

The value of remote marine aerosol measurements for constraining radiative forcing uncertainty

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Abstract. Aerosol measurements over the Southern Ocean are used to constrain aerosol-cloud interaction radiative forcing (RF_{aci}) uncertainty in a global climate model. Forcing uncertainty is quantified using one million climate model variants that sample the uncertainty in nearly 30 model parameters. Measurements of cloud condensation nuclei and other aerosol properties from an Antarctic circumnavigation expedition strongly constrain natural aerosol emissions: default sea spray emissions need to be increased by around a factor of 3 to be consistent with measurements. Forcing uncertainty is reduced by around 7% using this set of several hundred measurements, which is comparable to the 8% reduction achieved using a diverse and extensive set of over 9000 predominantly Northern Hemisphere measurements. When Southern Ocean and Northern Hemisphere measurements are combined, uncertainty in RF_{aci} is reduced by 21% and the strongest 20% of forcing values are ruled out as implausible. In this combined constraint, observationally plausible RF_{aci} is around 0.17 W m^{-2} weaker (less negative) with 95% credible values ranging from -2.51 to -1.17 W m^{-2} (standard deviation -2.18 to -1.46 W m^{-2}). The Southern Ocean and Northern Hemisphere measurement datasets are complementary because they constrain different processes. These results highlight the value of remote marine aerosol measurements.

1 Introduction

The uncertainty in the magnitude of the effective radiative forcing caused by aerosol-cloud interactions (ERF_{aci}) due to changing emissions over the industrial period is around twice that for CO_2 (Stocker et al., 2013). It is essential to reduce this uncertainty if global climate models are to be used to robustly predict near-term changes in climate (Andreae et al., 2005, Myhre et al., 2013, Collins et al., 2013, Tett et al., 2013, Seinfeld et al., 2016).

Aerosol forcing uncertainty has persisted in climate models since the 1990s partly because there are no measurements covering the industrial period that can be used to directly constrain simulations of long-term changes in aerosol and cloud properties (Gryspeerd et al., 2017; McCoy et al., 2017). Estimates of aerosol forcing over the industrial period therefore rely on models that have been evaluated against measurements made in the present-day atmosphere. However, it is known that the aerosol forcing (in particular the component caused by aerosol-cloud interactions) depends sensitively on the state of aerosols in the pre-industrial period (Carslaw et al., 2013; Wilcox et al. 2015) when natural aerosols were dominant (Carslaw et al., 2017). Observations of natural aerosols in the present-day atmosphere are therefore expected to help constrain the simulated forcing unless there have been significant changes in natural aerosol processes over the industrial period, for which there is little evidence (Carslaw et al., 2010).

50 In this paper we address the questions: i) To what extent can measurements of aerosols in pristine (natural)
51 environments help to constrain model simulations and thereby reduce the large uncertainty in aerosol forcing?
52 ii) What is the relative importance of measurements in remote and polluted environments for constraining the
53 forcing uncertainty? It is known that the abundance of natural aerosols affects the magnitude of forcing in a
54 model (Spracklen and Rap, 2013; Carslaw et al., 2013). However, to assess the effect on the *uncertainty* in
55 forcing it is necessary to explore how the spread of predictions of a set of models changes when constrained by
56 measurements. The 5th Coupled Model Intercomparison Project is inadequate for this purpose because of
57 insufficient aerosol diagnostics (Wilcox et al., 2015). Here we use large perturbed parameter ensembles (PPEs)
58 of the UK Hadley Centre General Environment Model HadGEM3 (Hewitt et al, 2011). The PPEs were created
59 by systematically perturbing numerous model parameters related to natural and anthropogenic emissions and
60 physical processes (Yoshioka et al., 2019). The simulated aerosol forcings have uncertainty ranges that exceed
61 those of multi-model ensembles (Yoshioka et al., 2019; Johnson et al., 2019). Instantaneous radiative forcing
62 (RF) is quantified using the 26-parameter AER PPE in which just aerosol-related parameters were varied, and
63 the effective radiative forcing (ERF) is quantified using the 27-parameter AER-ATM PPE in which aerosol and
64 physical atmosphere parameters were varied (Yoshioka et al., 2019). We use these PPEs to quantify how the
65 constraint provided by pristine aerosol measurements affects the spread of aerosol forcings simulated by the
66 ensembles.

67

68 Previous analysis of HadGEM3 PPEs showed that measurements of the present-day atmosphere in regions
69 affected by anthropogenic emissions help to constrain the uncertainty in aerosol-radiation interaction forcing
70 (RF_{ari}) but not the component due to aerosol-cloud interactions (RF_{aci}). For example, Regayre et al. (2018)
71 showed that top-of-the-atmosphere shortwave radiation flux measurements reduce ERF_{aci} uncertainty by only
72 around 10%, despite the fluxes in the present-day and early-industrial environments sharing multiple causes of
73 uncertainty. Johnson et al. (2019) showed that a much larger dataset of over 9000 (predominantly Northern
74 Hemisphere) aerosol measurements reduced the uncertainty in global, annual mean aerosol RF_{ari} (neglecting
75 rapid adjustments) by 35%, but RF_{aci} uncertainty by only around 7%. These measurements reduce the
76 uncertainty in a small number of parameters related to anthropogenic emissions and aerosol processing in
77 polluted environments. However, important causes of uncertainty in RF_{aci} , such as natural aerosol emission
78 fluxes, were largely unconstrained.

79

80 The Southern Ocean is one of the few regions on Earth (along with some boreal forests) in which the same
81 processes are expected to affect cloud-active aerosol concentrations in the present-day and early-industrial
82 atmospheres (Hamilton et al., 2014). In this study we make use of aerosol measurements from the Antarctic
83 Circumnavigation Expedition: Study of Preindustrial-like Aerosols and Their Climate Effects (ACE-SPACE)
84 campaign (Schmale et al., 2019). They offer a unique opportunity to constrain the early-industrial aspects of
85 aerosol forcing uncertainty because the Southern Ocean is a source of natural aerosols that are relevant at the
86 global scale and remains largely unaffected by anthropogenic aerosol and precursor emissions.

87

88 We use near-surface measurements of cloud condensation nuclei concentrations at 0.2% and 1.0%
89 supersaturations ($CCN_{0.2}$ and $CCN_{1.0}$; Tatzelt et al., 2019), as well as mass concentrations of non-sea-salt sulfate
90 particles with dry aerodynamic diameters less than 10 μm and number concentrations of particles with dry
91 aerodynamic diameter larger than 700 nm (N_{700} ; corresponds to volume equivalent diameter larger than around
92 500 to 570 nm; Schmale et al., 2019a). The measurements are compared to output from 1 million variants of the
93 HadGEM3 model that sample combinations of parameter settings in the model. These model variants are used to
94 represent aerosol forcing uncertainty in our model using probability density functions (pdfs) and were generated
95 by sampling from Gaussian Process emulators that were trained on the PPE model outputs (see SI Methods).
96 Model variants that were judged to be observationally implausible against the measurements were rejected,
97 resulting in a set of plausible variants from which the uncertainty in aerosol forcing could be computed (see SI
98 Methods). In the results shown below, we retained approximately 3% of model variants (following Johnson et
99 al., 2019) that best match all four measured aerosol properties.

100 2 Results

101

102 Fig. 1 shows the $CCN_{0.2}$ mean and standard deviation from the unconstrained and constrained model variants to
103 exemplify the effect of constraint on model output. The mean concentrations in the unconstrained sample are
104 much smaller than measured concentrations. However, the range of $CCN_{0.2}$ values in the unconstrained sample
105 spans the measurements in most locations (Fig. 1b). The measurement constraint increases $CCN_{0.2}$

106 concentrations (more than double the unconstrained mean in many locations; Fig. 1c) and greatly reduces the
107 CCN_{0.2} uncertainty (by more than half everywhere to less than 50 cm⁻³; Fig. 1d).

108
109 Fig. 2 shows pdfs of the output from the model for the four variables used as constraints, calculated as means
110 over the locations where measurements were taken. The constraint reduces the uncertainty in all measurement
111 types (narrower pdfs) and the central tendency of the pdfs is closer to the regional mean of measurements after
112 constraint. Rejecting around 97% of model variants as implausible compared to measurements greatly improves
113 the model-measurement comparison.

114
115 After constraint, the remaining model variants inhabit specific parts of the 26-dimensional parameter uncertainty
116 space used to quantify the model uncertainty. We explore the effect of constraints on parameter values using 1-
117 dimensional marginal probability distributions (described in detail in Johnson et al., 2019) – see Fig. 3 and Fig.
118 S2 for equivalent AER-ATM results. The magnitude of the marginal probability distribution after constraint
119 reflects the number of ways in which a particular value of a parameter can be combined with settings of all the
120 other parameters to produce an observationally plausible model. The white space in the marginal pdfs shows
121 where parameter value density has decreased.

122
123 The relative simplicity of aerosol emissions and processes over the Southern Ocean (compared to polluted
124 continental regions) means that measurements can be used to tightly constrain uncertainty in the associated
125 parameters. Two parameters (sea spray emissions and dry deposition velocity) are tightly constrained such that
126 some parameter values are ruled out as implausible even when combined with uncertainties in all other
127 parameters. Several other parameters (related to cloud droplet pH, dimethylsulfide (DMS) emissions and wet
128 deposition) are more modestly constrained. These joint constraints (see also Fig. S3) suggest the model-
129 measurement comparison is improved when aerosol number concentrations and mass are relatively high.

130
131 Sea spray emissions are tightly constrained to be around 3 times larger than the default model value.
132 Observationally plausible values of the sea spray scaling parameter range from around 1.6 to 5.1 and all other
133 values (including the default emission calculated in the model) are ruled out as implausible. This suggests that
134 sea spray emissions in our model need to be significantly higher than those calculated using the wind speed
135 dependent Gong (2003) parametrisation in the Southern Hemisphere summer. The higher flux is consistent with
136 Revell et al. (2019), who showed that a more recent version of our model simulates cloud droplet concentrations
137 and aerosol optical depth values that are lower than observed over the Southern Ocean in the Southern
138 Hemisphere summer. However, in the Southern Hemisphere winter Revell et al., (2019) simulated higher
139 aerosol optical depths than observed, which they corrected by reducing the dependence of sea spray emissions
140 on wind speed. Hence, our constraint on sea spray emission fluxes may only be appropriate for Southern
141 Hemisphere summer when wind speeds are relatively low. We do not make any assumptions about the
142 composition of these additional summertime sea spray particles. They may be rich in organic material as
143 proposed by Gantt et al. (2011) which would alter the CCN activity of emitted particles. However, the
144 consistency of constraint of CCN_{0.2} and N₇₀₀ towards higher values (Fig. 2, table S3) implies that a general
145 scaling of the existing sea spray flux is consistent with the measurements from December to April, without the
146 need for an additional source of fine-mode, organic-rich particles.

147
148 The dry deposition velocity of accumulation mode aerosols (Dry_Dep_Acc) has an 84% likelihood of being
149 lower than the default model value after applying the constraint. Furthermore, deposition velocities larger than
150 around 3 times the default value are effectively ruled out. This constraint is consistent with the higher aerosol
151 concentrations implied by constraint of the sea spray emission parameter.

152
153 Other parameters are more modestly constrained. The constraint on the aerosol precursor DMS emission flux
154 scale factor is two-sided, reducing the credible range of DMS emission scalings from 0.5 to 2.0 down to 0.54 to
155 1.9. This constraint suggests the default surface sea water concentration (Kettle and Andreae, 2000) and
156 emission parameterisation (Nightingale, et al., 2000) are consistent with measurements (including aerosol
157 sulfate) and do not benefit from being scaled. Furthermore, ACE-SPACE measurements are consistent with
158 less-efficient aerosol scavenging (55% likelihood of Rain_Frac, the parameter that controls the fractional area of
159 the cloudy part of model grid boxes where rain occurs, being below the unconstrained median value 0.5) and
160 less aqueous phase sulfate production (pH of cloud droplets has a 62% likelihood of being lower than the
161 unconstrained median value). These combined constraints suggest, in agreement with sea spray and deposition
162 parameter constraints, higher aerosol number and mass concentrations are consistent with measurements.

163
164 The effects of measurement constraint on pdfs of RF_{aci} and ERF_{aci} are shown in Fig. 4. Removing implausible
165 model variants has reduced the uncertainty in several parameters including natural aerosol emission fluxes,

166 which translates into a reduction in RF_{aci} uncertainty (Carslaw et al., 2013). The measurement constraints have
167 two important effects on aerosol forcing. Firstly, the magnitude of median RF_{aci} weakens from -1.99 W m^{-2} to -
168 1.88 W m^{-2} (-1.64 to -1.49 W m^{-2} for ERF_{aci}). A weaker forcing is consistent with higher natural aerosol
169 emissions, increased aerosol load and higher cloud droplet number concentrations in the early-industrial period.
170 Table 1 shows that our constraint on natural emission parameters also constrains Southern Ocean cloud droplet
171 number concentrations towards higher values, reducing the credible interval by around 50% and bringing mean
172 values into closer agreement with MODerate Imaging Spectroradiometer (MODIS; Salomonson et al., 1989)
173 instrument data (note that droplet number concentrations were not used to constrain the model). Thus, we
174 conclude that the constraint on aerosol forcing towards weaker values is a genuine constraint and not the result
175 of an arbitrary tuning. Secondly, the constrained forcing pdfs are approximately symmetric but have shorter tails
176 (lower kurtosis). This suggests the constraints are selectively ruling out model variants that are outliers. The
177 95% credible range of RF_{aci} values is reduced by around 9% (from -2.84 to -1.15 W m^{-2} down to -2.64 to -1.10
178 W m^{-2}) and around 9% for ERF_{aci} (from -2.69 to -0.62 W m^{-2} down to -2.43 to -0.54 W m^{-2}). The consistency of
179 forcing constraint across two distinct PPEs suggests the results are insensitive to differences in meteorology,
180 parameters perturbed in the PPEs, and the inclusion of rapid atmospheric adjustments. These results are also
181 insensitive to additional constraint to ensure energy balance at the top of the atmosphere (Fig. S5).

182
183 Johnson et al. (2019) reduced the global, annual mean RF_{aci} uncertainty by constraining multiple anthropogenic
184 emission and model process parameters (as well as some natural aerosol parameters) using over 9000
185 predominantly Northern Hemisphere measurements of aerosol optical depth, $PM_{2.5}$, particle number
186 concentrations and mass concentrations of organic carbon and sulfate. We used the same methodology as
187 Johnson et al. (2019) to rule out implausible model variants from the same original sample of one million model
188 variants, so we can readily combine these constraints. Around 700 model variants (0.07%) are observationally
189 plausible in both the Southern Ocean (ACE-SPACE) and Johnson et al. (2019) constraints. Although this is a
190 relatively small percentage of the original sample, 700 observationally-plausible model variants is far more than
191 are typically used to quantify model uncertainty or multi-model diversity (e.g. around 30 for CMIP6). The
192 marginal parameter pdfs from this 700-member sample are shown in Fig. 5. Because Johnson et al. (2019)
193 studied only the AER PPE (from which RF_{aci} can be computed) we are unable to explore the effect of the
194 combined constraint on ERF_{aci} .

195
196 The two measurement datasets constrain distinct groups of parameters. There are a few cases where the same
197 parameters are constrained by both datasets and in these cases the parameter values are constrained consistently
198 (e.g. cloud droplet pH) or more strongly through ACE-SPACE (e.g. sea spray emissions). The complementary
199 nature of these constraints means that the combined constraint marginal parameter pdfs (Fig. 5) are remarkably
200 similar to those in our Fig. 3e (for sea spray and DMS emission fluxes, as well as deposition and pH parameters)
201 and in figure 6 of Johnson et al. (2019) for other parameters.

202
203 The Johnson et al. (2019) constraint reduced the RF_{aci} uncertainty by around 6% and our ACE-SPACE
204 measurement constraint reduced the uncertainty by around 9%. However, the RF_{aci} uncertainty is reduced by
205 around 21% (Fig. 6a) after applying both constraints, meaning the combined constraint is stronger than the sum
206 of individual constraints.

207
208 The Johnson et al. (2019) constraint strengthened the RF_{aci} by around 0.3 W m^{-2} (more negative) because the
209 largest sea spray emission flux scaling and largest new particle formation rates were ruled out (Fig. 6 in Johnson
210 et al., 2019). Our ACE-SPACE constraint rules out the same large sea spray emission fluxes, but also rules out
211 all emission flux scale factors lower than around 1.6 (Fig. 3), which increases the baseline aerosol concentration
212 in the early-industrial atmosphere. The ACE-SPACE measurements also constrain several other parameters that
213 collectively weaken the median RF_{aci} by around 0.18 W m^{-2} . Therefore, using the combined measurement
214 dataset, the strongest RF_{aci} values have been ruled out as implausible and the credible range of observationally
215 plausible RF_{aci} values is reduced to around -2.51 to -1.17 W m^{-2} (-2.18 to -1.46 W m^{-2} , when using one standard
216 deviation to quantify the uncertainty). Uncertainty in RF_{ari} is reduced by around 48% with observationally
217 plausible values ranging from -0.27 to -0.09 W m^{-2} (-0.23 to -0.13 W m^{-2} , when using one standard deviation),
218 because the strongest RF_{ari} values are ruled out as observationally implausible.

219 3 Discussion

220
221 Our results show, as hypothesised from previous sensitivity analyses, that remote marine measurements are
222 valuable for constraining the natural aerosol state of the atmosphere (Carslaw et al., 2013; Regayre et al., 2014;

223 Regayre et al., 2018). Remote marine aerosol measurements provide new information about plausible model
224 behaviour because they are closely related to model emissions and processes that measurements in polluted
225 environments do not constrain.

226
227 For the first time we have achieved a meaningful reduction of 21% in the RF_{aci} uncertainty by constraining the
228 aerosol properties in the model. The reduction in forcing uncertainty can still be improved by considering the
229 following: Firstly, using measurements of cloud properties and cloud-aerosol relations, as well as measurements
230 associated with primary sulfate and carbonaceous particle emission sizes, could constrain model parameters that
231 cause RF_{aci} uncertainty but are not constrained by a combination of Northern Hemisphere and pristine Southern
232 Ocean measurements. Secondly, even within the considerably reduced volume of multi-dimensional parameter
233 space there still exist many compensating parameter effects (Fig. S3), which limit the constraint on individual
234 parameter ranges (Lee et al., 2016; Regayre et al., 2018). The impact of these compensating effects could be
235 greatly reduced by perturbing uncertain emissions regionally rather than globally as we do here. Our results are
236 based on uncertainty in a single climate model. The model is structurally consistent in our experiments, so
237 neglects uncertainty caused by choice of microphysical and atmospheric process representations. Our model
238 also neglects some potentially important sources of remote marine aerosol, such as primary marine organic
239 aerosol (Mulcahy et al., 2020) and methane-sulfonic acid (Schmale et al., 2019; Hodshire, et al., 2019; Revell et
240 al., 2019). Model inter-comparison projects (such as CMIP6) can be used to quantify the diversity of RF (or
241 ERF) output from models, but they lack information about single model uncertainty. Ideally, multi-model
242 ensembles would contain a perturbed parameter component, so that model diversity and single model forcing
243 uncertainty could be quantified simultaneously. But, computational costs prevent many modelling groups from
244 engaging with this important aspect of uncertainty quantification, limiting our shared knowledge about the
245 causes of aerosol forcing uncertainty. Studies like ours that quantify the remaining uncertainty in aerosol forcing
246 and its components after constraint using multiple measurement types fill an important knowledge gap. This
247 knowledge can be used to form a more complete understanding of the importance of historical and near-term
248 aerosol radiative forcing which would reduce the diversity in equilibrium climate sensitivity across models.
249

250 **Data availability**

251 The ACE-SPACE data are accessible from: <https://zenodo.org/communities/spi-ace>. The basis for our cloud
252 droplet number concentration data are available from
253 <http://catalogue.ceda.ac.uk/uuid/cf97ccc802d348ec8a3b6f2995dfbfbff>. Simulation output data for both AER and
254 AER-ATM PPEs are available on the JASMIN data infrastructure (<http://www.jasmin.ac.uk>). Some of the
255 climate-relevant fields are derived and stored in netCDF files (.nc) containing data for all ensemble members
256 and made available as a community research tool as described in Yoshioka et al. (2019). Model data and
257 analysis code can be made available from the corresponding author upon request.

258 **Author Contribution**

259 LR applied the statistical methodology and generated results. LR and MY created the PPEs. LR and JJ designed
260 the experiments and elicited probability density functions of all aerosol parameters. KC and MY participated in
261 the formal elicitation process. JS, AB, MG, CT, SH and FS collected and processed the ACE-SPACE
262 measurements. DG processed the cloud droplet number concentration data. LR, KS, JS and JJ analysed the
263 results. LR and KS wrote the manuscript with contributions from all authors.

264 **Competing Interests**

265 Author KC is an executive editor of ACP.

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442 Table 1. Annual and monthly mean cloud drop number concentrations over the Southern Ocean (over the region between
443 50°S and 60°S at around 1km altitude above sea level) in the original unconstrained sample and the sample of model variants
444 constrained to ACESPACE campaign measurements. Mean values and 95% credible interval values are shown for each
445 sample, with interquartile ranges in brackets. For comparison, we show cloud drop concentrations calculated from MODIS
446 instrument data following Grosvenor et al., (2018) for the year 2008 (SI Methods: Measurements).

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	Annual	December	January	February	March	April
MODIS (cm ⁻³)	73	89	91	90	82	63
Unconstrained mean (cm ⁻³)	38	39	39	41	42	39
Unconstrained credible interval (cm ⁻³)	7-125 (112)	8-115 (103)	8-117 (109)	7-122 (115)	7-129 (122)	7-118 (111)
Constrained mean (cm ⁻³)	66	67	69	72	76	70

Constrained credible interval (cm^{-3})	41-96 (55)	43-96 (53)	44-99 (55)	45-105 (60)	47-111 (64)	44-101 (57)
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Fig. 1. a,c) Mean and b,d) standard deviation of $\text{CCN}_{0.2}$ concentrations from the a,b) unconstrained sample and c,d) the sample constrained using concentration measurements of $\text{CCN}_{0.2}$, $\text{CCN}_{1.0}$, non-sea-salt sulfate and particles with dry aerodynamic diameters larger than 700 nm. Measured $\text{CCN}_{0.2}$ values are plotted as dots. Means and standard deviations were calculated using samples taken from emulators trained using monthly mean values. December to March sample values were combined based on longitudinal agreement with measurements.

Fig. 2. Unconstrained (black) and observationally constrained (red) pdfs of aerosol properties: a) $\text{CCN}_{0.2}$, b) $\text{CCN}_{1.0}$, c) N_{700} and d) aerosol sulfate. The pdfs were calculated at locations where measurements were used for constraint across the months December to March. Densities for each sample of model variants are scaled so that the area under the curve integrates to one. The green dashed line shows the median of the measurements and the dotted green lines show the approximate uncertainty ranges due to multiple model-measurement comparison uncertainties that were accounted for in the constraint (See SI Methods).

Fig. 3. Marginal probability distributions for the 26 aerosol parameters after constraint using ACE-SPACE measurements. The density of parameter values in the unconstrained sample are shown as horizontal dashed lines (uniform sampling over the parameter space). Densities of constrained samples are shown in colour and are scaled so that the maximum densities in the constrained and unconstrained samples are aligned. The 25th, 50th and 75th percentiles of each marginal distribution are shown in the central boxes. Parameter values on the x-axes correspond to values used in the model (Yoshioka et al., 2019, table S3).

Fig. 4. Probability distributions of a) RF_{aci} and b) ERF_{aci} . The distributions of the unconstrained sample of one million model variants from statistical emulators of each PPE are in black. Red lines show the distributions after constraint using ACE-SPACE measurements (around 3% of the unconstrained sample). The 25th, 50th and 75th percentiles of each sample are shown as shaded boxes and dashed lines span the 2.5th and 97.5th percentiles.

Fig. 5. Marginal probability distributions for the 26 aerosol parameters after constraint using around 250 Southern Ocean measurements and more than 9000 aerosol measurements in Johnson et al. (2019). Plotting features of this figure are identical to Fig. 3.

Fig. 6. Probability distributions of a) RF , b) RF_{aci} and c) RF_{ari} from the unconstrained (black line) and constrained (red line) samples of model variants. The constrained sample includes model variants that agree with our ACE-SPACE measurement constraint and the Johnson et al. (2019) constraint. Plotting features are identical to Fig. 4.