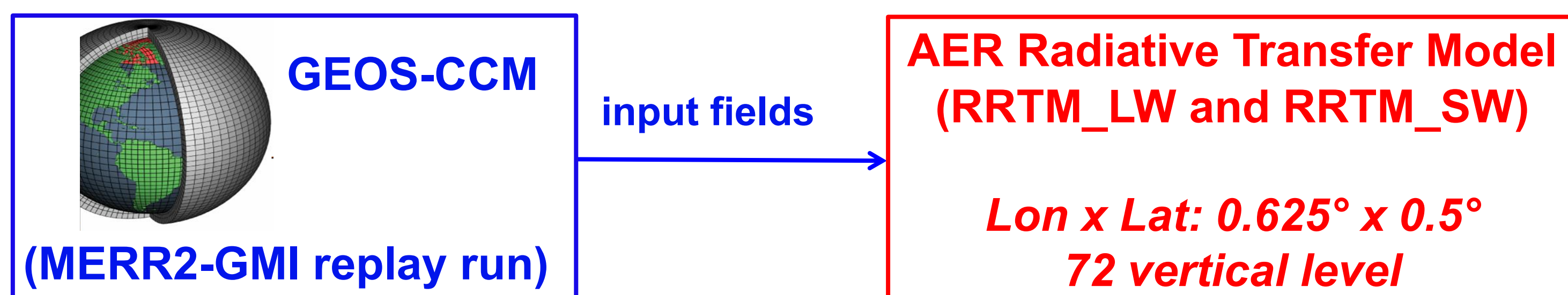


1: Summary

Liu et al. (2017) shows that strong stratospheric influence on the interannual variation of O₃ in the upper troposphere, where its radiative impact is largest. Given the observed and predicted net global decrease in emissions and the predicted increase in O₃ stratosphere-troposphere-exchange, distinguishing the role of stratospheric contribution from that of emissions to tropospheric O₃ variations and resulting changes in radiative forcing is critical to understand how they will change in the future.

In this study, we quantify the stratospheric and emissions contributions on tropospheric O₃ radiative effect (RE) using sensitivity runs of the AER offline radiative transfer model (RRTM) with different O₃ fields, including stratospheric ozone tracer (StratO₃) and O₃ simulated from the high-resolution GEOS-5 Replay run (~50 km). The StratO₃ is used to quantify O₃ of stratospheric origin in the troposphere at all locations and times. The offline module is set up so that all input fields except O₃ remain fixed. Our study shows that RE from tropospheric O₃ with stratospheric origin accounts for significant contributions to the global tropospheric O₃ RE.

2. Model and Approach



Input fields are reading from outputs from MERRA2-GMI replay run:

- Concentrations of molecular absorbers include water vapor, carbon dioxide, ozone, nitrous oxide, methane, oxygen, nitrogen
- Surface conditions including surface emissivity, skin temperature etc. (Reflected and emitted radiation from surface)
- View geometry (local zenith angle, polarization)
- Cloud influence: ice or water fraction and size, cloud-water-path (CWP)

Sensitivity runs with different O₃ input fields (O_{3_in}):

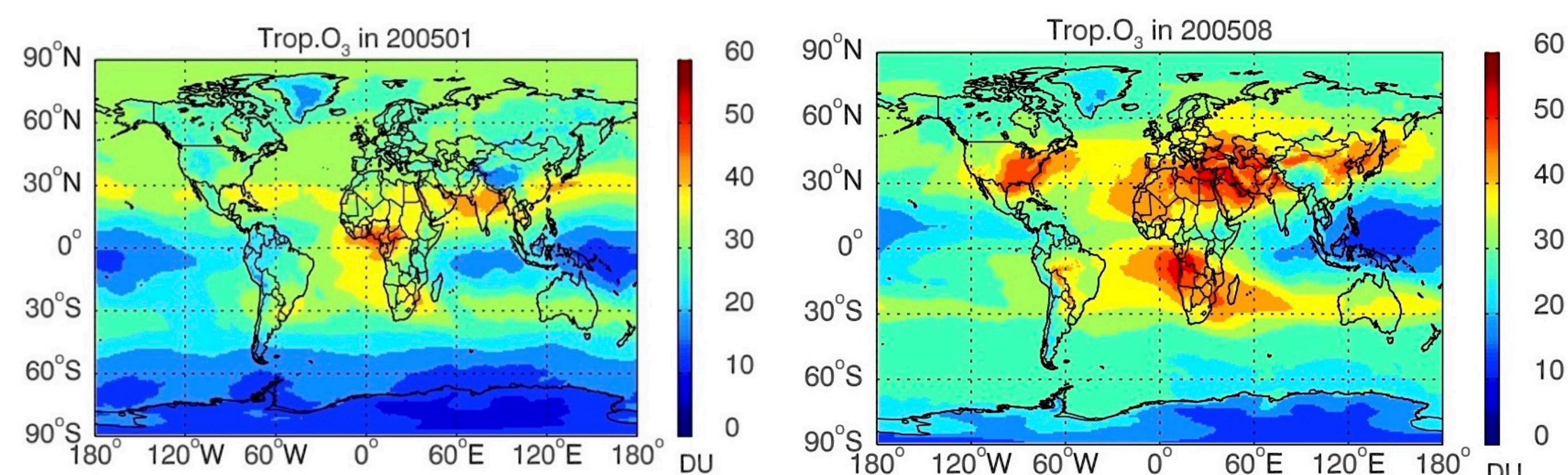
In the stratosphere: O_{3_in} = O₃

In the troposphere:

1. O_{3_in} = 0 ppb
2. O_{3_in} = O₃
3. O_{3_in} = StratO₃ (an artificial stratospheric O₃ tracer)
4. O_{3_in} = O₃ - StratO₃ (O₃ from emissions)

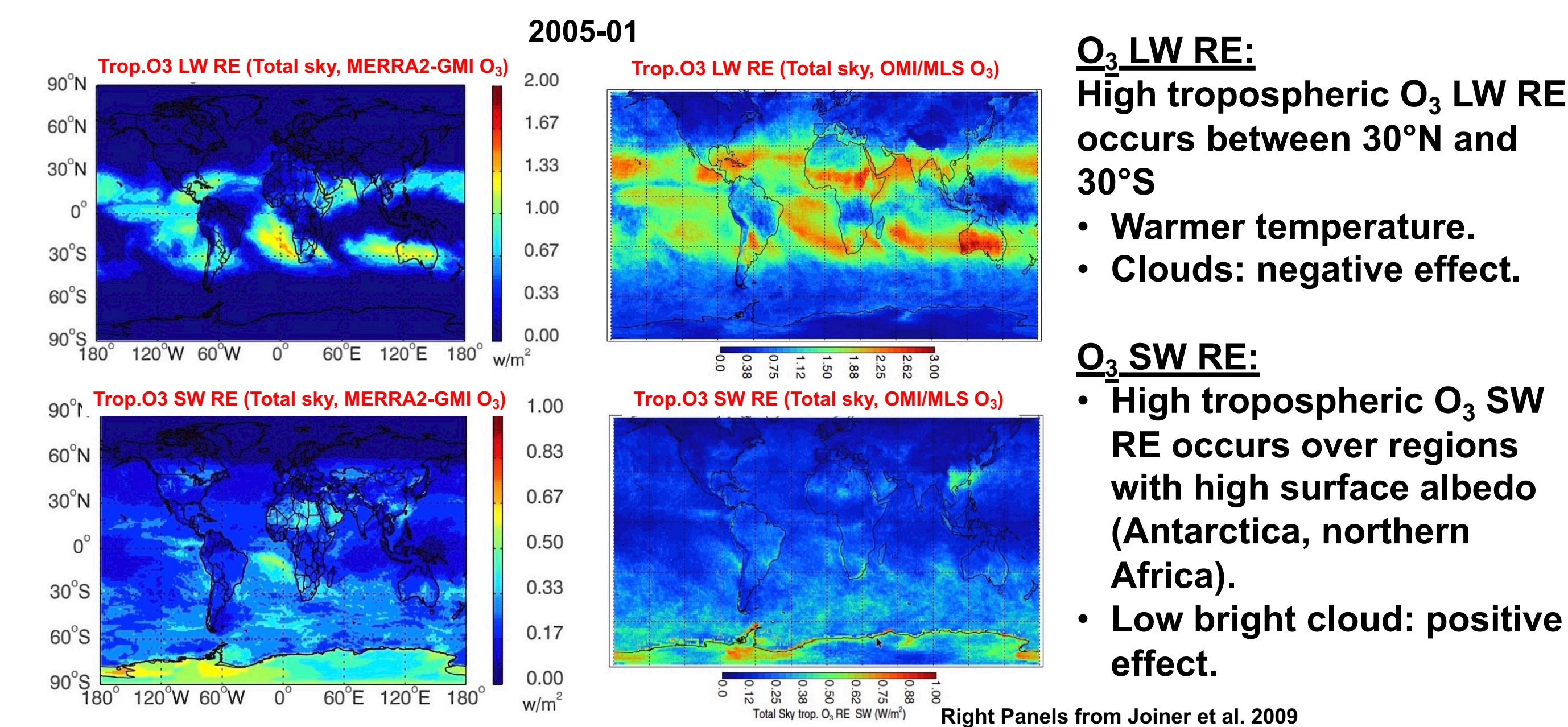
Tropospheric O ₃ RE	Irradiance difference (run 2 - run 1)
Tropospheric O ₃ RE from stratosphere	Irradiance difference (run 3 - run 1)
Tropospheric O ₃ RE from emissions	Irradiance difference (run 4 - run 1)

3.1. Tropospheric O₃ column (TOC)

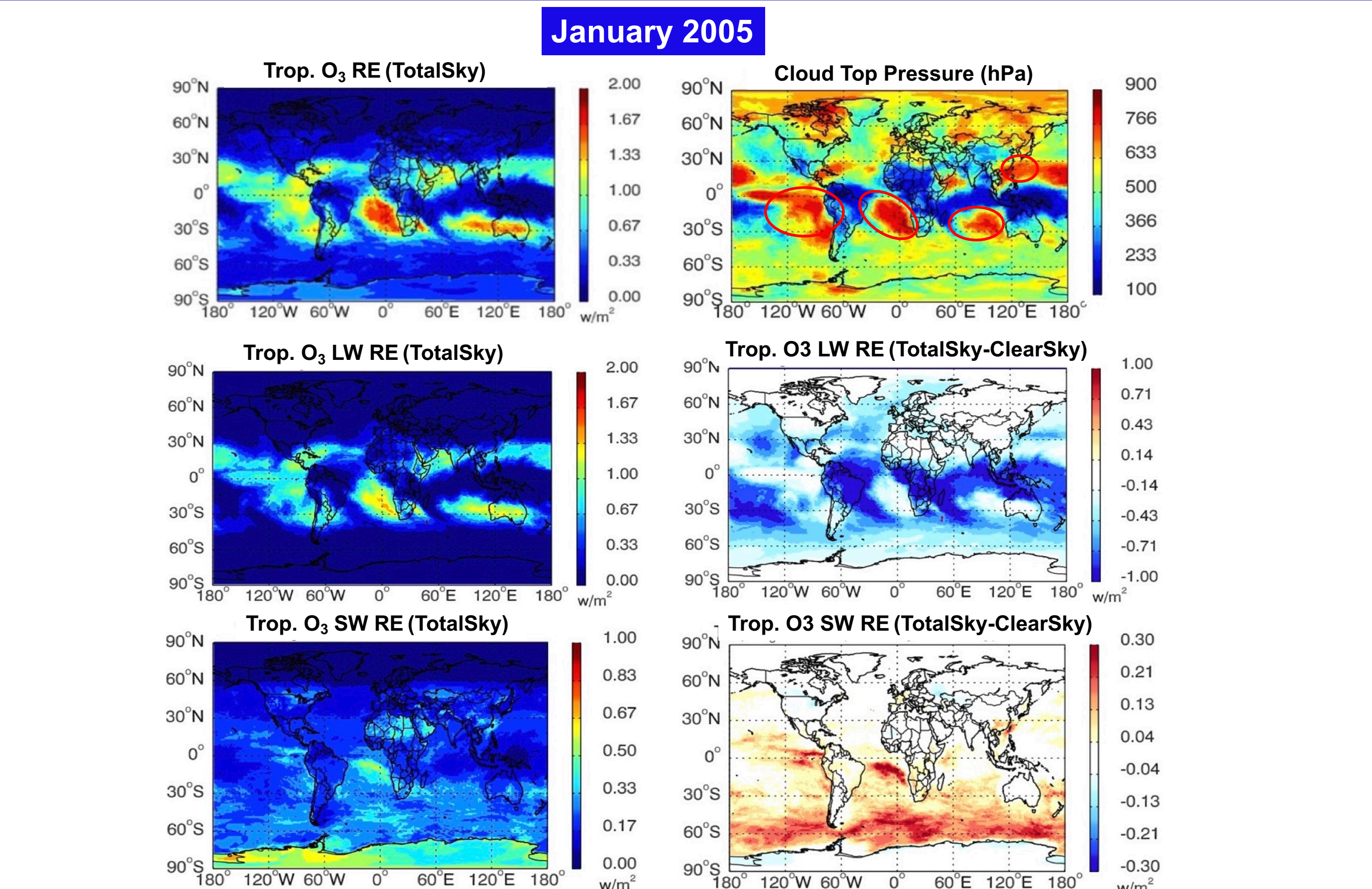


- Significant spatial variations of TOC, with O₃ minimum in west Pacific warm pool.
- January 2005: Elevated TOC occurs near subtropical jets with increased stratospheric O₃ influence. TOC maximum over western Africa.
- August 2005: TOC shows regional maximum over the downwind of polluted N.H. continents, and tropical south Atlantic (wave-1 pattern).

3.2. Model agrees with tropospheric O₃ RE calculation based on satellite observations

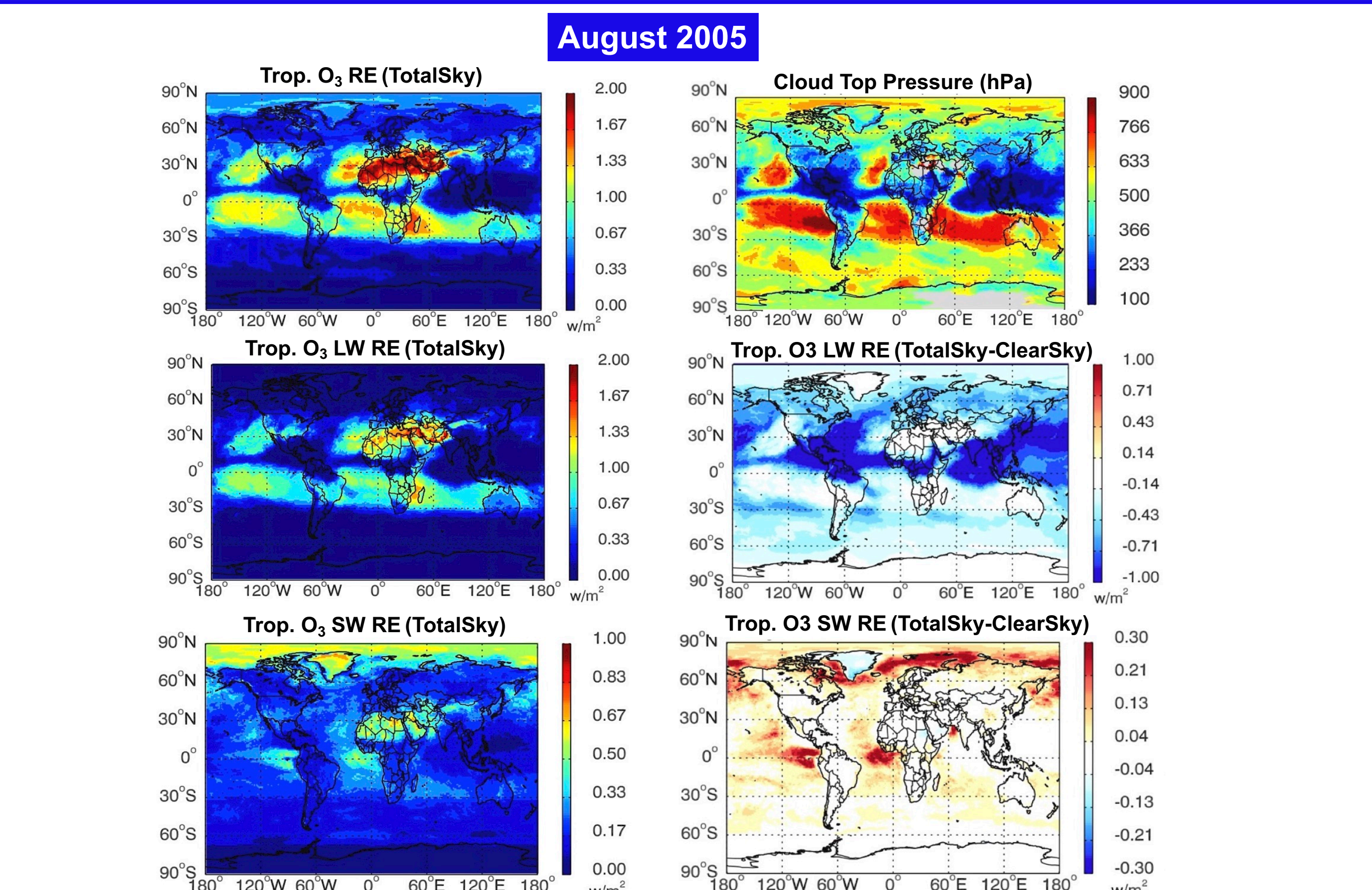


4. Tropospheric O₃ radiative effect and clouds effect

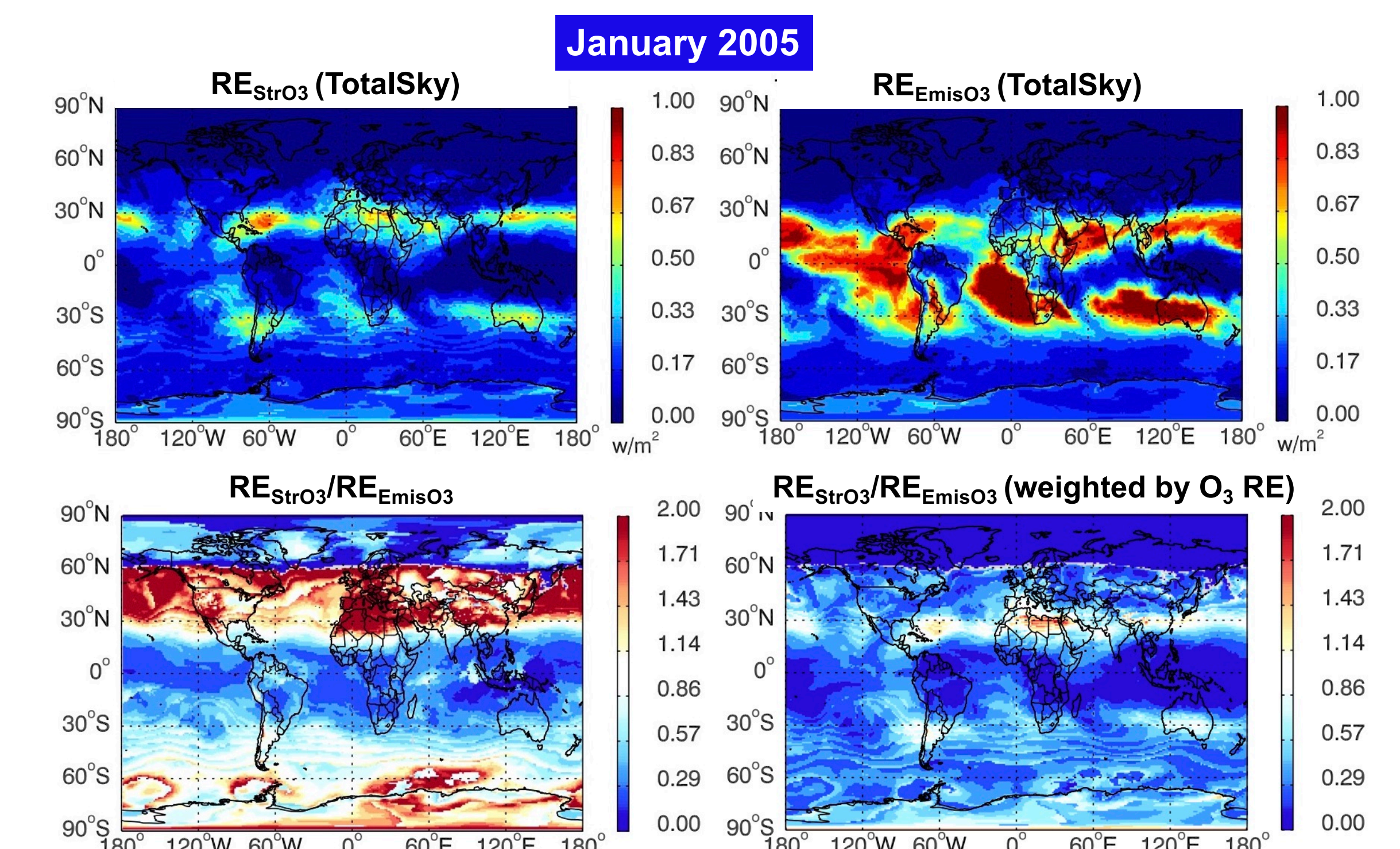


High tropospheric O₃ RE occurs over:

- Regions around subtropical Jets: Elevated O₃ + small cloud negative effect.
- Regions with low clouds: much weaker negative effects on LW RE + positive effects on SW RE.

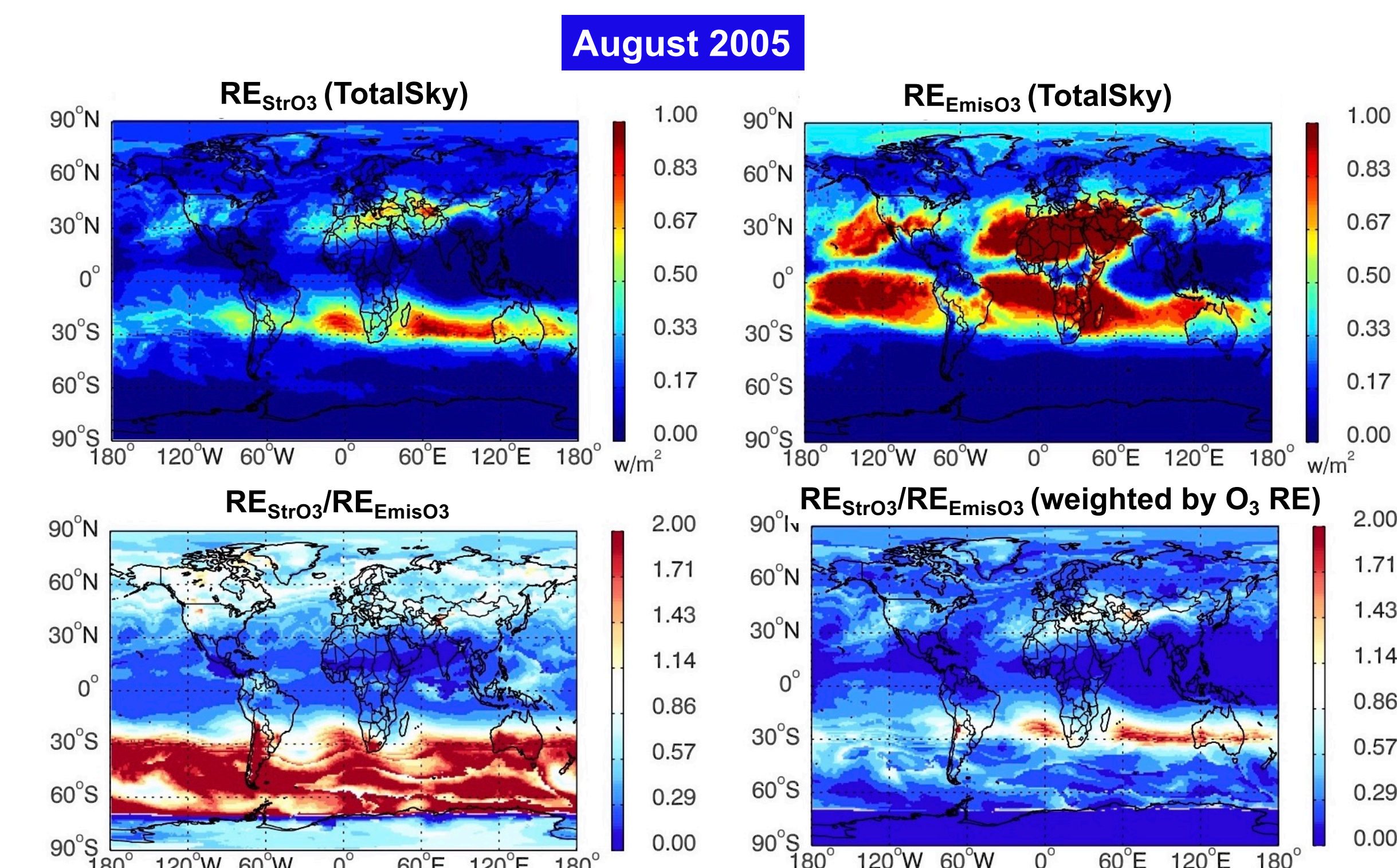


5. Stratospheric and emission components of tropospheric O₃ radiative effect



The StrO₃/EmisO₃ RE ratio is weighted by trop. O₃ RE to highlight regions where stratospheric contribution is important to the global radiation budget.

RE from trop. O₃ with stratospheric origin accounts for ~40% of the global trop. O₃ RE, with regional maximum around subtropical jets, terrain area of Antarctica in January 2005.



Stratospheric O₃ accounts for ~36% of the global trop. O₃ RE, with regional maximum over S.H. subtropical jet regions, northern Africa, eastern Mediterranean and the Middle East in August 2005.

Key Findings:

1. RE from tropospheric O₃ with stratospheric origin accounts for ~40% and 36% of the global tropospheric O₃ RE in January and August 2005.
2. On regional scale, RE from tropospheric O₃ with stratospheric origin exceeds that from emissions and shows dominant contributions to tropospheric O₃ RE over following regions:
January: around subtropical jets, terrain area of Antarctica, and southern ocean.
August: SH subtropical jet regions, the eastern Mediterranean and the Middle East.

REFERENCE:

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Liu, J., Rodriguez, J. M., Steenrod, S. D., Douglass, A. R., Logan, J. A., Olsen, M. A., Wargan, K., and Ziemke, J. R.: Causes of interannual variability over the southern hemispheric tropospheric ozone maximum, *Atmos. Chem. Phys.*, 17, 3279-3299, <https://doi.org/10.5194/acp-17-3279-2017>, 2017.

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