Responses to Reviewer Comments

General – We are grateful to the reviewers for their interest and helpful critiques. We appreciate their time and effort in helping to improve this manuscript. Responses to the review comments are highlighted in yellow.

**Reviewer 1**

Major Comment from Reviewer 1

Overall, I am very pleased with how thoroughly the authors addressed my first comments, and I am ready to support the paper for publication, as long as one major comment is addressed. As an aside, I congratulate the authors in that I have overheard sincere community interest in the ACPD paper from my colleagues.

Major comment based on Specific comment 4b: Given that 62 different clouds were examined, it was not clear to me why a select few were chosen for more detailed analysis. Please provide this information.

Response – There were 62 cloud "samples". The numbers of individual clouds were fewer, since more than one profile was conducted through a cloud layer. We have added a few words to clarify this and the reason for the selections on lines 281-290.

New comment from the reviewer: Ah, I clearly misunderstood that there were 62 independent, separate clouds sampled (to me, the terms "cloud averaged data point" or "Forty-five cloud samples" (e.g., line 650) imply independent samples, taken from separate clouds). Now that I know that the points are not averages from single clouds, but rather averages of penetrations within clouds, I suggest replacing the terminology of "cloud-averaged data points" and "cloud samples" in the paper. Despite the new text added on lines 281-290, these terms may mislead other readers (particularly, but not only, if they are skimming the abstract and conclusions), and there are more accurate terms that can be used. At minimum, I suggest rephrasing to something like "the average of individual cloud penetrations".

However, I strongly believe more analysis may be required here. Individual cloud penetrations within the same cloud (e.g., Fig. 5), should not be treated as independent samples, as they are in the text. Failing to do so will bias the results (e.g., Tables 1 and 2) toward the characteristics of clouds with the most profiles sampled.

As it was not stated how many independent clouds were sampled, particularly for the HA cases, it is hard to fully evaluate the impact of this issue on the results. It would be helpful if the authors could provide as much information as they can about how many independent clouds they actually sampled. Something along the lines of what the authors show with the new supplemental Figure 7 is helpful, but alone it is not sufficient. I also strongly recommend that the authors redo the statistics with independent clouds separately. If the findings are not still the same, further discussion in the text will be needed.

Author response - We have further clarified our approach using the phrasing "averages of individual cloud penetrations", as suggested by the reviewer. We did consider that our approach biased the statistics in the manner the reviewer discusses, but we believe our approach is reasonable in terms of the effects of aerosols on clouds for climate. Cloud coverage is a major factor in terms of cloud influence on climate. If we generate one point from one large cloud and plot it with one point from one small cloud, the "climate-impact" of a regression is biased inappropriately to the small cloud. It is difficult to conduct these measurements in an ideal manner. Our approach was to sample every liquid cloud that was available to us as much as possible, and it was random in so much as we sampled cloud as the opportunity presented itself along our flight path without purposely seeking cloud. Although we are not saying this approach is perfect, in some way it takes into account clouds that are more extensive than

others, and that bias we feel is appropriate from a climate perspective. We have added some discussion to this effect on lines 266-271 of the revised manuscript.

Minor comments from Reviewer 1

• Regarding the following newly added sentence: "Variations in particle chemistry will induce some variance in these results, but because activation diameters are estimated starting with larger particles and moving to smaller sizes, changes in chemistry only offer the possibility of activation of particles still smaller than estimated here." My understanding is that size and hygroscopicity are two opposing constraints on activation, and thus depending on the situation, activation could depend on either. I suggest the authors rephrase this sentence.

Author response – We are not of the opinion that size and hygroscopicity are "opposing constraints", since they tend to act in the same way; nor do we feel that treating them as independent (which some people seem to do) is appropriate. Regardless, our comment is simply about number conservation. Given the nature of the particle number distribution, if we start from the largest size and move towards smaller sizes to estimate an activation diameter, the true activation diameter can only be larger than our estimate if there is a significant error in our number measurements; hopefully, we have demonstrated otherwise. Chemistry can only dictate that particles smaller than our estimate activated. That of course would happen at the expense of some larger particles, and we have added that point of clarification to the sentence you mention.

• Why add in the Koellner reference at all? It is fairly long and not as descriptive as simply saying, as the authors did in their response, that: 1) back trajectories suggested that the air arrived from Canadian forest fires, 2) there was an [~20 ppbv?] increase in CO concentrations compared to other days sampled that month, and 3) the size distribution transitioned to larger sizes, which suggests some BB influence?

Author response – Thank you for the comment, but it is our preference not to make this change.

• Line 83: "Lohmann and Leck (2005) hypothesized the need for highly surface- active particles to explain CCN [activity?] at S less than 0.3%."

Author response – Corrected.

**Reviewer 2 (Dr. Hudson)**

Minor comments from Reviewer 2 (Dr. Hudson)

The authors have satisfactorily answered my comments. As will be shown I have some problems with interpretations or explanations of Figs. 3 and 4. Nonetheless, this manuscript should definitely be published. Field results of such important topics as these need to be made available.

I did detailed editing for the first half of the manuscript. This was intended and resulted mostly in text reductions. In almost every instance maximum numbers of words were used. This was not just the definite article as mentioned in my first review but also prepositions such as of and in. However, it occurred to me that perhaps for some reasons these authors do not want to reduce the length of the manuscript. Thus, since my editing might be in vain I stopped most

editing half way through the manuscript. If this notion is incorrect the authors can follow my examples of the first half to edit the 2nd half accordingly.

Author response – We thank Dr. Hudson for his conscientious editing of the manuscript. As below, we have considered all suggestions, and we have adopted most.

L32. Delete the.
L38. Delete the.
L39. And to with. Change are used to infer to imply.
L40. Delete respectively. The two sets are in order.
L43. Delete the twice.
L44. Delete the. Delete in the. Change changes to variations. Move aerosol in front of variations.
L45. In to within.
L46. From to between. To to and.
L47. Delete the.
L52-3. Change and lower during to than in.
L53. Delete of. Particles singular. Move transport after particle.
L54. Delete the. Delete 2nd of. Move focus after chemistry.
L55. Delete in the. Move Arctic in front of research. Delete the. Add time to spring. Delete period. Delete from. Move transition after summer.
L56. Delete 1st 2 thes. Opportunity plural. Delete in. move changes after chemistry.
L57. Delete last the.
L59. Delete of the. Move Arctic on front of warming.
L72. Change aerosols to particles.
L73. Delete sources. Delete for. Move of after nuclei.
L74. Drop s of towards.
L76. Delete much. Change sometimes to often.
Author response – L32.–L76.: most of the changes have been made.

L82. Explain surface-active. Does this mean hygroscopic?

Author response – Components that lower the surface tension. Details are in the reference.

L83. Delete 1st the.
L84-85. Move summertime and Arctic in front of environment. Delete of the.
L85. Particles singular and move in front of concentrations. Delete of. Above to larger than.
L87. Delete nm.
L89. Delete much.
L91. Period after parentheses. And to Moreover.
L92. At to to. ing to e.
L96. Delete made.
L97. Delete that.
L98. Delete the. Delete in the. Move CDNC after model.
L99. Delete will. Above to for.
L100. Insert greater than before 10. Delete in. move increases after CCN. Delete of the. Change phere to pheric. Move cooling after atmospheric. Change the to This.
L101. Move threshold after concentration.

L101-2. Change and it is noted that the value of 10 cm-3 to although this. Insert a after not. Insert limit after universal.
L103. Delete 2nd the. Add time to summer and move in front of microphysics. Clouds singular and move in front of microphysics.
L107. Insert Yum and Hudson, 2001 and Wylie and Hudson, 2002.
L108. Delete 1st the. Add time to spring.
L109. Aerosols to particles.
L111. Delete of. Aerosols singular and move in front of observations.
L112. Delete during. Add time to summer and move in front of Arctic.
L113. Delete of. Aerosols singular and move in front of measurements. Add time to summer. Move Arctic in front of clouds.
L114. Period after forcing. Insert They to begin next sentence. ing to ed.
L115. Delete about. Move forcing from these plumes after maximum. Half is approximate.
L117. Among to between. Insert and after coupled. Move uncoupled in front of to. delete versus those. Period after surface. Change but to They. Delete of the. Move observations after microphysics. Insert their after and.
L119. Characterization plural. Was to were.
L119-20. Move June in front of low. Delete in.
L121. Delete the. Delete in the. Tops singular. Move cloud top in front of CDNC.
L121-2. Remove quotes. Move aerosols to end of sentence.
L122. Delete the twice. Bases singular. Insert cloud in front of base.
L123. Delete does. Add s to influence. Delete the.
L124. Delete the twice. Delete in. add time to summer. Cloud plural and move after Arctic.
L127. Aerosols and clouds singular and move in front of observations. Remove of.
L128. The to this.
L153. Delete are.
L157. The to these.
L162. The to these.
L165. Add s to detect. Move detects particles right after that. Uses to using. Delete of. Move scattering after light. Delete to.
L167. Delete a reduced pressure of.
L168. Last The to This.
L169. Delete of. Move measurement in front of stability.
L170. Delete the. Delete of. Particles singular. Move hygroscopicity to end of sentence.
L171. Using to with.
L173. Period after parentheses. Delete and.
L176. Dimensions to dimensional. Move two dimensional in front of Cloud. Delete in. sized from about to between. To to and.
L177. Using to with. Delete For. Delete present. The to this.
L177-8. Move this study to the end of this sentence. insert from after phase.
L179. Period after parentheses. Delete and.
L182. Delete The.
L183. Use CO.
L184. Delete at. Move 150nm in front of excitation. The to This.
L191. Measured to done.
L192. The to This.
L198. Semicolon to period. Insert However before the.
L199. Insert or below after in.

L200. Are to was.
L201. Are to was. Is to was.
L205. Delete of.
L206. Delete the.
L207. Delete of. Particles singular. Move transmission after particle. Approximately to near.
L211. Delete at.
L211-2. Move exhaust tube in front of flow.
L212. Move flow after tube. Delete the four times. Delete of twice. Delete at. Delete allowed. Delete last flow. Delete ly.
L213. Move flow after intake. Delete at the. Period after TAS. Delete and.
L216. Delete 1st the. Delete of the. Move aircraft in front of forward.
L217. Change lowered to reduced. 1st the to this.
L220. Delete to. Move Analysis in front of Approach.
L223. Beginning to between. Delete ending July. The to These.
L224. Delete relatively. Delete the. Relatively and distinct are opposites.
L225. Delete The.
L226. Change calm to light. Change varying to variable. Insert the before south. Delete to north.
L233. Period after parentheses. Then insert This was. Delete in part. Possibly is enough of a hedge.
L237. The to this.
L239. Delete of. Flying to legs.
L240. Above to altitude.
L241. Delete the surface. Surface here must me sea level.
L245. Delete surrounding. Period after surfaces. Change as well as to Furthermore. Delete because. Insert was after 2. Insert by after marked.
L246. Put Fig. 2 in front of panel. Delete in.
L247. Flight plural. Delete plans were. Change towards sampling to on.
L251. Change greater to larger. Delete for.
L252. Delete The.
L253. Over to between. Dash to and. Delete data, which are.
L253-4. Move Fig. S3 in front of example.
L254. Change shown in to and. Particle singular and move in front of number. Delete of. Delete 1st nm.
L258. In to within. Move study in front of area. Delete of. Move Within the study area to beginning of sentence. Delete when they. Change ideally to mostly. Ascending to ascents.
L259. Or to and. Descending to descents. Delete through them. Delete the. Base plural. Delete of. Clouds singular. Move bases after cloud.
L260-1. Move only liquid phase clouds after μm.
L262. With the caveat to except.
L264. May to might.
L265. Period to whereas.
L266. Move July 7 in front of stratocumulus. Delete sampled on. But to though.
L270. Insert and when after where. Change was clear and achievable to could be observed. Clear is a poor word choice to describe cloud base. Change semicolon to period.
L272. In to within. Comma to and.
L272-3. Move in flight in front of indistinguishable.
L273. Period after parentheses. And to Thus,. The to such. Cloud plural.

L278. Are to were.
L281-2 and elsewhere. Points is not the best word choice unless you are referring to elements of a figure. Sections or segments are alternatives. Or just refer to data without another word.
L282. Is to was. The to These.
L288. Delete the.
L289. Move valid in front of in-cloud. Delete considered. Delete inside of cloud.
L290. Thermodynamic plural. Delete measurements, it is used twice in this sentence. Delete the. In to within.
L293. On to upon. Insert Hudson and Frisbie [1991] and Hallett and Christensen [1984].
L303. Delete on July 7 sampled. Delete 1st the.
L304. Change and the to while.
L305. Period after altitude. Insert These before features. Change common to the to characterize. Move formation after cloud. Delete of. Change and indicating to so.
L306. Delete cloud. Delete in air rising. Insert below after from.
L308. Delete 1st the.
L309. Delete the. In to Within.
L313. Delete 3rd the. Insert Hudson [1993] at end of sentence. Change The to Thus.
L315. Delete 2nd the.
L317. Insert corresponding before N5.
L318. Delete the. Move closure after concentration (singular). Delete of.
L320. Change down to about to as small as.
L320-1. Move based on maximum CDNC to beginning of sentence.
L321. Delete the. Delete of. Move particles after sulphate.
L322. Change the to that. Delete of the. Clouds singular and move in front of bases.
L324. At to of.
L325. And to while.
L326. Change but there are to except that. Insert is before more. Insert broken after more.
L326-7. Move the July 17 profile right after except that.
L327. Delete what is left of this line.
L328. Delete adiabatic lifting. Insert lower LWC before intervals. Delete with decreasing LWC. Change associated with to due to.
L329. Change the to cloud. Delete of the stratocumulus. Profile plural. Last the to this.
L330. Move LWC in front of peak. Delete in the. Change below to from.
L331. Move CO in front of increase. Delete in. insert at before about.
L332. Move the in front of erosion. Change was to went. In to into. Change case to cloud or clouds.

Author response – L83.-L332: most of the changes have been made.

L333. What aerosol increase above cloud? Aerosol decreases at many levels. Delete the.
Author response – The N100 and CCN, shown in Fig. 3c are higher above cloud than below cloud. We have re-written it to say “indicating that the higher concentrations of N50-100, N100 and CCN above cloud relative to below cloud did not enhance the CDNC”

L334. This is true at a greater altitude range.

Author response – “about” changed to “at least”

L335. This is not shown in the figure. Delete last the.

Author response – N50 is derived from the sum of N50-100 and N100, both of which are shown in the figure. Last “the” deleted.

L336. Larger to higher. Of to between.
L340. 1st the to a.
L342. Last the to this.
L343. Period after study. Delete and. Insert that before BB.
L347. Add ed to reach.
L348. Period after layer. Insert This is.
L349. Reduced to lower. Increased to higher.
L350. Semicolon to comma. Insert which is. Increase to higher. Delete in. insert concentrations above than after aerosol. Delete between. The to this. Delete and.

Author response - L336.-L350.: most of the changes have been made.

L351. Delete above the layer. N50 is apparently not shown in the graph! I do not see these numbers.

Author response – They are derived from N50-100 added to N100.

L354. Delete sized. Delete 1st nm. Insert diameter after nm.
L355. Insert comma which after CDNC. Down to about to as small as.
L356. Reduced to lower.
L357. Insert below cloud after of. And to with. Above cloud CCN does not show this.
L358. Delete case of a. delete in.
L358-9. Move LWC in front of variations.

Author response - L354.-L358-359.: most of the changes have been made.

L359. Delete the. Suggests to suggest. Looks higher than 49.

Author response – Changes made. Because it is narrower, the lower CDNC at the boundaries have a larger influence on the average.

L361. The to These.

Author response – Changed.

L363. I do not see this in the figure.

Author response – N50 is derived from the sum of N50-100 and N100.

L365. Pre to below.

Author response – “pre” is correct.

L365-7. It is problematic to get valid aerosol measurements of any kind in the narrow layer between these clouds.
L366. 44?
L367. 52?, seems higher than 34. Seem higher than 66. Seems higher than 35.
L369. For the lower cloud layer yes, but not so sure about the other two.

Author response – Estimating the aerosol for such layers is problematic, and we make a statement to this effect regarding this case. Again the N50, which is quoted in the text is derived from the sum of N50-100 and N100, which we think explains the confusion over the N50 number concentrations.

L372. Delete 2nd July.
L373. Delete and July.

Author response – Second change made.

L377. Looks much higher in Fig. 4a.

Author response – The mean is influenced by many low points.

L378. Apparently below cloud is to the right? 16:45 to 17:09 and beyond? This needs to be stated. N100 appears to be 2. 0.6% ammonium sulfate is 40 nm diameter. 100 µm ammonium sulfate is 0.1%.

Author response – The cloud (CDNC represented by blue points) is to the left. Aerosol is to the right, and the wind direction, as indicated, is right to left. The CCN at 0.6% are higher than the CDNC except at 130 m where the CDNC reaches 12/cc.

L385. Explain these 7 samples. This is not obvious from the figure.

Author response – Explanation added by re-writing the segment as "Seven samples were identified over the period 16:06-16:29 based on the LWC above 0.01 g m-3. The CDNC are overall higher than on July 5 with sample averages ranging from 4 cm-3 to 13 cm-3; the one-second CDNC are as high as 34 cm-3 and the mean VMD (not shown) range from 19.6 um to 22.8 um."

L389. Where is N100 this low? To the right and left sides of Fig. 4b N100 is 50 or more! Lower values are seen within cloud. All measurements where altitude is shown seem to be within cloud and thus invalid for aerosol measurements.

Author response – As stated in the text, "In the air nearly free of cloud and below 120 m".

L390-1. Delete due to instrument problems.

Author response – deleted.

L391-2. These are not the numbers that appear on Fig. 4b.

Author response – The results have been checked several times. They are correct based on our defined approach.

L400. Delete from. East to easterly. Delete to west.
L404. The to This.
L414. The to These.
L418. The to these.
L427. The to these.
L467. The to these.
L533. Insert S after %.
L572. Reach above to exceed.
L578. Insert that before 1st the.

Author response – L400.-L578.: appropriate changes have been made.

Hallett, J., and L. Christensen, The splashing and penetration of raindrops into water, J. Rech. Atmos. 18, 226-242, 1984.
Hudson, J.G., 1993: Cloud condensation nuclei near marine cumulus. J. of Geophys. Res., 98, 2693-2702.
Hudson, J.G. and P.R. Frisbie, 1991: Cloud condensation nuclei near marine stratus. J. of Geophys. Res., 96, D11, 20,795-20,808.
Wylie, D., and J.G. Hudson, 2002: Effects of long range transport and clouds on cloud condensation nuclei in the Springtime Arctic. J. Geophys. Res., 107(D16), 4318, doi:10.1029/2001JD000759
Yum, S.S., and J.G. Hudson, 2001: Vertical distributions of cloud condensation nuclei spectra over the springtime Arctic Ocean. J. Geophys. Res., 106, 15045-15052.

Author response – All above references are included.

# Effects of 20-100 nanometre particles on liquid clouds in the clean summertime Arctic

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**Abstract.** Observations addressing effects of aerosol particles on summertime Arctic clouds are limited. An airborne study, carried out during July, 2014 from Resolute Bay, Nunavut, Canada, as part of the Canadian NETCARE project, provides a comprehensive in-situ look into some effects of aerosol particles on liquid clouds in the clean environment of the Arctic summer. Median cloud droplet number concentrations (CDNC) from 62 cloud samples are 10 $cm^{-3}$ for low-altitude cloud (clouds topped below 200 m) and 101 $cm^{-3}$ for higher-altitude cloud (clouds based above 200 m). The lower activation size of aerosol particles is ≤ 50 nm diameter in about 40% of the cases. Particles as small as 20 nm activated in the higher-altitude clouds consistent with higher supersaturations (S) for those clouds inferred from comparison of the CDNC with cloud condensation nucleus (CCN) measurements. Over 60% of the low-altitude cloud samples fall into the CCN-limited regime of Mauritsen et al. (ACP, 2011) within which increases in CDNC may increase liquid water and warm the surface. These first observations of that CCN-limited regime indicate a positive association of the liquid water contents (LWC) and CDNC, but no association of either the CDNC or LWC with aerosol variations. Above the Mauritsen limit, where aerosol indirect cooling may result, changes in particles with diameters from 20 nm to 100 nm exert a relatively strong influence on the CDNC. Within this exceedingly clean environment, as defined by low CO and low concentrations of larger particles, the background CDNC are estimated to range between 16 $cm^{-3}$ and 160 $cm^{-3}$, where higher values are due to activation of particles ≤ 50 nm that likely derive from natural sources. These observations offer the first wide-ranging reference for the aerosol cloud albedo effect in the summertime Arctic.

## 1. Introduction

Mass concentrations of the atmospheric aerosol in the Arctic are higher during winter than in summer due to differences in transport of anthropogenic particles and wet scavenging (e.g. Barrie, 1986; Stohl, 2006). Atmospheric chemistry and aerosol-cloud Arctic research has largely focussed on the springtime. The winter to summer transition offers the opportunity to examine changes in chemistry as the sun rises over the polluted polar atmosphere (e.g. Barrie et al., 1988) and to study impacts of anthropogenic aerosol on the Arctic solar radiation balance (e.g. Law and Stohl, 2007; Quinn et al., 2008). Greater-than-expected warming of the Arctic (e.g. Christensen et al., 2013) and rapidly diminishing Arctic sea ice extent (e.g. Maslanik et al., 2011) have drawn considerable attention to the role of anthropogenic and biomass burning particles as warming agents for the Arctic (e.g. Law and Stohl, 2007; Quinn et al., 2008; Shindell et al., 2008; Brock et al., 2011; Jacob et al., 2010; UNEP, 2011; Stohl et al., 2013). Recent evidence indicates that the net impact of aerosol particles on the Arctic over the past century has been one of cooling rather than warming (Najafi et al., 2015).

Low-level liquid water clouds are frequent in the sunlit Arctic summer (e.g. Intrieri et al., 2001), and these clouds can have a net cooling effect (e.g. Brenner et al., 2001; Garret et al., 2004; Lubin and Vogelmann, 2010; Zhao and Garrett, 2015; Zamora et al., 2015;). Knowledge of the influence of the atmospheric aerosol on climatic aspects of these clouds is complicated by the relatively large potential differences in the albedo of the underlying surface (e.g. Herman, 1977; Lubin and Vogelmann, 2010) and the fact that the Arctic is relatively free of anthropogenic influence in summer, which means that particles from natural sources can be the most significant nuclei for cloud droplets. Those sources shift the number distribution toward

70 particles smaller than 100 nm (e.g. Heintzenberg and Leck, 1994; Ström et al., 2003;
71 Heintzenberg et al., 2006; Engvall et al., 2008; Tunved et al., 2013; Leaitch et al., 2013;
72 Heintzenberg et al., 2015). Particles smaller than 100 nm are often dismissed as being too small
73 to nucleate cloud droplets due to the assumption that the cooling mechanisms are too slow to
74 generate the supersaturation (S) required to activate the smaller particles in Arctic liquid clouds
75 (e.g. Garret et al., 2004; Lubin and Vogelmann, 2010; Browse et al., 2014; Zhao and Garrett,
76 2015). That assumption may lead to reduced estimates from natural feedbacks to climate and
77 increased estimates of aerosol indirect forcing from anthropogenic sources. Lohmann and Leck
78 (2005) hypothesized the need for highly surface-active particles to explain CCN activity at S less
79 than 0.3%. However, cloud S is also strongly constrained by the concentrations of particles
80 larger than 100 nm, and in the clean summertime Arctic environment with relatively low
81 concentrations of particles larger than 100 nm, there is some evidence that higher S may be
82 achieved and smaller particles activated (e.g. Hudson et al., 2010; Korhonen et al., 2010; Leaitch
83 et al., 2013). Further, the suggestion that the minima between 50 and 100 nm in Arctic particle
84 size distributions results from cloud processing implies consistent activation sizes less than 100
85 nm (Heintzenberg et al., 2015). The effect of the background aerosol on liquid clouds has been
86 identified as one of the most important factors for reducing uncertainty in the aerosol cloud
87 albedo effect (Carslaw et al., 2013). Moreover, the effectiveness of particles smaller than 100 nm
88 for cloud droplet nucleation is a large factor in that uncertainty.
89 Effects of pollution on clouds may also lead to warming, but a reference to clean clouds
90 is still required (e.g. Garrett et al., 2009). Mauritsen et al. (2011) modeled cloud radiative
91 forcing for low clouds using CCN number concentrations derived from shipborne observations
92 over the Arctic Ocean (Tjernström et al., 2004; Tjernström et al., 2014). They found the impact

93 from changes in CCN for ultra-low values (< 10 $cm^{-3}$), where CCN concentrations are equivalent
94 to model CDNC, results in a net warming due to associated longwave changes, whereas for
95 concentrations greater than 10 $cm^{-3}$ CCN increases are estimated to produce a net atmospheric
96 cooling. This CCN concentration threshold is referred to here as the "Mauritsen limit", although
97 this value of 10 $cm^{-3}$ is not a universal limit (Mauritsen et al., 2011). In the clean summertime
98 Arctic, knowledge of the natural aerosol and its influence on cloud microphysics is critical to the
99 assessment of aerosol effects on Arctic climate.

100 Past studies of Arctic aerosols and clouds have emphasized the areas of the Beaufort and
101 Chukchi Seas (e.g. Hobbs and Rango, 1998; Curry et al., 2001 and references therein; Lohmann
102 et al., 2001; Yum and Hudson, 2001; Peng et al., 2002; Wylie and Hudson, 2002; Earle et al.,
103 2011; Lance et al., 2011; Jouan et al., 2014; Klingebiel et al., 2014). Most of those studies have
104 focused on springtime when the aerosol can be influenced by anthropogenic or biomass burning
105 sources. As well, there has been considerable interest in mixed-phase clouds in the lower Arctic
106 troposphere (e.g. Shupe et al., 2004; Sandvik et al., 2007; Morrison et al., 2012), but a notable
107 lack of in-situ aerosol observations in combination with liquid water clouds over the summertime
108 Arctic. Among the studies that have considered in-situ aerosol measurements and summertime
109 Arctic clouds, Zamora et al. (2015) examined the efficiency of biomass burning (BB) plumes on
110 indirect forcing. They estimated half of the possible maximum forcing from these plumes,
111 mostly due to the reduction in cloud-base S by higher concentrations of larger particles that
112 control water uptake. Shupe et al. (2013) discussed some differences between clouds coupled
113 and uncoupled to the surface. They did not conduct in-situ cloud microphysics observations, thei
114 and vertical aerosol characterizations were constrained to particles >300 nm. Hobbs and Rango
115 (1998) found that droplets in June low clouds over the Beaufort Sea occasionally contained drops

as large as 35 μm diameter. They also found that cloud-top CDNC correlated significantly with cloud base "aerosols". They suggested that cloud-top entrainment did not control CDNC, although there may be times when entrainment influences Arctic CDNC (e.g. Klingebiel et al., 2014).

Motivated by limited knowledge of aerosol effects on summertime Arctic clouds and particle activation details, the Canadian Network on Climate and Aerosols: Addressing Key Uncertainties in Remote Canadian Environments (NETCARE - http://www.netcare-project.ca/), conducted airborne aerosol and cloud observations during July, 2014 in the area around Resolute Bay, Nunavut, Canada. The observations from this study are used here to characterize CDNC, LWC, and the volume-weighted mean droplet diameter (VMD). Further, aerosol particle size distributions (5 nm and larger; CCNC(0.6%)) from outside of clouds are compared with droplet number concentrations from inside of clouds. Specifically, in the indicated sections, the following questions are addressed.

1) Given the scarcity of data, what are the characteristics of clouds in the summertime Arctic, and do clouds near the surface have characteristics different from those aloft? (Sect. 3.2)
2) What are the sizes of particles that act as nuclei for cloud droplets? This will allow a closer connection between aerosol processes, particle sizes and climate effects? (Sect. 3.3)
3) What is the relationship between droplet size and droplet number? In particular, what is the aerosol influence on cloud below the Mauritsen-limit, and is it possible to assess a background influence of the aerosol on clouds in the Arctic summer? (Sect. 3.4)

## 2. Methodologies

The instrument platform was the Alfred Wegener Institute (AWI) Polar 6 aircraft, a DC-3 aircraft converted to a Basler BT-67 (see Herber, A., Dethloff, K., Haas, C., Steinhage, D., Strapp, J. W., Bottenheim, J., McElroy, T. and Yamanouchi, T.; POLAR 5 - a new research aircraft for improved access to the Arctic, ISAR-1, Drastic Change under the Global Warming, Extended Abstract, pp. 54-57, 2008).

## 2.1 Instrumentation

The following measurements are relevant to this discussion:

a) Particle number concentrations >5 nm diameter were measured with a TSI 3787 water-based ultrafine condensation particle counter (UCPC), sampling at a flow rate of 0.6 L $min^{-1}$. Hereafter, these measurements are referred to as N5.

b) Aerosol particle size distributions from 20 nm to 100 nm (45 s up scans and 15 s down scans) were measured using a Brechtel Manufacturing Incorporated (BMI) Scanning Mobility System (SMS) coupled with a TSI 3010 Condensation Particle Counter (CPC). The sheath and sample flows were set to 6 L $min^{-1}$ and 1 L $min^{-1}$. BMI software was used to process these distributions.

c) Aerosol particle size distributions from 70 nm to 1 μm were measured using a Droplet Measurement Technology (DMT) Ultra High Sensitivity Aerosol Spectrometer (UHSAS) that detect particles using scattering of 1054 nm laser light (e.g. Cai et al., 2008).

d) CCNC(0.6%) were measured using a DMT CCN Model 100 counter operating behind a DMT low pressure inlet at approximately 650 hPa. For the nominal water S of 1%, the effective S at 650 hPa was found to be 0.6% as discussed below. This S was held

constant throughout the study for greater measurement stability, improved response, and to examine the hygroscopicity of smaller particles.

e) Droplet size distributions from 2-45 μm were measured with a Particle Measuring Systems (PMS) FSSP-100. This FSSP-100 had been modified with new tips to reduce shattering artifacts (Korolev et al., 2011). It was mounted in a canister under the port-side wing. The CDNC, VMD and LWC are calculated from the measured droplet distributions.

f) Two-dimensional cloud particle images from about 50 μm to 800 μm were measured using a PMS 2DC Grey-scale probe. These observations are used here only to ensure the absence of the ice phase. This 2DC-Grey was also modified with new tips to reduce shattering artifacts (Korolev et al., 2011). It was mounted in a canister beside the FSSP-100.

g) Carbon monoxide (CO) is used here as a relative indicator of aerosol influenced by pollution sources and as a potential tracer for aerosol particles entering cloud. CO was measured with an Aerolaser ultra-fast carbon monoxide monitor model AL 5002 based on VUV-fluorimetry, employing the excitation of CO at 150 nm. This instrument was modified such that in-situ calibrations could be conducted in flight.

Details of the instrument calibration and evaluations are given in the Supplement (S1).

## 2.1 State parameters and Winds

State parameters and meteorological measurements were made with an AIMMS-20, manufactured by Aventech Research Inc. This instrument consists of three modules: 1) an Air Data Probe that measures the three-dimensional aircraft-relative flow vector (true air speed, angle-of-attack, and sideslip), temperature and relative humidity, and includes a three-axis accelerometer pack for turbulence measurement; 2) an Inertial Measurement Unit that consists of three gyros and three accelerometers providing the aircraft angular rate and acceleration; 3) a Global Positioning System for aircraft 3D position and inertial velocity. Horizontal and vertical wind speeds were measured with accuracies of 0.50 and 0.75 m/s, respectively. However, the vertical resolution was insufficient to measure gusts in the sampled clouds. The accuracy and resolution for temperature measurement are 0.30 and 0.01 C. The accuracy and resolution for relative humidity measurement are 2.0 and 0.1 %. The sampling frequency is 1 Hz.

## 2.2 Inlets

Aerosol particles were sampled through a shrouded inlet diffuser (diameter 0.35 cm at intake point), which is the same inlet discussed by Leaitch et al. (2010). For the airspeeds during this study, particle transmission by the inlet is near unity for particles from 20 nm to <1 μm. The intake was connected inside the cabin to a 1.9 cm OD stainless steel manifold off of which sample lines were drawn to the various instrument racks using angled inserts. Total flow at the intake point was approximately isokinetic at 55 L $min^{-1}$ based on the sum of flows drawn by the instrumentation (35 L $min^{-1}$) and the measured manifold exhaust flow. The manifold exhaust flowed freely into the back of the cabin such that the intake flow varied with aircraft TAS and the manifold was not significantly over pressured.

CO was sampled through a separate inlet consisting of a 0.40 cm OD Teflon tube using aircraft forward motion to push air into the line in combination with a rear-facing 0.95 cm OD Teflon exhaust line that reduced the line pressure. The continuously measured sample flow was approximately 12 L $min^{-1}$.

## 2.3 Data Analysis Approach

Eleven research flights were conducted from Resolute Bay, Nunavut (74°40'48"N 94°52'12"W) from July 4, 2014 to July 21, 2014, inclusive. These measurements were associated with two distinct weather regimes. During period 1 (July 4-12), weather conditions around Resolute Bay were affected by an upper low (Supplement Fig. S4). The wind speeds at 500 hPa were mostly light and variable. The surface (1000 hPa) was dominated by weak high-pressure with generally clear skies, light winds, and occasional scattered to broken stratocumulus. Low-cloud or fog was at times present in association with open water, and the air was relatively clean as discussed below. There was a transition period from July 13-16 when flights were not possible due to fog at Resolute Bay. During period 2 (July 17-21), the area came under the influence of a deep low pressure system to the south (Supplement Fig. S5) that brought more wind and higher cloud. The air was not as clean as during period 1, based on the measured aerosol mass and CO concentrations (see Table 1). This was possibly due to transport of BB aerosol from the Northwest Territories; further discussion in Section 2.3.1. Based on the bulk Richardson number and data from radiosondes, Aliabadi et al. (2015) estimated boundary-layer heights at 254 m (±155 m) across the study.

A summary of all flight tracks is shown in Fig. 1. Flights mostly consisted of vertical profiles and low level transits over ice, water and melt ponds that contributed to the formation of low cloud, where low cloud is defined here as cloud tops below 200 m-msl. Higher level cloud was also sampled during the profiles and transits. The polynyas that were sampled over are shown in the top center of each panel of Fig. 2. Cloud was sampled on eight of the 11 flights, more frequently during period 1 because of overall better visual contrast between clouds and surfaces. Furthermore, period 2 was marked by the presence of the Canadian Coast Guard Ship Amundsen in Lancaster Sound (bottom center of each Fig. 2 panel) when the flights were focused on sampling of the ship's plume (e.g. Aliabadi et al., 2016).

All aerosol number concentrations are given in terms of standard atmospheric pressure and temperature (STP: 1 atm and 15$^{o}$C). The CDNC are also referenced to STP where comparisons are made with the aerosol number concentrations. Number concentrations of particles larger than 100 nm (N100) are taken from the UHSAS. All data, except the SMS, are 1 second averages that represent a sampling path length of 60–80 m. Size distributions between 20 and 100 nm are from the SMS are 1-minute averages. Except for the Fig. S3 example, all particle number concentrations smaller than 100 nm are from the SMS. Nx-100 refers to the number concentration within the interval "x-100" where x ranges between 20 and 90. Values of Nx with $x < 100$ are derived from the sum of Nx-100 (SMS) + N100 (UHSAS).

Clouds were sampled during a flight whenever possible, mostly by ascending or descending through them. It was not possible to sample below the low-altitude cloud bases. Most clouds were liquid phase, based on the 2DC-Grey images of cloud particles >50 μm, and only liquid phase clouds are discussed here. In addition, none of the liquid clouds exhibited detectable precipitation, except that droplets in a couple of the lowest altitude clouds were very

low in number and relatively large in size (30-40 μm); considering the settling speeds of such droplets, they may be viewed as precipitating. The higher-altitude clouds were either stratus or stratocumulus, whereas the low-level clouds were fog or stratus. Although still light, turbulence appeared to be the greatest in the July 7 stratocumulus. Cloud droplet sizes are represented by the volume-weighted mean diameter (VMD), which has the property that the VMD can be used with CDNC to calculate LWC.

The pre-cloud aerosol for the HA clouds is mostly derived from averages of values collected within about 50 m of cloud base when a cloud base was visible and achievable. In some cases, as discussed in Section 2.3.1 and 2.3.2, the pre-cloud aerosol concentrations include contributions from above cloud (July 19) or are from similar or lower altitudes in the clear air upwind of the cloud. For the aerosol measurements made with the 1-minute averaged number concentrations from the SMS, values from further below-cloud are necessary in some cases. These values are however consistent with the 1-second aerosol measurements closer to cloud base.

Every possible liquid cloud was sampled along a flight path, and some cloud layers were sampled more than once. That will bias the sample numbers to clouds of greater spatial extent. However, that bias is appropriate from a climate perspective since cloud extent is a major factor for the impacts of clouds on climate. A total of 62 liquid water cloud samples, or averages of individual cloud penetrations, were averaged with the constraint that the mean LWC is > 0.01 g $m^{-3}$. The samples are integrations over periods ranging from 11 to 1000 seconds with a median sample time of 65 seconds that is equivalent to a horizontal path length of about four kilometers. In sections 2.3.1 and 2.3.2, a range of detailed examples are used to show how the aerosol observations relate to the cloud observations for the higher-altitude (HA) cloud (clouds based

above 200 m) and low-altitude (LA) cloud (clouds topped below 200 m), and to 1) demonstrate how the pre-cloud aerosol concentrations were assessed for the 62 samples and 2) note where effects of entrainment may be a factor and how multiple cloud layers are considered. At 200 m or below, the LA clouds were in the boundary layer, in flight indistinguishable from the surface (i.e. some were possibly fog). Thus, sampling below such clouds was not possible due to proximity to the surface. Besides cloud microphysics, the only in-cloud measurements considered valid are the CO and thermodynamics. For completeness, the aerosol measurements within cloud are included in the plots associated with sections 2.3.1 and 2.3.2, but such measurements, including the CCN, are unreliable due to issues of drying and partial drying associated with the inlet and a particular instrument as well as droplet shattering on the inlet (e.g. Hudson and Frisbie [1991] and Hallett and Christensen [1984]). The in-cloud aerosol measurements are not part of the subsequent analysis.

### 2.3.1 Higher Altitude (HA) Cloud Examples

Four examples of profiles through HA clouds are shown in Fig. 3. There are two panels for each profile: the left-hand panel shows CO, CDNC and particle number concentrations (N5, Nx-100, N100, CCNC(0.6%)); the right-hand panel shows temperature, equivalent potential temperature ($\theta_e$), LWC and VMD. The temperatures, $\theta_e$ and VMD are scaled as indicated.

July 7 Case (Fig. 3 a, b): One of several similar profiles through a stratocumulus layer on during the transits to and from the polynyas north of Resolute Bay. The CDNC (at STP) are relatively constant with altitude while LWC and VMD both increase steadily with altitude. These features characterize cloud formation by lifting of air, and they indicate the cloud droplets were

298 nucleated on particles from below cloud base. The cloud top is relatively sharply capped by a
299 temperature inversion of about 2$^{o}$C at 2350 m, and the particle profiles along with $\theta_e$ and CO are
300 relatively constant below cloud base. There is no indication that entrainment, based on the LWC
301 profile, does anything other than reduce the CDNC. Within cloud, the number concentrations of
302 larger particles (N100) is reduced due to nucleation scavenging; although such particles are not
303 completely eliminated as smaller droplets can enter the inlet and dry in the sampling lines.
304 Smaller particles can be artificially increased in cloud due to the shattering of larger droplets on
305 the aerosol intake (e.g. Hudson, 1993), as indicated by the increase in the corresponding N5
306 higher in cloud. Thus, in-cloud aerosol measurements are shown here only for completeness, but
307 they are not used in the subsequent analysis. CDNC range up to 265 cm$^{-3}$ and the mean value is
308 199 cm$^{-3}$. Below cloud base, the N5, N20-100, N30-100, N50-100, N100 and CCNC(0.6%) are
309 approximately 235 cm$^{-3}$, 167 cm$^{-3}$, 145 cm$^{-3}$, 94 cm$^{-3}$, 67 cm$^{-3}$ and 117 cm$^{-3}$, respectively. The
310 below-cloud N20 of 234 cm$^{-3}$ approximately equals the N5 offering confidence in terms of
311 number concentration closure. The N30 (N30-100 + N100) compare most closely with the mean
312 CDNC leading to the conclusion that on average cloud droplets nucleated on particles down to
313 about 30 nm. Based on the maximum CDNC, it is possible that particles as small as 20 nm
314 contributed to the CDNC in this cloud; for 20 nm particles of ammonium sulphate to activate,
315 Köhler equilibrium theory indicates that S in the cloud bases would have had to reach above
316 1.5%.

317 July 17 Case (Fig. 3 c, d): The maximum and mean CDNC (STP), of about 75 cm$^{-3}$ and
318 55 cm$^{-3}$, respectively, are lower while the VMD peak of 20 μm is higher compared with the July
319 7 profile. The LWC are generally similar between July7 and 17, except that are more breaks in
320 the July 17 profile. Many of those breaks are due to the aircraft passing through the edges of

cloud during this profile. The inversion topping the cloud is weaker and the LWC peak occurs further from cloud top in the July 17 case versus the July 7 case. That LWC feature in combination with the general CO increase, beginning at about 660 m, suggests that the erosion of cloud top by entrainment went deeper into the July 17 cloud. Above 660m, the CDNC also decrease, suggesting the higher concentrations of N50-100, N100 and CCN above cloud relative to below cloud did not enhance the CDNC. Continuity from at least 100 m below cloud base is indicated by the CO and $\theta_e$ profiles, and the N50 approximates the mean CDNC and possibly maximum CDNC. The CCNC(0.6%) are 30-40 $cm^{-3}$ below cloud, indicating a S higher than 0.6%. The comparison between the July 7 and 17 cases is a specific example of the potential importance of smaller particles for the cloud albedo effect.

July 19 Case (Fig. 3 e, f): The July 19 profile includes two cloud layers, one from 1200-1400 m and a second from 1400-1500 m. The layer separation appears in the CO concentrations, which are approximately uniform through the lower layer and increasing in the upper layer. The CO levels of 100+ ppbv in this case are among the highest observed during this study. Transport patterns suggest that BB contributed to this aerosol (Köllner et al., Pollution in the summertime Canadian High Arctic observed during NETCARE 2014: Investigation of origin and composition, in Geophysical Research Abstracts, 17, EGU2015-5951, European Geophysical Union General Assembly 2015, Vienna, Austria, 2015). The mean CDNC (STP) in the lower and upper layers are 239 $cm^{-3}$ and 276 $cm^{-3}$ respectively. The VMD reached 15 μm in the lower layer. The VMD are overall smaller and decrease with altitude in the upper layer, consistent with the lower LWC and higher CDNC. In the upper layer, the CDNC increase from cloud bottom to near cloud top consistent with the increase in aerosol from below the layer to above the layer. The N50 and N100 estimated for the lower (upper) layer are 269 (334) $cm^{-3}$ and 197 (221) $cm^{-3}$

344 respectively, where the upper layer values are an average of the aerosol at 1400 m and just above
345 cloud top. Thus, on average the CDNC in both layers are approximated by the activation of
346 particles sized between 50 and 100 nm, and the maximum CDNC is approximated by the
347 activation of 50 nm particles. The CCNC(0.6%) are slightly below the N100, which would be
348 consistent with the lower hygroscopicity of BB particles. Comparison of below-cloud
349 CCNC(0.6%) with CDNC suggests cloud S above 0.6%.

350 July 20 Case (Fig. 3 g and h): This is a more complex cloud with substantial LWC
351 variations that suggest three cloud layers. The values of mean CDNC at STP are 45 $cm^{-3}$, 49 $cm^{-3}$
352 and 65 $cm^{-3}$ in the upper, middle and lower layers respectively. The VMD reach about 20 μm in
353 the lower layer and 26 μm in the upper layer with the lower CDNC. These layers are relatively
354 stable with CO and $\theta_e$ increasing slightly from below the cloud to above the top cloud layer.
355 N50 just below the lower layer approximately equals CDNC in that layer. It is more difficult to
356 estimate the pre-cloud aerosol for the middle and upper layers, but particles at least as small as
357 50 nm were apparently activated. For the summary statistics, the respective pre-cloud N100,
358 N50 and CCNC(0.6%) are estimated at 24 $cm^{-3}$, 44 $cm^{-3}$ and 24 $cm^{-3}$ for the upper cloud layer,
359 32 $cm^{-3}$, 52 $cm^{-3}$ and 32 $cm^{-3}$ for the middle layer and 34 $cm^{-3}$, and 66 $cm^{-3}$ and 35 $cm^{-3}$ for the
360 lower layer. Comparison of the CCNC(0.6%), which are in approximately the same
361 concentration as the N100, and CDNC suggests S near or in excess of 0.6%.
362
363 **2.3.2 Low-Altitude (LA) Examples**
364
365 July 5 and July 7 Cases: The two examples in Fig. 4 are for cloud or fog over the polynyas north
366 of Resolute Bay on July 5 and 7. Four cloud samples were collected on July 5 at altitudes below

200 m. The time series in Fig. 4a covers the period of collection of the two lowest samples: 16:18:02-16:21:57 at 130 m and 16:39:35-16:40:18 at 88 m. In the air upwind of the cloud or fog, the N100, N30 and CCNC(0.6%) are estimated at 3 $cm^{-3}$, 10-14 $cm^{-3}$ and 5 $cm^{-3}$. The mean values of the CDNC of 2.8 $cm^{-3}$ at 130 m and 0.7 $cm^{-3}$ at 88 m are explained by the N100 and S less than 0.6%. The maximum CDNC of 12 $cm^{-3}$ at 130 m suggests the activation of smaller particles, possibly as small as 30 nm, and S exceeding 0.6% perhaps due to some uplift influenced by orographic features north of the north polynya. At 88 m, the mean VMD (not shown) was 29 μm and ranged up to 35 μm giving those droplets potential to deposit over an hour or more, thereby potentially transferring water from the polynya to the downwind ice. On July 7, cloud or fog was present below 120 m and thicker towards the north edge of the north polynya and again to the north over the ice. Seven samples were identified over the period 16:06-16:29 based on the LWC above 0.01 g $m^{-3}$. The CDNC are overall higher than on July 5 with sample averages ranging from 4 $cm^{-3}$ to 13 $cm^{-3}$; the one-second CDNC are as high as 34 $cm^{-3}$ and the mean VMD (not shown) range from 19.6 μm to 22.8 μm. The CO mixing ratio is slightly higher within the cloud (81 ppbv) than above (79 ppbv); although this difference may not be significant. In the air nearly free of cloud and below 120 m, the N100 are 4-5 $cm^{-3}$, the N50 are 8-11 $cm^{-3}$ and the N20 are variable between 17 $cm^{-3}$ and 130 $cm^{-3}$; CCN are unavailable for this part of the flight. Mean values of CDNC/N100 and CDNC/N50 for seven cloud samples are 4.8 and 1.0, respectively, indicating that on average particles of about 50 nm were activated in this LA cloud. Based on the overall relationship between CCNC(0.6%) and N50, which is discussed in section 3.3, the mean S in the LA cloud of July 7 is estimated at 0.6%. Comparison with the maximum CDNC suggests that particles as small as 20 nm may have participated in the nucleation of droplets.

July 8 Case: Fig. 5 shows a time series of altitude, CO, N100, N80-100, N90-100, CCNC(0.6%) and CDNC from the sampling above and in the low cloud over Lancaster Sound on July 8. The cloud over the open water of the Sound is visible in the satellite picture in Fig. 2b. Cloud was also present over the ice to the west, but it was much thinner and reached only to about 150 m above the surface. Over the water, the cloud was sampled as high as 230 m by descending into it down to about 150 m between 17:27 UT and 17:43 UT as shown in Fig. 5. Observations in profiles from two of five samples are shown in Fig. 6. This cloud deepened as the aircraft approached the ice edge from over the water, and thinned abruptly over the ice with tops below 150 m as shown in Fig. 5 (time 17:47). The thicker cloud was associated with a shift in wind direction to more southerly suggesting an influence of the Prince Regent Inlet and surrounding terrain on the flow as well as possibly circulations influenced by the water-ice transition. The cloud layer was relatively stable and the $\theta_e$ profiles suggest a surface heat sink (Fig. 6a). Profiles of LWC and VMD in Fig. 6 (b, c) do not show increases with altitude characteristic of vertical mixing, such as for some of the HA clouds (Fig. 3); the change in the VMD per 50 m increase in height is about 1.7 μm for the well mixed cloud of July 7 (Fig. 3 a, b), whereas it is about 0.2 μm per 50 m for the LA cloud of flight 8 in Fig. 6. The CO mixing ratio shows little variation with time and altitude. The pre-cloud aerosol concentrations are more difficult to assess. Based on concentrations just above the cloud, particles >90 nm explain the CDNC. Based on the concentrations downwind at 150 m (approximately 17:47), activation of particles >80 nm is needed to explain the CDNC. The CCNC(0.6%) are about 129 $cm^{-3}$ downwind and between 157 $cm^{-3}$ and 234 $cm^{-3}$ just above cloud. It is concluded that in this case the droplets likely nucleated on particles mostly larger than 80-95 nm and the S in the clouds were less than 0.6%. For the purposes of summary statistics discussed next, the N100, N50 and

413 CCNC(0.6%) have been selected as an average of the downwind and immediately above cloud
414 concentrations: 73 $cm^{-3}$, 319 $cm^{-3}$ and 168 $cm^{-3}$, respectively.
415
416 **3. Summary Observations and Discussion**
417
418 Summary statistics for the cloud and aerosol samples are discussed in 3.1, the microphysics of
419 low-altitude and higher-altitude clouds are contrasted in 3.2, particle activation is summarized in
420 3.3 and in section 3.4 the relationship between VMD and CDNC is used to consider the
421 transition of aerosol indirect effects from potential warming to potential cooling. All analyses
422 are based on the 62 cloud samples discussed in section 2.3. The LA cloud subset is comprised of
423 24 samples and the HA cloud subset consists of 38 samples.
424
425 **3.1 Summary of mean observations**
426
427 The mean and median values of the microphysical properties of the cloud and pre-cloud aerosols
428 as well as the altitudes and temperatures derived from the 62 cloud samples are given in Table 1,
429 separated between periods 1 and 2. Values of the CDNC and the LWC are given relative to in-
430 situ volumes as well as STP. As discussed above, the pre-cloud CCNC(0.6%), N50, and N100
431 are averages of those values collected within about 50 m of cloud base where a cloud base was
432 clear and achievable. In other cases the pre-cloud CCNC(0.6%), N50 and N100 are the values at
433 the similar or lower altitudes in the clear air upwind of the cloud, except in the case of July 8
434 when the pre-cloud aerosol is based on the measurements in the area downwind plus those
435 immediately above cloud. The CCNC(0.6%) samples in Table 1 are limited to 44 due to
436 instrument problems, all of which occurred during the early part of July 7.

Cloud liquid water paths (LWP) are estimated for 36 of the samples when a complete profile between cloud base and cloud top was possible. The LWP are shown at the bottom of Table 1. Of the 36 LWP estimates, 34 are above 200 m, and the mean and median altitudes are 1044 m and 862 m, respectively. Not included in the summary statistics are the samples from July 8 shown in Figs. 5 and 6. For the minimum altitude reached in that cloud, the LWP ranged from 12 to 25 and thus the total LWP for that cloud exceeded 25.

During period 1, the median sampling altitude is lower and the temperatures are slightly below freezing compared with just above freezing during period 2. The CO mixing ratios are overall low and at approximately background values during period 1. The median CDNC are higher during period 1 than period 2, but the mean values are similar. The CDNC compare more closely with the N50 during period 1, while during period 2 the CDNC are about equally between the N50 and N100. The CCNC(0.6%) equated with particles between 50 nm and 100 nm during period 1, whereas during period 2 they were closer to the N100 values. The reduction in particle hygroscopicity during period 2 may be due to an increased presence of organics in the aerosol during that time (Willis et al., 2016).

### 3.2 Comparison of LA and HA cloud

The LA clouds were close to the surface, and all associated with open water; some or all may be technically fogs. They may be formed by advection of warmer moist air over a cooler surface (the July 8 LA cloud that moved from Baffin Bay westward along Lancaster Sound was likely dominated by that process), by radiation cooling or by the passage of very cold air over a warm moist surface. The latter, also known as sea smoke, is the likely explanation for the clouds over

460 the polynyas; also, it is possible that there was an advection component associated with the sea
461 smoke moving from the polynyas over the ice surfaces. In general, the LA clouds are associated
462 with low-level horizontal advection and heat and water exchange with the underlying ice or
463 water surface. In contrast, vertical motions are responsible for some of the HA clouds, and none
464 of the HA clouds interact so closely with the underlying surface. Due to those differences, the
465 characteristics of the LA and HA clouds are considered separately. Table 2 shows the mean and
466 median values for the samples separated between LA and HA clouds; vertical profiles of CDNC,
467 LWC and VMD samples are shown in Supplement Fig. S7. On average, the LA samples have
468 lower CDNC and higher VMD compared with the HA cases, and the LA clouds are activating on
469 larger particles relative to the HA clouds (e.g. CDNC/N50). The values of the
470 CDNC/CCNC(0.6%) indicate that the S are <0.6% for the LA clouds and close to 0.6% for the
471 HA clouds.

472 Variations in LWC are correlated with those of CDNC for the LA samples (Fig. 7a). The
473 coefficient of determination ($R^2$) rises from 0.57 to 0.98 if the one LA point at (137, 0.032) is
474 removed. In contrast, the correlation of the LWC with the CDNC for the HA samples is low
475 ($R^2$=0.12). There is no correlation of the LWC with the VMD for the LA points ($R^2$=0.04), and
476 for the HA clouds there is a modest correlation of LWC with MVD ($R^2$=0.26). Variations in
477 LWC with VMD within a cloud system are consistent with lifting of air from below, i.e.
478 nucleation of droplets at cloud base followed by their growth with increasing altitude, such as the
479 case shown in Fig. 3a and 3b. Variations of LWC with VMD can also result from homogeneous
480 mixing (i.e. entrainment of dry air that reduces LWC by partial evaporation of droplets without
481 reducing CDNC). The strong dependence of the variations in LWC with those of the CDNC in
482 the LA clouds may reflect changes in rate of cooling, collision-coalescence or inhomogeneous

483 mixing along the cloud transport pathway. For example, increases in the rate of cooling within or
484 between clouds will increase condensation rates, and potentially S, resulting in increased LWC
485 and CDNC. Changes in collision-coalescence will affect the CDNC and LWC in similar ways:
486 more collision-coalescence, lower CDNC and lower LWC due to precipitation. Inhomogeneous
487 mixing, the entrainment of dry air parcels into a cloud without mixing with the cloud droplets,
488 will reduce the CDNC averaged across the cloud and at the same time reduce the mean LWC.
489 Changes in the aerosol that are interactive with some of the cloud processes may contribute to
490 the CDNC and potentially the LWC through their influence on collision-coalescence.
491 The LWC-CDNC correlation is identifiable for individual flights with sufficient LA
492 samples: four flights, comprising 20 of the 24 LA samples, had three or more points as shown in
493 Fig. 8. The regressions for each of the July 7, 8 and 17 cases are approximately linear, and the
494 respective mean VMD are 20.8 μm, 18.8 μm and 18.2 μm. The mean LWC are 0.05 g $m^{-3}$, 0.3 g
495 $m^{-3}$ and 0.07 g $m^{-3}$. The VMD are relatively close together confirming similarities in the
496 relationships, even if not purely linear. For comparison, the mean VMD for the July 5 samples is
497 29.2 μm and the LWC is 0.02 g $m^{-3}$, which indicates that the July 5 case does not fit the linear
498 relationship shown in Fig. 8. The reasons behind the similarity of the VMD for the July 7, 8 and
499 17 are unknown, but it occurs despite the varied pre-cloud N50 and N100: N50 range of 5-272
500 $cm^{-3}$; N100 range of 1.1-73 $cm^{-3}$. The consistencies among the three flights for greatly differing
501 aerosol and CDNC imply a much smaller role for the aerosol in terms of the LWC. The
502 distributions of droplets extend above 20 μm in these cases, but few are of sufficient size to
503 initiate collision-coalescence (about 30 μm) (e.g. Rosenfeld et al., 2001) unless some fall out
504 already had occurred. Greater temporal and spatial coverage are needed to assess the
505 microphysical processes in these clouds.

506

507 ### 3.3 Particle Activation Sizes

508

509 Here, the sizes and CCN activity of particles that acted as nuclei for cloud droplets are examined.
510 The CDNC are plotted versus N100 in Fig. 9a, separated between LA and HA samples. The
511 CDNC are most often higher than the N100 and more so for the HA samples, which indicates
512 that particles smaller than 100 nm activated in most cases and most often in the HA clouds. The
513 mean and median values of CDNC(STP)/N100 are 2.2 and 1.8 for all 62 samples, and the $30^{th}$
514 percentile of the CDNC/N100 is 1.2, which means that in about 70 % of the cases droplets
515 nucleated on particles significantly smaller than 100 nm. Fig. 9a can be compared with the
516 results of Hegg et al. (2012) who showed a linear fit of CDNC to N100 for marine stratocumulus
517 with a slope of 0.72 for which the N100 in 94% of the samples was >150 $cm^{-3}$. Here, a slope
518 larger than unity is indicated, and the N100 are <100 $cm^{-3}$ in 90% of the samples. The
519 comparison indicates that relationships derived for higher concentration environments do not
520 necessarily apply to those of lower concentration environments. In the clean environment often
521 found in the Arctic during summer, the absence of larger particles may lower water uptake rates
522 during droplet nucleation, which will increase the S, enabling cloud droplets to nucleate on
523 smaller particles; the absence of larger particles may also help increase the concentrations of
524 smaller particles in the Arctic during summer, by promoting new particle formation through a
525 reduced condensation sink (e.g. Tunved et al., 2013; Leaitch et al., 2013). The CDNC are plotted
526 against the N50 in Fig. 9b showing that the mean activation size of the HA clouds was often
527 close to 50 nm. The median value of CDNC/N50 is 0.78 for all samples indicating that, based on
528 the averaged CDNC, cloud droplets nucleated on particles near or smaller than 50 nm about 40%

529 of the time. That percentage will increase if particle activation is considered relative to the
530 maximum CDNC.

531 The mean and median values of the CCNC(0.6%) associated with all cloud samples (84
532 $cm^{-3}$ and 47 $cm^{-3}$) are generally consistent with previous Arctic CCNC measurements. For
533 example, during the summer above 85°N, Martin et al. (2011) measured a mean CCNC at 0.73%
534 S of 47 $cm^{-3}$ with a standard deviation of 35 $cm^{-3}$, Yum and Hudson (2001) measured CCNC at
535 0.8% S below 1700 m over the Beaufort Sea during May, 1998 that ranged from 41 $cm^{-3}$ to 290
536 $cm^{-3}$, and Radke et al. (1976) measured a mean CCNC at 1% S of 90 $cm^{-3}$ in June near Barrow,
537 Alaska. Considering the median values of CDNC/CCNC(0.6%) for the LA and HA samples
538 (Table 2) and the slopes of linear regressions of CDNC versus CCNC(0.6%) (Fig. 10a), the
539 average inferred S for the HA clouds is about 0.6%, consistent with the overall activation of
540 smaller particles in those clouds. The mean S inferred for the LA clouds is significantly lower
541 than 0.6%. Based on the activation of a 90 nm particle (July 8 case; CCNC(0.6%) of 168 $cm^{-3}$ in
542 Fig. 10a) of low-moderate hygroscopicity, a reasonable estimate is 0.3% for the mean of the LA
543 clouds with some higher values indicated by the points near a CCNC(0.6%) of 25 $cm^{-3}$ in Fig.
544 10a. The S for these clean clouds are in contrast to polluted marine environments for which
545 estimates for these types of clouds are 0.2% or less (e.g. Modini et al., 2015). Consistent with the
546 present results, Hudson et al. (2010) found that effective S in marine stratus tended to increase
547 with a decrease in the CCNC, and for CCNC smaller than about 200 $cm^{-3}$ their effective S ranged
548 between 0.3% and 1.2%.

549 Variations in the measured CCNC(0.6%) are explained well by variations in smaller
550 (N50) and larger (N100) particles as shown in Fig. 10b. The slopes of the power-law fits, for

551 which the exponents are both close to unity, indicate that the CCNC(0.6%) at 0.6% S on average
552 fall between 50 nm and 100nm.
553
554 **3.4 Aerosol Influences on Warming to Cooling**
555
556 The relationship between the VMD and CDNC shown in Fig. 11 exhibits a scattered but clear
557 tendency for smaller VMD with increasing CDNC. The solid black curve is a reference line
558 based on the study-mean LWC of 0.12 g $m^{-3}$ (Table 1); points falling above or below the black
559 curve have higher or lower LWC, respectively. The vertical dashed green line represents our best
560 estimate of the Mauritsen limit below which Mauritsen et al. (2011) showed the cloud may
561 produce a net warming for an increase in the CDNC. The net warming is a consequence of an
562 increase in longwave absorption due to an increase in the LWC, where the latter results from a
563 reduction in deposition for the smaller droplets associated with increased CDNC. A value of 16
564 $cm^{-3}$ is our best estimate of the Mauritsen limit for this data set because all points with CDNC
565 below that value fall well below the mean LWC, therefore offering greater potential for changes
566 in the CDNC to increase the LWC. Above the estimated Mauritsen limit, an increase in CDNC
567 may produce a net cooling due to the cloud albedo effect, since at that point the longwave
568 forcing does not change significantly as the effects of deposition are minimized and the cloud
569 effectively behaves as a black body.
570 The aerosol influence on clouds with CDNC below the Mauritsen-limit is considered in
571 section 3.4.1. In section 3.4.2, the potential background influence of the aerosol on clouds with
572 CDNC above the Mauritsen-limit is examined.
573
574 **3.4.1 Below the Mauritsen limit**

Seventeen of the 62 samples fall at or below our best estimate of the Mauritsen limit. Fifteen of those 17 samples are from LA clouds with median pre-cloud N50 and N100 estimates of 8.2 $cm^{-3}$ and 3.0 $cm^{-3}$ respectively. The lower number concentrations contribute to overall larger VMD; although some of the points below the estimated Mauritsen limit have VMD values much less than 20 μm. Increases in small particles, potentially from particle nucleation or fragmentation (e.g. Leck and Bigg, 1999 and 2010), are hypothesized to increase the CDNC thereby enhancing longwave warming by these clouds, at least until the CDNC exceed the estimated Mauritsen limit. The LA points from the July 5 and the July 7 cases, identified in Fig. 11, offer one insight. The median CDNC for July 5 is six times lower than the July 7 CDNC: 1.3 $cm^{-3}$ and 7.8 $cm^{-3}$, for July 5 and 7, respectively. The median N50 are 6 $cm^{-3}$ and 8.3 $cm^{-3}$ for July 5 and 7, respectively, and the median N100 are 3 $cm^{-3}$ and 2.2 $cm^{-3}$ for July 5 and July 7, respectively. The CDNC are similar to N50 in the July 7 case, but lower than both the N50 and N100 in the July 5 case indicating that the aerosol was not a limiting factor in the July 5 case. Consistent with the discussion in section 3.2, all 15 LA points show a correlation of LWC with the CDNC ($R^2$=0.57), but correlations of CDNC with N50 and N100 are weak: $R^2$=0.19 and 0.06, respectively. The CCN are not used here because only seven points with CCNC(0.6%) are available; the seven do, however, correlate well with the N50. If the limit of 10 $cm^{-3}$ of Mauritsen et al. (2011) is applied, reducing the number of points to 12, the assessment does not change: the LWC-CDNC correlation improves slightly and the correlations of the CDNC with the N100 and the N50 weaken.

The LWC do not correlate with either the N50 or the N100 (Supplement Fig. 8). In this low CDNC environment, where cloud droplets may grow large enough to be gravitationally

598 removed from the cloud without the support of collision-coalescence, the absence of a positive
599 correlation of either the CDNC or LWC with the aerosol indicates that small changes in the
600 aerosol did not contribute significantly to the changes in the LWC. Variations in other
601 processes, such as mixing or the rate of cooling, may be responsible for the correlation of CDNC
602 and LWC. It can be argued that some aerosol must exist for these clouds to form, but these
603 observations show no association of changes in either the CDNC or LWC with changes in the
604 aerosol.
605
606 **3.4.2 Background aerosol influence on clouds**
607
608 Above the estimated Mauritsen limit, the general reduction in the VMD with the CCNC(0.6%)-
609 associated increase in CDNC reflects the impact of increases in aerosol on clouds. In Fig. 11,
610 samples are identified between those associated with lower CO (green circles; <81 ppbv, the
611 median CO value of all samples) and those with highest CO (red circles; >90 ppbv); six samples
612 have no CO measurement and the remaining points have CO falling within 81-90 ppbv. Five of
613 the seven higher-CO samples are from the July 19 case (e.g. Fig. 3e, 3f) that has been linked with
614 BB (Köllner et al., 2015; reference above), and the highest CDNC point (273 $cm^{-3}$; no CO
615 measurement) is also from July 19 and likely influenced by BB. The higher-CO samples cover a
616 range of CDNC from 16 $cm^{-3}$ to at least 238 $cm^{-3}$ with CO reaching up to 113 ppbv. The higher
617 CO samples are associated with larger particles (N50/N100=1.5), consistent with a BB influence,
618 compared with the lower CO samples (N50/N100=3.2). These values for BB fall at the low end
619 of the observations from Zamora et al. (2015), but their CO concentrations are much higher than
620 those measured in this study. The lower-CO samples may be dominated by regional biogenic

emissions (Willis et al., 2016). The lower- and higher-CO points overlap over a CDNC range of 16 $cm^{-3}$ to 160 $cm^{-3}$, consistent with the range of pre-industrial CDNC from global models of 30 $cm^{-3}$ to 140 $cm^{-3}$ (Penner et al., 2006; Korhonen et al., 2008). In this clean environment, the contributions from 20-100 nm particles have a broad impact on the range of CDNC, affirming the large uncertainty associated with estimating a baseline for the cloud albedo effect discussed by Carslaw et al. (2013).

## 4. Summary and Conclusions

Aerosol particle size distributions, CCNC(0.6%) at 0.6% water S, carbon monoxide (CO) and cloud microphysics were measured from an airborne platform based out of Resolute Bay, Nunavut from July 4 to July, 21, 2014 as one part of the Canadian NETCARE project. The flights were conducted over ice and water surfaces from about 60 m above the surface to about 6000 m. Sixty-two (62) cloud-averaged samples were derived, each constrained for the mean LWC >0.01 g $m^{-3}$ or the cloud threshold used here. The analysis separates the cloud samples between 24 low-altitude (LA: <200 m) samples and 38 higher altitude (HA: >200 m) samples as well as situations of lower and higher CO and observations above and below the Mauritsen et al., (2011) CCNC(0.6%) (or CDNC) limit.

The median pre-cloud N100 of 33 $cm^{-3}$ and the median CO mixing ratio of 81 ppbv indicate that the aerosols supporting the sampled clouds were relatively clean, and particularly during the first part of the study many of the aerosol particles may have been derived from regional natural sources. The median CDNC at STP is 10 $cm^{-3}$ for the LA clouds (24 samples) and 101 $cm^{-3}$ for the HA clouds (38 samples), which correspond with the median pre-cloud N50

644 of 11 $cm^{-3}$ for the LA samples and 133 $cm^{-3}$ for the HA samples. The lower sizes of particles
645 activated in cloud varied from about 20 nm to above 100 nm. In 40% of cases, the average lower
646 size of activation was 50 nm or smaller. Overall, smaller particles were activated more often in
647 the HA clouds. Variations in particle chemistry will induce some variance in these results, but
648 because activation diameters are estimated starting with larger particles and moving to smaller
649 sizes, changes in chemistry only offer the possibility of activation of particles still smaller than
650 estimated here; although that would have to occur at the expense of larger particles.

651 From the median values of CDNC/CCNC(0.6%) (1.2 for the HA clouds and 0.6 for the
652 LA clouds) and the linear regression of CDNC and CCNC(0.6%), it is inferred that the average S
653 were approximately 0.6% for the HA clouds and 0.3% for the LA clouds. Higher estimates will
654 be obtained if the maximum CDNC are taken into consideration rather than the mean CDNC.
655 The relatively high S for these clean Arctic stratus and stratocumulus have similarities with the
656 observations of Hudson et al. (2010) for relatively clean stratus off the coast of California.

657 In 17 cases, 15 of which are LA clouds, the CDNC fell at or below the CCN limit
658 discussed by Mauritsen et al. (2011), which is estimated here as 16 $cm^{-3}$. These are the first
659 collection of simultaneous observations of the microphysics of aerosols and clouds in this unique
660 regime in which the net radiative impact of increases in the CDNC is hypothesized to be
661 warming due to changes in the LWC. The LWC of the points below the Mauritsen limit all fall
662 below the study-mean LWC, and the LWC increases with the CDNC. Neither the CDNC nor the
663 LWC are positively correlated with the pre-cloud aerosol (N50 or N100). In this environment of
664 low cloud or fog and ultra-low CDNC, variations in cloud processes such as mixing or the rate of
665 cooling may be responsible for the correlation of CDNC and LWC. These observations show no

666 association of changes in either the CDNC or LWC with changes in the aerosol within the
667 Mauritsen limit.

668 Forty-five cloud samples with CDNC above the Mauritsen limit exhibit a clear influence
669 of changing aerosol. The cloud microphysics for the clouds formed in cleaner air (smaller
670 particles and lower CO: <81 ppbv) overlap with clouds formed in what was likely more polluted
671 air (larger particles and higher CO: >90 ppbv) covering a CDNC range of 16-160 $cm^{-3}$. It is
672 concluded that 20-100 nm particles from natural sources can have a broad impact on the range of
673 CDNC in clean environments, affirming a large uncertainty in estimating a baseline for the cloud
674 albedo effect.

675

676

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**Table 1.** Summary of averaged cloud observations with LWC>0.01 g $m^{-3}$ for study periods 1 and 2. Values without parentheses are referenced to ambient volumes and values in parentheses are referenced to STP. 5;95 are the 5th and 95th percentiles.

| Measurement | Period 1 (July 5-11): 35 samples; 1.2 hours in cloud | | | Period 2 (July 11-21): 27 samples; 0.4 hours in cloud | | |
|---|---|---|---|---|---|---|
| | Mean | Median | 5;95 | Mean | Median | 5;95 |
| Altitude (m-msl) | 920 | 178 | 88;2272 | 1011 | 835 | 97;2608 |
| Temperature ($^{o}$C) | -1.9 | -0.4 | -6.5;2.2 | +1.2 | +2.2 | -4.9;3.5 |
| CDNC (STP) ($cm^{-3}$) | 75 (85) | 93 (91) | 1.1;154 (1.1;185) | 73 (83) | 52 (55) | 13;228 (14;265) |
| LWC (STP) (g $m^{-3}$) | 0.12 (0.13) | 0.10 (0.12) | 0.014;0.32 (0.013;0.32) | 0.12 (0.13) | 0.12 (0.13) | 0.025;0.26 (0.024;0.31) |
| VMD (μm) | 17.2 | 18.7 | 9.9;30.0 | 15.0 | 14.5 | 9.1;21.4 |
| CCNC(0.6%) ($cm^{-3}$): (17 P-1; 27 P-2) | 90 | 120 | 2;168 | 81 | 43 | 18;227 |
| N50 ($cm^{-3}$) | 113 | 134 | 4.8;319 | 126 | 68 | 29;334 |
| N100 ($cm^{-3}$) | 35 | 47 | 1.3;73 | 81 | 31 | 13.8;274 |
| CDNC(STP)/CCNC(0.6%) | 0.75 | 0.56 | 0.18;1.50 | 1.18 | 1.22 | 0.47;1.87 |
| CDNC(STP}/N50 | 0.82 | 0.90 | 0.16;1.40 | 0.73 | 0.68 | 0.28;1.08 |
| CDNC(STP)/N100 | 2.78 | 2.63 | 0.28;7.94 | 1.37 | 1.25 | 0.58;2.15 |
| CCNC(0.6%)/N50 | 0.64 | 0.63 | 0.50;0.84 | 0.64 | 0.64 | 0.52;0.87 |
| CCNC(0.6%)/N100 | 1.92 | 1.79 | 0.67;3.11 | 1.27 | 1.0 | 0.75;2.28 |
| CO (ppbv) | 79 | 80 | 77;81 | 90 | 87 | 81;108 |
| LWP (g $m^{-2}$); (13 P-1; 23 P-2) | 30 | 27 | 1.5;4 | 22 | 13 | 1.0;70.5 |

**Table 2**. Summary of averaged observations for low-altitude (LA) and higher-altitude (HA) clouds. Values without parentheses are referenced to ambient volumes and values in parentheses are referenced to STP. 5, 95 are the 5th and 95th percentiles.

| Measurement | LA (<200m): 24 samples; 0.89 hours in cloud | | | HA (>200m): 38 samples; 0.72 hours in cloud | | |
|---|---|---|---|---|---|---|
| | Mean | Median | 5;95 | Mean | Median | 5;95 |
| Altitude (m-msl) | 129 | 127 | 79;178 | 1485 | 1481 | 457;2391 |
| Temperature ($^{o}$C) | +0.6 | +0.2 | -2.5;2.9 | -1.2 | +0.9 | -6.5;2.7 |
| CDNC (STP) (cm$^{-3}$) | 31 (30) | 11 (10) | 1;106 (1;102) | 101 (118) | 91 (101) | 28;211 (31;245) |
| LWC (STP) (g m$^{-3}$) | 0.10 (0.10) | 0.05 (0.05) | 0.01;0.34 (0.01;0.33) | 0.13 (0.15) | 0.13 (0.15) | 0.04;0.25 (0.04;0.30) |
| VMD (μm) | 20.7 | 20.1 | 14.6;31 | 13.4 | 12.5 | 9.1;19.4 |
| CCNC(0.6%) (cm$^{-3}$); (16 LA; 28 HA) | 74 | 24 | 2;184 | 90 | 58 | 21;217 |
| N50 (cm$^{-3}$) | 91 | 11 | 4.2;319 | 136 | 133 | 41;334 |
| N100 (cm$^{-3}$) | 26 | 4 | 1.3;73 | 73 | 47 | 20;232 |
| CDNC(STP)/CCNC(0.6%) | 0.61 | 0.57 | 0.18;1.3 | 1.2 | 1.2 | 0.6;1.9 |
| CDNC(STP}/N50 | 0.61 | 0.44 | 0.14;1.5 | 0.91 | 0.93 | 0.5;1.3 |
| CDNC(STP)/N100 | 2.3 | 1.4 | 0.35;9.0 | 2.1 | 1.9 | 0.7;3.7 |
| CCNC(0.6%)/N50 | 0.66 | 0.71 | 0.52;0.7 | 0.68 | 0.64 | 0.5;0.9 |
| CCNC(0.6%)/N100 | 1.8 | 1.6 | 0.96;2.6 | 1.5 | 1.1 | 0.8;3.4 |
| CO (ppbv) | 81 | 80 | 78;82 | 86 | 83 | 77;107 |

**Figure. Captions**

Figure 1. Compilation of the flight tracks. All flights originated from Resolute Bay (74°40'48"N 94°52'12"W).

Figure 2. Satellite images from July 5 when LA clouds were sampled over the two polynyas to the north and from July 8 when LA clouds were sampled along Lancaster Sound (July 8). Lancaster Sound is cloud free on July 5 and mostly covered by cloud on July 8. Resolute Bay is marked with a "X". Images are courtesy of NASA Worldview: https://earthdata.nasa.gov/labs/worldview/.

Figure 3. Four examples of profiles through HA clouds. a) Case from July 7 showing CO, CDNC, CCNC(0.6%) and particle number concentrations, where Nx-100, N100 and N5 are for particles sized between "x" nm and 100 nm, >100 nm and >5 nm respectively. b) Case from July 7 showing LWC, VMD, $\theta_e$ and temperature, where VMD, $\theta_e$ and temperature have been scaled as indicated in the legend. c) As in a), but case from July 17 and without N5. d) As in b), but case from July 17. e) As in a), but case from July 19. f) As in b) but case from July 19. g) As in a) but case from July 20 and without N5. H) as in b), but case from July 20. The CDNC are all referenced to STP, and $\theta_e$ is given in degrees Centigrade before scaling.

Figure 4. Time series during the sampling of low (LA) cloud or fog over the polynyas north of Resolute Bay. a) July 5 time series showing CO, CDNC, CCNC(0.6%) and particle number concentrations, where N30-100 is for particles sized between 30 nm and 100 nm and N100 is for particles sized >100 nm. b) July 7 time series showing CO, CDNC and particle number concentrations, where N20-100, N50-100 and N100 are for particles sized between 20 nm and 100 nm, between 50 nm and 100 nm and >100 nm respectively. CCNC(0.6%) measurements are unavailable for this period on July 7. Wind direction and relative position of polynyas are indicated in both panels. CDNC are referenced to STP.

Figure 5. Time series of altitude, CO, N80-100, N90-100, N100, CCNC(0.6%) and CDNC from low cloud (LA) cloud sampling over Lancaster Sound on July 8. The cloud was deeper over the open water of the Sound (see satellite picture in Fig. 2b). Over the ice to the west, the cloud was not as deep and could not be sampled. Segments over water and ice are indicated at the top of the figure.

Figure 6. Profiles down into cloud showing a) $\theta_e$, b) LWC and c) VMDData for periods 17:27-17:29 UT and 17:38-17:39 UT during July 8. d) shows CDNC, N100, CO and CCNC(0.6%) for the 17:27-17:29 UT profile, and e) shows CDNC, N100, CO and CCNC(0.6%) for the 17:38-17:39 UT profile.

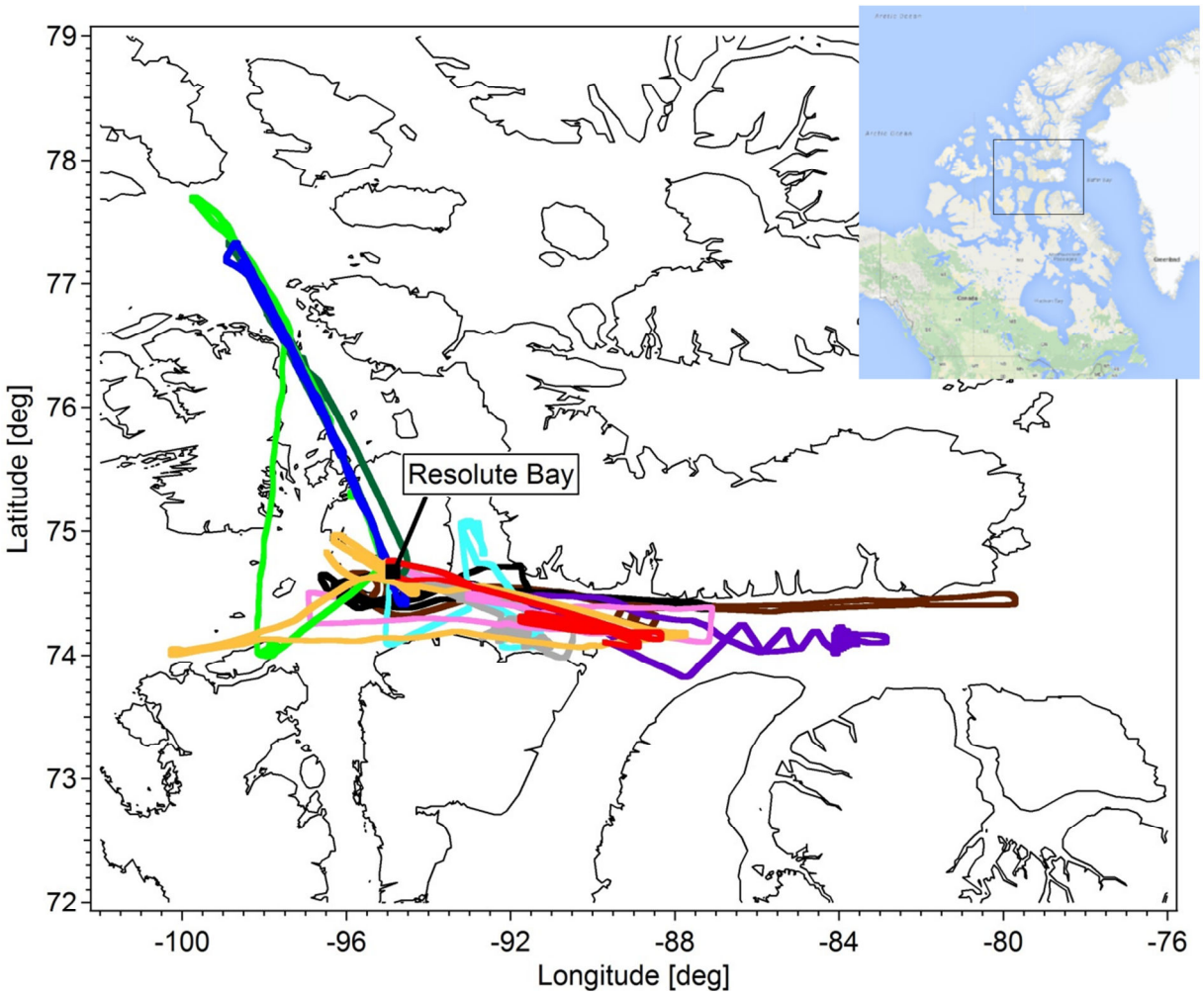
Figure 7. The LWC plotted as a function of the CDNC (a) and VMD (b) for the LA (orange) and HA (blue) samples. Linear regressions for each of the LA and HA samples are also plotted, and the coefficients of determination are given in the legends.

Figure 8. As in Fig. 7a, but only for four identified LA cases (July 5, 7, 8 and 11). Linear regressions for each set of samples are also plotted, and the coefficients of determination are given in the legends.

Figure 9. Plots of CDNC versus a) N100 and b) N50. Points are identified between LA (yellow) and HA (blue) samples, and the 1:1 lines are for reference.

Figure 10. a) CDNC plotted versus the CCNC(0.6%) measured at 0.6% supersaturation; points are identified between LA (yellow) and HA (blue) samples, and linear regressions through the origin are shown. b) CCNC(0.6%) plotted versus N50 and N100; power law fits to each are provided for reference.

Figure 11. The mean VMD of all cloud samples plotted versus the CDNC. All CDNC are referenced to the in-situ pressure. The dashed vertical green line represents the "CCN-limited" division discussed by Mauritsen et al (2011) and estimated here as 16 $cm^{-3}$. The solid black line is another reference showing the relationship between VMD and CDNC for a constant LWC: the study mean LWC of 0.12 g $m^{-3}$ (Table 1). Samples with higher CO (>90 ppbv) are identified by the open red circles. Also highlighted for the discussion are LA samples from July 5 (red dots) and July 7 (orange dots). The median CDNC are 1.3 $cm^{-3}$ and 7.8 $cm^{-3}$, for July 5 and 7, respectively; the N50 are 6 $cm^{-3}$ and 8.3 $cm^{-3}$ for July 5 and 7, respectively; the N100 are 3 $cm^{-3}$ and 2.2 $cm^{-3}$ for July 5 and July 7, respectively.

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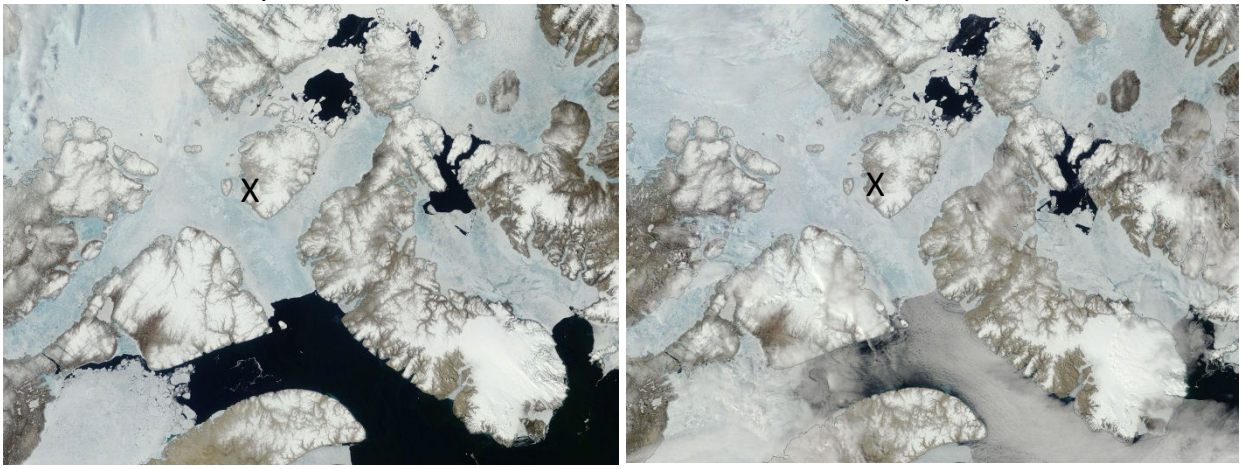
July 5, 2014 July 8, 2014

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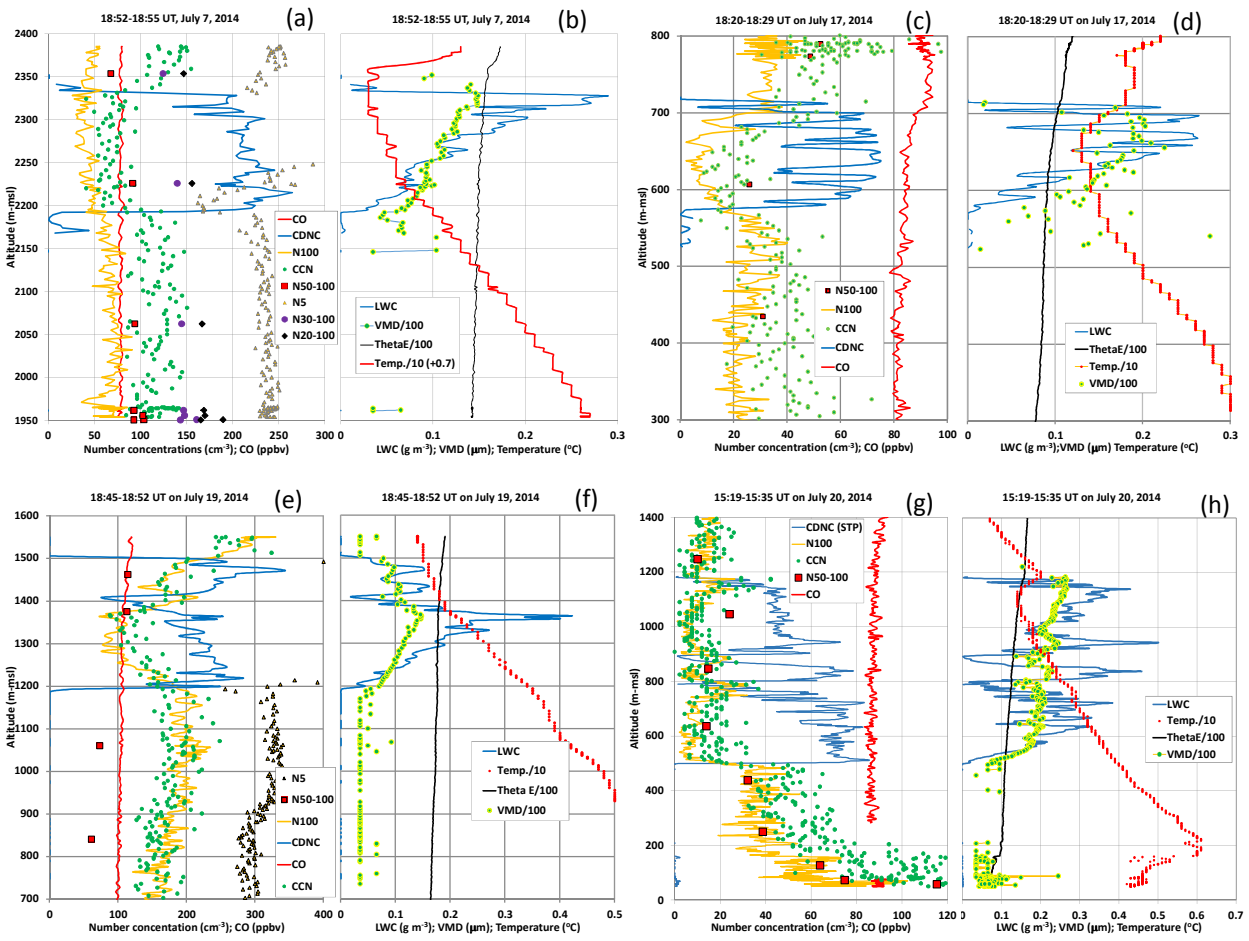


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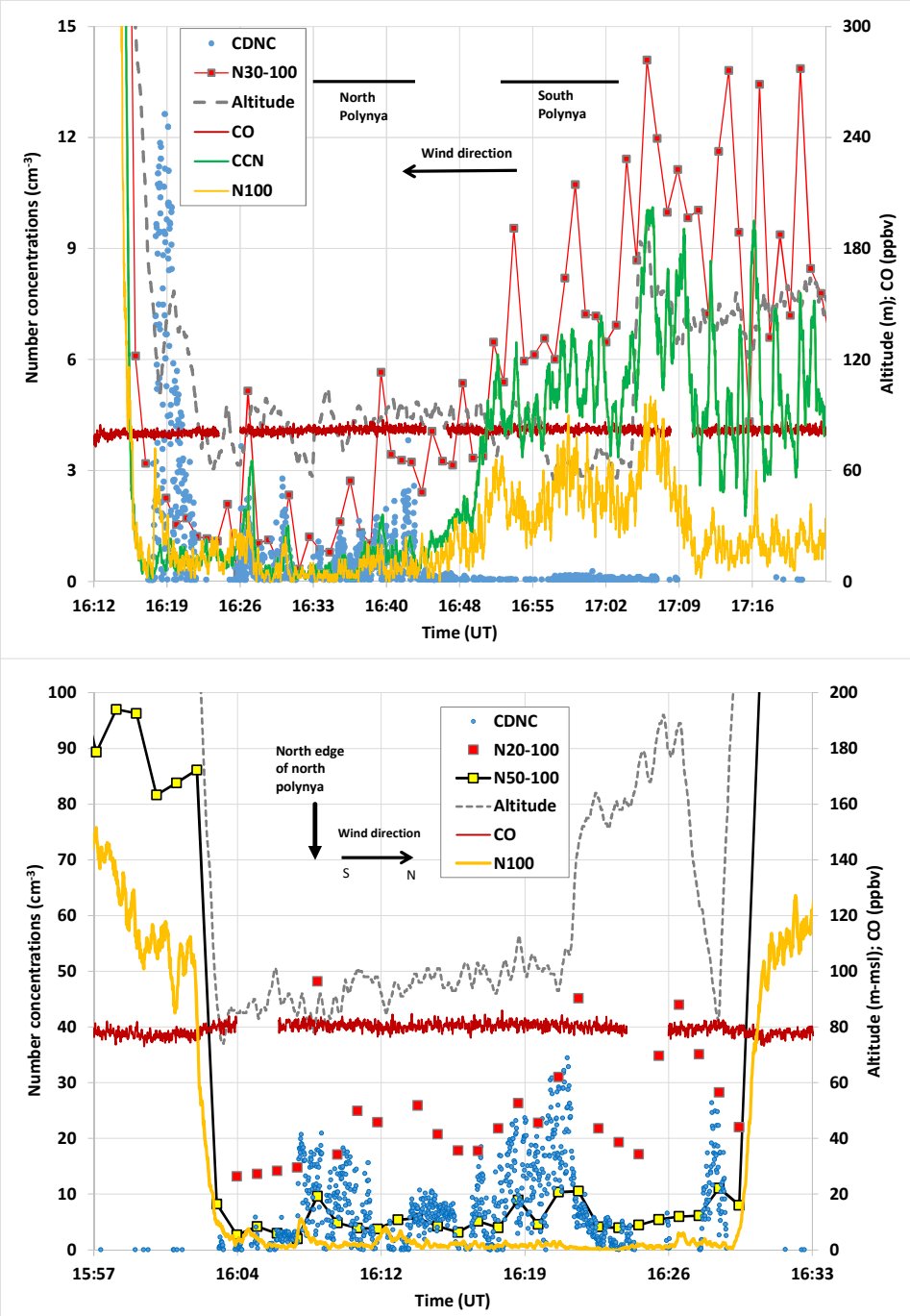


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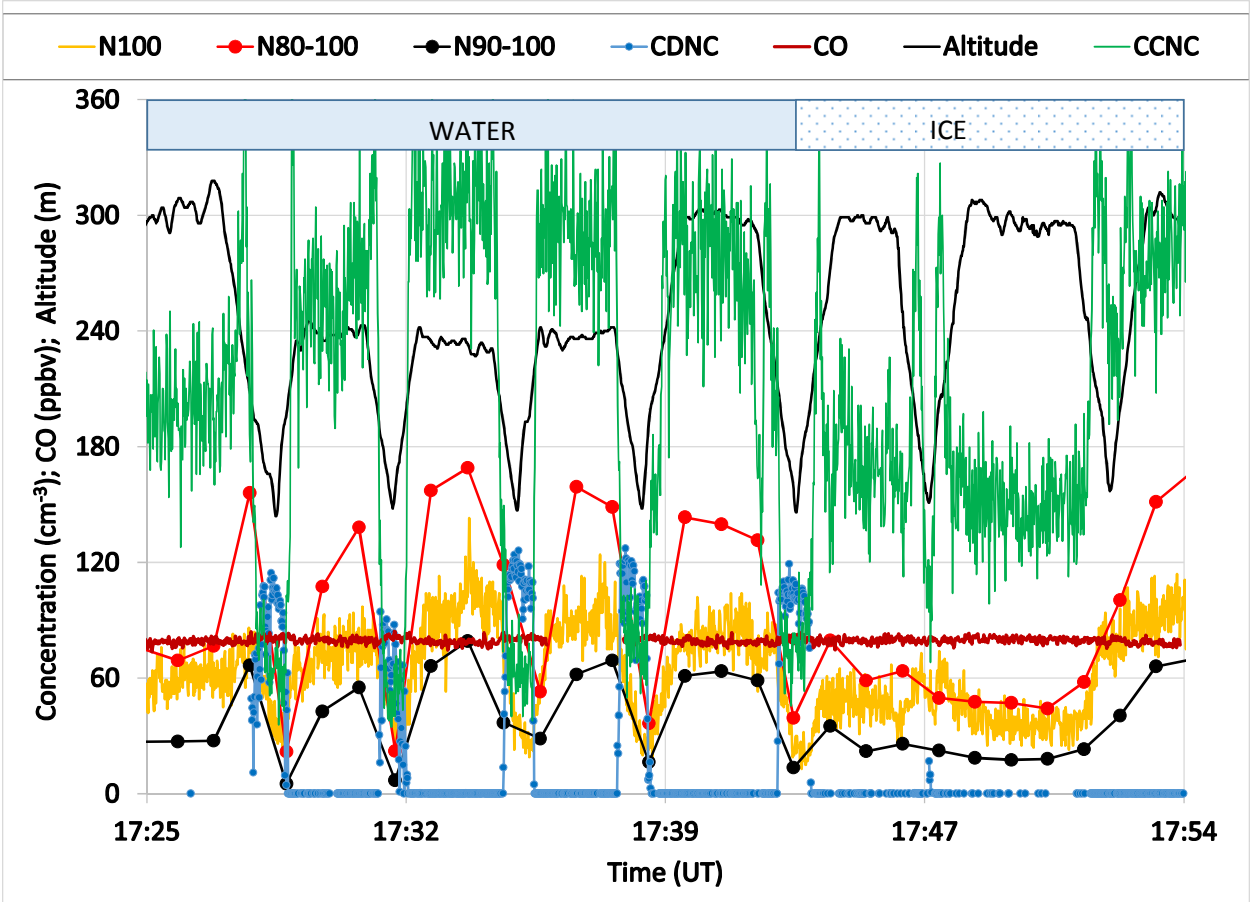


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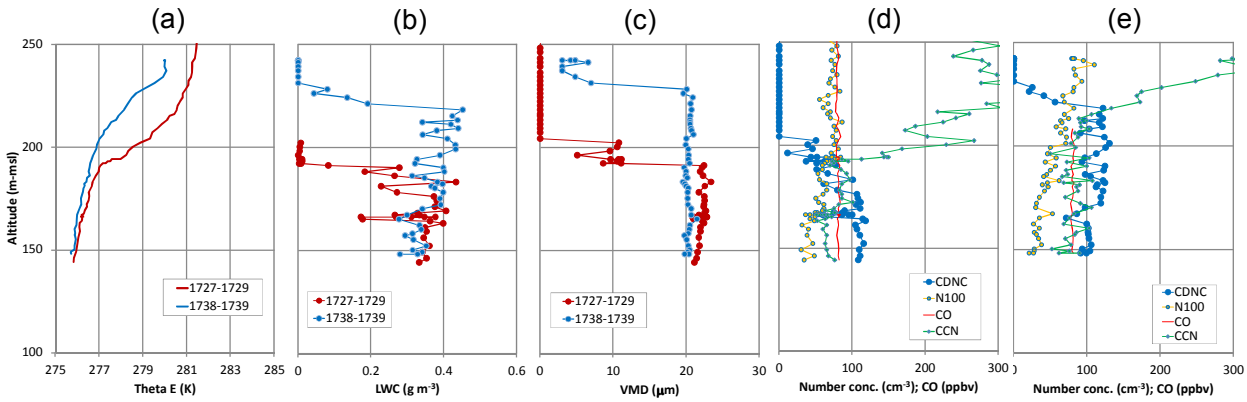


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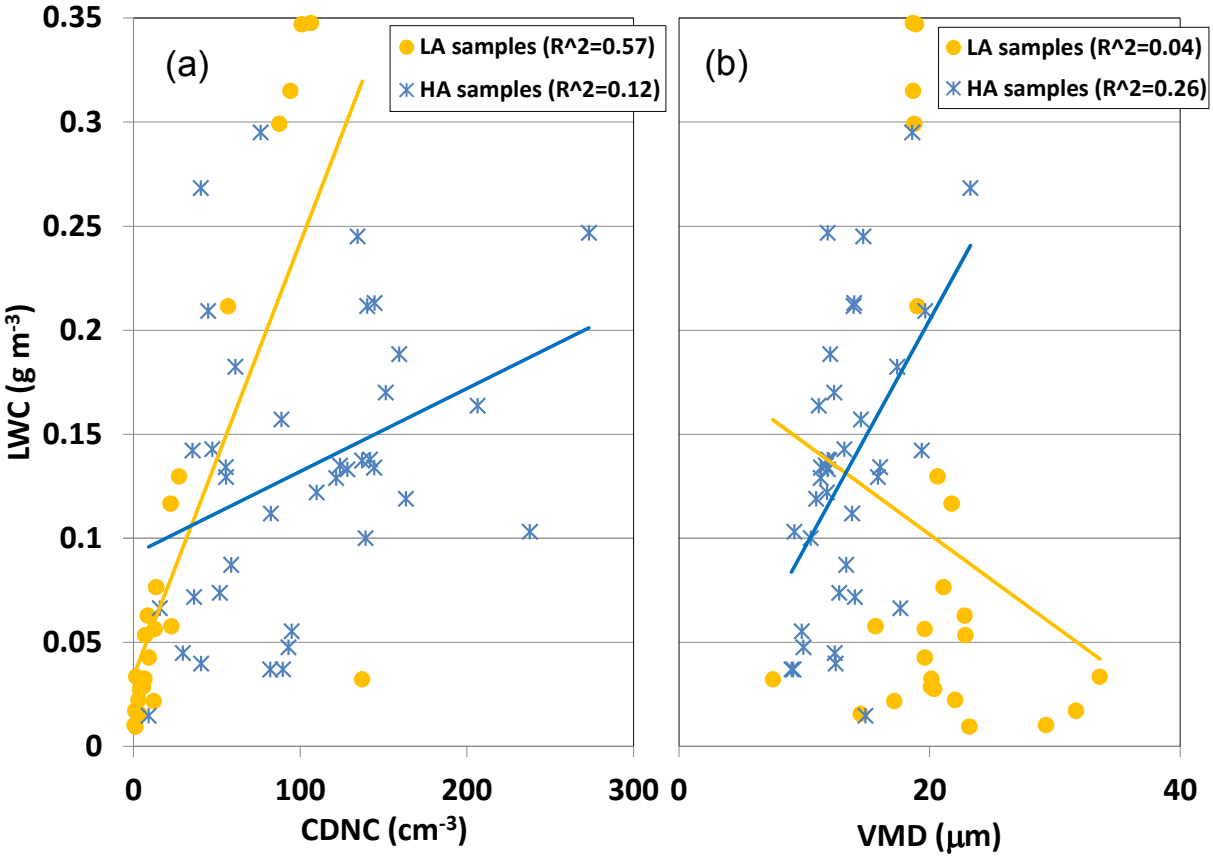



Figure 7. The LWC plotted as a function of the CDNC (a) and VMD (b) for the LA (orange) and HA (blue) samples. Linear regressions for each of the LA and HA samples are also plotted, and the coefficients of determination are given in the legends.

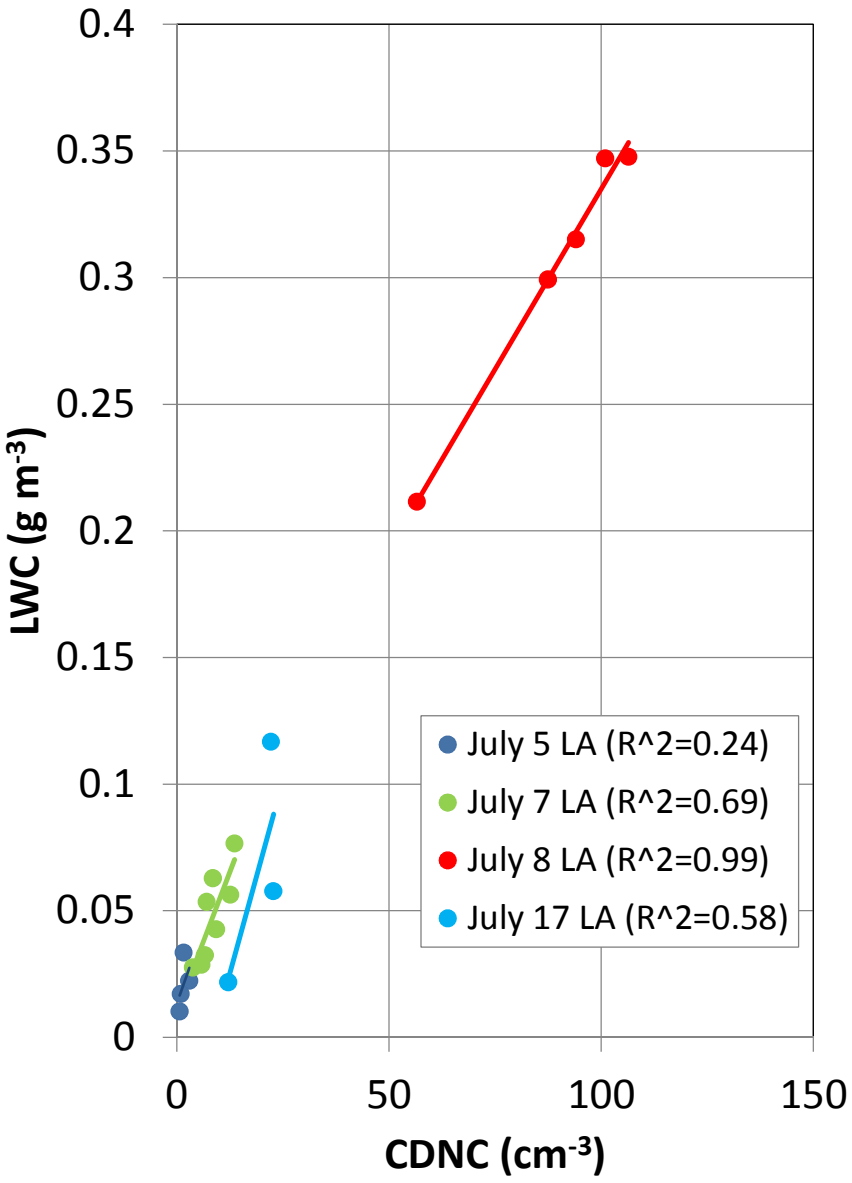



Figure 8. As in Fig. 7a, but identifying the specific LA cases of July 5, 7, 8 and 17. Linear regressions for each set of samples are also plotted, and the coefficients of determination are given in the legends. The slopes are significant at a 95% confidence level within ±30% for July 7 and within 60% for July 8. The slopes in the July 5 and 17 cases are not significant at a 95% confidence level.

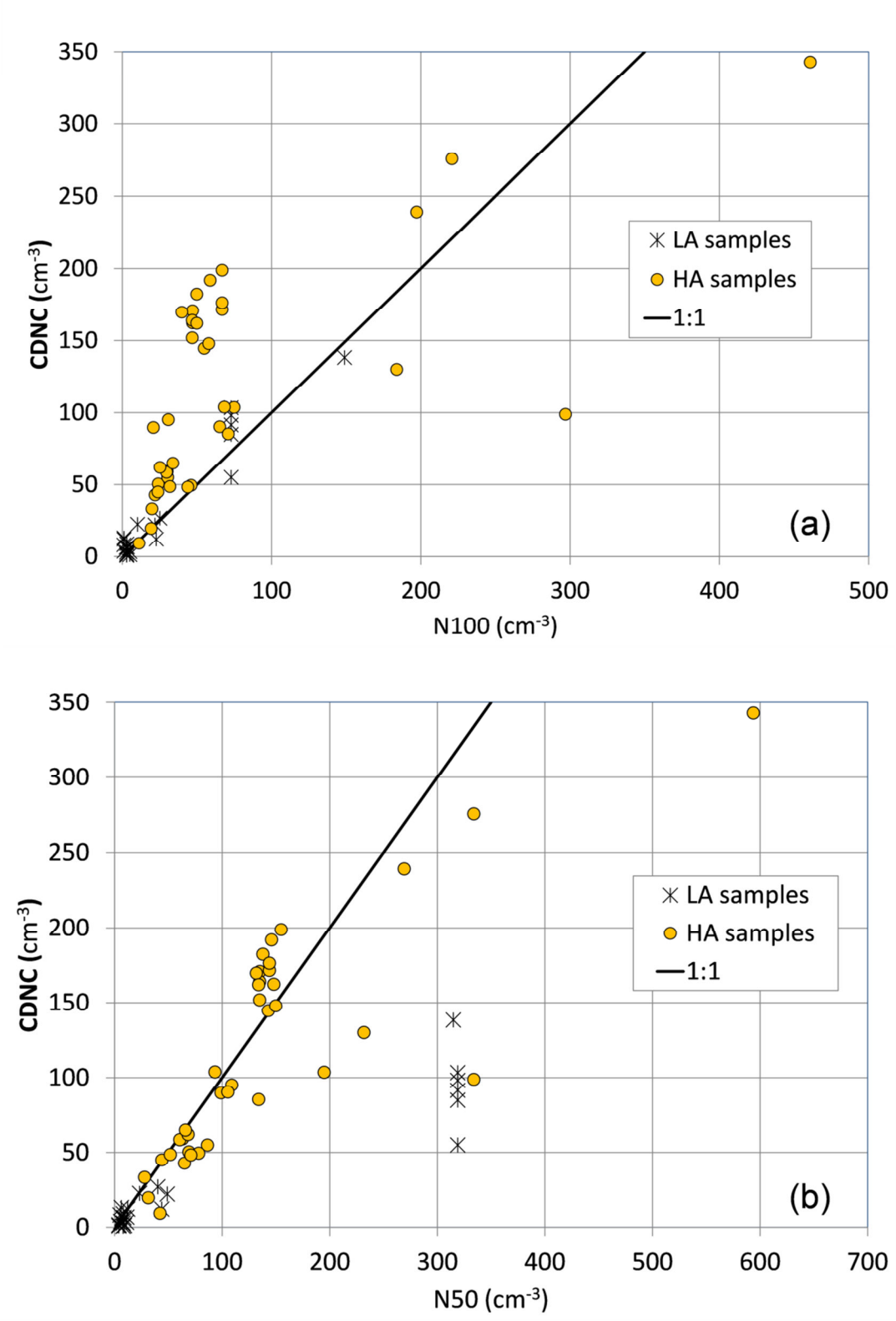



Figure 9. Plots of CDNC versus a) N100 and b) N50. Points are identified between LA (yellow) and HA (black asterisk) samples, and the 1:1 lines are for reference.

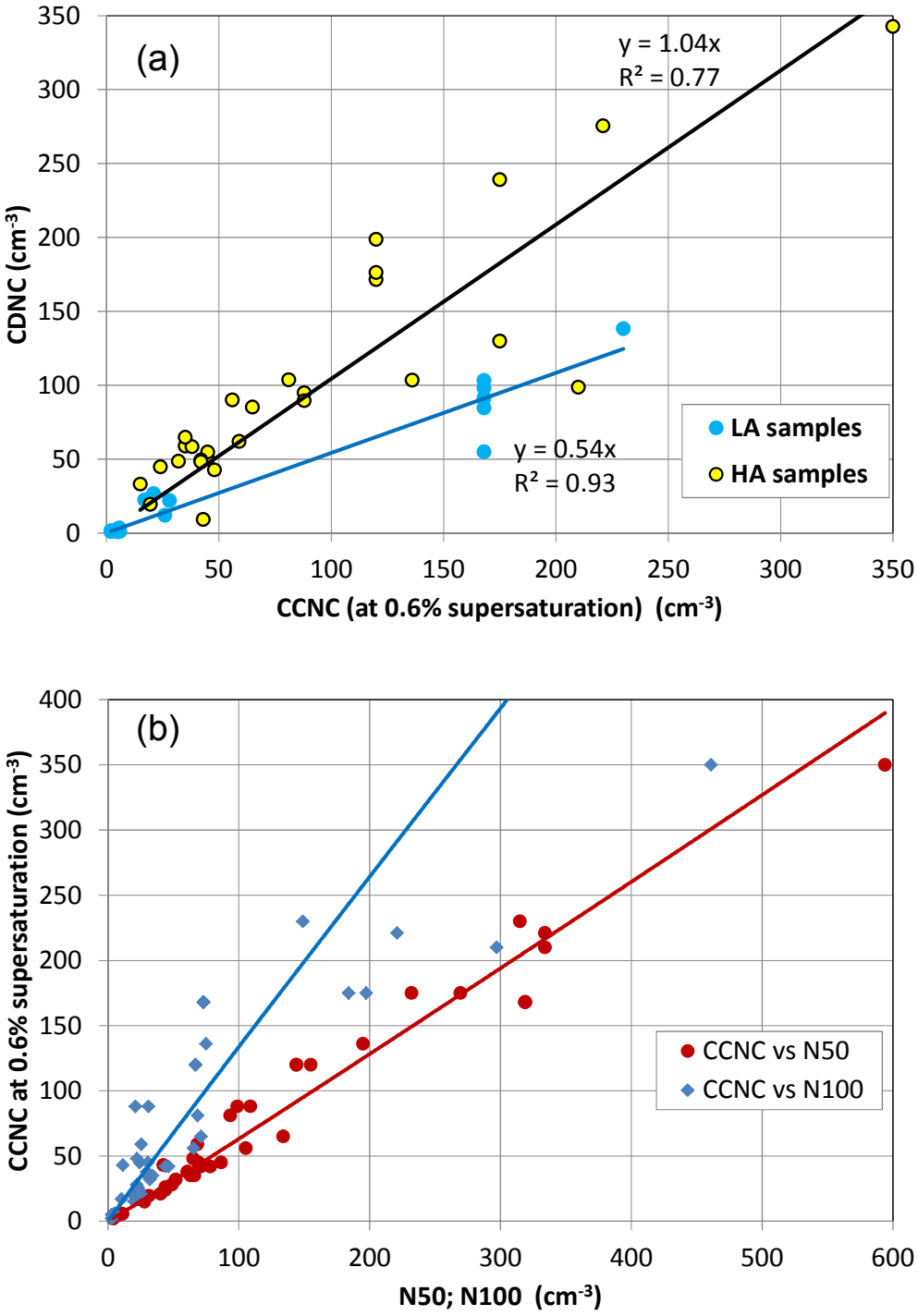
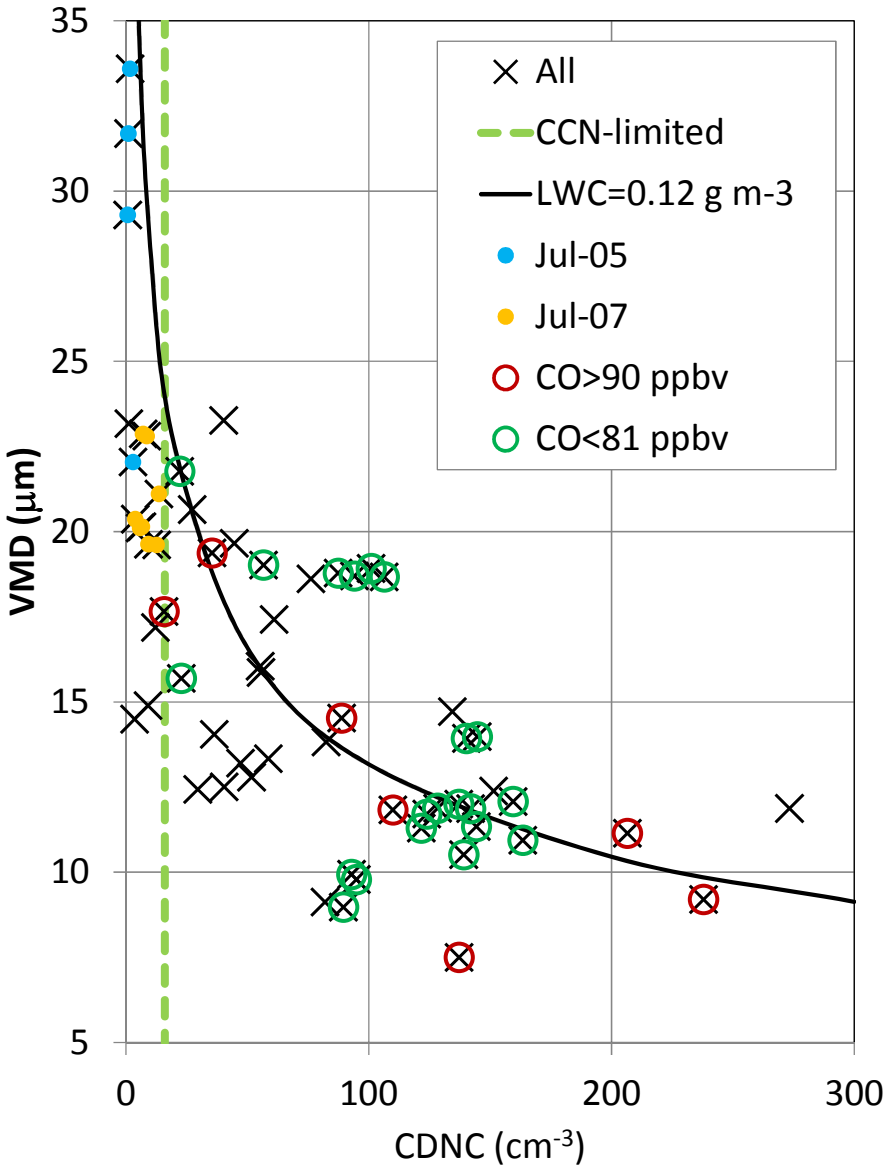



Figure 10. a) CDNC plotted versus the CCNC measured at 0.6% supersaturation; points are identified between LA (yellow) and HA (blue) samples, and linear regressions through the origin are shown; the CCNC(0.6%) points are limited to 44 of the 62 total, due to problems with the CCN measurement; the 44 are split 16 and 28 between LA and HA,. b) CCNC(0.6%) (44 points) plotted versus N50 and N100; power law fits to each are provided for reference.

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