- 1 Mesoscopic surface roughness of ice crystals pervasive across
- 2 a wide range of ice crystal conditions
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7 Abstract

8 Here we show high-magnification images of hexagonal ice crystals acquired by

9 Environmental Scanning Electron Microscopy (ESEM). Most ice crystals were grown and
10 sublimated in the water vapor environment of an FEI-Quanta-200 ESEM, but crystals grown

11 in a laboratory diffusion chamber were also transferred intact and imaged via ESEM. All of

12 these images display prominent mesoscopic topography including linear striations, ridges,

- 13 islands, steps, peaks, pits, and crevasses; the roughness is not observed to be confined to
- 14 prism facets. The observations represent the most highly magnified images of ice surfaces yet

15 reported and expand the range of conditions where the rough surface features are known to be

- 16 conspicuous. Microscale surface topography is seen to be ubiquitously present at temperatures
- 17 ranging from -10°C to -40°C, at super-saturated and sub-saturated conditions, on all crystal

18 facets, and irrespective of substrate. Despite the constant presence of surface roughness, the

19 patterns of roughness are observed to be dramatically different between growing and

20 sublimating crystals, and transferred crystals also display qualitatively different patterns of

- 21 roughness. Crystals are also demonstrated to sometimes exhibit inhibited growth in
- 22 moderately supersaturated conditions following exposure to near-equilibrium conditions, a
- 23 phenomena interpreted as evidence of 2-D nucleation. New knowledge of the characteristics
- 24 of these features could affect the fundamental understanding of ice surfaces and their physical
- 25 parameterization in the context of satellite retrievals and cloud modeling. Links to
- supplemental videos of ice growth and sublimation are provided.
- 27

281Introduction

29 It has been broadly recognized (IPCC 2007; Bony et al. 2006) that cloud-climate feedbacks

30 are the most weakly constrained radiative forcings for general circulation models of a

changing future climate. Among the variety of cloud feedbacks that have been identified and 1 2 parameterized, cirrus clouds are notable for being especially uncertain — even the sign of the likely feedback effect is subject to debate (Mitchell et al. 2008; Burkhardt 2011). Baran 3 (2012) presents a compelling case that the complex radiative scattering properties of 4 5 hetereomorphic ice crystals must be accounted for to reach a physically consistent parameterization of ice clouds in climate models. In addition to cloud-climate feedbacks, it is 6 7 expected that prominent mesoscopic topography (defined here as variation scale between 100 8 nm and 5 µm) would affect active research questions in stratospheric ozone chemistry 9 (McNeill et al. 2007) and thunderstorm electrification (Dash et al. 2001). Despite recognition 10 that the microscale structures of ice are critically important to consider, very few images of 11 ice are available at this resolution.

While many experimental research efforts have characterized the effects of temperature,
supersaturation, and pressure on particle-scale ice crystal growth rates and morphology under

conditions relevant to atmospheric processes (Lamb and Hobbs 1971; Kuroda and Lacmann,
1982; Bailey and Hallett 2004; Nelson and Knight 1998; Magee et al. 2006), to date the limits
of light microscopy have prevented a thorough analysis of the surface details of ice crystals

17 beyond the microscale. Several studies have employed interference techniques or ellipsometry

to suggest the presence of surface structure and disorder of ice (Bryant et al. 1960; Furukawa
et al. 1987; Sazaki et al. 2010), but no highly-resolved images of mesoscopic surface structure

1) et al. 1907, Suzaki et al. 2010), out no inginy resorved images of mesoscopic surface structure

have been available. Early SEM studies examined formvar replicas of ice crystals, and in
some instances were successful in revealing crystal structures beyond the resolution of light

22 microscopy (Shaefer 1941; Truxby 1955; Kuroiwa 1969, Goodman et al. 1989). The use of

23 chemical etchants also revealed new information on the nature of lattice defects in ice and

24 provide tantalizing images that are reminiscent of the some of the mesoscopic structures seen

25 in the present work (Truxby 1955; Kuroiwa 1969). Despite the availability of sub-micron

26 details that can be preserved by proper application of formvar replication, several authors

27 have pointed out that the effects of the solvents on the surface are hard to define (e.g.

28 Takahashi 1988). The first scanning electron micrographs of non-replicated ice were made

29 from platinum sputter-coated natural snowflakes by Wergin et al. (1995); their striking

30 images show a variety of complex structures, but due to warm collection conditions and the

31 sputtering processes, they are ambiguous with respect to presence of mesoscopic surface

32 structure (Wergin et al. 1995). Recently, Neshyba et al. (2013), Ulanowski et al. (2014),

33 Petersen et al. (2011), Pfalzgraff et al. (2010), Kuhs et al. (2010), and Zimmerman et al.

(2007) have used uncoated, in-situ growth of ice crystals in ESEM or variable-pressure SEM 1 2 to identify surface roughness in the form of ridges or "trans-prismatic" strands. Through images resolved at up to 10,000x magnification (compared with previous maximum 3 magnifications of $\sim 1000x$), our observations underscore this newly emerging view of the ice 4 5 surface, demonstrating that mesoscopic surface roughness is a non-uniform condition present on a wide array of ice crystals, and is not confined to a narrow range of macroscopic 6 7 morphology, substrate, temperature, humidity, or growth rates. 8 Over fifty years have passed since the first account of a relationship between surface 9 roughness and specular reflectivity (Bennett and Porteus, 1961). A growing recognition is 10 now emerging that divergence away from the classical smooth-facet assumption for ice will 11 affect the way that light is scattered. A broad suite of recent measurements and models 12 describe the potential effect of surface roughness and complex micro-scale geometry in 13 smoothing the peaks of the ice-scattering phase function (Baran 2012; Baum et al. 2010; 14 Baum et al. 2011; Cotton et al. 2010; Heymsfield et al. 2006; Ulanowski et al. 2006; 15 Ulanowski et al. 2010; Ulanowski et al. 2012; Um and McFarquhar, 2011). Despite acknowledged uncertainty regarding the details of real roughened ice surfaces, a 16 17 variety of recent studies have concluded that cirrus radiative measurements and models come 18 into better agreement when attempting to account for complicated shapes and rough ice 19 surfaces. For example, Kahnert et al. (2008) estimated that the effect of varying 20 parameterizations of ice surface microphysics can affect the modeled radiative influence of cirrus by a factor of two. Baum et al. (2011) recently determined that assumption of a 21 22 roughened ice surface results in better fits to ice particle data retrievals from the CALIOP 23 lidar instrument on board the NASA-CALIPSO A-train satellite. Polarized ice-cloud 24 reflectances have also been shown to have high sensitivity to ice particle geometry and texture (Chepfer et al. 2001; Baran and Labonnote 2006). Polarized ice-cloud reflectances measured 25 26 by PARASOL have been found to agree best with a mixed-habit model of severely-roughened 27 ice particles (Cole et al. 2013) and retrievals indicate the greatest average roughness in ice 28 clouds associated with tropical deep convection (Cole et al. 2014). Mauno et al. also point 29 out discrepancies between measured and modeled cirrus-modulated shortwave radiative 30 fluxes that would be improved by assumption of surface roughness on ice crystals (Mauno, 31 2011). In addition to passive satellite measurements of cirrus ice microphysics, it is also likely 32 that ice scattering functions assumed in radar and lidar imaging of cirrus and mixed-phase 33 cloud properties would be affected (Sun et al. 2011). Because satellite observations, aircraft

1 measurements, and modeling of cirrus and mixed-phase clouds have often been at odds

2 (Baran 2012; Garrett and Gerber 2003; Gierens et al. 2003; Kramer et al. 2009; Harrington et

3 al. 2009; Jensen et al. 2009; McFarquhar et al. 2007; Peter et al. 2006; Connolly et al. 2007),

4 potential artifacts of surface roughness on both satellite retrievals and modeling of ice

5 microphysics must be carefully evaluated.

6 2 Methods

We employed an FEI Quanta-200 FEG environmental scanning electron microscope (ESEM) to observe ice crystals at temperatures from -10° C to -45° C and through a wide range of under-saturated, equilibrium, and super-saturated vapor pressures. Most experiments examined in-situ nucleation, growth, and sublimation of ice crystals in the pure water vapor environment of the ESEM. Several experiments also included imaging of ice crystals grown in an external diffusion chamber and then transferred into the ESEM.

13 **2.1 In-situ growth and sublimation**

14 In-situ growth and sublimation experiments were conducted on the ESEM stage, with the ice 15 nucleating on a substrate mounted to a Peltier cooling block. An aluminum stub fitted to the 16 Peltier block was machined to expose a 1 mm diameter surface to the chamber environment 17 such that the entire surface could be contained in the ESEM field of view at low 18 magnification. An insulating vinyl mask was used to ensure that ice growth was limited to the 19 intended cold surface. Growth and sublimation experiments were conducted on bare 20 aluminum as well as on a variety of thin substrates that were attached to the aluminum by 21 thermally conductive epoxy. Substrates that were tested included stainless steel, aluminum, 22 and copper, as well as mineral substrates including Covellite, Muscovite, Bismuth, Magnesite, 23 Galena, and Quartz.

24 Once the substrate was mounted onto the Peltier stage, the ambient air in the ESEM chamber 25 was evacuated and back-filled with water vapor, after which the ice growth experiments were 26 initiated. After cooling to a temperature at or above -45°C, ice nucleation, growth, and 27 sublimation was controlled through direct adjustment of the water vapor pressure or surface temperature. Vapor pressure was controlled at 0.5 Pa increments between 0 and 500 Pa via 28 automatic differential pumping. Vapor measurements were made to 0.1 Pa resolution. In a 29 typical experiment, vapor pressure was first adjusted to the equilibrium frost point and 30 subsequently increased in small increments until ice crystal nucleation was observed. A 31

magnification of 100x was employed until ice crystal growth became visible, after which 1 2 variable resolutions (as high as about 10,000x) were used in to capture detailed images and movies of the growing ice crystals. Maximum image acquisition size for the FEI Quanta200 is 3 4 4096 x 3775 pixels (~15.5 MP). At 2500x magnification, this results in a single pixel with 5 dimension 25.4 nm x 25.4 nm. At 2500x magnification, we observed that typical imaging conditions resulted in linear features that could be resolved at limits of 2-3 pixels (50-75 nm). 6 7 With increasing magnification, signal-to-noise reduction partially (but not completely) offset 8 resolution gains, such that we estimate a resolution limit of approximately 25 nm for these 9 methods.

10 In a typical experiment, the differential pumping and thermoelectric cooling of the substrate 11 required approximately 10 minutes to reach equilibrium values of vapor pressure near 65 Pa 12 and -25°C. Temperature of the Peltier block is automatically reported through the FEI 13 instrument software to 0.1°C resolution and the water vapor pressure is reported to 0.1 Pa 14 resolution. Based upon drop freezing measurements, and trials of substrate-free growth, we 15 are confident that vapor pressure values were precisely and accurately controlled and reported; however, we determined that the surface temperature of the substrate was typically 16 several degrees warmer (+0.5 to +6.5 °C) than the temperature indicated for the Peltier base. 17 This offset was determined by observing the (equilibrium) vapor pressure at which ice neither 18 19 grew nor sublimated and inferring surface temperature from the Murphy and Koop vapor pressure formulation (2005); we observed this thermal offset to vary based on the 20 21 temperature, the thickness of the substrate, and its thermal conductivity. Additional detail for 22 the calibration of temperature offset and calculated saturation ratio is provided in Appendix 23 A. Following the approach to equilibrium, vapor pressure was raised in 0.5 Pa increments until nucleation of one or several ice crystals occurred. It was frequently challenging to 24 25 nucleate a single crystal, even with incremental increases in vapor pressure. Once a crystal of 26 interest was developed, it was examined and photographed at a variety of magnifications, 27 ranging from 100x to 10000x and at a variety of vapor densities, including super- and sub-28 saturated conditions, and through multiple cycles of growth and sublimation.

29 2.2 Transported Crystals

In order to image crystals grown in the presence of background air pressure and to minimize
the role of the growth substrate, we also developed a new technique to grow, capture, and
transfer ice crystals from a controlled diffusion chamber into the ESEM. Ice crystals were

grown in a diffusion chamber contained within a large-volume, ultra-low temperature freezer. 1 2 The full freezer volume is held at -55 ° C and all walls are coated with ice to achieve a saturated environment. The diffusion chamber generates a small region of supersaturation 3 along a thin vertical fiber between two parallel, horizontal ice plates, with the ice surface 4 5 temperatures maintained by thermoelectric modules. Ice crystals were grown at ambient lab pressure (~1000 hPa), and at low supersaturation (<110% RH_i), with temperature near -50° C, 6 7 usually with many crystals in close proximity along the fiber. Within the saturated freezer, 8 several dozen of these ice crystals were then captured in a pre-chilled small-volume (~1 cm³) 9 containment cell. The transfer and sealing process from diffusion chamber to cryo-cell 10 requires just a few seconds, and intentionally avoids any exposure to ambient room air. The 11 base of the containment cell was formed from an aluminum stub to allow direct transfer into 12 the ESEM cooling stage. Upon capture, the cell was sealed closed and transferred to a 13 specially-designed cryogenic dewer (filled with liquid nitrogen or crushed dry ice) for 14 transport to the ESEM. The cell was then removed from the dewer and quickly placed onto the pre-chilled cooling stage of the ESEM. Throughout the capture and transfer process, the 15 primary objective was to allow ESEM imaging of the ice surfaces as they had been growing 16 17 in the diffusion chamber and without additional sublimation or growth. To prevent unintended 18 sublimation, the small-volume containment cell was filled with many crystals and quickly 19 sealed so that equilibrium vapor pressures would preserve the crystal surfaces. The 20 containment cell was also enveloped by materials with significant heat capacity and the cryo-21 dewer designed and handled to maintain near isothermal conditions. Once the sealed 22 containment cell was transferred to the ESEM cooling stage, the microscope chamber was 23 evacuated and adjusted to equilibrium vapor pressures matching the cold stage and 24 containment cell temperature. After reaching equilibrium vapor pressure in the ESEM 25 chamber, the containment cell was then opened using electronically driven stage movement to 26 mechanically pull off the top seal. Following unsealing of the containment cell, ESEM 27 imaging could proceed as in the in-situ growth and sublimation experiments.

28

29 3 Results and discussion

The overriding impression of our experiments suggests that surface structures on the 0.1 to 10 micron scale are a ubiquitous feature of ice crystal facets. The morphology of many of these structures agree with observations recently reported by Neshyba et al. (2013), Ulanowski et

al. (2014), Pfalzgraff et al. (2010), Petersen et al. (2011), and Zimmerman et al. (2007). Our 1 2 observations provide new evidence of roughening on basal crystal facets, as well as images and videos at high magnifications (1000x -10000x) that reveal smaller scale roughening that 3 is not readily apparent at magnification below 1000x. Our observations also show strongly 4 5 different morphology between textures of growing surfaces as compared to sublimating surfaces. The results also include images of ice crystals grown in air-vapor mixtures and then 6 7 transferred into the ESEM. These images do show mesoscopic surface roughness, but suggest 8 that the presence of air or different modes of internal heat transfer may significantly affect the 9 character of surface texture development.

10 **3.1 Growing Crystals**

As described in section 2.1, ice crystals where nucleated and grown on an ESEM cold state at 11 12 a variety of temperatures and pressures. Crystals typically nucleated at just a few percent 13 above equilibrium vapor pressures and continued to grow steadily in proportion to the 14 magnitude of ice supersaturation. At high supersaturations >150% RH_i, nucleation and growth proceeded so quickly that it was difficult to isolate and follow the progression of a 15 16 single crystal, as the entire stage would be overtaken by intersecting crystals within a few 17 seconds. Therefore, most growth experiments occurred at modest supersaturation, usually 18 105-125% RH_i. While we did observe clear surface morphology differences between growing 19 and sublimating crystals, we did not detect a systematic dependence on the degree of 20 supersaturation, the rate of growth, nor the composition of the underlying substrate.

21 **3.1.1** Roughness morphologies and scales

In our ESEM growth experiments, we observed mesoscopic linear striations on prism faces (Figure 1e-g) similar to those described by Neshyba et al. (2013) and Pfalzgraff et al (2010) as well as a variety of other mesoscopic roughness structures. Figure 1 provides a multi-panel view of characteristic growing surface features across the range of our roughness observations. The panels are organized with temperature decreasing toward the bottom and magnification increasing to the right of the figure. The magnification, temperature, pressure, and humidity information from the databars are available for easier viewing in the appendix

29 table A.1.

30 Particularly in panels 1a, 1c, 1d, 1f, and 1g, distinct roughness can be observed on the basal

31 facets of growing crystals as well as the prism facets. The basal plane roughness usually

1 appears less linear and less symmetrically organized than the prismatic strands. Image

2 sequences (typically around 1 frame/s) reveal the dynamic progress of roughening features as

3 they migrate across the ice surface (videos available in supplemental data, see especially

4 BasalRoughnessLayers.mp4 and SingleCrystal.mp4). Figure 1 panels 1g-1i also display

5 evidence of microfaceting on basal and prism facets. Microfaceting could be induced by

6 cycles of growth and sublimation at all temperatures, but was only observed to develop during

7 steady growth at temperatures below -35°C.

8 These data have also confirmed the idea that surface roughness can be enhanced in proximity

9 to a grain boundary between two neighboring crystals (e.g. Pedersen et al. 2011). In panel 1b,

10 it is apparent that the ridges (in the centers of the images) become larger just as the two

11 advancing facets collide, with ridges radiating outward from the impact point. A video of

12 ridge topographic intensification precipitated by colliding ice crystals is available in the

13 supplementary material (ImpactWave.mp4). It is possible that this effect might enhance

14 roughness in aggregates of ice particles or in cloud regions with high collision frequencies,

15 both of which have been found to be particularly common near deep convection, a region that

16 has also been associated with satellite retrievals indicative of severe particle roughening (Cole

17 et al. 2014). Blackford (2007) and Pedersen et al. (2011) also point out that the

18 microstructure of ice along grain boundaries can play an important role in advancing the

19 understanding of the mechanical properties of snowpacks that are susceptible to avalanche, as

20 well as the dynamics of glaciers.

21 **3.1.2 Magnification and contrast effects**

22 Figure 2 shows a four-panel plot of a growing hexagonal ice crystal starting at 762x 23 magnification in panel a. This crystal was grown near -21°C and is being held near 24 equilibrium in this sequence of panels; the crystal was not visibly growing or sublimating across the 57 seconds separating the first and final panel. Mesoscopic topography is not 25 26 visible in panel a, so the magnification was increased to 2155x (panel b), which made 27 roughness apparent on the prism facets of the ice crystals. However, the basal facet is tilted 28 almost perpendicular to our viewing angle; therefore, we could not see whether any complex 29 surface architecture existed on the basal facet. By increasing the magnification further and adjusting the contrast and brightness of the ESEM, panel c shows that rough surface 30 31 topography was actually present on the basal facet (at 3625x magnification). A measurement 32 was made of the depth of a prominent terrace in panel d (1.59 microns); this terrace is among the thickest observed during experiments, suggesting that most topographic features have heights well below 1 micron. This pattern was typical of all experiments; only the most prominent surface structures were typically visible at a magnification of 500x-1000x, while closer examination at 1000-5000x would reveal smaller-scale structures and topography on surfaces that had not been resolved at lower magnification, or at sub-optimal aspect, brightness, and contrast.

We attempted to extract profiles of surface height from our ESEM micrographs in to calculate
the roughness measure <r> recently defined by Neshyba et al (2013) in the context of
carefully measured mesoscopic striations observed on ice prism facets growing in VPSEM at
-45°C:

11
$$r = 1 - \sqrt{\frac{1}{1 + \left(\frac{dy}{dz}\right)^2}}.$$
 (1)

12 The average roughness <r> is determined along the measured profile. In several circumstances, we succeeded in retrieving an approximate profile, but, as indicated by 13 14 Ulanowski et al. (2014), in most routine SEM imaging circumstances, we found that it was 15 not straightforward to extract this height profile with confidence. Where this profile could be 16 estimating on growing crystals imaged at magnifications similar to those analyzed by 17 Neshyba et al. (~300-500x magnification), we retrieved similar values of roughness near <r> 18 ≈ 0.05 . Height profiles along several sections of more highly magnified crystals indicated by 19 red rectangles in panels 1f and 1i yielded <r> between .10 and .40. Height profiles, and thereby <r> values, can be retrieved by careful attention during the imaging process to SEM 20 21 stage and crystallographic orientation, resolving of crystal edges, or potentially by 3-D 22 reconstruction of equilibrium crystals captured from multiple angles. We found that it was 23 usually not possible to determine height profiles when structures are visible, but near the limit 24 of the image resolution, or where pronounced roughness did not intersect a resolved crystal 25 edge, a common feature in our observations. We also find that the *<*r> metric, much like the 26 subjective perception of roughness, is affected by image magnification and resolution, and in-27 turn by the minimum sampling interval of the height profile. If, as our observations imply, 28 surface roughness is significant at sub-micron scales that are not well-resolved at 500x 29 magnification, <r> values obtained from profiles retrieved at 500x may be underestimating 30 total roughness.

3.1.3 Inhibited growth observations

2 As expected, with growth experiments occurring on large, rough substrates, we observed that 3 most crystals nucleated at low supersaturation and then grew steadily and at rate in proportion to ice supersaturation values. At temperatures below -30°C, we repeatedly observed crystals 4 where growth became completely stalled, even at moderate supersaturations (>115% RH_i) 5 6 that had previously induced growth in that crystal and continued to lead to growth in adjacent 7 crystals. This stalled growth was only observed following a specific cycle of humidity 8 adjustment: the excess vapor supply of a steadily growing crystal was gradually reduced until 9 reaching equilibrium, with observable growth ceasing, and with no sublimation apparent. This equilibrium condition was held for 1-5 minutes, after which the vapor pressure was gradually 10 11 increased (or temperature decreased) until RH_i exceeded equilibrium by a few percent. In 12 many instances, the original crystal would not resume growth, even though adjacent crystals continued to nucleate and grow. Figure 3 provides time-separated panels illustrating one such 13 event were a crystal previously held at equilibrium failed to grow following a 0.3°C decrease 14 15 in temperature, despite several nearby crystals nucleating and growing (please see 16 supplemental videos Inhibited1.mp4, Inhibited2.mp4, and Inhibited3.mp4 for additional 17 examples). We interpret this an example of side-by-side ice crystals subject to 2 different 18 surface conditions: a) one where steady growth continues at emergent dislocations or 19 stacking faults, likely partly induced by the underlying substrate and b) one where a 20 previously rough surface has been reconditioned by the momentary maintenance of 21 equilibrium vapor pressure, leaving the surface to grow only by 2-D nucleation requiring 22 vapor in excess of a critical supersaturation. The stalled crystal surfaces do not appear to be 23 completely absent of mesoscopic roughness, nor are they observably different than the 24 growing surfaces, implying that the surface condition differentiating separate growth 25 mechanisms is determined at a smaller scale than observable here. The observation of growth 26 at dislocations and 2D nucleations would not be unprecedented -- Sazaki et al. (2010) used 27 interference-contrast microscopy to demonstrate depositional growth in ice via both 2-D layer 28 nucleation and spiral growth steps at screw dislocations. Several recent studies have also 29 suggested that ice up to 243 K does not have a well ordered hexagonal crystal structure. 30 Instead, ice can contain a mixture of cubic and hexagonal sequences which can give rise to 31 roughness on the prismatic faces (Malkin et al., 2012; Kuhs et al. 2012). It has also been 32 shown that the proportion of cubic sequences decreases as ice is heated and the ice tends 33 towards perfect hexagonal ice (Murray and Bertram, 2006; Kuhs et al., 2012). Both of these

1 findings are consistent with the steady growth we observe at low supersaturation as well as

2 the occurrence of inhibited growth when the surface is given the opportunity to anneal

3 emergent stacking faults.

4 **3.2** Sublimating Crystals

5 In many instances, cycling of vapor pressure was conducted to observe the sensitivity of surface topography to ambient humidity. In the portions of the cycle below equilibrium vapor 6 pressure, a significantly different character to the surface roughness was observed. Instead of 7 8 regular or spreading ridges, plateaus, and steps, we observed concave, scalloped depressions 9 developing away from the original surface. In the case of poly-crystalline examples (panel 10 4a), the scalloped depressions took on especially dramatic shapes near former grain 11 boundaries, with sharp peaks often evident during advanced sublimation. However, even 12 single-crystal examples did produce marked scalloping (panels 4b-d), which often initiated at 13 the site of roughness produced during previous growth (see supplemental video 14 GrowandSublime.mp4). It appears that the sublimation occurs outward from multiple centers, often at the site of a former ridge or ledge. In fact, on the basal plane, the sublimation can 15 even be seen to produce hexagonal pits (visible in GrowandSublime.mp4 at ~14s) and the 16 17 scalloping develops partly from the intersection of multiple spreading pits. Furthermore, if the 18 supersaturation was once again increased above equilibrium, the crystal would typically 19 exhibit micro-faceting that initiated along the sharp ridges bounding adjacent sublimation-20 scallops. The scale of the scalloping can become fairly large (e.g. panel 4a), so it would be surprising that these structures wouldn't be seen by optical microscopy if they were indeed 21 22 characteristic of sublimation in diffusion-limited regimes—we speculate that this mode of 23 sublimation may be unique to the kinetic attachment regime, and perhaps significant only for 24 very small ice atmospheric ice particles.

25 **3.3 Transported Crystals**

As described in section 2.2, ice crystals where grown at low supersaturation, at -50° C, and at ambient lab pressure in an external, freezer-based diffusion chamber and subsequently captured and transported to the ESEM cold stage in a sealed small-volume cell, with conditions maintained at ice/vapor equilibrium until imaging could commence. The goal of this test was to compare the character of surface roughness observed in experiments described in 3.1 and 3.2 with ice crystals grown in conditions more closely approximating cirrus clouds.

While not completely isolated like an ice crystal floating in air, the crystals grown in the 1 2 freezer-chamber grew outward from a fiber, differentiating them from the close substrate contact seen with ESEM-grown crystals. The presence of large partial pressures of Nitrogen 3 and Oxygen provides the other important departure from ESEM-grown crystals. It is 4 5 important to note that crystals grown within the chamber (Figure 1-5) are firmly within the attachment kinetics growth regime due to the absence of air and therefore low resistance to 6 7 diffusion. By contrast, the transported crystals were grown at high pressure to fairly large 8 size, representing growth that was within the diffusion-limited regime. The particle-scale 9 habits and aspect ratios are in agreement with measurements by Bailey and Hallett (2004). 10 The habits are qualitatively more elongated than the nearly isometric crystals typically 11 observed in ESEM. The surfaces themselves (Figure 5) clearly exhibit signatures of 12 mesoscopic roughening that are similar to examples of roughness seen in ESEM-grown 13 crystals. However, for most transported crystals, the crystal edges are more intricate that those 14 observed in ESEM and portions of surfaces of some transported crystals do appear smooth 15 even at magnification greater than 1000x, a rare observation in our ESEM-grown crystals. 16 While we took efforts to maintain equilibrium conditions between capture and imaging, we 17 still cannot say with certainty that the crystals were not exposed to some variation in RH_i 18 during transport. The transported crystals hint at some significant differences in roughness 19 morphology, but they do provide evidence that crystals grown in air/water mixtures and with 20 minimal support also exhibit mesoscopic roughness with similarity to that observed in ESEM-21 grown crystals.

22

23 **4.** Conclusions

The ESEM images of ice in section 3 confirms the observations on the nature of mesoscopic 24 roughness recently described by several studies (Neshyba et al. 2013, Ulanowski et al., 2014; 25 26 Pfalgraff et al. 2010, Pedersen et al. 2011, Zimmerman 2007) and also reveals additional 27 patterns of roughness morphology at high magnification, across a wide range of temperatures, 28 on crystal basal facets, sublimating crystals, and crystals grown in air/water vapor mixtures. 29 Light microscopy has generated highly-detailed images of ice crystals (Libbrecht, 2005; 30 Walden et al., 2003), but a combination of working distance constraints, diffraction limits to resolution, and transmitted light illumination have conspired to prevent visible imaging of 31 32 mesoscopic surface features. We suggest that submicron scale roughness can also be obscured

to detection by ESEM imaging, where the probability of resolving mesoscopic surface 1 2 topography depends on a combination of factors including the size and depth of the features themselves, the magnification and resolution of the micrograph, the brightness and contrast, 3 and the orientation of the ice crystal with respect to the viewing angle. When brightness and 4 5 contrast are adjusted to optimal levels, increasing the magnification and resolution serves to create a detailed portrait of ice crystal mesoscopic surfaces which in turn can increase the 6 7 calculated roughness measure <r>. Furthermore, high magnification images near 10,000x 8 (Figure 6) depict significant roughness on spatial scales below 200 nm, indicating that for 9 modeling of roughness-induced scattering changes, it is not sufficient to consider light 10 scattering only in the geometric optics regime.

11 Because the ice crystals shown here were grown both inside the pure vapor environment of 12 the ESEM and in external air/vapor mixtures, the ubiquitous presence of mesoscopic 13 roughness indicates that the roughening mechanisms are not caused by the ESEM 14 environment alone and are likely to be significant in atmospheric ice. Despite strong 15 similarity in the subjective appearance of roughness morphologies in crystals grown in diffusion-limited vs. attachment kinetics regimes, there do also appear to be significant 16 17 differences in overall crystal habit, intricacy of crystal edges, as well as some difference in the patterns of mesoscopic roughness. To increase the utility of these observations, these 18 19 differences should be investigated further and analyzed quantitatively.

20 The current microscopy observations appears to be well-aligned with a growing body of evidence (Baran 2012; Baum et al. 2011; Ulanowski et al. 2006; Yang et al. 2008; Cole et al. 21 22 2013) suggesting that measured scattering from ice crystals fit a rough surface model more successfully than crystals with presumed smooth-faceted surfaces. Several of the satellite 23 24 studies also point out that while roughened ice particles are strongly indicated by radiative 25 retrievals, these results have not been able to diagnose the physical morphology or scale of the apparent roughness (Cole et al. 2013; Ulanowski et al. 2014) and that further progress will 26 27 depend on a combination of in-situ observations and laboratory measurements, such that realistic representations of surface roughness can be integrated into scattering models. 28 29 We believe that these new observations of prevalent surface topography in ice crystals

30 warrant careful consideration in the scattering models that are used for satellite retrievals of

31 cirrus ice microphysics, and in turn, affect the radiative modeling of cirrus clouds in climate

32 models. Furthermore, the ubiquity of a complex mesoscopic landscape on the surface of ice

- 1 crystals also has potential wide-ranging impacts to theories of charge transfer in
- 2 thunderstorms, the heterogeneous chemistry of stratospheric ozone, and the sintering of ice
- 3 crystals in snowpacks and glaciers. We suggest that the next steps should focus on efforts to
- 4 examine and quantify roughness in crystals transported from cirrus-analog environments and
- 5 development of a mixed air/vapor capability for ice crystal growth in ESEM.

1 Appendix A.

2 This appendix is intended to provide additional detail regarding thermodynamic conditions

3 and temperature calibration processes that have been used to infer approximate saturation

4 ratios. Appendix Table A1. also displays the magnification and thermodynamic data

5 measured in the image panels from Figures 1-6.

6 Vapor pressure measurements within the chamber were observed to be repeatable to within 7 approximately 0.4 Pa. Triple-point measurements and ice growth without substrates also 8 showed no discernible systematic error in pressure readings. Reported temperature 9 measurements are made by a thermocouple imbedded in the Peltier cooling sub-stage, several 10 mm below the substrate surface. The surface was consistently observed to be several degrees 11 warmer than the sub-stage thermocouple reading. The magnitude of this difference was not constant – it was observed to vary between +0.6 $^{\circ}$ C and +6.5 $^{\circ}$ C. The magnitude of 12 13 temperature offset appeared to be increased by the following factors: lower overall 14 temperature, increased substrate thickness, lower substrate thermal conductivity, weaker 15 coolant flow through Peltier block, and shorter thermal equilibration time. Despite this variation, in each experiment, once a working temperature was established, the vapor pressure 16 17 could be gradually increased to induce ice nucleation and growth and then gradually decreased until growth ceased and sublimation was first observed – this balance point was set 18 19 as the equilibrium vapor pressure and used to calculate the inferred surface temperature based 20 on the Murphy and Koop (2005) formulation for saturated vapor pressure over ice:

21 $\ln e_i = 9.550426 - 5723.265/T + 3.53068 \ln(T) - 0.00728332 T$ (A1)

22 where e_i is the saturation vapor pressure over ice in [Pa] and T is the ice temperature in [K].

23 Because most experiments involved growth and sublimation induced by vapor pressure

24 adjustments, the variable experimental saturation ratios could be determined relative to the

25 pre-established equilibrium point. In several instances, it was possible to detect slight

26 temperature drift over an hour-long experiment, which could introduce errors of several tenths

27 of a degree in calculated temperatures. Based on these considerations, along with (much

smaller) uncertainty in vapor pressure, the calibrated surface temperatures are deemed to be

29 valid +/- 0.3 °C and ice saturation ratio calculations are approximately +/- 4 % confidence.

30 Introduction of additional thermometry points will be undertaken in future work in order to

- 1 reduce uncertainty in surface temperatures and saturation ratio, enabling quantitative
- 2 modeling of growth and sublimation rates.

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Increasing magnification



- 2 Figure 1: Image panels at a variety of magnification, temperatures, and pressures -- all
- 3 showing various examples of mesoscopic surface topography. Red rectangles in panels 1f
- 4 and 1i show sites used for calculating roughness measure <r>. Data detail available in
- 5 appendix table A.1.
- 6



- 1 2
- 3 Figure 2. Demonstration of increased magnification effect upon perceived roughness is
- 4 shown here, as well as a measurement of ridge depth (panel d). Data detail available in
- 5 appendix table A.1.
- 6



Figure 3: Series of frames separated by ~10s (20 s between panel d and e as the acquisition was re-centered to capture growing crystals at bottom-left in panel d and e). The original 2 crystals at equilibrium in panel a. do not grow when subjected to a temperature decrease of 0.3° C (equivalent to ~105 RH_i), with vapor pressure held constant. Data detail available in appendix table A.1.



Figure 4: Sublimating ice crystals displaying scalloped depressions and sharp ridges and
peaks of roughness. Panel a. was originally composed of a polycrystalline ice particle, with
peaks and ridges prominent at former grain boundaries. Data detail available in appendix table
A.1.



Figure 5: Transported ice crystals, showing portions of 4 different crystals grown at -50 °C in a static diffusion chamber between 100 and 105% RH_i. The crystals were maintained in cryogenic equilibrium during transport into the ESEM for imaging under vapor-matched saturation conditions. Data detail available in appendix table A.1.

1



Figure 6: Ice crystals are observed at high magnification, the highest we were able to attain without sacrificing the quality of the image. Inset image (1903x magnification) shows the location of the high-magnification zoom, located on the basal facet of a growing crystal, and near to the intersection of the adjoining prism facet with another crystal's basil facet. Data detail available in appendix table A.1.

| Panel | Magnification | Vapor Pressure (Pa) {+/- 0.5 Pa} | Sub-stage Temp. (°C) {+/- 0.1 °C } | Calibrated surface Temp. (°C) {+/- 0.3 °C } | Ice Saturation Ratio {+/- 4%} |
|----------|---------------|----------------------------------------|------------------------------------------|---------------------------------------------------|-------------------------------------|
| Figure 1 | | | | | |
| а | 350 | 73.3 | -25.0 | -24.0 | 1.05 |
| b | 694 | 73.3 | -25.0 | -24.0 | 1.05 |
| С | 1061 | 75.2 | -25.0 | -23.9 | 1.06 |
| d | 834 | n/a | n/a | n/a | n/a |
| е | 1559 | 29.1 | -37.6 | -33.8 | 1.14 |
| f | 2249 | 25.3 | -39.2 | -35.2 | 1.13 |
| g | 669 | 25.3 | -40.6 | -35.5 | 1.20 |
| h | 1031 | 22.9 | -41.0 | -36.0 | 1.14 |
| i | 2453 | 22.9 | -41.0 | -36.0 | 1.14 |
| Figure 2 | | | | | |
| а | 762 | 93.6 | -25.0 | -22.0 | 1.00 |
| b | 2155 | 93.2 | -25.0 | -22.0 | 1.00 |
| С | 3625 | 93.6 | -25.0 | -22.0 | 0.99 |
| d | 3625 | 93.6 | -25.0 | -22.0 | 1.00 |
| Figure 3 | | | | | |
| а | 2858 | 29.5 | -37.5 | -32.6 | 1.02 |
| b | 2858 | 29.5 | -37.8 | -32.9 | 1.05 |
| С | 2858 | 29.5 | -37.8 | -32.9 | 1.05 |
| d | 2858 | 29.1 | -37.8 | -32.9 | 1.04 |
| е | 2858 | 29.5 | -37.8 | -32.9 | 1.05 |
| f | 2858 | 29.5 | -37.8 | -32.9 | 1.05 |
| Figure 4 | | | | | |
| а | 519 | n/a | n/a | n/a | n/a |
| b | 927 | 29.1 | -36.5 | -32.0 | 0.94 |
| С | 669 | 25.3 | -39.0 | -33.5 | 0.96 |
| d | 1532 | 27.6 | -40.0 | -33.0 | 0.99 |
| Figure 5 | | | | | |
| а | 228 | 40.1 | -30.0 | -29.4 | 1.00 |
| b | 673 | n/a | n/a | n/a | n/a |
| С | 1671 | 50.4 | -30.0 | -27.3 | 1.00 |
| d | 766 | 60.0 | -29.3 | -25.5 | 1.00 |
| Figure 6 | | | | | |
| а | 10533 | 84.0 | -25.0 | -22.4 | 1.05 |

Table A.1 Magnification, temperature, vapor pressure, and calculated saturation ratios for
images figures.