

Author's response for manuscript acp-2019-834:

'Robust observational constraint of uncertain aerosol processes and emissions in a climate model and the effect on aerosol radiative forcing'

In our response, reviewer comments are marked in bold, our responses and original text in plain text, and altered text in the paper in bold italic. Additionally, altered text in the tracked changes manuscript attached at the end of this response is highlighted with yellow.

Response to reviewer 1 (Anonymous reviewer)

This manuscript investigates the constraint on aerosol forcing uncertainties by observations of aerosol mass, number, and optical depth. The manuscript builds upon earlier work from the authors using statistical fits of climate model simulations in order to fully sample the uncertainty space of many (27) uncertain aerosol parameters in the model. The authors use the observations to determine which portions of the parameter space provide poor agreement with the observations and can be ruled out. They then determine how much this narrows the direct and cloud-albedo indirect forcing uncertainty ranges.

I feel this paper is ready for publication in ACP once several minor comments have been addressed.

We thank the reviewer for their interesting and useful comments on our manuscript, and respond in full to the comments below. We would like to clarify here that the PPE used in this study perturbed only 26 aerosol parameters. We did simultaneously create a 27 parameter PPE alongside this PPE (see Yoshioka et al., 2019), but that ensemble has not been analysed here.

Comment 1:

P6 L5: Have the authors thought about how much additional information would be added if they had the simulations/emulators built over all of the years with observations such that observations could be compared with the same month-year? What if the exact days/times could be compared? Certainly each of these steps requires a jump in computational investment, so it doesn't make sense to add here. But if the authors have ideas about how much it could help, it would be an interesting thing to discuss (perhaps in the conclusions?).

This is an interesting point and we have considered this. Accounting for all sources of model-measurement uncertainty is important, and our strategy here is to account for as many sources as possible in our definition of the implausibility metric. The representativeness error terms in the metric would be smaller if we had model output for several years spanning the measurement data, and at a higher temporal and spatial resolution, which would give us more confidence in ruling out implausible model variants.

We have found that inter-annual variability can account for around 30 to 90% of the model-measurement comparison representation uncertainty. Hence, making direct year-on-year comparisons between all measurements and the PPE could significantly reduce the effect of inter-annual variability on the comparison process. However, the large set of observations used in this study contains campaign data from many different years, and so this strategy is not computationally feasible here, as the reviewer remarks.

We have included some additional discussion about this issue in the conclusions section of our manuscript (section 5), in point 1 of our list of future recommendations for future directions and requirements to achieve better constraint. The added text is:

'Alternatively, representation errors could be reduced by increasing the temporal and spatial resolution of our model, and the effect of inter-annual variability could be reduced if we always compare the model to measurements for the correct year. In our calculations of model variant implausibility, inter-annual variability accounts for around 30 to 90% of the model-measurement comparison representation uncertainty. With smaller representation errors in the model-measurement comparison, we could more confidently reject implausible model variants. However, this approach is not currently computationally feasible.'

We appreciate the reviewer pointing out the importance of reducing the causes of model-measurement uncertainty and considering these when designing experiments. We are currently creating a PPE (on the NERC-funded A-CURE project) with output at a much higher temporal resolution that spans the period covered by multiple important flight campaigns.

Comment 2:

P6: At least some of the PM2.5 networks where sulfate concentrations were taken also have OC or OA. Why was OA from those sites not used (e.g. IMPROVE)?

For this study we have used all OC/OA measurement data that had been processed into a 'model-ready' format for comparison through the NERC GASSP project, and stored in the GASSP database, at the time of analysis. Unfortunately, this did not include OA data from some of the measurement networks like IMPROVE. More measurements are being added to the GASSP database and processed to a format for model comparison, and we anticipate that such data will be available for future work. In the next phases of our model-measurement comparison work, we will endeavour to ensure that more OA data from networks such as IMPROVE are incorporated to further strengthen our resulting comparisons with this variable.

Comment 3:

P12 L11-12: The incorrect model year relative the measurement year could also cause this mismatch, right? I could imagine this being a big issue for regions with significant biomass burning emissions, which have significant interannual variability.

Yes, the inter-annual variability could also potentially cause such a mismatch. This text in Section 2.4.3 has been adjusted to also include this possibility. It now reads:

*'...We assume that this large implausibility for the significant majority of variants indicates either there is a structural error in the model or that the model is unable to represent these point measurements because of its low spatial and temporal resolution. **An alternative explanation is a mismatch in the model's meteorological year to the year of the measurement** (section 2.4.1). We flag these measurements for further investigation of potential structural errors or underestimated error terms (these are not examined further in this study).'*

Comment 4:

Figure 6: I appreciate the challenge in creating this figure. Is there any way that it can be made more crisp. The combination of very small fonts and rough resolution makes it very hard to read (at least when printed).

We have generated an improved version of Figure 6 (in section 3.1) with a crisper resolution and as large as possible font size for the labelling within it.

Comment 5:

P18 L31: Need to say “northern-hemisphere winter” here (it’s said in the next sentence), but it should be clear when “winter” is first used.

We have corrected this in the manuscript. The sentence (near the beginning of Section 3.2) now reads:

‘Cloud pH is constrained more by AOD in ***northern-hemisphere*** winter (Figure 7a) when in-cloud oxidation of SO₂ by ozone dominates sulphate production.’

Comment 6:

P19 L13: Is nitrate not included at all in the simulations? It seems like that could throw things off a lot (which is I guess what you’re saying). It would be good to make it clear that nitrate was not included (if it’s not) or if it’s just incorrect.

The model does not contain a scheme for nitrate aerosol. This is stated in our description of the HadGEM3-UKCA climate model version we have used in Section 2.1. However, we agree with the reviewer that this is not clear at this point in the results section, and we have therefore revised the text to give more clarity in the point we are making. This text (in Section 3.2) now reads:

‘However, it may also indicate a structural deficiency. ***The low deposition rates in winter imply that PM2.5 has missing sources in winter but not in the summer, such as nitrate. Our model does not include aerosol nitrate, which (if included) would increase northern-hemisphere winter PM2.5 concentrations and weaken the constraint on dry deposition towards lower values in northern-hemisphere winter.***’

Have the authors looked at how much the observations impact the direct aerosol *effect* (the radiative effect of natural and anthro aerosols relative to no aerosols) rather than the radiative forcing? I wonder how different the level of constraint would be. This isn’t necessary for the paper unless the authors can do this quickly and agree that it’s interesting.

We had not looked into how our constraint using this large set of aerosol observations would impact the direct aerosol radiative effect (the radiative effect of natural and anthropogenic aerosols relative to no aerosols).

Figure 1 (below) shows the effect of the observational constraint using all measurement types on the distribution of the present-day global annual mean clear-sky aerosol radiative effect (RE; relative to no aerosol radiative effects). The distribution is constrained towards weaker (less negative) values and the 95% CI range ratio [Constrained/Unconstrained] is 0.53, which translates to a reduction in the 95% CI of 47% on constraint. This constraint is stronger than the constraints we obtain on the

pre-industrial to present-day radiative forcing (RF), which we expected as the RE is more closely related to our set of measurements (all aerosol properties) than RF, for which the uncertainty depends on both aerosol and cloud-related properties (Regayre et al, 2018).

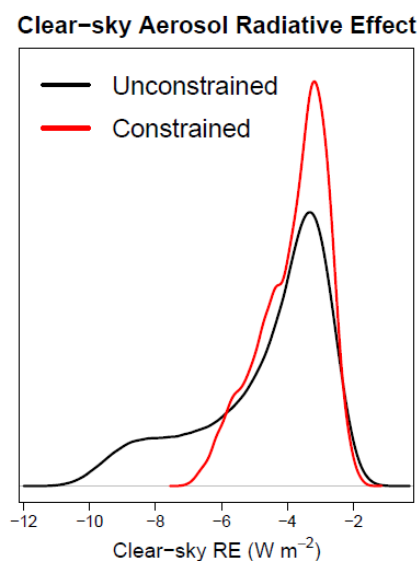


Figure 1: Effect of observational constraint using all measurement types on the probability distribution of the present-day global annual mean clear-sky aerosol radiative effect (relative to no aerosol radiative effects). The black line shows the prior (unconstrained) distribution and the red line shows the constrained distribution.

We have added the following text to the manuscript in Section 3.6 to reflect that the constraint is stronger for the direct aerosol radiative effect, however we do not include the figure as we want the focus of the paper to remain on the effect of the observational constraint on the radiative forcing.

‘Furthermore, we have found that the observational constraint on present-day (2008) global annual mean aerosol radiative effects (RE; relative to no aerosol radiative effects) is stronger than the constraint on aerosol radiative forcing. The uncertainty in the clear-sky aerosol RE is reduced by 47%. Industrial-period aerosol radiative forcing has distinct sources of uncertainty from aerosol radiative effects in the present-day atmosphere (Regayre et al., 2018), so a stronger constraint on present-day radiative effects is in line with expectations.’

References:

- Yoshioka, M., Regayre, L. A., Pringle, K. J., Johnson, J. S., Mann, G. W., Partridge, D. G., Sexton, D. M. H., Lister, G. M. S., Schutgens, N., Stier, P., Kipling, Z., Bellouin, N., Browse, J., Booth, B. B. B., Johnson, C. E., Johnson, B., Mollard, J. D. P., Lee, L. A. and Carslaw, K. S.: Ensembles of Global Climate Model Variants Designed for the Quantification and Constraint of Uncertainty in Aerosols and their Radiative Forcing, *J. Adv. Model. Earth Syst.*, 11(11), 3728–3754, doi:10.1029/2019MS001628, 2019.
- Regayre, L. A., Johnson, J. S., Yoshioka, M., Pringle, K. J., Sexton, D. M. H., Booth, B. B. B., Lee, L. A., Bellouin, N. and Carslaw, K. S.: Aerosol and physical atmosphere model parameters are both important sources of uncertainty in aerosol ERF, *Atmos. Chem. Phys.*, 18(13), 9975–10006, doi:10.5194/acp-18-9975-2018, 2018.

Response to reviewer 2 (Anonymous reviewer)

This manuscript investigates the constraint on aerosol forcing uncertainties by observations of AOD, PM_{2.5}, particle number concentrations, sulphate and organic mass concentrations. The manuscript is scientifically sound, and of a high scientific quality. In my opinion, the manuscript is ready for publication in ACP, since it presents not significant shortcomings in the scientific significance or quality.

We thank the reviewer for their support to our manuscript. As the reviewer has said our manuscript is ready for publication, no amendments have been made in response to this review.

Robust observational constraint of uncertain aerosol processes and emissions in a climate model and the effect on aerosol radiative forcing

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Abstract. The effect of observational constraint on the ranges of uncertain physical and chemical process parameters was explored in a global aerosol-climate model. The study uses 1 million variants of the HadGEM3-UKCA climate model that sample 26 sources of uncertainty, together with over 9000 monthly aggregated grid-box measurements of aerosol optical depth, PM_{2.5}, particle number concentrations, sulphate and organic mass concentrations. Despite many compensating effects in the model, the procedure constrains the probability distributions of parameters related to secondary organic aerosol, anthropogenic SO₂ emissions, residential emissions, sea spray emissions, dry deposition rates of SO₂ and aerosols, new particle formation, cloud droplet pH and the diameter of primary combustion particles. Observational constraint rules out nearly 98% of the model variants. On constraint, the $\pm 1\sigma$ (standard deviation) range of global annual mean direct radiative forcing, RF_{ari}, is reduced by 33% to -0.14 to -0.26 W m⁻², and the 95% credible interval (CI) is reduced by 34% to -0.1 to -0.32 W m⁻². For the global annual mean aerosol-cloud radiative forcing, RF_{aci}, the $\pm 1\sigma$ range is reduced by 7% to -1.66 to -2.48 W m⁻², and the 95% CI by 6% to -1.28 to -2.88 W m⁻². The tightness of the constraint is limited by parameter cancellation effects (model equifinality) as well as the large and poorly defined ‘representativeness error’ associated with comparing point measurements with a global model. The constraint could also be narrowed if model structural errors that prevent simultaneous agreement with different

measurement types in multiple locations and seasons could be improved. For example, constraints using either sulphate or PM_{2.5} measurements individually result in $RF_{\text{ari}} \pm 1\sigma$ ranges that only just overlap, which shows that emergent constraints based on one measurement type may be over-confident.

1 Introduction

5 Global model simulations of aerosols and their climatic effects are very uncertain. Different global aerosol models have large spread in their simulations of aerosol microphysics, radiation and forcing (Mann et al., 2014; Myhre et al., 2013; Shindell et al., 2013; Tsigaridis et al., 2014). This multi-model spread can be due to different model structures, missing processes, parameter settings, algorithms or coding errors. Individual climate models are also very uncertain because the values of parameters related to physical processes and emissions are often poorly defined (Johnson et al., 2018; Lee et al., 2011b, 2013; 10 Regayre et al., 2018). The uncertainty in the aerosol effective radiative forcing (ERF) over the industrial period caused by aerosol processes, physical atmosphere model processes and emissions could be as large as the multi-model spread (Johnson et al., 2018; Regayre et al., 2014, 2018).

There are two main methods to reduce model uncertainty, often called bottom-up and top-down approaches. The bottom-up 15 approach involves improving the representation of model processes and refining estimates of the associated parameter values through experiment and theory. This approach is necessary to improve model fidelity, but it does not provide an estimate of the model uncertainty, and the uncertainty may grow if the increase in model complexity requires a large number of new and poorly defined parameters. To reduce model uncertainty, bottom-up model development needs to be combined with top-down approaches in which numerous uncertain process-related parameters and emissions are adjusted to improve the agreement of 20 models with measurements.

The difficulty with top-down model adjustments (in its simplest form, model tuning) is that the uncertainty stems from large combinations of uncertain model input parameters. This means that the adjustment of small sets of parameters to improve model agreement with measurements will not produce robust results (Carslaw et al., 2018). For example, a model simulation 25 of particle concentrations could be improved by adjusting particle formation rates, but many other combinations of parameters related to emissions, chemistry or deposition might be able to achieve similar model skill (Carslaw et al., 2013b). Models that are narrowly tuned in this way can therefore produce a wide range of results when used to make predictions outside the range of conditions under which they were tuned. This is likely to be a cause of the large uncertainty in aerosol radiative forcing, which is a predicted rather than observable quantity.

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If other aerosol-climate models are comparable with our own HadGEM-UKCA model, then they contain at least 20 important uncertain parameters related to emissions and processes, although fewer than about 10 parameters will dominate the

uncertainty in a particular model variable in any one environment and time of year (Lee et al., 2016; Regayre et al., 2014, 2018). Therefore, to define and reduce the model uncertainty it is necessary to find from within 10 dimensions of parameter space all the parameter combinations that produce plausible agreement with different aerosol properties observed across all seasons and global environments. A single well-configured version of a model produced by parameter tuning tells us nothing about the combinations of parameter values that can achieve consistency with measurements within their uncertainty range, nor does it tell us anything about the model output uncertainty.

In this paper we address the question: To what extent do extensive and diverse aerosol measurements enable the plausible range of model parameters to be constrained if the full range of their compensating effects is accounted for? By ‘constrain’ we mean a narrowing of the probability distribution of a parameter (and potentially the absolute range) compared to the uncertainty range that was assumed when the model was built. We also quantify how the identification of observationally plausible parameter ranges feeds through to a reduction in the uncertainty in predictions of aerosol radiative forcing over the industrial period. The study focuses on model constraint using measurements of aerosol properties rather than cloud properties, therefore we emphasise the effect on aerosol-radiation interaction forcing rather than aerosol-cloud interaction.

2 Methods

2.1 The HadGEM3-UKCA climate model

We use the Global Atmosphere 4 (GA 4.0; Walters et al., 2014) configuration of the Hadley Centre General Environment Model version 3 (HadGEM3; Hewitt et al., 2011), which incorporates the UK Chemistry and Aerosol (UKCA) model at version 8.4 of the UK Met Office's Unified Model (UM). UKCA simulates trace gas chemistry and the evolution of the aerosol particle size distribution and chemical composition using the GLObal Model of Aerosol Processes (GLOMAP-mode; Mann et al., 2010) and a whole-atmosphere chemistry scheme (Morgenstern et al., 2009; O'Connor et al., 2014). The model has a horizontal resolution of 1.25x1.875 degrees and 85 vertical levels.

The aerosol size distribution is defined by seven log-normal modes: one soluble nucleation mode as well as soluble and insoluble Aitken, accumulation and coarse modes. The aerosol chemical components are sulphate, sea salt, black carbon (BC), organic carbon (OC) and dust. The model does not include any representation of nitrate aerosols. Secondary organic aerosol (SOA) material is produced from the first stage oxidation products of biogenic monoterpenes under the assumption of zero vapour pressure. SOA is combined with primary particulate organic matter after kinetic condensation.

GLOMAP simulates new particle formation, coagulation, gas-to-particle transfer, cloud processing and deposition of gases and aerosols. The activation of aerosols into cloud droplets is calculated using globally prescribed distributions of sub-grid vertical velocities (West et al., 2014) and the removal of cloud droplets by autoconversion to rain is calculated by the host

model. Aerosols are also removed by impaction scavenging of falling raindrops according to the parametrisation of clouds and precipitation collocation (Boutle et al., 2014; Lebsock et al., 2013). Aerosol water uptake efficiency is determined by kappa-Kohler theory (Petters and Kreidenweis, 2007) using composition-dependent hygroscopicity factors.

5 Anthropogenic emission scenarios prepared for the Atmospheric Chemistry and Climate Model Inter-comparison Project (ACCMIP) and prescribed in some of the CMIP Phase 5 experiments are used here. Biomass burning emissions for recent decades were prescribed using a ten year average of 2002 to 2011 monthly mean data from the Global Fire and Emissions Database (GFED3; van der Werf et al., 2010) and according to Lamarque et al. (2010) for 1850. Volcanic SO₂ emissions are prescribed in the model by combining emissions from the Andres and Kasgnoc (1998) dataset for continuously erupting and
10 sporadically erupting volcanoes and the Halmer et al. (2002) dataset for explosive volcanoes.

A full description of the set-up for our model simulations can be found in Yoshioka et al. (2019), which we summarise here. The base model simulation was subject to a multi-year spin-up period. Parameter perturbations were then applied distinctly to individual ensemble members (which branch from the base model) and spun-up for a further month. We then ran each
15 simulation for a further 12 months to produce the data used here. Horizontal winds and temperatures in the simulations are nudged towards European Centre for Medium-Range Weather Forecasts (ECMWF) ERA-Interim reanalyses for 2008 between approximately 1.2 and 80 km using a 6-hour relaxation timescale. Nudging means that pairs of simulations have identical synoptic-scale features, which enables the effects of perturbations to aerosol and chemical processes to be quantified using single-year simulations, although the magnitude of forcing will vary with the chosen year (Fiedler et al., 2019; Yoshioka et
20 al., 2019).

2.2 Creation of perturbed parameter model variants

Our method to determine observational constraint on the model parameters and radiative forcings involves producing a very large set of ‘model variants’, each with a different combination of parameter values, and then ruling out model variants for which a set of model outputs are judged to be implausible against measurements (see section 2.4). The model variants were
25 generated using a perturbed parameter ensemble (PPE) of 235 model simulations of HadGEM3-UKCA (the ‘AER PPE’ detailed in Yoshioka et al., 2019) that samples 26 sources of uncertainty in the aerosol model (Carslaw et al., 2017; Yoshioka et al., 2019) – see Table A1 in Appendix A.

A set of 235 simulations alone is much too small to allow statistical analysis of model performance across 26 dimensions of parameter space. We therefore built Gaussian Process emulators (surrogate models) using the PPE simulations as training data
30 (Lee et al., 2011b), which define how the model outputs vary continuously over the 26-dimensional parameter space and enable dense sampling over parameter uncertainty. Separate emulators were built describing the monthly mean value of each model output in each model grid cell. We then used Monte Carlo sampling from these emulators to produce output for a set of 1

million model variants (parameter input combinations). Uniform distributions were assumed for each parameter in this sampling. The emulator is not a perfect representation of a model output, but its uncertainty can be estimated and accounted for in the model-measurement comparison. In the rest of this paper we refer to the emulator-derived values of model outputs at each sampled 26-d input combination as a ‘model variant’.

- 5 The AER PPE samples only uncertainties in the aerosol component of the model and the radiative forcing does not account for atmospheric and cloud adjustments – i.e., it is a radiative forcing (RF) rather than an effective radiative forcing, which we analysed in previous papers (Johnson et al., 2018; Regayre et al., 2018). The prior (unconstrained) 95% credible interval of global mean aerosol RF is $-2.23 \pm 0.94 \text{ W m}^{-2}$. However, because of the way that multiple parameters compensate (Lee et al., 2016; Regayre et al., 2018), the forcing uncertainty in this PPE is similar to the AER-ATM PPE in which additional physical
10 atmosphere model parameters were perturbed and cloud adjustments accounted for (Yoshioka et al., 2019). Because the AER PPE analysed here samples only aerosol uncertainties, we restrict the constraints to measurements of aerosol properties. In future work we will extend the analysis to radiation, precipitation and cloud measurements that are relevant to the wider range of parameters in the AER-ATM PPE.

- The choice of the 26 perturbed parameters and their uncertainty ranges were defined using expert elicitation (Yoshioka et al.,
15 2019). The parameters (Table A1; full descriptions given in Yoshioka et al., 2019) relate to natural and anthropogenic emission fluxes of aerosol precursor gases and primary particles, the properties of primary particles (size), aerosol processes, aerosol hygroscopicity, removal rates and cloud droplet formation (updraft speed). The list of parameters is not exhaustive, but one-at-a-time parameter perturbation tests were used to show that any other parameters have a smaller effect regionally and globally in our model than the set we chose. Finally, we note that the evaluated uncertainty in global annual mean RF in this study
20 differs from that shown in Yoshioka et al. (2019) as we have used uniform parameter distributions when sampling over the parameter uncertainty space, while elicited parameter distributions were used in Yoshioka et al. (2019). Our choice to use uniform distributions here means that the constraint can be fully attributed to the model-measurement comparison.

2.3 Measurements

- We use aerosol measurements from ground stations, ship campaigns and aircraft campaigns covering the following aerosol
25 properties: aerosol optical depth (AOD), PM_{2.5} concentrations, sulphate mass concentrations, organic carbon mass concentrations, and number concentrations of particles larger than 3 nm dry diameter (N_3) and 50 nm dry diameter (N_{50}) – see Appendix B and Table S1 in the supplementary data file. All measurements used are from within the boundary layer, which we define to be at an atmospheric pressure greater than 800hPa. We do not attempt to constrain aerosol properties above the boundary layer.

- 30 The measurements were all made at specific locations and times (i.e. they are ‘point measurements’) in the period from October 1995 to December 2015, and we use measurements from all years within this period regardless of whether the year of the

measurement matches the year of the PPE model simulations. (We take account of the inter-annual differences by incorporating an error term in the constraint process, see Section 2.4). The measurements were aggregated to monthly mean values in grid cells of size 2.50° longitude by 3.75° latitude (4 model grid boxes of the N96 model grid). In cases where there is more than one measurement in a model grid cell, the observed values were averaged. This processing resulted in 9464 monthly-
5 aggregated grid-box measurements (over 6 aerosol properties and 12 months). Figure 1 shows the global spatial coverage of the gridded measurements for each aerosol property, along with the monthly temporal coverage for each measurement, which is indicated by the colour-scale. Table 1 shows the breakdown of the number of grid-box measurements by variable and month.

The AOD data are level 2.0 (quality assured) monthly-mean data at 440 nm wavelength from the AERONET (Aerosol Robotic
10 Network) network (Giles et al., 2019; Holben et al., 1998). Our dataset includes an average of 312 aggregated grid-box measurements for comparison in each month. Figure 1 shows that the measurements are well distributed across all continental regions except Antarctica. The coverage at high northern latitudes is relatively sparse, and there are only a small number of island measurement that are representative of marine aerosol environments. The temporal coverage is very good, with the majority of stations providing measurements in all months of the year.

15 The PM_{2.5} and sulphate concentration data come from multiple large networks. The sulphate concentration data are from the Interagency Monitoring of Protected Visual Environments (IMPROVE) network (USA), the European Monitoring and Evaluation Programme (EMEP) network and the Acid Deposition Monitoring Network in East Asia (EANET). For PM_{2.5}, we use data from the IMPROVE network, the World Data Centre for Aerosols (WDCA) (European sites), the Asia-Pacific
20 Aerosol Database (A-PAD) and the Canadian National Air Pollution Surveillance Program (NAPS). Other PM_{2.5} measurements are included from smaller networks and individual stations in Australia, South America, Taiwan and South Africa, as well as sulphate and PM_{2.5} data recorded at the Station for Observing Regional Processes of the Earth System (SORPES) in Nanjing, East China. The PM_{2.5} data (except for the SORPES site) were obtained, processed and gridded to the N96 model grid as described in Browse et al. (2019). Figure 1 shows that these PM_{2.5} and sulphate measurements are highly
25 clustered over polluted land areas of the Northern hemisphere, mostly in North America and Europe with limited coverage elsewhere, especially in remote and marine areas. Nearly all stations in these data sets have full temporal coverage, leading to approx. 150 and 170 aggregated grid-box measurements for comparison in each month for PM_{2.5} and sulphate respectively (Table 1).

For N₅₀ particle concentrations and OC concentrations we have a mixture of measurements from a small number of land-based ground stations along with measurements taken over marine environments from ship and aircraft campaigns (see Table A2).
30 The N₅₀ concentration data were mainly derived from size distribution measurements and gridded to the N96 model grid as described in Browse et al. (2019). The amount of campaign data, and hence global spatial coverage in the gridded data, is greater for N₅₀ than for OC (Figure 1), and the number of aggregated grid-box measurements is variable between months

(Table 1). Due to the nature of field campaigns, the temporal coverage is much sparser for these variables, with each campaign only measuring for 1-3 months of the year, shown by the blue colours for the data of these variables in Figure 1.

The measurement data for N₃ particle concentration has the smallest number of grid-box measurements over the year and spatially is the sparsest data set included here. The data for this aerosol property comes from only 12/13 ground stations (ACTRIS; Asmi et al., 2013), which are mostly located in Europe, with one in the Arctic, one in Antarctica and one in Northern India. The N₃ concentrations at each site were derived directly by integrating size distribution measurements. This data was then averaged over multiple years for each month and location by the authors.

	AOD	Sulphate	PM2.5	OC	N ₃	N ₅₀
Jan	294	149	168	6	13	77
Feb	301	148	168	14	13	90
Mar	309	151	170	82	13	148
Apr	316	151	170	74	12	199
May	322	149	167	23	12	64
Jun	320	150	170	23	12	96
Jul	323	148	172	23	13	115
Aug	326	148	169	23	13	109
Sep	321	147	166	22	13	133
Oct	315	147	165	41	13	119
Nov	309	146	168	37	13	155
Dec	298	147	169	15	12	67
Total	3754	1781	2022	383	152	1372

Table 1. The number of monthly aggregated grid-box measurements for each variable in each month. The total number over all months and all variables is 9464.

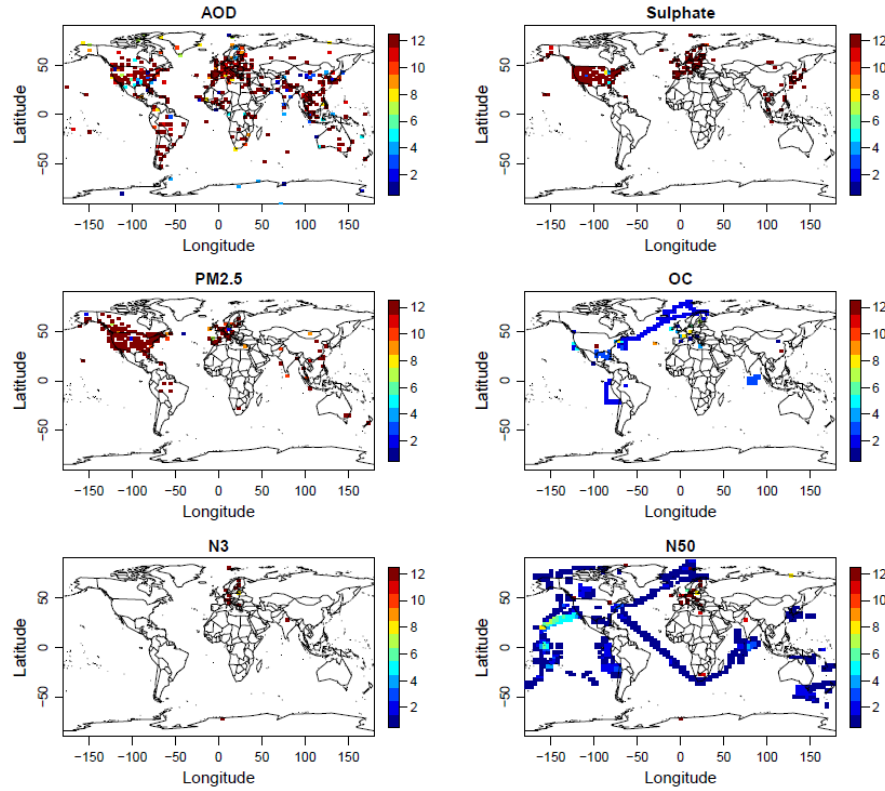


Figure 1. The distribution of measurements used in the constraint. The colours indicate the number of months covered by the measurements (although the data may not cover all days within a month).

2.4 Constraint methodology

We apply the statistical methodology of history matching, which has been applied to complex models in a range of fields, including epidemiological modelling of virus transmission (Andrianakis et al., 2017), risk assessment for oil field developments (Craig et al., 1997), modelling galaxy formation (Rodrigues et al., 2017) and climate modelling (Edwards et al., 2011; McNeill et al., 2016; Williamson et al., 2013). The methodology is described in detail in previous papers (Johnson et al., 2018; Regayre et al., 2018), which built upon our earlier study (Lee et al., 2016). We therefore describe the overall methodology only briefly here, but present a full description of the new aspects related to using real measurements rather than ‘synthetic’ measurements (Johnson et al., 2018).

In the comparison of the model and measurements we account for emulator uncertainty, measurement uncertainty (instrument error), representativeness uncertainties (caused by spatial and temporal mismatches in resolution and sampling between model and measurements), and potential structural model uncertainty. The model-measurement difference together with these measures of uncertainty is incorporated into an ‘implausibility measure’ and our model constraint procedure in order to identify implausible parts of parameter space (model variants).

2.4.1 Implausibility measure

The implausibility metric $I(\mathbf{x})$ is calculated for each of the 1 million model variants \mathbf{x} , for each gridded measurement. $I(\mathbf{x})$ weights the difference between the model and measurements by the uncertainties associated with the comparison (Craig et al., 1996; Williamson et al., 2013):

$$I(\mathbf{x}) = \frac{|M - O|}{\sqrt{[Var(M) + Var(O) + Var(R) + Var(S)]}}, \quad (1)$$

- 5 where M is the estimate of model output calculated using the emulator and O is the observed value (the measurement). In the denominator $Var(M)$ is the variance in the model estimate (associated with replacing the model with the emulator), $Var(O)$ is the variance in the measurement (i.e., instrument or retrieval uncertainty), $Var(R)$ is the variance associated with the comparison of the model with the measurements, called the representativeness error (Schutgens et al., 2017, 2016a, 2016b), and $Var(S)$ is a model structural error term.
- 10 A low value of the implausibility metric indicates either the model-measurement difference is small (i.e., the model is skilful) or that the uncertainty in the denominator is large (i.e., we cannot tell whether the model is skilful because the uncertainties are too large). Therefore, the implausibility metric allows model variants to be ruled out if the model-measurement difference is large and we can be confident that it is large.

- The representativeness error $Var(R)$ has three components. $Var(R_{sp})$ (sp = spatial) accounts for uncertainty associated with spatial variability below the grid scale of the model, which means that a point measurement may not be representative of the grid-box mean (Schutgens et al., 2016b). $Var(R_{temp})$ ($temp$ = temporal) accounts for the temporal sampling of a measurement, which may not match the temporal sampling of the model (e.g. a ship track through the grid-box over a short time period which is compared with a monthly-mean model value (Schutgens et al., 2016a). $Var(R_{iav})$ (iav = inter-annual variability) accounts for the fact that we sometimes match measurements and the model for the correct calendar month but not for the correct year.
- 20 This is necessary in cases where we use measurements from years for which we have not run the model. We assume that

$$Var(R) = Var(R_{sp}) + Var(R_{temp}) + Var(R_{iav}) \quad (2)$$

The magnitude of these errors is discussed in section 2.4.2.

- The structural error term $Var(S)$ has been included in previous studies using the implausibility metric. It is intended to represent an estimate of the potential structural error in the model. Practically, however, we have no way to estimate this term for all variables at all times and geographical locations. We therefore set it to zero, and instead use very large values of implausibility to point us towards potential structural errors in the model-observation comparison and constraint procedure, as
- 25 described in section 2.4.3.

2.4.2 Estimation of uncertainty terms

Our estimates of the uncertainty terms in Equation 1 are preliminary and are designed to test our approach. We discuss in the conclusions the need to refine our understanding of these uncertainty terms.

- 5 For all aerosol properties we assume an instrument uncertainty of 10%, a spatial co-location uncertainty of 20%, and a temporal sampling uncertainty of 10% on the measured value. The spatial sampling uncertainty for monthly mean aerosol properties is estimated based on Schutgens et al. (2017, 2016b). These studies examined a typical spatially heterogeneous continental environment where the sampling error is dominated mainly by local aerosol sources that are not resolved by the global model. The magnitude of uncertainty is likely to vary globally (especially between land and ocean), with surface measurements typically having larger errors than column measurements and the magnitude of error also depending on the location of a ground site with respect to the grid-box centre, but we do not account for these variations. We base our estimate of the temporal sampling uncertainty on Schutgens et al. (2016a) who quantified the error associated with the different temporal sampling of models and measurements (e.g., daily measurements or temporally sporadic measurements versus monthly mean model, etc.). The emulator uncertainty is taken from the Gaussian error on the emulator mean prediction, which is known for every parameter combination (i.e., each of the 1 million model variants).

- The inter-annual uncertainty was defined to be the standard deviation of monthly mean aerosol properties in each grid cell over a 30-year period. We take information from an analysis of the trend and variation of gridded aerosol properties in a HadGEM3-UKCA hindcast simulation over the period of 1980-2009 (Turnock et al., 2015). For each month and grid-box, the monthly mean output of the aerosol variable of interest for each year of the simulation was obtained. These values were de-trended using linear regression and the resulting residuals were then analysed. We use a relative measure of monthly mean uncertainty defined by the standard deviation of these residuals divided by the de-trended mean. As an example, Figure 2 shows the relative standard deviation for the surface-level N_{50} concentration in July.

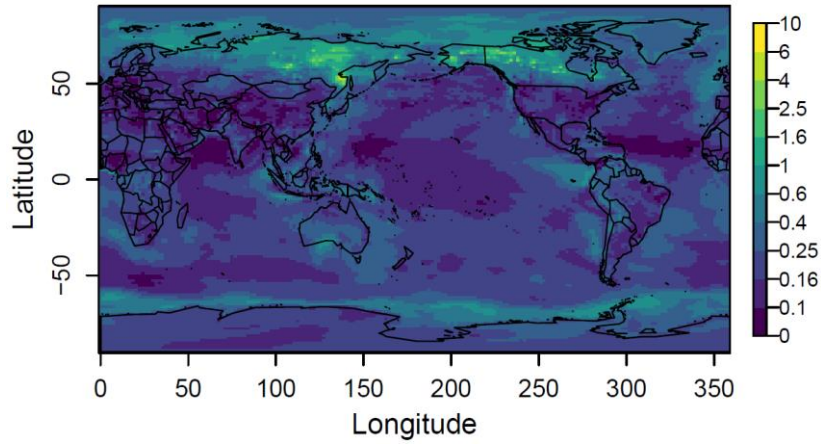


Figure 2. The relative standard deviation for surface-level N_{50} conc. in July, used in the estimation of the inter-annual variability component of representativeness error R_{lav} .

2.4.3 Methodology for ruling out observationally implausible parts of parameter space

5 There is an element of subjectivity involved in comparing a model with point measurements and reaching a conclusion about the fidelity of the model. The comparison may indicate either: a) the model seems to be structurally adequate, but the parameters need to be adjusted to optimise agreement, or b) the model is structurally deficient (i.e., there are missing or incorrect process representations in the model). Structural deficiencies may be apparent, for example, because the model skill is particularly poor in one region or at one time of year, or it is not possible to obtain good skill across multiple variables simultaneously.

Our use of 1 million model variants and more than 9000 monthly aggregated grid-box measurements means that we need to automate the model-measurement comparison processes and detection of potential structural errors while also using the measurements to rule out implausible parts of parameter space. The difficulties for us in detecting structural errors are: a) we cannot inspect each of the 1 million model variants individually, so we need to rely on summary statistics; b) many of the aerosol point measurements are spatially and temporally sparse, so we cannot easily detect spatial and temporal changes in model skill that might indicate structural error; c) the measurements do not have the same spatial distribution in all months (because of brief, localised field campaigns) so spatial-temporal biases are hard to detect; d) the uncertainty in each measurement (particularly the representativeness error, section 2.4.2) is spatially and temporally heterogeneous and often very poorly defined.

20 Our approach is summarised in Figure 3. It is designed to rule out implausible parts of parameter space while avoiding doing so in cases where the biases shared by many model variants could be caused by structural errors in the model.

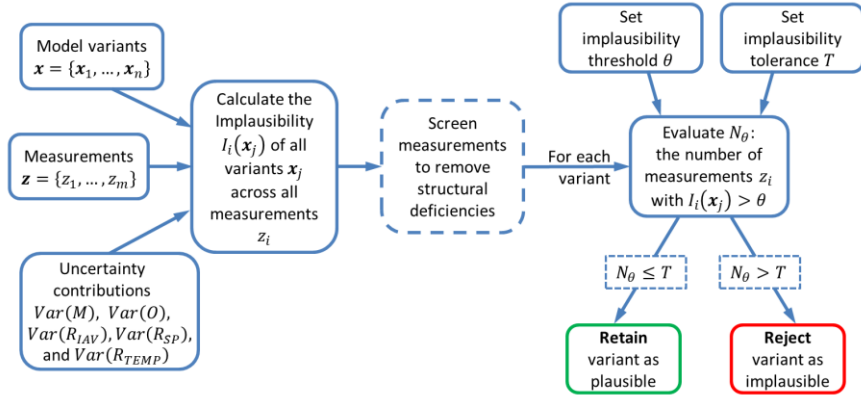


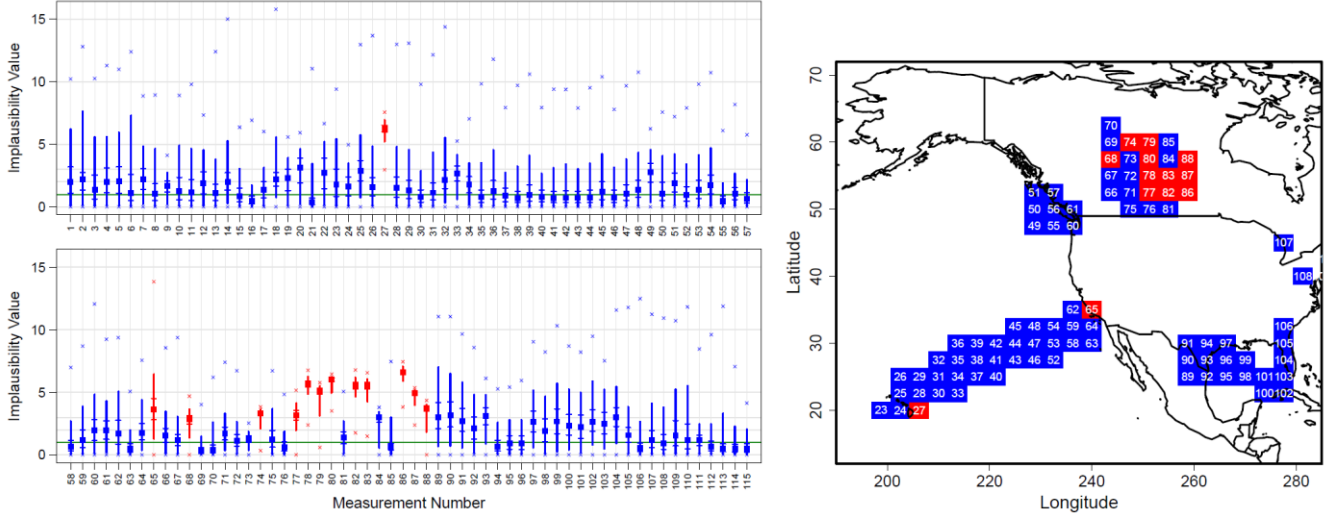
Figure 3. Flow chart detailing the process followed for each model variant x_j , in using the calculated implausibility over a set of m measurements $z = \{z_1, z_2, \dots, z_m\}$ (for a single output variable y) simultaneously to constrain the model uncertainty.

The steps are:

- 1) The implausibility is quantified for each of the 1 million variants across all measurements of a single type in a particular month. Figure 4 shows an example for the measurements of N_{50} in July. For each measurement (numbered on the horizontal axis, left plot), the distribution of the implausibility over the variants is shown by the bar representing the 95% credible interval.
- 2) Measurements are identified for which 97.5% of the model variants have an implausibility $I > 1$. These measurements are excluded from the constraint procedure (shown in red in Figure 4). We assume that this large implausibility for the significant majority of variants indicates either there is a structural error in the model or that the model is unable to represent these point measurements because of its low spatial and temporal resolution. An alternative explanation is a mismatch in the model's meteorological year to the year of the measurement (section 2.4.1). We flag these measurements for further investigation of potential structural errors or underestimated error terms (these are not examined further in this study).
- 3) Using all other measurements, (where more model variants have lower implausibility, shown in blue in Figure 4), we use the implausibility metric values to decide whether to rule each variant out as implausible, or retain it as plausible. If we ruled out all model variants with high implausibility for each measurement in turn (treating the measurements independently, as in many emergent constraint studies), we could end up ruling out all parts of parameter space. Our criterion is therefore to rule out a model variant if more than a defined fraction (or number) of the measurements (tolerance T) exceeds a defined implausibility threshold (θ). For example, we might rule out a model variant if more than 20% of measurements exceed an implausibility of 3.5 (i.e., bias is 3.5 times the expected error).

We apply this approach to the set of measurements for each variable (measurement type) in each month and then combine the constraints to a joint constraint over months and/or over variables such that if a variant is ruled out for any single month/variable combination, then it is also ruled out in the joint constraint. This method allows us to identify the set of model variants that

capably represent measurements of a range of variables and across multiple locations and seasons. We extensively explored various choices of the tolerance and threshold values in each variable/month case and found that the final constrained parameter ranges were reasonably robust, except when the number of measurements was small.



- 5 **Figure 4.** Left: The distribution of implausibility calculated over the 1 million model variants for each measurement in the July N_{50} concentration set, shown vertically. For each measurement, numbered along the x-axis, the range of the implausibility distribution is shown by the outer crosses, the bar corresponds to the 95% credible interval (2.5% to 97.5% empirical quantiles), the horizontal markers through the bar show the inter-quartile range, and the square point is the median implausibility. Here we assume no structural error term in the implausibility calculations and use the implausibility distribution to identify potential structural errors. Measurements coloured red are ruled out as potential structural error cases (as the lower 95% credible interval bound is >1), and those coloured blue are retained and used in our constraint procedure. Right: Corresponding map to show the locations of the rejected July N_{50} measurements (red) and those retained for constraint (blue), over the North Pacific and North America region (outside this region, all measurements were retained). We hypothesise that the red points over the Pacific correspond to ports with localised pollution sources while the red points over Canada correspond to localised fire emissions that are not represented at the resolution of the model.
- 10
- 15 Our choices of the threshold and tolerance for each measurement type are given in Table A2 in Appendix A. A wide range of values were tested in each case, starting with a set threshold of $\theta = 3.5$ and iterating through increasing tolerances T up to a maximum of $T = 33\%$ (1/3 of the measurements), before further increasing θ by 0.5 (to a maximum of $\theta = 4.5$) and re-iterating over T in order to retain (approximately) a chosen percentage of model variants. Approximately the same percentage of variants was attained for all months of a variable type and combined for an ‘all months’ constraint. Our final choices for each variable
- 20 type on its own (left column in Table A2) were relaxed for the joint all-variables-months constraint (retaining a larger percentage of variants in each month for each variable, so a weaker constraint; right column in Table A2), in order to retain a reasonable number of model variants and avoid over-constraining on any one observational type.

Our assumption of zero structural error ($Var(S) = 0$) in the implausibility calculations means that structural errors in the model can easily come to light in our constraint process. This occurs either in the calculated implausibility values for a

measurement (where large values are consistently produced over the 1 million variants covering the model uncertainty, indicating a large model-measurement discrepancy, e.g. Figure 4), or when bringing together the constraint effects of different sets / types of measurements (where very few, if any, model variants that lead to plausible model output in all cases/measurement types simultaneously can be identified and retained). Even though we do not directly account for structural errors in the implausibility measure itself, our constraint approach offsets the effects of such errors on the achieved constraint as much as possible. This is accomplished by screening out observations with large model-measurement discrepancies from the constraint process (step 2; Figure 4) and by relaxing the constraint criteria for the joint all-variables-months constraint. Through this approach we are able to produce as robust a constraint as possible, given the limitations we have in specifying structural and representational errors.

10 2.5 Interpretation of constrained parameter probability distributions

Observationally plausible parts of parameter space exist in 26 dimensions. We show the results as *1-dimensional marginal probability distributions*, which are 1-dimensional projections of the 26-dimensional parameter probability distribution. Figure 5 shows an idealised representation for a 2-dimensional parameter constraint. The white parts of the joint distribution are ruled out, leaving the shaded region of joint parameter space as observationally plausible. The effect on the marginal probability distribution of parameter 1 is to entirely rule out the lowest and highest values (i.e., there is no combination of these values of parameter 1 with parameter 2 that produces an observationally plausible model). Where some values of parameter 1 are ruled out over the range of parameter 2, the likelihood of parameter 1 having those values is reduced.

In the results below, the parameter probability distributions therefore reflect the relative likelihood of the parameter having particular values, with lower probabilities indicating that there are fewer ways in which the parameter can be combined with the other 25 parameters to produce a plausible model. For conciseness in the results section we say, for example, that “a measurement constrains the parameter to low values”, which means that we retain a larger proportion of model variants with low values.

Figure 5 also shows the separate and joint effects of two observational constraints. We show this conceptually because it arises in the results. Measurement 1 rules out the lowest values of parameter 1 and suggests that parameter 1 is likely to be at the high end of the sampled range. Conversely, measurement 2 suggests that parameter 1 is likely to be at the low end of the range. However, the correct interpretation of this situation is that intermediate values of the parameter are consistent with *both* measurements (measurement 1 is consistent with the model for all but the lowest values of the parameter and measurement 2 is consistent for all but the highest values). Only in cases where the two separate constrained parameter pdfs do not overlap can we conclude with certainty that there is likely to be a structural deficiency in the model. However, to obtain multivariate constraint we prevent this happening by screening out measurements with large model-measurement discrepancies and relaxing the constraint criteria with each measurement type.

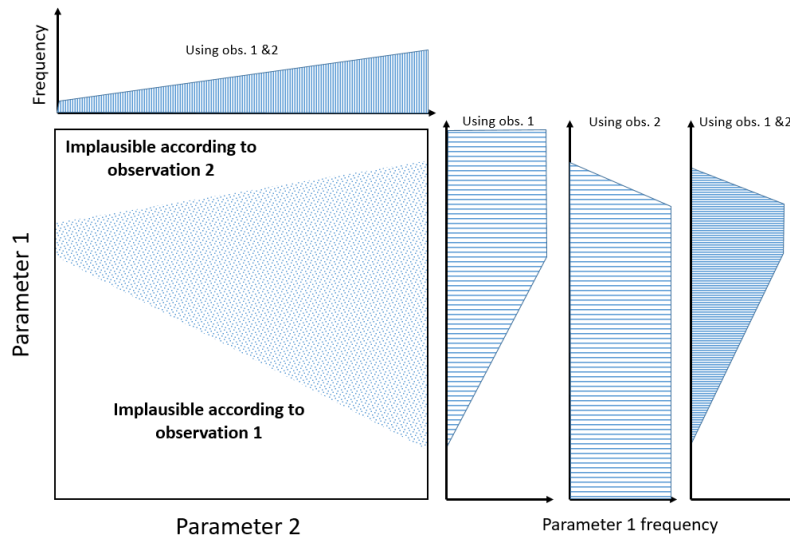


Figure 5. Schematic of parameter constraint in two dimensions using two measurements.

3 Results

3.1 Constraint using individual measurement types

Figure 6 shows the constrained marginal parameter distributions for all parameters based on using individual measurement types (each column on left) and all measurement types together (right column).

AOD measurements constrain aerosol and precursor emissions to low values and removal rates to high values. These constraints imply that the PPE produces generally too-high AODs across the sampled parameter space, which is the case (section 3.4). In particular, sea spray emissions higher than about 3.6 times the baseline emissions are ruled out, but emissions down to as low as 0.125 times the baseline emissions are plausible. For anthropogenic SO₂ emissions, the likelihood of the emissions scale factor being below 1 (corresponding to the default value from the inventory) increases from 55% to 70% on constraint. For BVOC emissions, the likelihood of the emissions (or effectively the production of SOA) being more than 3 times the default emission of 46 Tg/y (=138 Tg/y) is reduced from 31% to 13%, but all lower values from the default emission down to our lower bound of 37 Tg/y are equally plausible.

The AOD measurements also constrain the model to low values of other parameters: more variants with higher cloud droplet pH values are ruled out (judged implausible) and as a result cloud droplet pH is nearly 3 times as likely to be below the central value of its range of 5.8 as above it, which is consistent with a higher likelihood of slower production of sulphate aerosol from in-cloud SO₂ oxidation. The hygroscopicity of OC in the particles (κ_{OC}) is also weakly constrained to low values, which reduces the water content of aerosol and reduces AOD. The rate of aerosol scavenging by precipitating raindrops (the Rain_Frac parameter) is weakly constrained to high values.

Sulphate measurements strongly constrain SO_2 emissions to low values, which is consistent with the AOD constraint. Given this constraint, the SO_2 emissions have a 78% likelihood of being below the default value from the inventory and the median emission is reduced to 0.78 times the default. Also consistent with the AOD constraint, the deposition rate of accumulation mode particles is constrained strongly to high values, with an 87% likelihood of the rate being above the default value.

5 Likewise, the SO_2 dry deposition rate is constrained fairly weakly to higher values, with a 60% likelihood of the scaled value being above the default value. Each of these constraints is consistent with too-high sulphate concentrations in many of the sampled model variants across the parameter uncertainty space (section 3.4).

PM_{2.5} measurements have a similar effect to AOD on some parameters, but for others there are differences. Emissions of sea spray and BVOC emissions are constrained similarly (to low values). However, SO_2 emissions and cloud droplet pH are

10 weakly constrained to higher values and the dry deposition rate of accumulation mode particles is weakly constrained to low values, opposite to the AOD and sulphate constraints for these parameters. The PM_{2.5} measurements also weakly constrain the residential combustion emissions to high values. PM_{2.5} and AOD are strongly correlated in the PPE (Johnson et al., 2018), so differences in the constrained parameters most likely reflect differences in the spatial distribution of the measurements (Figure 1) and how that maps on to the spatial variations in sensitive parameters. As described in section 2.5, these apparently

15 opposing constraints are not necessarily inconsistent: for AOD and PM_{2.5} there may be other parameter settings that can be combined with low SO_2 emissions to achieve agreement with the measurements (so the space is not ruled out).

Organic carbon (OC) measurements strongly constrain the scaled magnitude of residential carbonaceous emissions to a narrower credible interval of about 0.3-1.8 centred near the default value specified in the emissions. Emissions above 2.0 times the default value are effectively ruled out and there is only a 13% likelihood of the emissions being below half the default

20 value. Fossil fuel emissions have a 70% likelihood of being above the default emission value. The OC measurements also constrain the scaled BVOC emissions in a similar way to PM_{2.5} and AOD, with scaled emissions above about 2.1 times (97 Tg/y) having only a 31% likelihood (compared to 50% prior to constraint). OC measurements also constrain the lowest values of BVOC emissions, which was not achieved with PM_{2.5} and AOD. The likelihood of the scaled emissions being below 1 (46 Tg/y) is 6% (compared with 11%). The dry deposition rate of Aitken mode particles is constrained to the low part of the range,

25 which will tend to increase OC concentrations in the atmosphere consistent with the constraint of fossil fuel emissions to high values. There is also a weak constraint of the ageing rate towards higher values, which has a 55% likelihood of being in the upper half of the range. The rate of aerosol scavenging by precipitating raindrops (Rain_Frac parameter) is constrained similarly, but to lower values. Again, although weak, these two constraints imply slower ageing, slower removal rates, longer OC lifetime and higher atmospheric concentrations. Biomass burning emissions are only very weakly constrained towards

30 lower emissions. The lack of constraint on the biomass burning emissions from OC measurements here is likely a result of the limited coverage, if any, of the OC measurements in regions important for biomass burning such as Africa and South East Asia (Figure 1).

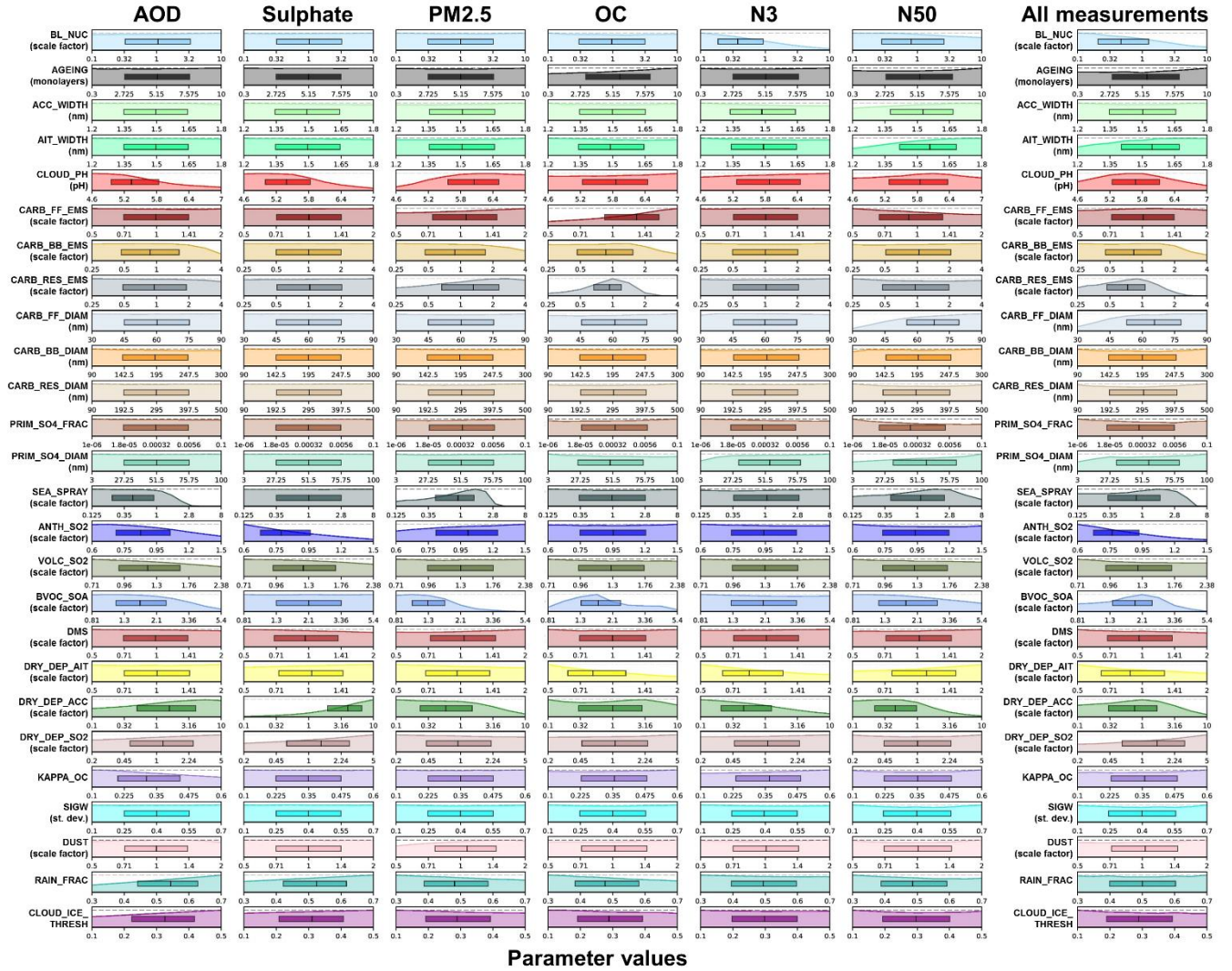


Figure 6. Marginal parameter distributions after constraint using individual measurement types over all months (6 columns on the left) and after using all measurement types over all months together (right column). The 25th, 50th and 75th percentiles of each constrained distribution are shown in the central boxes, and the parameter values on the x-axes correspond to values as they are used in the model [parameters that are multiplicative scaling factors are shown on the log₁₀ scale], covering the full parameter ranges (Yoshioka et al., 2019). The corresponding choices of threshold θ and tolerance T that were applied in the constraint process to generate these results are given in Table A2 (left column for each individual measurement type; right column for the joint measurement-types constraint), along with the percentage of model variants that is retained in the constrained sample in each case. See section 2.5 for a definition of marginal parameter distributions.

Particle concentration (N₃ and N₅₀) measurements constrain a wider range of parameters than the measurements of mass-related properties. The rate of boundary layer nucleation is strongly constrained to the low part of the sampled range by the N₃ measurements (a 77% likelihood of being below the default rate), suggesting N₃ concentrations are generally too high across

the PPE. N_3 also weakly constrains the dry deposition of Aitken and accumulation mode particles to low values. Low deposition rates of accumulation mode particles (hence higher atmospheric concentrations) will result in a higher condensation sink and more removal of sulphuric acid that participates in particle nucleation, so this is consistent with the constraint of nucleation rates to low values. The constraint of Aitken mode deposition to low values is less obvious. Aitken mode particles can contribute substantially to N_3 , so low deposition rates would enhance N_3 (opposite to the constraint on nucleation rates). However, nucleation rates are constrained to very low values, so in such a situation Aitken particles can begin to act as a sink term for nucleation by affecting the condensation sink and by growing into accumulation mode particles. BVOC emissions are not constrained by N_3 measurements, even though SOA enters into the nucleation rate expression. This is most likely because high BVOC emissions also enhance total SOA, which acts as a condensation sink for nucleation, so the two effects cancel (Carslaw et al., 2013b).

For N_{50} , the constraints are consistent with shifting the N_{50} concentrations in the ensemble towards lower values (section 3.4). N_{50} has very little effect on the range of boundary layer nucleation rate. In contrast, a previous study found that boundary layer nucleation made a statistically significant difference to model skill at about half of the ground sites they analysed (Reddington et al., 2011) – although that study tested the effect of including or not including boundary layer nucleation rather than perturbing the rate as we do here. Without boundary layer particle formation the model was structurally deficient and had poor skill at around half the sites analysed. However, our results show that uncertainty in the parameter value itself is unimportant when other parameter uncertainties are considered. This parameter is unconstrained by N_{50} measurements because there are many alternative ways of achieving model-measurement agreement.

N_{50} measurements also tend to constrain primary particle emissions to the lower end of the range (fossil fuel and primary sulphate emissions), albeit weakly. Residential particle emissions are not constrained, but the measurements we used are not well located to achieve this. It also constrains the emitted particle diameters to the high end of their ranges (fossil fuel, primary sulphate), which is again consistent with low number concentrations (since we perturb emission diameter independently of the mass, so number concentration is affected). The constraint of particle emission sizes is consistent with a previous study that showed CCN concentrations are sensitive to the assumed size (Reddington et al., 2011). Our results show that N_{50} measurements allow the emission size to be constrained, even though there are many other compensating factors that can affect CCN concentrations. N_{50} weakly constrains cloud pH to higher values, consistent with greater production of sulphate aerosol and a higher sink for nucleation. BVOC emissions are constrained to the low end, which is consistent with reduced growth of nucleation mode particles into the Aitken and accumulation modes. N_{50} also constrains deposition rates: accumulation mode deposition is constrained to low values and Aitken mode deposition to high values, suggesting a shift in the aerosol size distribution towards larger aerosols is consistent with N_{50} measurements.

3.2 Seasonal variations in constraint

Many of the parameter constraints vary seasonally, which can be linked to seasonal variations in emissions and parameter sensitivity. Some examples are shown in Figure 7. Cloud pH is constrained more by AOD in northern-hemisphere winter (Figure 7a) when in-cloud oxidation of SO₂ by ozone dominates sulphate production. BVOCs are constrained by AOD only in northern-hemisphere summer when the emissions are strong (Figure 7b). There are several other seasonal variations in the constraint effect from AOD measurements that we do not show. For example, anthropogenic SO₂ emissions are constrained by AOD more in winter because the AOD uncertainty in summer is dominated by the uncertainty in SOA. The hygroscopicity of OC is also constrained more in summer when OC is a larger component of the aerosol. Biomass burning emissions are constrained in NH summer as expected from wildfire emission seasonality and the northern-hemisphere bias of our measurements dataset. Residential emissions are only constrained in winter when emissions are high. Microphysical process rates (dry deposition of accumulation mode and wet scavenging rates) are consistently constrained throughout the year.

For PM_{2.5}, the seasonality of constraint is very similar to AOD with one notable exception. The dry deposition rate of accumulation mode particles is constrained to high values in summer (consistent with AOD and sulphate), but to low values in the winter (Figure 7c). This may occur just because of the way in which the combinations of parameters control PM_{2.5} – for example BVOCs can account for PM_{2.5} in summer so high dry deposition rates cannot be ruled out. However, it may also indicate a structural deficiency, ~~with the low deposition rates in winter implying that PM_{2.5} has missing sources in winter but not in the summer, such as nitrate. The low deposition rates in winter imply that PM_{2.5} has missing sources in winter but not in the summer, such as nitrate. Our model does not include aerosol nitrate, which (if included) would increase northern-hemisphere winter PM_{2.5} concentrations and weaken the constraint on dry deposition towards lower values in northern-hemisphere winter.~~

For N₃, we find that the boundary layer nucleation rate is constrained only in summer when photochemical production of the nucleating vapours is fast (Figure 7d). This is consistent with previous studies that have examined the seasonal cycle of organic-mediated nucleation (Riccobono et al., 2014). Similarly, N₃ measurements constrain SO₂ emissions and cloud droplet pH in summer when nucleation is most active. This is in contrast to the AOD and sulphate measurements, which constrained these two parameters in winter when their relative contribution to aerosol mass is greater.

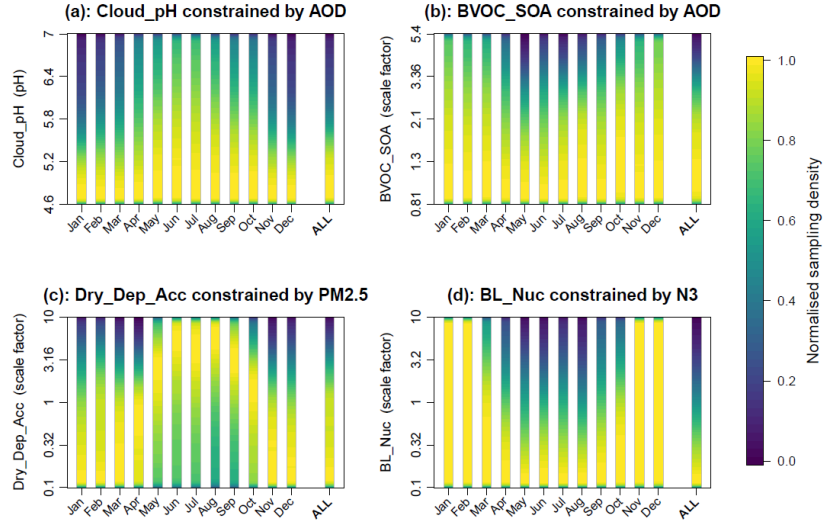


Figure 7. Seasonal variation in the constraint of parameter marginal probability distributions. The examples are a) constraint of the pH of cloud droplets (Cloud_pH) parameter using global AOD measurements, b) constraint of SOA production from BVOCs (BVOC_SOA) using AOD measurements, c) constraint of the dry deposition rate of accumulation mode particles (Dry_Dep_Acc) using global PM2.5 measurements, d) constraint of boundary layer nucleation rates (BL_Nuc) using N₃ measurements mainly over Europe

For N₅₀ we find that parameter constraints do not vary smoothly through the year (not shown). This is because the N₅₀ measurements we have used are primarily from campaigns, which move around the globe, resulting in constraint of regionally important parameters. This is one indication that we need to add long-term network measurements of N₅₀ to the dataset.

3.3 Constraint using all measurement types

The multi-variate constraint is shown as the right-hand column of pdfs in Figure 6 and Table 2 shows corresponding parameter distribution statistics from this constraint. For each individual variable / month constraint that feeds into this multi-variate constraint, the implausibility threshold and tolerance criteria (θ and T), were relaxed from the individual measurement constraints to retain approximately 75% of the 1 million model variants (Table A2). This relaxed criteria leads to measurements that provide stronger constraint being down-weighted and individual parameter constraints becoming weaker, but it means that we are able to avoid over-constraining on any one measurement type. Using all measurement types together leads to retention of only 2.1% of the original 1 million model variants as plausible models (nearly 98% rejected; Table A2). In most cases the marginal parameter distributions from this constraint can be understood in terms of the combination of individual constraints described above.

Boundary layer nucleation rates are constrained to the low end of the range, which can be attributed almost entirely to the N₃ measurements. However, the constraint is slightly weaker than when just N₃ measurements are used because of the need to relax the tolerances and thresholds applied when ruling out model variants using multiple measurement types (section 2.4.3). The nucleation rate is constrained such that the likelihood of it being in the lower half of the range (0.1-1 times the default value) is 70% - more than twice the likelihood of it being in the upper half of the range (1-10 times the default value).

The pH of cloud droplets, which controls aqueous-phase oxidation of SO₂ to form sulphate aerosol, is constrained to be more likely in the middle of our elicited range. This results from a combination of AOD and sulphate measurements constraining it to the lower end of the range and PM_{2.5} measurements constraining it to the higher end. Observational constraint is unable to rule out any of the pH values between 4.6 and 7.0, although there is reduction of 0.13 in the 95% credible interval to 4.69-6.84 (from 4.66-6.94 before constraint) and a larger reduction of 0.32 in the interquartile range to 5.24-6.12 (from 5.2-6.4 before constraint).

Biomass burning emissions are weakly constrained. The likelihood of emissions being more than a factor 2 above the default value is reduced to 14% (from 25%), but all values below this down to 0.25 times the default value are still equally likely, as they were before constraint.

Residential carbonaceous emissions are constrained primarily through a combination of PM_{2.5} and OC measurements. This emissions scaling parameter is constrained to be most likely near the middle of its range around the default setting, with emissions higher than about 2.7 times the default emission rate ruled out completely and also some weaker constraint at the lower end of the range. The 95% credible interval has significantly shifted towards lower values, from 0.27-3.73 times the default value before constraint to 0.27-1.85 times the default value after constraint, with the constrained interquartile range being 0.46-1.06 (Table 2).

The diameter of fossil fuel particles is constrained mainly through the N₅₀ measurements towards larger diameters, with a likelihood of being in the upper half of our elicited range (60-90 nm diameter) of 61% and the median of this parameter distribution shifting to a larger diameter on constraint, increasing from 60nm to 65.63nm

Sea spray emissions are constrained through a combination of AOD and PM_{2.5} measurements, and to a lesser extent by N₅₀. The multivariate constraint is slightly weaker than was achieved by AOD and PM_{2.5} individually, although we are still able to rule out emissions in the range 4.7-8 times the default value. Emissions in the range 0.125-2.8 times the default value are not strongly constrained by any of the measurements.

Anthropogenic SO₂ emissions are strongly constrained to the lower part of the elicited range by a combination of AOD and sulphate measurements. The emissions are most likely to be at the lower end of our elicited range (0.6 times the default value) and the likelihood of the emissions being in the range 0.6-1 times the default value is 82%. Our interquartile range after constraint is 0.67-0.93 times the baseline emission value of 98 Tg/y from the MACCity inventory, so 65-91 Tg/y. Our constrained range therefore lies largely below the baseline value, with only an 18% probability of it being above the baseline value. Liu et al. (2018) have developed a new SO₂ emission inventory based on OMI measurements. They did not provide a global estimate of SO₂ emissions, but over the US and Europe, where most of our sulphate measurements are located, their revised emissions are 40% lower than in HTAP, which is in the same direction as our constraint. In their inverse model study, Lee et al. (2011a) estimated global land SO₂ emissions of 100-105 Tg/y (with an estimated uncertainty of 20%), in agreement with MACCity emissions, but their central value is around our 85th percentile.

BVOC emissions are constrained to a central value that corresponds to a global annual SOA production of about 86.5 Tg/y. No values in the parameter range (corresponding to an emissions range of 37-250 Tg/y) are ruled out, although the likelihood of SOA production being in either the upper (above 150Tg/y) or lower (below 60Tg/y) quadrants of the scaling range is significantly reduced and the interquartile range of the parameter distribution has reduced from 60-155 Tg/y to 62-111 Tg/y.

5 BVOCs were constrained in Spracklen et al. (2011) using global Aerosol Mass Spectrometer measurements (which we also used) and a set of model runs that perturbed combinations of biogenic monoterpene and isoprene emissions as well as an anthropogenic VOC. Here we have used a combination of AOD, PM2.5, OC, N₅₀ and N₃ measurements, all of which are influenced by SOA. Their best estimate of the global SOA source was 140 Tg/y with an uncertainty range of 50–380 Tg/y. This included 100 Tg/y from anthropogenic sources (which they called anthropogenically controlled SOA), which we do not
10 include in our set of perturbed parameters. When we use just global OC from AMS measurements we find a 95% range on BVOC SOA of 42-195 Tg/y. Measurements of PM2.5, AOD and, to a lesser extent N₅₀, provide additional constraint, resulting in a 95% interval of 40-172 Tg/y and a median of 86.5 Tg/y. This range accounts for potential compensating effects of uncertainty in deposition rates and other parameters that were not considered in Spracklen et al. (2011).

The dry deposition rate of Aitken mode particles is weakly constrained to low values, which comes mainly from the OC
15 and N₃ observational constraint. The likelihood of the deposition rate being in the range 0.5-1.0 times the default value (1.0) is increased from 50% to 60% on constraint.

The dry deposition rate of accumulation mode particles is constrained to the middle of the range. This is likely because sulphate measurements constrain the deposition rate to be towards the high end while the other measurements constrain it towards the low end. The multivariate constraint is weaker than when individual measurement types are used (AOD, sulphate,
20 PM2.5, N₅₀, N₃), which results from relaxing the individual constraints in order to retain a reasonable number of model variants when multiple variables do not agree on the best value of the deposition rate.

The dry deposition rate of SO₂ is constrained to the upper part of the elicited range, with the likelihood of it being in the range 1-5 times the default value (i.e., an increase in SO₂ emissions) now 62% after constraint.

Parameter	Median		95% Credible Interval		95% CI Range Ratio (Constrained/Unconstrained)
BL_Nuc*	1.00	0.47	(0.11 , 8.91)	(0.11 , 6.79)	0.94
Ageing	5.15	5.51	(0.54 , 9.76)	(0.55 , 9.81)	1.00
Acc_Width	1.50	1.50	(1.21 , 1.79)	(1.21 , 1.79)	1.00
Ait_Width	1.50	1.55	(1.21 , 1.79)	(1.23 , 1.79)	0.97
Cloud_pH	5.80	5.67	(4.66 , 6.94)	(4.69 , 6.84)	0.94
Carb_FF_Ems*	1.00	1.01	(0.52 , 1.93)	(0.52 , 1.93)	1.00
Carb_BB_Ems*	1.00	0.83	(0.27 , 3.73)	(0.26 , 3.28)	0.97

Carb_Res_Ems*	1.00	0.73	(0.27 , 3.73)	(0.27 , 1.85)	0.73
Carb_FF_Diam	60.00	65.63	(31.50 , 88.50)	(35.16 , 88.80)	0.94
Carb_BB_Diam	195.00	194.97	(95.25 , 294.75)	(94.89 , 295.27)	1.00
Carb_Res_Diam	295.00	299.73	(100.25 , 489.75)	(99.26 , 492.03)	1.01
Prim_SO4_Frac*	3.16×10^{-4}	2.41×10^{-4}	(1.33×10^{-6} , 7.50×10^{-2})	(1.26×10^{-6} , 7.46×10^{-2})	1.00
Prim_SO4_Diam	51.50	56.43	(5.43 , 97.58)	(7.06 , 98.04)	0.99
Sea_Spray*	1.00	0.82	(0.14 , 7.21)	(0.14 , 3.69)	0.83
Anth_SO2*	0.95	0.77	(0.61 , 1.47)	(0.61 , 1.35)	0.90
Volc_SO2*	1.30	1.25	(0.73 , 2.31)	(0.73 , 2.30)	1.00
BVOC_SOA*	2.09	1.88	(0.85 , 5.15)	(0.86 , 3.74)	0.82
DMS*	1.00	0.97	(0.52 , 1.93)	(0.52 , 1.92)	1.00
Dry_Dep_Ait*	1.00	0.88	(0.52 , 1.93)	(0.51 , 1.90)	1.00
Dry_Dep_Acc*	1.00	0.76	(0.11 , 8.91)	(0.11 , 5.73)	0.90
Dry_Dep_SO2*	1.00	1.45	(0.22 , 4.61)	(0.23 , 4.76)	1.00
Kappa_OC	0.35	0.36	(0.11 , 0.59)	(0.11 , 0.59)	1.00
Sig_W	0.40	0.40	(0.12 , 0.68)	(0.11 , 0.69)	1.04
Dust*	1.00	1.03	(0.52 , 1.93)	(0.52 , 1.94)	1.00
Rain_Frac	0.50	0.50	(0.31 , 0.69)	(0.31 , 0.69)	1.00
Cloud_Ice_Thresh	0.30	0.29	(0.11 , 0.49)	(0.11 , 0.49)	1.00

Table 2. Marginal parameter distribution statistics (median and 95% credible interval) for the unconstrained sample of 1 million model variants in black and the constrained sample of model variants from the constraint using all measurement types simultaneously in red. (*Parameter values given as a multiplicative scaling). The final column shows the ratio of the constrained to the unconstrained 95% credible interval range, accounting for the nature of the parameter (absolute or multiplicative) by using the \log_{10} scale for the calculation when the parameter is a multiplicative scaling.

3.4 Model-measurement comparison

Figure 8 compares the unconstrained (black) and constrained distributions of model outputs with the measurements (green). We show the results when single measurement types are used for constraint (blue) and when all measurement types are used together (red). The constraint procedure clearly rules out wide ranges of model outputs that are inconsistent with the measured values, shown by the vertical green lines. For example, the unconstrained distribution of mean global sulphate concentration (at the measurement sites) extends up to about $6 \mu\text{g m}^{-3}$ in January, but the tail of the distribution is limited to $3 \mu\text{g m}^{-3}$ after constraint.

The constrained model distribution sometimes agrees much better with the measurements when only a single measurement type is used compared to when all measurements are used. The weaker multi-variate constraint is because we relax the

constraint on individual variables so as not to rule out all model variants. This effect is most apparent for sulphate and PM2.5. The mean of the constrained PM2.5 distribution using all measurements is about 40% lower than the mean of the measurements in January but the mean of the sulphate distribution is about 50% higher than the mean of the measurements. This is likely to indicate a structural error in the model that prevents good model-measurement agreement with both quantities in the same parts of model parameter space. One explanation could be that the model is missing sources of PM2.5 mass (e.g. nitrate aerosols in winter), which forces a compromise in which the constraint methodology rules in sulphate concentrations that are at the upper end of the uncertainty range to minimise the error for PM2.5. Although relaxing our constraint criteria offsets many effects of such structural errors, the shifting of these all-measurement constraint distributions away from the measurements indicates some structural error is still not fully accounted for. It is possible that our constraint would adjust better to account for this structural deficiency if we could directly specify a structural error term in the implausibility measure through $Var(S)$.

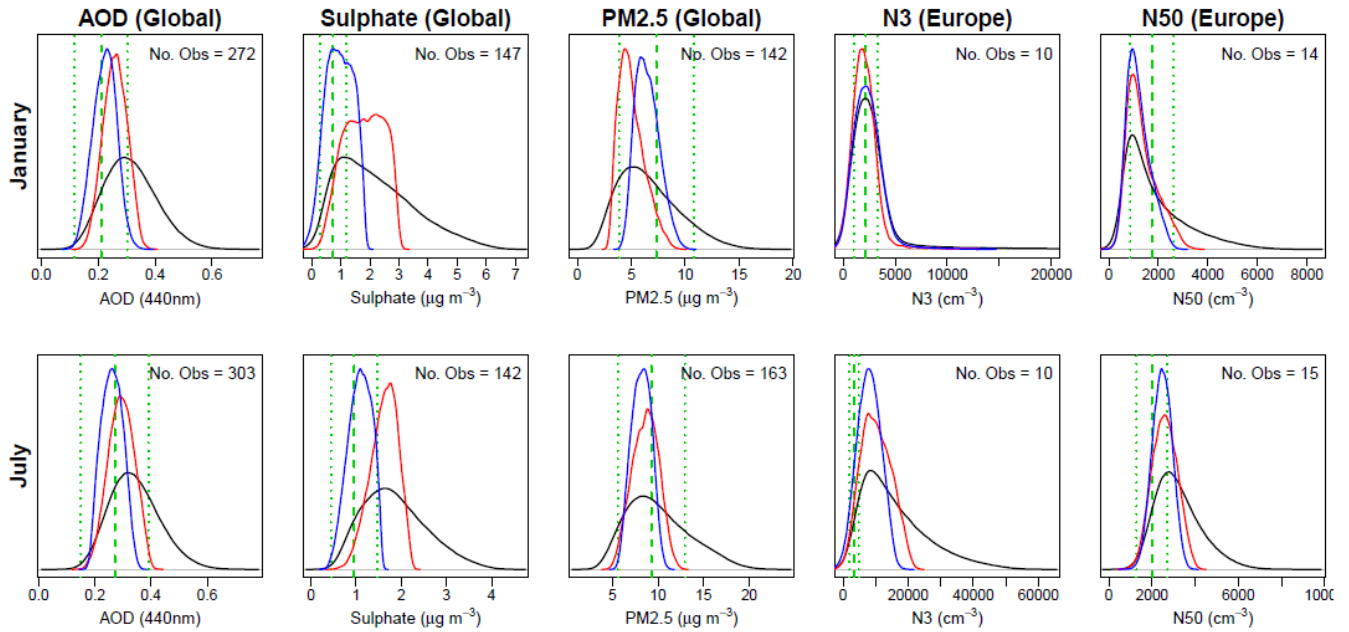


Figure 8. Comparison of the constrained model with the measurements for January (top) and July (bottom). The distributions were calculated as a mean over model grid boxes containing measurements. AOD, Sulphate and PM2.5 are global comparisons; N₃ and N₅₀ are Europe only comparisons due to the limited global coverage of these measurements in each month. The black line shows the prior (unconstrained) probability distribution of the model. The blue line shows the constrained model distribution when only measurements of each type are used in the constraint. The red line shows the constrained model distribution when all measurement types are used. The green dashed line shows the mean of the measurements and the dotted lines show the approximate 95% uncertainty range on an average observation that was accounted for in the constraint.

3.5 Unconstrained parameters

Several parameters are barely constrained or not constrained at all using all the measurements. Unconstrained microphysical processes or assumptions are the ageing rate of insoluble into soluble particles, the width of the lognormal accumulation mode,

the hygroscopicity of organic material (κ_{OC}), the updraft speed and wet deposition rates. Among the emissions, unconstrained parameters are the emission rates of fossil fuel particles, degassing volcanic SO_2 , DMS and dust emissions.

There are several potential reasons for the lack of constraint. It is possible that parts of the joint parameter space are ruled out, but with a negligible effect on the marginal parameter distribution (i.e., the ruled out parameter space is uniform across the parameter of interest). For example, wet deposition rates are directly compensated by emission rates and the ageing rate affects the wet removal rate. Another reason is that we did not include measurements in regions where the six variables are sensitive to these parameters. This is likely to be the case for DMS, volcanic and dust emissions given the relative lack of measurements over remote ocean regions and downwind of dust sources, which means these regions are not strongly weighted in the overall constraint process. Furthermore, some parameters may be more related to other aerosol properties that we have not used for constraint. For example, aging rates in the model are not constrained, likely because the ageing process predominantly affects the black carbon concentration which is not included as a measurement type in this study.

3.6 Implications for constraint of aerosol forcing

Figure 9 shows the nine most-important parameters for the uncertainty in global mean aerosol forcing in the PPE in terms of the forcing uncertainty they account for (Yoshioka et al., 2019). Some of these parameters are fairly strongly constrained by the measurements, but others are unconstrained. Within the joint parameter space of just these nine parameters there is considerable potential for model variants that compensate, thereby reducing the effectiveness of the constrained parameters on the forcing. It also needs to be borne in mind that global mean forcing is the sum of regional forcings, and in each region a different set of parameters is being constrained and may be constraining the same parameters to different parts of their range (Lee et al., 2016; Regayre et al., 2015).

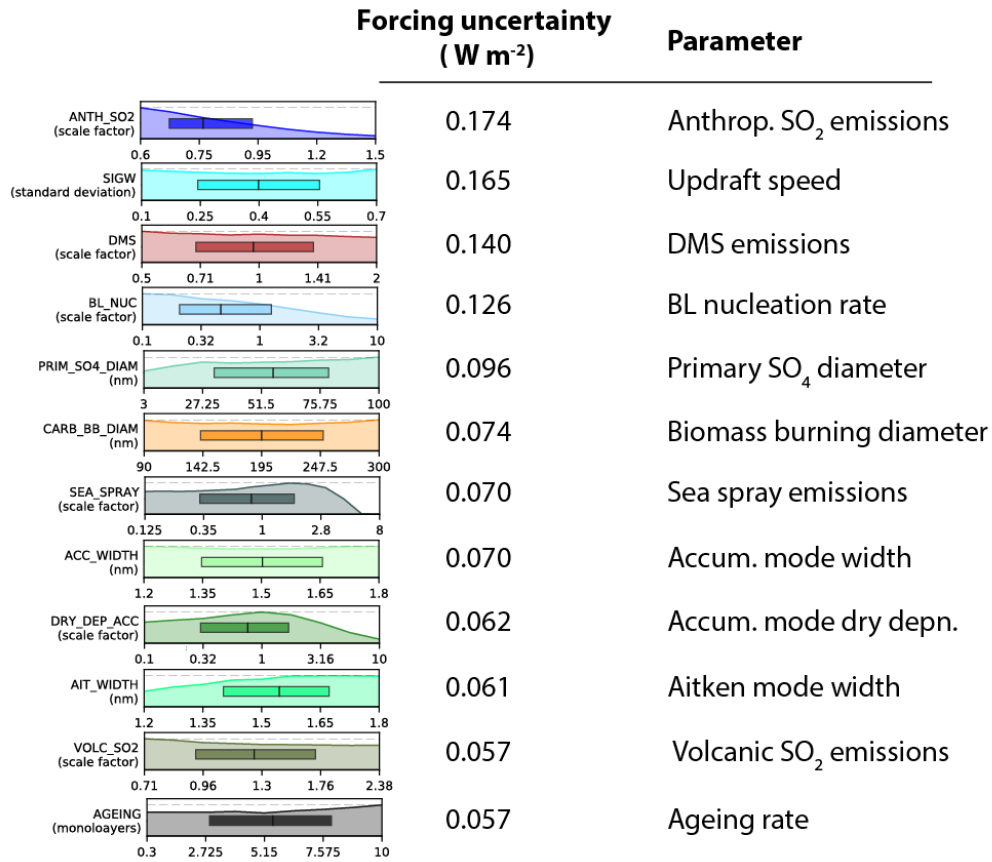


Figure 9. Ranked list of parameters that dominate the uncertainty in global mean aerosol radiative forcing in the ensemble (Yoshioka et al., 2019).

Figure 10 shows how the constrained parameters affect the uncertainty in predicted global annual mean net RF and its component parts due to aerosol-cloud interactions (RF_{aci}) and aerosol radiation interactions (RF_{ari}). (Note that this calculation of RF differs from that shown in Yoshioka et al. (2019), which used elicited parameter distributions when sampling over the parameter uncertainty space, while we use uniform distributions for the sampling here). Table 3 shows the corresponding parameter distribution statistics (median, inter-quartile range, $\pm 1\sigma$ range (on mean value) and 95% credible interval) for these forcing constraints.

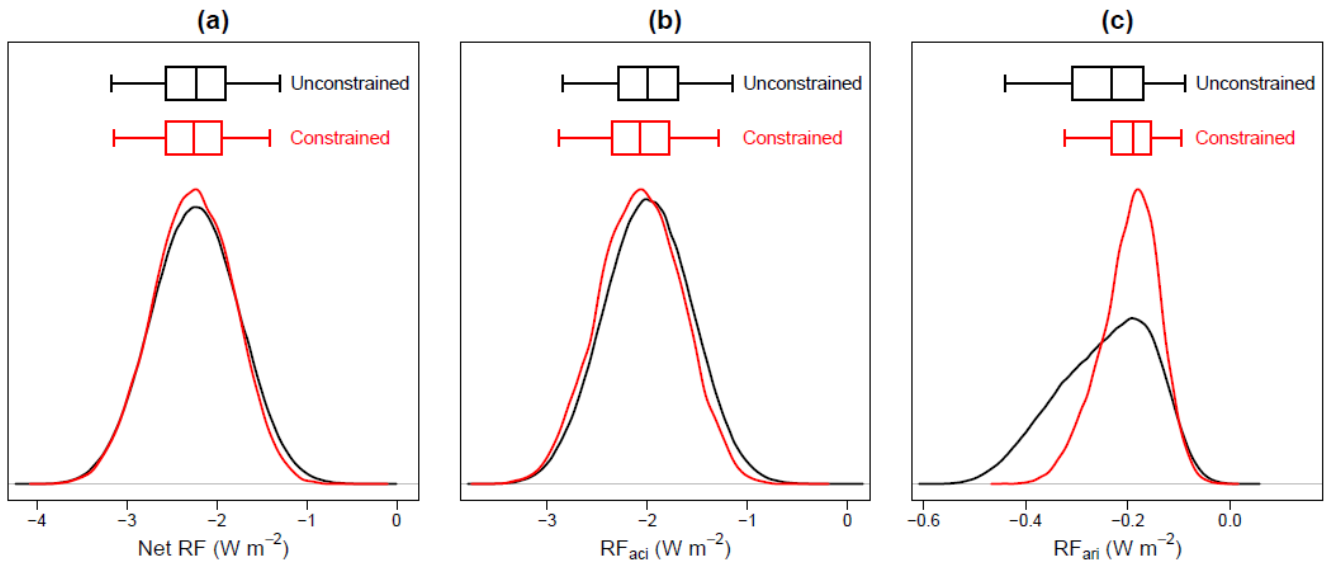


Figure 10. Effect of observational constraint using all measurement types on the probability distribution of global annual mean aerosol radiative forcing: (a) Net RF, (b) RF_{aci} (aerosol-cloud interaction), and (c) RF_{ari} (aerosol-radiation interaction). The black line shows the prior (unconstrained) distribution and the red line shows the constrained distribution. For each box and whisker plot, the box represents the interquartile range split at the median forcing (the line inside the box), and the whiskers extend to the lower and upper bounds of the 95% credible interval of the distribution.

	Median	Interquartile range	$\pm 1\sigma$ Range	95% Credible Interval	95% CI Range Ratio (Constrained/Unconstrained)	95% CI Reduction
Net RF	-2.23 -2.26	(-2.56 , -1.90) (-2.57 , -1.95)	(-2.71 , -1.75) (-2.71 , -1.81)	(-3.17 , -1.29) (-3.15 , -1.41)	0.92	8%
RF_{aci}	-1.99 -2.07	(-2.29 , -1.69) (-2.36 , -1.78)	(-2.43 , -1.55) (-2.48 , -1.66)	(-2.84 , -1.15) (-2.88 , -1.28)	0.94	6%
RF_{ari}	-0.23 -0.19	(-0.31 , -0.17) (-0.23 , -0.16)	(-0.33 , -0.15) (-0.26 , -0.14)	(-0.44 , -0.09) (-0.32 , -0.10)	0.66	34%

Table 3. Uncertainty distribution statistics (median, interquartile range, $\pm 1\sigma$ range (on mean value) and 95% credible interval) for the global annual mean aerosol radiative forcing (Net RF, RF_{aci} and RF_{ari}) from the unconstrained sample of 1 million model variants in black and the constrained sample of model variants from the all variables constraint in red. The final two columns show the ratio of the constrained to the unconstrained 95% credible interval and corresponding percentage reduction in this interval on constraint.

The net RF is dominated by RF_{aci} , which is only weakly constrained (Figure 10b) by 6%, in line with the net RF (Figure 10a). This occurs because our constraint uses measurements of aerosol properties rather than cloud properties. Although the overall reduction in the RF_{aci} uncertainty is weak, the pdfs in Figure 10b show a slight shift in RF_{aci} to stronger forcings, with the median RF_{aci} shifting from -1.99 W m^{-2} in the prior (unconstrained) distribution to -2.07 W m^{-2} after constraint. The likelihood of the strength of RF_{aci} being weaker than -1.5 W m^{-2} (less negative) is reduced by 38% and the likelihood of it being stronger (more negative) than -2.5 W m^{-2} is increased by 20%. In general, although the anthropogenic emissions were constrained to lower values (which should weaken the forcing), the sea spray emissions were constrained to lower values, which acts to

strengthen the forcing (Carslaw et al., 2013a; Regayre et al., 2014). The 95% credible interval for the net RF is reduced by 8%.

The 95% credible interval of direct forcing, RF_{ari} , is reduced by 34% (Figure 10c) and the $\pm 1\sigma$ range is reduced by 33%. RF_{ari} is constrained most strongly by the PM2.5, AOD and sulphate measurements (not shown) but insignificantly by the OC, N_3 and N_{50} measurements. The inconsistency between the constraints on PM2.5, AOD and sulphate (Figure 8) leads to inconsistency in the constraint on RF_{ari} . In particular, using just sulphate measurements results in an $RF_{ari} \pm 1\sigma$ range of -0.10 to -0.22 $W m^{-2}$, but using just PM2.5 measurements results in a range -0.20 to -0.36 $W m^{-2}$. This highlights the importance of detecting and fixing structural deficiencies in the model as well as the limitations of using single-variable emergent constraints.

Furthermore, we have found that the observational constraint on present-day (2008) global annual mean aerosol radiative effects (RE; relative to no aerosol radiative effects) is stronger than the constraint on aerosol radiative forcing. The uncertainty in the clear-sky aerosol RE is reduced by 47%. Industrial-period aerosol radiative forcing has distinct sources of uncertainty from aerosol radiative effects in the present-day atmosphere (Regayre et al., 2018), so a stronger constraint on present-day radiative effects is in line with expectations.

It is important to note that the probability distribution of net aerosol RF includes values that, with current knowledge, would produce a net negative (greenhouse gas plus aerosol) forcing over the industrial period. The forcing is dominated by aerosol-cloud processes, which we have not attempted to constrain here. Nevertheless, the lack of constraint shows, as in Lee et al. (2016), a well-configured global aerosol model has little bearing on the uncertainty in RF_{aci} . In contrast, the constraint of RF_{ari} , which has not been attempted in our previous studies, is significant and encouraging.

4 Conclusions

We have used extensive point measurements of aerosol optical depth (AOD), PM2.5, sulphate mass, organic carbon mass, and the concentrations of particles larger than 50 nm dry diameter (N_{50}) and 3 nm (N_3) from surface sites, aircraft and ships to constrain uncertain aerosol parameters in a global aerosol-climate model. Twenty-six parameters related to aerosol emissions and processes were varied in a perturbed parameter ensemble and statistical emulators were used to generate a set of 1 million model variants that represent the model outputs for combinations of parameter values across the 26-dimensional uncertainty space. The plausibility of each model variant was tested against each measurement type in turn and then in combination using a history matching procedure based on an implausibility metric (Craig et al., 1996; Williamson et al., 2013). The resulting probability distributions of aerosol forcings can be considered as the “observationally plausible” ranges for the HadGEM3-UKCA model.

Observational constraint ruled out almost 98% of the 26-dimensional parameter space and the probability distributions of many parameter values were effectively constrained. Fourteen of the parameters were constrained to some extent, despite the fact that there are many ways in which parameter values can be combined to produce plausible results within the uncertainties.

Constraint of a parameter means that the probability distribution of a parameter (and potentially its absolute range) is narrowed, and hence the likelihood of the parameter taking a particular range of values within its absolute range is increased; i.e., there are more ways to combine these parameter values with values of the other 25 parameters to produce a plausible model. For two parameters, some of the individual prior elicited parameter ranges were ruled out entirely: very high sea spray emissions and very high residential carbonaceous emissions. The very highest BVOC emissions were nearly ruled out. However, for the remaining parameters it was not possible to entirely rule out any part of the prior range.

Parameter constraints are mostly, but not always, consistent across multiple measurement types, even though the different measurement types were made at very different locations and sometimes at different times of the year. For example, we often found consistent constraint of parameters related to the production of aerosol mass or number (emissions, nucleation, secondary aerosol mass) and parameters related to removal (condensation sink, deposition rates of gases and aerosols). There is also a very clear seasonal variation in the magnitude of constraint related to variations in the dominant processes.

The multivariate constraint has the following effect on the parameter probability distributions, which were assumed before constraint to be equally plausible between lower and upper bounds defined by expert elicitation:

1. Boundary layer nucleation rates based on a sulphuric acid-organic mechanism (Metzger et al., 2010) are constrained to the low end of the elicited range, mainly from the N_3 measurements.
2. The pH of cloud droplets, which controls aqueous-phase oxidation of SO_2 to form sulphate aerosol, is constrained to be more likely in the middle of our elicited range but we were unable to rule out any of the pH values between 4.6 and 7.0. The constraint led to a reduction of 0.32 in the interquartile range to 5.24-6.12 (from 5.2-6.4 before constraint) and a reduction of 0.13 in the 95% credible interval to 4.69-6.84 (from 4.66-6.94 before constraint).
3. Biomass burning emissions are weakly constrained by the $PM_{2.5}$, AOD and OC measurements, with a reduced likelihood of the emissions exceeding a factor 2 above the default value. All lower emissions down to 0.25 times the default value are equally likely as they were before constraint.
4. Residential carbonaceous emissions are constrained primarily through a combination of $PM_{2.5}$ and OC measurements. Emissions higher than about 2.7 times the default emission rate were effectively ruled out by the OC measurements as observationally implausible.
5. The diameter of fossil fuel particles is constrained by N_{50} measurements, with a reduced likelihood of being in the range 30-45 nm, but diameters in the range 60-90 nm are unconstrained by any measurements.
6. Sea spray emissions are constrained through a combination of AOD, $PM_{2.5}$ and N_{50} measurements. Emissions in the range 4.7-8.0 times the default value are ruled out but emissions in the range 0.125-2.8 times the default value are not strongly constrained by any of the measurements.

7. Anthropogenic SO₂ emissions are strongly constrained to low values by AOD and sulphate measurements, with an 82% likelihood of being below the default value from the emission inventory.
 8. BVOC emissions are constrained by AOD, PM_{2.5}, OC, N₅₀ and N₃ measurements. The likelihood of either high or low emissions is reduced. On constraint, our median estimate corresponds to 86.5 Tg/y SOA production, with a 95% credible interval of 40-172 Tg/y. However, no values in the range 37-250 Tg/y are ruled out entirely.
 9. The dry deposition rate of Aitken mode particles is weakly constrained to low values using OC and N₃ measurements and the dry deposition rate of accumulation mode particles is constrained to the middle of the range by AOD, sulphate, PM_{2.5}, N₅₀ and N₃ measurements. The dry deposition rate of SO₂ is constrained to the upper part of the elicited range, with a 62% likelihood of it being above the default value.
 10. Several parameters of importance to aerosol forcing were not well constrained, in particular parameters related to microphysical processes (primary sulphate particle diameter, the diameter of biomass burning particles, the width of the accumulation mode and the ageing rate). Dimethyl sulphide emissions were also not strongly constrained.
- The prior (unconstrained) uncertainty (95% credible interval) in the pre-industrial to present-day net aerosol RF is reduced by 8%. The radiative forcing in the ensemble accounts for direct and indirect (cloud albedo) effects, but not cloud adjustments.
15. RF_{ari} uncertainty (95% credible interval) is reduced by 34%, but the net RF uncertainty is dominated by the RF_{aci} uncertainty, which is reduced by only 6%. The recent assessment of aerosol forcing (Bellouin et al., 2019) adopted $\pm 1\sigma$ ranges to define the uncertainty. Our equivalent $\pm 1\sigma$ ranges are -0.14 to -0.26 W m⁻² for RF_{ari} (reduction of 33% due to constraint) and -1.66 to -2.48 W m⁻² for RF_{aci} (reduction of 7% due to constraint). The reduction in uncertainty is much larger for RF_{ari} than RF_{aci} because our constraints focus on aerosol properties rather than cloud properties.
 20. Our results highlight the importance of using multiple measurement types to constrain aerosol-climate models. We have shown that use of a single measurement type, as is done in emergent constraint studies, would lead to an over-confident constraint. This is because potential structural deficiencies in our model prevent consistently good constraint across several measurement types. In particular, we showed that constraint using PM_{2.5} or sulphate aerosol measurements lead to probability distributions of RF_{ari} that barely overlap. The final multivariate constraint on forcing is therefore a compromise that achieves reasonable
 25. agreement with all observations rather than being over-confidently constrained by one metric.

In terms of future directions and requirements to achieve better constraint, we make the following recommendations:

1. We need to understand and quantify model-measurement representativeness errors. The biggest challenge (and the factor that most limits the constraint, other than model structural error) is quantification of the representativeness error associated with comparing point measurements with a global model (Reddington et al., 2017; Schutgens et al., 2017, 2016a, 2016b). The ambiguity in deciding whether a model-measurement bias is related to structural error in the model or underestimation of the uncertainty terms was the main limiting factor in our constraint procedure. The representativeness uncertainties have been estimated based on model simulations at a few locations (Schutgens et al.,

2016b), but they have not been measured, and we have no information about these uncertainties at other locations. Small campaigns to characterise the space-time variability around existing sites and for potential new sites should be considered.

Alternatively, representation errors could be reduced by increasing the temporal and spatial resolution of our model, and the effect of inter-annual variability could be reduced if we always compare the model to measurements for the correct year. In our calculations of model variant implausibility, inter-annual variability accounts for around 30 to 90% of the model-measurement comparison representation uncertainty. With smaller representation errors in the model-measurement comparison, we could more confidently reject implausible model variants. However, this approach is not currently computationally feasible.

2. We need to understand and quantify structural deficiencies in the model. Our current approach, to assume zero structural error in our implausibility calculations, enables some structural errors in the model to be detected by comparing single and multi-measurement constraints. Ideally, to produce the most robust constraint possible, these structural errors should be accounted for directly using $Var(S)$ in the implausibility measure – or, better still, corrected in the model.
3. We should prioritise expansion of measurements to cover more long-term well-characterised sites. Such measurements will have much lower (or at least well-defined) representativeness error when used for model evaluation and constraint. Sites should be characterised in terms of how they represent a typical model grid box over a long period of time. These sites should be in diverse locations so that they help to constrain a wide range of model uncertainties (Reddington et al., 2017).
4. We should aim to dedicate part of field campaigns to routine, unbiased (or effectively random) sampling of aerosols across the scale of model grid boxes. Such measurements will also have much lower representativeness error than measurements that target specific processes or aerosol environments. Many field campaigns (particularly using aircraft) often prioritise measurements to explore aerosol processes, or to characterise particular aerosol environments (e.g., pollution plumes). Others, like the Atmospheric Tomography Mission emphasised the effective sampling of air masses in an unbiased way. A greater emphasis on such tomography missions, even just as part of a larger field campaign, would benefit model uncertainty reduction.
5. Additional aerosol measurements could be used to further constrain the parameters and forcing. We expect the following (already available) measurements would provide further constraint: i) black carbon measurements to help constrain the aerosol absorption component of radiative forcing, but also as a measure of aerosol removal rates; ii) SO_2 concentrations to avoid ambiguity between sulphate and $PM_{2.5}$ constraints; iii) measurements from biomass burning regions to help constrain both the emissions and the size distribution of the particles; iv) vertical profiles of aerosol (Watson-Parris et al., 2019); v) more particle number size distribution information instead of just N_3 and N_{50} , which have not constrained the size distribution well enough.

6. Measured or derived process rates would be very useful because they would help to constrain model parameters directly, rather than relying on indirect constraint through measured state variables. For example, dry and wet deposition rates (Emerson et al., 2018) are required on the scale of model grid boxes (10's to 100's km). Similarly, direct estimates of particle formation and growth rates (Kerminen et al., 2018) would be useful, rather than just relying on integral particle number concentrations averaged over long periods.

5 It is very challenging to constrain model uncertainty using a large set of perturbed parameter model variants and extensive measurements of different types. However, the resulting ranges of model parameters and outputs (like radiative forcings) estimated in this way are much more robust than those based on a very small number of models. Although our study is incomplete (not all parameters were perturbed and not all measurements were used) the outcome is an estimate of the

10 “observationally plausible” range of aerosol forcings for the HadGEM3-UKCA aerosol-climate model. The ranges might be wider if we accounted for more sources of uncertainty and directly accounted for structural errors in the implausibility calculations, but they could also be narrowed if we improved model structural deficiencies, reduced model-measurement-representativeness errors and used a wider set of measurement types.

Appendix A

Index	Parameter Name	Description
1	BL_Nuc	Boundary layer nucleation rate
2	Ageing	Ageing of hydrophobic aerosols (no of monolayers of soluble material)
3	Acc_Width	Modal width of accumulation modes (nm)
4	Ait_Width	Modal width of Aitken modes (nm)
5	Cloud_pH	pH of cloud droplets (used to calculate the conversion of SO ₂ into sulphate)
6	Carb_FF_Ems	Carbonaceous fossil fuel emissions scale factor
7	Carb_BB_Ems	Carbonaceous biomass burning emissions scale factor
8	Carb_Res_Ems	Carbonaceous residential (biofuel) emissions scale factor
9	Carb_FF_Diam	Carbonaceous fossil fuel emission diameter (nm)
10	Carb_BB_Diam	Carbonaceous biomass burning emission diameter (nm)
11	Carb_Res_Diam	Carbonaceous residential (biofuel) emission diameter (nm)
12	Prim_SO4_Frac	Mass fraction of SO ₂ converted to new sulphate particles in power plant plumes
13	Prim_SO4_Diam	Mode diameter of new sub-grid sulphate particles (nm)
14	Sea_Spray	Sea spray aerosol scale factor
15	Anth_SO2	Anthropogenic SO ₂ emission scale factor
16	Volc_SO2	Volcanic SO ₂ emission scale factor
17	BVOC_SOA	Biogenic secondary aerosol formation from volatile organic compounds scale factor
18	DMS	Dimethyl sulphide surface ocean concentration scale factor
19	Dry_Dep_Ait	Aitken mode dry deposition velocity scale factor
20	Dry_Dep_Acc	Accumulation mode dry deposition velocity scale factor
21	Dry_Dep_SO2	SO ₂ dry deposition velocity scale factor
22	Kappa_OC	Kappa-Kohler coefficient of organic carbon
23	Sig_W	Updraft vertical velocity standard deviation (used to calculate the activation of aerosols into cloud drops)
24	Dust	Dust emission scale factor
25	Rain_Frac	Fraction of cloud-covered area in large-scale clouds where aerosol scavenging by rain drops occurs
26	Cloud_Ice_Thresh	Threshold of cloud ice fraction above which nucleation scavenging is suppressed (restricting further activation of aerosols into cloud drops)

Table A1. The 26 aerosol parameters included in the AER PPE. Further details are provided in a separate publication (Yoshioka et al., 2019).

AOD		All-months constraint choices (LHS Fig. 56)			Joint all-variables-months constraint choices (RHS Fig 56)		
Month	No. Obs used in constraint	Threshold Implausibility	Tolerance (No. Obs)	% Variants Retain	Threshold Implausibility	Tolerance (No. Obs)	% Variants Retain
Jan	272 / 294	3.5	33	40.03	4	76	75.60
Feb	274 / 301	3.5	27	42.37	3.5	82	75.71
Mar	284 / 309	3.5	23	40.54	3.5	80	75.78
Apr	289 / 316	3.5	17	39.70	3.5	52	76.47
May	292 / 322	3.5	12	34.55	3.5	47	76.17
Jun	295 / 320	3.5	24	43.10	3.5	53	74.13
Jul	303 / 323	3.5	24	43.38	3.5	48	75.46
Aug	310 / 326	3.5	19	35.51	3.5	50	74.06
Sep	303 / 321	3.5	24	43.09	3.5	61	74.65
Oct	285 / 315	3.5	23	39.57	3.5	80	75.97
Nov	273 / 309	3.5	27	41.91	3.5	93	74.75
Dec	267 / 298	3.5	43	41.44	4	91	75.82
ALL	3447 / 3754			19.01			60.42
Sulphate		All-months constraint choices (LHS Fig. 56)			Joint all-variables-months constraint choices (RHS Fig 56)		
Month	No. Obs used in constraint	Threshold Implausibility	Tolerance (No. Obs)	% Variants Retain	Threshold Implausibility	Tolerance (No. Obs)	% Variants Retain
Jan	147 / 149	3.5	3	38.76	4	50	75.42
Feb	164 / 148	3.5	3	38.79	4	38	74.64
Mar	144 / 151	3.5	6	42.70	4	46	74.99
Apr	149 / 151	3.5	6	38.66	4	51	75.22
May	146 / 149	3.5	9	41.38	4.5	35	74.60
Jun	142 / 150	3.5	6	38.19	3.5	48	74.40
Jul	142 / 148	3.5	9	41.75	4	40	75.19
Aug	143 / 148	3.5	9	38.97	4	46	75.00
Sep	144 / 147	3.5	3	39.13	4	46	74.77
Oct	143 / 147	3.5	3	38.03	4	46	75.02
Nov	143 / 146	3.5	3	40.21	4	46	75.52
Dec	144 / 147	3.5	3	41.52	4	49	72.42
ALL	1733 / 1781			27.73			64.90
PM2.5		All-months constraint choices (LHS Fig. 56)			Joint all-variables-months constraint choices (RHS Fig 56)		
Month	No. Obs used in constraint	Threshold Implausibility	Tolerance (No. Obs)	% Variants Retain	Threshold Implausibility	Tolerance (No. Obs)	% Variants Retain
Jan	142 / 168	3.5	20	41.52	3.5	40	74.02
Feb	145 / 168	3.5	20	36.27	3.5	38	73.51

Mar	154 / 170	3.5	15	39.14	3.5	34	75.09
Apr	152 / 170	3.5	9	43.70	3.5	24	72.93
May	154 / 167	3.5	9	43.16	3.5	22	75.83
Jun	158 / 170	3.5	13	34.89	3.5	32	74.91
Jul	163 / 172	3.5	16	40.27	3.5	36	74.08
Aug	161 / 169	3.5	13	36.70	3.5	39	75.68
Sep	154 / 166	3.5	9	42.71	3.5	31	75.37
Oct	151 / 165	3.5	12	45.71	3.5	24	75.59
Nov	142 / 168	3.5	14	38.76	3.5	31	73.69
Dec	148 / 169	3.5	24	40.51	3.5	44	75.52
ALL	1824 / 2022			10.80			48.89
OC		All-months constraint choices (LHS Fig. 56)			Joint all-variables-months constraint choices (RHS Fig 56)		
Month	No. Obs used in constraint	Threshold Implausibility	Tolerance (No. Obs)	% Variants Retain	Threshold Implausibility	Tolerance (No. Obs)	% Variants Retain
Jan	3 / 6	3.5	0	63.60	4.5	0	76.25
Feb	10 / 14	3.5	3	58.59	3.5	4	86.56
Mar	41 / 82	3.5	2	57.69	3.5	3	72.06
Apr	49 / 74	3.5	2	68.29	3.5	3	76.20
May	20 / 23	3.5	1	56.91	3.5	2	76.69
Jun	22 / 23	3.5	3	71.67	3.5	4	76.64
Jul	18 / 23	3.5	2	57.40	3.5	4	73.36
Aug	21 / 23	3.5	2	65.05	3.5	3	72.19
Sep	17 / 22	4	4	67.45	3.5	5	77.65
Oct	24 / 41	3.5	4	72.35	3.5	4	72.35
Nov	15 / 37	4.5	0	58.57	3.5	1	81.81
Dec	11 / 15	3.5	1	71.62	3.5	1	71.62
ALL	251 / 383			6.94			20.00
N ₃		All-months constraint choices (LHS Fig. 56)			Joint all-variables-months constraint choices (RHS Fig 56)		
Month	No. Obs used in constraint	Threshold Implausibility	Tolerance (No. Obs)	% Variants Retain	Threshold Implausibility	Tolerance (No. Obs)	% Variants Retain
Jan	12 / 13	3.5	0	95.74	3.5	0	95.74
Feb	12 / 13	3.5	0	96.60	3.5	0	96.60
Mar	12 / 13	3.5	0	85.80	3.5	0	85.80
Apr	11 / 12	3.5	0	68.78	3.5	1	75.35
May	11 / 12	3.5	0	56.05	3.5	4	77.67
Jun	12 / 12	3.5	1	58.61	3.5	4	75.70
Jul	13 / 13	3.5	1	57.15	3.5	4	73.41
Aug	13 / 13	3.5	1	53.12	3.5	4	73.59
Sep	13 / 13	3.5	0	67.29	3.5	1	74.97

Oct	12 / 13	3.5	0	76.94	3.5	0	76.94
Nov	12 / 13	3.5	1	97.44	3.5	1	97.44
Dec	10 / 12	3.5	0	94.03	3.5	0	94.03
ALL	143 / 152			44.03			61.58
N ₅₀		All-months constraint choices (LHS Fig. 56)			Joint all-variables-months constraint choices (RHS Fig 56)		
Month	No. Obs used in constraint	Threshold Implausibility	Tolerance (No. Obs)	% Variants Retain	Threshold Implausibility	Tolerance (No. Obs)	% Variants Retain
Jan	74 / 77	3.5	3	55.41	3.5	9	76.09
Feb	82 / 90	3.5	7	53.78	3.5	13	75.28
Mar	132 / 148	3.5	8	50.40	3.5	16	74.89
Apr	177 / 199	3.5	25	52.59	3.5	32	71.65
May	62 / 64	3.5	2	53.39	3.5	7	74.25
Jun	88 / 96	3.5	4	48.43	3.5	9	73.21
Jul	102 / 115	3.5	4	48.30	3.5	10	74.10
Aug	97 / 109	3.5	4	58.29	3.5	8	73.51
Sep	114 / 133	3.5	7	52.34	3.5	14	76.95
Oct	112 / 119	3.5	9	45.93	3.5	16	72.77
Nov	113 / 155	3.5	20	49.80	3.5	27	77.18
Dec	62 / 67	3.5	9	46.17	3.5	14	77.70
ALL	1215 / 1372			9.99			40.45
Final % variants retain for the joint all-variables-months constraint:							2.085

Table A2. The choices of the number of measurements (tolerance, T) allowed to exceed a threshold implausibility (θ) for constraint on each of the monthly observed aerosol properties, along with the corresponding percentage of the large sample of 1 million model variants (that covers the PPE parametric uncertainty) that is retained on constraint for each choice. The choices on the left correspond to those used for the all-months constraints shown on the left of Figure 6. The choices on the right correspond to those used for the joint all-variables-months constraint shown on the right in Figure 6. The percentage of variants retained is also shown for the combined all-months constraints for each variable (ALL) as well as for the joint all-variables-months constraint (final row).

Appendix B: Description of the data sets used in this study

For the AERONET Aerosol Optical Depth (AOD) data used in this study, we thank the PIs of the AERONET sites used for maintaining their instruments and providing their data to the community. We also acknowledge AERONET for their continuous efforts in providing high-quality measurements and derivative products. All data used in this work can be accessed through the AERONET web page: <http://aeronet.gsfc.nasa.gov/>.

For the sulphate data used in this study, we acknowledge the EMEP (<http://ebas.nilu.no/>; Tørseth et al., 2012), IMPROVE (<http://views.cira.colostate.edu/fed/>) and EANET (<http://www.eanet.asia/product/index.html>) measurement networks for making their measurement data available, along with all data managers involved in data collection. Additional ground station observations from the SORPES (Station for Observing Regional Processes of the Earth System) monitoring station in Nanjing, China (Ding et al., 2016) are also included. Data on the Acid Deposition in the East Asian Region was provided

from the Network Center for EANET, <https://monitoring.eanet.asia/document/public/index> (last accessed: 7 June 2018).

IMPROVE is a collaborative association of state, tribal, and federal agencies, and international partners. US Environmental Protection Agency is the primary funding source, with contracting and research support from the National Park Service. The Air Quality Group at the University of California, Davis is the central analytical laboratory, with ion analysis provided by Research Triangle Institute, and carbon analysis provided by Desert Research Institute.

For the PM_{2.5} data used in this study, we acknowledge the IMPROVE (<http://views.cira.colostate.edu/fed/>), WMO GAW–WDCA (<https://www.gaw-wdca.org>; <http://ebas.nilu.no/>; Tørseth et al., 2012), A-PAD (Atanacio et al., 2016), and NAPS (Galarneau et al., 2016; <http://maps-cartes.ec.gc.ca/rnspa-naps/data.aspx?lang=en>) measurement networks for making their measurement data available, along with all data managers involved in data collection. Further ground station measurements are included from sites in Australia (ANSTO stations: Cohen and Atanacio, 2015), South America (Artaxo et al., 2013), Taiwan (Fang and Chang, 2010), South Africa (Vakkari et al., 2013) and Nanjing, China (SORPES station; Ding et al., 2016). The PM_{2.5} data for Europe was obtained from the world data centre for aerosol (WDCA), and we thank the following data providers to this network: Adamos Adamides (Cyprus), Jacobus P. J. Berkhout (Netherlands), Elke Bieber (Germany), Tanja Bolte (Slovenia), Geoff Broughton (United Kingdom), Darius Ceburnis (Ireland), Anna Degorska (Poland), Iveta Dubakova (Latvia), Fermin Elizaga (Spain), Marina Froehlich (Austria), Marina Frolova (Latvia), Robert Gehrig (Switzerland), Alberto Gonzalez (Spain), C. Gruening (Italy), Savvas Kleanthous (Cyprus), Manuel Lambas (Spain), Maj Britt Larka Abellan (Spain), Marijana Murovec (Slovenia), Jaroslav Pekarek (Czech Republic), Noemi Perez (Spain), Cinzia Perrino (Italy), Jean-Philippe Putaud (Italy), Xavier Querol (Spain), Stephane Sauvage (France), Karin Sjoberg (Sweden), Andre Sonntag (Germany), Gerald Spindler (Germany), D. P. J. Swart (Netherlands), Karin Uhse (Germany), Milan Vana (Czech Republic), Keith Vincent (UK), P. Aalto (Finland), M. Kulmala (Finland), Anne-Gunn Hjellbrekk, (Norway) and Aas Wenche (Norway).

The N₃ data used in this study was obtained from the EBAS ACTRIS database (Asmi et al., 2013; <https://www.actris.eu/>; <http://ebas.nilu.no/>), collated via the Global Aerosol Synthesis and Science Project, GASSP (Reddington et al., 2017) and public data on the EBAS database. The EBAS database has largely been funded by the UNECE CLRTAP (EMEP) and AMAP and through NILU internal resources. Specific developments have been possible due to projects like EUSAAR (EU-FP5; EBAS web interface), EBAS Online (Norwegian Research Council INFRA; upgrading the database platform) and HTAP (European Commission DG-ENV; import and export routines to build a secondary repository in support of <http://www.htap.org>; last access: 4 April 2019). A large number of specific projects have supported development of data and meta data reporting schemes in dialog with data providers (EU; CREATE, ACTRIS and others). Through ACTRIS, the research leading to these results has received funding from the European Union's Horizon 2020 research and innovation programme under grant agreement No 654109. For a complete list of programmes and projects for which EBAS serves as a database, please consult the information box in the Framework filter of the web interface. These are all highly acknowledged for their support.

The observations of OC used in this study are collated from 7 observational campaigns and supplemented by additional ground station observations. The campaign data were collated via GASSP and are derived from size distribution measurements taken during the following campaigns: VOCALS (NERC Grant: NE/F019874/1; Allen et al., 2011; Hawkins et al., 2010; Wood et al., 2011), CalNex (Ryerson et al., 2013), WACS (Quinn et al., 2014), ICEALOT (Frossard et al., 2011), DYNAMO (DeWitt et al., 2013), NEAQS-2004 (Quinn et al., 2006; Wang et al., 2007), TEXAQS06 (Bates et al., 2008), RHAMBLE (NERC Grants NE/D006570/1, NE/E011454/1; Allan et al., 2009) and ACCACIA (NERC Grant NE/I028696/1; Allan et al., 2015). The OC ground station observations used are from the AMS Global Database which has a worldwide coverage (Zhang et al.), along with data for further European sites from ACTRIS (<https://www.actris.eu/>; <http://ebas.nilu.no/>; collated via GASSP) and data from individual stations including Chilbolton (England; Crippa et al., 2014), COPS (NERC Grant NE/E016200/1; Hornisgrinde, Germany; Irwin et al., 2010; Jones et al., 2011), Holme Moss (England; Liu et al., 2011), OP3 (NERC Grant NE/D004624/1; South-East Asia; Hewitt et al., 2010) and MC4 (NERC Grant NE/H008136/1; Weybourne, England; Liu et al., 2013) collated via GASSP, the SORPES site in Nanjing, China (Ding et al., 2016), and AMF stations in the USA and north-east Atlantic (Atmospheric Radiation Measurement (ARM) user facility, 2014a, 2014b). The OC data at the AMF stations (USA and north-east Atlantic) were obtained from the Atmospheric Radiation Measurement (ARM) Program sponsored by the U.S. Department of Energy, Office of Science, Office of Biological and Environmental Research, Climate and Environmental Sciences Division. The AMS Global Database ground station data covers the following sites: Barcelona (Minguillón et al., 2011; Mohr et al., 2012), Beijing (Sun et al., 2010), Blodgett Forest (Farmer et al., 2011), Boulder (Nemitz et al., 2008), Cape Hedo, Chebogue Point, Cheju Island (Topping et al., 2004), Chilbolton, Cool (Setyan et al., 2012), Duke Forest (Stroud et al., 2007), Edinburgh (Allan et al., 2003a), Finokalia (Hildebrandt et al., 2010), Fukue Island (Takami et al., 2005), Helsinki (Timonen et al., 2013), Houston (Canagaratna et al., 2007), Hyytiälä (Allan et al., 2006), Jungfraujoch (Ng et al., 2010), Komaba (Takegawa et al., 2006), K-Puszt, Mace Head (Dall'Osto et al., 2010), Mainz (Ng et al., 2010), Manaus (Chen et al., 2009), Manchester (Allan et al., 2003b), Melpitz (Poulain et al., 2011), Mexico City (Aiken et al., 2009), Montseny, New York City (Drewnick et al., 2004; Sun et al., 2011; Weimer et al., 2006), Pasadena (Hayes et al., 2013), Pinnacle State Park (Bae et al., 2007), Pittsburgh (Zhang et al., 2005), Point Reyes National Seashore | ARM Mobile Facility (AMF) (Ervens et al., 2010), Puy de Dome (Freney et al., 2011), Riverside | SOAR field site (Docherty et al., 2011; Williams et al., 2010), San Pietro Capofiume, Storm Peak, Trinidad Head (Millet et al., 2004), Vancouver (Alfarra et al., 2004; Boudries et al., 2004), Weybourne Atmospheric Observatory, Whistler Mountain (Sun et al., 2009), Whiteface Mountain (Hogrefe et al., 2004), and Writtle Agricultural College.

The observations of N₅₀ used in this study are collated from 19 observational campaigns and supplemented by additional ground station observations. The campaign data, collated via GASSP, and derived from size distribution measurements taken during the following campaigns: ACE1 (Bates et al., 1998; Clarke et al., 1998), VOCALS (NERC Grant: NE/F019874/1; Allen et al., 2011; Hawkins et al., 2010; Wood et al., 2011), DOE ARM MAGIC (Lewis and Teixeira, 2015), CalNex (Ryerson et al., 2013), WACS (Quinn et al., 2014), NEAQS-2002 (Bates et al., 2005; Quinn and Bates, 2005), ARCTAS (McNaughton et

al., 2011), ASCOS (Heintzenberg and Leck, 2012; Tjernström et al., 2014), ICEALOT (Frossard et al., 2011), AEROSOL99 (Bates et al., 2001), DYNAMO (DeWitt et al., 2013), INDOEX (Quinn and Bates, 2005; Ramanathan et al., 2001), PEM-Tropics-A (Fenn et al., 1999), PEM-Tropics- B (Raper et al., 2001), PASE (Hudson and Noble, 2009), NAURU99 (Long and McFarlane, 2012), ACE-ASIA (Bates et al., 2004; Huebert et al., 2003), NEAQS-2004 (Quinn et al., 2006; Wang et al., 2007) and TEXAQS06 (Bates et al., 2008). The N_{50} ground station observations used, collated via GASSP and public data on the EBAS database, are from Canada (Jeong et al., 2010; Leaitch et al., 2013; Takahama et al., 2011), South Africa (Vakkari et al., 2013), the Russian Arctic (Asmi et al., 2016), India (Hyvärinen et al., 2010), Antarctica (Fiebig et al., 2009) and European sites (Asmi et al., 2011). ASCOS (the Arctic Summer Cloud Ocean Study) was funded by the Knut and Alice Wallenberg Foundation and DAMOCLES (EU 6th Framework Program). The Swedish Polar Research Secretariat provided access to the icebreaker Oden and logistical support.

Data Availability

Measurement data can be sourced from the databases and data providers as outlined in Appendix B and Table S1 in the supplementary data file. Model data and analysis code can be made available upon request from the corresponding author. The authors welcome use of the perturbed parameter ensemble for advancing climate research.

Author Contributions

JSJ applied the statistical methodology and generated and analysed the results. JSJ, LAR and KSC wrote the article. JSJ, LAR, KSC, KJP, STT, JB, DMHS, JWR and NAJS contributed to the analysis and interpretation of results. KJP, MY, LAR, NAJS, DGP, KSC and JSJ helped prepare the model configuration that served as the template for the PPE. LAR and JSJ designed the experiments. All PPE simulations were created by MY. LAR and JSJ elicited probability density functions of all aerosol parameters, and KSC, KJP, MY, and STT participated (alongside many other experts) in the formal elicitation process. DL, JDA, HC, AD, DDC, AA, VV and EA contributed measurement data to the study from individual measurement campaigns.

Competing Interests

The authors declare that they have no conflict of interest.

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