



Observations and comparisons of properties in Arctic stratocumulus during ACCACIA

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Observations and comparisons of cloud microphysical properties in spring and summertime Arctic stratocumulus during the ACCACIA campaign

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Abstract

Measurements from four case studies in spring and summer-time Arctic stratocumulus clouds during the Aerosol–Cloud Coupling And Climate Interactions in the Arctic (AC-CACIA) campaign are presented. We compare microphysics observations between cases and with previous measurements made in the Arctic and Antarctic. During AC-CACIA, stratocumulus clouds were observed to consist of liquid at cloud tops, often at distinct temperature inversions. The cloud top regions precipitated low concentrations of ice into the cloud below. During the spring cases median ice number concentrations ($\sim 0.5 \text{ L}^{-1}$) were found to be lower by about a factor of 5 than observations from the summer campaign ($\sim 3 \text{ L}^{-1}$). Cloud layers in the summer spanned a warmer temperature regime than in the spring and enhancement of ice concentrations in these cases was found to be due to secondary ice production through the Hallett–Mossop (H–M) process. Aerosol concentrations during spring ranged from $\sim 300\text{--}400 \text{ cm}^{-3}$ in one case to lower values of $\sim 50\text{--}100 \text{ cm}^{-3}$ in the other. The concentration of aerosol with sizes, $D_p > 0.5 \mu\text{m}$, was used in a primary ice nucleus (IN) prediction scheme, DeMott et al. (2010). Predicted IN values varied depending on aerosol measurement periods, but were generally greater than maximum observed median values of ice crystal concentrations in the spring cases, and less than the observed ice concentrations in the summer due to the influence of secondary ice production. Comparison with recent cloud observations in the Antarctic summer (Grosvenor et al., 2012), reveals lower ice concentrations in Antarctic clouds in comparable seasons. An enhancement of ice crystal number concentrations (when compared with predicted IN numbers) was also found in Antarctic stratocumulus clouds spanning the Hallett–Mossop (H–M) temperature zone, but concentrations were about an order of magnitude lower than those observed in the Arctic summer cases, but were similar to the peak values observed in the colder Arctic spring cases, where the H–M mechanism did not operate.

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1 Introduction

The Arctic is a region that has experienced rapid climate perturbation in recent decades, with warming rates there being almost twice the global average over the past 100 years (ACIA, 2005; IPCC 2007). The most striking consequence of this warming has been the decline in the extent and area of sea ice, especially in the warm season. The lowest sea ice extent and area on record were both observed on 13 September 2012 (Parkinson and Comiso, 2013) and despite some uncertainty, ice-free Arctic summers could become a reality by 2030 (Overland and Wang, 2013). The underlying warming is very likely caused by increasing anthropogenic greenhouse gases and arctic amplification, which is a well-established feature of global climate models (see for example IPCC 5th Assessment Report 2014). However, the details of Arctic climate are complex with interactions between the atmospheric boundary layer, cloud, overlying sea-ice and water leading to a number of feedback mechanisms. These interactions are not well understood due to variability in the spatial and temporal extent of feedback mechanisms, and the fact that those that are included in Global Climate Models (GCMs) may not be accurately parameterised (Callaghan et al., 2011). Clouds play an important role in a number of proposed feedback processes that may be active in the Arctic (Curry et al., 1996; Walsh et al., 2002), Arctic clouds are the dominant factor controlling the surface energy budget, producing a mostly positive forcing throughout the year, apart from a brief cooling period during the middle of summer (Intrieri et al., 2002a). These clouds affect both the long-wave (year-round) and short-wave (summer-only) radiation budgets, and influence turbulent surface exchange. Cloud microphysical influence on cloud radiative properties depends on the amount of condensed water and the size, phase and habit of the cloud particles (Curry et al., 1996). These factors are controlled in part by the Cloud Condensation Nuclei (CCN) and Ice Nuclei (IN) concentrations and properties. Very low aerosol concentrations in the Arctic can result in clouds with properties differing greatly from those at mid-latitudes (Tjernström et al., 2008). A paucity of observations in the Arctic means that neither the aerosol processes,

nor cloud properties are well understood or accurately represented within models, with the result that aerosol and cloud-forcing of Arctic climate is poorly constrained.

In the Arctic lower troposphere low cloud dominates the variability in Arctic cloud cover (Curry et al., 1996), with temperature and humidity profiles showing a high frequency of one or more temperature inversions (Kahl, 1990) below which stratocumulus clouds form. During the Arctic summer, therefore, these low clouds often consist of multiple layers, with a number of theories describing their vertical separation (Her-
man and Goody, 1976; Tsay and Jayaweera, 1984; McInnes and Curry, 1995a). Such cloud layers have been observed during different seasons but the relationship between temperature and the formation of ice in them is not well understood. Jayaweera and Ohtake (1973) observed very little ice above -20°C , but Curry et al. (1997) observed ice to be present in clouds at temperatures between $-8^{\circ}\text{C} < T < -14^{\circ}\text{C}$ during the Beaufort Arctic Storms Experiment (BASE). It is possible that the large variation in temperature at which glaciation is observed is caused by changes in the concentration and composition of aerosol (Curry, 1995). Recent work, such as in the Arctic Cloud Experiment (ACE) (Uttal et al., 2002) has improved our knowledge of Arctic mixed-phase clouds, which dominate in the coldest 9 months of the Arctic year. ACE reported that clouds were mainly comprised of liquid tops, tended to be very long lived and continually precipitated ice. The longevity of these clouds might be considered unusual as the formation of ice leads to loss of water through the Wegener–Bergeron–Findeison process. More recently the Mixed-Phase Arctic Cloud Experiment (M-PACE, 2004) investigated the Arctic autumn transition season. M-PACE was conducted on the North slope of Alaska, in the area to the east of Barrow (Verlinde et al., 2007). Again predominantly mixed-phase clouds were observed with liquid layers present at temperatures as low as -30°C . Remote sensing studies also showed that ice was generally present in low concentrations, mostly associated with precipitation shafts, however, there was also evidence of light snow below thicker layer clouds. IN concentrations were also measured and observed to be low, consistent with liquid water being observed down to very low temperatures. Here we present detailed airborne microphysical and aerosol

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measurements made in stratocumulus cloud regions in the European Arctic during the recent Aerosol–Cloud Coupling And Climate Interactions in the Arctic (ACCACIA) campaigns. We present data from two aircraft during early spring, in March and April 2013, and from a single aircraft during the following Arctic summer, in July 2013.

The objectives of this paper are:

1. to report the microphysics and cloud particle properties of Arctic clouds, and the properties, number and size distributions of aerosols in the vicinity of these,
2. to identify the origin of the ice phase in these clouds and to compare ice crystal number concentrations with the parameterisation of primary Ice Nucleus (IN) concentrations of DeMott et al. (2010),
3. to compare the cloud physics in spring and summer conditions and to identify any contributions of secondary ice particle production,
4. to compare and contrast the mixed phase cloud microphysics of Arctic clouds with clouds observed in the Antarctic.

2 Methodology

The ACCACIA campaigns took place during March–April 2013 and July 2013. They were conducted in the region between Greenland and Norway mainly in the vicinity of Svalbard (and further afield to the south and west of the archipelago). The overarching theme of the project was to reduce the large uncertainty in the effects of aerosols and clouds on the Arctic surface energy balance and climate. Key to the work presented here is an understanding the microphysical properties of Arctic clouds and their dependence on aerosol properties. To this end the FAAM BAe-146 aircraft performed a number flights incorporating profiled ascents, descents and constant altitude runs below, within and above cloud during the spring period. This provided high-resolution

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measurements of the vertical structure of the cloud microphysics and the aerosol properties in and out of cloud regions. The British Antarctic Survey (BAS) Twin Otter aircraft flew during both campaign periods, providing a subset of the BAe-146 measurements. It was the only aircraft present during the summer period. A total of 9 science flights were conducted during the spring period with complementary flights from the BAS twin otter and 6 flights by the BAS twin otter alone during the summer period.

Two case studies are selected from both the early spring and summer campaigns. The spring campaign case studies were selected for having quite different aerosol loadings within the boundary layer. One was in Arctic air with low total aerosol numbers, while the second had higher aerosol loadings in the boundary layer. Summer flight cases were selected for being the cases with higher cloud layer temperatures in comparison to the spring cases. Summer case cloud layer temperatures were significantly higher than in the spring cases, and were observed to be in the temperature zone, -3°C to -9°C , where a powerful mechanism of secondary ice particle production through rime-splintering, the Hallett–Mossop mechanism, (H–M) (Hallett and Mossop, 1974), is known to operate under particular conditions, and so could greatly enhance ice crystal number concentrations. Temperature profiles in the spring cases revealed stratocumulus cloud temperatures generally between $-10^{\circ}\text{C} < T < -20^{\circ}\text{C}$, outside of the H–M zone.

2.1 Instrumentation

Instrumentation onboard the Facility for Airborne Atmospheric Measurements (FAAM) British Aerospace-146 (BAe-146, or 146) aircraft used for making measurements of the cloud and aerosol microphysics reported in this paper included: the Cloud Imaging Probe models 15 and 100 (CIP-15 and CIP-100, Droplet Measurement Technologies (DMT), Boulder, USA) (Baumgardner et al., 2001), the Cloud Droplet Probe (CDP-100 Version 2, DMT) (Lance et al., 2010) and the Two Dimensional-Stereoscopic Probe (2D-S, Stratton Park Engineering Company Inc. Boulder, USA) (Lawson et al., 2006). The CIP-15 and CIP-100 are optical array shadow probes consisting of 64 element

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photodiode arrays providing image resolutions of 15 and 100 μm respectively. The 2D-S is a higher resolution optical array shadow probe which consists of a 128 element photodiode array with image resolution of 10 μm . The CDP measures the liquid droplet size distribution over the particle size range $3 < d_p < 50 \mu\text{m}$. The intensity of forward scattered laser light in the range $4\text{--}12^\circ$ is collected and particle diameter calculated from this information using Mie scattering solutions (Lance et al., 2010).

A Cloud Aerosol Spectrometer (CAS, DMT) and a Passive Cavity Aerosol Spectrometer Probe (PCASP-100X, DMT) were both used to measure aerosol size distributions onboard the 146. The CAS measures particles in the size range $0.51 < d_p < 50 \mu\text{m}$ using forward scattered light from single particles in the $4\text{--}13^\circ$ range and backscattered light in the $5\text{--}13^\circ$ range. Particle size can be determined from both the forward and back-scattered light intensity using Mie scattering solutions (Baumgardner et al., 2001). The PCASP is another Optical Particle Counter (OPC) and measures aerosol particles in the size range $0.1 < d_p < 3 \mu\text{m}$. In this instrument, particles are sized through measurement of the intensity of laser light scattered within the $35\text{--}120^\circ$ range (Rosenberg et al., 2012). All the above instruments were mounted externally on the FAAM aircraft. Non refractory aerosol composition measurements were provided using an Aerodyne Compact-Time of Flight Aerosol Mass Spectrometer (C-ToF-AMS) whilst aerosol black carbon measurements were provided by a single particle soot photometer (SP-2, DMT). Results from these will be reported elsewhere. Examples of additional core data measurements that were also used in this paper include temperature (Rosemount/Goodrich type 102 temperature sensors) and altitude measured by the GPS-aided Inertial Navigation system (GIN).

Instrumentation on board the Twin Otter Meteorological Airborne Science Instrumentation (MASIN) aircraft, relevant to measurements reported in this paper included: a CDP-100 for drop size distributions; a 2D-S (summer only), both similar to those on the FAAM aircraft; a CIP-25 (as on FAAM except consisting of a 64 element photodiode array providing an image resolution of 25 μm) and core data including temperature measured by Goodrich Rosemount Probes (models; 102E4AL and 102AU1AG for non-

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to 2 μm , although the extra contribution to the aerosol concentrations used in the calculations is likely to be small. Grosvenor et al. (2012) demonstrated that the scheme is not particularly sensitive to small changes in total aerosol concentrations $> 0.5 \mu\text{m}$ in clean Antarctic regions. Measurements from the higher resolution PCASP were selected from the size range 0.5 to 1.6 μm , in keeping with the D10 scheme. The D10 predicted IN concentrations were then compared directly as a function of temperature with the observed ice crystal concentrations. The minimum observed median temperature was input to D10 and predicted IN numbers compared with the maximum observed median ice crystal number concentrations (Fig. 11) for the clouds during each of the 4 cases. The results are shown in Table 2.

The results of this comparison from all 4 cases can be compared with previous observations of Arctic clouds and with recent aircraft measurements of clouds over the Antarctic Peninsula in the summer (Grosvenor et al., 2012).

3 Spring case 1 – Friday 22 March 2013 (FAAM flight B761)

On this day the FAAM aircraft first flew from Kiruna, Sweden (67.85° N, 20.21° E) to Svalbard, Norway landing at Longyearbyen, (78.22° N, 15.65° E) to refuel. After take-off at $\sim 11:45$ UTC a ~ 2 h science flight was undertaken to the south east of Svalbard (Fig. 1a) before returning to Kiruna. The objective was to investigate stratocumulus cloud in this area, near to the ice edge, and from over ice to open ocean (moving from N to S in the target area). The flight focused on a series of profiled descents and ascents to enable measurements to be made of the cloud layer from below cloud base to above cloud top and into the inversion layer above. During the flight there were 3 significant penetrations through the inversion at cloud top and in each case there was a marked temperature increase of $\sim 5^\circ\text{C}$. Microphysical time series data for this case are presented, with the relevant runs highlighted in Fig. 2. A description of one cloud profile is given here, with further profiles described in Appendix A. For this case, boundary layer aerosol number concentrations (from the PCASP) were found to be

relatively low at $\sim 50\text{--}100\text{ cm}^{-3}$. Widespread low cloud was observed south and east of Svalbard (Fig. 1) with winds from the north advecting from over the sea-ice towards open sea. Earlier dropsonde measurements (on the transit into Longyearbyen prior to refuelling) showed surface winds of $\sim 3\text{ m s}^{-1}$ increasing to 15 m s^{-1} at 500 mbar.

3.1 Profiled descent A1

During profile A1 the aircraft (now travelling north) descended from the inversion layer. Cloud top was encountered at 1650 m ($T = -18.6^\circ\text{C}$). The highest values of N_{ice} were observed in the cloud top region, at $\sim 4\text{ L}^{-1}$ with peaks up to 7 L^{-1} where IWCs were 0.15 gm^{-3} . Particles here consisted of small irregular ice particles (mean size $\sim 360\text{ }\mu\text{m}$) that showed evidence of riming, together with small droplets. CDP LWC increased to 0.3 gm^{-3} with $N_{\text{drop}} \sim 55\text{ cm}^{-3}$ (mean diameter $\sim 17\text{ }\mu\text{m}$). At an altitude of around 1400 m a.s.l. ($\sim 250\text{ m}$ below cloud top) N_{ice} decreased to $\sim 1\text{ L}^{-1}$, and mean ice particle size increased to $\sim 395\text{ }\mu\text{m}$. N_{drop} increased to $\sim 70\text{ cm}^{-3}$, while mean size decreased slightly ($\sim 16\text{ }\mu\text{m}$). LWCs generally decreased somewhat to $\sim 0.2\text{ gm}^{-3}$. As the aircraft descended to an altitude of $\sim 1150\text{ m}$, N_{ice} increased by approximately a factor of 2 (to $\sim 2\text{ L}^{-1}$). At around 13:15 UTC a number of rapid transitions from liquid to predominantly glaciated conditions were observed in the mid cloud region at 730 m and $T = -12^\circ\text{C}$. The initial phase change occurred as LWC decreased from 0.2 to 0.01 gm^{-3} while IWCs increased to a peak value of 0.2 gm^{-3} and peak N_{drop} fell close to 1 cm^{-3} . 2D-S imagery (Fig. 3c) highlights these changes taking place as small droplets are quickly replaced by small irregular ice crystals and eventually larger snow particles (mean diameter $\sim 610\text{ }\mu\text{m}$) that consisted of heavily rimed ice crystals and aggregates, some of which can be identified as exhibiting a dendritic habit. Observations of dendritic ice are consistent with the ice crystal growth habit expected at this temperature level (-12°C). Three further swift phase transitions were observed as the aircraft approached cloud base. LWC in the liquid dominated regions was between ~ 0.15 and

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0.25 g m^{-3} while N_{drop} peaked at $\sim 130 \text{ cm}^{-3}$. During the ice phase sections of the transition cycle, mean particle sizes were $\sim 615 \mu\text{m}$ and N_{ice} peaked at up to 5 L^{-1} . The contribution of these glaciated cloud regions to the IWC was considerable, with values up to 0.1 g m^{-3} recorded. These transitions ended as the aircraft descended below cloud base ($T = -12 \text{ }^\circ\text{C}$) at 700 m.a.s.l., and precipitating snow was observed (mean size $\sim 710 \mu\text{m}$).

4 Spring case 2 – Wednesday 3 April 2013 (FAAM flight B768)

The FAAM aircraft departed Longyearbyen at around 11:00 UTC and conducted measurements to the NW of Svalbard to investigate low-level clouds over sea ice as well as the transition to deeper more convective type cloud as the aircraft moved away from the ice edge and over warmer water (moving from NW to SE in the target area – Fig. 1b). A low pressure (1004 mbar) region was centred south of Svalbard with an associated band of cloud and precipitation. To the NW of Svalbard, within the measurement area, surface winds were E–NE and $< 10 \text{ ms}^{-1}$. Measurements revealed an air mass containing significantly more aerosol than in Spring case 1, with PCASP concentrations typically $\sim 300\text{--}400 \text{ cm}^{-3}$ in the boundary layer. During the flight the aircraft made two distinct saw tooth profiles through the cloud layer and into the inversion above cloud top where temperatures in each instance increased by $\sim 2 \text{ }^\circ\text{C}$. Figure 4 shows time series of the microphysical measurements made during this science flight. Further profile descriptions can be found in Appendix B.

4.1 Profiled descent B1

Flying NW, the aircraft performed a profiled descent from the inversion layer ($T = -16.5 \text{ }^\circ\text{C}$) into cloud top, $\sim 1550 \text{ m.a.s.l.}$, where the measured temperature was $-17 \text{ }^\circ\text{C}$. LWCs rose to a peak value of $\sim 0.9 \text{ g m}^{-3}$ and N_{drop} (mean diameter $\sim 15 \mu\text{m}$) peaked at $\sim 320 \text{ cm}^{-3}$. The highest values of N_{ice} never exceeded 0.5 L^{-1} in this cloud top re-

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measure aerosol concentrations and composition in the vicinity of cloud, together with the microphysical properties of the clouds by undertaking a combination of profiles and straight and level runs through stratocumulus cloud layers to capture the microphysical structure. Time series of data collected during this flight are presented in Fig. 6. Profile

C2 is described below, with details of the measurements made during C1 found in Appendix C.

5.1 Profile C2

The aircraft performed a sawtooth profile, descending from cloud top at ~ 3300 m down to a minimum altitude of ~ 2300 m followed by a profiled ascent to complete the sawtooth. During the descent into cloud top ($T = -9^\circ\text{C}$) LWCs rose sharply to peak values of 0.3 gm^{-3} and N_{drop} (mean diameter $19\ \mu\text{m}$) increased to 155 cm^{-3} . N_{ice} in the cloud top regions peaked at 1 L^{-1} . With decreasing altitude, LWC declined gradually to values close to 0.01 gm^{-3} . As the temperature increased to above -8°C , ice crystal number concentrations (mean diameter $210\ \mu\text{m}$) increased to 5 L^{-1} , with peaks to $\sim 12\text{ L}^{-1}$. 2D-S imagery revealed the presence of small columnar ice crystals together with small liquid droplets (CDP mean diameter $8.5\ \mu\text{m}$) and some irregular ice particles. At 2880 m ($T = -6.5^\circ\text{C}$) the cloud dissipated until the next cloud layer was encountered 200 m below ($T = -5^\circ\text{C}$). In this region CDP LWC and N_{drop} were more variable than in the cloud layer above. Generally LWCs were $< 0.1\text{ gm}^{-3}$ with peaks in N_{drop} to $\sim 155\text{ cm}^{-3}$ and transitions between liquid cloud and predominantly glaciated cloud were observed. N_{ice} peaked at 25 L^{-1} and IWCs peaked at 0.15 gm^{-3} . 2D-S imagery showed many columnar ice crystals, typical of the growth regime at this temperature ($\sim -5^\circ\text{C}$) and consistent with the enhancement of N_{ice} through the H–M process. The aircraft reached its minimum altitude ($T = -3^\circ\text{C}$) before beginning a profiled ascent to complete the sawtooth. The cloud microphysics of the lower cloud layer were the same as encountered in the descent leg, but with LWCs at times higher (peaks up to 0.2 gm^{-3}). Transitions between liquid and glaciated phases were observed again, with a notable period of high

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0.3 g m⁻³ and N_{drop} rose to a maximum of $\sim 120 \text{ cm}^{-3}$ (mean diameter 14 μm). The aircraft penetrated cloud top at 3700 m ($T = -4.5^\circ\text{C}$).

After climbing above cloud top, the aircraft performed a profiled descent back into the cloud layer to begin another SLR at 3400 m ($T = -4.5^\circ\text{C}$). At cloud top LWCs were $\sim 0.2 \text{ g m}^{-3}$ N_{drop} peaked at 115 cm^{-3} . N_{ice} values were greater than in the previous cloud top region. There were two peaks of up to 15 L^{-1} with particle mean particle diameters of $\sim 370 \mu\text{m}$. Images show columnar particles, some of which had aggregated, were present together with small liquid droplets (CDP mean diameter 11.5 μm). The second peak contained columnar ice crystals of a similar size (mean diameter 400 μm). The largest spike in ice concentrations occurred in close proximity to the first peak, with values as high as 20 L^{-1} observed, while IWCs peaked at 0.15 g m^{-3} . Images showed irregular and columnar ice particles (mean diameter 260 μm) present together with small liquid droplets (CDP mean diameter 12 μm) (Fig. 9b). After these highs in ice number, concentrations declined to $\sim 2.5 \text{ L}^{-1}$ before the aircraft made a short profiled ascent and concentrations rose again to peak values of 10 L^{-1} . At 3550 m cloud dissipated and the aircraft descended through a predominantly clear region before reaching another significant cloud layer at 3450 m ($T = -4^\circ\text{C}$). CDP N_{drop} and LWCs were variable in this region with 10 s mean values rising to 145 cm^{-3} and 0.1 g m^{-3} respectively. The droplets were small (mean diameter 8 μm) and ice was almost completely absent during this part of the profile. After an SLR at 3400 m, the aircraft descended as the cloud layer dissipated but encountered another, more significant layer around 3250 m ($T = -2.5^\circ\text{C}$). LWCs increased to peak values of 0.4 g m^{-3} and droplet concentrations (mean diameter 10.5 μm) increased to a peak of 410 cm^{-3} . This cloud layer was again predominantly liquid. A spike in 2D-S concentrations was observed which imagery revealed was again due to drizzle droplets. These data were removed from the ice dataset.

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7 Primary IN parameterization comparison

Ice number concentrations as a function of altitude for science flight periods have been presented and here these observations are compared to calculations of the primary IN concentrations predicted using the D10 scheme, using aerosol concentrations (diameter $> 0.5 \mu\text{m}$) that were measured on each flight as input. DeMott et al. (2010) analysed datasets of IN concentrations over a 14 year period from a number of different locations and found that these could be related to temperature and the number of aerosol $> 0.5 \mu\text{m}$. The parameterisation provided an improved fit to the datasets and predicted 62% of the observations to within a factor of 2. Table 2 shows mean aerosol concentrations for measurement periods during each case, the input temperature to D10, the maximum median ice concentration used for comparison and the predicted IN concentration based on both the PCASP and CAS aerosol measurements (where available). During the spring measurement campaign it was possible to compare the CAS and PCASP probe data sets. Despite some variation in concentrations reported between the two instruments, D10 predicted IN values were found to be fairly insensitive to these differences. Grosvenor et al. (2012) highlighted that changes of about a factor of 4 produced a very limited change in the IN concentrations predicted by the scheme.

In spring case 1 the maximum median ice value reached 0.61 L^{-1} so predicted IN values were generally higher (between a factor of 2 and 4) than this median ice concentration observation. However peaks in ice concentrations of up to $\sim 10 \text{ L}^{-1}$, were also observed (Fig. 2) so on these occasions D10 significantly under predicts observed ice number concentrations when compared to these peak values. During spring case 2, maximum median ice concentration values were similar to spring case 1. Secondary ice production was observed close to the sea surface in this case so these higher median concentrations have been disregarded for the purposes of the D10 primary IN comparison. Aerosol measurements from the CAS were lower than from the PCASP but predicted IN values were in good agreement (less than a factor of 2) with the ob-

served maximum median concentration. The peak concentrations observed during the flight were $\sim 5 \text{ L}^{-1}$ (Fig. 4) and as in the first spring case D10 under predicted these peak concentrations by about a factor of 10.

During summer case 1 the minimum cloud temperatures were higher ($T = -10^\circ\text{C}$) than in the spring cases. Maximum median ice concentrations observed were also higher (3.35 L^{-1}). The origin of these enhanced concentrations is attributed to SIP, making a direct comparison with the D10 primary IN scheme difficult. Predicted IN concentrations from D10 were found to underestimate the maximum median ice concentrations observed in this summer case (due to secondary ice production), but were in agreement with the concentrations observed near cloud top, where the ice phase is likely to represent primary heterogeneous ice nucleation. Observed ice concentrations in summer case 2 were also higher than in the previous spring cases and similar to the first summer case. The second case had higher minimum cloud temperatures than in the first summer case ($T = -4.3^\circ\text{C}$). Due to effect of SIP at this temperature, it was not possible to compare D10 with the concentrations of ice observed in these clouds.

8 Discussion

Summaries of typical profiles during each case have been presented, with microphysics data encompassing all cloud penetrations during the science flights presented as a function of altitude shown in Figs. 10, 11 and 12. Figure 10 shows the cloud liquid droplet parameters, Fig. 11 the ice crystal concentration statistics and Fig. 12 the ice mass and diameter parameters. In each case (a) is spring case 1, (b) spring case 2, (c) summer case 1 and (d) summer case 2. The yellow lines on the ice plots (Fig. 8) show the approximate location of cloud top and cloud base altitudes deduced from liquid water content measurements exceeding 0.01 gm^{-3} from the CDP. It is notable that droplet concentrations (Fig. 10) are much higher in the second spring case than in the first spring case (max median values ~ 60 and $\sim 400 \text{ cm}^{-3}$ for spring case 1 and 2 respectively) and this is attributed to differences in aerosol concentrations. N_{drop} are

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regions with few (generally $< 1 \text{ L}^{-1}$) ice crystals, formed through heterogeneous ice nucleation at these temperatures, were observed in both cases (Fig. 11c and d). LWCs in summer case 1 were lower than the spring cases (median values $< \sim 0.1 \text{ g m}^{-3}$) and similar in shape to the uniform profiles seen in the spring cases. The second summer case had higher median LWCs (up to 0.35 g m^{-3}) and showed much more variability with a number of increases and decreases in median LWC values with altitude (Fig. 10d). Median cloud top ice concentrations in summer case 1 were similar to the spring cases ($\sim 0.2 \text{ L}^{-1}$) (Fig. 11d), however maximum median values lower down in the cloud reached 3.35 L^{-1} (Table 2), about a factor of 14 higher than in the spring cases. Peaks in ice number concentrations around the -5°C level reached between $30\text{--}40 \text{ L}^{-1}$. During the summer, the clouds spanned the temperature range -3 to -8°C , where a well-known mechanism of secondary ice production operates through splintering during riming; the Hallett–Mossopp process (H–M). The observations in this case, of liquid water together with ice particles at temperatures around -5°C , are consistent with this process being active and enhancing ice number concentrations (Figs. 7 and 9). Time series (Figs. 6 and 8) showed more variation than in the spring cases. Distinct liquid cloud tops were still evident, but at lower altitudes significant variations in LWCs, droplet number concentrations and ice number concentrations were seen together with gap regions where little or no cloud was present. On a number of occasions predominantly liquid conditions were swiftly replaced by regions of high concentrations of columnar ice crystals. Some of these transitions took place over ~ 1 s or horizontal distance of the order 60 m. These rapid fluctuations were attributed to the contributions from the H–M process. The process of glaciation through secondary enhancement of ice number concentrations is likely to have caused some of this increased variability in cloud properties too, with liquid droplets quickly being removed through depletion of liquid water by the ice phase. The cloud layers during summer case 2 spanned a higher temperature range than summer case 1. Cloud tops were around -4°C , and median ice number concentrations reached maximum values of 2.5 L^{-1} , about an order of magnitude higher than in the spring cases. Time series (Fig. 8) and percentile

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plots (Fig. 11d) showed peaks in ice number concentrations to $\sim 25 \text{ L}^{-1}$ and in these regions probe imagery revealed distinctive columnar ice crystals likely to have grown from splinters produced via H–M, into habits typical of growth at these temperatures around -4°C . In addition, the formation of high ice concentrations may have led to the dissipation of some liquid cloud regions below cloud top due to consumption of the liquid phase by ice crystals growing by vapour diffusion (i.e. ice crystal growth via the Bergeron–Findeisen (B-F) process (Bergeron, 1935). This is consistent with the observed summer clouds being more broken than the clouds observed during spring. However, as discussed in the introduction, it is also recognised that cloud-radiation interactions may lead to the separation of cloud layers during the Arctic summer.

Comparison of the observed N_{ice} with the D10 parameterization of primary ice nuclei numbers revealed that during the spring case 1, maximum median N_{ice} was lower than the primary IN concentrations predicted by D10, but similar in spring case 2. Peaks in N_{ice} were much higher than the D10 IN predictions, by an amount depending on the aerosol measurement period used as input to D10 (Table 2). In the summer cases the enhancement of N_{ice} through the H–M process made a realistic comparison difficult. Despite this difficulty, the first summer case had cloud top temperatures that were just outside the H–M temperature zone (-10°C) and median N_{ice} in this region was $\sim 0.2 \text{ L}^{-1}$, which is within a factor of 2 of values predicted by D10 (Table 2). At lower altitudes the increase in cloud temperatures allowed rime-splintering to enhance concentrations to above what would be expected via primary heterogeneous ice nucleation. In the second summer case cloud top temperatures were higher (-4°C), and enhancement of the ice crystal number concentrations through SIP prevented observations of any first ice by primary nucleation being made. Ice crystal number concentrations were thus enhanced to values above what was predicted by D10 throughout the depth of the cloud. Whilst primary ice nucleation is identified as the most important ice forming process in the spring clouds, the summer stratocumulus ice concentrations were dominated by secondary ice production via the H–M process as discussed. Due to this SIP

enhancement, ice concentrations in summer reached much higher values than those observed anywhere in the spring cases.

The microphysical structure of the spring and summer stratocumulus layers was found to be consistent with previous observations of arctic clouds. We observed generally low droplet number concentrations with increased concentrations during incursions of higher aerosol loadings. This is consistent with observations by Verlinde et al. (2007). During spring cases, LWCs and liquid droplet size increased uniformly to cloud top, however during summer months the vertical structure of cloud layers was more variable (e.g. Hobbs and Rangno, 1998). During spring cases in particular, liquid cloud tops at distinct temperature inversions continually precipitated low concentrations of ice into the cloud below, which has been observed previously in the Arctic. During the Arctic summer, Hobbs and Rangno (1998) observed generally higher ice concentrations with columnar and needle ice crystals in concentrations of “tens per litre” where stratocumulus cloud top temperatures were between -4 and -9°C . The summer cases we observed contained median values of N_{ice} that were 4–6 times greater than we observed in the spring cases. In the spring, the cloud layers were colder than the temperature range within which H–M is active, and accordingly contained peak concentrations of ice closer to predictions from D10. In the summer cases, the clouds spanned a warmer temperature range between about 0 and -10°C , leading to low concentrations of primary ice that when conditions became suitable, were then enhanced through rime-splintering. During the spring we also observed cloud that penetrated into the inversion layer, rather than being capped below it. On average the cloud top was seen to extend ~ 30 m into the inversion layer over which range the mean temperature increase was $\sim 1.6^{\circ}\text{C}$.

Changes in aerosol concentrations and composition have been suggested as a possible factor in explaining previous observations of the glaciation of arctic clouds at different temperatures (Curry et al., 1996). During spring case 2 higher concentrations of aerosol were observed when compared to spring case 1. Droplet number concentrations were also much higher in spring case 2, generally $300\text{--}400\text{ cm}^{-3}$ in comparison

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to spring case 1 where concentrations were generally $\sim 50\text{--}100\text{ cm}^{-3}$. Despite this, no significant difference was observed in the ice number concentrations. However, it should be noted that despite the higher total concentrations, the population of aerosol $> 0.5\ \mu\text{m}$ was not significantly enriched in spring case 2 compared to the spring case 1. D10 has a dependency only on this portion of the aerosol size distribution, so may explain the similar primary ice number concentrations for both spring case studies.

Grosvenor et al. (2012) studied stratocumulus clouds in the Antarctic over the Larsen C ice shelf. These observations contained periods where temperatures were comparable to those in the spring cases studied here. The lower layers of Antarctic cloud were also reported to contain higher concentrations of ice produced via the H–M process, similar to the summer cases that we have discussed. A summary of some of the measurements reported from the Antarctic in Grosvenor et al. (2012) can be found in Table 3. Measurements of cloud regions outside the H–M temperature zone revealed very low ice number concentrations, with maximum values about 2 orders of magnitude lower than those observed in the spring cases reported here. Aerosol concentrations from a CAS probe (similar to the one deployed in this study) reported generally lower concentrations of aerosol particles $D_p > 0.5\ \mu\text{m}$. The D10 IN predictions in the Antarctic were reported to compare better with maximum, rather than mean ice values. A similar result was found in this study where predicted primary IN values were greater than observed median values. However, when comparing with peak ice concentration values the scheme significantly under-predicted these. Grosvenor et al. (2012) discussed the possibility that due to the D10 parameterisation being based on mean IN concentrations from many samples, the finding that IN predictions compared well with the maximum values rather than mean values may suggest the scheme was over predicting IN concentrations generally in the Antarctic (for these particular cases at least). In the H–M layer in the Antarctic over Larsen C, ice crystal number concentrations were found to be higher than those observed in colder temperature regimes (not spanning the H–M temperature range), in keeping with the findings from the Arctic presented this paper. However the concentrations produced by the H–M process in the Antarctic were

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generally only a few per litre, approximately an order of magnitude lower than those observed during the summer cases in the Arctic.

9 Conclusions

Detailed microphysics measurements made in Arctic stratocumulus cloud layers during the early spring and summer, have been presented.

- Two spring and two summer cases were presented. The cloud layers during summer cases spanned a warmer temperature range ($\sim 0^{\circ}\text{C} \geq T > -10^{\circ}\text{C}$) than in spring (generally $\sim -10^{\circ}\text{C} \geq T > -20^{\circ}\text{C}$).
- Spring case 2 had significantly higher aerosol concentrations ($\sim 300\text{--}400\text{ cm}^{-3}$) compared to the first spring case ($\sim 50\text{--}100\text{ cm}^{-3}$). Despite this difference, ice number concentrations were found to be similar in both spring cases, suggesting the source of the increased aerosol concentrations was not providing additional IN that were efficient over the temperature range $-10^{\circ}\text{C} > T > -20^{\circ}\text{C}$.
- In the spring cases, cloud layers appeared more uniform with steady increases in LWC and cloud droplet size to cloud top, where low concentrations ($< 1\text{ L}^{-1}$) of ice were frequently observed to precipitate through the depth of the cloud layer. The small irregular particles observed at cloud top grew to a median diameter $\sim 500\text{ }\mu\text{m}$ in both cases with peaks in diameter $> 1000\text{ }\mu\text{m}$ as the crystals descended through the cloud. 2D-S imagery revealed the dominant growth habit to be dendritic in nature. The summer cases consisted of multiple cloud layers that were observed to be more variable than in the spring. However, liquid cloud top regions were still evident and ice was again observed to precipitate into the cloud layers below.
- The maximum median ice number concentrations observed within cloud layers during the summer cases were approximately a factor of 5 (or more) higher than in

the spring cases. This enhancement in the ice number concentrations is attributed to the contribution of secondary ice production through the H–M process.

- This finding suggests that low level summer stratocumulus clouds situated in the H–M temperature zone in the Arctic may contain significantly higher ice number concentrations than in spring clouds due to the temperature range of the former spanning the active H–M temperature zone.
- Predicted values from the DeMott et al. (2010) scheme of primary ice nuclei, using aerosol measurements obtained during the science flights as input, tended to overpredict IN concentrations compared to the observed maximum median ice crystal number concentrations during the spring, but under-predict IN when compared to peak ice crystal concentrations. During the summer cases, due to contributions from secondary ice production, the scheme predicted significantly lower values of ice particles than those observed.
- Grosvenor et al. (2012) observed lower concentrations of aerosol $> 0.5 \mu\text{m}$ in the Antarctic when compared to similar measurements made in the Arctic. They found that IN predictions using D10 agreed better with their observed peak ice concentration values rather than their maximum mean values. They measured approximately an order of magnitude lower primary ice concentrations in summer Antarctic clouds than in our spring Arctic cases, but did observe enhancement through SIP in warmer cloud layers where concentrations increased to a few per litre. These were still about an order of magnitude less than the enhanced concentrations observed in the Arctic summer cases presented here, but were similar to the peak values observed in spring cases over the Arctic (where no SIP was observed).

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Appendix A:

A1 Profiled ascent A1

During profile A1 the aircraft (travelling south) made a profiled ascent from 300 m above the sea surface, reaching cloud base at 650 m, identified using a Liquid Water Content threshold of $LWC > 0.01 \text{ gm}^{-3}$, as derived from CDP data. Below cloud base the 2D-S probe revealed low concentrations ($< 0.5 \text{ L}^{-1}$) of irregular snow (Fig. 3d) particles (mean size $\sim 530 \mu\text{m}$) that had precipitated from the cloud layer above. As the aircraft climbed through cloud base, temperatures decreased to -11°C . CDP droplet concentrations (N_{drop}) (10 s averaged values) increased to $\sim 80 \text{ cm}^{-3}$, LWCs peaked at $\sim 0.2 \text{ gm}^{-3}$ and mean droplet diameters were $\sim 8 \mu\text{m}$. Measurements from the 2D-S showed ice crystals with mean size $\sim 415 \mu\text{m}$ in low concentrations, $\sim 1 \text{ L}^{-1}$. Images from the 2D-S revealed irregular snow particles with some dendritic habits coexisting with small liquid droplets. As the ascent continued the aircraft encountered a layer containing higher N_{ice} at -14°C . Ice crystals consisted of snow particles (mean size $350 \mu\text{m}$) in concentrations $\sim 4 \text{ L}^{-1}$. Probe imagery showed these to be a mixture of large irregular ice crystals, small, more pristine plate-like crystals and some crystals with columnar habits. The highest 10 s mean N_{ice} , reached $\sim 6 \text{ L}^{-1}$ with peak values $\sim 15 \text{ L}^{-1}$. These were observed in a region approximately 500 m below cloud top. Maximum 10 s averaged Ice Water Content (IWC) reached 0.2 gm^{-3} with peaks up to 0.3 gm^{-3} in the same region. Particle images here revealed (Fig. 3b) irregular ice crystals together with a few smaller pristine plates. The mid region of this stratocumulus deck also consisted of liquid droplets (mean diameter $\sim 13 \mu\text{m}$) in concentrations $\sim 75 \text{ cm}^{-3}$, and $LWC \sim 0.3 \text{ gm}^{-3}$, with some 1 s integration periods being as high as 0.5 gm^{-3} . As the aircraft approached cloud top, where the lowest temperature recorded was -19.5°C , N_{ice} reduced to $\sim 0.5 \text{ L}^{-1}$ with mean sizes of $285 \mu\text{m}$, however this region was dominated by liquid droplets (mean diameter $17 \mu\text{m}$) with N_{drop} up to 95 cm^{-3} , and LWC values peaking at 0.7 gm^{-3} . Imagery from the 2D-S revealed many small

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droplets together with numerous small irregular ice crystals in this cloud top region. After measuring the vertical structure of the cloud layer, which was approximately 1 km in depth, the aircraft penetrated cloud top at 1675 m and passed through an inversion layer where the temperature increased to -13°C .

5 A2 Profiled descent A3

Following another ascent, the aircraft performed a profiled descent (A3) from the inversion layer, $T = -13^{\circ}\text{C}$, penetrating cloud top at 1569 m a.s.l. where $T = -16^{\circ}\text{C}$. As the aircraft descended, LWC increased rapidly to 0.9 gm^{-3} at 30 m below cloud top, the highest LWC recorded at any point during the flight. Mean droplet diameters in this region were $\sim 23\text{ }\mu\text{m}$ in concentrations of $\sim 90\text{ cm}^{-3}$. 2D-S images revealed many small liquid droplets with a few small (mean diameter $190\text{ }\mu\text{m}$) irregular ice crystals (Fig. 3a) with $N_{\text{ice}} \sim 1\text{ L}^{-1}$. The region immediately below this cloud top layer, between 1520 and 1275 m, exhibited a steady decline in LWC while droplet concentrations and N_{ice} maintained similar values to those observed in the cloud top region. Mean ice crystal diameters increased markedly to $520\text{ }\mu\text{m}$ before LWCs eventually fell to below the threshold value (0.01 gm^{-3}), marking the base of an upper layer of cloud. A subsequent cloud layer, 750 m below, was then encountered. In the clear air region separating these two cloud layers temperatures rose by around 5 to -11°C and large ($\sim 760\text{ }\mu\text{m}$) irregular snow particles, some of which exhibited dendritic growth habits, were observed. Precipitation concentrations were generally $< 0.5\text{ L}^{-1}$. Mean IWCs in this precipitation zone were $\sim 0.01\text{ gm}^{-3}$. The particles observed falling from the higher cloud layer descended into the cloud layer below at 1275 m a.s.l. In the top of this lower cloud layer ($T = -11^{\circ}\text{C}$) LWCs rose to 0.4 gm^{-3} with N_{drop} (mean diameter $15\text{ }\mu\text{m}$) increasing to $\sim 120\text{ cm}^{-3}$ while N_{ice} increased to $\sim 1\text{ L}^{-1}$, 2D-S probe imagery in this region revealed the presence of larger snow particles (mean diameters $\sim 815\text{ }\mu\text{m}$). As the aircraft descended further, LWCs gradually decreased while N_{drop} remained fairly constant before

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reaching cloud base at 280 m, (much closer to sea level than in profiles A1 and A2). Below cloud base precipitating snow (mean particle size $\sim 625 \mu\text{m}$) was observed.

Appendix B:

B1 Profiled ascent B2

5 During profiled ascent B2 (prior to profile descent B1 above) the aircraft climbed from below cloud base at 190 m ($T = -5^\circ\text{C}$) travelling initially through snow precipitation in concentrations peaking at $\sim 3 \text{L}^{-1}$ (mean diameter $420 \mu\text{m}$). Images revealed dendritic ice crystals that had descended from the cloud layer above (Fig. 5c). IWCs in this region peaked at 0.025g m^{-3} . Cloud base during this profile was less well defined than in
10 later ascents with variable LWCs and droplet number concentrations before a more defined cloud base was encountered at 1010 m. N_{drop} then increased rapidly to 270cm^{-3} (mean diameter $\sim 12.5 \mu\text{m}$) while LWCs increased more gradually to $\sim 0.4 \text{g m}^{-3}$. N_{ice} through this region showed a decline to $< 0.1 \text{L}^{-1}$, and consisted of precipitating snow particles with a mean diameter of $430 \mu\text{m}$. Closer to cloud top (1410 m) ice crystal
15 number concentrations increased, to peak values of $\sim 1 \text{L}^{-1}$. Images (Fig. 5b) showed smaller crystals (mean diameter $\sim 370 \mu\text{m}$) at this higher altitude, with evidence of hexagonal habits and peak values of IWC $\sim 0.04 \text{g m}^{-3}$. Droplet concentrations towards cloud top were similar to lower in the cloud, while LWCs increased to 0.6g m^{-3} and mean droplet diameter increased to $\sim 15 \mu\text{m}$. The coldest temperature reached within
20 the cloud layer was -18°C , but cloud top (at $\sim 1530 \text{m}$) was warmer by 1°C . A further increase of 1°C was observed as the aircraft ascended through the inversion layer. The depth of this cloud layer (520 m) was significantly less than that observed during the previous spring case cloud layer penetrations.

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B2 Constant altitude runs B3 and B4

During straight and level run (SLR) B3 the aircraft flew below cloud base at 390 m a.s.l. to characterise precipitation. During B3 the aircraft briefly traversed a region of low cloud with high N_{drop} (peaking at $\sim 520 \text{ cm}^{-3}$) but generally low LWCs ($< 0.1 \text{ gm}^{-3}$). These cloud droplets were small (mean diameter $\sim 6 \mu\text{m}$). 2D-S imagery also revealed small drops were present together with snow crystals (mean diameter $\sim 370 \mu\text{m}$) that were precipitating into these brief regions of low cloud. During B3 temperatures increased from -12 to -10°C . Crystal habits in the out of cloud regions were dominated by aggregates of dendrites and some pristine ice crystals ($\sim 0.5 \text{ L}^{-1}$). Here, LWCs were below 0.01 gm^{-3} , although the 2D-S also detected drizzle droplets precipitating from the cloud layer above (mean concentration $\sim 0.2 \text{ L}^{-1}$). Later in B3 the aircraft left its constant altitude and descended to 80 m a.s.l. ($T = -8.5^\circ\text{C}$). Mean N_{ice} increased to $\sim 2 \text{ L}^{-1}$ with peaks up to 4 L^{-1} . There was a corresponding increase in 2D-S droplet concentrations to a mean of $\sim 1 \text{ L}^{-1}$. 2D-S imagery shows the presence of small columnar shaped ice crystals (similar to those shown in Fig. 5d), together with larger snow particles and drizzle droplets. CDP LWC was $< 0.01 \text{ gm}^{-3}$ in this region, since the larger drizzle droplets measured by the 2D-S were outside the CDP size range. In this region of enhanced N_{ice} , just above the sea surface, IWCs, which were generally $< 0.01 \text{ gm}^{-3}$ in the below cloud base region, increased to peak values of 0.04 gm^{-3} .

At the start of run B4, prior to undertaking a mainly straight and level run (SLR) initially to the NW, the aircraft first descended from the inversion layer ($T \sim -14^\circ\text{C}$) into the cloud top (1050 m a.s.l.). LWC initially rose sharply to a peak of 0.5 gm^{-3} before gradually falling away to a mean value $\sim 0.3 \text{ gm}^{-3}$. Mean droplet concentrations over a ~ 5 min period were 340 cm^{-3} (mean diameter $11 \mu\text{m}$) and the 2D-S imagery revealed the presence of small droplets together with large snow crystals (mean diameter $730 \mu\text{m}$) in concentrations $< 0.1 \text{ L}^{-1}$ and IWCs of 0.03 gm^{-3} . At 12:40 UTC a generally cloud free region was encountered and sampled for ~ 4 min before re-entering cloud again. During this period the aircraft was turned onto a reciprocal heading at the NW

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single pixel ($10\ \mu\text{m}$) particles were also measured. These likely represent the small droplets detected by the CDP in this region (mean diameter $13.5\ \mu\text{m}$) in concentrations of $125\ \text{cm}^{-3}$. Later during C1.2, N_{ice} fell to values $< 0.25\ \text{L}^{-1}$. The aircraft performed a profiled descent at the start of C1.3, descending 200 m to $\sim 2720\ \text{m}$ ($T = -4\ ^\circ\text{C}$).

5 During the descent, LWCs and droplet number concentrations fell to near zero values while N_{ice} increased to peak values of $5\ \text{L}^{-1}$ (and IWC peaked at $0.02\ \text{g m}^{-3}$). 2D-S images again revealed the presence of small (mean diameter $255\ \mu\text{m}$) rimed irregular ice crystals and ice crystals of columnar habit. In the temperature range spanned by this cloud, these observations are consistent with the contribution of secondary ice production (SIP) through rime-splintering. During C1.3 further N_{ice} peaks up to $5\ \text{L}^{-1}$ consisting of columnar particles and irregular ice crystals were observed (Fig. 7b). The liquid phase of the cloud in this region was much more variable than nearer to cloud top. Increases in peak LWCs to $0.01\ \text{g m}^{-3}$ were seen together with an increase in droplet number concentrations to $\sim 150\ \text{cm}^{-3}$ (mean diameter $13.5\ \mu\text{m}$). These occurred between periods where LWC values were near zero and the cloud was predominantly glaciated.

15 During C1.4 the aircraft descended 300 m to 2450 m ($T = -3\ ^\circ\text{C}$). During this run the time between peaks in N_{drop} increased, while the highest N_{ice} measured during this science flight were observed (peaking at $N_{\text{ice}} = 35\ \text{L}^{-1}$). IWCs peaked at $0.2\ \text{g m}^{-3}$, which is significantly greater than values observed elsewhere in this cloud system. 2D-S imagery (Fig. 7c) reveals that these high ice crystal number concentrations were dominated by columns (mean diameter $260\ \mu\text{m}$), which at times were seen together with small liquid droplets. These observations are consistent with SIP through the H-M process.

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Appendix D:

D1 Profiled descent D1

Well into the flight, the BAS aircraft performed a profiled descent from cloud top at 3700 m to 2400 m over the temperature range -5.2 to 3°C . At cloud top, LWCs rose to a peak of 0.3 g m^{-3} , with peak N_{drop} (mean diameter $12.5\text{ }\mu\text{m}$) up to 270 cm^{-3} . N_{ice} , initially close to zero, rose to peaks of 6 L^{-1} with IWCs up to 0.1 g m^{-3} . 2D-S images (Fig. 9a) showed columnar ice crystals (mean diameter $350\text{ }\mu\text{m}$) in this region, together with liquid droplets. At times swift transitions between predominantly liquid and glaciated conditions were observed. At 3500 m ($T = -3.5^{\circ}\text{C}$) the CDP stopped measuring significant values of LWC ($> 0.01\text{ g m}^{-3}$) and this appeared to mark a gap region in the cloud layer of approximately 100 m in depth. The 2D-S did detect low N_{ice} in this region. These were generally below $< 0.5\text{ L}^{-1}$. When the aircraft descended into the lower cloud layer ($T = -2^{\circ}\text{C}$) LWCs increased to peak values of 1 g m^{-3} , where N_{drop} (mean diameter $13.5\text{ }\mu\text{m}$) increased to values as high as 250 cm^{-3} . 2D-S imagery revealed few ice crystals in this region but high drizzle drop concentrations.

At 2800 m ($T = 0^{\circ}\text{C}$) a further period of drizzle droplets was observed in the 2D-S imagery. These again appeared stretched and made it impossible to separately identify ice in the data set, so there is no reliable ice crystal mass and number concentration data in this region. At this time, CDP LWCs peaked at 0.4 g m^{-3} and droplet concentrations varied from close to zero to up to $\sim 350\text{ cm}^{-3}$. The mean diameter of the droplets measured by the CDP was $10\text{ }\mu\text{m}$. As the aircraft descended towards its minimum descent altitude large variations in LWCs and droplet concentrations continued to be observed with peaks up to 0.2 g m^{-3} and 420 cm^{-3} respectively.

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Table 1. Flight numbers, run numbers, and their associated time intervals, altitude and temperature range for the four ACCACIA case studies presented.

Flight	Run Number	Time (UTC)	Altitude (m)	Temperature (°C)
B761	A1	13:13:26–13:16:43	1850–50	–19 to –5
B761	A2	13:04:40–13:10:33	300–1850	–8 to –19
B761	A3	13:23:20–13:33:19	1700–50	–19 to –7
B768	B1	11:45:16–11:54:02	1600–50	–17 to –9
B768	B2	11:38:39–11:44:59	50–1600	–17 to –4
B768	B3	12:01:30–12:19:08	400–50	–12 to –9
B768	B4	12:32:20–12:48:14	1300–1050	–16 to –14
M191	C1.1	08:53:45–09:00:00	~ 2950	~ –7
M191	C1.2	09:00:00–09:06:50	~ 2900	~ –6
M191	C1.3	09:06:50–09:13:35	~ 2750	~ –5
M191	C1.4	09:13:35–09:21:09	2750–2250	–4 to –2
M191	C2	10:14:58–10:33:51	3350–2300	–7 to –3
M192	D1	12:58:58–13:06:02	3100–3750	–5 to –1
M192	D2	12:19:10–12:48:16	3100–3750	–5 to –1

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Table 2. Measurements of: aerosol concentrations $> 0.5 \mu\text{m}$ from the CAS and PCASP probes, together with predicted primary IN number using the DeMott et al. (2010) (D10) scheme (with either CAS or PCASP aerosol concentration data as input). Observed minimum median cloud temperatures were input to D10, and IN predictions were compared with observed maximum median ice concentrations.

Flight	Max Median Ice (L^{-1})	Min Median Temp ($^{\circ}\text{C}$)	Max RH (%)	CAS Aerosol Conc (cm^{-3})	PCASP Aerosol Conc (cm^{-3})	Predicted CAS IN value (L^{-1})	Predicted PCASP IN value (L^{-1})
Case 1a	0.61	−18.7	90.3	0.99 ± 0.25	3.13 ± 1.74	$1.02 \pm 1.14/0.88$	$1.80 \pm 2.25/1.20$
Case 1b	0.61	−18.7	22.16	0.14 ± 0.1	4.94 ± 2.22	$0.38 \pm 0.50/0.21$	$2.26 \pm 2.72/1.68$
Case 1c	0.61	−18.7	85.43	1.48 ± 0.37	4.04 ± 2.25	$1.24 \pm 1.34/1.08$	$2.05 \pm 2.55/1.37$
Case 2a	0.47	−16.2	69.68	1.50 ± 0.30	3.23 ± 1.68	$0.76 \pm 0.82/0.69$	$1.05 \pm 1.26/0.77$
Case 2b	0.47	−16.2	92.60	2.40 ± 0.32	4.96 ± 2.28	$0.93 \pm 0.98/0.87$	$1.27 \pm 1.49/0.97$
Case 2c	0.47	−16.2	93.86	2.07 ± 6.57	3.07 ± 1.86	$0.87 \pm 1.61/$	$1.03 \pm 1.26/0.69$
Case 3a	3.35	−10	89.37	0.06 ± 0.07	–	$0.06 \pm 0.07/$	–
Case 3b	3.35	−10	59.66	0.15 ± 0.11	–	$0.08 \pm 0.09/0.05$	–
Case 3c	3.35	−10	89.79	0.33 ± 0.76	–	$0.10 \pm 0.13/$	–
Case 3d	3.35	−10	89.70	0.48 ± 0.21	–	$0.11 \pm 0.12/0.09$	–
Case 4a	2.50	−4.3	79.70	3.73 ± 1.03	–	$0.009 \pm 0.009/0.009$	–
Case 4b	2.50	−4.3	73.46	4.03 ± 0.58	–	$0.009 \pm 0.009/0.009$	–
Case 4c	2.50	−4.3	31.57	0.24 ± 0.14	–	$0.007 \pm 0.007/0.006$	–

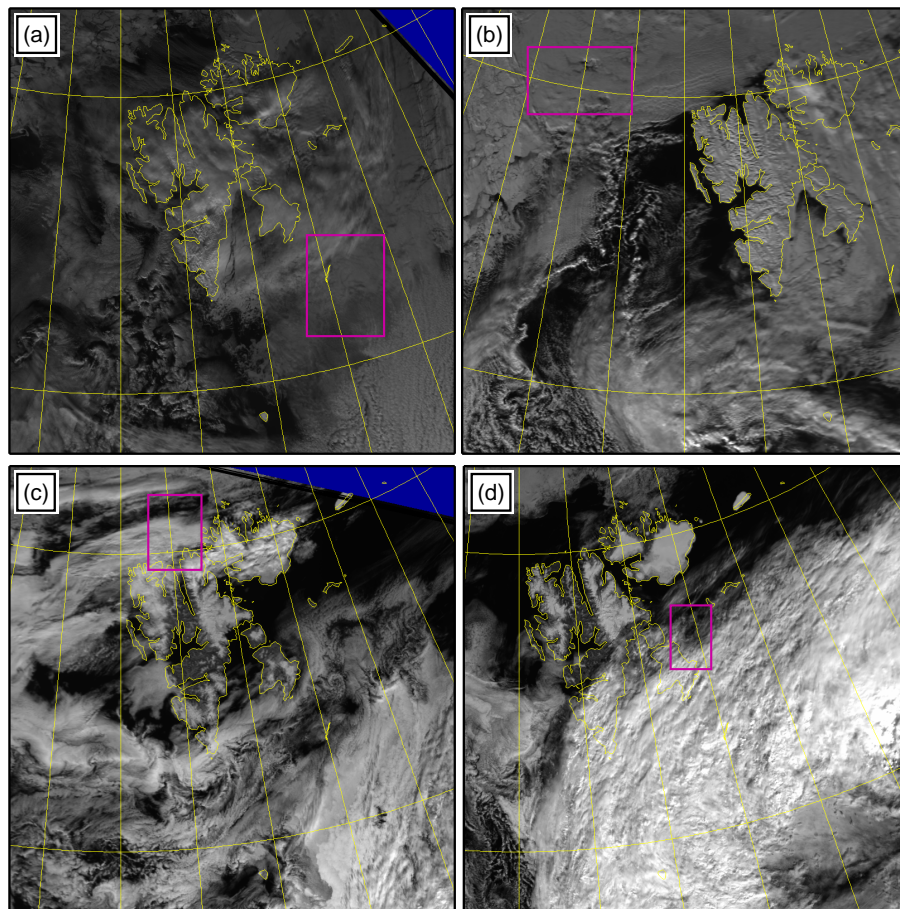


Figure 1. AVHRR visible satellite imagery for spring case 1 (a), spring case 2 (b), summer case 1 (c) and summer case 2 (d). Science flight area highlighted by purple boxes in each figure.

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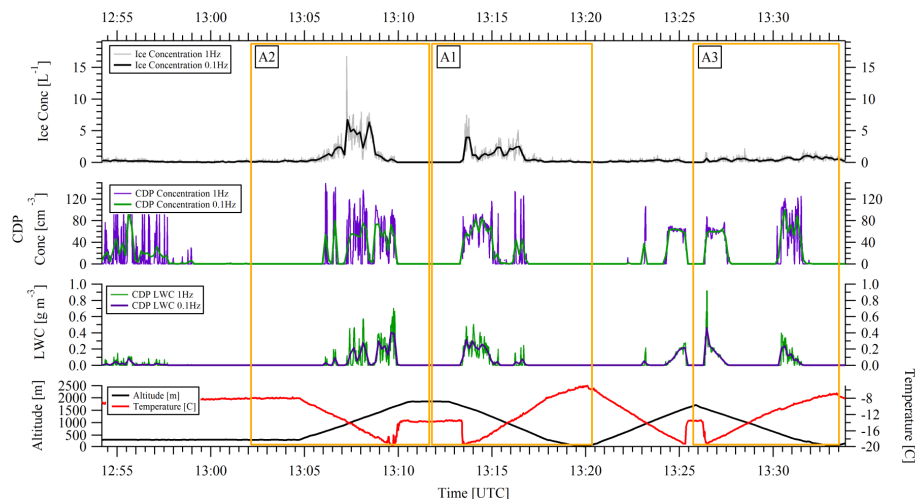


Figure 2. Microphysics time series for spring case 1. Data includes temperature ($^{\circ}\text{C}$) and altitude (m) (lower panel) together with 1 and 10 s data sets for CDP liquid water content (g m^{-3}) (panel 2 from bottom), CDP cloud particle number concentration (cm^{-3}) (panel 3), and ice water content (g m^{-3}) and ice number concentrations (L^{-1}) (top panel). Profiles A2 and A3 are described in Appendix A.

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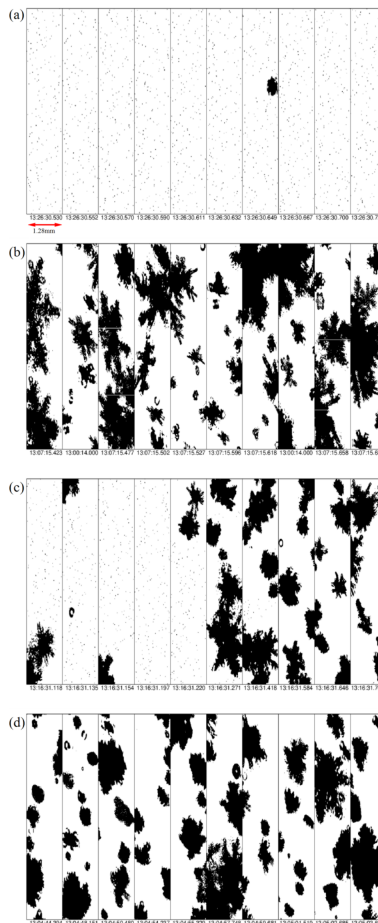


Figure 3. Images from the 2D-S cloud probe during spring case 1 from: (a) a cloud top region during A1; (b) 500 m below cloud top during A2; (c) region of swift transitions between ice and liquid and (d) precipitation region below cloud base.

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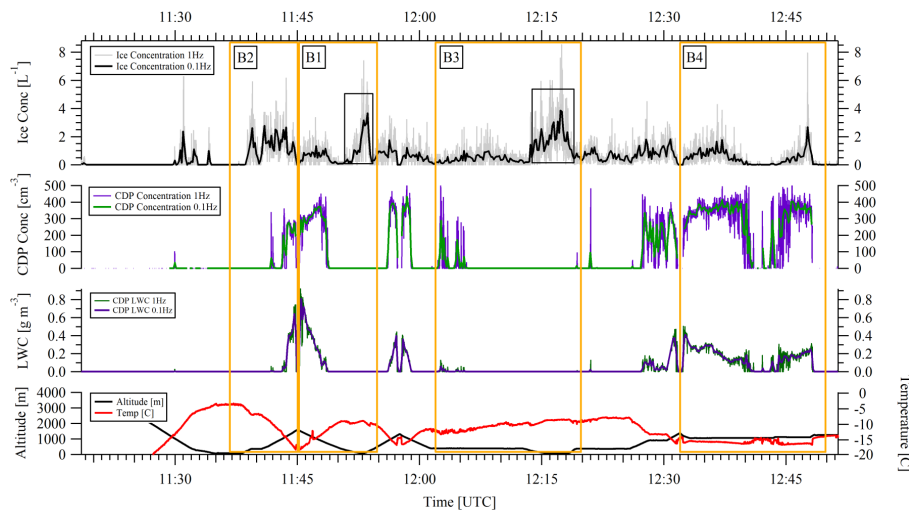


Figure 4. Microphysics time series data for spring case 2. Data includes temperature ($^{\circ}\text{C}$) and altitude (m) (lower panel) 1 and 10 s data sets for CDP liquid water content (g m^{-3}) and CDP concentration (cm^{-3}) (middle panels), and ice water content (g m^{-3}) and ice number concentrations (L^{-1}) (top panel). Profiles B2, B3 and B4 are described in Appendix B.

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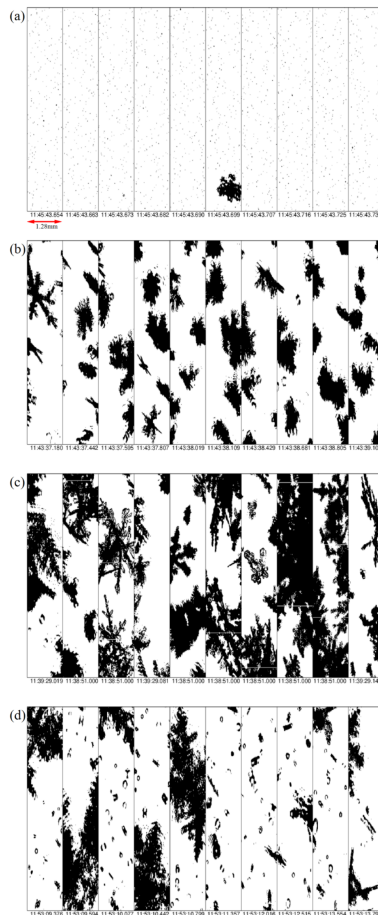


Figure 5. Images from the 2D-S cloud probe from spring case 2 for: **(a)** cloud top during B1; **(b)** profiled ascent during B2; **(c)** dendritic ice in the cloud base region during B2 and **(d)** columnar ice above the sea surface during B2.

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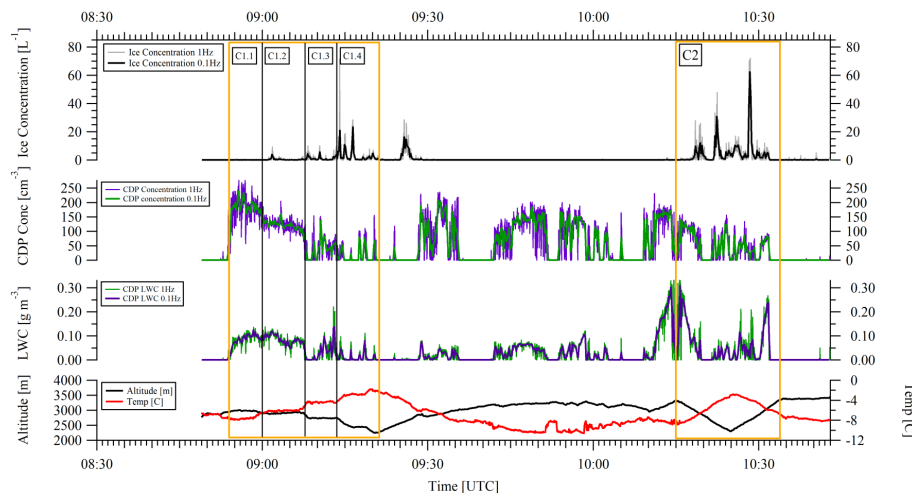


Figure 6. Microphysics time series data for summer case 1. Data includes temperature ($^{\circ}\text{C}$), altitude (m) (lower panel) together with 1 and 10 s data sets for CDP liquid water content (g m^{-3}) (second panel up), CDP concentration (cm^{-3}), ice water content (g m^{-3}) and ice number concentrations (L^{-1}) (top panel). Flight segments C1.1, C1.2, C1.3 and C1.4 are described in Appendix C.

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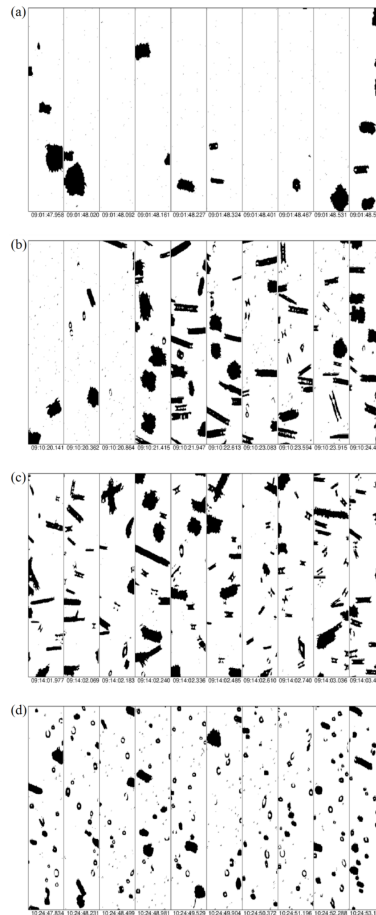


Figure 7. Images from the 2D-S cloud probe from summer case 1 for: **(a)** small irregular ice during C1.2; **(b)** and **(c)** secondary ice production during C1.3 and C1.4 respectively, and **(d)** ice together with drizzle during C2.

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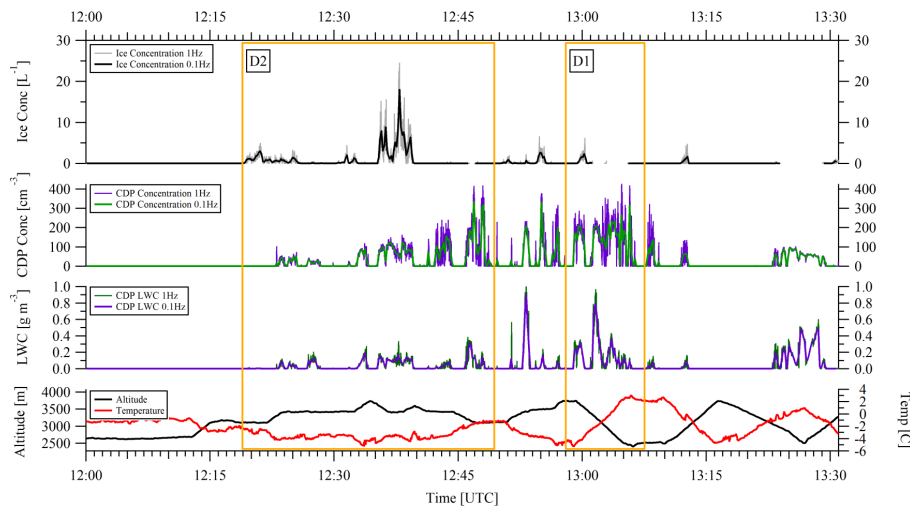


Figure 8. Microphysics time series data for summer case 2. Data includes temperature ($^{\circ}\text{C}$), altitude (m) (lower panel) together with 1 and 10 s data sets for CDP liquid water content (g m^{-3}), CDP concentration (cm^{-3}) (middle panels), ice water content (g m^{-3}) and ice number concentrations (L^{-1}) (top panels). Profile D1 is described in Appendix D.

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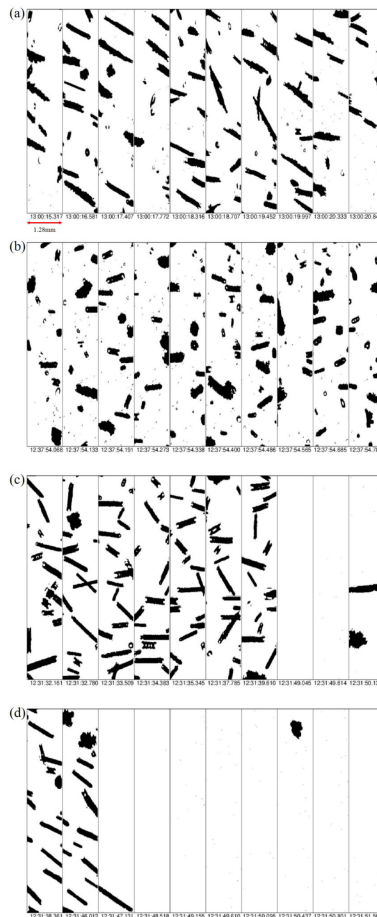


Figure 9. 2D-S cloud probe imagery for summer case 2 showing: **(a)** columnar ice during D1; **(b)** images of columns together with liquid during D2 and swift transitions between **(c)** glaciated and **(d)** liquid phases during D2.

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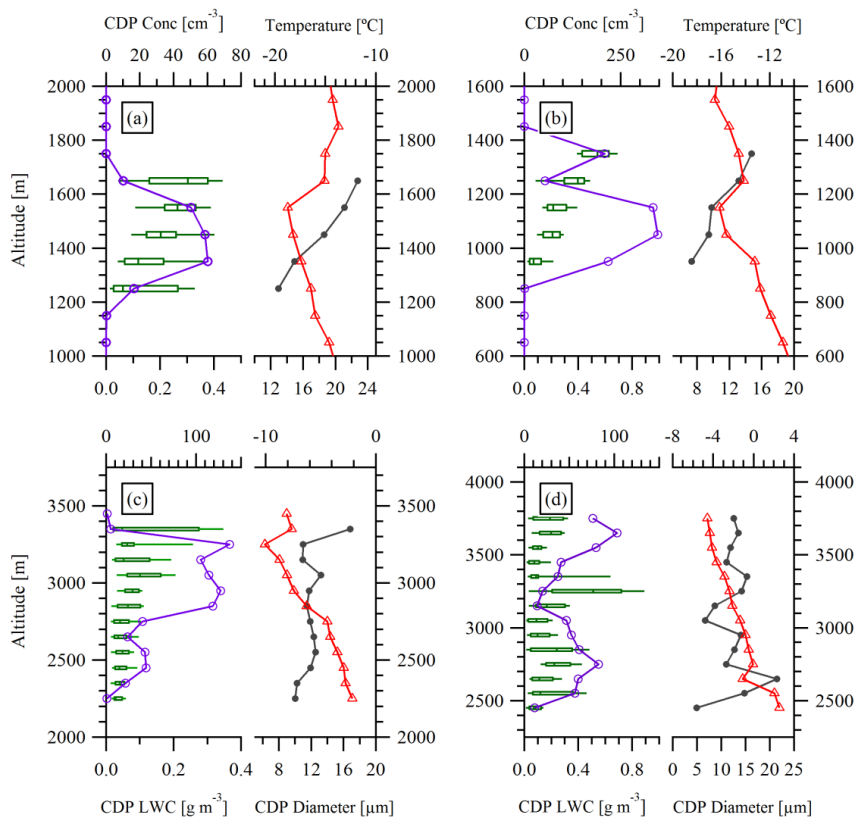


Figure 10. Percentile plots (50th, 25th, 75th percentiles, whiskers to 10 and 90 %) as a function of altitude for LWC from CDP (green), and median droplet number concentration (purple), median droplet diameter (grey) and median temperature (red). Data are averaged over 100 m deep layers. **(a–d)** are for Spring Case 1, Spring Case 2, Summer Case 1 and Summer Case 2 respectively.

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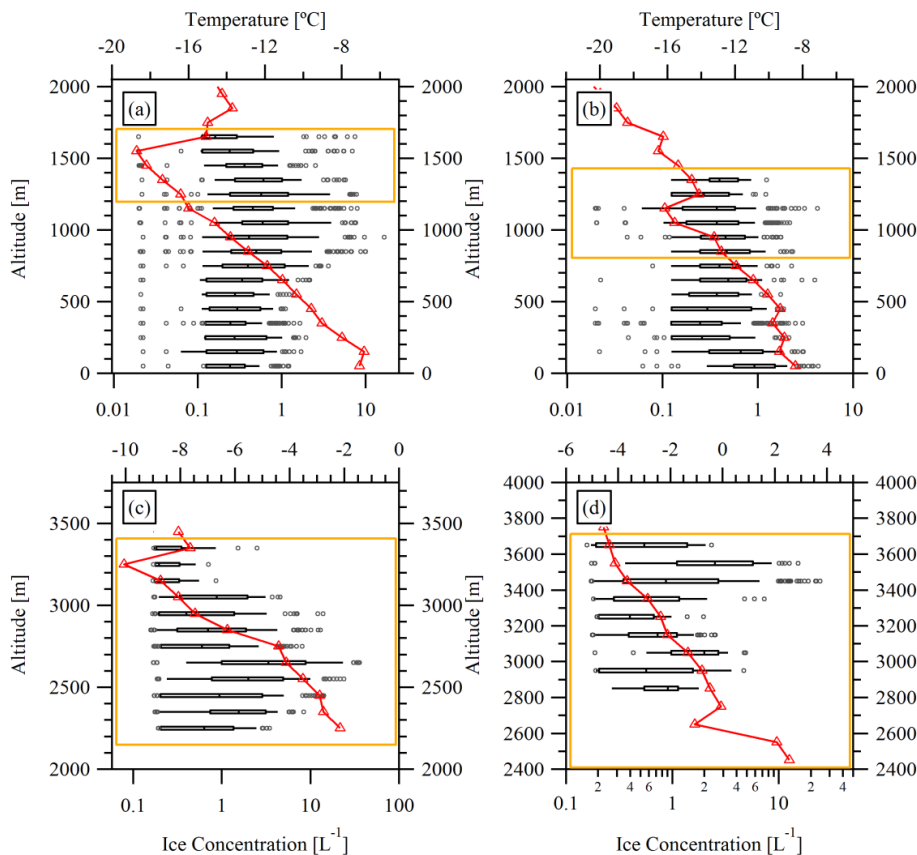


Figure 11. Box and whisker plots with 50th, 25th, 75th percentiles, whiskers to 10 and 90 % and outliers between 95 and 100 % as a function of altitude for ice number concentrations (black) and median temperature (red) (a–d) and altitude averages as in Fig. 10 above). The box in yellow provides an indication of the full extent of cloud layers investigated. (a–d) are for Spring Case 1, Spring Case 2, Summer Case 1 and Summer Case 2 respectively.

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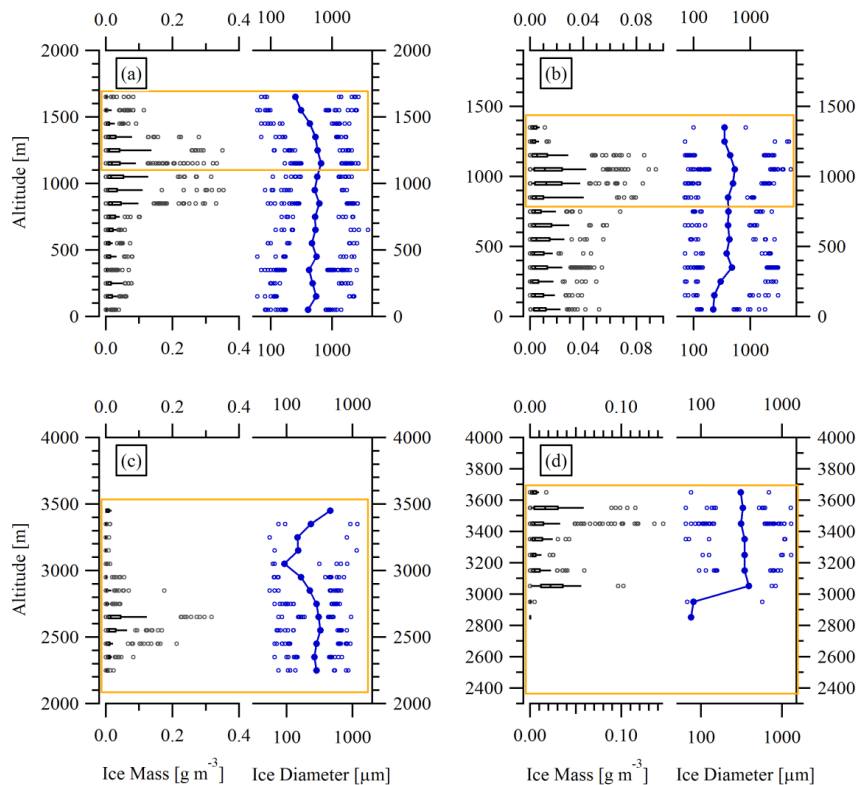


Figure 12. Box and whisker plots with 50th, 25th, 75th percentiles, whiskers to 10 and 90 % and outliers between 95 and 100 % as a function of altitude for ice mass (black) and median ice crystal diameter with outliers between 95 and 100 % (blue). (**a–d** and altitude averages as in Fig. 10 above). The box in yellow provides an indication of the full extent of cloud layers investigated. (**a–d**) are for Spring Case 1, Spring Case 2, Summer Case 1 and Summer Case 2 respectively.