correspondence

Icelandic volcanic emissions and climate

To the Editor — On 31 August 2014, a large non-explosive fissure eruption of more than 1 km³ of lava began in Iceland, about 45 km from the Bárðarbunga volcano^{1,2}. For more than 100 days the 'Nornahraun' eruption emitted an average of about 35 kilotons of sulphur dioxide (SO₂) per day into the atmosphere¹; almost three times the daily anthropogenic emissions of SO_2 by the 28 European Union member states in 2010³. Our global climate model simulations of the eruption (Supplementary Information) suggest that if the eruption were to continue into the spring and summer of 2015, its radiative forcing over the North Atlantic and Western Europe would exceed, in absolute value, that of all current human aerosol emissions in the same region.

During September 2014, peak SO₂ emissions reached 120 kilotons per day, although there is substantial uncertainty associated with the SO₂ emission fluxes. So far, the eruption has not emitted any significant amounts of volcanic ash. Even though SO₂ emissions from the Nornahraun eruption do not reach the stratosphere, this type of eruption can have significant effects on regional climate by increasing the albedo of low-level liquid clouds^{4–6}, thereby reflecting more solar radiation back to space (Fig. 1).

Our model simulations assume emissions of 40 kilotons per day into an altitude range of 1,500 m to 3,000 m, and account for different meteorological conditions (Supplementary Information). Our simulations predict significant regional radiative forcing over Europe and the North Atlantic (45-70° N, 50° W–15° E), estimated from differences with control simulations (without the eruption). In the autumn months (September-November), radiative forcing amounts to -0.21 W m⁻² $(\sigma = 0.05)$ (Fig. 1), of which a striking 80% $(\sigma = 5\%)$ is attributable to albedo effects of sulphur on liquid clouds in solar wavelengths. Simulations indicate that should this level of emissions occur in summer (June-August), -7.4 W m^{-2} ($\sigma = 1.0$) of radiative forcing would result, with 94% attributable to indirect effects ($\sigma = 2\%$).

By comparison, based on similar simulations, we estimate -1.5 W m⁻² and -5.8 W m⁻² of radiative forcing from all anthropogenic aerosol emissions in autumn and summer, respectively, in this same region. During summer the radiative effects are larger due to a greater solar flux and a higher burden of sulphates from gas-phase oxidation. The simulated change in regional aerosol optical depth (Δ AOD, Fig. 1) due



Figure 1 | Simulated changes due to the Nornahraun eruption in Iceland. **a,b**, Differences in aerosol optical depth (Δ AOD, unitless) at visible wavelengths (550 nm) between simulations with and without volcanic sulfur dioxide emissions during September-November (**a**) and June-August (**b**). **c,d**, Differences in radiative flux at the top of the atmosphere (Δ TOA) for the same simulations, for September-November (**c**) and June-August (**d**).

to the eruption would be hard to detect in autumn (mean $\triangle AOD = 0.005$, $\sigma = 0.001$), but may be detectable in satellite data during boreal summer for similarly large eruptions (mean $\triangle AOD = 0.03$, $\sigma = 0.006$, but local $\triangle AOD$ up to 0.1). At present, there are limited ground-based aerosol data in this region.

The Nornahraun eruption provides an unprecedented opportunity to observe aerosol-cloud interactions induced by continuous volcanic degassing into a region of high cloud susceptibility⁷. This magnitude of indirect forcing by aerosols may significantly alter regional modes of climate variability, even without reaching the stratosphere. Sensitivity tests (Supplementary Information) indicate that the radiative forcing is sensitive to different atmospheric flow conditions in the region during the eruption. The radiative perturbation of the Northern Hemisphere (Fig. 1) may result in atmospheric circulation anomalies. Similar climate impacts may have occurred during historic Icelandic volcanic eruptions (for example, Laki in AD 1783-1784)8.9.

Local SO₂ and particle emissions are currently being analysed, but more detailed regional measurements are needed of the various gaseous and particulate species from both ground-based and satellite platforms^{4,5,10}. Permanent ground-based instrumentation in Iceland would be valuable to augment existing global networks. This would enable detailed evaluation of climate simulations of the impacts of this and similar eruptions.

References

- 1. 100 Days of Continuous Eruptive Activity in Holuhraun (Icelandic Met Office, 2014); http://en.vedur.is/media/jar/bb100days_ens.pdf
- 2. Sigmundsson, F. et al. Nature 517, 191–195 (2014).
- 3. Emission Trends of Sulphur Oxides in EU28 Countries (European Environment Agency, 2014); http://go.nature.com/OBGTcT
- Yuan, T., Remer, L. A. & Yu, H. Atmos. Chem. Phys. 11, 7119–7132 (2011).
- 5. Gassó, S. J. Geophys. Res. 113, D14S19 (2008).
- 5. Schmidt, A. et al. Atmos. Chem. Phys. 12, 7321-7339 (2012).
- 7. Twomey, S. Atmos. Environ. 25, 2435-2442 (1991).
- 8. Thordarson, T. & Self, S. J. Geophys. Res. 108(D1), 4011 (2003).
- 9. Schmidt, A. et al. Atmos. Chem. Phys. 10, 6025-6041 (2010).
- Ebmeier, S. K., Sayer, A. M., Grainger, R. G., Mather, T. A. & Carboni, E. Atmos. Chem. Phys. 14, 10601–10618 (2014).
- Additional information

Supplementary information is available in the online version of the paper.

Andrew Gettelman^{1*}, Anja Schmidt² and Jón Egill Kristjánsson³

¹National Center for Atmospheric Research, PO Box 3000, Boulder, Colorado 80305, USA. ²School of Earth and Environment, University of Leeds, Leeds LS2 9JT, UK. ³Department of Geosciences, University of Oslo, PO Box 1022, Blindern, 0315 Oslo, Norway. *e-mail: andrew@ucar.edu

Published online: 16 February 2015 Corrected online: 25 February 2015