Aerosol indirect effects on the nighttime Arctic Ocean surface from thin, predominantly liquid clouds

Lauren M. Zamora[1,2]*, Ralph A. Kahn[2], Sabine Eckhardt[3], Allison McComiskey[4], Patricia Sawamura[5,6], Richard Moore[5], Andreas Stohl[3]

[1]former NASA Postdoctoral Program Fellow, Universities Space Research Association; Now at Earth System Science Interdisciplinary Center, University of Maryland, College Park, College Park, MD, USA

[2]NASA Goddard Space Flight Center, Greenbelt, MD, USA

[3]NILU - Norwegian Institute for Air Research, Kjeller, Norway

[4]NOAA Earth System Research Laboratory, Boulder, CO, USA

[5]NASA Langley Research Center, Hampton, VA, USA

[6]Science Systems and Applications, Inc., Greenbelt, MD, USA

Correspondence to: Lauren M. Zamora (lauren.m.zamora@nasa.gov)

Abstract. Aerosol indirect effects have potentially large impacts on the Arctic Ocean surface energy budget, but model estimates of regional-scale aerosol indirect effects are highly uncertain and poorly validated by observations. Here we demonstrate a new way to quantitatively estimate aerosol indirect effects on a regional scale from remote sensing observations. In this study, we focus on nighttime, optically thin, predominantly liquid clouds. The method is based on differences in cloud physical and microphysical characteristics in carefully selected clean, average and aerosol-impacted conditions. The cloud subset of focus covers just ~5% of cloudy Arctic Ocean regions, warming the Arctic Ocean surface by ~1-1.4 W m^{-2} regionally during polar night. However, within this cloud subset, aerosol and cloud conditions can be determined with high confidence using CALIPSO and CloudSat data and model output. This cloud subset is generally susceptible to aerosols, with a polar nighttime estimated maximum regionally integrated indirect effect of ~0.11 W m^{-2} at the Arctic sea ice surface. Aerosol presence is related to reduced precipitation, cloud thickness, and radar reflectivity, and in some cases, an increased likelihood of cloud presence in the liquid phase. These observations are consistent with a thermodynamic indirect effect hypothesis and are inconsistent with a glaciation indirect effect. However, this cloud subset shows large differences in surface and meteorological forcing in shallow and higher altitude clouds and between sea ice and open ocean regions. For example, optically thin, predominantly liquid clouds are much more likely to overlay another cloud over the open ocean, which may reduce aerosol indirect effects on the surface. Also, shallow clouds over open ocean do not appear to respond to aerosols as strongly as over stratified sea ice environments, indicating a larger influence of meteorological forcing over aerosol microphysics in these types of clouds over the rapidly changing Arctic Ocean.

1 Introduction

Aerosol indirect effects on clouds are among the biggest uncertainties in climate models (Boucher et al., 2013). It is particularly important to reduce these uncertainties in the Arctic, where warming is occurring at a faster rate than in other locations (Serreze et al., 2009), and where local aerosol indirect effects can be large (Garrett et al., 2004; Garrett and Zhao, 2006; Lubin and Vogelmann, 2006; Zhao and Garrett, 2015). Understanding aerosol indirect effects is also important because aerosol emissions within and in the vicinity of the Arctic are changing, and perhaps more importantly, the major aerosol removal processes and transport pathways to the Arctic may be changing as well (Jiao and Flanner, 2016).

Unfortunately, accurate observation-based estimates of regional mean forcings are very difficult to obtain at most locations around the planet due to a variety of confounding factors and errors. These include: 1) a reliance on proxies for cloud condensation nuclei (CCN) and ice nucleating particles (INP), 2) meteorological co-variability and other synoptic-scale surface and atmospheric factors, including the aerosol spatial distribution, 3) the complexity of cloud responses to aerosol type and amount (Fan et al., 2016), 4) spatial and temporal limitations of the datasets, and 5) an insufficient understanding of cloud characteristics even in the absence of anthropogenic aerosols (Ghan et al., 2016; Wilcox et al., 2015). Knowledge of this last factor is difficult to obtain because pristine conditions are rare in most locations globally (Hamilton et al., 2014). To quantify mean regional aerosol indirect effects using observations, one would need datasets that cover the large spatial and temporal scales required to include the full range of natural heterogeneity, plus a way to correctly identify clean background conditions. As a result, current estimates of regional indirect aerosol impacts on the surface radiation rely predominantly upon models that still cannot accurately represent many relevant Arctic processes (e.g., Morrison et al. (2012); Ovchinnikov et al. (2014)).

In some ways, isolating aerosol indirect effects over the Arctic Ocean can be even more challenging than in other regions. Sampling conditions at the ground are harsh, there is low thermal and visible contrast between sea ice and clouds, and observations are limited by the frequent presence of multi-layer clouds. The very cold temperatures that characterize the Arctic affect chemical reactions and physical processes (e.g., the development of frost flowers, diamond dust, and blowing snow), making comparisons with lower latitude systems more challenging. However, the Arctic Ocean is ideal for the study of indirect effects in other ways. For example, the surface and meteorological conditions over sea ice are highly homogenous compared to many other regions of the world. Moreover, pristine conditions still occur in this region with relatively high frequency, despite periodic episodes of combustion-derived aerosol transport from lower latitudes. Current day observations in clean background conditions are among our best proxies for pre-industrial conditions (Hamilton et al., 2014), and a better understanding of pre-industrial conditions is in turn key to the ability to determine current-day indirect aerosol impacts on a regional scale (e.g., Gettleman (2015); Ghan et al. (2016; 2013); Carslaw et al. (2013); Wilcox et al. (2015); Kiehl et al. (2000)).

Here we present a method for identifying spatially distributed properties in a subtype of Arctic Ocean clean background clouds using a combination of the CALIPSO and CloudSat active remote sensing instruments and an atmospheric transport model. We use the difference between average cloud characteristics gathered across the Arctic Ocean and average clean background clouds over the same region to estimate the maximum regional indirect aerosol impacts on the surface. This calculation provides an estimate of the actual regional impact of aerosol indirect effects on the surface including aerosol-meteorological co-variability after stochastic meteorological effects have been taken into account. We also examine differences between the cloud characteristics under various aerosol conditions to assess cloud formation mechanisms in the presence of aerosol.

One goal of this work is to illustrate one way that regional-scale aerosol indirect effects on the surface can be obtained quantitatively from observational data. In the past, such estimates have primarily been supplied only by models. We focus on the subset of Arctic Ocean clouds where aerosol impacts can be identified with the greatest certainty: optically thin (cloud optical depth, COD < 3), predominantly liquid clouds during polar night. Optically thin, liquid-containing clouds are generally common over this region (Bennartz et al., 2013; Shupe and Intrieri, 2004). Such clouds are also effective at radiating longwave (LW) radiation downward (e.g., Garrett and Zhao (2006)), thus having a potentially large contribution to surface forcing (Shupe and Intrieri, 2004). Moreover, models tend to under-predict the formation of these optically thin clouds at supercooled temperatures (Cesana et al., 2012), making aerosol influences on droplet characteristics and ice nucleation of particular interest. Within the larger liquid-containing cloud group, this study focuses on predominantly liquid clouds, where aerosol conditions can be assessed with highest certainty. The analysis is also limited to nighttime samples both to improve CALIPSO aerosol-condition assessments and to reduce confounding impacts from direct and semi-direct effects.

2 Methods

2.1 Sample selection

To describe aerosol impacts on Arctic Ocean clouds with high confidence using CALIPSO and CloudSat data, it was vital to our method that we be able to identify clean background cases accurately. We selected a specific group of clouds where non-background aerosol (hereafter referred to simply as "aerosol") conditions and cloud properties could be ascertained with the greatest confidence. The main Arctic Ocean cloud subset of focus consists of clouds that are Optically thin (COD < ~3), Nighttime, predominantly Liquid clouds, henceforth referred to as "ONLi" clouds for brevity. Because the ONLi cloud profiles were taken only at night, the majority of them were collected during the winter when there are relatively high aerosol inputs from lower latitudes (Shaw, 1995). Within the full ONLi cloud group, we identified the subsets of clouds present in clean background and aerosol-influenced conditions. Results were also compared with an internal subset of clouds where aerosol conditions and cloud properties could be ascertained with even higher confidence (i.e., those clouds that were

Measured > One km above the surface, Optically thin (COD < ~3), collected at Nighttime, predominantly Liquid, and from the Top-layer, henceforth referred to as "MOONLiT" clouds). The criteria for the cloud groups and aerosol classifications are summarized in Table 1. Justification for these criteria and descriptions of the individual datasets used for sample selection are described in more detail below.

2.1.1 CALIPSO

Aerosol vertical distribution, cloud top height, cloud base height, cloud optical depth, and initial approximate cloud phase were obtained from the polar-orbiting CALIPSO satellite lidar v. 3.01 level 2, 5-km aerosol profile and cloud layer products at 532 nm. These data have a vertical resolution of 30 m within the vertical region where most predominantly liquid Arctic Ocean clouds were found (up to 8 km asl). Before averaging, along-track cloud profile data were collected at a horizontal resolution of 1/3 km. Averaged aerosol data have a horizontal resolution of between 5-80 km, with the horizontal resolution decreasing with aerosol concentration. For example, in clear air with no detected aerosols, the horizontal resolution is 80 km; in strong aerosol layers, the horizontal resolution providing adequate signal-to-noise can be as low as 5 km (Vaughan et al., 2009).

Because our samples were taken at night, Moderate Resolution Imaging Spectroradiometer (MODIS) optical depths were not available. Instead, the CALIPSO product was used to measure CODs, as it offers substantially higher data availability in the optical thickness range of interest (COD < 3) than CloudSat (Christensen et al., 2013). Only non-quality-flagged (i.e., the highest quality) CALIPSO COD data were used. CALIPSO cloud optical depth uncertainties rise with COD due to uncertainties in the lidar ratio in liquid clouds with COD > 1 (CALIPSO Quality Statements: Lidar Level 2 Cloud and Aerosol Layer Products, Version releases: 3.01, 3.02). We excluded COD data with uncertainties $\geq$ 75% of the COD value (these constituted ~5% of all cases).

Because it can be difficult to accurately separate Arctic aerosol from diamond dust and thin ice clouds using backscatter data (*M. Vaughan*, pers. comm.; Grenier and Blanchet (2010)), we focused on CALIPSO liquid-containing clouds. To gain greater confidence in the aerosol classification within the MOONLiT subset, ice clouds were not allowed in those profiles. Note that CALIOP cloud "phase" indicates only whether the cloud predominantly contained liquid or ice; there is no mixed-phase designation. At a later step, CloudSat data were used to further refine cloud phase information.

CALIPSO data were obtained over the Arctic Ocean between 60-82°N and between 1 January 2008 – 7 December 2009 (during the latter part of CloudSat epoch 2). To obtain the lowest possible comparable detection limit, the analysis was restricted to nighttime clouds. Here, nighttime profiles are taken in the CALIPSO orbit over the hemisphere of Earth that is dark at any given time, and so the borders of this hemisphere may include some low-light conditions. MOONLiT clouds were additionally restricted to upper-layer clouds only. We focused on ONLi clouds present between 0.2 and 8 km above the surface to enable better below-cloud aerosol detection. MOONLiT cloud cases were further restricted to above 1 km for

better comparison to high-quality CloudSat data. Very few predominantly liquid clouds are expected above 8 km. Clouds were included only when the feature's optical properties scored between 70 and 100 in the cloud-aerosol discrimination (CAD) algorithm (a high confidence cloud determination) (Liu et al., 2009). The lidar-determined presence of a below-cloud aerosol layer was a key criterion in identifying clean background clouds with confidence, as discussed further in Sect. 3.1.

5 Thus, the analysis was limited to non-opaque clouds (COD < ~3), as determined by the 532 nm Extinction Quality Control flag.

The "clean, background" cloud subset met the above criteria, but no aerosol features were permitted above or below cloud, even when air masses had been horizontally averaged across 80-km resolution in the CALIPSO aerosol detection algorithm,

10 which is the resolution that detects weak aerosol layers with highest confidence. Given these constraints, the backscatter aerosol detection limit for "clean background" clouds is as low as possible, and should have only negligible variations based on detector noise and background molecular and O_3 densities above cloud (Vaughan et al., 2009). Because CALIPSO cannot always detect dilute aerosols (Di Pierro et al., 2013; Kacenelenbogen et al., 2014; Rogers et al., 2014; Winker et al., 2013), particularly below-cloud where the lidar signal has been reduced, "clean background" clouds were also required to have

15 modeled above and below-cloud FLEXPART ("FLEXible TRAjectory model", (Stohl et al., 1998, 2005)) black carbon concentrations of < 30 ng C m^{-3} (see Sect. 2.1.3 and 3.1 for further discussion). The "aerosol-influenced" subset had aerosols with CAD scores between -100 and -70 (high confidence aerosol classification) above or below the cloud and FLEXPART modeled below-cloud black carbon concentrations of > 30 ng C m^{-3}. The geographical distributions of the all-cloud, clean-cloud, and aerosol-influenced cloud sets are shown in Fig. 1.

20 **2.1.2 CloudSat**

CloudSat cloud profiling radar data are collected at a vertical resolution of 240 m. CloudSat has a wider swath than CALIPSO (1.4x1.8 km) and it takes measurements on the same polar orbit, only seconds ahead of CALIPSO. Because the CloudSat radar does not accurately estimate cloud properties below ~0.7-1 km agl (Huang et al., 2012; Mioche et al., 2015). CloudSat data were provided only for clouds with bases ≥ 0.75 km agl. Some of the very thin clouds detected by CALIPSO

25 had radar reflectivities that were too low to be detected by CloudSat, and CloudSat may sometimes mistakenly assign precipitating ice as a cloud (de Boer et al., 2008). Therefore, radar reflectivity data and CloudSat reflectivity-derived cloud parameters, where available, were obtained from the height bins closest to where CALIPSO detected a cloud.

Average reflectivity between the CALIPSO-determined cloud top and base was obtained from the CloudSat 2B-GEOPROF version R04 dataset. Cloud phase and precipitation occurrence were acquired from 2B-CLDCLASS-LIDAR version R04

30 estimates (Wang, 2013). In this product, cloud phase is determined from a combination of CALIPSO water layer detection and integrated backscattering coefficient, temperature, CloudSat reflectivity, and an assumed temperature-dependent reflectivity threshold for ice particles (Zhang et al., 2010). This phase classification is uncertain for clouds with reflectivities

of < -29 dBZ (the CloudSat sensitivity limit), and for very thin clouds due to the coarse vertical resolution of the instrument. As we focused on cold, optically thin clouds in this study, many (~29%) of our samples were below the CloudSat detection limit. Thus, phase was only assessed in clouds with cloud phase certainty values of > 5 and with reflectivity values of > -29 dBZ. Infrequently, clouds that met the CALIPSO criterion in Table 1 were classified as predominantly ice phase by the 2B-

5   CLDCLASS-LIDAR product; these cases were excluded from the analysis for simplicity, despite the potential for supercooled water to be misclassified as ice particles (Van Tricht et al., 2016).

Estimated mean liquid cloud droplet effective radii (r_{el}) were obtained from the CloudSat 2B-CWC-RO version R04 product (LO_RO_effective_radius) (Austin and Stephens, 2001). We chose this CloudSat r_{el} product, which assumes that all particles are liquid, for two reasons: 1) CALIPSO had independently assigned the clouds a predominantly liquid phase, and 2)

10   uncertainties in the other liquid r_{el} data product available for nighttime samples (RO_liq_effective_radius) may be fairly high because of a reliance on an overly-simplistic, temperature-dependent phase partitioning scheme (e.g., de Boer et al. (2008); Lee et al. (2010)). Where available, r_{el} data were averaged over vertical regions within the CALIOP-determined "liquid" phase cloud base and top. Sometimes the corresponding CloudSat-determined cloud base and top were slightly different. In these cases, CALIOP heights were used because of its better ability to detect liquid droplets, and because CloudSat may

15   sometimes misclassify precipitating ice as part of the cloud (de Boer et al., 2008), which can lead to overestimation of r_{el}. Quality-flagged data were excluded, such as observations from precipitating clouds, as determined from the CloudSat 2B-CLDCLASS-LIDAR version R04 product. Note: although we counted the number of cases where precipitation occurred for comparison at a different step, precipitating cases were otherwise excluded from most other derived cloud parameters in the analysis. These cases were excluded in order to obtain comparable data across cloud characteristics, which was particularly

20   important for the longwave emissions calculations detailed in Sect. 2.2 that included the r_{el} as one of several input parameters.

We present some limited CloudSat-derived r_{el} data here, but it is important to note the fairly high uncertainties in some of these data. Aside from the assumption of liquid phase, there is a known bug in the CloudSat code that might cause r_{el} in liquid clouds to be overestimated, and to our knowledge there has been no extensive validation of the CloudSat 2B-CWC-

25   RO r_{el} product in the Arctic. de Boer et al. (2008) found fairly reasonable agreement, with perhaps some overestimation, between CloudSat-determined r_{el} in mixed-phase clouds compared to r_{el} measured from ground-based instruments. However, only a few samples were collected with the in-cloud constraint in that study. The cumulative uncertainties in r_{el} on the radiative impact results are discussed further in Sect. 3.5.

2.1.3 FLEXPART

30   The locations of combustion aerosol plumes were modeled using BC from the FLEXPART model (Stohl et al., 1998, 2005). The FLEXPART model has been used extensively to study pollution and smoke transport in the Arctic, and is well-validated

for this purpose (Damoah et al., 2004; Eckhardt et al., 2015; Forster et al., 2001; Paris et al., 2009; Sodemann et al., 2011; Stohl et al., 2002, 2003, 2015). We chose BC as a combustion aerosol tracer because it represents aerosol removal better than a gaseous tracer like carbon monoxide, and because FLEXPART can largely capture the Arctic BC seasonal cycle (Eckhardt et al., 2015) that is driven by a combination of seasonal changes in emissions, atmospheric transport patterns and removal processes. In some cases, wildfires can emit large amounts of light absorbing organic carbon aerosols (or "brown carbon") without emitting large amounts of BC (e.g., Chakrabarty et al. (2016)). In these cases, FLEXPART BC may not represent smoke aerosols well.

For this study, as in Eckhardt et al. (2015), FLEXPART was driven with meteorological analysis data from the European Centre for Medium-Range Weather Forecasts (ECMWF) at a resolution of 1˚ longitude and 1˚ latitude. BC emissions were based on the ECLIPSE emission inventory (Stohl et al., 2015), which also includes emissions from gas flaring, and biomass burning emissions. In the model simulations, BC was removed from the atmosphere through dry deposition, and wet scavenging both below and within clouds. However, no transformation of BC from a hydrophobic to a hydrophilic state was considered and removal parameters were chosen as typical for a hydrophilic aerosol. FLEXPART-modeled BC concentrations were calculated for the years 2008 and 2009 at a horizontal resolution of 1° latitude and 2° longitude and at 0.05, 0.2, 1, 2, 3, 5, 7, and 10 km agl. Below-cloud BC concentrations were taken to be the closest modeled concentration available to 0.5 km below cloud base. When there were multi-layer clouds and the next cloud top was < 1 km away, the concentration closest to the middle distance between the two clouds was used instead.

2.2 Ancillary datasets

Aircraft out-of-cloud black carbon data were obtained from NASA's Arctic Research of the Composition of the Troposphere from Aircraft and Satellites (ARCTAS) campaign (Fuelberg et al., 2010; Jacob et al., 2010; Kondo et al., 2011). The aircraft data with highest aerosol particle concentrations were clustered between 50-60° N during this campaign. Thus, we included aircraft data from between 50-82° N (subarctic + Arctic) in order to assess comparable ranges of dilute and concentrated aerosols expected to be present over the Arctic. Submicron aerosol dry size distributions between 0.06–1 μm were measured from a DMT Ultra-High Sensitivity Aerosol Spectrometer (UHSAS) between 0-2.1 km (2.9 km for springtime samples). Submicron aerosol scattering data at 532 nm were obtained from a Radiance Research (RR) nephelometer and were corrected for truncation errors. Submicron aerosol scattering coefficients at 450 and 700 nm were estimated as the difference between total scattering from a TSI 3563 Integrating Nephelometer and the RR nephelometer when the fine mode aerosol fraction exceeded 0.6. Ambient total scattering coefficients at the three wavelengths were obtained from the TSI nephelometer, and were corrected for truncation errors following Anderson and Ogren (1998). Aerosol absorption coefficients at 450, 532, and 700 nm were measured with a RR three-wavelength Particle Soot Absorption Photometer (PSAP).

An aircraft-derived, 180° backscatter coefficient is calculated following Sawamura et al. (2017) in order to compare the in situ data to that from CALIOP (units of Mm^{-1} sr^{-1}). First, the measured dry, submicron aerosol size distribution, scattering coefficient, and absorption coefficient at 532 nm are input into a Mie theory model to determine the aerosol effective dry refractive index. Next, a hygroscopic growth factor was applied to the dry size distribution in the Mie theory model to reproduce observed humidified light scattering and thus derive the aerosol refractive index at ambient relative humidity. The 180° backscatter coefficient then follows from Mie theory using the adjusted size distribution and refractive index. This method is best suited for spherical particles, which we assume dominate the ARCTAS samples based on the main aerosol sources during the campaign (non-dust background aerosols, anthropogenic pollution and smoke (Jacob et al., 2010)).

Several other supplemental datasets were used for cloud environmental context. ETOPO1 Bedrock GMT4 data (Amante and Eakins, 2009) were used to identify cloud profiles over the Arctic Ocean region. NOAA/NSIDC Climate Data Record of Passive Microwave Sea Ice Concentration, version 2 data (Meier et al., 2013; Peng et al., 2013) were used to approximate the fractional sea ice cover over ocean at the specific month and location of each profile. A sample was classified as being primarily over sea ice or open ocean when the sea ice fraction at the given location and month was > 80% or < 20%, respectively.

Lastly, integrated surface longwave (4-30 μm) radiation was calculated with an updated Santa Barbara DISTORT Atmospheric Radiative Transfer program (SBDART, (Ricchiazzi et al., 1998)). Shortwave effects are not expected to be significant during nighttime conditions. Following McComiskey and Feingold (2008), the calculations assume homogeneous cloud cover and spectrally uniform surface albedo. Median surface longwave reflectivity (R) for open ocean and sea ice in clear conditions with no clouds or aerosols (0.64 and 0.69, respectively) was calculated from MERRA 2 output (GMAO, 2015) based on the times and locations of the data and the following formula (Josey, 2003):

$$(1) \qquad R = 1 - \frac{E - A}{I},$$

where E is the emitted longwave radiation from the surface, A is the net longwave flux into the surface from the atmosphere, and I is the downwelling longwave radiation from the atmosphere. Note: the A parameter above is proxied by the closest available parameter in the MERRA2 output, surface absorbed longwave radiation, and thus it does not include factors such as transmission, latent heat, or conduction and convection. Because even a 50% change in R would lead to < 1% error in the cloud longwave surface flux calculations, we expect the resulting uncertainty in R to have negligible impact on our results.

3 Results and Discussion

3.1 Correct identification of clean background conditions

To accurately characterize clean background conditions, it is necessary to detect combustion-related aerosol layers with confidence. For CALIPSO, dilute aerosols are least likely to be detected below-cloud due to signal attenuation inside the cloud (Di Pierro et al., 2013), but CALIOP can sometimes miss dilute aerosol layers even in clear air above clouds (Di Pierro et al., 2013; Kacenelenbogen et al., 2014; Rogers et al., 2014; Sheridan et al., 2012; Winker et al., 2013). Most previous works focused either on daytime samples, which have comparatively low signal-to-noise ratios, or on extinction data, which are more uncertain because they assume a prescribed lidar ratio. To begin quantifying the false negative rate relevant to this study, we used two independent methods to estimate the fraction of the time when nighttime Arctic CALIPSO data would not detect above-cloud aerosols when actually present.

First, we estimated the fraction of air masses containing various observed concentrations of aerosol tracers that would be detected at the reported theoretical 80 km resolution nighttime backscatter detection limit from Winker et al. (2009). This analysis is based on co-located aircraft backscatter, particle number, and BC data from the ARCTAS aircraft campaign (Fig. 2a). The results suggest that CALIOP would miss ~36% of slightly polluted air masses (i.e., BC concentrations > 30 ng m^{-3}, or CN$_{PCASP}$ concentrations > 127 particles cm^{-3}) at 80 km resolution in nighttime air masses not below another feature. This estimate might be affected by errors from assuming Mie theory and a theoretical detection limit that may not be perfectly representative in the field, as well as errors caused by a limited amount of field data from scattered locations.

As an independent consistency check, we next determined the frequency at which aerosols were detected by both FLEXPART and CALIOP. To do so, we compared the fraction of observed clear sky (no-cloud) CALIOP profiles that were expected to contain aerosols at different simulated FLEXPART aerosol concentrations for January 2008 (Fig. 2b). These results suggested that CALIOP may not have detected up to ~33% of slightly polluted air masses (BC > 31 ng m^{-3}) above cloud, although this value likely overestimates the actual false negative rate given inherent model errors. This independent estimate is fairly similar to the previously estimated false negative rate, and so we expect the real-world above-cloud CALIOP false negative rate for dilute aerosols to be ~33-36%. Below-cloud errors would be higher, but are more difficult to quantify because of the variability of in-cloud attenuation.

Based on CALIPSO criteria alone, the above estimates suggest that aerosol detection uncertainties may be higher than desireable, particularly below cloud. We address this issue in two ways. First, we apply the criteria for determining clean background cloud that depend not only on aerosol-free CALIPSO profiles, but also on modeled above- and below-cloud BC concentrations of < 30 ng m^{-3} (see Sect. 2.1.3). We expect the model aerosol-occurrence criterion to substantially improve the classification confidence because coincidences of false negatives in both the CALIOP data and the model are likely to be rare (they are most likely to occur in dilute aerosol conditions). As such, this method should correctly identify clean

background clouds much more frequently than 64-67% of the time. Unfortunately, further quantification in the classification confidence is difficult because both model accuracy and the degree of below-cloud lidar attenuation are variable in time and space. Secondly, we assess the MOONLiT cloud subset along with ONLi cloud results. MOONLiT clouds are a subset of ONLi clouds that, among other criteria meant to enhance certainty in aerosol layer identification, are in the top layer (see

5   Sect. 2.1 and Table 1 for more details). Trends in MOONLiT cloud results are mainly noted only if they are dissimilar to those in the larger ONLi cloud group, and are otherwise provided in the supplementary material. To our knowledge, the combined CALIPSO and model criteria used here allow the most confident classification of background conditions currently possible for remote sensing studies of the Arctic.

3.2 Notes on limitations imposed by the methods

10   In order to have greater confidence in quantifying the regional scale aerosol indirect effects, this study is limited to ONLi clouds and their MOONLiT cloud subset. It is important to emphasize that the ONLi cloud group is not representative of all Arctic clouds. During our study period, ONLi clouds were present in only 5.3% of all total comparable nighttime cloudy profiles over the Arctic Ocean ("comparable clouds" defined as having a satisfactory in-cloud CAD score of 70-100 and with cloud bases > 200 m to exclude fog). Liquid-dominated clouds tend to be found at lower altitudes than thicker opaque clouds

15   and thus may not always be identified in multi-layer clouds using CALIPSO. However, even though the actual prevalence of these clouds may be somewhat underestimated, it is clear that ONLi clouds represent just a small fraction of all Arctic clouds. Thus, we emphasize that the aerosol indirect responses described in this paper are not necessarily representative of Arctic clouds in general.

Moreover, the cloud-selection criteria imposed by our methods may induce some uncertainties in the analysis. For example,

20   due to the low COD constraint, it is possible that some fraction of the cloud subset influenced by aerosols may be selected from a different group of cloud types than some fraction of the clean background cloud subset. As an illustration, in a subarctic aircraft case study presented in Zamora et al. (2016) (see Appendix A for further details), cumulus clean background clouds with an observed cloud thickness of ~250 m had CODs of ~5. These clouds would have been too optically thick for the CALIOP lidar to penetrate. However, highly comparable nearby clouds in a smoke plume had CODs

25   of only ~2, and the cloud-property differences were likely driven by the aerosol (Zamora et al., 2016). In this example, only the subset of clouds influenced by smoke aerosols would have met this study's COD criterion and not the clean background cloud counterparts. Median reductions in COD were fairly minor for aerosol-impacted clouds relative to background clouds, and were not significant over open ocean, and so we do not expect this effect to have a large impact on our study.

30   Similarly, any aerosol-driven phase changes that shifted clouds between predominantly ice- and liquid-containing clouds (e.g., Girard et al. (2013)) could have eliminated or added samples from/to our study, also potentially adding some bias to our results. These uncertainties are difficult to quantify, but are likely to be much smaller than the error that would be

introduced by expanding the dataset to include other, non-ONLi cloud subsets that would be characterized with greater uncertainty.

3.3 ONLi cloud characteristics in clean marine background conditions

In our study, sampled clouds were thin by definition and were thus unlikely to occur under very turbulent conditions. The range in turbulence covered in the sample set was also likely limited during polar night due to the lower variability in external heating and generally high static stability of the Arctic atmosphere. Nonetheless, we expect that clouds over the open ocean are impacted more by thermodynamic coupling with the surface (Shupe et al., 2013) than over sea ice, where surface-based inversions occur more frequently (Ganeshan and Wu, 2015). In this study, we stratify clouds into these two regimes, to distinguish the effects of systematic differences in atmospheric stability and large-scale atmospheric and surface forcing between the two systems (Curry et al., 1996; Jaiser et al., 2012; Taylor et al., 2015).

ONLi clouds were much more likely to overlay another cloud layer over open ocean than over sea ice, as demonstrated by the average height of the next below-cloud feature (Fig. 3b, Table 2). A similar result was also observed previously at the SHEBA ship-based observatory (Intrieri et al., 2002). There are also differences between shallow and higher clouds. Shallow clouds are defined here as having cloud bases < 1.1 km asl, based on the lower quartile range of the cloud base height data. Over both open ocean and sea ice, shallow clouds are warmer and are more likely to have a liquid- vs. mixed-phase CloudSat designation (Tables S1 and S2). Shallow clouds are on average optically thicker, but geometrically thinner, than higher clouds. They are also less likely to be observed in multi-layer cloud conditions in both regimes ($p < 0.05$, permutation test), which may be due in part because they are systematically less observable due to lidar attenuation in higher thick cloud layers.

It is possible that some of the differences between shallow and high ONLi clouds are due to differences in cloud formation mechanisms. For example, previous studies suggest that shallow liquid-containing Arctic clouds might form from the advection of warm, moist air over a cool surface, whereas higher liquid-containing clouds might form from a longwave radiative flux divergence (Smith and Kao, 1995) or partial dissolution of a higher-level stratus cloud (Herman and Goody (1976). One previous model sensitivity study linked shallow liquid-containing clouds in a 3-day Arctic multi-layer cloud system with surface turbulent heat fluxes, and overlying liquid-containing clouds with large-scale advection and maintenance by radiative cooling (Luo et al., 2008).

The different probabilities of cloud-layering occurrence over sea ice vs. open ocean and in cloud properties over different heights complicates comparisons between the two regimes. However, comparing only single-layer clouds with bases above 1.1 km, the median cloud base height of open ocean clouds is ~240 m higher (~480 m for MOONLiT clouds) than for clouds over the sea ice ($p < 0.05$, permutation test). Autumn ship-based cloud observations in the Chukchi and Beaufort Seas also show higher cloud bases over the open ocean [*Sato et al.*, 2012; *Young et al.*, 2016]. Over sea ice, the lower cloud heights

and the presence of fewer multi-layer ONLi clouds compared to the open ocean (Table 2) are likely related to the lower height and greater frequency of surface-based inversions over Arctic sea ice, which can reduce surface moisture fluxes to higher altitudes (Bradley et al., 1992; Ganeshan and Wu, 2015; Zhang et al., 2011). Below 1.1 km, cloud base heights for single-layer clouds are not significantly different between regimes.

5   Over the open ocean, clouds were also warmer than over sea ice, and a higher fraction of ONLi clouds were observed with very low layer mean reflectivity (Z_m), defined as $Z_m < -29$ dBZ (the CloudSat detection limit) (Table 2). The very low Z_m clouds are geometrically and optically very thin (Table 2). Previously observed relationships between Z_m and r_{el} suggest that the very low Z_m clouds also likely have smaller r_{el} values (Frisch et al., 2002).

Because reflectivity was fairly low within the thin, predominantly liquid cloud profiles that fit our criteria, and temperatures
10   were generally between -1 to -28 ^{o}C, in many cases it was difficult to know for certain which clouds were of mixed vs. liquid phase. Of the clouds that were assigned a high-confidence phase classification by CloudSat, most contained some ice particles (93%, n=5238 for sea ice, and 79%, n=2992 for open ocean). We believe it likely that a comparatively higher fraction of the very low Z_m clouds were present in the liquid-only phase. First, these clouds had very low Z_m values (indicative of small particles), and at the same time they were independently assigned a predominantly liquid phase by
15   CALIPSO. Secondly, their median temperatures were warmer than clouds with higher Z_m (by ~1-3°C over sea ice, and nearly 1-7°C over comparable altitudes over open ocean, Table 2). Relatedly, low Z_m clouds were more than two times more likely to be found over open ocean than over sea ice (Table 2). Further study would be needed to fully verify phase for this cloud subset, but the indications that these clouds have higher liquid fractions are consistent with the observations that a) Arctic liquid clouds tend to have smaller r_{el} values than mixed-phase clouds (Hobbs and Rangno, 1985; Lance et al., 2011;
20   Lebo et al., 2008; Rangno and Hobbs, 2001), and b) clouds over the open ocean (which were more likely to have very low Z_m values (Fig. 4 a,d)) are also more likely to be liquid-containing (Cesana et al., 2012).

3.4 Aerosol impacts on clouds over sea ice

We expect that the greater uniformity in surface and meteorological conditions over sea ice will increase the likelihood of being able to isolate aerosol impacts from meteorological noise, compared to the situation over the open ocean, and cloud
25   characteristics were indeed fairly uniform over sea ice. We observed only minor differences in cloud base height between ONLi clouds present in clean background conditions and all ONLi clouds (Table 2). Above 1.1 km, the cloud base temperatures in clean background conditions were not significantly different from those in all air mass conditions. Below 1.1 km, clean background clouds appear to be found in slightly warmer conditions (by ~2 ^{o}C) (Table S1).

30   Clean background clouds were significantly more likely to be precipitating than other clouds in both height bins (Table 2). This observation falls in line with aerosol-driven reductions in snowfall that have been predicted and observed previously,

inside and outside of the Arctic (Albrecht, 1989; Borys et al., 2000, 2003; Girard et al., 2005; Lance et al., 2011; Lohmann et al., 2003; Mauritsen et al., 2011; Morrison et al., 2008). These observed reductions in precipitation are inconsistent with the glaciation indirect effect, in which ice formation would be expected to increase due to higher concentrations of combustion-related INP (Lohmann and Feichter, 2005). The presence of aerosols is also correlated with a significant reduction in radar

5   reflectivity, generally associated with smaller particles on theoretical grounds (Fig. 4, Table 2). Correspondingly, there is also a significantly higher probability that clean background clouds detected by CALIPSO would also be detected by CloudSat than in all clouds or in aerosol-impacted clouds (Table 2).

The r_{el} values are derived from radar reflectivity, and as such, aerosol-related decreases in reflectivity suggest smaller r_{el}

10   values. This observation follows expectations based on the Twomey effect, and is in line with previous studies in the Arctic that have observed smaller r_{el} correlated with increasing influence of aerosols (Coopman et al., 2016; Lubin and Vogelmann, 2006; Peng et al., 2002; Tietze et al., 2011; Zamora et al., 2016; Zhao and Garrett, 2015). Here, the cloud droplet effective radius decreased systematically as expected aerosol influence rose, and the estimated mode r_{el} was respectively 10.3, 10.0, and 9.8 μm for the ONLi clean cloud, all cloud, and the aerosol-influenced cloud subsets. This reduction was similar in the

15   MOONLiT subset, at 10.5, 10.3, and 10.0 μm, respectively (Table S1). Unfortunately, the differences in r_{el} are available only for the thicker clouds that CloudSat was able to observe, and in some cases, data were available only for the middle sections of clouds, which are expected to have higher relative r_{el} values. Thus, the estimated mean r_{el} values presented here might be skewed higher than would be derived from a dataset that more fully sampled the cloud fields, and the differences compared to clean background cases could underestimate actual differences. The difference in estimated ONLi r_{el} is about half of a

20   previously reported, regionally integrated value for all Arctic clouds. Using MODIS r_{el} estimates in thicker clouds (median COD ~ 11) with temperatures between 0-2 ^{o}C, Tietze et al. (2011) saw an ~1 μm difference between the very cleanest clouds and median clouds. Note that these regionally averaged net changes in r_{el} are much smaller than would be expected locally in very polluted clouds (e.g., Zamora et al. (2016)).

25   There are differences between cloud thicknesses in clean background air and other air masses that suggest the potential for meteorological co-variability in the samples. Clean ONLi clouds are optically and geometrically thinner than the other cloud groups. Lower moisture associated with continental airflow that carries the aerosol might explain this difference (Lohmann and Feichter, 2005), if recent surface contact with warmer mostly mid-latitude regions did not enhance moisture. However, in two related remote sensing studies where Arctic clouds were tightly binned within related meteorological groups, COD

30   differences still appeared, and thus the authors attributed these differences to aerosol-driven changes in liquid water path (LWP) (Coopman et al., 2016; Tietze et al., 2011).

It is difficult to say whether the aerosol-related impacts on precipitation and radar reflectivity observed here are simply related to Twomey effects on liquid droplets, or whether some more complex mixed-phase and/or meteorological dynamics

are also involved. One possibility is that the expected aerosol-driven reductions in r_{el} may hinder the transition from liquid to mixed-phase clouds due to preferential freezing of larger particles (Lohmann and Feichter, 2005; Morrison et al., 2012). This possibility is supported by a small but significant increase in the portion of detected liquid phase clouds within sea ice clouds above 1.1 km (Tables 2 and S1). This observation was not significant in the MOONiT cases, perhaps due to very small sample size (Table S3). One previous aircraft-based study offered some evidence to suggest that the thermodynamic indirect effect is important in the Arctic, particularly for thin clouds (Jackson et al., 2012). However, low sample number and surface/ meteorological variability made this mechanism difficult to conclusively demonstrate on a larger scale. In a different study using CloudSat and CALIPSO, no strong evidence of this process was found (Grenier and Blanchet, 2010). That study was also inconclusive because of high uncertainties related to the reliance on an above-cloud sulfate aerosol proxy, and a focus on ice phase clouds where it is more difficult for CALIPSO to accurately separate aerosols from ice particles. If the very low Z_m ONLi clouds in our study do indeed contain fewer cases with ice particles (see Sect. 3.3 above), the greater presence of very low Z_m clouds in aerosol-influenced conditions (Fig. 4) would also support the possibility of the thermodynamic indirect effect dominating within the ONLi cloud subset. As more information is needed to verify phase in very low Z_m clouds, for now this possibility remains a conjecture.

Other possible mechanisms that could explain the observed aerosol-related impacts on cloud properties are that polluted air might contain fewer ice nucleating particles (INP) than clean background air (Borys, 1989), that solutes might lower the homogeneous freezing temperature and reduce INP efficiency (Girard and Asl, 2014; Koop et al., 1998), that differential contact nucleation could play some role (Ladino Moreno et al., 2013; Morrison et al., 2005), and/or that riming efficiency could be reduced (Lohmann and Feichter, 2005).

3.5 Aerosol impacts on clouds over the open ocean

Whereas cloud properties over sea ice were relatively tightly constrained, there was a much larger range in cloud properties over the open ocean (Table 2) that may in part reflect the greater variability and higher magnitudes of surface turbulent heat and moisture fluxes over open ocean (e.g., (Morrison et al., 2008; Strunin et al., 1997; Taylor et al., 2015)). Variability reduced our ability to compare clouds within this regime, as did the uneven vertical distribution of aerosols. CALIPSO-detected aerosols in the Arctic are most frequently found at altitudes below 2 km (Devasthale et al., 2011b; Di Pierro et al., 2013; Kafle and Coulter, 2013; Winker et al., 2013). At 2.1 km, the median ONLi cloud base was above this level over the open ocean, and the 2.6 km median cloud base in the clean background cloud subset was even higher. Thus, the difference in median cloud altitude between the different subsets likely induces a categorical bias in the cloud properties shown in Table 2.

To both account for aerosol height differences and retain a sample size from our 2-year dataset that was as informative as possible, we separated clouds found over open ocean into three cloud-base-height bins (Table S2), and summarized the

resulting information in Table 2. As over sea ice, the first bin includes clouds with base heights between 0.2 and 1.1 km. This range encompasses the lower quartile range of all open ocean clouds, isolating the shallow clouds that were observed to have different characteristics from the higher clouds in clean background conditions (Sect. 3.3). This range also happens to coincide with the lower quartile range of sea ice clouds so that these two bins are more or less comparable to each other with respect to cloud-base height. The second bin covers 1.2-3.2 km (the interquartile range of open ocean clouds). The last bin includes clouds with bases > 3.2 km. Although aerosol-influenced clouds still appear most often near the bases of their bins, the median cloud height and temperature differences within bins are fairly small (Table S1).

There are some significant differences between clouds with and without aerosol influence in ONLi clouds with bases above 1.1 km. Similarly to clouds over sea ice, radar reflectivity is reduced with higher aerosol influence, and the fraction of low Z_m clouds increases (Table 2). Median r_{el} dropped by 0.4 μm in aerosol-impacted cases vs. clean background cases, compared to a 0.5 μm reduction over sea ice. Clouds with bases > 1.1 km, especially those at higher altitudes, are also thinner.

The reflectivity and r_{el} trends were not consistently observed in the MOONLiT subset, likely because smaller sample size caused the lack of statistical confidence in the binned samples (see Table S3). However, in a similar study using MODIS data for liquid clouds over the Arctic, Coopman et al. (2016) found significant trends in r_{el} with greater predicted aerosol concentrations when they stratified their results by lower tropospheric stability (LTS), which is much greater over sea ice than over open ocean (Taylor et al., 2015). Like us, they found that the trends were weaker for regions with less expected LTS (which in our case would be over open ocean). The MOONLiT subset also had a significantly greater fraction of clouds that were assigned a liquid phase in aerosol-influenced samples compared to clean background samples for clouds where high quality CloudSat phase information was available above 3.2 km. This trend was not observed in the ONLi cloud subset, potentially because the differences between clean and aerosol-influenced cases were more ambiguous than in the MOONLiT cloud subset, but the trend toward more liquid clouds in aerosol-influenced conditions was also observed in the higher ONLi cloud bin over sea ice. It is unclear whether a similar trend in phase would remain if more of the samples had contained high-quality phase data, so we can only remark that the association between aerosols and liquid phase clouds is not inconsistent with the thermodynamic indirect effect.

In contrast to clouds found at higher levels, there were not many significant differences associated with aerosol-influence in ONLi clouds with bases below 1.1 km. Moreover, some of the differences that were significant were small enough to not be very meaningful (e.g., a 20 m reduction in mean cloud base height with a corresponding 0 m difference in median cloud base height for clean clouds compared to all ONLi clouds). This observation suggests that dynamics might be overwhelming any aerosol changes to cloud microphysics in this regime, although our sample size for CloudSat derived parameters was reduced by only assessing those clouds that were > 750 m above the surface to avoid ground clutter of the instrument. Median cloud

base heights in aerosol-influenced clouds were slightly higher (120 m) than clean clouds, which might have contributed to slightly colder cloud top heights.

3.6 Upper bounds on regional surface radiative impacts

5   Over our two-year time period, we identified tens of thousands of predominantly liquid ONLi clouds over the Arctic Ocean (Table 2). This sample size and regional spread of the data are large enough that we make the assumption that the cloud characteristics provided in Table 2 approximate the net nighttime cloud characteristics that exist for this cloud subset after exposure to the full spectrum of environmental conditions in each regime (sea ice and open ocean). We calculated the *maximum regional* radiative impact of clean background ONLi clouds on the nighttime surface, based on the regional

10   frequency of occurrence of observable ONLi clouds in nighttime profiles over the entire (cloudy or clear) Arctic Ocean during our time period (2.52% and 4.84% over sea ice and open ocean, respectively; 3.23% over the full Arctic Ocean domain). Table 2 clean background cloud characteristics were used to calculate longwave flux changes to the surface compared to clear air, assuming cloud homogeneity and a single cloud layer, estimated at 56.05-58.44 W m^{-2} and 20.86-21.48 W m^{-2} for sea ice and open ocean regions, respectively. Maximum regional radiative impacts were estimated by

15   multiplying these longwave fluxes by the ONLi cloud regional frequency of occurrence. Note that the presence of lower-level clouds will reduce the regional impact of ONLi clouds on the surface. Variable input parameters for the radiative impact calculations included cloud base height, cloud thickness and COD, and r_{el} for clouds over sea ice and open ocean. Parameter values were taken from Table 2 median values, except for r_{el}, where the interquartile range was used to reflect the larger uncertainty in that parameter.

The estimated maximum regional radiative impact of clean background ONLi clouds during polar night was between 1.41-1.47 W m^{-2} over sea ice and 1.01-1.04 W m^{-2} over open ocean. Maximum regional ONLi cloud impacts on the surface were smaller over the open ocean in part due to lower cloud temperatures associated with higher median cloud altitudes (an effect also seen during the SHEBA campaign (Shupe and Intrieri, 2004)). This effect occurred despite there being more ONLi

25   cloud cover over open ocean than over sea ice (a general trend that is also observed in total cloud fraction (Kay and L'Ecuyer, 2013)). Also, the higher open ocean clouds are expected to have lower liquid water paths (based on thinner CODs, Table 2), which influences longwave cloud forcing in very thin clouds that are not opaque in the infrared (Turner, 2007). For reference, using the CloudSat 2B-FLEXHR-LIDAR product, Kay and L'Ecuyer (2013) estimated the annual mean longwave forcing at the surface due to all clouds over sea ice and open ocean to be ~24-36 and 32-56 W m^{-2}, respectively, depending

30   on location. Barton et al. (2014) model-mean estimates for cloud impacts on surface longwave downwelling radiation during polar night over sea ice above 70 ^{o}N (within the 95% confidence interval for surface temperatures) were ~15-30 W m^{-2}. These published estimates included the impacts of non-ONLi clouds, which the current study does not.

We also estimated the maximum regional surface indirect radiative effect of aerosols on ONLi clouds over sea ice. To do so, we subtracted the maximum regional surface radiative impacts of the clean background cloud subset from the impacts expected of all observed ONLi clouds. Radiative calculations were not made for aerosol-driven effects on ONLi clouds over the open ocean due to the lack of significant differences in most relevant parameters and the altitude-based bias in the full

5 open ocean dataset. As with background clouds, aerosol-indirect radiative effect estimates were made using the median cloud base and top heights, the median COD, and the r_{el} interquartile range for sea ice clouds presented in Table 2. Based on this information, we estimate that excluding changes in cloud fraction, aerosols could have indirectly decreased current-day surface downwelling longwave fluxes during polar night over sea ice, from ONLi clouds specifically, by no more than 0.11 W m^{-2}, integrated over sea ice across the Arctic and for all aerosol concentrations. As with the background cloud estimates,

10 this spatially integrated estimate assumes single layer cloud conditions. Estimated regional aerosol indirect impacts specifically from the shallow (base height < 1.1 km) sea ice ONLi clouds accounted for about half of this effect. In this instance, holding all other variables equal, aerosol-related changes in cloud optical depth were an order of magnitude more important for radiative effects than the changes in cloud droplet effective radius, and the changes in geometric thickness had nearly no impact on the longwave impacts. It is important to note that because this range is spatially integrated across the

15 Arctic, local aerosol impacts in strong haze layers can be much higher (e.g., Garrett et al. (2004); Carrió et al. (2005); Zhao and Garrett (2015)). For example, Zhao and Garrett (2015) found that the local cloud indirect longwave forcing in single-layer stratus clouds at Barrow, Alaska in the upper quartile of combustion aerosol concentrations was 8.1-9.9 W m^{-2} greater than in clouds associated with the lower quartile of combustion aerosol concentrations. In a similar study at Barrow, Lubin and Vogelmann (2006) used the lower and upper quartile of aerosol particle concentrations to show that downwelling flux

20 for high CN cases was 3.4 W m^{-2} higher than for low CN cases.

To be clear, in estimating mean aerosol indirect effects in this section, we did not isolate absolute or local indirect aerosol effects from the confounding effects of meteorology and meteorological co-variability. Instead, we estimated the current-day impact of combustion-derived aerosols on the regional indirect effect that ultimately influences the current-day surface

25 radiation (which includes any meteorological co-variability present during these two years). This study was limited to only two years of data; future studies with more data might provide a better representation of the full range in aerosol and meteorological conditions the Arctic experiences over longer timescales.

As a final note, in this study we did not account for any aerosol-driven changes in cloud fraction. Aerosol-driven changes in

30 cloud fraction may have occurred, given the reduced precipitation and the shift in CloudSat-estimated cloud type from predominantly altocumulus to predominantly stratocumulus in increasingly aerosol-impacted conditions over sea ice (Table 2). If aerosols do increase cloud fraction, this effect could be the most important indirect impact that aerosols have on the Arctic's surface radiation budget, because the presence of cloud where there otherwise would not be one has more of a local

impact on surface radiation than does a change to a cloud that is already present (Feingold et al., 2016; Sedlar and Devasthale, 2012; Shupe and Intrieri, 2004). Addressing these issues will require further study with additional types of data.

4 Summary and Conclusions

Aerosol indirect effects have uncertain, but potentially large, impacts on the Arctic Ocean surface energy budget. As a step
5   toward reducing uncertainty in current-day aerosol regional indirect effects on the surface, here we have better constrained the characteristics of a small subset of clean, average and aerosol-impacted clouds for which we have relatively strong constraints on cloud properties and the associated aerosol environment. We focused on optically thin (COD<~3), predominantly liquid clouds collected at nighttime, which we termed "ONLi" clouds; they cover about 3% of the nighttime Arctic Ocean (5% of total non-fog cloudy regions). However, within the ONLi cloud subset, it was possible to gain a high
10 confidence in classification of clean background conditions with existing satellite remote sensing data. Using combined CALIPSO, CloudSat, and model output, we identify clean background clouds with a frequency that is much better than 64-67% of the time for top-layer clouds. Although the exact frequency of confident identification of clean background conditions beyond this range is difficult to quantify, particularly for clouds beneath another cloud layer, the level of confidence in clean background classification represents a substantial improvement compared to any previous remote
15 sensing study of the Arctic region, as best we know.

Within the ONLi cloud subset, we observed clear differences between clouds over open ocean and over sea ice, consistent with different surface and meteorological conditions in these two regimes. For example, when the surface is open ocean compared to sea ice, ONLi clouds are much more likely to overlay another cloud and to be present in liquid phase. A greater
20 frequency of multi-layer clouds over the open ocean might affect the retreat of sea ice, and in turn, how this changes the impact of clouds on surface radiation of the Arctic Ocean. However, further study is needed to expand this observation beyond just conditions that contain ONLi clouds. There were also noticeable differences between shallow ONLi clouds (cloud bases < 1.1 km) and higher ONLi clouds. As expected, shallow clouds were warmer and more likely be assigned a liquid- rather than mixed-phase CloudSat designation; they were also optically thicker and geometrically thinner. These
25 differences in cloud properties may be in part to due the differing cloud formation mechanisms for shallow clouds. Previous studies support this hypothesis (e.g., Herman and Goody (1976); Smith and Kao (1995); Luo et al. (2008)), as does the observation, from the present study, that shallow ONLi clouds are less sensitive to aerosols.

Except in shallow, open ocean clouds, we observed that ONLi clouds are susceptible to aerosols. Consistent with other
30 studies, the presence of aerosols exceeding background levels in clouds over sea ice is associated with reductions in r_{el}, cloud geometric and optical thickness, precipitation, radar reflectivity, and COD. Perhaps due to greater boundary layer turbulent fluxes, clouds over the open ocean appear to be less susceptible to the influence of aerosols, although some changes in phase and thickness were observed in the altitude-binned samples presented here. Due to aerosol-induced ONLi cloud changes

over sea ice, we estimate that the region-wide maximum surface radiation impact during polar night is ~0.11 W m^{-2}, with shallow clouds contributing about half of this signal. It is unclear from the current work what the impact over open ocean might be. In comparison, the maximum region-wide direct radiative impact of clean ONLi clouds at night is estimated to be 1.0 W m^{-2} and 1.4 W m^{-2} over sea ice and open ocean regions, respectively. Note that the presence of multi-layer clouds and cloud patchiness will reduce the radiative impact of ONLi clouds on the surface. Also, these maximum regional indirect effect estimates do not include any potential aerosol-driven changes in cloud extent, which could be important for estimating ONLi-cloud overall regional indirect effects. Thus, aerosol-driven changes in cloud fraction dominate the uncertainty in estimates of the overall indirect aerosol radiative impact on the nighttime Arctic surface energy balance, based on this method. Unfortunately, the cloud fraction over the Arctic Ocean is particularly difficult to constrain over short time scales with passive remote sensing, given the low contrast between clouds and sea ice and long polar nighttime conditions, and due to very limited spatial coverage for active remote-sensing..

We find no evidence to suggest that the glaciation indirect effect is important within the ONLi cloud subset. Beyond that, we have no strong support for aerosol impacts on mixed-phase cloud dynamics, although we see some tantalizing evidence to suggest that large liquid particles need be present for ice formation in non-shallow ONLi clouds, in a mechanism similar to the thermodynamic indirect effect. These findings are in line with and expand upon previous aircraft observations (Jackson et al., 2012). Aerosols were associated with higher fractions of liquid phase clouds than in clean background cases in both sea ice ONLi clouds > 1.1 km and in open ocean MOONLiT clouds > 3.2 km (for which additional cloud-selection criteria were applied; Table 1), for cases when high quality phase data were available. Above 1.1 km, open ocean and sea ice clouds influenced by aerosol were less reflective at 94-GHz. Where high quality CloudSat data were available, these clouds also had noticeably smaller estimated median r_{el} values, which is in line with previous studies. Over sea ice, aerosol-influenced clouds were less likely to be precipitating. Moreover, the fraction of low Z_m clouds increases with aerosol presence in both regimes and at all altitudes except in shallow open ocean clouds. These low Z_m clouds are more likely to be liquid-dominated, based on their lower radar reflectivity combined with their independently assigned predominantly liquid phase designation by CALIPSO, their warmer median cloud temperatures, and relatedly, their > 2 times higher relative fraction over open ocean compared to sea ice. Together, these observations suggest that aerosols could play an important role in ice nucleation and nighttime radiative heating via the thermodynamic indirect effect in ONLi clouds. However, more information on cloud phase in low-reflectivity clouds is necessary to more fully explore this possibility.

Although we limited this study to carefully describing average and clean background clouds within only a subset of remotely sensed Arctic Ocean clouds, we were able to provide a first observation-based estimate of regional scale aerosol indirect effects on the surface for such clouds, demonstrating one way in which remote sensing observations can be used to quantitatively assess aerosol-cloud interactions on a regional scale in other conditions and locations as well. Given that so far only models have been able to estimate regional aerosol indirect effects on the surface energy balance, this study lays an

important foundation for improving the quantification of aerosol indirect effects. The tradeoff for selecting a small subset of clouds in this study is the low representativeness of ONLi clouds. To constrain observation-based aerosol impacts and nucleation processes on a larger scale for the Arctic Ocean, optically thick and ice-containing clouds must also be included. Expanding this study to a longer time period would help better incorporate the natural variability in Arctic meteorology and

5 aerosols that might not be represented during this 2-year period. Including daylit or summertime air masses would also be useful; mid-summer air masses tend to be cleaner than wintertime Arctic air masses and have a higher fraction of liquid-containing clouds (Van Tricht et al., 2016). Moreover, it would enable the use of MODIS data to examine cloud phase (e.g., via the DARDAR data product (Delanoë and Hogan, 2010)) and droplet distribution.

Acknowledgements

10 We recognize and thank the efforts and funding from the large number of people and agencies involved in making the following datasets available, including: the NASA Langley Research Center Atmospheric Science Data Center, which provided the CALIPSO data, the *CloudSat* Data Processing Center run by the Cooperative Institute for Research in the Atmosphere (CIRA), which provided the CloudSat data, and the NASA ARCTAS program and members. Specifically we would like to thank Y. Kondo and B. Anderson for making their ARCTAS data publically available. We also thank G. de

15 Boer, J. Creamean, G. Feingold, K.B. Huebert, J. Limbacher, S. Platnick, A. Solomon, H. Telg, M. Vaughan, D.L. Wu, Y. Yang, H. Yu, and T.L. Yuan for helpful discussions. The research of L. Zamora was supported by the NASA ACMAP program, via an appointment to the NASA Postdoctoral Program at the NASA Goddard Space Flight Center, administered by Universities Space Research Association. The work of R. Kahn is supported in part by NASA's Climate and Radiation Research and Analysis Program under H. Maring, and NASA's Atmospheric Composition Program under R. Eckman. The

20 NILU team was supported by funding from NordForsk in the framework of eSTICC (eScience Tools for Investigating Climate Change at High Northern Latitudes).

Appendix A

In Zamora et al. (2016), the case study CODs were not presented. Here, we calculated the relevant CODs from the following relationship:

$$(2) \quad COD = \frac{3}{2} \frac{LWC \ (z_t - z_b)}{r_{el}},$$

where LWC is the liquid water content, z_t and z_b are cloud top and base height, respectively, and r_{el} is the cloud droplet effective radius.

Supplement link (to be provided by Copernicus)

Competing Interests:

5   The authors declare that they have no conflict of interest.

References

Albrecht, B. A.: Aerosols, Cloud Microphysics, and Fractional Cloudiness, Science, 245(4923), 1227–1230, doi:10.1126/science.245.4923.1227, 1989.

10   Amante, C. and Eakins, B. W.: ETOPO1 1 Arc-Minute Global Relief Model: Procedures, Data Sources and Analysis. NOAA Technical Memorandum NESDIS NGDC-24. National Geophysical Data Center, NOAA., , doi:10.7289/V5C8276M, 2009.

Anderson, T. L. and Ogren, J. A.: Determining Aerosol Radiative Properties Using the TSI 3563 Integrating Nephelometer, Aerosol Sci. Technol., 29(1), 57–69, doi:10.1080/02786829808965551, 1998.

15   Austin, R. T. and Stephens, G. L.: Retrieval of stratus cloud microphysical parameters using millimeter-wave radar and visible optical depth in preparation for CloudSat: 1. Algorithm formulation, J. Geophys. Res. Atmospheres, 106(D22), 28233–28242, doi:10.1029/2000JD000293, 2001.

Barton, N. P., Klein, S. A. and Boyle, J. S.: On the Contribution of Longwave Radiation to Global Climate Model Biases in Arctic Lower Tropospheric Stability, J. Clim., 27(19), 7250–7269, doi:10.1175/JCLI-D-14-00126.1, 2014.

20   Bennartz, R., Shupe, M. D., Turner, D. D., Walden, V. P., Steffen, K., Cox, C. J., Kulie, M. S., Miller, N. B. and Pettersen, C.: July 2012 Greenland melt extent enhanced by low-level liquid clouds, Nature, 496(7443), 83–86, doi:10.1038/nature12002, 2013.

de Boer, G., Tripoli, G. J. and Eloranta, E. W.: Preliminary comparison of CloudSAT-derived microphysical quantities with ground-based measurements for mixed-phase cloud research in the Arctic, J. Geophys. Res. Atmospheres, 113(D8),

25   D00A06, doi:10.1029/2008JD010029, 2008.

Borys, R. D.: Studies of ice nucleation by Arctic aerosol on AGASP-II, J. Atmospheric Chem., 9(1–3), 169–185, doi:10.1007/BF00052831, 1989.

Borys, R. D., Lowenthal, D. H. and Mitchell, D. L.: The relationships among cloud microphysics, chemistry, and precipitation rate in cold mountain clouds, Atmos. Environ., 34(16), 2593–2602, doi:10.1016/S1352-2310(99)00492-6,

30   2000.

Borys, R. D., Lowenthal, D. H., Cohn, S. A. and Brown, W. O. J.: Mountaintop and radar measurements of anthropogenic aerosol effects on snow growth and snowfall rate, Geophys. Res. Lett., 30(10), 1538, doi:10.1029/2002GL016855, 2003.

Boucher, O., Randall, D. A., Artaxo, P., Bretherton, C. S., Feingold, G., Forster, P. M., Kerminen, V.-M., Kondo, Y., Liao, H., Lohmann, U., Rasch, P., Satheesh, S. K., Sherwood, S., Stevens, B. and Zhang, X. Y.: Clouds and Aerosols, in Climate

Change 2013: The Physical Science Basis. Contribution of Working Group I to the Fifth Assessment Report of the Intergovernmental Panel on Climate Change, edited by T. F. Stocker, D. Qin, G. K. Plattner, M. Tignor, S. K. Allen, J. Boschung, A. Nauels, Y. Xia, V. Bex, and P. M. Midgley, Cambridge University Press, Cambridge, UK and New York, NY, USA., 2013.

5   Bradley, R. S., Keimig, F. T. and Diaz, H. F.: Climatology of surface-based inversions in the North American Arctic, J. Geophys. Res. Atmospheres, 97(D14), 15699–15712, doi:10.1029/92JD01451, 1992.

Carslaw, K. S., Lee, L. A., Reddington, C. L., Pringle, K. J., Rap, A., Forster, P. M., Mann, G. W., Spracklen, D. V., Woodhouse, M. T., Regayre, L. A. and Pierce, J. R.: Large contribution of natural aerosols to uncertainty in indirect forcing, Nature, 503(7474), 67–71, doi:10.1038/nature12674, 2013.

10   Carrió, G. G., Jiang, H. and Cotton, W. R.: Impact of Aerosol Intrusions on Arctic Boundary Layer Clouds. Part II: Sea Ice Melting Rates, J. Atmospheric Sci., 62(9), 3094–3105, doi:10.1175/JAS3558.1, 2005.

Cesana, G., Kay, J. E., Chepfer, H., English, J. M. and Boer, G. de: Ubiquitous low-level liquid-containing Arctic clouds: New observations and climate model constraints from CALIPSO-GOCCP, Geophys. Res. Lett., 39(20), L20804, doi:10.1029/2012GL053385, 2012.

15   Chakrabarty, R. K., Gyawali, M., Yatavelli, R. L. N., Pandey, A., Watts, A. C., Knue, J., Chen, L.-W. A., Pattison, R. R., Tsibart, A., Samburova, V. and Moosmüller, H.: Brown carbon aerosols from burning of boreal peatlands: microphysical properties, emission factors, and implications for direct radiative forcing, Atmos Chem Phys, 16(5), 3033–3040, doi:10.5194/acp-16-3033-2016, 2016.

Christensen, M. W., Stephens, G. L. and Lebsock, M. D.: Exposing biases in retrieved low cloud properties from CloudSat:

20   A guide for evaluating observations and climate data, J. Geophys. Res. Atmospheres, 118(21), 12,120-12,131, doi:10.1002/2013JD020224, 2013.

Coopman, Q., Garrett, T. J., Riedi, J., Eckhardt, S. and Stohl, A.: Effects of long-range aerosol transport on the microphysical properties of low-level liquid clouds in the Arctic, Atmos Chem Phys, 16(7), 4661–4674, doi:10.5194/acp-16-4661-2016, 2016.

25   Curry, J. A., Schramm, J. L., Rossow, W. B. and Randall, D.: Overview of Arctic Cloud and Radiation Characteristics, J. Clim., 9(8), 1731–1764, doi:10.1175/1520-0442(1996)009<1731:OOACAR>2.0.CO;2, 1996.

Damoah, R., Spichtinger, N., Forster, C., James, P., Mattis, I., Wandinger, U., Beirle, S., Wagner, T. and Stohl, A.: Around the world in 17 days - hemispheric-scale transport of forest fire smoke from Russia in May 2003, Atmos Chem Phys, 4(5), 1311–1321, doi:10.5194/acp-4-1311-2004, 2004.

30   Delanoë, J. and Hogan, R. J.: Combined CloudSat-CALIPSO-MODIS retrievals of the properties of ice clouds, J. Geophys. Res. Atmospheres, 115(D4), D00H29, doi:10.1029/2009JD012346, 2010.

Devasthale, A., TjernströM, M., Karlsson, K.-G., Thomas, M. A., Jones, C., Sedlar, J. and Omar, A. H.: The vertical distribution of thin features over the Arctic analysed from CALIPSO observations, Part I: Optically thin clouds, Tellus B, 63(1), 77–85, doi:10.1111/j.1600-0889.2010.00516.x, 2011a.

Devasthale, A., Tjernström, M. and Omar, A. H.: The vertical distribution of thin features over the Arctic analysed from CALIPSO observations. Part II: Aerosols, Tellus B, 63(1), doi:10.3402/tellusb.v63i1.16190, 2011b.

Di Pierro, M., Jaeglé, L., Eloranta, E. W. and Sharma, S.: Spatial and seasonal distribution of Arctic aerosols observed by the CALIOP satellite instrument (2006–2012), Atmos Chem Phys, 13(14), 7075–7095, doi:10.5194/acp-13-7075-2013, 2013.

Eckhardt, S., Quennehen, B., Olivié, D. J. L., Berntsen, T. K., Cherian, R., Christensen, J. H., Collins, W., Crepinsek, S., Daskalakis, N., Flanner, M., Herber, A., Heyes, C., Hodnebrog, Ø., Huang, L., Kanakidou, M., Klimont, Z., Langner, J., Law, K. S., Lund, M. T., Mahmood, R., Massling, A., Myriokefalitakis, S., Nielsen, I. E., Nøjgaard, J. K., Quaas, J., Quinn, P. K., Raut, J.-C., Rumbold, S. T., Schulz, M., Sharma, S., Skeie, R. B., Skov, H., Uttal, T., von Salzen, K. and Stohl, A.: Current model capabilities for simulating black carbon and sulfate concentrations in the Arctic atmosphere: a multi-model evaluation using a comprehensive measurement data set, Atmos Chem Phys, 15(16), 9413–9433, doi:10.5194/acp-15-9413-2015, 2015.

Feingold, G., McComiskey, A., Yamaguchi, T., Johnson, J. S., Carslaw, K. S. and Schmidt, K. S.: New approaches to quantifying aerosol influence on the cloud radiative effect, Proc. Natl. Acad. Sci., 201514035, doi:10.1073/pnas.1514035112, 2016.

Forster, C., Wandinger, U., Wotawa, G., James, P., Mattis, I., Althausen, D., Simmonds, P., O'Doherty, S., Jennings, S. G., Kleefeld, C., Schneider, J., Trickl, T., Kreipl, S., Jäger, H. and Stohl, A.: Transport of boreal forest fire emissions from Canada to Europe, J. Geophys. Res. Atmospheres, 106(D19), 22887–22906, doi:10.1029/2001JD900115, 2001.

Frisch, S., Shupe, M., Djalalova, I., Feingold, G. and Poellot, M.: The Retrieval of Stratus Cloud Droplet Effective Radius with Cloud Radars, J. Atmospheric Ocean. Technol., 19(6), 835–842, doi:10.1175/1520-0426(2002)019<0835:TROSCD>2.0.CO;2, 2002.

Fuelberg, H. E., Harrigan, D. L. and Sessions, W.: A meteorological overview of the ARCTAS 2008 mission, Atmos Chem Phys, 10(2), 817–842, doi:10.5194/acp-10-817-2010, 2010.

Ganeshan, M. and Wu, D. L.: An investigation of the Arctic inversion using COSMIC RO observations, J. Geophys. Res. Atmospheres, 120(18), 2015JD023058, doi:10.1002/2015JD023058, 2015.

Garrett, T. J. and Zhao, C.: Increased Arctic cloud longwave emissivity associated with pollution from mid-latitudes, Nature, 440(7085), 787–789, doi:10.1038/nature04636, 2006.

Garrett, T. J., Zhao, C., Dong, X., Mace, G. G. and Hobbs, P. V.: Effects of varying aerosol regimes on low-level Arctic stratus, Geophys. Res. Lett., 31(17), L17105, doi:10.1029/2004GL019928, 2004.

Gettelman, A.: Putting the clouds back in aerosol–cloud interactions, Atmos Chem Phys, 15(21), 12397–12411, doi:10.5194/acp-15-12397-2015, 2015.

Ghan, S., Wang, M., Zhang, S., Ferrachat, S., Gettelman, A., Griesfeller, J., Kipling, Z., Lohmann, U., Morrison, H., Neubauer, D., Partridge, D. G., Stier, P., Takemura, T., Wang, H. and Zhang, K.: Challenges in constraining anthropogenic aerosol effects on cloud radiative forcing using present-day spatiotemporal variability, Proc. Natl. Acad. Sci., 201514036, doi:10.1073/pnas.1514036113, 2016.

Ghan, S. J.: Technical Note: Estimating aerosol effects on cloud radiative forcing, Atmos Chem Phys, 13(19), 9971–9974, doi:10.5194/acp-13-9971-2013, 2013.

Girard, E. and Asl, N. S.: Relative importance of acid coating on ice nuclei in the deposition and contact modes for wintertime Arctic clouds and radiation, Meteorol. Atmospheric Phys., 123(1–2), 81–92, doi:10.1007/s00703-013-0298-9, 2014.

Girard, E., Blanchet, J.-P. and Dubois, Y.: Effects of arctic sulphuric acid aerosols on wintertime low-level atmospheric ice crystals, humidity and temperature at Alert, Nunavut, Atmospheric Res., 73(1–2), 131–148, doi:10.1016/j.atmosres.2004.08.002, 2005.

Girard, E., Dueymes, G., Du, P. and Bertram, A. K.: Assessment of the effects of acid-coated ice nuclei on the Arctic cloud microstructure, atmospheric dehydration, radiation and temperature during winter, Int. J. Climatol., 33(3), 599–614, doi:10.1002/joc.3454, 2013.

GMAO: MERRA-2 tavgM_2d_rad_Nx: 2d,Monthly mean,Time-Averaged,Single-Level,Assimilation,Radiation Diagnostics V5.12.4, version 5.12.4, Greenbelt, MD, USA, Goddard Earth Sciences Data and Information Services Center (GES DISC), Accessed June 20, 2016, , doi:10.5067/OU3HJDS973O0, 2015.

Grenier, P. and Blanchet, J.-P.: Investigation of the sulphate-induced freezing inhibition effect from CloudSat and CALIPSO measurements, J. Geophys. Res. Atmospheres, 115(D22), D22205, doi:10.1029/2010JD013905, 2010.

Hamilton, D. S., Lee, L. A., Pringle, K. J., Reddington, C. L., Spracklen, D. V. and Carslaw, K. S.: Occurrence of pristine aerosol environments on a polluted planet, Proc. Natl. Acad. Sci., 111(52), 18466–18471, doi:10.1073/pnas.1415440111, 2014.

Herman, G. and Goody, R.: Formation and Persistence of Summertime Arctic Stratus Clouds, J. Atmospheric Sci., 33(8), 1537–1553, doi:10.1175/1520-0469(1976)033<1537:FAPOSA>2.0.CO;2, 1976.

Hobbs, P. V. and Rangno, A. L.: Ice Particle Concentrations in Clouds, J. Atmospheric Sci., 42(23), 2523–2549, doi:10.1175/1520-0469(1985)042<2523:IPCIC>2.0.CO;2, 1985.

Huang, Y., Siems, S. T., Manton, M. J., Protat, A. and Delanoë, J.: A study on the low-altitude clouds over the Southern Ocean using the DARDAR-MASK, J. Geophys. Res. Atmospheres, 117(D18), D18204, doi:10.1029/2012JD017800, 2012.

Intrieri, J. M., Shupe, M. D., Uttal, T. and McCarty, B. J.: An annual cycle of Arctic cloud characteristics observed by radar and lidar at SHEBA, J. Geophys. Res. Oceans, 107(C10), SHE 5-1, doi:10.1029/2000JC000423, 2002.

Jackson, R. C., McFarquhar, G. M., Korolev, A. V., Earle, M. E., Liu, P. S. K., Lawson, R. P., Brooks, S., Wolde, M., Laskin, A. and Freer, M.: The dependence of ice microphysics on aerosol concentration in arctic mixed-phase stratus clouds during ISDAC and M-PACE, J. Geophys. Res., 117(D15), D15207, doi:10.1029/2012JD017668, 2012.

Jacob, D. J., Crawford, J. H., Maring, H., Clarke, A. D., Dibb, J. E., Emmons, L. K., Ferrare, R. A., Hostetler, C. A., Russell, P. B., Singh, H. B., Thompson, A. M., Shaw, G. E., McCauley, E., Pederson, J. R. and Fisher, J. A.: The Arctic Research of the Composition of the Troposphere from Aircraft and Satellites (ARCTAS) mission: design, execution, and first results, Atmos Chem Phys, 10(11), 5191–5212, doi:10.5194/acp-10-5191-2010, 2010.

Jaiser, R., Dethloff, K., Handorf, D., Rinke, A. and Cohen, J.: Impact of sea ice cover changes on the Northern Hemisphere atmospheric winter circulation, Tellus A, 64(0), doi:10.3402/tellusa.v64i0.11595, 2012.

Jiao, C. and Flanner, M. G.: Changing black carbon transport to the Arctic from present day to the end of 21st century, J. Geophys. Res. Atmospheres, 121(9), 2015JD023964, doi:10.1002/2015JD023964, 2016.

5   Josey, S. A., Pascal, R. W., Taylor, P. K. and Yelland, M. J.: A new formula for determining the atmospheric longwave flux at the ocean surface at mid-high latitudes, J. Geophys. Res. Oceans, 108(C4), 3108, doi:10.1029/2002JC001418, 2003.

Kacenelenbogen, M., Redemann, J., Vaughan, M. A., Omar, A. H., Russell, P. B., Burton, S., Rogers, R. R., Ferrare, R. A. and Hostetler, C. A.: An evaluation of CALIOP/CALIPSO's aerosol-above-cloud detection and retrieval capability over North America, J. Geophys. Res. Atmospheres, 119(1), 230–244, doi:10.1002/2013JD020178, 2014.

10   Kafle, D. N. and Coulter, R. L.: Micropulse lidar-derived aerosol optical depth climatology at ARM sites worldwide, J. Geophys. Res. Atmospheres, 118(13), 7293–7308, doi:10.1002/jgrd.50536, 2013.

Kay, J. E. and L'Ecuyer, T.: Observational constraints on Arctic Ocean clouds and radiative fluxes during the early 21st century, J. Geophys. Res. Atmospheres, 118(13), 7219–7236, doi:10.1002/jgrd.50489, 2013.

Kiehl, J. T., Schneider, T. L., Rasch, P. J., Barth, M. C. and Wong, J.: Radiative forcing due to sulfate aerosols from 15   simulations with the National Center for Atmospheric Research Community Climate Model, Version 3, J. Geophys. Res. Atmospheres, 105(D1), 1441–1457, doi:10.1029/1999JD900495, 2000.

Kondo, Y., Matsui, H., Moteki, N., Sahu, L., Takegawa, N., Kajino, M., Zhao, Y., Cubison, M. J., Jimenez, J. L., Vay, S., Diskin, G. S., Anderson, B., Wisthaler, A., Mikoviny, T., Fuelberg, H. E., Blake, D. R., Huey, G., Weinheimer, A. J., Knapp, D. J. and Brune, W. H.: Emissions of black carbon, organic, and inorganic aerosols from biomass burning in North America 20   and Asia in 2008, J. Geophys. Res. Atmospheres, 116(D8), D08204, doi:10.1029/2010JD015152, 2011.

Koop, T., Ng, H. P., Molina, L. T. and Molina, M. J.: A New Optical Technique to Study Aerosol Phase Transitions: The Nucleation of Ice from H2SO4 Aerosols, J. Phys. Chem. A, 102(45), 8924–8931, doi:10.1021/jp9828078, 1998.

Ladino Moreno, L. A., Stetzer, O. and Lohmann, U.: Contact freezing: a review of experimental studies, Atmos Chem Phys, 13(19), 9745–9769, doi:10.5194/acp-13-9745-2013, 2013.

25   Lance, S., Shupe, M. D., Feingold, G., Brock, C. A., Cozic, J., Holloway, J. S., Moore, R. H., Nenes, A., Schwarz, J. P., Spackman, J. R., Froyd, K. D., Murphy, D. M., Brioude, J., Cooper, O. R., Stohl, A. and Burkhart, J. F.: Cloud condensation nuclei as a modulator of ice processes in Arctic mixed-phase clouds, Atmospheric Chem. Phys., 11(15), 8003–8015, 2011.

Lebo, Z. J., Johnson, N. C. and Harrington, J. Y.: Radiative influences on ice crystal and droplet growth within mixed-phase stratus clouds, J. Geophys. Res. Atmospheres, 113(D9), D09203, doi:10.1029/2007JD009262, 2008.

30   Lee, S., Kahn, B. H. and Teixeira, J.: Characterization of cloud liquid water content distributions from CloudSat, J. Geophys. Res. Atmospheres, 115(D20), D20203, doi:10.1029/2009JD013272, 2010.

Li, J., Huang, J., Stamnes, K., Wang, T., Lv, Q. and Jin, H.: A global survey of cloud overlap based on CALIPSO and CloudSat measurements, Atmos Chem Phys, 15(1), 519–536, doi:10.5194/acp-15-519-2015, 2015.

Liu, Z., Vaughan, M., Winker, D., Kittaka, C., Getzewich, B., Kuehn, R., Omar, A., Powell, K., Trepte, C. and Hostetler, C.: The CALIPSO Lidar Cloud and Aerosol Discrimination: Version 2 Algorithm and Initial Assessment of Performance, J. Atmospheric Ocean. Technol., 26(7), 1198–1213, doi:10.1175/2009JTECHA1229.1, 2009.

Lohmann, U. and Feichter, J.: Global indirect aerosol effects: a review, Atmos Chem Phys, 5(3), 715–737, doi:10.5194/acp-5-715-2005, 2005.

Lohmann, U., Zhang, J. and Pi, J.: Sensitivity studies of the effect of increased aerosol concentrations and snow crystal shape on the snowfall rate in the Arctic, J. Geophys. Res. Atmospheres, 108(D11), 4341, doi:10.1029/2003JD003377, 2003.

Lubin, D. and Vogelmann, A. M.: A climatologically significant aerosol longwave indirect effect in the Arctic, Nature, 439(7075), 453–456, doi:10.1038/nature04449, 2006.

Luo, Y., Xu, K.-M., Morrison, H., McFarquhar, G. M., Wang, Z. and Zhang, G.: Multi-layer arctic mixed-phase clouds simulated by a cloud-resolving model: Comparison with ARM observations and sensitivity experiments, J. Geophys. Res. Atmospheres, 113(D12), D12208, doi:10.1029/2007JD009563, 2008.

Mauritsen, T., Sedlar, J., Tjernström, M., Leck, C., Martin, M., Shupe, M., Sjogren, S., Sierau, B., Persson, P. O. G., Brooks, I. M. and Swietlicki, E.: An Arctic CCN-limited cloud-aerosol regime, Atmos Chem Phys, 11(1), 165–173, doi:10.5194/acp-11-165-2011, 2011.

McComiskey, A. and Feingold, G.: Quantifying error in the radiative forcing of the first aerosol indirect effect, Geophys. Res. Lett., 35(2), L02810, doi:10.1029/2007GL032667, 2008.

Meier, W., Fetterer, F., Savoie, M., Mallory, S., Duerr, R. and Stroeve, J. C.: NOAA/NSIDC Climate Data Record of Passive Microwave Sea Ice Concentration, Version 2. [G02202], NSIDC: National Snow and Ice Data Center, Boulder, Colorado, USA. [online] Available from: doi: http://dx.doi.org/10.7265/N55M63M1 (Accessed 23 May 2016), 2013.

Mioche, G., Jourdan, O., Ceccaldi, M. and Delanoë, J.: Variability of mixed-phase clouds in the Arctic with a focus on the Svalbard region: a study based on spaceborne active remote sensing, Atmos Chem Phys, 15(5), 2445–2461, doi:10.5194/acp-15-2445-2015, 2015.

Morrison, H., Shupe, M. D., Pinto, J. O. and Curry, J. A.: Possible roles of ice nucleation mode and ice nuclei depletion in the extended lifetime of Arctic mixed-phase clouds, Geophys. Res. Lett., 32(18), L18801, doi:10.1029/2005GL023614, 2005.

Morrison, H., Pinto, J. O., Curry, J. A. and McFarquhar, G. M.: Sensitivity of modeled arctic mixed-phase stratocumulus to cloud condensation and ice nuclei over regionally varying surface conditions, J. Geophys. Res. Atmospheres, 113(D5), D05203, doi:10.1029/2007JD008729, 2008.

Morrison, H., de Boer, G., Feingold, G., Harrington, J., Shupe, M. D. and Sulia, K.: Resilience of persistent Arctic mixed-phase clouds, Nat. Geosci., 5(1), 11–17, doi:10.1038/ngeo1332, 2012.

Ovchinnikov, M., Ackerman, A. S., Avramov, A., Cheng, A., Fan, J., Fridlind, A. M., Ghan, S., Harrington, J., Hoose, C., Korolev, A., McFarquhar, G. M., Morrison, H., Paukert, M., Savre, J., Shipway, B. J., Shupe, M. D., Solomon, A. and Sulia,

K.: Intercomparison of large-eddy simulations of Arctic mixed-phase clouds: Importance of ice size distribution assumptions, J. Adv. Model. Earth Syst., 6(1), 223–248, doi:10.1002/2013MS000282, 2014.

Paris, J.-D., Stohl, A., Nédélec, P., Arshinov, M. Y., Panchenko, M. V., Shmargunov, V. P., Law, K. S., Belan, B. D. and Ciais, P.: Wildfire smoke in the Siberian Arctic in summer: source characterization and plume evolution from airborne measurements, Atmos Chem Phys, 9(23), 9315–9327, doi:10.5194/acp-9-9315-2009, 2009.

Peng, G., Meier, W. N., Scott, D. J. and Savoie, M. H.: A long-term and reproducible passive microwave sea ice concentration data record for climate studies and monitoring, Earth Syst. Sci. Data, 5(2), 311–318, doi:10.5194/essd-5-311-2013, 2013.

Peng, Y., Lohmann, U., Leaitch, R., Banic, C. and Couture, M.: The cloud albedo-cloud droplet effective radius relationship for clean and polluted clouds from RACE and FIRE.ACE, J. Geophys. Res. Atmospheres, 107(D11), AAC 1-1, doi:10.1029/2000JD000281, 2002.

Rangno, A. L. and Hobbs, P. V.: Ice particles in stratiform clouds in the Arctic and possible mechanisms for the production of high ice concentrations, J. Geophys. Res. Atmospheres, 106(D14), 15065–15075, doi:10.1029/2000JD900286, 2001.

Ricchiazzi, P., Yang, S., Gautier, C. and Sowle, D.: SBDART: A Research and Teaching Software Tool for Plane-Parallel Radiative Transfer in the Earth's Atmosphere, Bull. Am. Meteorol. Soc., 79(10), 2101–2114, doi:10.1175/1520-0477(1998)079<2101:SARATS>2.0.CO;2, 1998.

Rogers, R. R., Vaughan, M. A., Hostetler, C. A., Burton, S. P., Ferrare, R. A., Young, S. A., Hair, J. W., Obland, M. D., Harper, D. B., Cook, A. L. and Winker, D. M.: Looking through the haze: evaluating the CALIPSO level 2 aerosol optical depth using airborne high spectral resolution lidar data, Atmos Meas Tech, 7(12), 4317–4340, doi:10.5194/amt-7-4317-2014, 2014.

Sawamura, P., Moore, R. H., Burton, S. P., Chemyakin, E., Müller, D., Kolgotin, A., Ferrare, R. A., Hostetler, C. A., Ziemba, L. D., Beyersdorf, A. J., and Anderson, B. E.: HSRL-2 aerosol optical measurements and microphysical retrievals vs. airborne in situ measurements during DISCOVER-AQ 2013: an intercomparison study, Atmos. Chem. Phys. Discuss., doi:10.5194/acp-2016-1164, in review, 2017.

Sedlar, J. and Devasthale, A.: Clear-sky thermodynamic and radiative anomalies over a sea ice sensitive region of the Arctic, J. Geophys. Res., 117(D19), doi:10.1029/2012JD017754, 2012.

Serreze, M. C., Barrett, A. P., Stroeve, J. C., Kindig, D. N. and Holland, M. M.: The emergence of surface-based Arctic amplification, The Cryosphere, 3(1), 11–19, doi:10.5194/tc-3-11-2009, 2009.

Shaw, G. E.: The Arctic Haze Phenomenon, Bull. Am. Meteorol. Soc., 76(12), 2403–2413, doi:10.1175/1520-0477(1995)076<2403:TAHP>2.0.CO;2, 1995.

Sheridan, P. J., Andrews, E., Ogren, J. A., Tackett, J. L. and Winker, D. M.: Vertical profiles of aerosol optical properties over central Illinois and comparison with surface and satellite measurements, Atmos Chem Phys, 12(23), 11695–11721, doi:10.5194/acp-12-11695-2012, 2012.

Shupe, M. D. and Intrieri, J. M.: Cloud Radiative Forcing of the Arctic Surface: The Influence of Cloud Properties, Surface Albedo, and Solar Zenith Angle, J. Clim., 17(3), 616–628, doi:10.1175/1520-0442(2004)017<0616:CRFOTA>2.0.CO;2, 2004.

Shupe, M. D., Walden, V. P., Eloranta, E., Uttal, T., Campbell, J. R., Starkweather, S. M. and Shiobara, M.: Clouds at Arctic
5   Atmospheric Observatories. Part I: Occurrence and Macrophysical Properties, J. Appl. Meteorol. Climatol., 50(3), 626–644, doi:10.1175/2010JAMC2467.1, 2011.

Shupe, M. D., Persson, P. O. G., Brooks, I. M., Tjernström, M., Sedlar, J., Mauritsen, T., Sjogren, S. and Leck, C.: Cloud and boundary layer interactions over the Arctic sea ice in late summer, Atmos Chem Phys, 13(18), 9379–9399, doi:10.5194/acp-13-9379-2013, 2013.

10   Smith, W. S. and Kao, C.-Y. J.: Numerical Simulations of Observed Arctic Stratus Clouds Using a Second-Order Turbulence Closure Model, J. Appl. Meteorol., 35(1), 47–59, doi:10.1175/1520-0450(1996)035<0047:NSOOAS>2.0.CO;2, 1996.

Sodemann, H., Pommier, M., Arnold, S. R., Monks, S. A., Stebel, K., Burkhart, J. F., Hair, J. W., Diskin, G. S., Clerbaux, C., Coheur, P.-F., Hurtmans, D., Schlager, H., Blechschmidt, A.-M., Kristjánsson, J. E. and Stohl, A.: Episodes of cross-polar transport in the Arctic troposphere during July 2008 as seen from models, satellite, and aircraft observations, Atmos
15   Chem Phys, 11(8), 3631–3651, doi:10.5194/acp-11-3631-2011, 2011.

Stohl, A., Hittenberger, M. and Wotawa, G.: Validation of the lagrangian particle dispersion model FLEXPART against large-scale tracer experiment data, Atmos. Environ., 32(24), 4245–4264, doi:10.1016/S1352-2310(98)00184-8, 1998.

Stohl, A., Eckhardt, S., Forster, C., James, P. and Spichtinger, N.: On the pathways and timescales of intercontinental air pollution transport, J. Geophys. Res. Atmospheres, 107(D23), 4684, doi:10.1029/2001JD001396, 2002.

20   Stohl, A., Forster, C., Eckhardt, S., Spichtinger, N., Huntrieser, H., Heland, J., Schlager, H., Wilhelm, S., Arnold, F. and Cooper, O.: A backward modeling study of intercontinental pollution transport using aircraft measurements, J. Geophys. Res. Atmospheres, 108(D12), 4370, doi:10.1029/2002JD002862, 2003.

Stohl, A., Forster, C., Frank, A., Seibert, P. and Wotawa, G.: Technical note: The Lagrangian particle dispersion model FLEXPART version 6.2, Atmos Chem Phys, 5(9), 2461–2474, doi:10.5194/acp-5-2461-2005, 2005.

25   Stohl, A., Aamaas, B., Amann, M., Baker, L. H., Bellouin, N., Berntsen, T. K., Boucher, O., Cherian, R., Collins, W., Daskalakis, N., Dusinska, M., Eckhardt, S., Fuglestvedt, J. S., Harju, M., Heyes, C., Hodnebrog, Ø., Hao, J., Im, U., Kanakidou, M., Klimont, Z., Kupiainen, K., Law, K. S., Lund, M. T., Maas, R., MacIntosh, C. R., Myhre, G., Myriokefalitakis, S., Olivié, D., Quaas, J., Quennehen, B., Raut, J.-C., Rumbold, S. T., Samset, B. H., Schulz, M., Seland, Ø., Shine, K. P., Skeie, R. B., Wang, S., Yttri, K. E. and Zhu, T.: Evaluating the climate and air quality impacts of short-
30   lived pollutants, Atmos Chem Phys, 15(18), 10529–10566, doi:10.5194/acp-15-10529-2015, 2015.

Strunin, M. A., Postnov, A. A. and Mezrin, M. Y.: Arctic HazeMeteorological potential for contamination of arctic troposphere: Boundary layer structure and turbulent diffusion characteristics, Atmospheric Res., 44(1), 37–51, doi:10.1016/S0169-8095(97)00008-2, 1997.

Taylor, P. C., Kato, S., Xu, K.-M. and Cai, M.: Covariance between Arctic sea ice and clouds within atmospheric state regimes at the satellite footprint level, J. Geophys. Res. Atmospheres, 120(24), 2015JD023520, doi:10.1002/2015JD023520, 2015.

Tietze, K., Riedi, J., Stohl, A. and Garret, T. J.: Space-based evaluation of interactions between aerosols and low-level Arctic clouds during the Spring and Summer of 2008, Atmospheric Chem. Phys., 11, 3359–3373, NaN-11-3359–2011, 2011.

Turner, D. D.: Improved ground-based liquid water path retrievals using a combined infrared and microwave approach, J. Geophys. Res. Atmospheres, 112(D15), D15204, doi:10.1029/2007JD008530, 2007.

Van Tricht, K., Lhermitte, S., Lenaerts, J. T. M., Gorodetskaya, I. V., L'Ecuyer, T. S., Noël, B., van den Broeke, M. R., Turner, D. D. and van Lipzig, N. P. M.: Clouds enhance Greenland ice sheet meltwater runoff, Nat. Commun., 7, 10266, doi:10.1038/ncomms10266, 2016.

Vaughan, M. A., Powell, K. A., Winker, D. M., Hostetler, C. A., Kuehn, R. E., Hunt, W. H., Getzewich, B. J., Young, S. A., Liu, Z. and McGill, M. J.: Fully Automated Detection of Cloud and Aerosol Layers in the CALIPSO Lidar Measurements, J. Atmospheric Ocean. Technol., 26(10), 2034–2050, doi:10.1175/2009JTECHA1228.1, 2009.

Wang, Z.: B-CLDCLASS-LIDAR Interface Control Document, [online] Available from: http://cswww.cira.colostate.edu/icd_pdf.php?avid=36&pvids=12 (Accessed 11 August 2016), 2013.

Wilcox, L. J., Highwood, E. J., Booth, B. B. B. and Carslaw, K. S.: Quantifying sources of inter-model diversity in the cloud albedo effect, Geophys. Res. Lett., 42(5), 2015GL063301, doi:10.1002/2015GL063301, 2015.

Winker, D. M., Vaughan, M. A., Omar, A., Hu, Y., Powell, K. A., Liu, Z., Hunt, W. H. and Young, S. A.: Overview of the CALIPSO Mission and CALIOP Data Processing Algorithms, J. Atmospheric Ocean. Technol., 26(11), 2310–2323, doi:10.1175/2009JTECHA1281.1, 2009.

Winker, D. M., Tackett, J. L., Getzewich, B. J., Liu, Z., Vaughan, M. A. and Rogers, R. R.: The global 3-D distribution of tropospheric aerosols as characterized by CALIOP, Atmos Chem Phys, 13(6), 3345–3361, doi:10.5194/acp-13-3345-2013, 2013.

Zamora, L. M., Kahn, R. A., Cubison, M. J., Diskin, G. S., Jimenez, J. L., Kondo, Y., McFarquhar, G. M., Nenes, A., Thornhill, K. L., Wisthaler, A., Zelenyuk, A. and Ziemba, L. D.: Aircraft-measured indirect cloud effects from biomass burning smoke in the Arctic and subarctic, Atmos Chem Phys, 16(2), 715–738, doi:10.5194/acp-16-715-2016, 2016.

Zhang, D., Wang, Z. and Liu, D.: A global view of midlevel liquid-layer topped stratiform cloud distribution and phase partition from CALIPSO and CloudSat measurements, , 115, D00H13, doi:10.1029/2009JD012143, 2010.

Zhang, Y., Seidel, D. J., Golaz, J.-C., Deser, C. and Tomas, R. A.: Climatological Characteristics of Arctic and Antarctic Surface-Based Inversions, J. Clim., 24(19), 5167–5186, doi:10.1175/2011JCLI4004.1, 2011.

Zhao, C. and Garrett, T. J.: Effects of Arctic haze on surface cloud radiative forcing, Geophys. Res. Lett., 2014GL062015, doi:10.1002/2014GL062015, 2015.

Table 1: Criteria used for cloud and air mass classification.

Data source type	ONLi clean clouds	ONLi all clouds	ONLi aerosol-influenced clouds	MOONLiT clean clouds	MOONLiT all clouds	MOONLiT aerosol-influenced clouds	Clear air
CALIPSO v. 3.01 L2 532 nm aerosol profile data							
Latitude: 60-82°N	x	x	x	x	x	x	x
Nighttime	x	x	x	x	x	x	x
Uppermost cloud layer only				x	x	x	
Cloud top altitude < 8 km asl	x	x	x	x	x	x	
Cloud base altitude > 0.2 km asl	x	x	x				
Cloud base altitude > 1 km asl				x	x	x	
COD < ~3 (no extinction QC flag)	x	x	x	x	x	x	
In-cloud CAD score between 70-100	x	x	x	x	x	x	
CALIPSO "liquid"-phase only	x	x	x	x	x	x	
No cloud phase quality control flags	x	x	x	x	x	x	
No aerosol above cloud	x			x			
Aerosol observed above or below cloud			x			x	
No aerosol between cloud base and surface or next cloud top, whichever comes first	x			x			
Aerosol CAD score between -100 and -70			x			x	
No clouds or aerosol anywhere in profile							x
No absolute profile CAD score values <70				x	x	x	
No ice allowed anywhere in profile				x	x	x	
FLEXPART model output							
BC ≤ 30 ng C m^{-3}	x			x			
BC ≥ 30 ng C m^{-3}			x			x	
CloudSat 2B-CLDCLASS-LIDAR data[a]							
>750 m above ground	x	x	x	x	x	x	
Non precipitating clouds	x	x	x	x	x	x	
Liquid- or mixed-phase only	x	x	x	x	x	x	
Liquid-phase only (for r_{el} measurements)	x	x	x	x	x	x	

[a]As available for clouds with radar reflectivities above the detection limit of -29 dBZ

Table 2: Median (interquartile range) and sample number (n) of Arctic Ocean ONLi cloud properties as classified by the criteria in Table 1, separated by reflectivity above and below detection limit (DL, -29 dBZ) and surface regime. Red (grey) color indicates significant (not significant) differences compared to clean background clouds, as determined at 95% confidence using a permutation test. Blue indicates that significance was lost[d] with altitude binning. An asterisk indicates that the trend observed without binning was still observed in altitude bins > 1.1 km (see Supplementary Tables 1 and 2 for more details).

		Sea ice						Open ocean					
Attribute	Zm	Clean background	n	All clouds	n	Aerosol-impacted[a]	n	Background	n	All	n	Aerosol-impacted[a]	N
Base T (°C)	> DL	-18.9 (-21.8 to -16.0)	5804	-19.3 (-22.3 to -16.1)	19504	-19.3 (-22.9 to -14.8)	800	-13.2 (-18.7 to -7.8)	3681	-11.7 (-17.6 to -6.7)	11339	-13.8 (-18.6 to -8.7)	487
	< DL	-18.6 (-22.2 to -15.0)	897	-18.4 (-21.5 to -15.1)	4594	-18.5 (-22.1 to -15.1)	391	-8.4 (-17.0 to -3.4)	1548	-7.3 (-15.7 to -2.7)	6206	-9.8 (-17.0 to -4.9)	346
	All	-18.8 (-21.8 to -15.8)	6975	-19.1 (-22.2 to -15.8)	25140	-18.9 (-22.7 to -14.9)	1261	-11.7 (-18.2 to -6.0)	5487	-10.0 (-16.9 to -4.8)	18499	-12.3 (-17.7 to -6.6)	879
Top T (°C)	> DL	-23.6 (-27.4 to -20.1)	5804	-23.2 (-27.1 to -18.7)*	19504	-23.0 (-27.1 to -18.7)*	800	-20.2 (-25.8 to -13.0)	3681	-17.5 (-24.0 to -11.7)	11339	-20.0 (-24.8 to -14.4)	487
	< DL	-21.2 (-25.6 to -17.8)	897	-20.9 (-24.2 to -17.7)*	4594	-21.4 (-24.5 to -18.2)	391	-11.8 (-21.5 to -6.6)	1548	-10.7 (-19.6 to -6.0)	6206	-13.0 (-20.9 to -8.2)	346
	All	-23.3 (-27.2 to -19.6)	6975	-22.7 (-26.6 to -19.0)*	25140	-22.3 (-26.4 to -18.4)*	1261	-18.0 (-24.2 to -10.0)	5487	-15.1 (-22.4 to -8.6)	18499	-17.8 (-23.1 to -10.2)	879
Altitude, base (km)	> DL	1.72 (1.30-2.38)	5804	1.60 (1.12-2.20)*	19504	1.78 (1.24-2.44)	800	2.74 (1.36-3.70)	3681	2.26 (1.18-3.40)	11339	2.50 (1.60-3.40)	487
	< DL	2.02 (1.42-2.86)	897	1.78 (1.12-2.50)*	4594	2.08 (1.54-2.68)	391	2.32 (1.36-3.58)	1548	2.02 (1.30-3.16)*	6206	2.26 (1.54-2.98)	346
	All	1.78 (1.30-2.38)	6975	1.60 (1.12-2.26)	25140	1.90 (1.30-2.56)	1261	2.62 (1.36-3.64)	5487	2.14 (1.18-3.82)	18499	2.38 (1.54-3.22)	879
Thickness (km)	> DL	0.96 (0.66-1.32)	5804	0.78 (0.60-1.20)	19504	0.72 (0.60-0.96)*	800	0.84 (0.60-1.32)	3681	0.78 (0.60-1.32)*	11339	0.72 (0.60-1.11)*	487
	< DL	0.60 (0.48-0.72)	897	0.60 (0.48-0.72)*	4594	0.54 (0.48-0.66)*	391	0.06 (0.48-0.78)	1548	0.06 (0.48-0.72)	6206	0.06 (0.48-0.72)	346
	All	0.84 (0.60-1.26)	6975	0.72 (0.60-1.08)	25140	0.66 (0.54-0.84)	1261	0.72 (0.54-1.08)	5487	0.66 (0.54-1.08)*	18499	0.66 (0.54-0.84)*	879
COD	> DL	1.14 (0.65-1.85)	4160	1.00 (0.60-1.63)	16234	0.84 (0.53-1.40)	772	0.82 (0.39-1.54)	3286	0.88 (0.44-1.51)	10474	0.81 (0.48-1.26)	463
	< DL	0.55 (0.30-1.11)	816	0.63 (0.36-1.07)	4372	0.53 (0.34-0.89)	387	0.49 (0.23-1.09)	1427	0.62 (0.29-1.21)	5885	0.61 (0.33-1.12)	339
	All	1.03 (0.55-1.72)	5195	0.90 (0.52-1.51)	21533	0.73 (0.42-1.15)	1227	0.69 (0.29-1.41)	4952	0.77 (0.35-1.40)	17265	0.72 (0.37-1.18)	847
Multi-layer clouds	> DL	75%	5804	79%	19504	91%	800	90%	3681	89%	11339	94%	487
	< DL	85%	897	85%	4594	95%*	391	92%	1548	91%	6206	96%	346
	All	77%	6975	80%	25140	92%	1261	90%	5487	90%	18499	95%	879
BC at base (ng m⁻³)	> DL	15 (10-21)	5804	26 (14-54)	19504	60 (42-94)	800	13 (8-20)	3681	18 (10-36)	11339	61 (42-95)	487
	< DL	15 (11-21)	897	24 (13-48)	4594	54 (38-94)	391	13 (8-19)	1548	17 (9-36)	6206	61 (40-105)	346
	All	15 (10-21)	6975	26 (14-52)	25140	59 (41-94)	1261	13 (8-19)	5487	18 (10-37)	18499	61 (41-102)	879
% < CloudSat DL[b]	All	15%	6194	21%	21841	36%	1163	33%	4950	40%*	16612	44%*	850
% Mixed-phase[b]	> DL	95%	4795	93%*	15698	91%*	681	79%	2992	75%*	9153	80%	417
% precipitating[b,c]	> DL	18%	5916	13%	18125	11%*	737	8%	3283	8%	10077	11%	454
r_{el} (μm)[b]	> DL	10.3 (9.4-11.2)	4917	10.0 (9.2-11.0)*	15414	9.8 (9.1-10.7)*	650	10.0 (9.2-11.2)	2729	10.0 (9.1-11.2)	8420	9.7 (9.0-10.9)*	368
Reflectivity (dBZ)[b]	> DL	-20.4 (-24.3 to -16.7)	5294	-21.5 (-25.3 to -17.6)	17287	-22.8 (-26.4 to -18.8)	745	-21.7 (-25.7 to -16.9)	3680	-22.2 (-26.2 to -17.0)*	11329	-23.5 (-26.7 to -19.1)*	487

[a]Aerosol-impacted, as determined in the third column of Table 1.

[b]For clouds with bases >750 m asl

[c]Precipitating clouds were included in this metric only; for all other attribute classifications, clouds were required to have no observed precipitation in order to be comparable with r_{el} estimates that were most reliable in non-precipitating clouds.

[d]Significance was presumed to be lost across altitude bins when there were multiple cases of non-significance among altitude bins or different trends in significance between altitude bins.
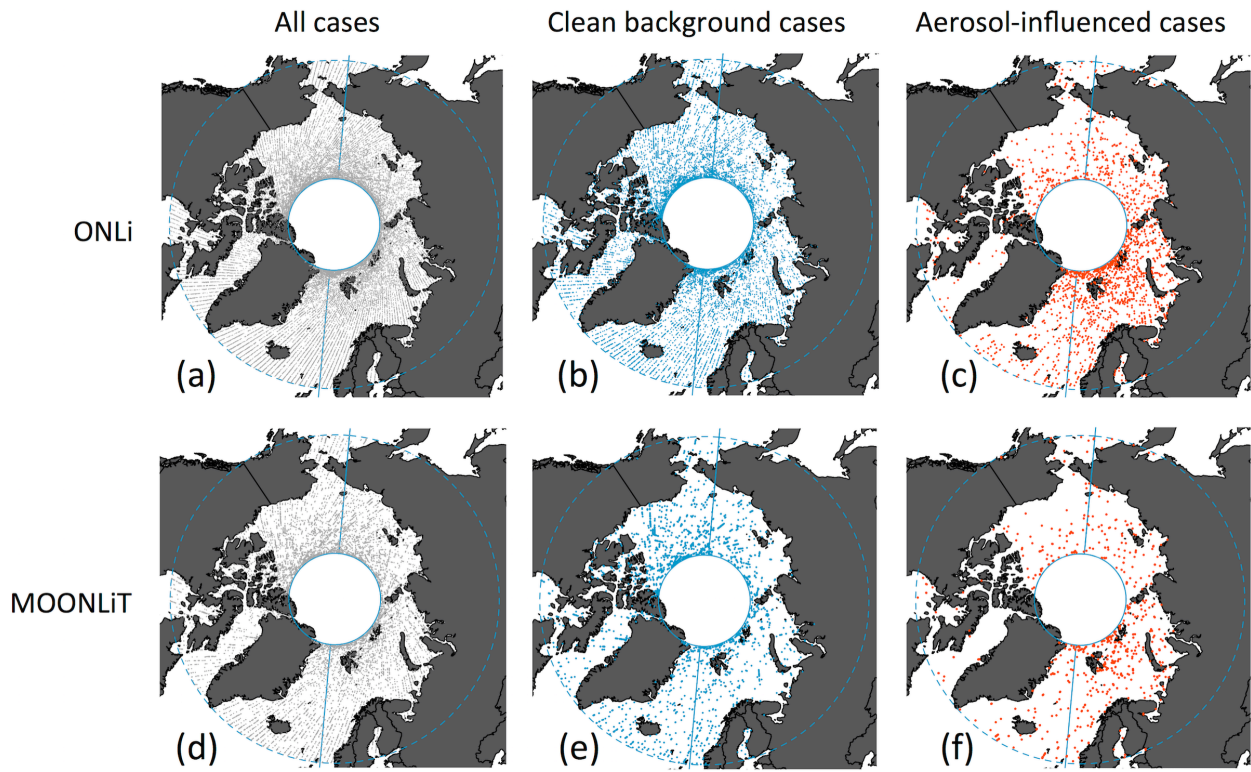
Figure 1: The geographical distribution of ONLi and MOONLiT cloud profiles, where (a,d) grey indicates all cases, (b,e) blue indicates clean background cases, and (c,f) red indicates aerosol-influenced cases.
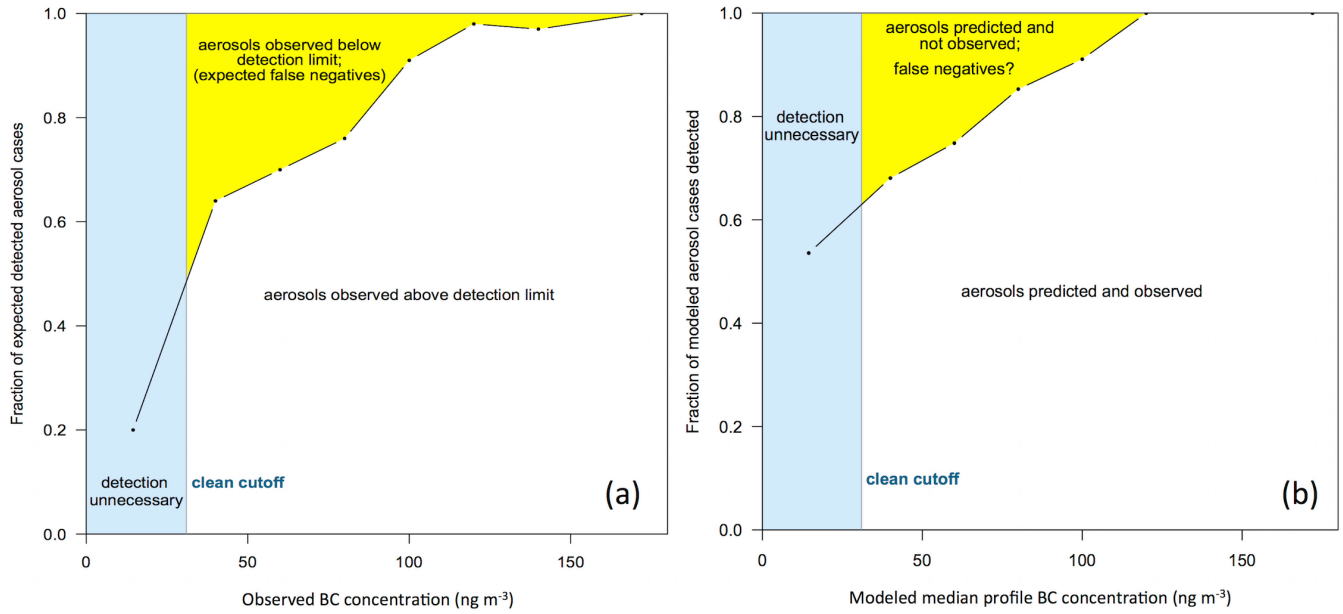
Figure 2: Based on CALIPSO Arctic profiles under non-cloudy conditions, we compare a) the expected fraction and b) possible maximum fraction of false negatives (aerosol present but not detected) for different aerosol concentrations and/or combustion tracers. Tracers include black carbon (BC, ng C m^{-3}) and the concentration of aerosols with diameters > 0.12 μm (CN$_{PCASP}$, cm^{-3}). The expected fraction of false negatives in panel a) was determined by comparing binned out-of-cloud 2008 ARCTAS-A and -B BC concentrations and ISDAC CN$_{PCASP}$ concentrations with the fraction of the total number of samples between 1-5 km that had converted backscatter values (Mm^{-1} sr^{-1}) above the CALIPSO clear-sky nighttime backscatter detection limit from Winker et al. (2009) (see text for more details). Possible maximum false negative values in panel b) were determined by comparing the FLEXPART model's median BC concentrations between 0-10 km with the fraction of the total CALIPSO profiles under non-cloudy conditions during January, 2008 where aerosols were not detected. The clean cutoff below which air is taken as "clean" is assumed to be 31 ng C m^{-3} and 127 cm^{-3} for BC and CN$_{PCASP}$, respectively.
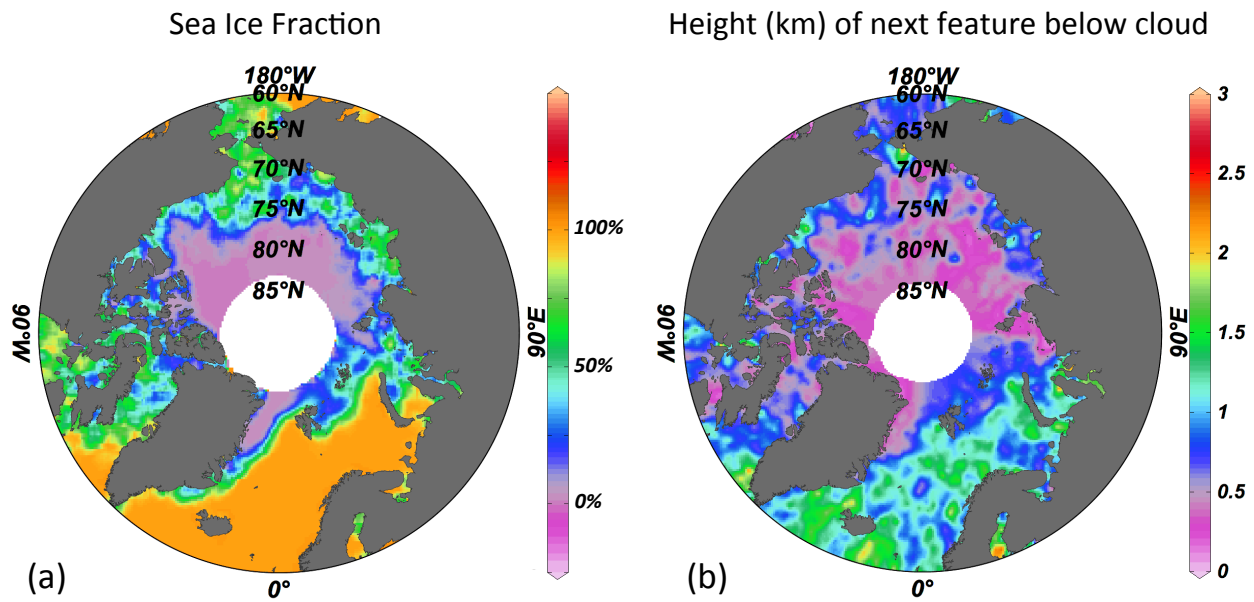
Figure 3: The weighted-average gridded maps of features below individual cloud points from Fig. 1b for a) sea ice fraction, and b) height of the next lowest feature associated with individual cloud profiles, where a value of 0 indicates that the ocean surface or sea ice was the next lowest feature. Over open ocean, ONLi clouds were much more likely to overlay another cloud than over sea ice.
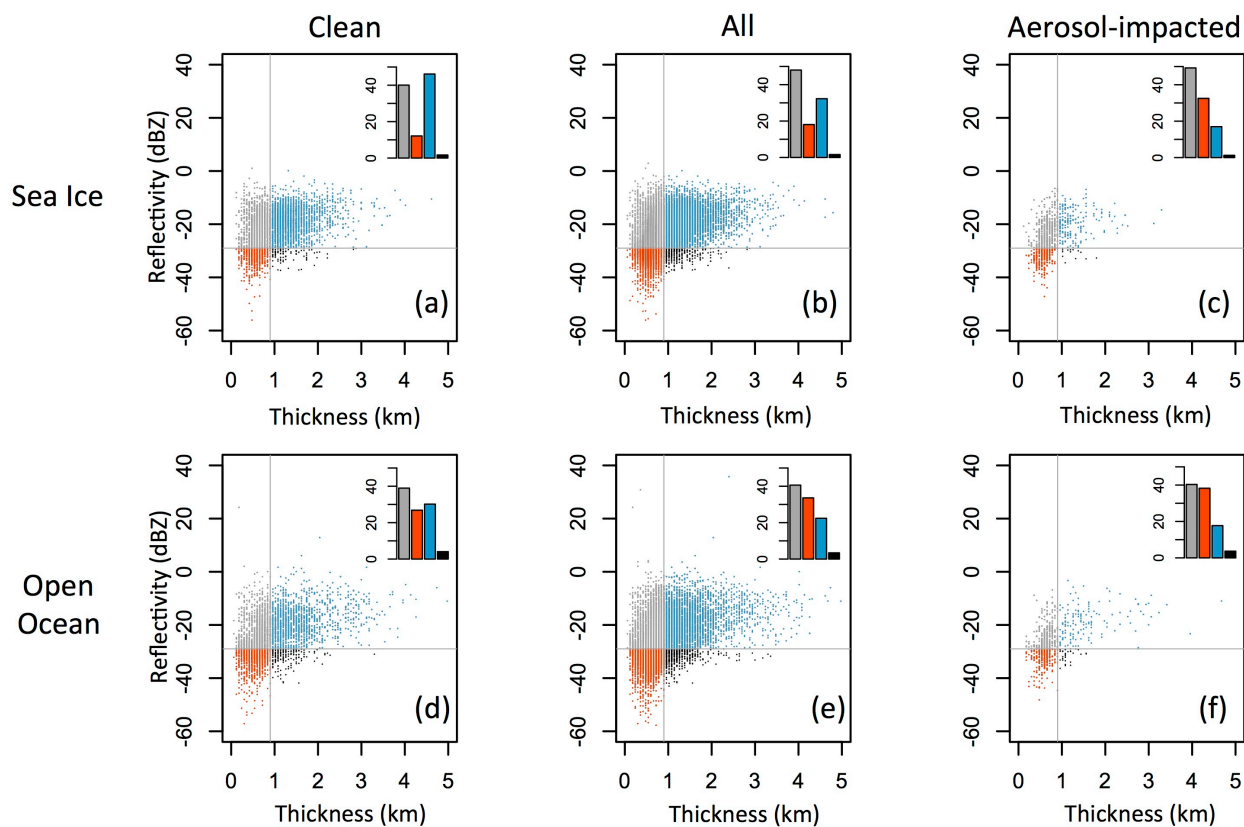
Figure 4: A comparison of CALIPSO ONLi cloud thickness (km) with CloudSat reflectivity (dBZ), as separated by sea ice and open ocean regimes, and by clouds found in conditions labeled as clean background, all conditions, and aerosol-impacted conditions. To better show changes in the two parameters, plots have been divided into four quadrants (above (grey and blue) and below (orange and black) the CloudSat reflectivity detection limit of -29 dBZ), and above (blue and black) and below (orange and grey) a thickness of 0.9 km. In the upper right of each plot is shown the percent of cases within each quadrant, following the quadrant color scheme. Points represent clouds > 750 m asl.