, and Y. L. Yung, 2013b: Influence of stratospheric sudden warming on AIRS mid-tropospheric CO₂. *J. Atmos. Sci.*, **70**, 2566–2573, doi:[10.1175/JAS-D-13-064.1](https://doi.org/10.1175/JAS-D-13-064.1).
- Keeling, C. D., and R. Revelle, 1985: Effects of El Niño/Southern Oscillation on the atmospheric content of carbon dioxide. *Meteoritics*, **20**, 437–450.
- , T. P. Whorf, M. Wahlen, and J. Vanderpligt, 1995: Interannual extremes in the rate of rise of atmospheric carbon dioxide since 1980. *Nature*, **375**, 666–670, doi:[10.1038/375666a0](https://doi.org/10.1038/375666a0).
- , J. F. S. Chin, and T. P. Whorf, 1996: Increased activity of northern vegetation inferred from atmospheric CO₂ measurements. *Nature*, **382**, 146–149, doi:[10.1038/382146a0](https://doi.org/10.1038/382146a0).
- Keppel-Aleks, G., P. O. Wennberg, and T. Schneider, 2011: Sources of variations in total column carbon dioxide. *Atmos. Chem. Phys.*, **11**, 3581–3593, doi:[10.5194/acp-11-3581-2011](https://doi.org/10.5194/acp-11-3581-2011).
- Kistler, R., and Coauthors, 2001: The NCEP–NCAR 50-Year Reanalysis: Monthly means CD-ROM and documentation. *Bull. Amer. Meteor. Soc.*, **82**, 247–267, doi:[10.1175/1520-0477\(2001\)082<0247:TNNYRM>2.3.CO;2](https://doi.org/10.1175/1520-0477(2001)082<0247:TNNYRM>2.3.CO;2).
- Li, K. F., B. Tian, D. E. Waliser, and Y. L. Yung, 2010: Tropical mid-tropospheric CO₂ variability driven by the Madden-Julian oscillation. *Proc. Natl. Acad. Sci. USA*, **107**, 19 171–19 175, doi:[10.1073/pnas.1008222107](https://doi.org/10.1073/pnas.1008222107).

- Miller, C. E., and Coauthors, 2007: Precision requirements for space-based X_{CO_2} data. *J. Geophys. Res.*, **112**, D10314, doi:[10.1029/2006JD007659](https://doi.org/10.1029/2006JD007659).
- Olsen, E. T., and S. J. Licata, 2015: AIRS Version 5 release tropospheric CO_2 products. Jet Propulsion Laboratory, 38 pp. [Available online at http://disc.sci.gsfc.nasa.gov/AIRS/documentation/v5_docs/AIRS_V5_Release_User_Docs/AIRS-V5-Tropospheric-CO2-Products.pdf.]
- Pagano, T. S., E. T. Olsen, M. T. Chahine, A. Ruzmainkin, H. Nguyen, and X. Jiang, 2011: Monthly representations of mid-tropospheric carbon dioxide from the Atmospheric Infrared Sounder. *Imaging Spectrometry XVI*, S. Shaklan, Ed., International Society for Optical Engineering (SPIE Proceedings, Vol. 8158), 81580C, doi:[10.1117/12.894960](https://doi.org/10.1117/12.894960).
- , —, H. Nguyen, A. Ruzmaikin, X. Jiang, and L. Perkins, 2014: Global variability of midtropospheric carbon dioxide as measured by the Atmospheric Infrared Sounder. *J. Appl. Remote Sens.*, **8**, 084984, doi:[10.1117/1.JRS.8.084984](https://doi.org/10.1117/1.JRS.8.084984).
- Pearman, G. I., and P. Hyson, 1980: Activities of the global biosphere as reflected in atmospheric CO_2 records. *J. Geophys. Res.*, **85**, 4457–4467, doi:[10.1029/JC085iC08p04457](https://doi.org/10.1029/JC085iC08p04457).
- Press, W., S. Teukolsky, W. Vetterling, and B. Flannery, 1992: *Numerical Recipes in Fortran 77: The Art of Scientific Computing*. 2nd ed. Cambridge University Press, 933 pp.
- Rienecker, M. M., and Coauthors, 2011: MERRA: NASA's Modern-Era Retrospective Analysis for Research and Applications. *J. Climate*, **24**, 3624–3648, doi:[10.1175/JCLI-D-11-00015.1](https://doi.org/10.1175/JCLI-D-11-00015.1).
- Su, H., J. H. Jiang, C. Zhai, T. J. Shen, J. D. Neelin, G. L. Stephens, and Y. L. Yung, 2014: Weakening and strengthening structures in the Hadley circulation change under global warming and implications for cloud response and climate sensitivity. *J. Geophys. Res. Atmos.*, **119**, 5787–5805, doi:[10.1002/2014JD021642](https://doi.org/10.1002/2014JD021642).
- Susskind, J., J. Blaisdell, L. Iredell, and F. Keita, 2011: Improved temperature sounding and quality control methodology using AIRS/AMSU data: The AIRS Science Team Version 5 retrieval algorithm. *IEEE Trans. Geosci. Remote Sens.*, **49**, 883–907, doi:[10.1109/TGRS.2010.2070508](https://doi.org/10.1109/TGRS.2010.2070508).
- Taylor, K. E., R. J. Stouffer, and G. A. Meehl, 2012: An overview of CMIP5 and the experiment design. *Bull. Amer. Meteor. Soc.*, **93**, 485–498, doi:[10.1175/BAMS-D-11-00094.1](https://doi.org/10.1175/BAMS-D-11-00094.1).
- Wang, C., 2002: Atlantic climate variability and its associated atmospheric circulation cells. *J. Climate*, **15**, 1516–1536, doi:[10.1175/1520-0442\(2002\)015<1516:ACVAIA>2.0.CO;2](https://doi.org/10.1175/1520-0442(2002)015<1516:ACVAIA>2.0.CO;2).
- Wang, J., and Coauthors, 2011: The influence of tropospheric biennial oscillation in mid-tropospheric CO_2 . *Geophys. Res. Lett.*, **38**, L20805, doi:[10.1029/2011GL049288](https://doi.org/10.1029/2011GL049288).