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LETTER

Diverse policy implications for future ozone and surface UV in a changing climate

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Abstract

Due to the success of the Montreal Protocol in limiting emissions of ozone-depleting substances, concentrations of atmospheric carbon dioxide, nitrous oxide, and methane will control the evolution of total column and stratospheric ozone by the latter half of the 21st century. As the world proceeds down the path of reducing climate forcing set forth by the 2015 Conference of the Parties to the United Nations Framework Convention on Climate Change (COP 21), a broad range of ozone changes are possible depending on future policies enacted. While decreases in tropical stratospheric ozone will likely persist regardless of the future emissions scenario, extratropical ozone could either remain weakly depleted or even increase well above historical levels, with diverse implication for ultraviolet (UV) radiation. The ozone layer’s dependence on future emissions of these gases creates a complex policy decision space for protecting humans and ecosystems, which includes unexpected options such as accepting nitrous oxide emissions in order to maintain historical column ozone and surface UV levels.

1. Introduction

Now that the world has agreed to curtail global warming following the 2015 Conference of the Parties to the United Nations Framework Convention on Climate Change (COP 21) in Paris, implementation requires emissions reductions of various greenhouse gases, including carbon dioxide (CO2), methane (CH4), and nitrous oxide (N2O). These gases not only warm the Earth, but also affect ozone abundances (WMO (World Meteorological Organization) 2014). Consequently, changes in these gas emissions and their simultaneous impacts on climate and the ozone layer become important considerations for future policy decisions.

The stratospheric ozone layer protects Earth’s surface from the most damaging ultraviolet (UV) radiation. It has been substantially depleted over the last 35 years due to halocarbon emissions, which are both potent greenhouse gases and ozone-depleting substances (ODSs). An undisputed environmental policy-making success story, the Montreal Protocol and its amendments will return ODSs to near historical levels (1955–1975, prior to significant depletion) by the latter half of this century and prevent significant health consequences of excessive UV exposure (Newman and McKenzie 2011, Chipperfield et al 2015). However, the evolution of stratospheric ozone beyond the middle of the 21st century, when ODS emissions are minimal and their atmospheric concentrations continue to decrease, will largely be determined by CO2, N2O, and CH4 (Oman et al 2010, Fleming et al 2011, Portmann et al 2012, Revell et al 2012, Eyring et al 2013, Stolarksi et al 2015, Iglesias-Suarez et al 2016). Under some future emission scenarios, extratropical ozone far exceeds historical levels, reducing surface UV below their historical levels (Hegglin and Shepherd 2009, Watanabe et al 2011, Bais et al 2015). N2O destroys stratospheric ozone by increasing reactive odd-nitrogen species. N2O is currently the largest ozone-depletion-potential (ODP)-weighted emission (Ravishankara et al 2009) and a potent
greenhouse gas; its potential mitigation via the Mongreal Protocol has been examined (Kanter et al 2013). The ODP of N2O changes with CO2 and CH4 abundances (Ravishankara et al 2009, Wang et al 2014, Revell et al 2015). The dominant effect of increased CH4 is increased ozone production in the lower stratosphere and troposphere down to the surface (Fleming et al 2011, Eyring et al 2013). Increased CO2 cools the stratosphere and consequently increases ozone particularly in the upper stratosphere (Fleming et al 2011). Increased CO2 is expected to accelerate the Brewer-Dobson Circulation (BDC), leading to stratospheric ozone decreases in the tropics and increases in the extratropics (Butchart 2014). While CH4 and N2O affect ozone predominantly via chemical reactions, CO2 affects ozone indirectly via temperature and dynamical changes. Although N2O, CH4, and CO2 all exacerbate climate change, they have different, and possibly non-linear (Meul et al 2015), influences on both total column and stratospheric column ozone.

The Montreal Protocol’s regulations to mitigate ODSs are expected to reduce their radiative forcing while increasing global stratospheric ozone back towards historical levels. Conversely, protecting the future ozone layer under evolving CO2, N2O, and CH4 emissions may involve the conundrum of weighing benefits for climate change against maintaining the ozone layer and surface UV radiation near historical levels.

**2. Methods**

To demonstrate the complexities in future ozone changes when N2O or CH4 concentrations are increased or reduced in climates with varying CO2, we ran simulations using the National Center for Atmospheric Research (NCAR) Whole Atmosphere Community Climate Model (WACCM) for the Representative Concentration Pathways (RCPs), RCP 2.6, 4.5, and 8.5. WACCM version 4 (Marsh et al 2013) is an atmospheric extension to NCAR’s Community Earth System Model (CESM) with a high model top (∼140 km) and fully interactive chemistry in the middle atmosphere, though tropospheric chemistry has a limited representation CH4 and carbon monoxide oxidation). The horizontal resolution is 1.9 degrees latitude by 2.5 degrees longitude with 66 vertical levels. The quasi-biennial oscillation is represented by nudging the tropical winds to match observed interannual variability.

A historical transient simulation from 1955–2005 was performed using time-evolving observed forcings (surface concentrations of radiatively active species, solar spectral irradiance, and volcanic sulfates). The ocean was initialized from a reference case and allowed to freely evolve. Future RCP transient simulations were run from 2006–2095 using projected concentrations of radiatively active species (Meinshausen et al 2011). From these transient simulations, we used the sea surface temperature (SST) and sea ice climatologies for both the historical (1955–1975) and the future (2075–2095) climates to force 20-year ‘time-slice’ experiments (following 10 years of spin-up) with varying concentrations of N2O, CH4, and CO2 (table 1) based on the RCP scenarios. Time-slice runs simulate ‘slices’ of time rather than the full transient response; the atmosphere responds to fixed climatological SSTs and forcings held at a constant value for a particular year. In runs where we modulate N2O or CH4 concentrations, we assume that the associated changes in SSTs/sea ice are not large (i.e., they are dominated by the changes in CO2).

For comparison with values in table 1, 2016 global concentrations of CO2, N2O, and CH4 are ∼402 ppm, ∼328 ppb, and ∼1840 ppb, respectively. Note that in the end-of-the-century RCP 8.5 climate, CO2 and CH4 have increased substantially (roughly a factor of three), while N2O has increased only by ∼50%, compared to the historical climate. Total chlorine (Cl\textsubscript{2}) is higher for all future climates compared to Cl\textsubscript{2} in the historical run because the long-lived chlorofluorocarbons persist. Cly concentrations, aerosols, and tropospheric emission precursor concentrations in the time-slice experiments are based on corresponding historical or RCP scenario values, but we performed an additional sensitivity run to test the role of Cl\textsubscript{2} in the RCP 8.5

**Table 1. Concentrations of CO2, N2O, CH4, and Cl\textsubscript{2} used in each time-slice simulation, and the climatology period for sea surface temperatures (SSTs)/sea ice. Concentrations are from either the year 1957 (historical) or 2095 (future runs) RCPs.**

| Experiment             | CO2 (ppm) | N2O (ppb) | CH4 (ppb) | Cl\textsubscript{2} (ppb) | SSTs/Ice climatology |
|------------------------|-----------|-----------|-----------|--------------------------|----------------------|
| 1957 historical        | 314       | 291       | 1211      | 0.73                     | 1955–1975            |
| RCP 2.6                |           |           |           |                          |                      |
| RCP26_N2O_85           |           |           |           |                          |                      |
| RCP26_CH4_85           |           |           |           |                          |                      |
| RCP 4.5                |           |           |           |                          |                      |
| RCP 8.5                |           |           |           |                          |                      |
| RCP85_N2O_26           |           |           |           |                          |                      |
| RCP85_CH4_26           |           |           |           |                          |                      |


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scenario (not shown). Solar spectral irradiances (SSI), specified from the Lean et al. (2005) model and used for solar heating and photolysis reactions, vary on daily timescales in the transient runs, but are set constant in the historical time-slice run to 1960 values and in the future time-slice runs to 2095 values. The SSI integrated over all wavelengths is the total spectral irradiance (TSI); for comparison, the TSI is \( \sim 1362 \text{ W m}^{-2} \) in 1960 and \( \sim 1361 \text{ W m}^{-2} \) in 2095. For solar and geomagnetic parameters (e.g., the F 10.7 cm flux, \( K_p, A_p \)), a constant value (the solar cycle average) was used in all time-slice runs.

We also consider changes in stratospheric ozone in coupled climate models with interactive chemistry from the Fifth Coupled Model Intercomparison Project (CMIP5) (Taylor et al. 2012). Stratospheric column ozone is determined by summing ozone above 200 hPa (more complex methods using a latitudinally-varying tropopause yielded similar results).

3. Results

In agreement with previous studies (Oman et al. 2010, Fleming et al. 2011, Portmann et al. 2012, Revell et al. 2012, Eyring et al. 2013, Stolarksi et al. 2015, Iglesias-Suarez et al. 2016), there is a broad range of possible changes to the stratospheric ozone layer by the end of the century (2075–2095) compared to the historical period (1955–1975) (shown in figure 1 for CMIP5 models with interactive chemistry). In an aggressive greenhouse gas mitigation scenario (RCP 2.6; blue shading), stratospheric ozone at most latitudes remains slightly below historical levels by the end of the 21st century. The \( \sim 5–10 \) Dobson Unit ozone depletion is due to slightly elevated \( N_2O \) and anthropogenic halocarbons remaining in the atmosphere.

However, under continued greenhouse gas emissions (RCP 8.5; red shading), stratospheric ozone remains depleted in the tropics but substantially exceeds historical levels in the extratropics. Lower tropical stratospheric ozone relative to the historical period is mostly due to the accelerated BDC and partly due to \( N_2O \)-related loss processes (Fleming et al. 2011, Eyring et al. 2013). The extratropical ozone increases are primarily associated with the cooling effects of \( CO_2 \) in the mid- to upper stratosphere, the increased advection of ozone-rich air via the strengthened BDC, and enhanced production due to increased \( CH_4 \) (Fleming et al. 2011). In a moderate emission scenario (RCP 4.5; green shading), stratospheric ozone remains depleted in the tropics, but extratropical ozone returns to near historical levels, due to a balance between increases caused by \( CO_2 \) and \( CH_4 \), and depletion caused by ODSs and \( N_2O \). The changes in stratospheric ozone from the historical period (1955–1975) to the end of the century (2075–2095) in our WACCM RCP 2.6, 4.5, and 8.5 transient runs (dashed lines) are largely consistent with the CMIP5 simulations, with the exception of weaker ozone recovery in the Southern Hemisphere extratropics and greater ozone recovery in the Northern Hemisphere extratropics (figure 1; dashed lines).

Time-slice experiments were forced using SST and sea ice climatologies from the transient simulations.

![Figure 1. Comparison of CMIP5 model changes in stratospheric ozone [DU] between the mean 2075–2095 period and the mean 1955–1975 period as a function of latitude, for RCP 2.6 (blue), RCP 4.5 (green), and RCP 8.5 (red). The multi-model mean is shown with the solid line. The change in stratospheric ozone in the WACCM transient runs is shown in dashed lines. The CMIP5 models with interactive chemistry included here are: CESM1-WACCM, GFDL-CM3, GISS-E2-H-p2, GISS-E2-H-p2, and MIROC-ESM-CHEM.](image-url)
and the concentrations of CO$_2$, N$_2$O, CH$_4$, and total Cl$_y$ shown in table 1. Here we focus on the RCP 2.6 and 8.5 scenarios, since the stratospheric ozone response to RCP 4.5 falls in between these extreme scenarios (figure 1). For the RCP 2.6 future time-slice run (figures 2(A), (C); bold black line), ozone is slightly below historical levels at most latitudes by the end of the 21st century in agreement with the transient simulation (figure 1). We can then examine what would happen if either CH$_4$ or N$_2$O were not aggressively mitigated. If CH$_4$ were increased to RCP 8.5 concentrations for 2095, column ozone increases at all latitudes due to large tropospheric ozone increases (figures 2(A), (C); green shading). If N$_2$O were increased to RCP 8.5 concentrations for 2095, ozone decreases at all latitudes due to stratospheric ozone depletion (figures 2(A), (C); purple shading). Note that the tropical ozone depletion in this scenario would potentially be larger than during the period of maximum historic ozone layer depletion (1985–2005, dotted line). A key point is that if the world were to achieve reductions of CO$_2$ and CH$_4$ concentrations to RCP 2.6 levels, N$_2$O mitigation would become important to avoid exacerbation of both climate change and ozone layer depletion. On the other hand, if CO$_2$ and N$_2$O were reduced to RCP 2.6 levels but CH$_4$ concentrations increased, we expect stratospheric ozone increases towards historical levels and large increases in global tropospheric ozone.

In contrast, modulating N$_2$O or CH$_4$ in a climate with high greenhouse gas concentrations has different consequences for the total and stratospheric column ozone. In the RCP 8.5 future time-slice run, stratospheric ozone remains depleted in the tropics but substantially exceeds historical levels in the extratropics (figure 2(B); bold black line). Large CH$_4$ increases in the RCP 8.5 scenario (and thus large increases in tropospheric ozone) leads to increases in the total column ozone above historical levels at all latitudes (figure 2(D); bold black line). Note that increases in column ozone due to enhanced CH$_4$ occurs more in the troposphere than in the stratosphere, as to be

![Figure 2.](image-url)
expected. In this climate, a reduction in N₂O concentrations would reduce anthropogenic climate forcing and slightly reduce stratospheric ozone depletion at low latitudes, but would increase column ozone beyond historical levels in the extratropical stratosphere and at all latitudes (figures 2(B), (D); purple shading). A reduction in CH₄ concentrations would also reduce anthropogenic climate forcing and tropospheric ozone, and thus decrease extratropical ozone levels towards historical levels, but could exacerbate tropical stratospheric ozone decreases (figures 2(B), (D); green shading).

There are many potential impacts of changes in tropospheric and stratospheric ozone on human health and the ecosystem. There are large uncertainties in tropospheric ozone due to uncertainties in tropospheric precursor emissions, but the changes simulated in WACCM fall within the CMIP5 multi-model spread (Eyring et al. 2013, Young et al. 2013). We assess the latitudinal changes in the UV index (UVI) at local noon using an approximation based on calculated total column ozone for cloud-free, aerosol-free, low-surface albedo conditions (Madronich 2007):

\[ \text{UVI} \sim 12.5 \mu_\odot^{1.42} \left( \frac{\Omega}{300} \right)^{-1.23}, \]

where \( \mu_\odot \) is the cosine of the solar zenith angle at solar noon on the 15th of each month, and \( \Omega \) is the total column ozone. Note that the noon UVI ranges from zero at the poles to \(~12\) at the equator, and varies substantially with latitude and season, with the highest values in the extratropical summer hemisphere. Thus a 20% change in UVI at 45° latitude represents absolute increases in the UVI of \(~2\) in the summer but only \(~0.5\) in the winter.

Figure 3 shows the percent change in the UVI (based on total, not stratospheric, column ozone) compared to the historical period as a function of month for (A, C) the RCP 2.6 future climate and (B, D) the RCP 8.5 future climate for N₂O concentrations fixed at 344 ppbv (the 2095 RCP 2.6 value) or 428 ppbv (the 2095 RCP 8.5 value). Figures 3(A) and (D) represent the ‘control’ RCP 2.6 and 8.5 climates, respectively, while figures 3(B) and (C) correspond to the lower and upper boundary for potential future changes in ozone in figure 2. Clearly, changes in UVI by the latter half of the century depend more on future CO₂ concentrations than on N₂O concentrations. In the RCP 2.6 future climate, UVI remains only slightly larger than historical levels, except in the Southern Hemisphere (SH) spring where remaining halocarbons preferentially deplete polar ozone and increase surface UV exposure (highlighting the need to phase out these compounds). Conversely, in the RCP 8.5 future climate, UVI decreases everywhere relative to historical levels, and especially in the extratropics. These results are in general agreement with previous studies, though some studies find weak increases in UV exposure in the tropics (Bais et al. 2011, Watanabe et al. 2011). It is clear that lower concentrations of N₂O would reduce UVI towards historical levels (i.e., zero percent change) in the RCP 2.6 climate (figures 3(A) versus (C)) but further away from historical levels in the RCP 8.5 climate (figures 3(B) versus (D)). In the RCP 8.5 future climate with N₂O mitigated to RCP 2.6 levels (figure 3(B)), the UVI is reduced up to \(~20\%\) in the Northern Hemisphere (NH) extratropics. The UVI is actually closer to historical levels when N₂O is increased (figure 3(D)).

4. Discussion and conclusions

The negative health effects of enhanced UV exposure, namely increased incidence of skin cancers and cataracts (Williamson et al. 2014, Lucas et al. 2015), could continue to impact the Southern Hemisphere as well as much of the globe in an RCP 2.6 future climate, particularly if N₂O concentrations increase. This enhanced UV exposure could be especially detrimental in the tropics, where UV exposure is already high. In contrast, consequences of reduced UV exposure are not as clear and need to be better understood, particularly if greenhouse gas emissions continue to increase (Hegglin and Shepherd 2009, Bais et al. 2015).

For humans, one primary concern of lower UV is a reduction in vitamin D synthesis, which is associated with increased risks of osteoporosis, rickets, certain types of cancer, cardiovascular disease, multiple sclerosis, and rheumatoid arthritis (Autier et al. 2014). Some recent studies are unable to find evidence of vitamin D supplementation reducing risk of these diseases (Lucas et al. 2015). Extratropical UV reduction may be particularly important because it occurs in regions where vitamin D deficiencies are already prevalent (Correa et al. 2013, Lucas et al. 2015). Reductions in UV could also affect terrestrial and aquatic ecosystems, as well as biogeochemical and carbon cycles (Ballare et al. 2011, Williamson et al. 2014, Bornman et al. 2015, Erickson III et al. 2015, Hader et al. 2015).

The UV changes presented here are based on a number of assumptions, such as cloud-free and unpolluted conditions. Potential future increases in cloud cover and reductions in surface reflectivity over the Arctic would cause additional decreases in UV at northern high-latitudes, while projected decreases in aerosols may counter these surface UV reductions; however, these effects are uncertain (Watanabe et al. 2011, Correa et al. 2013, Bais et al. 2015). In addition, there are uncertainties in the tropospheric emission scenarios, and limitations of the model simulations themselves (such as limited tropospheric chemistry). Volcanic eruptions are not considered in our future simulations but can have substantial transient effects on stratospheric ozone and associated UV. Warmer temperatures and changing precipitation patterns may also cause behavioral changes, such as...
altering the time people spend outdoors and thus, possibly, their overall UV exposure (Lucas et al. 2015).

Uncertainties aside, it is reasonable to assert that a comprehensive scientific foundation for future policy decisions to protect the ozone layer will include (a) the collective impacts of expected CO$_2$, CH$_4$, and N$_2$O emissions on ozone, (b) the intricacies of balancing continued tropical stratospheric ozone decreases with potential extratropical ozone increases, and (c) the amount of total ozone increases or decreases that are deemed a concern for human and ecosystem health. Because of the large global warming influence of CO$_2$ and its enhancing effects on ozone, reducing CO$_2$ emissions may be the ideal policy for both reducing climate change and returning ozone to near historical levels at all latitudes by the latter half of the century. Reducing CH$_4$ emissions would have similar co-benefits. In contrast, reducing N$_2$O emissions increases stratospheric ozone, which is beneficial in the present-day climate and for certain future scenarios where ozone depletion persists; but decreasing N$_2$O may actually raise ozone above historical levels in the latter half of the century if CO$_2$ and CH$_4$ emissions continue unabated as in the RCP 8.5 scenario.

Reducing greenhouse gas emissions to at least the RCP 4.5 trajectory is required to obtain the goals set forth by COP 21, which aim to limit global temperature changes to less than 2°C above pre-industrial levels (Collins et al. 2013). If achieved, this would also return stratospheric ozone and UV to near historical levels globally by 2100. If we aim to limit global temperature changes to less than 1.5°C, CO$_2$, CH$_4$, and N$_2$O all need to be reduced to near RCP 2.6 levels, which would also benefit the ozone layer. If the world cannot achieve a reduction in CO$_2$ emissions to at least RCP 4.5 levels, the range of policy options to protect the ozone layer would necessarily broaden. One possibility includes allowing increased N$_2$O emissions in order to maintain historical ozone and UV levels in the extratropics. For this option, the benefits of reducing ozone levels in the extratropics would need to be carefully weighed against slightly larger depletion in the tropics and the greenhouse gas effects of N$_2$O. This again highlights the importance of meeting COP21
goals to minimize future climate change and maintain surface UV at historical levels for human health and ecosystems.

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