

Bulk properties of isentropic mixing into the tropics in the lower stratosphere

K. Minschwaner,¹ A. E. Dessler,² J. W. Elkins,³ C. M. Volk,^{3,4} D. W. Fahey,⁵ M. Loewenstein,⁶ J. R. Podolske,⁶ A. E. Roche,⁷ and K. R. Chan⁶

Abstract. Timescales for mixing of midlatitude air into the tropical lower stratosphere are deduced from observations of long-lived tracers N₂O and CCl₃F. Bulk mixing between tropical and midlatitude regions is assumed to be isentropic and relatively slow compared with local mixing within each region. The mean value of the mixing timescale ranges from 12 to 18 months near 20 km. There is a tendency for shorter mixing times at higher and lower altitudes, although vertical profiles of mixing cannot be definitively established by the data. A more robust quantity is given by the fraction of midlatitude air entrained into the tropical upwelling region. Implied mixing fractions exceed 50% above 22 km.

Introduction

It has long been recognized that the observed distributions of stratospheric water vapor and ozone imply upward transport across the tropical tropopause and a mean meridional circulation characterized by ascent in the tropics and subsidence at higher latitudes [Brewer, 1949; Dobson, 1956]. Early work by Newell [1963] on the dispersal of ¹⁸⁵W bomb debris and more recent studies of sulfate aerosol distributions [Trepte and Hitchman, 1992] show the persistence of maxima in tracer fields within 20° of the equator and between 21 and 28 km altitude. These observations indicate a barrier to mixing with the extratropics, at least in comparison to rapid irreversible mixing within midlatitude "surf zones" [McIntyre and Palmer, 1984]. Poleward transport across the subtropical barrier appears to be associated with planetary wave activity, such as the tongues of air stretching from the tropics into midlatitudes reported by Randel *et al.* [1993]. These features have been

identified also in transport model simulations [Waugh, 1993; Chen *et al.*, 1994]. At lower altitudes, measurements of N₂O and H₂O suggest that mixing out of the tropics is much more efficient, occurring on a timescale of 3 months or less below 22 km [Hintsa *et al.*, 1994].

The above studies emphasized transport from the tropics to midlatitudes, but an equally important issue is the mixing of midlatitude air into the tropics. The timescale associated with mixing into the tropics is a key factor in assessing the stratospheric effect of high-speed civil transport. Model calculations indicate that the sensitivity of ozone loss rates to perturbations in NO_x levels are much greater at 30 km than at 20 km [Considine *et al.*, 1995]. It is thus important to understand the extent to which aircraft exhaust emitted at midlatitudes may be mixed into the tropics and carried to higher altitudes where its effect on ozone is significant. Observations of long-lived gases and reactive compounds NO_y and O₃ indicate a small, but possibly important, amount of in-mixing from the extratropics [Avallo and Prather, 1996]. Here we examine the bulk properties of stratospheric mixing into the tropics through analysis of long-lived tracer fields obtained from balloon, aircraft, and satellite platforms. The observations are coupled to a diagnostic formulation for tropical tracer budgets which permits estimation of the mean rate of isentropic mixing from midlatitudes to the tropics in the lower stratosphere.

Observations

We focus on two long-lived species, N₂O (nitrous oxide) and CCl₃F (CFC-11). Sources for both gases are confined to the troposphere, and the sinks are due primarily to photodissociation in the stratosphere. Global, steady state lifetimes are calculated to be 123 and 44 years for N₂O and CCl₃F, respectively [Minschwaner *et al.*, 1993]. Local photochemical lifetimes at 22 km in the tropics are considerably smaller, of the order of 25 years

¹Department of Physics, New Mexico Institute of Mining and Technology, Socorro.

²NASA Goddard Space Flight Center, Greenbelt, Maryland.

³Climate Monitoring and Diagnostics Laboratory, NOAA Boulder, Colorado.

⁴Also at the Cooperative Institute for Research in Environmental Sciences, Boulder, Colorado.

⁵Aeronomy Laboratory, NOAA, Boulder, Colorado.

⁶NASA Ames Research Center, Moffett Field, California.

⁷Lockheed Palo Alto Research Laboratory, Palo Alto, California.

for N_2O and 1 year for CCl_3F in the annual mean. The wide range of photochemical time constants between these two species is useful for the problem at hand, as effects of isentropic mixing are more easily quantified if the mixing timescale is shorter than the local photochemical lifetime. Another consideration, however, is the mean difference between tropical and midlatitude mixing ratios along a given isentropic surface.

Figures 1 and 2 show balloon and aircraft measurements of tropical CCl_3F (Figure 1) and N_2O (Figure 2) spanning 5°S to 10°N by Goldan *et al.* [1980] and Vedder *et al.* [1978, 1981]. Midlatitude data shown in Figures 1 and 2 are from 41°N by Goldan *et al.* [1980]. The Goldan *et al.* N_2O data are adjusted according to the correction factor recommended by Goldan *et al.* [1981]. Uncertainties for these data range from 10% to 25%. All observations were made during the period 1976 to 1979 and are scaled to 1980 using a height-independent factor derived from observed trends in the troposphere [Elkins *et al.*, 1993]. The altitude coordinate of the original data is mapped to annual mean isentropes in Figures 1 and 2 using climatological pressure and temperature fields [Anderson *et al.*, 1986].

Curves drawn through the data in Figures 1 and 2 are least squares, quadratic exponential fits ($y = \exp(ax^2 + bx + c)$) which are used in the subsequent analysis. The limited amount of tropical data precludes the use of a nonparametric fit for this work. Choice of the quadratic exponential function is guided by the expected form of the tropical vertical profile in the absence of isentropic mixing, as discussed below. In addition, this is one of the lowest order functions that can reproduce the correct asymptotic limits (0 at ∞ , ap-

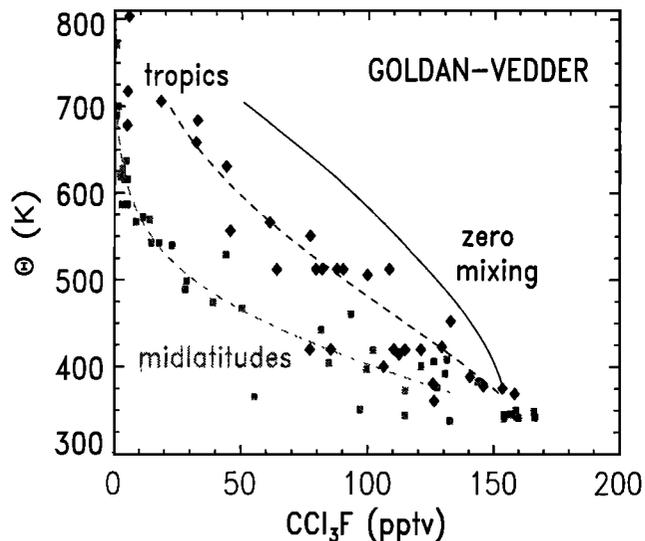


Figure 1. CCl_3F mixing ratios obtained from balloon and aircraft measurements by Goldan *et al.* [1980] and Vedder *et al.* [1978, 1981] mapped to climatological potential temperature as discussed in the text. Black diamonds are data obtained between 5°S and 10°N ; gray squares are for data from 41°N . Dashed curves are least squares fits to the data. The solid curve labeled “zero mixing” is calculated according to (2).

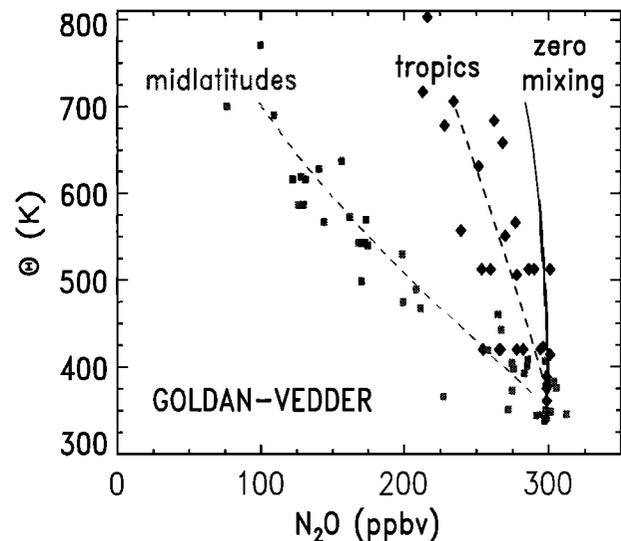


Figure 2. Same as Figure 1 but for mixing ratios of N_2O .

proximately constant mixing ratio below tropopause) and has been used previously by Ehhalt *et al.* [1983] to best describe their measurements of tracer vertical profiles at midlatitudes. All of our least-squares fits are constrained to match tropospheric values [Elkins *et al.*, 1993] at the tropopause.

The data shown in Figure 3 consists of CCl_3F measurements obtained by the National Oceanic and Atmospheric Administration (NOAA) Airborne Chromatograph for Atmospheric Trace Species (ACATS-4) onboard the NASA ER 2 aircraft [Elkins *et al.*, 1996]. The observations were made during the Airborne Southern Hemisphere Ozone Experiment Measurements for Assessing the Effects of Stratospheric Aircraft (ASHOE/MAESA) (March–November 1994), and the Stratospheric Photochemistry, Aerosol, and Dynamics Expedition (SPADE) using ACATS-2 (May 1993). Midlatitude data consist of measurements between 30° to 40° latitude. Discrimination against anomalously high or low data influenced by the short-term effects of planetary wave activity is accomplished using modified potential vorticity (MPV) [Lait, 1994] as a quasi-conserved tracer. The midlatitude data are constrained by values of MPV between 8×10^{-6} and $1.6 \times 10^{-5} \text{ K m}^2 \text{ kg}^{-1} \text{ s}^{-1}$. MPV is obtained from daily National Meteorological Center analyses and interpolated to the measurement location. Tropical CCl_3F data, defined here for $\pm 15^\circ$ latitude and $|\text{MPV}| < 6 \times 10^{-6} \text{ K m}^2 \text{ kg}^{-1} \text{ s}^{-1}$, were obtained only during ASHOE/MAESA. The uncertainty of each measurement is estimated to be $< 3\%$.

Figure 4 presents N_2O measurements obtained from the SPADE and ASHOE/MAESA ER 2 flights. During SPADE, N_2O was measured by the ATLAS tunable diode laser instrument [Podolske and Loewenstein, 1993]. The measurement uncertainty is 5% (1σ). Measurements of N_2O during ASHOE/MAESA are taken from the NOAA ACATS instrument. These data have a

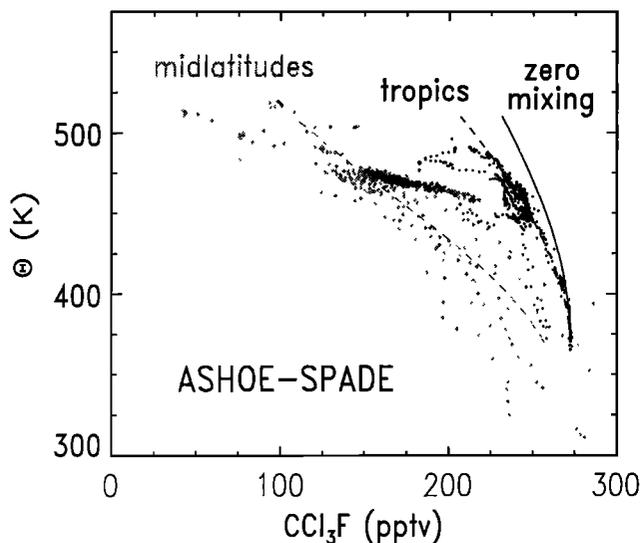


Figure 3. CCl_3F mixing ratios measured during ER 2 flights from the Airborne Southern Hemisphere Ozone Experiment - Measurements for Assessing the Effects of Stratospheric Aircraft (ASHOE/MAESA) and the Stratospheric Photochemistry, Aerosol, and Dynamics Expedition (SPADE) missions, as a function of potential temperature obtained by simultaneous measurements of pressure and temperature. Black points are from $\pm 15^\circ$ latitude with $|\text{MPV}| \leq 6 \times 10^{-6} \text{ K m}^2 \text{ kg}^{-1} \text{ s}^{-1}$; gray points are from $30^\circ\text{--}40^\circ\text{N}$ with $8 \times 10^{-8} < \text{MPV} < 1.6 \times 10^{-5} \text{ K m}^2 \text{ kg}^{-1} \text{ s}^{-1}$. As in Figure 1, dashed curves are fits to the data and the solid curve is the zero mixing solution given by (2).

measurement uncertainty of less than 3%. Nearly all of the ER 2 data lie below the 500 K isentropic level. At higher θ , the data shown in Figure 4 are from the Cryogenic Limb Array Etalon Spectrometer (CLAES) [Roche *et al.*, 1993] onboard the Upper Atmosphere Research Satellite, measured between February 12 and March 16, 1993. Typical uncertainties in these CLAES N_2O measurements are approximately 20%. Discrimination of all N_2O data into midlatitude and tropical bins is performed on the basis of latitude and MPV, as discussed above for CCl_3F .

Calculations

Inputs of CCl_3F and N_2O to the stratosphere are governed primarily by the cross-tropopause flux in the tropics. The local budget of these tracers in the tropical lower stratosphere is dominated by effects of upward transport, photochemical loss, and mixing with the extratropics. Focusing exclusively on the tropics, the continuity equation for the tropical mixing ratio χ of either gas is given by

$$\frac{\partial \chi}{\partial t} + \dot{\theta} \frac{\partial \chi}{\partial \theta} + L\chi + \frac{1}{\tau}(\chi - \chi') = 0 \quad (1)$$

where θ is potential temperature, $\dot{\theta}$ is the rate of radiative heating, L is the photochemical loss rate, and χ'

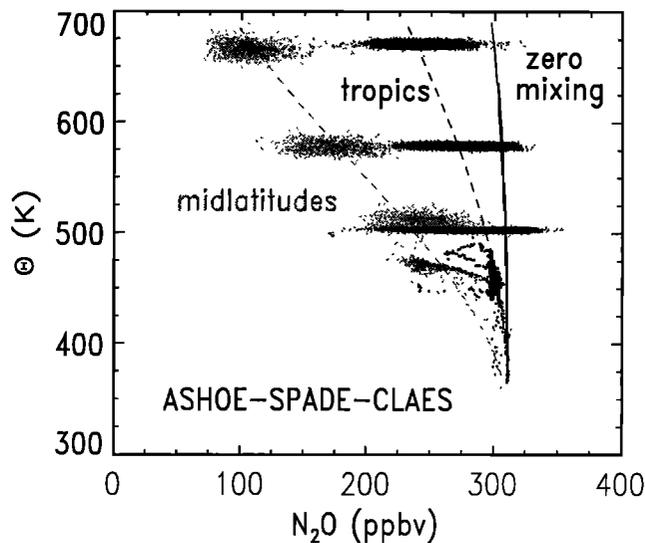


Figure 4. Same as Figure 3, except for N_2O from the ASHOE/MAESA and SPADE ER-2 missions. Data above about 500 K are from the CLAES satellite instrument on board the UARS satellite.

is the midlatitude mixing ratio. Quasi-horizontal mixing along surfaces of constant potential temperature is parameterized using a mixing timescale τ . This is the local timescale for mixing into the tropics on a given isentropic surface. Our approach assumes that isentropic mixing between tropics and midlatitudes is slow compared with mixing confined solely within each region. This point of view is supported by bomb and volcanic debris studies as well as observations of differences in tracer correlations between tropical and midlatitude regimes [Murphy *et al.*, 1993].

Using the analytic fits to the measured vertical profiles of χ , χ' , $\partial \chi / \partial \theta$, and calculated values of θ and L , we can apply (1) to determine τ as a function of θ . Photochemical loss (L) due to ultraviolet photodissociation is calculated using the high-resolution radiation code described by Minschwaner *et al.* [1993] and includes effects of scattering in the 190–210 nm spectral region. These loss rates give rise to calculated values of global and local lifetimes noted previously. The N_2O loss includes a small contribution (about 10%) due to reaction with $\text{O}(^1\text{D})$. Long-term atmospheric growth rates ($\partial \chi / \partial t$) are specified from analysis of tropospheric measurements (see Elkins *et al.* [1993] for discussion of methods). This term is small compared with either vertical advection or photochemical loss except in the case of CCl_3F data by Goldan *et al.* [1980] and Vedder *et al.* [1978, 1981] (hereinafter referred to as Goldan-Vedder), where 1980 growth rates were 5.7% per year.

Radiative heating ($\dot{\theta}$) is calculated using the Community climate Model 2 radiative transfer code [Briegleb, 1992a, b] initialized with tropical temperature, ozone, and water vapor profiles observed during the Central Equatorial Pacific Experiment [Dessler *et al.*, 1996]. Tropospheric clouds amounts are specified according to annual mean values from the International Satellite

Cloud Climatology Project [Rossow and Schiffer, 1991]. Longwave fluxes to space and shortwave albedo are in good agreement with measurements from the Earth Radiation Budget Experiment [Barkstrom, 1984]. Figure 5 shows the vertical profile of radiative heating. Implied ascent velocities in the tropical lower stratosphere (about $0.025 \text{ km day}^{-1}$) are consistent with the mean value determined by Holton [1990] on the basis of global mass balance.

Results

The zero mixing curves shown in Figures 1-4 are solutions to the homogeneous form of (1) using calculated values of θ and L . That is, we set $\tau = \infty$ and solve for χ under the assumption of no tropical-midlatitude mixing. The vertical profile thus takes the form of an exponential

$$\chi = \chi^{\circ} \exp \left(- \int_{\theta_o}^{\theta} \frac{L + \gamma}{\theta} d\theta \right) \quad (2)$$

where $\gamma = (1/\chi)(\partial\chi/\partial t)$ is the long-term growth rate, θ_o is the potential temperature of the tropopause, approximately 375 K, and χ° is the tropopause mixing ratio. The argument to the exponential function in (2) for both gases is well approximated by a quadratic polynomial, thus motivating the use of quadratic exponential fits to the observed vertical profiles.

The poor agreement shown in Figures 1-4 between the zero mixing solutions and tropical data is a clear indication of entrainment of midlatitude air into the tropical ascent region. The discrepancies are not related to errors in computed photochemical loss: uncertainties in photolysis frequencies are no larger than 20% [Minschwaner et al., 1993] which cannot explain the differ-

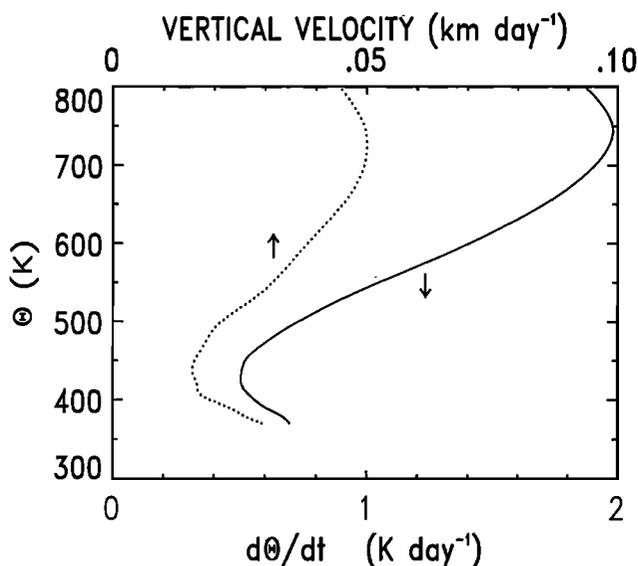


Figure 5. Calculated values of $\dot{\theta}$ used in (1) (solid curve, lower axis). Approximate vertical velocities are indicated by the dotted curve (upper axis).

ence between the zero mixing curve and the data. Possible errors in computed radiative heating also cannot be responsible, since it is easily shown that (2) gives rise to a power law relationship between CCl_3F and N_2O mixing ratios which is independent of the diabatic heating rate:

$$\chi_{\text{CCl}_3\text{F}} = \chi_{\text{CCl}_3\text{F}}^{\circ} \left(\frac{\chi_{\text{N}_2\text{O}}}{\chi_{\text{N}_2\text{O}}^{\circ}} \right)^K \quad (3)$$

where K (≈ 22 in the tropical lower stratosphere) is the ratio of CCl_3F to N_2O loss plus growth terms (the numerator of the integrand in (2) above). We found large differences between the power law relation and the tropical CCl_3F - N_2O correlations obtained from Figures 1 and 2, indicating that the zero mixing discrepancies cannot be due primarily to errors in computed radiative heating rates.

As mentioned above, timescales for isentropic mixing between the tropics and midlatitudes can be obtained from (1) using a combination of calculated quantities and mean vertical tracer profiles (analytic fits) based on the observations shown in Figures 1-4. Timescales derived in this fashion are shown in Figure 6. Owing to the higher measurement accuracies and availability of simultaneous observations of potential temperature, there is reason to place a higher weight on results from the ASHOE, SPADE, and CLAES (hereinafter referred to as A-S-C) data. These mixing time profiles are in fact bracketed by results from the Goldan-Vedder analyses shown in Figure 6.

Despite large differences in absolute magnitudes, a common feature of all data sets is the relatively slow mixing between about 430 and 500 K (19-21 km). Faster

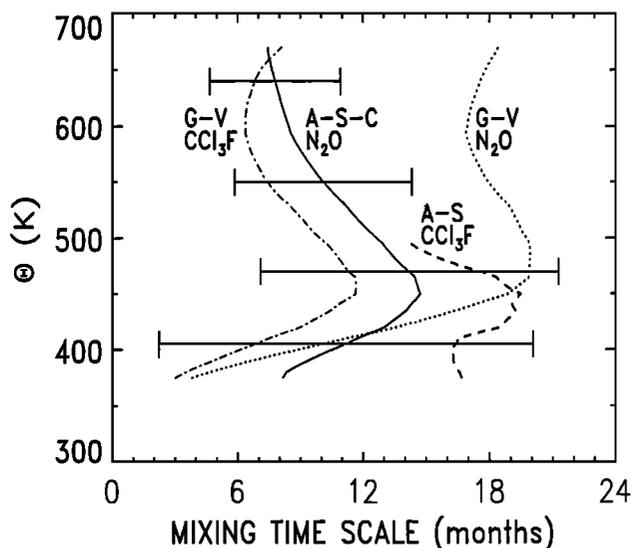


Figure 6. Timescale for mixing into the tropics based on fits to the data shown in Figure 1 (dash-dot, labeled G-V (Goldan-Vedder) CCl_3F), Figure 2 (dotted, labeled G-V N_2O), Figure 3 (dashed, labeled A-S (ASHOE-SPADE) CCl_3F), and Figure 4 (solid, labeled A-S-C (ASHOE-SPADE-CLAES) N_2O). Error bars represent the level of uncertainty for the A-S-C N_2O derived mixing time, as shown in Figure 7.

rates of mixing are indicated above and below this implied barrier, although these tendencies should be viewed with caution in light of the large level of uncertainty (see below). Nevertheless, it is of interest to note that the morphology of mixing suggested from Figure 6 is similar to that observed for export out of the polar vortices [e.g., Schoeberl *et al.*, 1992]; slow mixing in both cases is associated with strong gradients in potential vorticity [Trepte and Hitchman, 1992; Plumb *et al.*, 1994].

Uncertainties in the derived mixing times are estimated to be about 55% near 450 K, decreasing to 40%–46% above 500 K in the case of the A-S-C N₂O mixing time. Contributions to this uncertainty are displayed in Figure 7 which also includes the total uncertainty assuming that individual components add in quadrature. Calculated values of radiative heating ($\pm 20\%$ – 25% including effects of tropical cirrus) and photochemical loss ($\pm 15\%$ – 20% [Minschwaner *et al.*, 1993]) both make contributions to the overall uncertainty. Additionally, the large scatter of the data introduces uncertainties in our χ' , χ , and $\partial\chi/\partial\theta$ terms in (1). In regions where the quantity $\chi - \chi'$ is comparable to the uncertainties in χ and χ' , such as below 400 K, computed mixing times are uncertain by at least a factor of 2.

Owing to the large temporal range of the data, our results represent mean values averaged over at least 1 year. These mixing times thus represent the bulk effect of entrainment into the tropical lower stratosphere

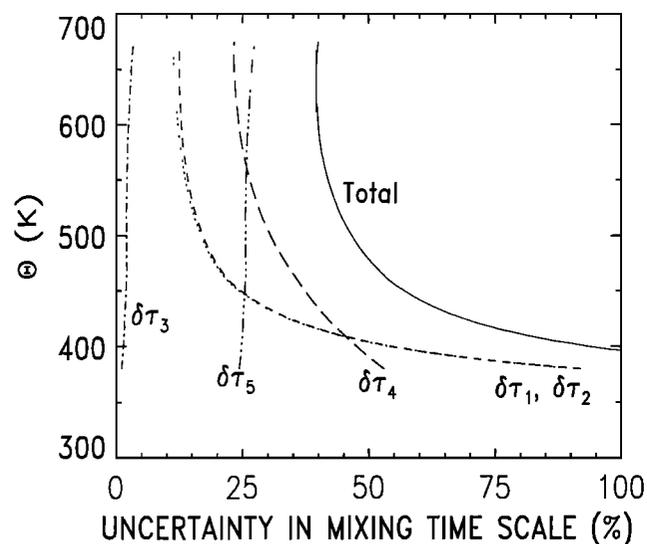


Figure 7. Components to the uncertainty for the A-S-C N₂O mixing timescale shown in Figure 6. The curves labeled $\delta\tau_1$ and $\delta\tau_2$ are contributions from uncertainties in χ and χ' , respectively, assuming these uncertainties both increase with θ from 7 to 15 ppb within the altitudes of interest; $\delta\tau_3$ is due to an 18% uncertainty in L ; $\delta\tau_4$ arises from a 50% uncertainty in $\partial\chi/\partial\theta$ at 380 K, decreasing to 20% at 650 K; and $\delta\tau_5$ originates from a 23% uncertainty in θ . Total uncertainty is calculated by adding the individual components in quadrature. Uncertainties for the other mixing profiles shown in Figure 6 are generally larger.

due to synoptic and planetary scale wave mixing. The strength of the subtropical barrier may also depend on season as well as the phase of the quasibiennial oscillation in tropical winds and temperature [Hitchman *et al.*, 1994], but these effects are beyond the scope of this paper. Model simulations also suggest hemispheric differences in mixing out of the tropics [Chen *et al.*, 1994]. We found small hemispheric differences in the CLAES midlatitude distributions, although the impact on computed mixing into the tropics was not significant.

Species used in this analysis must satisfy two criteria. First, the local photochemical lifetime in the tropics must be larger than the derived mixing time so that the effect of mixing is apparent in the tropical profile. Second, there must be a clear distinction between tropical and midlatitude profiles (given by $\chi - \chi'$ in (1)). We believe N₂O is better suited for this analysis because (1) the local photochemical lifetime exceeds 1 year in the tropics below about 28 km and (2) the reaction with O¹D becomes more important at middle and high latitudes in setting the total loss rate, thus producing a sufficiently large gradient between tropical and midlatitude profiles. Our highest confidence therefore lies in results from the A-S-C N₂O data. It is not clear why timescales derived from the Goldan-Vedder N₂O data are significantly larger above 500 K.

A quantity of interest is the fraction of midlatitude air ζ entrained into the tropical ascent region. If (1) is rewritten in terms of the difference Δ between the tropical and midlatitude mixing ratio of an infinitely long-lived tracer in steady state ($L = 0$, $\partial/\partial t = 0$) and we define $\zeta = (\Delta_o - \Delta)/\Delta_o$, where Δ_o is the initial difference at the isentropic surface intersecting the tropical tropopause, then ζ is determined by

$$\zeta(\theta) = 1 - \exp\left(-\int_{\theta_o}^{\theta} \frac{1/\tau}{\theta} d\theta\right) \quad (4)$$

Owing to the altitude dependence of mixing and tracer vertical profiles, this quantity is not the same as the apparent mixing fraction derived along surfaces of constant θ . For example, the apparent mixing fraction (obtained by simply comparing the observed and zero mixing profiles on a constant theta level) for the A-S-C N₂O data is about 30% at 650 K. As shown below, the actual fraction of midlatitude air is approximately 65%. The reason this value is much higher than the apparent mixing fraction is that much of the mixing is occurring at lower altitudes where the difference between the tropical and midlatitude mixing ratios is smaller.

Figure 8 shows the calculated profiles of ζ corresponding to mixing timescales from Figure 6. Differences between data sets are less than 40% above 500 K despite much larger disparities in profiles of τ ; apparently, ζ is a more robust quantity for the present method of analysis. As an indication of the effect of uncertainties in the vertical profile of τ , Figure 8 also includes a ζ profile corresponding to a fixed τ of 12 months. The fraction of midlatitude air entrained into the tropics increases from 0% at the tropopause to more than 50% above 550 K

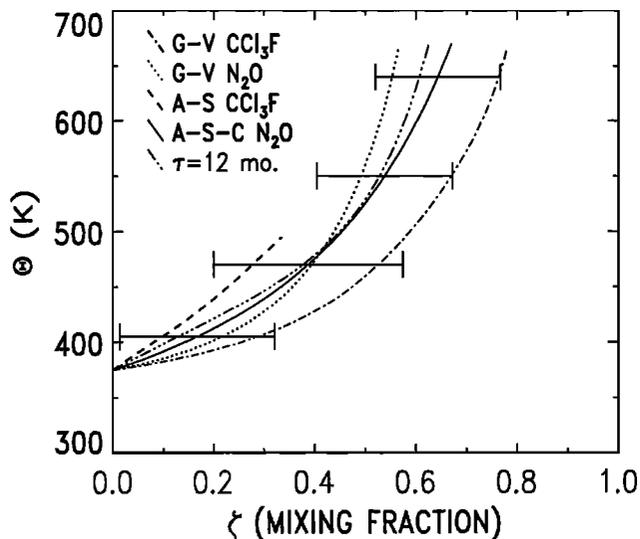


Figure 8. Fraction of midlatitude air entrained into the tropics based on the mixing time curves shown in Figure 6 (see (4) and discussion in text). Additionally, a ζ profile is shown that corresponds to a fixed τ of 12 months, independent of θ . Error bars represent the level of uncertainty for the mixing fraction derived from the A-S-C N_2O analysis which includes the uncertainty in τ and in θ . The latter uncertainty, however, very nearly cancels in the expression for ζ .

for all cases except the ASHOE-SPADE CCl_3F profile, which reaches 32% at 490 K. Note that the definition of midlatitude used here does not distinguish between the ultimate origins of midlatitude air parcels, only that they have spent sufficient time in the extratropics to acquire midlatitude characteristics of long-lived tracer amounts.

Our results for ascent with quasi-continuous mixing are qualitatively consistent with observations of the upward propagating seasonal signal of $2\text{xCH}_4 + \text{H}_2\text{O}$ presented by Mote *et al.* [1996]. They found that the amplitude of the seasonal signal (comparable to $1 - \zeta$ in the present analysis) diminished sharply between the tropopause and about the 500 K level but changed much less rapidly at higher altitudes, in agreement with the mixing fractions shown here in Figure 8. The signal attenuation below 500 K implies a lower limit to the mixing timescale of 7 to 9 months, also in agreement with our values of τ shown in Figure 6. However, the lower limit of 15-18 months derived by Mote *et al.* between 500 and 700 K is somewhat longer than our best estimate given by the A-S-C N_2O data in Figure 6.

Conclusion

The distributions of long-lived tracers N_2O and CCl_3F in the lower stratosphere indicate that a potential barrier exists for isentropic mixing of midlatitude air into the tropics. The tropical stratosphere may be effectively isolated for more than 1 year between 430 and 500 K. Although enhanced rates of mixing are inferred

above and below this region, the level of uncertainty does not preclude the possibility of a fixed value of τ , independent of θ . Additional data from the tropical lower stratosphere, for example, from the ongoing Stratospheric Tracers of Atmospheric Transport missions, would be highly desirable to more effectively constrain mean tropical profiles of long-lived tracers. It is unlikely, however, that the present method can be used to unambiguously derive a vertical profile of mixing in which variations with respect to θ will exceed the level of uncertainty. The mean timescales derived here suggest that further analysis would be best suited using N_2O or molecules with similar photochemical time constants, such as CCl_2F_2 (CFC-12). Recent work by Volk *et al.* (In situ measurements constraining exchange between the tropics and the middle latitudes in the lower stratosphere, submitted to *Science*, 1996) has provided mean mixing times below 490 K by employing tracer correlations and involving a more extensive suite of source gases and photochemically active species. Their mean in-mixing time of 13.6 months below 490 K agrees favorably with values reported here.

The implied entrainment of midlatitude air into the tropical ascent region is remarkably large; up to 50% of the air above the 550 K surface is of midlatitude origin. Recently, Plumb [1996] has described a conceptual "pipe" model of the tropical ascent region which successfully addresses shortcomings in the Plumb and Ko [1992] global mixing formulation. We find evidence here which supports a pipe-like behavior in the lower stratosphere. Owing to the integrated effect of isentropic mixing, however, the tropical import barrier appears to possess an altitude-dependent permeability.

Acknowledgments. We thank S. C. Wofsy and R. J. Salawitch for useful discussions on the analysis of the balloon tracer data, and Christos Zerefos for his efforts as guest editor of this manuscript. The radiative heating code was provided by J. T. Kiehl and B. P. Briegleb of the Climate and Global Dynamics Division of the National Center for Atmospheric Research. K.M. acknowledges support from the UARS Guest Investigator Program, grant NAG 5-2847. Support for A.E.D. was provided through a postdoctoral fellowship from the National Research Council.

References

- Anderson, G. P., S. A. Clough, F. X. Kneizys, J. H. Chetwynd, and E. P. Shettle, AFGL atmospheric constituent profiles (0-120km), *AFGL Tech. Rep., AFGL-TR-86-0110*, 43 pp., Air Force Phillips Lab., Hansom Air Force Base, Mass., 1986.
- Avallone, L. M., and M. J. Prather, Photochemical evolution of ozone in the lower tropical stratosphere, *J. Geophys. Res.*, **101**, 1457-1461, 1996.
- Barkstrom, B. R., The Earth Radiation Budget Experiment (ERBE), *Bull. Am. Meteorol. Soc.*, **65**, 1170-1185, 1984.
- Briegleb, B. P., Delta-Eddington approximation for solar radiation in the NCAR community climate model, *J. Geophys. Res.*, **97**, 7603-7612, 1992a.
- Briegleb, B. P., Longwave band model for thermal radiation in climate studies, *J. Geophys. Res.*, **97**, 11,475-11,485, 1992b.

- Brewer, A. W., Evidence for a world circulation provided by measurements of helium and water vapor distribution in the stratosphere, *Q. J. R. Meteorol. Soc.*, **75**, 351-363, 1949.
- Chen, P., J. R. Holton, A. O'Neill, and R. Swinbank, Isentropic mass exchange between the tropics and extratropics in the stratosphere, *J. Atmos. Sci.*, **51**, 3006-3018, 1994.
- Considine, D. B., A. R. Douglass, and C. H. Jackman, Sensitivity of two-dimensional model predictions of ozone response to stratospheric aircraft: An update, *J. Geophys. Res.*, **100**, 3075-3090, 1995.
- Dessler, A. E., K. Minschwaner, E. M. Weinstock, E. J. Hints, J. G. Anderson, and J. M. Russell, III, The effects of tropical cirrus clouds on the abundance of lower stratospheric ozone, *J. Atmos. Chem.*, **23**, 209-220, 1996.
- Dobson, G. M. B., Origin and distribution of polyatomic molecules in the atmosphere, *Proc. Roy. Soc. London, A*, **236**, 187-193, 1956.
- Ehhalt, D. H., E. P. Röth, and U. Schmidt, On the temporal variance of stratospheric trace gas concentrations, *J. Atmos. Chem.*, **1**, 27-51, 1983.
- Elkins, J. W., T. M. Thompson, T. H. Swanson, J. H. Butler, B. D. Hall, S. O. Cummings, D. A. Fisher, and A. G. Raffo, Decrease in the growth rates of atmospheric chlorofluorocarbons 11 and 12, *Nature*, **364**, 780-783, 1993.
- Elkins, J. W., et al., Airborne gas chromatograph for in situ measurements of long-lived species in the upper troposphere and lower stratosphere, *Geophys. Res. Lett.*, **24**, 347-350, 1996.
- Goldan, P. D., W. C. Kuster, D. L. Albritton, and A. L. Schmeltekopf, Stratospheric CFCl₃, CF₂Cl₂, and N₂O height profile measurements at several latitudes, *J. Geophys. Res.*, **85**, 413-423, 1980.
- Goldan, P. D., W. C. Kuster, A. L. Schmeltekopf, F. C. Fehsenfeld, and D. L. Albritton, Correction of atmospheric N₂O mixing ratio data, *J. Geophys. Res.*, **86**, 5385-5386, 1981.
- Hints, E. J., et al., SPADE H₂O measurements and the seasonal cycle of stratospheric water vapor, *Geophys. Res. Lett.*, **21**, 2559-2562, 1994.
- Hitchman, M. H., M. McKay, and C. R. Trepte, A climatology of stratospheric aerosol, *J. Geophys. Res.*, **99**, 20,689-20,700, 1994.
- Holton, J. R., On the global exchange of mass between the stratosphere and troposphere, *J. Atmos. Sci.*, **47**, 392-395, 1990.
- Lait, L. R., An alternative form for potential vorticity, *J. Atmos. Sci.*, **51**, 1754-1759, 1994.
- McIntyre, M. E., and T. N. Palmer, The "surf zone" in the stratosphere, *J. Atmos. Terr. Phys.*, **46**, 825-849, 1984.
- Mote, W. M., K. H. Rosenlof, M. E. McIntyre, E. S. Carr, J. C. Gille, J. R. Holton, J. S. Kinnersley, H. C. Pumphrey, J. M. Russell III, and J. W. Waters, An atmospheric tape recorder: The imprint of tropical tropopause temperatures on stratospheric water vapor, *J. Geophys. Res.*, **101**, 3989-4006, 1996.
- Minschwaner, K., R. J. Salawitch, and M. B. McElroy, Absorption of solar radiation by O₂: Implications for O₃ and lifetimes of N₂O, CFCl₃, and CF₂Cl₂, *J. Geophys. Res.*, **98**, 10,543-10,561, 1993.
- Murphy, D. M., et al., Reactive nitrogen and its correlation with ozone in the lower stratosphere and upper troposphere, *J. Geophys. Res.*, **98**, 5325-5332, 1993.
- Newell, R. E., Transfer through the tropopause and within the stratosphere, *Q. J. R. Meteorol. Soc.*, **89**, 167-204, 1963.
- Plumb, R. A., A "tropical pipe" model of stratospheric transport, *J. Geophys. Res.*, **101**, 3957-3972, 1996.
- Plumb, R. A., and M. K. W. Ko, Interrelationships between mixing ratios of long-lived stratospheric constituents, *J. Geophys. Res.*, **97**, 10,145-10,156, 1992.
- Plumb, R. A., et al., Intrusions into the lower stratospheric Arctic vortex during the winter of 1991-1992, *J. Geophys. Res.*, **99**, 1089-1105, 1994.
- Podolske, J. R., and M. Loewenstein, Airborne tunable diode laser spectrometer for trace-gas measurement in the lower stratosphere, *Appl. Opt.*, **32**, 5324-5333, 1993.
- Randel, W. J., et al., Stratospheric transport from the tropics to middle latitudes by planetary-wave mixing, *Nature*, **365**, 533-535, 1993.
- Roche, A. E., J. B. Kumer, J. L. Mergenthaler, G. A. Ely, W. G. Uplinger, J. F. Potter, T. C. James, and L. W. Sterritt, The Cryogenic Limb Array Spectrometer (CLAES) on UARS: experiment description and performance, *J. Geophys. Res.*, **98**, 10763-10776, 1993.
- Rossow, W. B., and R. A. Schiffer, ISCCP cloud data products, *Bull. Am. Meteorol. Soc.*, **72**, 2-20, 1991.
- Schoeberl, M. R., L. R. Lait, P. A. Newman, and J. E. Rosenfield, The structure of the polar vortex, *J. Geophys. Res.*, **97**, 7859-7882, 1992.
- Trepte, C. R., and M. H. Hitchman, Tropical stratospheric circulation deduced from satellite aerosol data, *Nature*, **355**, 626-628, 1992.
- Vedder, J. F., B. J. Tyson, R. B. Brewer, C. A. Boitnott, and E. C. Y. Inn, Lower stratosphere measurements of variation with latitude of CF₂Cl₂, CFCl₃, CCl₄, and N₂O profiles in the northern hemisphere, *Geophys. Res. Lett.*, **5**, 33-36, 1978.
- Vedder, J. F., E. C. Y. Inn, B. J. Tyson, and C. A. Boitnott, Measurements of CF₂Cl₂, CFCl₃, and N₂O in the lower stratosphere between 2°S 73°N latitude, *J. Geophys. Res.*, **86**, 7363-7368, 1981.
- Waugh, D. W., Subtropical stratospheric mixing linked to disturbances in the polar vortices, *Nature*, **365**, 535-537, 1993.

K. R. Chan, M. Loewenstein, and J. R. Podolske, NASA Ames Research Center, Moffett Field, CA 94035.

A. E. Dessler, NASA Goddard Space Flight Center, Greenbelt, MD 20771.

J. W. Elkins and C. M. Volk, Climate Monitoring and Diagnostics Laboratory, NOAA, Boulder, CO 80303.

D. W. Fahey, Aeronomy Laboratory, NOAA, Boulder, CO 80303.

K. Minschwaner, Department of Physics, New Mexico Institute of Mining and Technology, Socorro, NM, 87801. krm@kestrel.nmt.edu.

A. E. Roche, Lockheed Palo Alto Research Laboratory, Palo Alto, CA 94303.

(Received August 8, 1995; revised January 16, 1996; accepted January, 1996.)