

Assessment of an updated polar stratospheric cloud parameterisation for the UK Earth System Model (UKESM1.1) within the UK Met Office Unified Model (v13.9) using CALIOP and MLS observations

Isabelle Sangha^{1,2}, Nathan L. Abraham³, Andrew Orr¹, Hua Lu¹, Michael C. Pitts⁴, Lamont R. Poole⁵, and Michael Weimer⁶

¹British Antarctic Survey, High Cross, Madingley Road, Cambridge, CB3 0ET, UK

²Department of Chemistry, University of Cambridge, Cambridge, CB2 1EW, UK

³NCAS-Climate, University of Cambridge, CB2 1EW, UK

⁴NASA Langley Research Centre, 2 Langley Blvd, Hampton, VA 23681, USA

⁵Analytical Services & Materials, Inc., 107 Research Dr, Hampton, Virginia 23666, USA

⁶Institute of Environmental Physics, University of Bremen, Bremen, Germany

Correspondence: Isabelle Sangha (is568@cam.ac.uk)

Abstract. Accurately representing polar stratospheric clouds (PSCs) in global Chemistry-Climate Models and Earth System Models is important as they play a key role in springtime ozone depletion in polar regions by activating both chlorine and bromine species through heterogeneous reactions and denitrifying the stratosphere. Here, we present and evaluate an updated PSC parameterisation scheme implemented in the UK Earth System Model (UKESM1.1). The scheme includes the kinetic formation of nitric acid trihydrate (NAT) particles, the formation of supercooled ternary solution (STS) droplets assuming thermodynamic equilibrium, and accounts for ice and sulphate aerosols. To evaluate the new scheme, we compare modelled PSC production with satellite observations from the Cloud-Aerosol and Lidar with Orthogonal Polarization (CALIOP) instrument and model concentrations of gas-phase nitric acid, water vapour, and ozone concentrations with observations from the Aura Microwave Limb Sounder (MLS) instrument for the extended 2008 Antarctic and 2009/2010 Arctic winters. In comparison with the current, simpler scheme, the updated parameterisation increases both the range of PSC types that form and the seasonal variability, resulting in better agreement with CALIOP observations. However, there is still a gap in the model's ability to simulate ice PSCs with the full range of optical properties observed by CALIOP. The new scheme also slows the growth of NAT particles and enables nitric acid partitioning between gas, liquid, and solid phases, leading to improved agreement with gas-phase nitric acid MLS observations, although a substantial low-bias is still present in all model simulations. However, there is comparatively little impact on stratospheric ozone, with the exception near the edge of the polar vortex where the new scheme improves comparisons with MLS observations.

1 Introduction

Stratospheric ozone absorbs incoming solar radiation and its depletion leads to cooling of the polar stratosphere, influencing atmospheric circulation and surface climate (Randel and Wu, 1999; Polvani et al., 2011). Polar stratospheric clouds (PSCs) play a critical role in driving ozone depletion and shaping polar stratospheric chemistry. Heterogeneous reactions on the surfaces of PSCs convert reservoir chlorine and bromine species into active species during the polar winter, when the absence of solar radiation allows stratospheric temperatures to drop below the thresholds for PSC formation (Solomon et al., 1986; Tritscher et al., 2021). Once sunlight returns in spring, these active species participate in catalytic cycles that efficiently destroy ozone (O_3), resulting in rapid O_3 depletion (Molina and Molina, 1987). The threshold temperature for PSC formation depends on the composition of the PSC particles. Crystalline nitric acid trihydrate (NAT) are solid particles consisting of nitric acid (HNO_3) and water (H_2O) with formation temperatures of below $T_{NAT} \sim 195$ K under normal stratospheric ambient conditions at 20 km. Supercooled ternary solution (STS) droplets grow through the uptake of HNO_3 on the background sulphate aerosols as temperatures decrease. Under normal stratospheric ambient conditions (ie., 50 hPa with $HNO_3 \sim 10$ ppb and $H_2O \sim 5$ ppm), the typical temperature that HNO_3 rapidly condenses onto background sulphate aerosols is 191 K (although HNO_3 still condenses slowly on the background aerosols at higher temperatures if vapour pressures are high enough). Water ice particles form at the coldest temperatures of below $T_{ice} \sim 188$ K (Koop et al., 2000; Engel et al., 2013).

Particle composition also influences PSC surface reactivity (Carslaw et al., 1997). For many heterogeneous reactions occurring below T_{NAT} , liquid droplets (e.g., STS) are comparable to, or more effective than, solid NAT and ice particles in activating chlorine based on reactivity per unit surface area (Ravishankara and Hanson, 1996). Peter and Grooß (2012) showed that chlorine activation rates of NAT could only compete with the chlorine activation rates of STS in high number density ‘mother clouds’ that form downstream of ice clouds (Fueglistaler et al., 2002). Reaction rates on liquid binary H_2SO_4 - H_2O aerosol particles also reach substantial values before NAT formation, indicating that chlorine activation is mainly driven by liquid heterogeneous chemistry. However, large solid ice and NAT particles sediment from the stratosphere into the upper troposphere, causing stratospheric denitrification and dehydration. Removal of reactive nitrogen limits the amount of nitrogen available to deactivate chlorine and prolongs periods of O_3 depletion (Toon et al., 1986). This effect is especially important in the Arctic, where temperatures are comparably higher and denitrification can result in greater ozone loss (Chipperfield and Pyle, 1998). In Antarctica, chlorine and bromine activation, combined with an increase in stratospheric chlorine loading due to anthropogenic release of chlorofluorocarbons, has resulted in severe springtime O_3 depletion in recent decades, forming a region of anomalously low total column ozone values referred to as the Antarctic ozone hole (Cicerone et al., 1974; Farman et al., 1985). This has significantly affected Southern hemisphere circulation and climate (Orr et al., 2008; Previdi and Polvani, 2014; Polvani et al., 2018).

To accurately simulate stratospheric ozone depletion, PSC parameterisation schemes in global Chemistry-Climate Models (CCMs) and Earth System Models (ESMs) must represent PSC formation and associated ozone-loss chemistry. However, the complexity of the PSC schemes in these models varies. For example, the SOCOL (SOLar Climate Ozone Links) model uses a relatively simplistic approach that assumes thermodynamic equilibrium between the PSC particles and gas phases,

instantaneously forming PSCs when saturation vapour pressure is reached (Steiner et al., 2021). This requires assumptions about the particle size distribution, either number density or radius. By contrast, models such as EMAC (ECHAM5/MESSy Atmospheric chemistry model), ICON-ART (ICOsahedral Nonhydrostatic modelling framework and its Aerosol and Reactive Trace gases extension), and SD-WACCM-4.0/CARMA (Community Aerosol and Radiation Model for Atmospheres coupled to the Whole Atmosphere Community Climate Model with Specified Dynamics) use more complex methods to capture deviations from equilibrium conditions by simulating the growth and evaporation of particles, enabling a wider range of particle sizes to be represented (Kirner et al., 2011; Zhu et al., 2015; Weimer et al., 2021).

The UK Earth System Model (UKESM1.1) uses a relatively simple approach for PSC formation, accounting only for NAT, sulphate aerosols, and ice (Dennison et al., 2019). NAT particles are nucleated assuming thermodynamic equilibrium and are assumed to have a fixed radius of $1.0\ \mu\text{m}$ (Hanson and Mauersberger, 1988). Sulphate aerosols are included as a prescribed surface area density (SAD), and there is no mechanism for the formation of STS droplets (Sellar et al., 2019; Archibald et al., 2020). Ice particles are formed using the cloud microphysics scheme (Wilson and Ballard, 1999), but are assumed to have a fixed radius of $10\ \mu\text{m}$ when calculating the SAD for heterogeneous chemistry. While the exact mechanisms of NAT nucleation are still not fully understood, the growth of NAT particles occurs over hours to days (Voigt et al., 2005), and are not in equilibrium when they grow to larger sizes (Carslaw et al., 2002). Therefore, the inability of this scheme to simulate NAT outside of equilibrium conditions and to form STS particles limits the capability of UKESM1.1 to represent all PSC particle types and their evolution over the PSC season, hindering its ability to accurately simulate stratospheric ozone depletion, including projections of future ozone recovery and associated climate impacts (Austin et al., 2003; Eyring et al., 2006, 2013; Zeng et al., 2022).

PSCs are observed with satellite measurements (Pitts et al., 2013; Spang et al., 2018), ground-based lidar (Snels et al., 2019) and airborne and balloon-borne in situ observations (Fahey et al., 2001; Voigt et al., 2003). Ground-based lidars provide measurements with high temporal resolution and enhanced signal-to-noise ratios of many variables, but their effectiveness can be limited by tropospheric cloudiness (Tesche et al., 2021). Airborne and balloon-borne in situ measurements offer detailed observations of trace gases and aerosols, including measurements of size and number concentration, but have limited temporal and spatial coverage. Satellite-based measurements such as the Cloud-Aerosol Lidar with Orthogonal Polarization (CALIOP) onboard the Cloud-Aerosol Lidar and Infrared Pathfinder Satellite Observations (CALIPSO), which operated from June 2006 until August 2023, use the depolarization ratio to infer the particle shape and concentration, enabling distinction between liquid and solid particles (Pitts et al., 2013). Given its excellent temporal and spatial coverage and ability to differentiate PSC types, CALIOP provides a robust benchmarking dataset to evaluate PSC parameterisations. Additionally, the Microwave Limb Sounder (MLS) instrument on board the Aura Satellite flew in formation with CALIPSO in the A-Train constellation until 2018 (Stephens et al., 2002), providing nearly coincident measurements of atmospheric gas variables, such as HNO_3 , H_2O , and O_3 concentrations by observing passive thermal emissions on the millimetre and sub-millimetre wave length scale (Waters et al., 2006).

In this study, we address the limitations in the representation of NAT and STS particles in UKESM1.1 by replacing its current, simple representation of these particles with the more sophisticated representation employed in ICON-ART (Weimer et al.,

2021). This approach kinetically forms NAT particles and thermodynamically forms STS particles, making it more interactive and physically based. We evaluate the performance of this new scheme relative to the current scheme by comparing modelled PSCs with CALIOP PSC observations during the 2008 Antarctic and 2009/2010 Arctic winters (Pitts et al., 2011, 2018). Comparing over both Arctic and Antarctic PSC seasons enables evaluation of the new parameterisation under the different dynamical conditions of the northern and southern polar vortices. For the Antarctic winter, we additionally investigate how the updated scheme affects the temporal evolution of PSCs and gas-phase HNO_3 and O_3 concentrations with a comparison against MLS measurements. Through these analyses, we assess how the new scheme alters UKESM1.1 PSC composition diversity and seasonal progression, while identifying remaining limitations and possible future model developments.

2 Models, Data, and Methods

2.1 Model Descriptions

2.1.1 UKESM1.1

UKESM version 1 (UKESM1.1) is a state-of-the-art Earth system model that is built around the global climate model HadGEM3-GC3.1, which is composed of the Unified Model (UM) atmosphere model, NEMO ocean model, and CICE sea ice model (Sellar et al., 2019; Mulcahy et al., 2023). Here we use version 13.9 of the UM, with a resolution of 1.875° longitude by 1.25° latitude and 85 vertical levels extending to 85 km (N96L85). Aerosols and trace gas chemistry are simulated using the stratospheric-tropospheric (Strat-Trop) configuration of the United Kingdom Chemistry and Aerosol (UKCA) model (Archibald et al., 2020), which includes v8.0 of the two-moment modal aerosol microphysics module GLOMAP-mode (Mann et al., 2010; Dhomse et al., 2014; Mulcahy et al., 2020). Strat-Trop employs 84 species and accounts for 299 reactions, including additional heterogeneous bromine reactions and updated uptake coefficients on aerosols and solid-phase PSCs implemented in UKESM1.1 (for more details, see Dennison et al., 2019).

In the original PSC parametrisation used by UKESM1.1, NAT is calculated following Chipperfield (1999), assuming thermodynamic equilibrium with gas-phase HNO_3 and H_2O . NAT is formed every time step when the partial pressure of HNO_3 , p_{HNO_3} , exceeds the saturation vapour pressure of HNO_3 over NAT, $p_{\text{HNO}_3}^{\text{NAT}}$, calculated using the relationship defined in Hanson and Mauersberger (1988):

$$\log_{10}(p_{\text{HNO}_3}^{\text{NAT}}) = a_{\text{NAT}}(T) \times \log_{10}(p_{\text{H}_2\text{O}}) + b_{\text{NAT}}(T), \quad (1)$$

where T is the temperature, $p_{\text{H}_2\text{O}}$ is the partial pressure of H_2O , and

$$a_{\text{NAT}}(T) = -2.7836 - 0.00088(T), \quad (2)$$

$$b_{\text{NAT}}(T) = 38.9855 - \frac{11397.0}{T} + 0.009179(T). \quad (3)$$

This is only valid for the temperature range $180 \text{ K} < T < 200 \text{ K}$. For temperatures below 180 K, $p_{\text{HNO}_3}^{\text{NAT}}$ is calculated with $T = 180 \text{ K}$. All NAT particles are assumed to have a fixed-value radius, r_{NAT} , of $1.0 \mu\text{m}$. Denitrification is prescribed based on

Chipperfield (1999) with two different sedimentation velocities: one for pure NAT (0.46 mm s^{-1}) and one for mixed ice/NAT (17 mm s^{-1}) (Morgenstern et al., 2009).

Ice abundance is taken from the UM microphysics scheme which calculates the mass mixing ratio of ice using a single-moment scheme (Wilson and Ballard, 1999). The mass mixing ratio is then converted to a SAD by assuming a radius, r_{ice} , of 10.0 μm . STS formation is not explicitly included; instead, the effect of liquid droplets are accounted for with a prescribed SAD climatology from 1850–2014 prepared for the CMIP6 model intercomparison (Sellar et al., 2019; Archibald et al., 2020). Consequently, heterogeneous reaction rates are therefore calculated based on uptake on ice, NAT, and sulphate aerosols.

2.1.2 Updated PSC parameterisation

In the updated PSC scheme, the nucleation, growth, and evaporation of NAT particles is parameterised based on the analytic expression described in Carslaw et al. (2002). This is the same scheme implemented in ICON-ART, which was developed based on version 2.1 of ICON-ART (Schröter et al., 2018) and included in the published version 2025.04 (Hoshyaripour et al., 2025). Here, the time-dependent diffusive growth of the radius of a NAT particle, r_{NAT} , is estimated by:

$$\frac{dr_{\text{NAT}}}{dt} = \frac{G}{r_{\text{NAT}}}, \quad (4)$$

where t is time, and G is the surface growth factor computed using:

$$G = \frac{D_{\text{HNO}_3}^* M_{\text{NAT}}}{\rho_{\text{NAT}} R T} (p_{\text{HNO}_3} - p_{\text{HNO}_3}^{\text{NAT}}), \quad (5)$$

where M_{NAT} is the molar mass of NAT, ρ_{NAT} its density, and R the universal gas constant. $D_{\text{HNO}_3}^*$ denotes the diffusion coefficient of HNO_3 in air, accounting for non-continuum mass transfer effects of particles comparable in size to the mean free path and is computed using:

$$D_{\text{HNO}_3}^* = \frac{D_{\text{HNO}_3}}{1 + 4D_{\text{HNO}_3}/(\bar{c}_{\text{HNO}_3} r_{\text{NAT}})}, \quad (6)$$

where D_{HNO_3} is the diffusion coefficient of HNO_3 in air, and $\bar{c}_{\text{HNO}_3}$ is the mean molecular speed of HNO_3 .

To determine the new r_{NAT} at each time step as a result of growth or evaporation, Eq. (4) is integrated over time, t , which gives:

$$r_{\text{NAT}}^2(t + \Delta t) = r_{\text{NAT}}^2(t) + 2G\Delta t, \quad (7)$$

where Δt is the chemistry time step (equal to 1 hr in UKESM1.1).

To implement the kinetic NAT formulation in UKESM1.1, size bins with fixed radii are defined according to van den Broek et al. (2004). Nine bins, each with an average radius $r_{\text{NAT},b}$ and number density limit, $n_{\text{NAT},\text{lim},b}$, where b is the bin index, span radii from 0.1 μm to 22.5 μm , based on the observed range and distribution (Fahey et al., 2001). This results in a total maximum NAT number density limit, $n_{\text{NAT},\text{max}} = \sum_b n_{\text{NAT},\text{lim},b}$. In this study a range of values for $n_{\text{NAT},\text{max}}$ are used to assess the sensitivity to NAT number density. Regardless of the choice of value for $n_{\text{NAT},\text{max}}$, NAT initially nucleates in the smallest

bin (r_{NAT} of $0.1 \mu\text{m}$) with a number density of $1.5 \times 10^{-5} \text{ cm}^{-3}$ when $p_{\text{HNO}_3} > p_{\text{HNO}_3}^{\text{NAT}}$ (see Eq. 1) with a 3 K temperature threshold applied in line with similar NAT parameterisations to account for the supersaturation required for NAT formation (Groß et al., 2002; Kirner et al., 2011). The condition thus becomes:

$$p_{\text{HNO}_3} > p_{\text{HNO}_3}^{\text{NAT}}(T + 3\text{K}) \quad (8)$$

The change in particle radius within each size bin is calculated sequentially from the smallest to the largest bin using Eq. (4) and the bin-specific growth factor G_b following Eq. (5). Changes in radius are immediately converted to changes in number density, $n_{\text{NAT},b}$, by conserving mass, ensuring all particles in a bin have the same size. If $n_{\text{NAT},b}$ exceeds its limit $n_{\text{NAT},\text{lim},b}$, excess mass is transferred to the next larger bin. Evaporation occurs when $p_{\text{HNO}_3} < p_{\text{HNO}_3}^{\text{NAT}}$, making G_b negative in Eq. (5). At $T > 220 \text{ K}$, particles fully evaporate. Each size bin is treated as a separate tracer and NAT sedimentation is determined interactively, with the fall velocity calculated for each bin based on Stokes' law.

In the updated PSC scheme, STS droplets are formed by calculating the volume density of the sulphate aerosols or STS, V_{STS} , based on the uptake of HNO_3 and H_2O on aqueous sulphuric acid aerosols using the thermodynamic scheme documented in Carslaw et al. (1995). At temperatures above which HNO_3 uptake is negligible, V_{STS} corresponds to binary sulphate aerosols; as temperatures decrease, HNO_3 uptake increases and V_{STS} corresponds to ternary solution droplets. V_{STS} is then converted to a SAD, S_{STS} , for heterogeneous chemistry using the relationship described in Grainger et al. (1995):

$$S_{\text{STS}} = a_{\text{SV}}(V_{\text{STS}})^{b_{\text{SV}}}, \quad (9)$$

where $a_{\text{SV}} = 6.068$ and $b_{\text{SV}} = 0.671$ based on the empirical parameter fitting (Hervig and Deshler, 1998). The effective radius of the STS particles, r_{STS} , used when calculating the optical properties, is then determined by:

$$r_{\text{STS}} = \frac{3V_{\text{STS}}}{S_{\text{STS}}}. \quad (10)$$

To implement the new STS scheme in UKESM1.1, the concentration of aqueous sulphuric acid aerosols is provided each timestep from the interactive GLOMAP aerosol microphysical module (Mann et al., 2010; Dhomse et al., 2014; Mulcahy et al., 2020) and is used to calculate V_{STS} . Note that the STS routine is called before the NAT routine in UKCA. This means that when temperatures fall below T_{STS} , HNO_3 uptake on sulphate aerosols is calculated first. The formation, growth, or evaporation of NAT is then calculated based on the remaining gas-phase HNO_3 .

The new PSC scheme still computes ice PSCs using the UM microphysics scheme, but the conversion from mass mixing ratio to SAD has been modified. Previously, ice mass mixing ratio was converted to a SAD assuming a fixed radius ($r_{\text{ice}} = 10 \mu\text{m}$). With this assumption, the ice number density, n_{ice} , doesn't exceed 0.01 cm^{-3} , and the resulting SAD remains below $2.5 \mu\text{m}^2\text{cm}^{-3}$ for the periods analysed in this study. Although observations of ice PSC size distributions are limited, these values are relatively low compared to other PSC schemes (Zhu et al., 2017; Tritscher et al., 2019; Weimer et al., 2021). Therefore, following the approach of ICON-ART, in the updated PSC scheme the ice SAD is instead calculated assuming a fixed value for n_{ice} (Weimer, 2019). However, although ICON-ART uses $n_{\text{ice}} = 0.25 \text{ cm}^{-3}$, in this study we examine a range of n_{ice} values to assess the sensitivity of the results to this. For example, using higher n_{ice} values produce smaller ice particles with larger surface area densities.

175 With a prescribed n_{ice} , stratospheric r_{ice} is calculated as follows:

$$r_{\text{ice}} = \frac{1}{2} \sqrt[3]{\frac{m_{\text{ice}}}{130}}, \quad (11)$$

where m_{ice} is the mass of ice particles, defined as

$$m_{\text{ice}} = \frac{\rho W_{\text{ice}}}{n_{\text{ice}}}, \quad (12)$$

with W_{ice} the water mass mixing ratio of cloud ice and ρ the air density (Doms et al., 2011).

180 2.2 Observations

CALIOP was a dual-wavelength, polarization-sensitive lidar that observed profiles of the parallel and perpendicular components of the backscatter coefficients ($\beta_{\perp}$ and $\beta_{\parallel}$, respectively) at 532 nm and total backscatter at 1064 nm from mid-June 2006–August 2023 (Pitts et al., 2018). In this study, we use the CALIPSO lidar level 2 PSC mask version 2.0 (v2) which has a horizontal resolution of 5 km and vertical resolution of 180 m, and considers only nighttime measurements. This dataset was
 185 derived using the CALIOP v2 PSC Classification Algorithm (Pitts et al., 2018). For PSC detection, only the 532 nm backscatter coefficient measurements are used. The Modern-Era Retrospective analysis for Research and Applications (Gelaro et al., 2017) molecular number density is used in the theoretical calculation described in Hostetler et al. (2006) to determine the molecular backscatter, β_{mol} , which is then used to compute the 532 nm attenuated scattering ratio, R_{532} :

$$R_{532} = \frac{\beta_{\parallel} + \beta_{\perp}}{\beta_{\text{mol}}}. \quad (13)$$

190 PSCs are identified as statistical outliers in $\beta_{\perp}$ or R_{532} . Daily threshold values of $\beta_{\perp}$ and R_{532} are defined as the median plus one median absolute deviation ($\beta_{\perp, \text{threshold}}$ and $R_{\text{threshold}}$, respectively). A PSC occurrence is then defined as exceeding either background threshold by at least the measurement uncertainty, σ :

$$\beta_{\perp} > \beta_{\perp, \text{threshold}} + \sigma(\beta_{\perp}), \quad (14)$$

$$R_{532} > R_{\text{threshold}} + \sigma(R_{532}). \quad (15)$$

195 where $\sigma(\beta_{\perp})$ and $\sigma(R_{532})$ are computed using the noise-scale factor approach of Liu et al. (2006) as described for CALIOP by Hostetler et al. (2006). PSCs with $\beta_{\perp}$ exceeding the background threshold are assumed to contain non-spherical (i.e., solid) particles. R_{532} is used to further classify solid-containing PSCs as either liquid-NAT mixtures (NATmix), enhanced NAT mixtures with $r_{\text{NAT}} \lesssim 3 \mu\text{m}$ and NAT volume density $\gtrsim 1.0 \mu\text{m}^3\text{cm}^{-3}$ (enhNAT), ice, or high-number-density ice that forms in mountain waves (wave ice). The R_{532} boundary separating enhNAT and NATmix from ice, R_{NATice} , is calculated dynamically
 200 from coincident cloud-free MLS gas-phase HNO_3 and H_2O data and theoretical R_{532} values for fully developed STS. PSCs below $\beta_{\perp, \text{threshold}}$, but above $R_{\text{threshold}}$ are classified as STS. Classification boundaries are summarised in Table 1, though it is important to note that they are not all-inclusive, i.e., enhNAT might not capture all NAT mixture PSCs that are heterogeneously nucleated downstream of wave ice PSCs.

Table 1. Boundaries in R_{532} and $\beta_{\perp}$ values used to classify PSC observations using CALIOP v2 PSC classification. Threshold $\beta_{\perp,\text{threshold}}$ and $R_{\text{threshold}}$ are the median plus one median absolute deviation, and $\sigma(\beta_{\perp})$ and $\sigma(R_{532})$ are the measurement uncertainties of $\beta_{\perp}$ and R_{532} respectively. The boundary between ice and both NATmix and enhNAT, R_{NATice} , is calculated dynamically.

Classification	$\beta_{\perp}$ boundaries	R_{532} boundaries
Ice: mixture of liquid sulphate aerosols/STS and solid ice		
	$\beta_{\perp} > \beta_{\perp,\text{threshold}} + \sigma(\beta_{\perp})$	$R_{532} > R_{\text{NATice}}$
Wave ice: high number density ice particles presumably induced by mountain waves		
	$\beta_{\perp} > \beta_{\perp,\text{threshold}} + \sigma(\beta_{\perp})$	$R_{532} > 50$
NATmix: mixture of liquid sulphate aerosols/STS and solid NAT		
	$\beta_{\perp} > \beta_{\perp,\text{threshold}} + \sigma(\beta_{\perp})$	$R_{532} < R_{\text{NATice}}$
enhNAT: NAT mixtures with $r_{\text{NAT}} < 3\mu\text{m}$ and NAT volume density $> 0.1\mu\text{m}^3\text{cm}^{-3}$		
	$\beta_{\perp} > 2.5 \times 10^{-5} \text{ km}^{-2}\text{sr}^{-1}$	$2 < R_{532} < R_{\text{NATice}}$
STS: Super-cooled ternary solution particles		
	$\beta_{\perp} < \beta_{\perp,\text{threshold}} + \sigma(\beta_{\perp})$	$R_{532} > R_{\text{threshold}} + \sigma(R_{532})$

The Aura MLS passively measures microwave thermal emissions from the ground to 90 km to retrieve profiles of about 20 trace gases and cloud properties. It takes measurements at wavelengths in the millimetre to sub-millimetre range from the limb of Earth’s atmosphere from 82°S to 82°N every 25 s (Waters et al., 2006). In this study, model outputs are compared with MLS level 2 data (Livesey et al., 2006) for HNO_3 , H_2O , and O_3 . The MLS HNO_3 product is derived from the 240-GHz retrievals at pressures ≥ 22 hPa and 190-GHz retrievals for lower pressures, with a vertical resolution of HNO_3 data is 3–4 km in the region of interest. The MLS H_2O product is derived from the 190-GHz retrievals with a vertical resolution of 1.3–3.6 km in the region of interest. The MLS O_3 product uses the 240-GHz radiances and has a vertical resolution of 2.5 km.

2.3 Methodology

2.3.1 Comparing to observations

To compare PSCs simulated by UKESM1.1 with those observed by CALIOP, the particle size distributions from the model are used to calculate optical parameters as they would be observed by the satellite, i.e. $\beta_{\perp}$ and $\beta_{\parallel}$. The same classification algorithm used for CALIOP optical properties can then be applied to the model outputs. Comparing directly with the satellite optical properties is preferable to using the CALIOP SAD product derived from these properties, as these estimates are highly uncertain for NAT and ice (Pitts et al., 2018). These optical properties are derived from the particle size distribution using Mie theory and T-matrix calculations implemented in the scattering codes within the Zurich Optical Microphysical Model (ZOMM)

(Mishchenko et al., 1996; Hoyle et al., 2013; Engel et al., 2013). When applying this code to UKESM1.1 data, ice and NAT particles are represented as prolate spheroids with aspect ratios of 0.9, and a refractive index of 1.31 for ice and 1.48 for NAT (Engel et al., 2013; Tritscher et al., 2019).

To implement the Mie theory and T-matrix calculations, the ZOMM scattering code requires size-resolved number densities of STS, NAT, and ice as inputs. However, since UKESM1.1 outputs do not include the size distributions of all three PSC particle types, assumptions are necessary. Thus, to determine the size-resolved number densities, it is assumed that the total number density of PSC particles/sulphate aerosols is 10 cm^{-3} . The size-resolved number densities of NAT is determined inherently by the number density in each size bin for simulations using the new parameterisation. For the original parameterisation, the distribution is assumed to be monodisperse with a fixed radius of $1.0 \text{ } \mu\text{m}$. Sulphate aerosols and STS droplets are assumed to follow a lognormal distribution with a standard deviation of 1.6 and mean radius of r_{STS} estimated from Eq. (10). Ice is assumed to be a monodisperse distribution determined using either the prescribed radius or number density, depending on the parameterisation. To fully compare with CALIOP data, background $\beta_{\perp}$ and R_{532} values were added to the simulated optical properties, and noise, σ , for both $\beta_{\perp}$ and R_{532} is calculated using the CALIOP Noise Factor (CNF) defined in Tritscher et al. (2019):

$$\sigma(\beta_{\perp}) = \text{CNF} \times \sqrt{\beta_{\perp}}, \quad (16)$$

$$\sigma(R_{532}) = |R_{532}| \times \sqrt{\frac{\sigma^2(\beta_{\perp}) + \sigma^2(\beta_{\parallel})}{(\beta_{\perp} + \beta_{\parallel})^2} + (0.03)^2}, \quad (17)$$

with CNF taken as 0.00102 (corresponding to a CALIOP horizontal averaging scale of 135 km, the best case for detection). The last term in Eq. (17) accounts for the 3% relative uncertainty in molecular backscatter. To estimate $\beta_{\perp}$ and R_{532} with noise, a random value from a normal distribution with an expectation of the simulated $\beta_{\perp}$ or R_{532} value and a standard deviation of either $\sigma(\beta_{\perp})$ or $\sigma(R_{532})$, respectively, is chosen. As with the CALIOP observations, the optical properties of each model grid-cell must exceed $\beta_{\perp, \text{threshold}}$ or $R_{532, \text{threshold}}$ based on Eqns. 14 and 15, respectively, with σ calculated for each grid-cell using Eqns. 16 and 17.

To compare UKESM1.1-simulated HNO_3 , H_2O , and O_3 concentrations with MLS satellite observations, the model outputs for HNO_3 (solid, liquid, and gas phase), H_2O , and O_3 were interpolated to the same pressure levels as the MLS data. The MLS observations were then binned and averaged to match the UKESM1.1 horizontal grid, and the UKESM1.1 outputs were masked to only included grid-cells where MLS data were available. In the original PSC parameterisation, solid-phase HNO_3 in NAT is combined with gas-phase HNO_3 at the end of the chemical time step for transport. Similarly, in the new scheme the liquid-phase HNO_3 in STS is added back to the gas phase at the end of the chemical time step. Therefore, to determine the concentration of gas-phase HNO_3 concentration during PSC formation and heterogeneous reaction rate calculations, the mixing ratio of HNO_3 in solid phase or liquid phase (for old and new scheme, respectively) was subtracted from the gas-phase HNO_3 output offline. Finally, an area-weighted zonal mean was calculated across three 10-degree latitude bands for both the MLS and UKESM1.1 data.

Table 2. Average radius ($r_{\text{NAT},b}$) and number density limit ($n_{\text{NAT},\text{lim},b}$) of NAT particles in each size bin (b). The order of magnitude of the $n_{\text{NAT},\text{lim},b}$ values depends on the sensitivity simulation and is set to either 10^{-3} , 10^{-4} , or 10^{-5} .

Bin (b)	1	2	3	4	5	6	7	8	9
$r_{\text{NAT},b}$ (μm)	0.1	0.6	1.5	4.0	7.5	10.5	14.0	18.0	22.5
$n_{\text{NAT},\text{lim},b}$ ($\times 10^{-3}$, 10^{-4} , or 10^{-5} cm^{-3})	3.29	3.29	3.29	3.29	3.29	3.29	1.64	1.64	1.64

2.3.2 Simulations

UKESM1.1 simulations were run for five years from January 2007–December 2011, although we especially focus on the extended 2008 Antarctic and 2009/2010 Arctic winter seasons. For the Antarctic 2008 winter, this included the period July 10–18 2008, which represents typical winter stratospheric conditions with temperatures consistently below the PSC formation threshold (Pitts et al., 2018). By contrast, the Arctic 2009/2010 winter featured unusually cold stratospheric temperatures, i.e., up to 9 K below the climatological mean from early December 2009 to mid-January 2010, bringing regions below both T_{NAT} and T_{ice} . This was followed by a major sudden stratospheric warming in late January driven by strong planetary wave-1 forcing (Dörnbrack et al., 2012), resulting in four distinct periods of CALIOP PSC optical regimes (Pitts et al., 2011).

All UKESM1.1 simulations use the Coupled Model Intercomparison Project Phase 6 (CMIP6) historical emission data spanning from 1750–2014 (Feng et al., 2020), Reynold’s sea-surface temperature and sea-ice data (Reynolds et al., 2002), and are nudged to the fifth generation ECMWF atmospheric reanalysis of the global climate (ERA5-reanalysis) data (Hersbach et al., 2020). The optical comparison with CALIOP is examined for both winter seasons, while temporal PSC evolution and gas-phase HNO_3 , H_2O , and O_3 analysis focus on the Antarctic 2008 winter due to more consistent PSC formation.

Twelve simulations were performed using the updated PSC parameterisation with varying values defined for $n_{\text{NAT},\text{max}}$ and n_{ice} to conduct a sensitivity analysis assessing the effect of the prescribed parameters on PSC formation. In the implementation of the same kinetic NAT parameterisation used here, Kirner et al. (2011) and Weimer (2019) use $n_{\text{NAT},\text{lim},b}$ values of the order of 10^{-5} cm^{-3} based on measurements by Fahey et al. (2001). Here, we additionally investigate the effect of increasing the $n_{\text{NAT},\text{lim},b}$ values to the order of 10^{-4} and 10^{-3} cm^{-3} to cover the range of number densities observed (Fahey et al., 2001; Biele et al., 2001) and used to constrain CALIOP observations (Pitts et al., 2011). The size bins and different orders of magnitudes used for the $n_{\text{NAT},\text{lim},b}$ are summarized in Table 2 and result in values of the total maximum number density limit, $n_{\text{NAT},\text{max}} = \sum_b n_{\text{NAT},\text{lim},b}$, of 2.47×10^{-2} cm^{-3} , 2.47×10^{-3} cm^{-3} , and 2.47×10^{-4} cm^{-3} (for $n_{\text{NAT},\text{lim},b}$ values of the order of 10^{-3} , 10^{-4} , and 10^{-5} cm^{-3} , respectively). For ice PSCs, where observations are limited, n_{ice} was set to 0.001, 0.01, 0.1, and 0.25 cm^{-3} , spanning ranges used in other models (Tritscher et al., 2019; Steiner et al., 2021; Weimer et al., 2021; Zhu et al., 2017), as well as the ice number densities considered by Pitts et al. (2018). A summary of the simulations is presented in Table 3. To avoid redundancy, we focus on analysing results from RUN7, RUN8, RUN11, and RUN12 as they best match the optical properties from CALIOP observations while illustrating sensitivity to $n_{\text{NAT},\text{max}}$ and n_{ice} . These simulations have been renamed based on whether $n_{\text{NAT},\text{max}}$ or n_{ice} is relatively high (H) or low (L) with respect to the four selected simulations. For example,

Table 3. Parameter settings for total maximum NAT number density limit ($n_{\text{NAT,max}}$) and ice number density (n_{ice}) in cm^{-3} for each of the sensitivity simulations. RUN7, RUN8, RUN11, and RUN12 are selected for analysis in this study and are highlighted in bold and renamed according to whether $n_{\text{NAT,max}}$ or n_{ice} is relatively high (H) or low (L) with respect to the four chosen simulations.

	$n_{\text{ice}} (\text{cm}^{-3})$				
		0.001	0.01	0.1	0.25
$n_{\text{NAT,max}} (\text{cm}^{-3})$	2.47×10^{-4}	RUN1	RUN2	RUN3	RUN4
	2.47×10^{-3}	RUN5	RUN6	RUN7 (RUN_{LL})	RUN8 (RUN_{LH})
	2.47×10^{-2}	RUN9	RUN10	RUN11 (RUN_{HL})	RUN12 (RUN_{HH})

RUN8 has $n_{\text{NAT,max}} = 2.47 \times 10^{-3} \text{ cm}^{-3}$ which is relatively low and $n_{\text{ice}} = 0.25 \text{ cm}^{-3}$ which is relatively high, so it has been renamed RUN_{LH}. A control run (CONTROL) using the original PSC parameterisation was also performed for comparison.

280 Finally, the occurrence frequency and areal coverage of PSCs simulated by RUN12 (RUN_{HH}) and CONTROL are analysed. This requires defining SAD thresholds for NAT, ice, and STS to identify an occurrence of each PSC type at an individual grid-cell. SAD thresholds of $0.1 \mu\text{m}^2\text{cm}^{-3}$ and $1.0 \mu\text{m}^2\text{cm}^{-3}$ are used to identify ice and STS containing grid-cells, respectively. For NAT, the threshold is defined as the theoretical NAT SAD reached when particles grow to $r_{\text{NAT}} > 1.0 \mu\text{m}$ in the new parameterisation, corresponding to a particle size comparable to NAT PSCs in the CONTROL. This avoids counting grid-cells
285 with very small NAT SADs and results in a NAT threshold of $0.015 \mu\text{m}^2\text{cm}^{-3}$ for RUN_{HH}. Due to the known issues with the CALIOP SAD product mentioned above, observations are not included in this comparison.

3 Results

3.1 Comparison with optical properties from CALIOP

To assess the results from the simulations under conditions with temperatures consistently below the PSC formation thresholds,
290 Fig. 1 compares the simulated optical properties from CONTROL and the four selected sensitivity runs (RUN_{LL}, RUN_{LH}, RUN_{HL}, and RUN_{HH}) with CALIOP observations over the Antarctic from July 10–18 2008. Note that while the values of the counts of observations for CALIOP and grid-cells for the simulations in the 2D histograms cannot be directly compared, the relative abundance can be. During this period, CALIOP observations show distinct peaks in the NATmix and ice regions, with few points corresponding to the other three composition types (enhNAT, STS, and wave ice). In contrast, CONTROL shows
295 a distinct peak in the enhNAT region with a tail in the NATmix and no points classifying as ice. The CONTROL also has a narrower R_{532} range than CALIOP, resulting in fewer PSC types represented. (Note that since CONTROL includes no STS particles, the STS optical signal likely arises from NAT and ice particles.) The sensitivity runs generally capture the observed spread of the R_{532} values better than CONTROL, which results in comparatively more points being classified as STS. The sensitivity simulations with higher n_{ice} (RUN_{HH} and RUN_{LH}) have optical properties extending into the ice region, however
300 they don't cover the same range of optical properties in the ice region as the CALIOP observations. While CALIOP has a

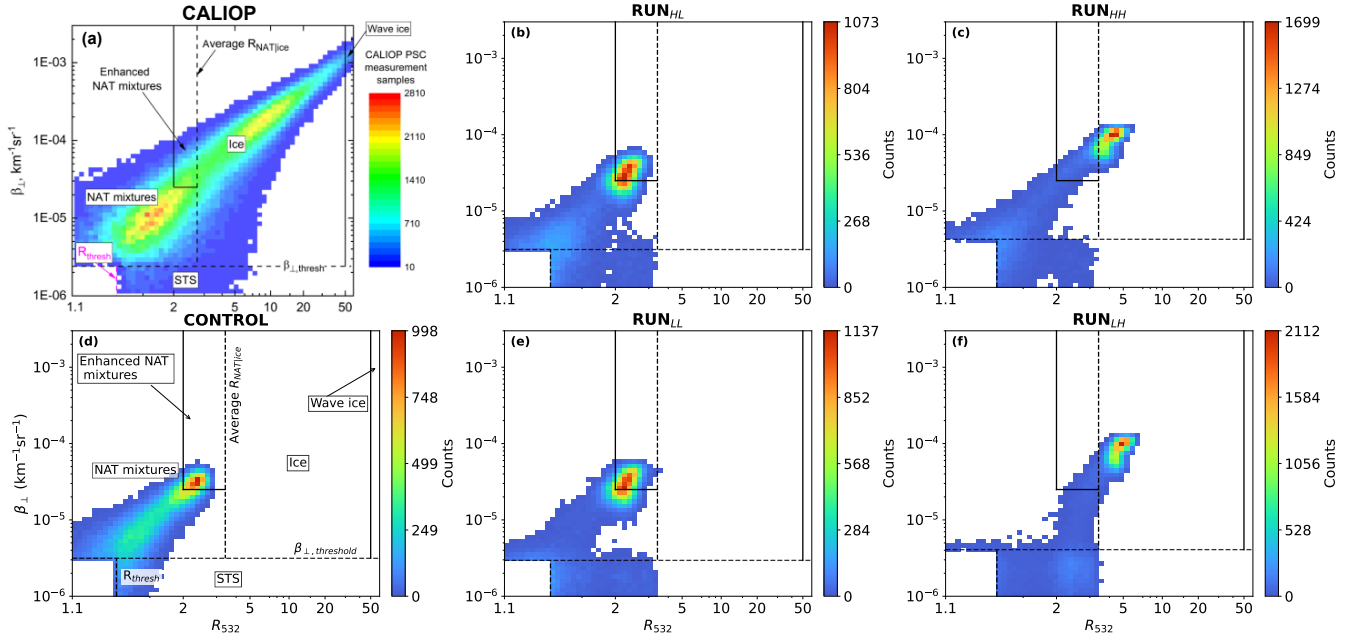


Figure 1. 2D histogram of optical properties from CALIOP (a), RUN_{HL} (b), RUN_{HH} (c), CONTROL (d), RUN_{LL} (e), and RUN_{LH} (f) over the Antarctic for the period July 10–18 2008, latitude range 65°S – 75°S , and potential temperature range 475–525 K. The optical properties are plotted in the CALIOP v2 classification coordinate system of R_{532} vs $\beta_{\perp}$, with regions of enhanced NAT mixtures (enhNAT), liquid-NAT mixtures (NATmix), STS, ice, and wave ice indicated. The shading indicates the number of CALIOP observations (a), or the number of model grid points (b–f). Dashed lines indicate the mean $\beta_{\perp}$ and R_{532} thresholds (including measurement uncertainty), as well as the mean R_{NATice} boundary, calculated for CALIOP and each simulation separately. Solid lines indicate the boundaries separating wave ice and ice and enhNAT and NATmix. The regions labelled for CONTROL (d) are the same for all the sensitivity simulations (b, c, e, f).

substantial number of observations with R_{532} values ranging from 5 to 20, the maximum R_{532} value achieved by the sensitivity simulations is around 6. Additionally, none of the sensitivity simulations have points with high enough R_{532} values to be classified as wave ice. Instead, they exhibit a relative maxima in either the enhNAT or ice regions, depending on the choice of n_{ice} . The sensitivity simulations with larger $n_{NAT,max}$ values (RUN_{HL} and RUN_{HH}) have more points in the NATmix region, whereas the lower $n_{NAT,max}$ sensitivity simulations (RUN_{LL} and RUN_{LH}) have slightly more points in the STS region. Optical properties from all 12 sensitivity runs demonstrate a relatively strong dependence on $n_{NAT,max}$ and n_{ice} , and are shown in Fig. S1 in the Supplementary Material.

To further evaluate the new parameterisation, Fig. 2 compares simulated optical properties from CONTROL and the four selected sensitivity runs (RUN_{LL} , RUN_{LH} , RUN_{HL} , and RUN_{HH}) with CALIOP observations during the 2009/2010 Arctic winter, with the season separated into four distinct periods. Over the first period, from 15–30 Dec 2009, CALIOP is dominated by NATmix with no ice observed. Over the next two periods from 31 Dec 2009–14 Jan 2010 and 15–21 Jan 2010, CALIOP shows an increase in NATmix, as well as the occurrence of enhNAT and STS particles, and wave ice. In the final period from

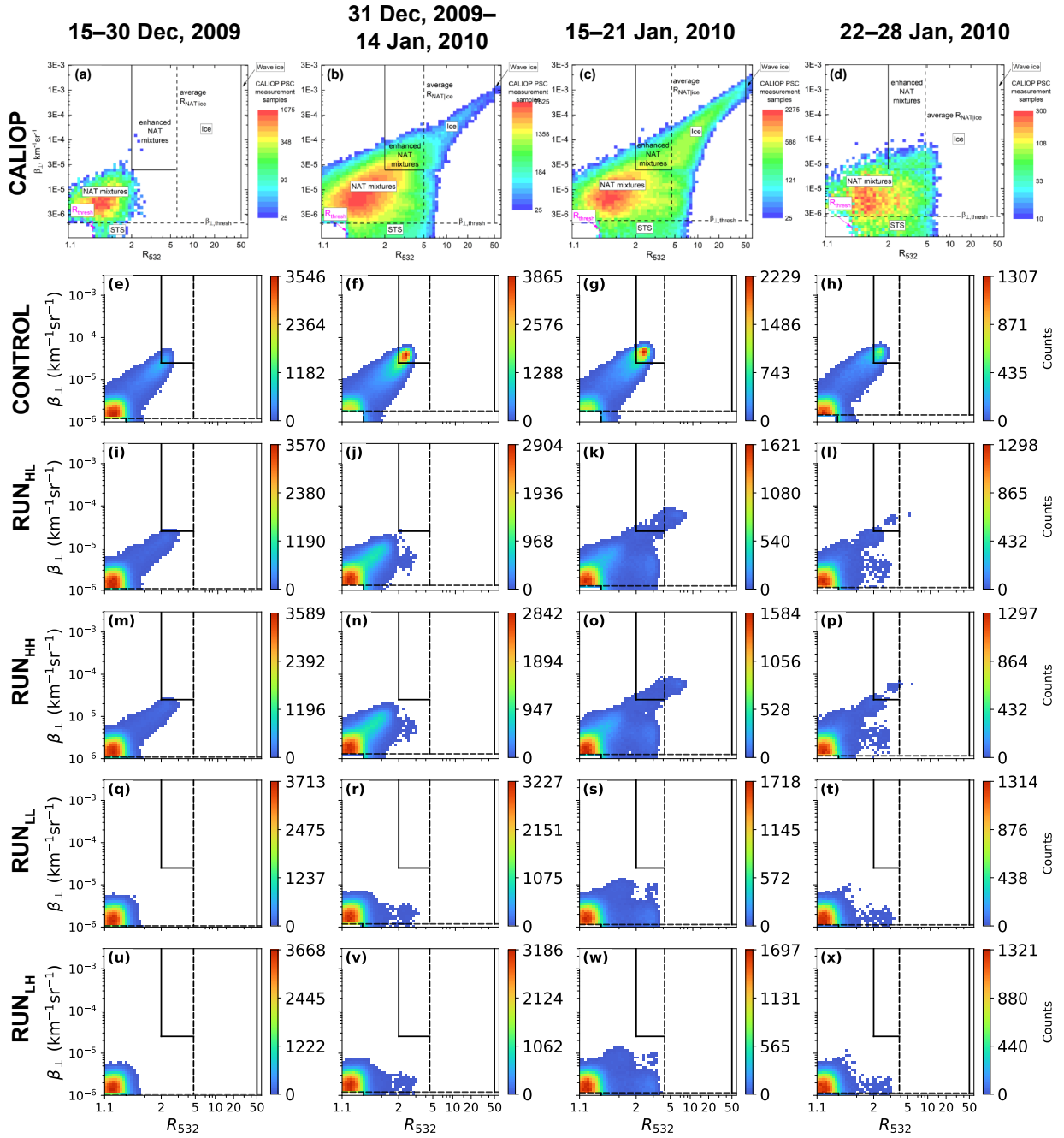


Figure 2. 2D histogram of optical properties from CALIOP (a–d), CONTROL (e–h), RUN_{HL} (i–l), RUN_{HH} (m–p), RUN_{LL} (q–t), and RUN_{LH} (u–x) over the Arctic for the 2009/2010 winter season, latitude range 65°N–75°N, and height range 15–30 km. The results are separated into four distinct periods, comprising 15–30 Dec 2009 (first column), 31 Dec 2009–14 Jan 2010 (second column), 15–21 Jan 2010 (third column), and 22–28 Jan 2010 (fourth column). The results are plotted with the same classification algorithm, shading, boundaries, and regions as Fig. 1.

22–28 Jan 2010, the vortex warmed above T_{ice} and widespread PSC observations from CALIOP declined appreciably. Most remaining PSCs are NATmix with only a few STS and ice particles. In contrast, CONTROL shows PSCs initially dominated by
 315 NATmix in the period from 15–30 Dec 2009, then dominated by enhNAT for the remainder of the season with a tail extending into the NATmix, exhibiting little variation throughout the season aside from the number of points classified as PSCs, i.e., little resemblance to CALIOP and almost no response to the changing atmospheric conditions.

Unlike CONTROL, the higher $n_{NAT,max}$ sensitivity runs (RUN_{HH} and RUN_{HL}) show broadly similar variations in PSC types over the Arctic winter, though not as pronounced as CALIOP observations (Fig. 2). During the first period (15–30 Dec 2009),
 320 RUN_{HL} and RUN_{HH} classify most PSCs as NATmix, in agreement with CALIOP. While during the second period (31 Dec 2009–14 Jan 2010), both these runs have a higher density of points in the NATmix region and extend into the STS region, again in agreement with CALIOP observations, but missing the points classifying as enhNAT and ice that are observed by CALIOP. However, during the third period (15–21 Jan 2010), both these runs spread into the ice and enhNAT region and further into the STS region, resulting in closer resemblance to CALIOP. Although, as with the Antarctic case, none of the points reach high
 325 enough values of R_{532} to be classified as wave ice and the extent of the points in the ice region is limited compared to CALIOP. During the fourth period (22–28 Jan 2010), both these runs show a decrease in NATmix, enhNAT, and STS, again similar to CALIOP. In contrast, RUN_{LL} and RUN_{LH} produce fewer PSCs that fall in the optical property classification range compared to RUN_{HL} and RUN_{HH} . However, while RUN_{LL} and RUN_{LH} show a similar temporal evolution of maximum R_{532} values over the season (i.e., an increase, followed by a slight decrease) that RUN_{HH} and RUN_{HL} (and CALIOP) also show, the equivalent
 330 evolution of maximum $\beta_{\perp}$ values is not as evident. This results in no points in the enhNAT region or ice region over the entire season in RUN_{LL} and RUN_{LH} .

Figures 1 and 2 show that increasing $n_{NAT,max}$ (i.e., comparing RUN_{LL} and RUN_{LH} to RUN_{HL} and RUN_{HH} ; see Table 3) shifts more points from the region below both the $\beta_{\perp}$ and R_{532} thresholds into the regions classified as NATmix, enhNAT, and ice. In contrast, increasing n_{ice} (i.e., comparing RUN_{LL} and RUN_{HL} to RUN_{LH} and RUN_{HH}) results in more points being
 335 classified as enhNAT or ice. The sensitivity to ice number density is more pronounced for the Antarctic case (Fig. 1), whereas sensitivity to NAT number density is more pronounced for the Arctic case (Fig. 2).

3.2 Temporal evolution of modelled PSCs

Figures 3 and 4 show the average PSC occurrence frequency over Antarctica at 20 km for NAT, ice, and STS for every month of the extended winter season over the entire 5-year simulation period of 2007–2011 for RUN_{HH} and CONTROL, respectively.
 340 RUN_{HH} was chosen as it shows the largest spread of optical properties compared with CALIOP (Fig. 1). Here, RUN_{HH} has high NAT occurrence frequency across Antarctica from June to August (values reaching 1), with lower frequencies in May and September (reaching 0.2). High occurrence frequency of ice (reaching 0.9) also occur during July and August, and are mainly concentrated over the centre of Antarctica. In contrast, STS only occurs from June to August in RUN_{HH} , particularly over Western Antarctica and the Antarctic Peninsula (i.e., near the edge of the stratospheric polar vortex), with much lower
 345 occurrence frequencies (reaching 0.2) - although this STS pattern is consistent with CALIOP observations (Pitts et al., 2018, Fig. 19 therein). CONTROL shows broadly similar patterns in both NAT and ice occurrence frequency as RUN_{HH} . The main

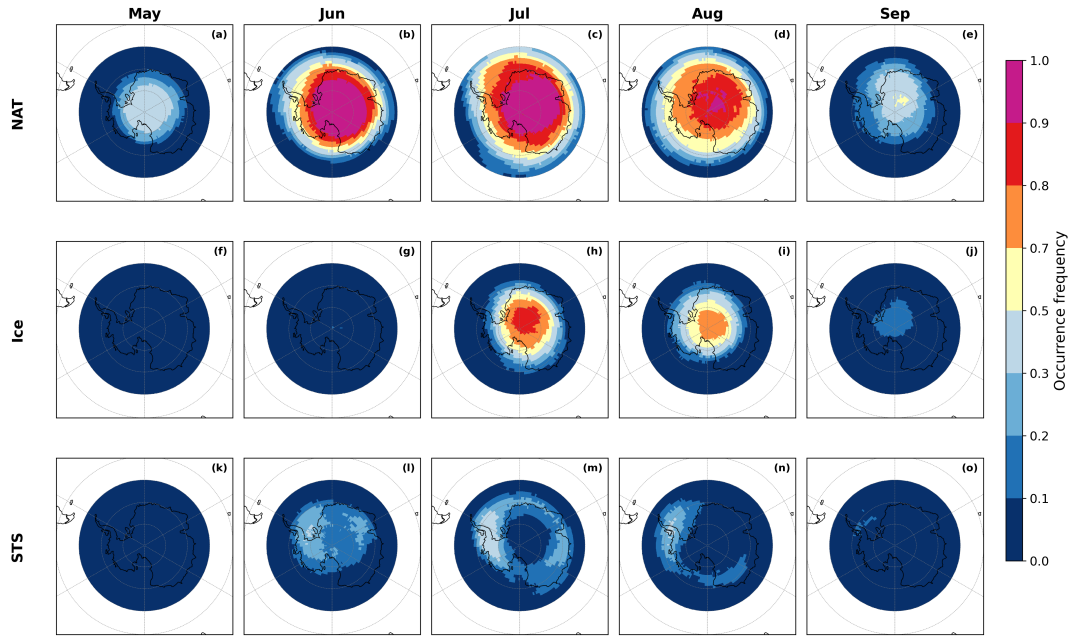


Figure 3. Monthly average PSC occurrence frequency over the Antarctic of NAT (a-e), ice (f-j) and STS (k-o) for May-September (left to right) over the 5-year period from 2007 to 2011 at an altitude of 20 km for RUN_{HH}. Occurrence frequency is defined as the number of days with PSC occurrence over the 5-year period for each month, divided by the total number of days over the 5-year period for each month.

differences are higher NAT frequency in May and September, reaching 0.7, and slightly lower ice frequencies in July and August. As CONTROL does not form STS, these results are not included in our comparison. The other sensitivity simulations show similar behaviour to RUN_{HH} (not shown). These results are further supported by the seasonal evolution of Antarctic PSC areal coverage for the extended 2008 winter season (see Fig. S2 in the Supplementary Material). Both CONTROL and RUN_{HH} show broadly similar areal coverage for NAT and ice PSCs, although there are slight differences in timing and vertical distribution. STS only forms in RUN_{HH} and starts having notable areal coverage later in the season and at a lower altitude than NAT. This results in both simulations showing broadly similar total PSC areal coverage over the entire winter season, although CONTROL has larger extent at higher altitude and RUN_{HH} at lower altitude.

To assess the seasonality and magnitude of SAD for the PSCs simulated by RUN_{HH} and CONTROL, Fig. 5 compares the temporal evolution of this quantity for NAT, ice, and liquid droplets (STS and sulphate aerosols) over the extended Antarctic 2008 winter. The smallest contribution to SAD for RUN_{HH} from the three PSC types is from NAT particles, which reach a maximum of around $0.36 \mu\text{m}^2\text{cm}^{-3}$ in mid-June from around 16 to 22 km (Fig. 5a), despite NAT showing the largest spatial coverage (Fig. 3). In contrast, the opposite occurs for liquid droplets and ice in RUN_{HH}, which have much larger SAD (Fig. 5b and c) despite smaller spatial coverage (Fig. 3). Additionally, the peak in SAD for ice particles occurs slightly after that for NAT and at a slightly lower altitude range, while the occurrence of liquid droplets occurs at even lower altitude but with no distinct peak. Compared to RUN_{HH}, the SAD for NAT particles simulated by CONTROL is up to an order of magnitude

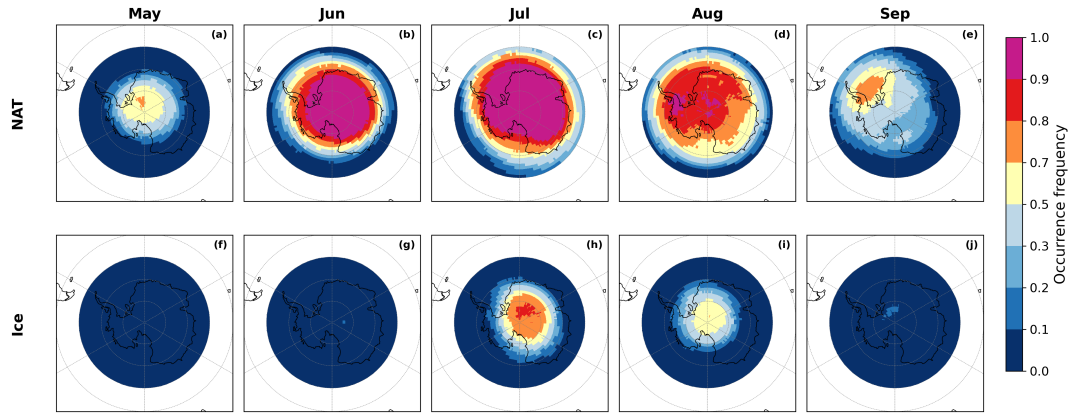


Figure 4. As in Fig. 3, but for CONTROL. Note that STS is not shown for CONTROL, as this is not simulated by the original PSC parameterisation.

higher (Fig. 5d). Differences in the distribution of SAD values for NAT particles simulated by CONTROL and RUN_{HH} across all grid-cells confirms this (Fig. S3 in the Supplementary Material), showing that CONTROL produces a narrow distribution of
 365 SAD for NAT particles concentrated at $1\text{--}10\ \mu\text{m}^2\ \text{cm}^{-3}$, while RUN_{HH} produces a broader distribution shifted to lower values of $0.01\text{--}1\ \mu\text{m}^2\ \text{cm}^{-3}$. The opposite is apparent for ice and liquid droplets (Fig. 5e and f), where CONTROL consistently has lower SAD than RUN_{HH}. Note that results for the other sensitivity simulations were broadly similar to RUN_{HH} (not shown), but with varying SAD magnitudes (see also Fig. S3).

3.3 Comparison of HNO₃ and H₂O from MLS

370 To further assess the simulations, Fig. 6 compares gas-phase HNO₃ at 68 hPa from CONTROL and the four selected sensitivity runs (RUN_{LL}, RUN_{LH}, RUN_{HL}, and RUN_{HH}) with MLS measurements over the extended 2008 Antarctic winter. Over the latitude band $60^\circ\text{S}\text{--}70^\circ\text{S}$ (i.e., near the edge of the stratospheric polar vortex), MLS shows a gradual drop in gas-phase HNO₃ concentration from about 8 ppbv in June to 4 ppbv by August (Fig. 6a). By contrast, at $70^\circ\text{S}\text{--}80^\circ\text{S}$ (i.e., within the polar vortex), gas-phase HNO₃ concentration drops sharply in June from 8–10 ppbv to ~ 2 ppbv and remains near this level until November
 375 (Fig. 6b), reflecting removal of HNO₃ from the gas-phase to form PSCs. While both the CONTROL and sensitivity runs have slight gas-phase HNO₃ depletion, they consistently underestimate gas-phase HNO₃ concentrations, especially in May before significant PSC formation. This likely reflects the known bias in low total reactive nitrogen (NO_y) in UKESM1.1 (Archibald et al., 2020). During the period of gas-phase HNO₃ depletion due to PSC formation, starting in mid-June, the underestimate of gas-phase HNO₃ concentration is most pronounced in CONTROL, with the sensitivity runs showing slightly higher gas-phase
 380 HNO₃ concentrations than CONTROL resulting in closer agreement to the MLS observations. A similar analysis at 100 hPa is presented in the Supplementary Material (Fig. S4), showing consistent results with Fig. 6. Additionally, the same analysis is presented in the Supplementary Material for the extended 2009/2010 Arctic winter at 68 hPa (Fig. S5). Here, there is not

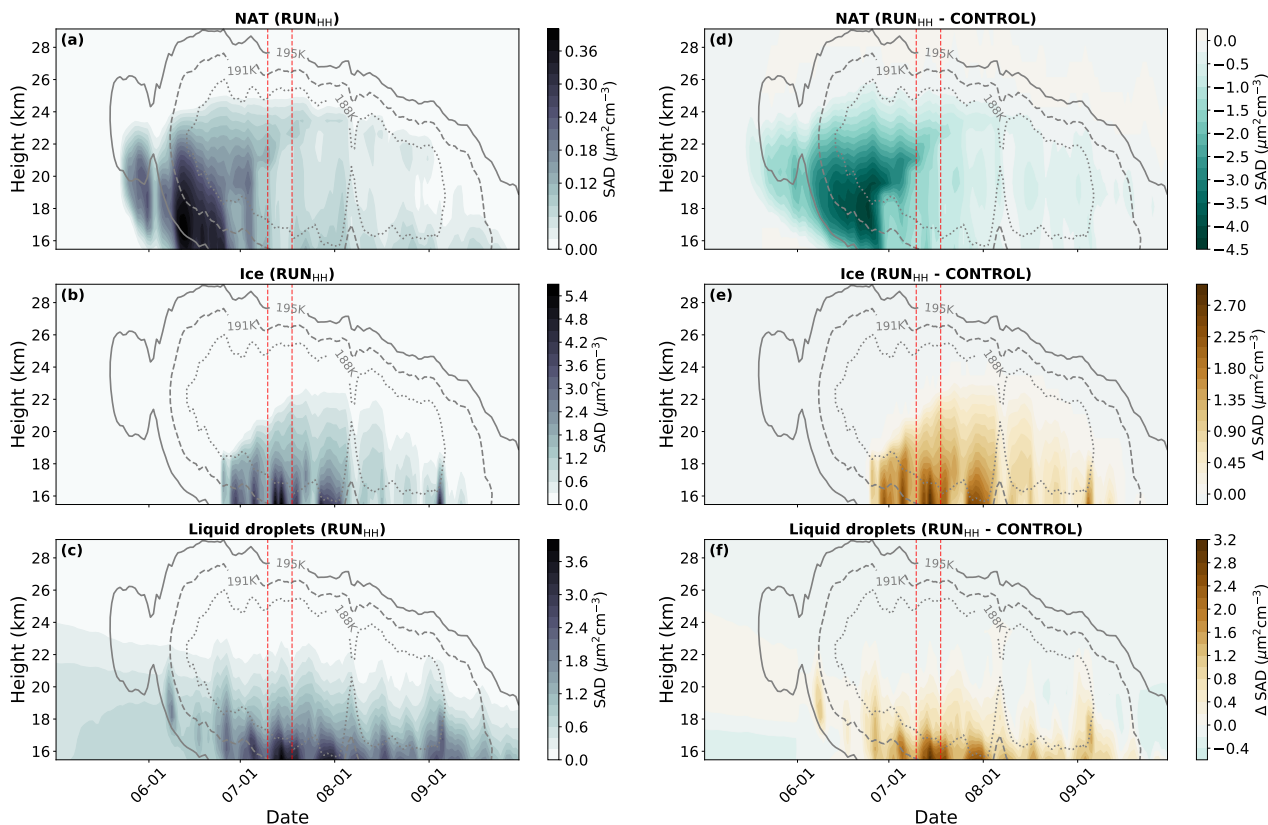


Figure 5. Time-height plot of zonal mean SAD (shading; $\mu\text{m}^2 \text{cm}^{-3}$) for NAT (a), ice (b) and liquid droplets (STS and sulphate aerosols, c) for RUN_{HH} over the Antarctic for the extended 2008 winter season (May to October), averaged across the latitude band 65°S - 90°S . Also shown are the differences between RUN_{HH} and CONTROL for NAT (d), ice (e), and liquid droplets (f). Grey contour lines indicate T_{NAT} (solid), T_{STS} (dashed), and T_{ice} (dotted). The red dashed vertical lines indicate the period July 10–18 2008 examined in Fig. 1. Note that the colour bar for all panels corresponds to different scales.

the same level of removal of gas-phase HNO_3 , however the same low-bias in the models compared to MLS at the start of the Antarctic winter season is present throughout the entire Arctic winter season.

385 The CONTROL and the sensitivity runs treat the removal of HNO_3 from the gas-phase differently: CONTROL converts all removed gas-phase HNO_3 to solid phase (NAT), whereas the sensitivity runs partition it between solid-phase and liquid-phase (STS). Figure 7 illustrates these differences for the extended 2008 Antarctic winter season by comparing the solid- and liquid-phase HNO_3 at 68 hPa from CONTROL and RUN_{HH} . The results show that RUN_{HH} starts partitioning HNO_3 from the gas-phase to solid-phase at roughly the same time as CONTROL (early June), but that the amounts of solid-phase HNO_3 simulated by CONTROL quickly surpasses RUN_{HH} . For example, over the latitude band 70°S – 80°S the amount of solid-phase HNO_3 simulated by CONTROL peaks in mid-June before declining in late June–early July (Fig. 7b). Conversely, RUN_{HH} has a less-pronounced peak of solid-phase HNO_3 in mid-June over 70°S – 80°S , followed by an increase in liquid-phase HNO_3 as

390

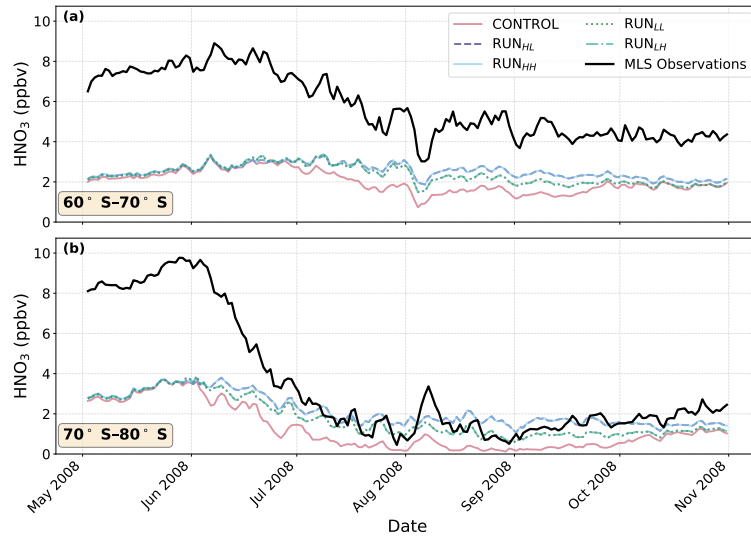


Figure 6. Zonal mean volume mixing ratio of gas-phase HNO₃ (ppbv) at 68 hPa for MLS (black line), CONTROL (pink line) and RUN_{LL}, RUN_{LH}, RUN_{HL}, and RUN_{HH} (purple, blue, green, and teal lines, respectively) over the Antarctic for the extended 2008 winter season (May to October), averaged across the latitude bands 60°S–70°S (a), and 70°S–80°S (b). Note that there is almost complete overlap of RUN_{HH} and RUN_{HL}, and RUN_{LL} and RUN_{LH}, resulting in only RUN_{HH} and RUN_{LL} being visible in the plot.

STS forms. At 70°S–80°S, the combined solid- and liquid-phase HNO₃ simulated by RUN_{HH} becomes comparable to solid-phase HNO₃ simulated by CONTROL by July and exceeds CONTROL during August, September, and October. These shifts in the amounts of solid- and liquid-phase HNO₃ throughout the winter season are consistent with the changes in SAD (Fig. 5). A similar analysis at 100 hPa is shown in the Supplementary Material (Fig. S6), which broadly agrees with Fig. 7.

Stratospheric gas-phase H₂O is additionally analysed in Fig. 8 for the extended 2008 Antarctic winter. All model simulations produce the same gas-phase H₂O concentrations since ice is formed as a part of UKESM1.1's microphysics and is unchanged in the new PSC parameterisation. The new scheme only affects how ice mass mixing ratio is converted to SAD for heterogeneous chemistry. Furthermore, the amount of H₂O that condenses to form STS particles is negligible compared to the gas-phase H₂O concentrations. Compared to MLS, the model slightly underestimates gas-phase H₂O in May and June, but shows weaker dehydration from July to November. As a result, simulated gas-phase H₂O is comparable to MLS over 60°S–70°S (Fig. 8a) and higher than MLS over the 70°S–80°S (Fig. 8b). The same analysis is presented in the Supplementary Material for the Arctic winter at 68 hPa (Fig. S7).

3.4 Comparison with ozone from MLS

To assess how the simulations represent O₃, Fig. 9 compares the simulated O₃ at 68 hPa from CONTROL and the four selected sensitivity runs (RUN_{LL}, RUN_{LH}, RUN_{HL}, and RUN_{HH}) with MLS observations during the extended 2008 Antarctic winter, extending into early springtime to capture ozone depletion. The MLS data shows distinct O₃ loss starting around mid-

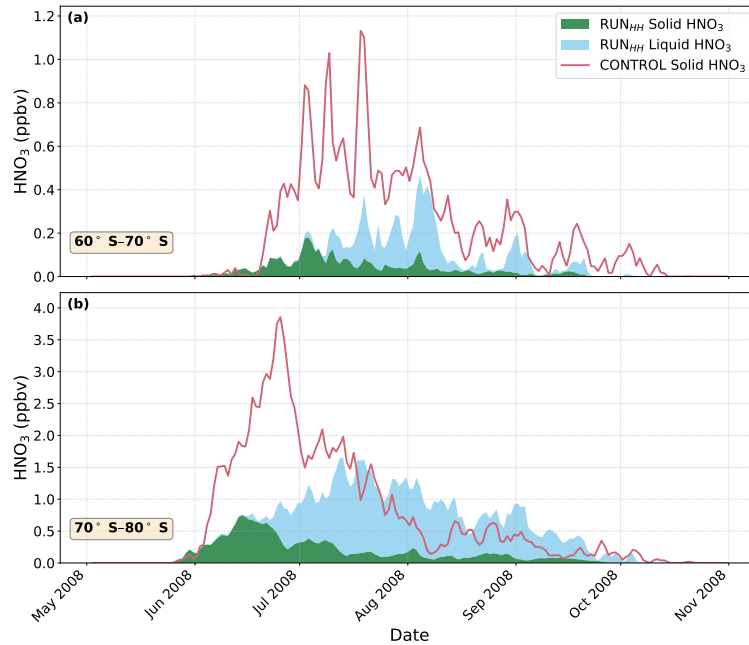


Figure 7. Zonal mean volume mixing ratio of solid- and liquid-phase HNO_3 (ppbv) at 68 hPa for CONTROL and RUN_{HH} over the Antarctic for the extended 2008 winter season (May to October), averaged across latitude bands 60°S – 70°S (a), and 70°S – 80°S (b). Solid-phase HNO_3 for CONTROL is shown with a pink line. The solid- and liquid-phase HNO_3 for RUN_{HH} are shown as a stacked area, with solid-phase in green and liquid-phase in blue. Note that liquid-phase HNO_3 is not shown for CONTROL, as this is not simulated by the original PSC parameterisation. Also, the panels have different scales.

August, reaching near-complete depletion over the latitude band 70°S – 80°S by early October, when the Antarctic ozone hole
 410 tends to form. While both the CONTROL and sensitivity runs capture the timing of O_3 depletion at 60°S – 70°S , CONTROL underestimates O_3 from September to October, whereas the sensitivity runs overestimate it from June to August (Fig. 9a). At 70°S – 80°S , both the CONTROL and the sensitivity runs underestimate O_3 from May to June, but overestimate it from August to September, with the same bias persisting into October for the sensitivity runs (Fig. 9b). All four sensitivity runs show comparable O_3 concentrations. More generally, there is often little difference between the amount of O_3 simulated by
 415 CONTROL and the sensitivity runs, with the exception being over the latitude band 60°S – 70°S (i.e., near the edge of the polar vortex) from September to October, where the sensitivity runs agree better with MLS. A similar analysis at 100 hPa is presented in the Supplementary Material (Fig. S8), with these results in general agreement with Fig. 9.

4 Summary

This study attempts to improve the representation of PSCs in UKESM1.1 by replacing its current, simple scheme with a kinetic
 420 scheme for NAT formation and a thermodynamic scheme for STS formation. These improvements enhance UKESM1.1's abil-

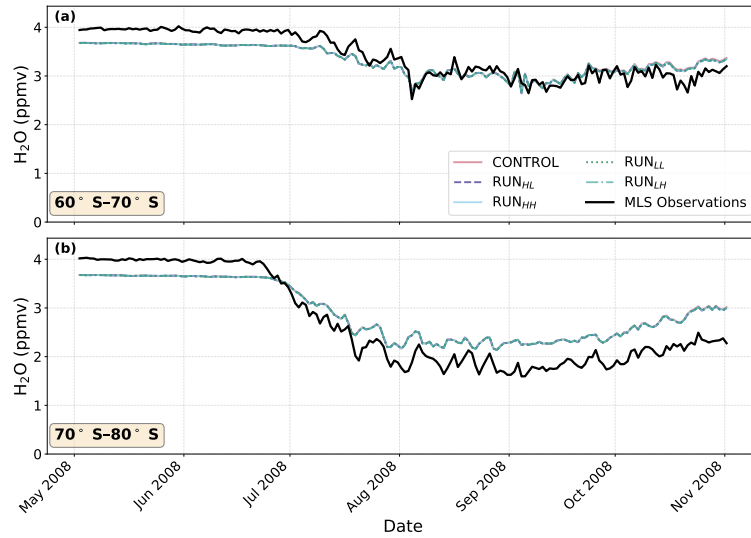


Figure 8. As Fig. 6, but for zonal mean gas-phase H_2O (ppmv). Note that there is complete overlap of all model runs (CONTROL and sensitivity simulations).

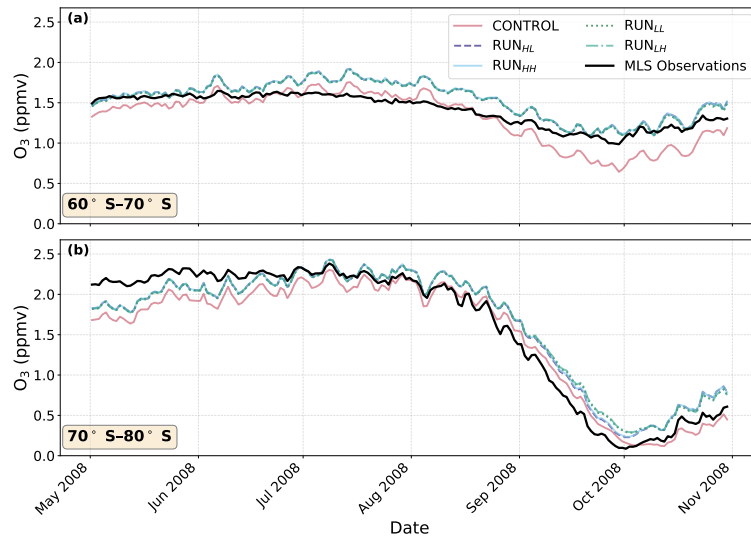


Figure 9. As Fig. 6, but for zonal mean O_3 (ppmv). Note that there is almost complete overlap of RUN_{HH} and RUN_{HL} , and RUN_{LL} and RUN_{LH} , resulting in only RUN_{HH} and RUN_{LL} being visible in the plot.

ity to represent the diversity and seasonal evolution of PSCs. Comparisons with CALIOP observations for the 2008 Antarctic and 2009/2010 Arctic winters demonstrates that the new parameterisation scheme achieves closer agreement with CALIOP observations by increasing the number of points classifying as STS and ice and capturing seasonal variability in PSC types

(Figs. 1 and 2). A sensitivity analysis using the new scheme reveals that $n_{\text{NAT,max}}$ and n_{ice} directly influences the relative abundance of each PSC type (Figs. 1 and 2). Larger values of these parameters favour the formation of smaller particles with greater SAD (Fig. S3). Additionally, the influence of $n_{\text{NAT,max}}$ is particularly strong in the Arctic, where warmer temperatures limit ice particle formation (Fig. 2). However, there is still a gap in the ability of the model to simulate high $\beta_{\perp}$ and high R_{532} PSCs that are observed by CALIOP (Figs. 1 and 2).

The improvement in the diversity of PSCs simulated by the new scheme is further supported by an examination of the temporal evolution of PSCs over the Antarctic winter season (Figs. 3 and 4). While overall NAT and ice occurrence frequencies remain similar to the original scheme, the new scheme produces STS in regions where it was previously absent, especially near the vortex edge. Furthermore, analysis of SAD over the 2008 Antarctic winter shows that the new scheme produces a greater diversity of PSC types by slowing NAT growth and enabling the formation of STS droplets, resulting in STS contributing a greater proportion to the total PSC SAD than NAT, especially later in the season. (Fig. 5).

The updated scheme leads to modest improvements in gas-phase HNO_3 compared to MLS, particularly within the polar vortex, although all model simulations significantly underestimate HNO_3 (Fig. 6). The overall impact on stratospheric O_3 is small, but differences emerge near the polar vortex edge, where reduced O_3 loss from August to October brings the sensitivity simulations closer to MLS observations (Fig. 9). Conditions near the vortex edge are more variable, meaning changes in PSC formation, such as slowing the growth of NAT, may have a stronger influence on heterogeneous chemistry and ozone depletion. In contrast, changes within the vortex core are limited because ozone depletion is already close to saturation under persistently cold and isolated conditions (Kirner et al., 2015; Tritscher et al., 2021).

The severe underestimation of HNO_3 at the beginning of the PSC season, likely due to a known low- NO_y bias in the UKESM1.1 stratosphere (Archibald et al., 2020), means that the improved comparison with observations of gas-phase HNO_3 from MLS using the new scheme does not necessarily imply an improvement in the ability of UKESM1.1 to accurately capture the full extent of HNO_3 depletion and denitrification. This is because UKESM1.1 begins the PSC/winter season with less HNO_3 than observed. This bias also affects NAT and STS formation since both depend on the partial pressure of HNO_3 . The previous thermodynamic equilibrium approach may have partly compensated for this bias in terms of NAT formation by converting all excess gas-phase HNO_3 into NAT, whereas the new scheme slows NAT growth, making the effect of low HNO_3 on NAT formation more apparent. As a result, the reduction in NAT SAD in the new scheme reflects both slower NAT growth and enhanced STS formation, but is likely also amplified by the background HNO_3 bias. The origin of this bias requires further investigation.

The CONTROL and sensitivity runs produced less ice than observed by CALIOP and failed to form wave ice (Figs. 1 and 2). This likely arises from the prescribed ice number density, which constrains the ice particles to a narrow range of SAD (Fig. S3). This may limit the model's ability to reproduce the broader range of optical properties observed by CALIOP. Similar findings have been noted in other models (Weimer et al., 2021), suggesting that reproducing high-number-density ice particles requires a method that does not have a fixed prescribed ice number density. Additionally, adiabatic cooling from fine-scale mountain waves propagating into the stratosphere can cause rapid, localised cooling in the stratosphere that triggers the formation of ice

PSCs, which in turn can seed NAT mother clouds downstream, resulting in high-number-density NAT and ice PSCs (Drdla et al., 2002; Voigt et al., 2003; Fueglistaler et al., 2003).

460 A parameterisation scheme that represented the cooling phase of these mountain-wave-induced temperature fluctuations was implemented in an earlier version of UKESM, and resulted in a considerable increase in the number of occasions that simulated stratospheric temperatures dropped below the threshold for ice PSCs (Orr et al., 2015, 2020). However, this scheme was not implemented in UKESM1.1, and so these effects are not represented in our simulations. This, combined with prescribing ice and NAT number densities, likely limits the formation of high-number-density PSCs in our simulations using the new
465 PSC parameterisation scheme. To address this, implementing the mountain-wave-induced temperature fluctuation scheme and allowing n_{ice} and $n_{\text{NAT,max}}$ to vary with cooling rate could improve the representation of PSC variability (Orr et al., 2015, 2020). Such development could further advance UKESM1.1’s capability to simulate PSC formation and variability, contributing to more accurate projection of future changes in polar ozone and climate (Polvani et al., 2011; Previdi and Polvani, 2014).

Code and data availability. All simulations used in this work were performed using version 13.9 of the Met Office Unified Model coupled to the United Kingdom Chemistry and Aerosol model (UM-UKCA). The UM code branch have been made available to reviewers
470 of this paper. Due to intellectual property copyright restrictions, the source code of the UM cannot be provided. The UKCA code used for this analysis can be accessed at DOI:10.5281/zenodo.17949939 (Sangha, 2025). The UM simulations are compiled and run in suites developed using the Rose suite engine (<https://metomi.github.io/rose/doc/html/index.html>, last accessed: 5 December 2025) and scheduled using the Cylc workflow engine (<https://cylc.github.io/cylc/>, last accessed 5 December 2025). Both Rose and Cylc are available under ver-
475 sion 3 of the GNU General Public License (GPL). The suite contains information required to extract and build the code, and configure and run the simulations. Every suite has a unique identifier and is held in the same revision-controlled repository service as the model’s code. All suites are available to any licensed user of the UM. The ZOMM optical model code is documented in Hoyle et al. (2013) and Engel et al. (2013). CALIPSO lidar level 2 polar stratospheric cloud mask version 2.0 (v2) is available at via DOI: 10.5067/CALIOP/CALIPSO/CAL_LID_L2_PSCMASK-STANDARD-V2-00 (https://asdc.larc.nasa.gov/project/CALIPSO/CAL_LID_L2_PSCMask-Standard-V2-00_V2-00). MLS HNO_3 and O_3 data products have been downloaded from <https://mls.jpl.nasa.gov/eos-aura-mls/data.php>, last accessed De-
480 cember 5, 2025 (Manney et al., 2020; Schwartz et al., 2020).

Author contributions. IS implemented the updated PSC scheme parameterisation into UKESM1.1, ran the simulations, analysed the data, and wrote the original manuscript. All authors provided guidance throughout the study and contributed to revising the manuscript. MW contributed the ICON-ART PSC code. MP and LP contributed figure panels based on CALIOP observations and ZOMM optical code.

485 *Competing interests.* The authors declare that they have no conflict of interests

Acknowledgements. We thank the editor and three anonymous reviewers for their valuable comments and suggestions, which have greatly improved the quality of this manuscript. IS was supported by the Natural Environment Research Council (NERC) grant number NE/S007164/1. NLA would like to thank NERC for financial support through the VISION project, NE/Z503393/1. AO was supported by NERC through the TerraFIRMA: Future Impacts, Risks and Mitigation Actions in a changing Earth System project, Grant reference NE/W004895/1. HL
490 was supported by the NERC-CANARI project (NE/W004984/1). MP and LP were supported by funding from the project Improved characterization of PSC processes derived from a third-generation CALIOP and MLD detection and composition classification algorithm, NASA Research Opportunities in Space and Earth Sciences. This work used Monsoon3, a collaborative High-Performance Computing facility funded by the Met Office and NERC, and JASMIN, the UK collaborative data analysis facility. We thank Peter Braesicke for providing ICON-ART PSC code, Paul Field for advice on the UKESM1.1 cloud microphysics scheme, and Alyn Lambert for support with running the
495 ZOMM optical code.

References

- Archibald, A. T., O'Connor, F. M., Abraham, N. L., Archer-Nicholls, S., Chipperfield, M. P., Dalvi, M., Folberth, G. A., Dennison, F., Dhomse, S. S., Griffiths, P. T., Hardacre, C., Hewitt, A. J., Hill, R. S., Johnson, C. E., Keeble, J., Köhler, M. O., Morgenstern, O., Mulcahy, J. P., Ordóñez, C., Pope, R. J., Rumbold, S. T., Russo, M. R., Savage, N. H., Sellar, A., Stringer, M., Turnock, S. T., Wild, O., and Zeng, G.: Description and evaluation of the UKCA stratosphere–troposphere chemistry scheme (StratTrop vn 1.0) implemented in UKESM1, *Geoscientific Model Development*, 13, 1223–1266, <https://doi.org/10.5194/gmd-13-1223-2020>, 2020.
- Austin, J., Shindell, D., Beagley, S. R., Brühl, C., Dameris, M., Manzini, E., Nagashima, T., Newman, P., Pawson, S., Pitari, G., Rozanov, E., Schnadt, C., and Shepherd, T. G.: Uncertainties and assessments of chemistry–climate models of the stratosphere, *Atmospheric Chemistry and Physics*, 3, 1–27, <https://doi.org/10.5194/acp-3-1-2003>, 2003.
- Biele, J., Tsias, A., Luo, B. P., Carslaw, K. S., Neuber, R., Beyerle, G., and Peter, T.: Nonequilibrium coexistence of solid and liquid particles in Arctic stratospheric clouds, *Journal of Geophysical Research: Atmospheres*, 106, 22 991–23 007, <https://doi.org/10.1029/2001JD900188>, 2001.
- Carslaw, K. S., Luo, B., and Peter, T.: An analytic expression for the composition of aqueous HNO₃–H₂SO₄ stratospheric aerosols including gas phase removal of HNO₃, *Geophysical Research Letters*, 22, 1877–1880, <https://doi.org/10.1029/95GL01668>, 1995.
- Carslaw, K. S., Peter, T., and Clegg, S. L.: Modeling the composition of liquid stratospheric aerosols, *Reviews of Geophysics*, 35, 125–154, <https://doi.org/10.1029/97RG00078>, 1997.
- Carslaw, K. S., Kettleborough, J. A., Northway, M. J., Davies, S., Gao, R.-S., Fahey, D. W., Baumgardner, D. G., Chipperfield, M. P., and Kleinböhl, A.: A vortex-scale simulation of the growth and sedimentation of large nitric acid hydrate particles, *Journal of Geophysical Research: Atmospheres*, 107, 43–1, <https://doi.org/10.1029/2001JD000467>, 2002.
- Chipperfield, M. P.: Multiannual simulations with a three-dimensional chemical transport model, *Journal of Geophysical Research: Atmospheres*, 104, 1781–1805, <https://doi.org/10.1029/98JD02597>, 1999.
- Chipperfield, M. P. and Pyle, J. A.: Model sensitivity studies of Arctic ozone depletion, *Journal of Geophysical Research: Atmospheres*, 103, 28 389–28 403, <https://doi.org/10.1029/98JD01960>, 1998.
- Cicerone, R. J., Stolarski, R. S., and Walters, S.: Stratospheric Ozone Destruction by Man-Made Chlorofluoromethanes, *Science*, 185, 1165–1167, <https://doi.org/10.1126/science.185.4157.1165>, 1974.
- Dennison, F., Keeble, J., Morgenstern, O., Zeng, G., Abraham, N. L., and Yang, X.: Improvements to stratospheric chemistry scheme in the UM-UKCA (v10.7) model: solar cycle and heterogeneous reactions, *Geoscientific Model Development*, 12, 1227–1239, <https://doi.org/10.5194/gmd-12-1227-2019>, 2019.
- Dhomse, S. S., Emmerson, K. M., Mann, G. W., Bellouin, N., Carslaw, K. S., Chipperfield, M. P., Hommel, R., Abraham, N. L., Telford, P., and Braesicke, P.: Aerosol microphysics simulations of the Mt.~Pinatubo eruption with the UM-UKCA composition–climate model, *Atmospheric Chemistry and Physics*, 14, 11 221–11 246, <https://doi.org/10.5194/acp-14-11221-2014>, 2014.
- Doms, G., Förstner, J., Heise, E., Herzog, H. J., Mironov, D., Raschendorfer, M., Reinhardt, T., Ritter, B., Schrodin, R., and Schulz, J.-P.: A description of the nonhydrostatic regional COSMO model. Part II: Physical parameterization, *Deutscher Wetterdienst*, Offenbach, Germany, 2011.
- Dörnbrack, A., Pitts, M. C., Poole, L. R., Orsolini, Y. J., Nishii, K., and Nakamura, H.: The 2009–2010 Arctic stratospheric winter – general evolution, mountain waves and predictability of an operational weather forecast model, *Atmospheric Chemistry and Physics*, 12, 3659–3675, <https://doi.org/10.5194/acp-12-3659-2012>, 2012.

- Drdla, K., Gandrud, B. W., Baumgardner, D., Wilson, J. C., Bui, T. P., Hurst, D., Schauffler, S. M., Jost, H., Greenblatt, J. B., and Webster, C. R.: Evidence for the widespread presence of liquid-phase particles during the 1999–2000 Arctic winter, *Journal of Geophysical Research: Atmospheres*, 107, 61–1, <https://doi.org/10.1029/2001JD001127>, 2002.
- Engel, I., Luo, B. P., Pitts, M. C., Poole, L. R., Hoyle, C. R., Grooß, J.-U., Dörnbrack, A., and Peter, T.: Heterogeneous formation of polar stratospheric clouds – Part 2: Nucleation of ice on synoptic scales, *Atmospheric Chemistry and Physics*, 13, 10 769–10 785, <https://doi.org/10.5194/acp-13-10769-2013>, 2013.
- Eyring, V., Butchart, N., Waugh, D. W., Akiyoshi, H., Austin, J., Bekki, S., Bodeker, G. E., Boville, B. A., Brühl, C., Chipperfield, M. P., Cordero, E., Dameris, M., Deushi, M., Fioletov, V. E., Frith, S. M., Garcia, R. R., Gettelman, A., Giorgetta, M. A., Grewe, V., Jourdain, L., Kinnison, D. E., Mancini, E., Manzini, E., Marchand, M., Marsh, D. R., Nagashima, T., Newman, P. A., Nielsen, J. E., Pawson, S., Pitari, G., Plummer, D. A., Rozanov, E., Schraner, M., Shepherd, T. G., Shibata, K., Stolarski, R. S., Struthers, H., Tian, W., and Yoshiki, M.: Assessment of temperature, trace species, and ozone in chemistry-climate model simulations of the recent past, *Journal of Geophysical Research: Atmospheres*, 111, <https://doi.org/10.1029/2006JD007327>, 2006.
- Eyring, V., Arblaster, J. M., Cionni, I., Sedláček, J., Perlwitz, J., Young, P. J., Bekki, S., Bergmann, D., Cameron-Smith, P., Collins, W. J., Faluvegi, G., Gottschaldt, K.-D., Horowitz, L. W., Kinnison, D. E., Lamarque, J.-F., Marsh, D. R., Saint-Martin, D., Shindell, D. T., Sudo, K., Szopa, S., and Watanabe, S.: Long-term ozone changes and associated climate impacts in CMIP5 simulations, *Journal of Geophysical Research: Atmospheres*, 118, 5029–5060, <https://doi.org/10.1002/jgrd.50316>, 2013.
- Fahey, D. W., Gao, R. S., Carslaw, K. S., Kettleborough, J., Popp, P. J., Northway, M. J., Holecek, J. C., Ciciora, S. C., McLaughlin, R. J., Thompson, T. L., Winkler, R. H., Baumgardner, D. G., Gandrud, B., Wennberg, P. O., Dhaniyala, S., McKinney, K., Peter, T., Salawitch, R. J., Bui, T. P., Elkins, J. W., Webster, C. R., Atlas, E. L., Jost, H., Wilson, J. C., Herman, R. L., Kleinböhl, A., and von König, M.: The Detection of Large HNO₃-Containing Particles in the Winter Arctic Stratosphere, *Science*, 291, 1026–1031, <https://doi.org/10.1126/science.1057265>, 2001.
- Farman, J. C., Gardiner, B. G., and Shanklin, J. D.: Large losses of total ozone in Antarctica reveal seasonal ClO_x/NO_x interaction, *Nature*, 315, 207–210, <https://doi.org/10.1038/315207a0>, 1985.
- Feng, L., Smith, S. J., Braun, C., Crippa, M., Gidden, M. J., Hoesly, R., Klimont, Z., van Marle, M., van den Berg, M., and van der Werf, G. R.: The generation of gridded emissions data for CMIP6, *Geoscientific Model Development*, 13, 461–482, <https://doi.org/10.5194/gmd-13-461-2020>, 2020.
- Fueglistaler, S., Luo, B. P., Voigt, C., Carslaw, K. S., and Peter, T.: NAT-rock formation by mother clouds: a microphysical model study, *Atmospheric Chemistry and Physics*, 2, 93–98, <https://doi.org/10.5194/acp-2-93-2002>, 2002.
- Fueglistaler, S., Buss, S., Luo, B. P., Wernli, H., Flentje, H., Hostetler, C. A., Poole, L. R., Carslaw, K. S., and Peter, T.: Detailed modeling of mountain wave PSCs, *Atmospheric Chemistry and Physics*, 3, 697–712, <https://doi.org/10.5194/acp-3-697-2003>, 2003.
- Gelaro, R., McCarty, W., Suárez, M. J., Todling, R., Molod, A., Takacs, L., Randles, C. A., Darmenov, A., Bosilovich, M. G., Reichle, R., Wargan, K., Coy, L., Cullather, R., Draper, C., Akella, S., Buchard, V., Conaty, A., da Silva, A. M., Gu, W., Kim, G.-K., Koster, R., Lucchesi, R., Merkova, D., Nielsen, J. E., Partyka, G., Pawson, S., Putman, W., Rienecker, M., Schubert, S. D., Sienkiewicz, M., and Zhao, B.: The Modern-Era Retrospective Analysis for Research and Applications, Version 2 (MERRA-2), *Journal of Climate*, 30, 5419–5454, <https://doi.org/10.1175/JCLI-D-16-0758.1>, 2017.
- Grainger, R. G., Lambert, A., Rodgers, C. D., Taylor, F. W., and Deshler, T.: Stratospheric aerosol effective radius, surface area and volume estimated from infrared measurements, *Journal of Geophysical Research: Atmospheres*, 100, 16 507–16 518, <https://doi.org/10.1029/95JD00988>, 1995.

- Groß, J.-U., Günther, G., Konopka, P., Müller, R., McKenna, D. S., Stroh, F., Vogel, B., Engel, A., Müller, M., Hoppel, K., Bevilacqua, R., Richard, E., Webster, C. R., Elkins, J. W., Hurst, D. F., Romashkin, P. A., and Baumgardner, D. G.: Simulation of ozone depletion in spring 2000 with the Chemical Lagrangian Model of the Stratosphere (CLaMS), *Journal of Geophysical Research: Atmospheres*, 107, 38–1, <https://doi.org/10.1029/2001JD000456>, 2002.
- 575 Hanson, D. R. and Mauersberger, K.: Laboratory studies of the nitric acid trihydrate: Implications for the south polar stratosphere, *Geophysical Research Letters*, 15, 855–858, <https://doi.org/10.1017/S0954102026100601>, 1988.
- Hersbach, H., Bell, B., Berrisford, P., Hirahara, S., Horányi, A., Muñoz-Sabater, J., Nicolas, J., Peubey, C., Radu, R., Schepers, D., Simmons, A., Soci, C., Abdalla, S., Abellan, X., Balsamo, G., Bechtold, P., Biavati, G., Bidlot, J., Bonavita, M., De Chiara, G., Dahlgren, P., Dee, D., Diamantakis, M., Dragani, R., Flemming, J., Forbes, R., Fuentes, M., Geer, A., Haimberger, L., Healy, S., Hogan, R. J.,
- 580 Hólm, E., Janisková, M., Keeley, S., Laloyaux, P., Lopez, P., Lupu, C., Radnoti, G., de Rosnay, P., Rozum, I., Vamborg, F., Villaume, S., and Thépaut, J.-N.: The ERA5 global reanalysis, *Quarterly Journal of the Royal Meteorological Society*, 146, 1999–2049, <https://doi.org/10.1002/qj.3803>, 2020.
- Hervig, M. E. and Deshler, T.: Stratospheric aerosol surface area and volume inferred from HALOE, CLAES, and ILAS measurements, *Journal of Geophysical Research: Atmospheres*, 103, 25 345–25 352, <https://doi.org/10.1029/98JD01962>, 1998.
- 585 Hoshyaripour, G. A., Baer, A., Bierbauer, S., Bruckert, J., Brunner, D., Foerstner, J., Hamzehloo, A., Hanft, V., Keller, C., Klose, M., Kumar, P., Ludwig, P., Metzner, E., Muth, L., Pauling, A., Porz, N., Reddmann, T., Reißig, L., Ruhnke, R., Satitkovitchai, K., Seifert, A., Sinnhuber, M., Steiner, M., Versick, S., Vogel, H., Weimer, M., Werchner, S., and Hoose, C.: The Atmospheric Composition Component of the ICON modeling framework: ICON-ART version 2025.04, *EGUsphere*, 2025, 1–54, <https://doi.org/10.5194/egusphere-2025-3400>, 2025.
- 590 Hostetler, C. A., Liu, Z., Reagan, J., Vaughan, M., Winker, D., Osborn, M., Hunt, W. H., Powell, K. A., and Trepte, C.: Cloud-Aerosol Lidar with Orthogonal Polarization (CALIOP) Algorithm Theoretical Basis Document, <https://doi.org/10.1109/IGARSS.2003.1294160>, 2006.
- Hoyle, C. R., Engel, I., Luo, B. P., Pitts, M. C., Poole, L. R., Groß, J.-U., and Peter, T.: Heterogeneous formation of polar stratospheric clouds – Part 1: Nucleation of nitric acid trihydrate (NAT), *Atmospheric Chemistry and Physics*, 13, 9577–9595, <https://doi.org/10.5194/acp-13-9577-2013>, 2013.
- 595 Kirner, O., Ruhnke, R., Buchholz-Dietsch, J., Jöckel, P., Brühl, C., and Steil, B.: Simulation of polar stratospheric clouds in the chemistry-climate-model EMAC via the submodel PSC, *Geoscientific Model Development*, 4, 169–182, <https://doi.org/10.5194/gmd-4-169-2011>, 2011.
- Kirner, O., Müller, R., Ruhnke, R., and Fischer, H.: Contribution of liquid, NAT and ice particles to chlorine activation and ozone depletion in Antarctic winter and spring, *Atmospheric Chemistry and Physics*, 15, 2019–2030, <https://doi.org/10.5194/acp-15-2019-2015>, 2015.
- 600 Koop, T., Luo, B., Tsias, A., and Peter, T.: Water activity as the determinant for homogeneous ice nucleation in aqueous solutions, *Nature*, 406, 611–614, <https://doi.org/10.1038/35020537>, 2000.
- Liu, Z., Hunt, W., Vaughan, M., Hostetler, C., McGill, M., Powell, K., Winker, D., and Hu, Y.: Estimating random errors due to shot noise in backscatter lidar observations, *Appl. Opt.*, 45, 4437–4447, <https://doi.org/10.1364/AO.45.004437>, 2006.
- Livesey, N. J., Van Snyder, W., Read, W. G., and Wagner, P. A.: Retrieval algorithms for the EOS Microwave limb sounder (MLS), *IEEE Transactions on Geoscience and Remote Sensing*, 44, 1144–1155, <https://doi.org/10.1109/TGRS.2006.872327>, 2006.
- 605 Mann, G. W., Carslaw, K. S., Spracklen, D. V., Ridley, D. A., Manktelow, P. T., Chipperfield, M. P., Pickering, S. J., and Johnson, C. E.: Description and evaluation of GLOMAP-mode: a modal global aerosol microphysics model for the UKCA composition-climate model, *Geoscientific Model Development*, 3, 519–551, <https://doi.org/10.5194/gmd-3-519-2010>, 2010.

- Manney, G., Santee, M., Froidevaux, L., Livesey, N., and Read, W.: MLS/Aura Level 2 Nitric Acid (HNO₃) Mixing Ratio V005, <https://doi.org/10.5067/Aura/MLS/DATA2511>, 2020.
- Mishchenko, M. I., Travis, L. D., and Mackowski, D. W.: T-matrix computations of light scattering by nonspherical particles: A review, *Journal of Quantitative Spectroscopy and Radiative Transfer*, 55, 535–575, [https://doi.org/10.1016/0022-4073\(96\)00002-7](https://doi.org/10.1016/0022-4073(96)00002-7), 1996.
- Molina, L. T. and Molina, M. J.: Production of chlorine oxide (Cl₂O₂) from the self-reaction of the chlorine oxide (ClO) radical, *Journal of Physical Chemistry*, 91, 433–436, <https://doi.org/10.1021/j100286a035>, 1987.
- Morgenstern, O., Braesicke, P., O'Connor, F. M., Bushell, A. C., Johnson, C. E., Osprey, S. M., and Pyle, J. A.: Evaluation of the new UKCA climate-composition model – Part 1: The stratosphere, *Geoscientific Model Development*, 2, 43–57, <https://doi.org/10.5194/gmd-2-43-2009>, 2009.
- Mulcahy, J. P., Johnson, C., Jones, C. G., Povey, A. C., Scott, C. E., Sellar, A., Turnock, S. T., Woodhouse, M. T., Abraham, N. L., Andrews, M. B., Bellouin, N., Browse, J., Carslaw, K. S., Dalvi, M., Folberth, G. A., Glover, M., Grosvenor, D. P., Hardacre, C., Hill, R., Johnson, B., Jones, A., Kipling, Z., Mann, G., Mollard, J., O'Connor, F. M., Palmiéri, J., Reddington, C., Rumbold, S. T., Richardson, M., Schutgens, N. A. J., Stier, P., Stringer, M., Tang, Y., Walton, J., Woodward, S., and Yool, A.: Description and evaluation of aerosol in UKESM1 and HadGEM3-GC3.1 CMIP6 historical simulations, *Geoscientific Model Development*, 13, 6383–6423, <https://doi.org/10.5194/gmd-13-6383-2020>, 2020.
- Mulcahy, J. P., Jones, C. G., Rumbold, S. T., Kuhlbrodt, T., Dittus, A. J., Blockley, E. W., Yool, A., Walton, J., Hardacre, C., Andrews, T., Bodas-Salcedo, A., Stringer, M., de Mora, L., Harris, P., Hill, R., Kelley, D., Robertson, E., and Tang, Y.: UKESM1.1: development and evaluation of an updated configuration of the UK Earth System Model, *Geoscientific Model Development*, 16, 1569–1600, <https://doi.org/10.5194/gmd-16-1569-2023>, 2023.
- Orr, A., Marshall, G. J., Hunt, J. C. R., Sommeria, J., Wang, C.-G., van Lipzig, N. P. M., Cresswell, D., and King, J. C.: Characteristics of Summer Airflow over the Antarctic Peninsula in Response to Recent Strengthening of Westerly Circumpolar Winds, *Journal of the Atmospheric Sciences*, 65, 1396–1413, <https://doi.org/10.1175/2007JAS2498.1>, 2008.
- Orr, A., Hosking, J. S., Hoffmann, L., Keeble, J., Dean, S. M., Roscoe, H. K., Abraham, N. L., Vosper, S., and Braesicke, P.: Inclusion of mountain-wave-induced cooling for the formation of PSCs over the Antarctic Peninsula in a chemistry–climate model, *Atmospheric Chemistry and Physics*, 15, 1071–1086, <https://doi.org/10.5194/acp-15-1071-2015>, 2015.
- Orr, A., Scott Hosking, J., Delon, A., Hoffmann, L., Spang, R., Moffat-Griffin, T., Keeble, J., Luke Abraham, N., Braesicke, P., and Orr, A.: Polar stratospheric clouds initiated by mountain waves in a global chemistry-climate model: A missing piece in fully modelling polar stratospheric ozone depletion, *Atmospheric Chemistry and Physics*, 20, <https://doi.org/10.5194/acp-20-12483-2020>, 2020.
- Peter, T. and GroöB, J.-U.: Polar stratospheric clouds and sulfate aerosol particles: microphysics, denitrification and heterogeneous chemistry, *Stratospheric ozone depletion and climate change*, pp. 108–144, 2012.
- Pitts, M. C., Poole, L. R., Dörnbrack, A., and Thomason, L. W.: The 2009–2010 Arctic polar stratospheric cloud season: a CALIPSO perspective, *Atmospheric Chemistry and Physics*, 11, 2161–2177, <https://doi.org/10.5194/acp-11-2161-2011>, 2011.
- Pitts, M. C., Poole, L. R., Lambert, A., and Thomason, L. W.: An assessment of CALIOP polar stratospheric cloud composition classification, *Atmospheric Chemistry and Physics*, 13, 2975–2988, <https://doi.org/10.5194/acp-13-2975-2013>, 2013.
- Pitts, M. C., Poole, L. R., and Gonzalez, R.: Polar stratospheric cloud climatology based on CALIPSO spaceborne lidar measurements from 2006 to 2017, *Atmospheric Chemistry and Physics*, 18, 10 881–10 913, <https://doi.org/10.5194/acp-18-10881-2018>, 2018.

- 645 Polvani, L., Waugh, D., Correa, G., and Son, S.-W.: Stratospheric Ozone Depletion: The Main Driver of Twentieth-Century Atmospheric Circulation Changes in the Southern Hemisphere, *Journal of Climate - J CLIMATE*, 24, 795–812, <https://doi.org/10.1175/2010JCLI3772.1>, 2011.
- Polvani, L. M., Abalos, M., Garcia, R., Kinnison, D., and Randel, W. J.: Significant Weakening of Brewer-Dobson Circulation Trends Over the 21st Century as a Consequence of the Montreal Protocol, *Geophysical Research Letters*, 45, 401–409, <https://doi.org/10.1002/2017GL075345>, 2018.
- 650 Previdi, M. and Polvani, L. M.: Climate system response to stratospheric ozone depletion and recovery, *Quarterly Journal of the Royal Meteorological Society*, 140, 2401–2419, <https://doi.org/10.1002/qj.2330>, 2014.
- Randel, W. J. and Wu, F.: Cooling of the Arctic and Antarctic Polar Stratospheres due to Ozone Depletion, *Journal of Climate*, 12, 1467–1479, [https://doi.org/10.1175/1520-0442\(1999\)012<1467:COTAAA>2.0.CO;2](https://doi.org/10.1175/1520-0442(1999)012<1467:COTAAA>2.0.CO;2), 1999.
- 655 Ravishankara, A. R. and Hanson, D. R.: Differences in the reactivity of type I polar stratospheric clouds depending on their phase, *Journal of Geophysical Research: Atmospheres*, 101, 3885–3890, <https://doi.org/10.1029/95JD03009>, 1996.
- Reynolds, R. W., Rayner, N. A., Smith, T. M., Stokes, D. C., and Wang, W.: An Improved In Situ and Satellite SST Analysis for Climate, *Journal of Climate*, 15, 1609–1625, [https://doi.org/10.1175/1520-0442\(2002\)015<1609:AIISAS>2.0.CO;2](https://doi.org/10.1175/1520-0442(2002)015<1609:AIISAS>2.0.CO;2), 2002.
- Sangha, I.: UKCA PSC branch, <https://doi.org/DOI:10.5281/zenodo.17949939>, 2025.
- 660 Schröter, J., Rieger, D., Stassen, C., Vogel, H., Weimer, M., Werchner, S., Förstner, J., Prill, F., Reinert, D., and Zängl, G.: ICON-ART 2.1: A flexible tracer framework and its application for composition studies in numerical weather forecasting and climate simulations, *Geoscientific Model Development*, 11, 4043–4068, <https://doi.org/10.5194/gmd-11-4043-2018>, 2018.
- Schwartz, M., Froidevaux, L., Livesey, N., and Read, W.: MLS/Aura Level 2 Ozone (O₃) Mixing Ratio V005, <https://doi.org/10.5067/Aura/MLS/DATA2516>, 2020.
- 665 Sellar, A. A., Jones, C. G., Mulcahy, J. P., Tang, Y., Yool, A., Wiltshire, A., O'Connor, F. M., Stringer, M., Hill, R., Palmieri, J., Woodward, S., de Mora, L., Kuhlbrodt, T., Rumbold, S. T., Kelley, D. I., Ellis, R., Johnson, C. E., Walton, J., Abraham, N. L., Andrews, M. B., Andrews, T., Archibald, A. T., Berthou, S., Burke, E., Blockley, E., Carslaw, K., Dalvi, M., Edwards, J., Folberth, G. A., Gedney, N., Griffiths, P. T., Harper, A. B., Hendry, M. A., Hewitt, A. J., Johnson, B., Jones, A., Jones, C. D., Keeble, J., Liddicoat, S., Morgenstern, O., Parker, R. J., Predoi, V., Robertson, E., Siahann, A., Smith, R. S., Swaminathan, R., Woodhouse, M. T., Zeng, G., and Zerroukat, M.:
- 670 UKESM1: Description and Evaluation of the U.K. Earth System Model, *Journal of Advances in Modeling Earth Systems*, 11, 4513–4558, <https://doi.org/10.1029/2019MS001739>, 2019.
- Snels, M., Scoccione, A., Di Liberto, L., Colao, F., Pitts, M., Poole, L., Deshler, T., Cairo, F., Cagnazzo, C., and Fierli, F.: Comparison of Antarctic polar stratospheric cloud observations by ground-based and space-borne lidar and relevance for chemistry–climate models, *Atmospheric Chemistry and Physics*, 19, 955–972, <https://doi.org/10.5194/acp-19-955-2019>, 2019.
- 675 Solomon, S., Garcia, R., Rowland, F., and Wuebbles, D.: On the Depletion of Antarctic Ozone, *Nature*, 321, 755–758, <https://doi.org/10.1038/321755a0>, 1986.
- Spang, R., Hoffmann, L., Müller, R., Grooß, J.-U., Tritscher, I., Höpfner, M., Pitts, M., Orr, A., and Riese, M.: A climatology of polar stratospheric cloud composition between 2002 and 2012 based on MIPAS/Envisat observations, *Atmospheric Chemistry and Physics*, 18, 5089–5113, <https://doi.org/10.5194/acp-18-5089-2018>, 2018.
- 680 Steiner, M., Luo, B., Peter, T., Pitts, M. C., and Stenke, A.: Evaluation of polar stratospheric clouds in the global chemistry–climate model SOCOLv3.1 by comparison with CALIPSO spaceborne lidar measurements, *Geoscientific Model Development*, 14, 935–959, <https://doi.org/10.5194/gmd-14-935-2021>, 2021.

- Stephens, G. L., Vane, D. G., Boain, R. J., Mace, G. G., Sassen, K., Wang, Z., Illingworth, A. J., O’connor, E. J., Rossow, W. B., Dur-
den, S. L., Miller, S. D., Austin, R. T., Benedetti, A., and Mitrescu, C.: THE CLOUDSAT MISSION AND THE A-TRAIN: A New
685 Dimension of Space-Based Observations of Clouds and Precipitation, *Bulletin of the American Meteorological Society*, 83, 1771–1790,
<https://doi.org/10.1175/BAMS-83-12-1771>, 2002.
- Tesche, M., Achtert, P., and Pitts, M. C.: On the best locations for ground-based polar stratospheric cloud (PSC) observations, *Atmospheric
Chemistry and Physics*, 21, 505–516, <https://doi.org/10.5194/acp-21-505-2021>, 2021.
- Toon, O. B., Hamill, P., Turco, R. P., and Pinto, J.: Condensation of HNO₃ and HCl in the winter polar stratospheres, *Geophysical Research
690 Letters*, 13, 1284–1287, <https://doi.org/10.1029/GL013i012p01284>, 1986.
- Tritscher, I., Grooß, J.-U., Spang, R., Pitts, M. C., Poole, L. R., Müller, R., and Riese, M.: Lagrangian simulation of ice particles and resulting
dehydration in the polar winter stratosphere, *Atmospheric Chemistry and Physics*, 19, 543–563, <https://doi.org/10.5194/acp-19-543-2019>,
2019.
- Tritscher, I., Pitts, M. C., Poole, L. R., Alexander, S. P., Cairo, F., Chipperfield, M. P., Grooß, J.-U., Höpfner, M., Lambert, A., Luo, B.,
695 Molleker, S., Orr, A., Salawitch, R., Snels, M., Spang, R., Woiwode, W., and Peter, T.: Polar Stratospheric Clouds: Satellite Observations,
Processes, and Role in Ozone Depletion, *Reviews of Geophysics*, 59, e2020RG000 702, <https://doi.org/10.1029/2020RG000702>, 2021.
- van den Broek, M. M. P., Williams, J. E., and Bregman, A.: Implementing growth and sedimentation of NAT particles in a global Eulerian
model, *Atmospheric Chemistry and Physics*, 4, 1869–1883, <https://doi.org/10.5194/acp-4-1869-2004>, 2004.
- Voigt, C., Larsen, N., Deshler, T., Kröger, C., Schreiner, J., Mauersberger, K., Luo, B., Adriani, A., Cairo, F., Di Donfrancesco, G., Ovarlez,
700 J., Ovarlez, H., Dörnbrack, A., Knudsen, B., and Rosen, J.: In situ mountain-wave polar stratospheric cloud measurements: Implications
for nitric acid trihydrate formation, *Journal of Geophysical Research: Atmospheres*, 108, <https://doi.org/10.1029/2001JD001185>, 2003.
- Voigt, C., Schlager, H., Luo, B. P., Dörnbrack, A., Roiger, A., Stock, P., Curtius, J., Vössing, H., Borrmann, S., Davies, S., Konopka, P.,
Schiller, C., Shur, G., and Peter, T.: Nitric Acid Trihydrate (NAT) formation at low NAT supersaturation in Polar Stratospheric Clouds
(PSCs), *Atmospheric Chemistry and Physics*, 5, 1371–1380, <https://doi.org/10.5194/acp-5-1371-2005>, 2005.
- 705 Waters, J. W., Froidevaux, L., Harwood, R. S., Jarnot, R. F., Pickett, H. M., Read, W. G., Siegel, P. H., Cofield, R. E., Filipiak, M. J., Flower,
D. A., Holden, J. R., Lau, G. K., Livesey, N. J., Manney, G. L., Pumphrey, H. C., Santee, M. L., Wu, D. L., Cuddy, D. T., Lay, R. R., Loo,
M. S., Perun, V. S., Schwartz, M. J., Stek, P. C., Thurstans, R. P., Boyles, M. A., Chandra, K. M., Chavez, M. C., Chen, G.-S., Chudasama,
B. V., Dodge, R., Fuller, R. A., Girard, M. A., Jiang, J. H., Jiang, Y., Knosp, B. W., LaBelle, R. C., Lam, J. C., Lee, K. A., Miller, D.,
Oswald, J. E., Patel, N. C., Pukala, D. M., Quintero, O., Scaff, D. M., Van Snyder, W., Tope, M. C., Wagner, P. A., and Walch, M. J.:
710 The Earth observing system microwave limb sounder (EOS MLS) on the aura Satellite, *IEEE Transactions on Geoscience and Remote
Sensing*, 44, 1075–1092, <https://doi.org/10.1109/TGRS.2006.873771>, 2006.
- Weimer, M.: Towards Seamless Simulations of Polar Stratospheric Clouds and Ozone in the Polar Stratosphere with ICON-ART, Ph.D.
thesis, <https://doi.org/10.5445/IR/1000100338>, 2019.
- Weimer, M., Buchmüller, J., Hoffmann, L., Kirner, O., Luo, B., Ruhnke, R., Steiner, M., Tritscher, I., and Braesicke, P.: Mountain-wave-
715 induced polar stratospheric clouds and their representation in the global chemistry model ICON-ART, *Atmospheric Chemistry and
Physics*, 21, <https://doi.org/10.5194/acp-21-9515-2021>, 2021.
- Wilson, D. R. and Ballard, S. P.: A microphysically based precipitation scheme for the UK meteorological office unified model, *Quarterly
Journal of the Royal Meteorological Society*, 125, 1607–1636, <https://doi.org/10.1002/qj.49712555707>, 1999.
- Zeng, G., Morgenstern, O., Williams, J. H. T., O’Connor, F. M., Griffiths, P. T., Keeble, J., Deushi, M., Horowitz, L. W., Naik, V., Emmons,
720 L. K., Abraham, N. L., Archibald, A. T., Bauer, S. E., Hassler, B., Michou, M., Mills, M. J., Murray, L. T., Oshima, N., Sentman, L. T.,

- Tilmes, S., Tsigaridis, K., and Young, P. J.: Attribution of Stratospheric and Tropospheric Ozone Changes Between 1850 and 2014 in CMIP6 Models, *Journal of Geophysical Research: Atmospheres*, 127, e2022JD036452, <https://doi.org/10.1029/2022JD036452>, 2022.
- 725 Zhu, Y., Toon, O. B., Lambert, A., Kinnison, D. E., Brakebusch, M., Bardeen, C. G., Mills, M. J., and English, J. M.: Development of a Polar Stratospheric Cloud Model within the Community Earth System Model using constraints on Type I PSCs from the 2010–2011 Arctic winter, *Journal of Advances in Modeling Earth Systems*, 7, 551–585, <https://doi.org/10.1002/2015MS000427>, 2015.
- Zhu, Y., Toon, O. B., Pitts, M. C., Lambert, A., Bardeen, C., and Kinnison, D. E.: Comparing simulated PSC optical properties with CALIPSO observations during the 2010 Antarctic winter, *Journal of Geophysical Research: Atmospheres*, 122, 1175–1202, <https://doi.org/10.1002/2016JD025191>, 2017.