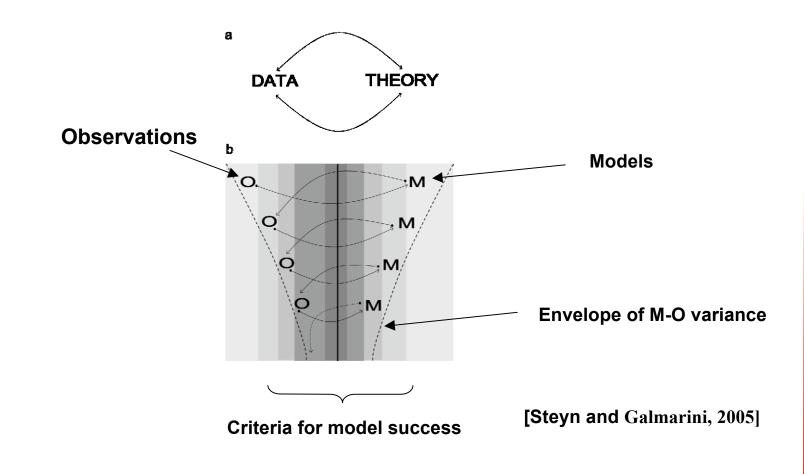
One-Dimensional Models

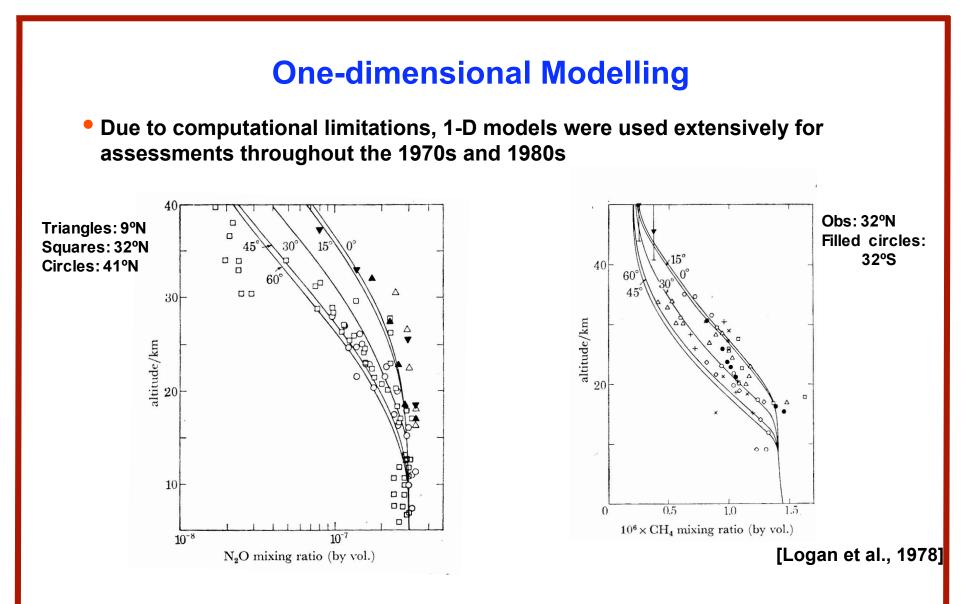
GCC Summer School, 2005 Banff, Alberta

> Dylan Jones Department of Physics University of Toronto

Observations and Models

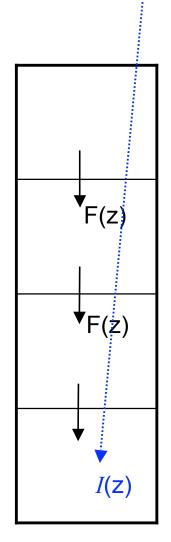


Choice of model should be guided by quality of observations and desired level of accuracy and precision of understanding or prediction of atmospheric phenomenon



 Despite their limitations, 1-D models adquately provided much of our early understanding of the trace constituents in the atmosphere - reflects limited spatio-temporal distribution of data in the pre-satellite era

One-Dimensional Framework



1-D models provide a latitude-longitude averaged (global or hemispheric) representation of the atmosphere

Continuity equation

$$\frac{\partial n(z)}{\partial t} = P - L - \frac{\partial}{\partial z} \left(K(z) M(z) \frac{\partial f(z)}{\partial z} \right)$$

M(z) = atmospheric number density n(z) = tracer number density f(z) = tracer mixing ratio K(z)= vertical

Radiation

$$I(z) = I_{o} \exp\left(-\frac{1}{\mu}\int_{z}^{\infty}\sigma n(z)dz\right)$$

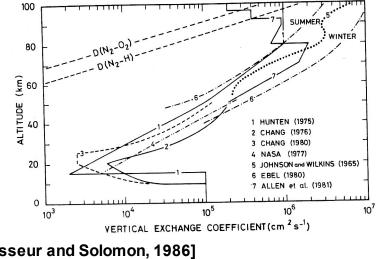
Vertical Diffusion Coefficients

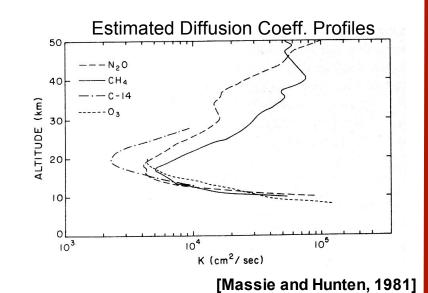
Use observed profiles of trace gases to empirically derive diffusion coefficient

vertical flux = -K(z)M(z)df(z)/dz

In steady state

$$K(z) = \left(M(z)\frac{df(z)}{dz}\right)^{-1} \left(\int_{z}^{\infty} (P(z) - L(z))dz\right)$$



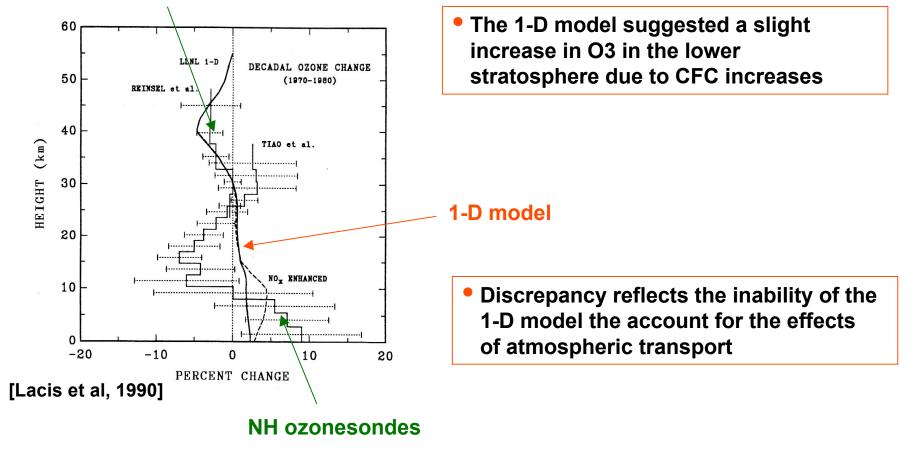


K profiles used in a range of 1-D models in 1960s, 1970s and 1980s

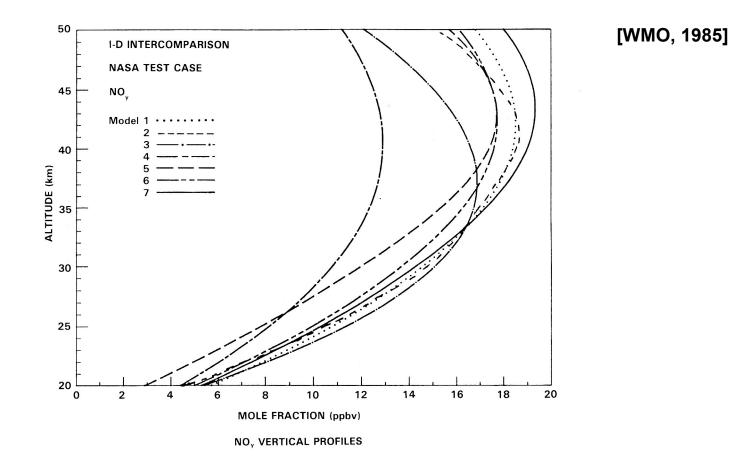
[Brasseur and Solomon, 1986]

Stratospheric Ozone Trends

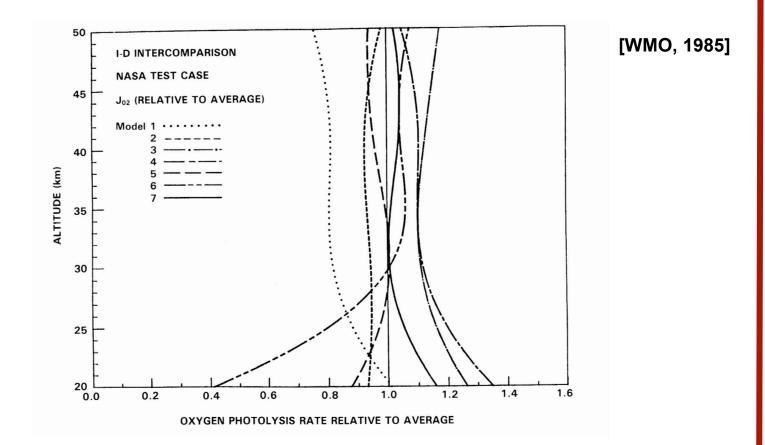
Umkehr measurements



Comparison of NO_y in 1-D Models

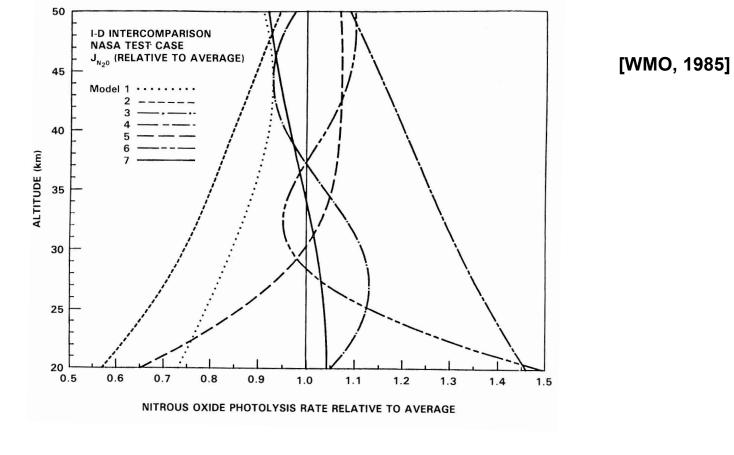


Comparison of O₂ Photolysis in 1-D Models



Photolysis of N₂O occurs in the region 173-240 nm \Rightarrow sensitive to Schumann-Runge bands (and Herzberg continuum) of O₂

Comparison of N₂O Photolysis in 1-D Models

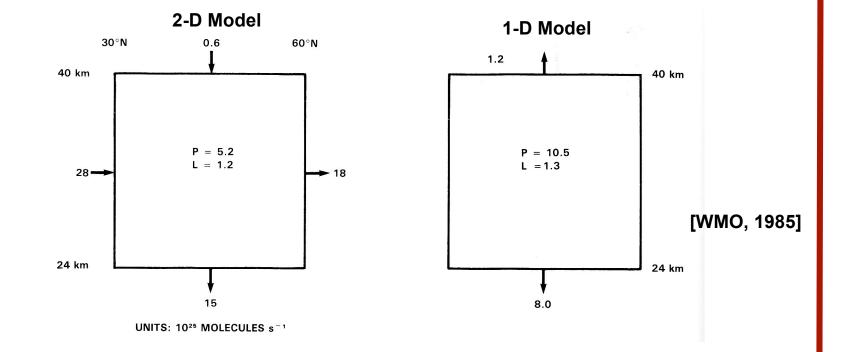


 $N_2O + hv \Rightarrow N_2 + O$

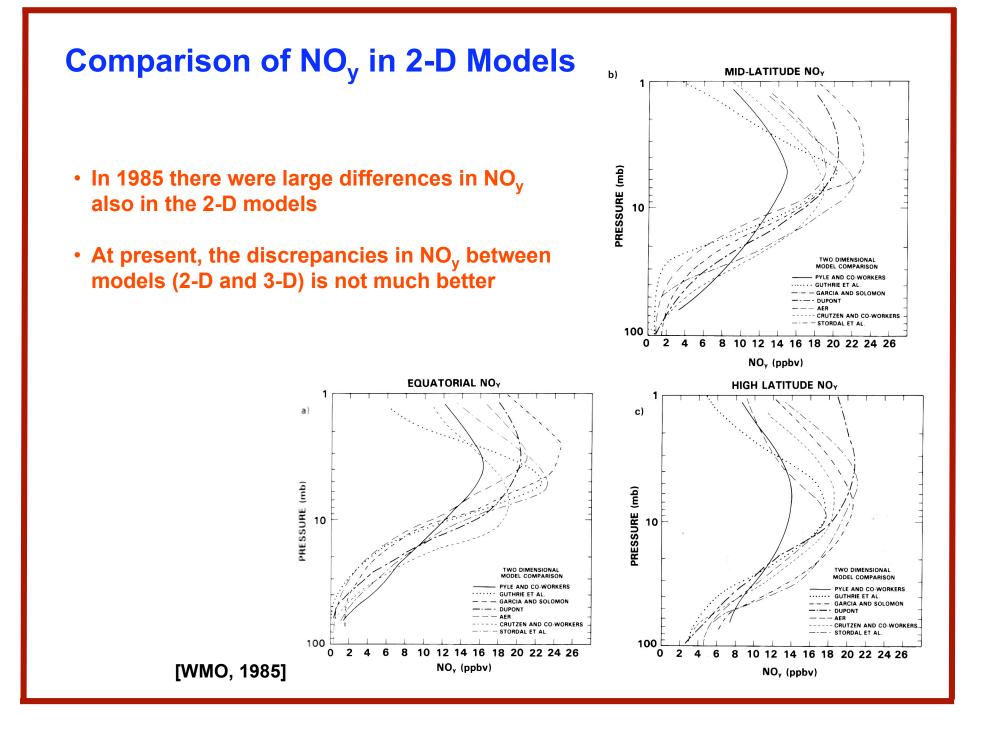
(90% of loss)

 $N_2O + O(^1D) \Rightarrow NO + NO$ (source of NO_y)

Comparison of NO_v budget in 1-D and 2-D Models

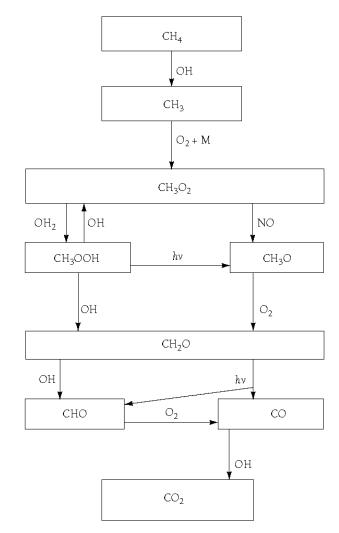


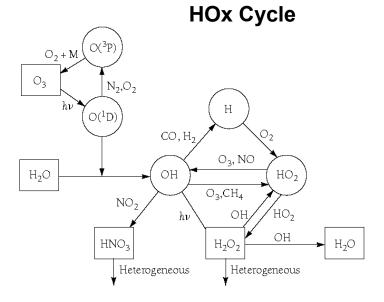
- In 2-D framework, meridional transport from source region in tropics important
- in 1-D model, balance is between in situ production (P) and vertical diffusive transport



Tropospheric chemistry

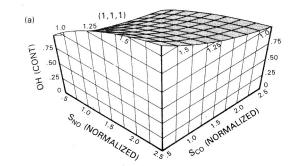
Methane Oxidation





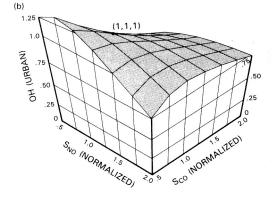
[M. B. McElroy, 2002]

Sensitivity of tropospheric OH to changes in emissions of NO and CO

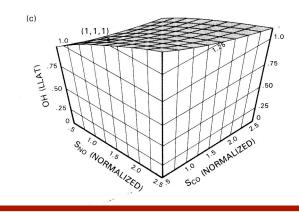


Continental

$S_{NO} = 8 \times 10^9 \text{ cm}^{-2} \text{s}^{-1}$	[NO _x]=0.2 ppb
$S_{CO} = 2.0 \times 10^{11}$	[CO] = 130 ppb
S _{CH4} = 6.5x10 ¹⁰	[CH ₄] = 1.7 ppm



Urban $S_{NO} = 7x10^{10} \text{ cm}^{-2}\text{s}^{-1}$ $[NO_x]=1.4 \text{ ppb}$ $S_{CO} = 8.0x10^{11}$ [CO] = 380 ppb $S_{CH4} = 7.8x10^{10}$ $[CH_4] = 1.7 \text{ ppm}$

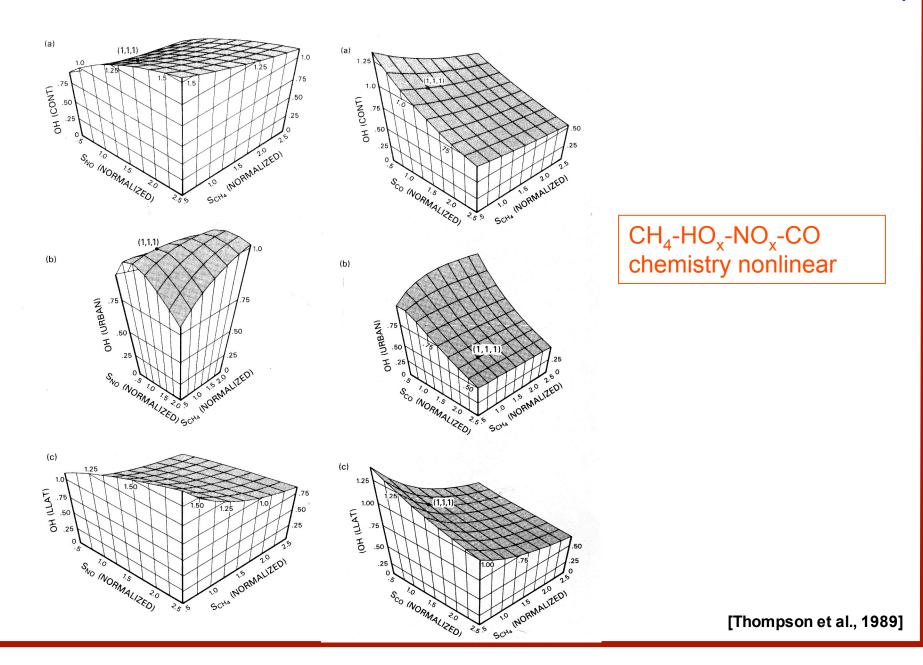


Low Latitude

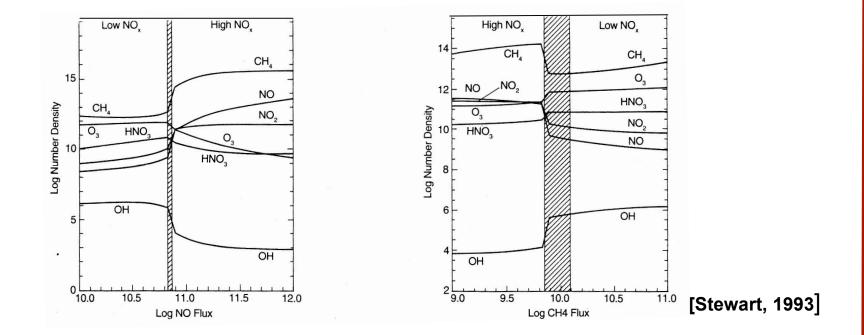
$$\begin{split} S_{\text{NO}} &= 6.7 \text{x} 10^8 \text{ cm}^{-2} \text{s}^{-1} & [\text{NO}_{\text{x}}] = 0.024 \text{ ppb} \\ S_{\text{CO}} &= 1.2 \text{x} 10^{11} & [\text{CO}] = 89 \text{ ppb} \\ S_{\text{CH4}} &= 1.0 \text{x} 10^{11} & [\text{CH}_4] = 1.7 \text{ ppm} \end{split}$$

[Thompson et al., 1989]

Sensitivity of tropospheric OH to changes in emissions of NO and CH₄



Instability in the CH_4 - NO_x - O_x - HO_x Chemical System (in a Box Model)

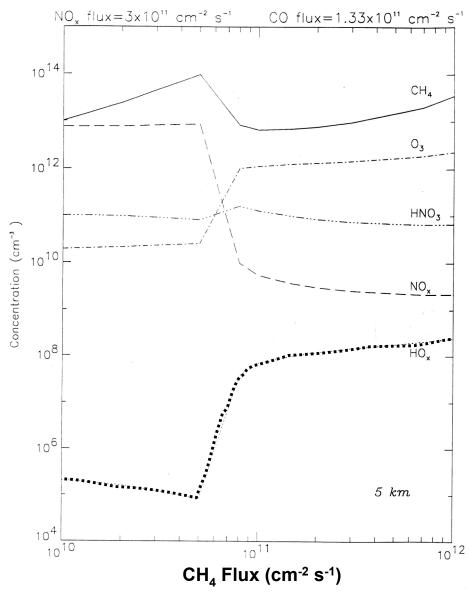


Abrupt transition between states characterized by low and high abundances of $\ensuremath{\mathsf{NO}_{\mathsf{X}}}$

Multiple Steady States NO OH 6.0 [..... 11.5 ______ 11.0 10.5 10.0 10.0 9.5 Log Number Density 2.0 4.5 Existence of MSS sensitive to abundance of other tracers e.g. no MSS if RH < 40%10.0 100000000000000000000 9.5 9.0 <u>10.00</u> 9.80 9.90 10.00 10.10 10.20 4.0 9.80 9.90 10.00 10.10 10.20 Log CH4 Flux Log CH4 Flux CH, H₂O₂ 9 Emminiari 14.5 mmmmmmmm 8 Atistic Territoria for the second sec Log Number Density 7 6 5 4 unstable states 13.5 stable states 13.0 12.5 <u>.....</u> 9.80 9.90 10.00 10.10 10.20 2 9.80 9.90 10.00 10.10 10.20 Log CH4 Flux Log CH4 Flux

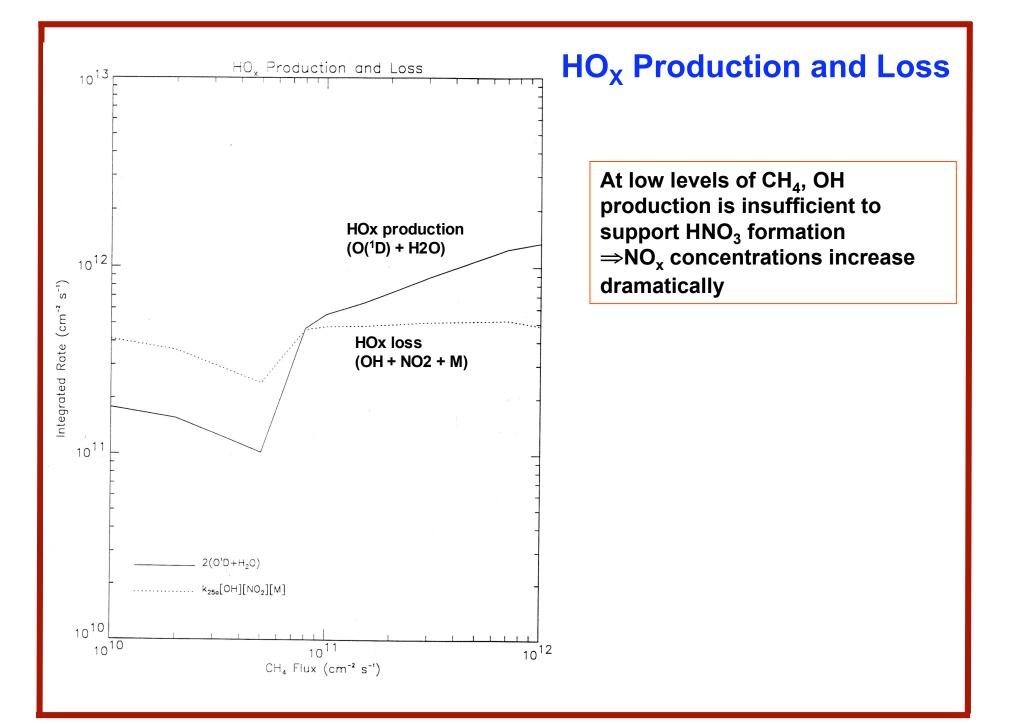
[Stewart, 1993]

1-D Simulation of Stewart's Results

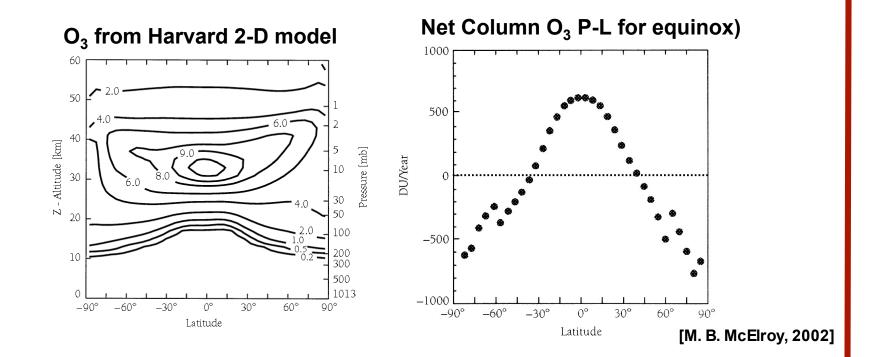


- Decreasing CH₄ emissions results in a transition to a high NO_x regime
- Total O₃ column decreases by a factor of 3 from the low to high NO_x regime (NO_x in lower strat ≈ 0.5 ppm)

- Results <u>suggest</u> the possibility of MSS in the 1-D model
- These abrupt transitions were not found in the 2-D model
- A more detailed search of state space is needed for the 1-D and 2-D models



Influence of Transport on Stratospheric O₃



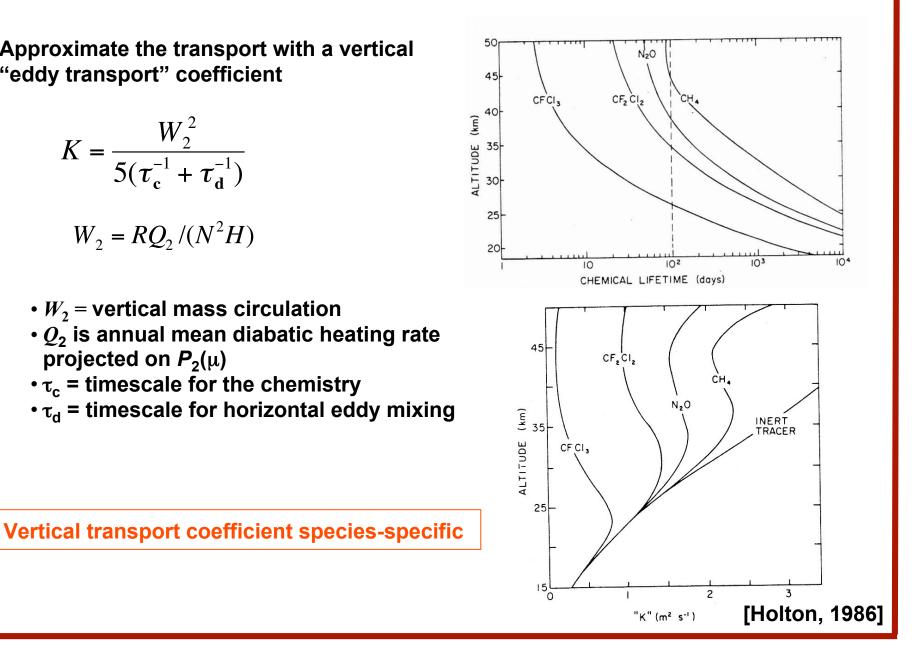
- Net O₃ production in tropical stratosphere reflects large flux of UV photons
- At high levels of CH4, oxidation shifts to tropical stratosphere which helps stabilize the system

A More Dynamically Consistent Approach

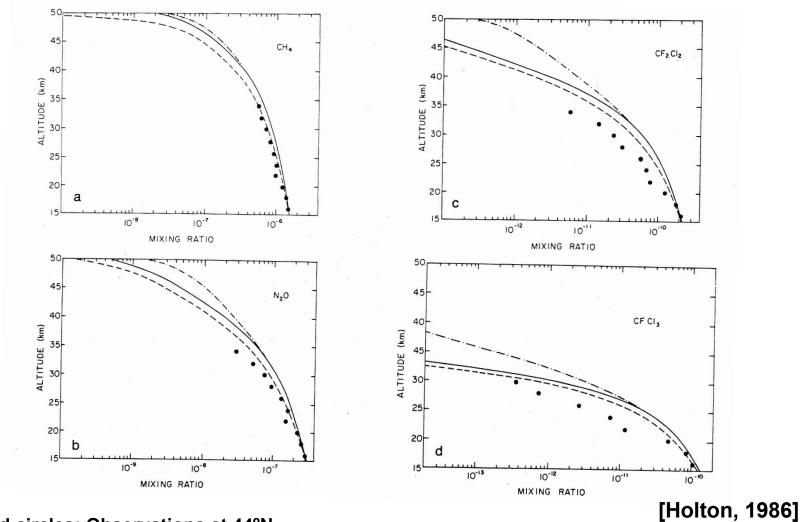
Approximate the transport with a vertical "eddy transport" coefficient

$$K = \frac{W_2^2}{5(\tau_c^{-1} + \tau_d^{-1})}$$
$$W_2 = RQ_2 / (N^2 H)$$

- W_2 = vertical mass circulation
- Q_2 is annual mean diabatic heating rate projected on $P_2(\mu)$
- τ_c = timescale for the chemistry
- τ_d = timescale for horizontal eddy mixing



Sensitivity of Tracer Profile to K_z



•Filled circles: Observations at 44°N
•Dashed-dotted line: global mean with K_z for inert tracer
•Solid line: global mean for species specific K_z
•Dashed line: simulated profiles for 45°N

Main Points of Lecture

- 1-D chemical-radiative-diffusive models were used extensively in 1970s and 1980s
- Modelling the vertical advective transport as a diffusive process provides a fundamentally flawed simulation of the distribution of the trace gases ⇒one K does not fit all
- More dynamically consistent approaches (e.g. Holton [1985]) were proposed but never adopted widely
- Despite these limitations, 1-D models provided much of our understanding of the trace constituents in the atmosphere during the pre-satellite era
- 1-D chemical-radiative models are currently used in 3-D model
 e.g., domain decomposition for parallel numerical simulations
- The limitations of 1-D models reflects importance of the meridional transport of tracers in the atmosphere ⇒ Brewer-Dobson circulation