1 **Title:** Drone data reveal heterogeneity in tundra greenness and phenology not captured by 2 satellites

3

4 Authors: Jakob J. Assmann<sup>1</sup>, Isla H. Myers-Smith<sup>2</sup>, Jeffrey T. Kerby<sup>3</sup>, Andrew M. Cunliffe<sup>4</sup>, 5 Gergana N. Daskalova<sup>2</sup>

6

0			
7	Affiliations:	<sup>1</sup> Department of Biology, Aarhus University, DK	
8		<sup>2</sup> School of GeoSciences, University of Edinburgh, UK	
9		<sup>3</sup> Aarhus Institute of Advanced Studies, Aarhus University, DK	
10		<sup>4</sup> Department of Geography, University of Exeter, UK	
11			
12	ORCID:	Jakob J. Assmann:	0000-0002-3492-8419
13		Isla H. Myers-Smith:	0000-0002-8417-6112
14		Jeffrey T. Kerby:	0000-0002-2739-9096
15		Andrew M. Cunliffe:	0000-0002-8346-4278
16		Gergana Daskalova:	0000-0002-5674-5322

17

#### Abstract: 18

## 19

20 Data across scales are required to monitor ecosystem responses to rapid warming in the 21 Arctic and to interpret tundra greening trends. Here, we tested the correspondence among 22 satellite- and drone-derived seasonal change in tundra greenness to identify optimal spatial 23 scales for vegetation monitoring on Qikiqtaruk - Herschel Island in the Yukon Territory, 24 Canada. We combined time-series of the Normalised Difference Vegetation Index (NDVI) 25 from multispectral drone imagery and satellite data (Sentinel-2, Landsat 8 and MODIS) with 26 ground-based observations for two growing seasons (2016 and 2017). We found high 27 cross-season correspondence in plot mean greenness (drone-satellite Spearman's p 28 0.67-0.87) and pixel-by-pixel greenness (drone-satellite R<sup>2</sup> 0.58-0.69) for eight one-hectare 29 plots, with drones capturing lower NDVI values relative to the satellites. We identified a 30 plateau in the spatial variation of tundra greenness at distances of around half a metre in the 31 plots, suggesting that these grain sizes are optimal for monitoring such variation in the two 32 most common vegetation types on the island. We further observed a notable loss of 33 seasonal variation in the spatial heterogeneity of landscape greenness (46.2 - 63.9%) when 34 aggregating from ultra-fine-grain drone pixels (approx. 0.05 m) to the size of medium-grain 35 satellite pixels (10 - 30 m). Finally, seasonal changes in drone-derived greenness were 36 highly correlated with measurements of leaf-growth in the ground-validation plots (mean 37 Spearman's p 0.70). These findings indicate that multispectral drone measurements can 38 capture temporal plant growth dynamics across tundra landscapes. Overall, our results 39 demonstrate that novel technologies such as drone platforms and compact multispectral 40 sensors allow us to study ecological systems at previously inaccessible scales and fill gaps 41 in our understanding of tundra ecosystem processes. Capturing fine-scale variation across 42 tundra landscapes will improve predictions of the ecological impacts and climate feedbacks of environmental change in the Arctic. 43

44

Arctic tundra, vegetation monitoring, landscape phenology, satellite, drones, Keywords: 45 46

UAV and RPAS, NDVI, scale

### 47 Introduction

48

Identifying the scales at which ecological processes operate is a fundamental, yet often neglected element of ecological research (1–3). Cross-scale ecological information can inform our understanding of the causes and consequences of global change (2). In tundra ecosystems, vegetation responses triggered by rapid Arctic warming could influence secosystem functions through altered carbon and nutrient cycles with potential feedbacks to the global climate system (4–8). Yet, challenging logistics have limited the extent of field-based observations in Arctic ecosystems (9–11). The grain sizes of global-extent satellite products (tens of meters to kilometres) are too coarse to capture the fine-scale dynamics of tundra plants (12–14) and to link vegetation change to key ecosystem functions (13). Thus, by bridging this "scale-gap", we can transform our understanding of pan-Arctic tundra vegetation change and associated global-scale climate feedbacks.

60

## 61 Satellites show greening of the tundra

62

63 Satellite observations indicate a 'greening' of tundra ecosystems (13,15–20) and shifts in 64 growing season phenology over recent decades (21–24). Observations of increasing tundra 65 greenness are often reported from surface-reflectance-derived Normalised Difference 66 Vegetation Index (NDVI) (16,18,25,26). Satellite-observed tundra greening has occurred 67 concurrently with ground-based observations of vegetation change in Arctic ecosystems (27) 68 including increased shrub cover (28–31) and taller community level plant height (32), as well 69 as earlier leaf emergence and flowering at some (33–36), but not all tundra sites (37–39). 70 However, mismatches between ground and satellite-based observations suggest the 71 potential for an observational scale gap (13).

72

## 73 Arctic vegetation change and phenology have been linked to warming

74

Satellite-observed Arctic greening trends have been linked directly to warming air temperatures (19,20,40–46) and indirectly to sea-ice declines (17,47–51). Ground-based observations of tundra vegetation change correspond with warming (27,32,52), but do not always co-occur with satellite greening trends in the regions around the ecological monitoring sites (13,53). While satellite-based phenology observations from the Arctic have been mainly linked to temperature (22,54,55), *in situ* phenology in the tundra has been shown to be influenced by a suite of interacting factors rarely tested in satellite-based analysis of Arctic phenology. These factors include, but are not limited to: snowmelt, temperature, day length, and the proximal influences of sea-ice on localised climate affect (34–36,38,56,57). Thus, ecological studies indicate greater complexity of drivers than analyses of satellite-derived greening trends to date.

86

87 Inconsistencies amongst satellite platforms and heterogenous greening trends

88

Greening trends and phenology measures derived from different satellite platforms do not always correspond with each other (13,18). Additionally, satellite-derived greening trends vary at global (18), continental (42,58–60) and regional scales (46–48,61–64). Many areas

92 of the Arctic show no trends in NDVI, with only around 20% of the Arctic spectrally greening

93 and around 1 - 4% of the Arctic spectrally browning (13,62,65,66). Recent analyses suggest 94 a slowdown of the Arctic-wide spectral greening trend over the past decade (43,67). 95 Furthermore, despite NDVI being related to the photosynthetically active biomass in the 96 tundra (14,68–70), geophysical, environmental and ecological factors, such as low solar 97 angle, atmospheric effects (including cloud and fog), snow cover, soil moisture and standing 98 water, in addition to the non-linearity of NDVI-biomass relationships, complicate the 99 interpretation of satellite-derived NDVI time-series at high latitudes (13,71). The growing 100 complexity highlighted in Arctic greening trends has led to repeated calls for ground 101 validation of satellite observations (11,18,59,60,66,72,73).

102

### 103 The scale discrepancy problem in Arctic greening

104

105 A major problem in linking satellite-derived trends of tundra spectral greenness and 106 phenology to *in situ* observations of ecological processes is the discrepancy in observational 107 scales (13,29,61,72,74). Satellite datasets with long-term records are limited by their 108 moderate- to coarse-grain sizes, ranging from 30 m (Landsat) to 250 m (MODIS) and 8 km 109 (AVHRR-GIMMS3g). *In situ* ecological monitoring in the Arctic is logistically challenging and 110 therefore restricted in extent to a limited number of sites and often metre-squared plots 111 (10,75). Only a few studies have linked on-the-ground vegetation or phenology change to 112 satellite trends in NDVI in Arctic tundra (13,14,47,48,53,76–78). However, drones equipped 113 with compact sensors now allow for the collection of ultra-fine-grain multispectral imagery at 114 landscape extents that can potentially bridge the scale-gap between satellite and 115 ground-based observations (14,79–82).

116

### 117 Novel drone data to study variation in greenness

118

119 Here, we set out to test whether drones can be used to identify the key ecological scales for 120 studying tundra greenness on Qikiqtaruk in the Canadian Arctic by bridging the scale gap 121 between satellite and in situ data. First, we tested whether satellite- and drone-derived 122 measures of mean landscape-scale greenness (NDVI) agree across two growing seasons 123 while controlling for the potentially confounding effects of topography and land cover. 124 Second, we identified the key spatial scales for ecological variation in landscape greenness 125 within the two most common vegetation types at our study site using variograms. Third, we 126 tested how the magnitude of seasonal variation in tundra greenness scales across grain 127 sizes from fine-resolution drone imagery to medium-grain satellite imagery. Finally, we 128 assessed whether drone-derived NDVI corresponds with on-the-ground measures of within growing season change in plant growth based on methods frequently used by long-term 129 130 field-based monitoring networks. Thus, in our analysis we validated satellite-derived landscape estimates of vegetation greenness with ultra-fine-grain drone data and described 131 spatial and temporal variation in tundra productivity at landscape extents (1-100 ha) with 132 grain sizes that were previously not accessible. 133

134

### 135 Methods

- 136
- 137 Site description: Qikiqtaruk Herschel Island
- 138

Qikiqtaruk (69.57 N, 138.91 W) is located in the Beaufort Sea along the coastline of the 139 140 North Slope of the Yukon Territory, Canada. The vegetation is characteristic moist acidic 141 shrub tundra (83) found in the Western Arctic regions of North America that has experienced 142 strong spectral greening in recent decades (13). The two most common plant communities 143 on the island are the tussock sedge ("Herschel") and Dryas-vetch ("Komakuk") vegetation 144 types (84,85). The tussock sedge vegetation is dominated by the name-giving tussock sedge 145 Eriophorum vaginatum L. with varying cover of Salix pulchra Cham. The top-soils of the 146 island are underlain by ice-rich permafrost and undergo frequent disturbance (85). The Dryas-vetch vegetation is particularly found on ground disturbed by soil creep and is 147 148 characterised by the near ubiquitous presence of Dryas integrifolia Vahl., the willow Salix arctica Phall., various grass species including Arctagrostis latifolia. (R.Br.) Griseb. and forb 149 species (86). The relative abundances of these species are shown in (Figure S1). Though 150 151 the two vegetation types are specific to the region, these plant communities would group 152 with tundra types S1, W2 and G3/4 of the Circumpolar Arctic Vegetation Map (87).

153

We established four study areas on the east end of the island, each with two co-located one-hectare plots in the two key vegetation cover types (Figure 1, Table S1). We selected plots with homogenous terrain and land cover to represent the two key vegetation types and to control for the potentially confounding effects of terrain and cover heterogeneity. The island harbours small herds of caribou (100s of individuals) and muskox (3 - 35 individuals in recent years) of fluctuating size, as well as cyclic populations of voles and lemmings (88). We estimate the overall impact of herbivory on the vegetation in our study plots to be low particularly in 2016 and 2017 when there were few muskox on the island.

162

### 163 Multispectral drone time-series

164

165 We analysed a total of 62 drone surveys from 21 dates; see Table S2 for a breakdown by 166 one-hectare monitoring plots. We collected multispectral drone imagery using Parrot Sequoia (Paris, France) compact multispectral sensors mounted on multi-rotor drone 167 168 platforms in June to August in 2016 and 2017. We used three different drone platforms: a Tarot 680 Pro hexacopter with camera sensor stabilisation in 2016, and a 3DR Iris+ and a 169 170 DJI Phantom 4 Pro without sensor stabilisation in 2017. Surveys were flown using parallel flight lines (a lawn-mower flight pattern) at an altitude of ca. 50 m, giving ground-sampling 171 distances of 0.04 m to 0.06 m. Images were acquired with 75% front- and side-lap as close 172 as possible to solar noon (mean absolute difference to solar noon 2.16 h, maximum 6-7 h). 173 See Table S2 and the methods section of the Supplementary Materials for further details on 174 175 the drone surveys, including additional information on radiometric calibration, as well as 176 temporal and spatial coverage.

177

We processed the Sequoia imagery using Pix4D Mapper v4.0.21 (Lausanne, Switzerland) with the *agMultispectral* template and the 'merge map tiles' option set to true to generate co-registered single-band surface reflectance maps. Radiometric calibration was carried out in Pix4D Mapper using pre- or post-flight imagery of calibrated reflectance panels; in 2016 we used a MicaSense (Seattle, USA) panel and in 2017 a SphereOptics (Herrsching, Germany) Zenith Lite panel. We measured panel reflectance pre- and post- season and used the mean values for radiometric calibration. We also calibrated for sensor properties and sun irradiance measured by the incident light sensor. We used four to six ground control points per survey precisely geolocated with a GNSS system to spatially constrain the reconstructions in Pix4D Mapper with an estimated accuracy of 1-2 pixels between bands and 2-6 pixels between surveys (81). We calculated the Sequoia NDVI as the normalised difference between the near-infrared (770 nm – 810 nm) and red (640 nm – 680 nm) bands of sensor.

191

192 Satellite time-series

193

194 Satellite time-series were obtained from three different satellite sensors: 1) the Moderate 195 Resolution Imaging Spectroradiometer (MODIS) on the USGS Terra satellite, 2) the 196 Multispectral Instrument (MSI) on Sentinel-2 A & B and 3) the Operational Land Imager (OLI) 197 on Landsat 8.

198

199 We obtained MODIS NDVI values for the time period from the 1st May to the 30th of 200 September in 2016 and 2017 for all 250 m MODIS pixels that contained the survey plots. 201 NDVI values were retrieved from the 16-day MOD13Q1 v6 Terra product (89) using the 202 Google Earth Engine (90). We discarded all values with a 'Summary QA' score of -1 (no 203 data) or 3 (cloudy). Table S3 lists the resulting MODIS-pixel-date pairs. The MODIS NDVI is 204 calculated as the normalised difference between bands 1 (841 nm – 876 nm) and band 2 205 (620 nm – 670 nm).

206

For the Sentinel-2 time-series, we gathered all Sentinel-2 MSI L1C scenes containing the tile covering Qikiqtaruk (T07WET) that were available on the Copernicus Open Access Hub (<u>https://scihub.copernicus.eu/</u>) for the same time period as the MODIS pixels. We processed all scenes to L2A using Sen2Cor 2.4.0 (91), retained all bands with 10 m resolution (2-4 & 8), applied the cloud mask and generated a true-colour image. We inspected all scenes visually and discarded all imagery with cloud contamination over the study area (78% of scenes for 2016 and 74% of scenes for 2017). The resulting set contained nine cloud-free Sentinel-2 L2A scenes of the study area from 2016 and fifteen scenes from 2017 (Table S4). Finally, the Sentinel NDVI was calculated as the normalised difference between band 8 (784.5 nm - 899.5 nm) and band 4 (650 nm - 680 nm).

217

218 Landsat 8 OLI Level-2 (surface reflectance) time-series were obtained using the USGS EarthExplorer website (https://earthexplorer.usgs.gov/) by guerying the search engine for all 219 220 scenes that covered the study site during the same time-period as the MODIS pixels (n = 94). The automatically generated cloud masks were of insufficient guality, so we manually 221 222 inspected all scenes and retained only the scenes cloud-free over the study site (n = 7 for 223 2016, n = 8 for 2017, Table S5). The Landsat 8 NDVI was then calculated as the normalised 224 difference between band 5 (845 - 885 nm) and band 4 (630 - 680 nm). The study plots were not designed with a Landsat 8 analysis in mind and did not naturally coincide with the 225 Landsat grid. We therefore calculated subsequent one-hectare plot NDVI averages as a 226 weighted mean, where each pixel was weighted by the proportion of the plot area covered 227 228 by the pixel.

229

230 Ground-based plant phenology measurements

232 We carried out ground-based phenology monitoring in eight 2 m x 2 m plots (Table S6), one adjacent to each one-hectare plot (mean distance = 23 m, max distance = 52 m). We placed 233 234 the ground-based monitoring plots adjacent to the drone-based survey plots to minimise the 235 effects of ecological disturbance and trampling in the drone-based survey plots caused by 236 the repeat visits necessary for the ground-based monitoring. Within these plots we monitored six individual plants from the most common species: E. vaginatum, D. integrifolia, 237 S. pulchra and A. latifolia in tussock sedge tundra; D. integrifolia, S. arctica and A. latifolia in 238 Dryas-vetch tundra. We measured the length of the longest leaf on each individual on the 239 240 survey date to the nearest millimetre. This approach is widely used in field-based phenology monitoring protocols (92), and will allow for NDVI to be directly related to phenological 241 242 changes in plant traits. We conducted ground-based surveys in tandem with the 243 drone-based surveys where logistical possible, resulting in a dataset of 52 drone and ground survey pairs spread over 20 different dates. The majority of ground-based phenology 244 surveys were carried out on the same day as the drone surveys (mean difference = 0.3245 days, maximum difference = 3 days, Table S7). 246

247

#### 248 Cross-sensor correspondence

249

250 To test cross-sensor correspondence, we first had to scale all datasets to a shared spatial grain and time-window. To achieve this, we first plotted the spatial mean NDVI for all 251 one-hectare plots, time-points and available sensors (MODIS = single pixel, Landsat 8 = 252 weighted mean) across both growing seasons (2016 and 2017). We then divided the two 253 growing seasons into 22 consecutive seven-day blocks starting on the 1st of May each year. 254 Next, we calculated the temporal mean of the spatial mean NDVI for each seven-day block 255 256 for all plot and sensor combinations where data was available. We then identified all matching seven-day block and study plot combinations for each drone-satellite and 257 satellite-satellite combination. We then tested cross-sensor correspondence by calculating 258 Spearman's rank correlation and mean sensor-to-sensor difference in the plot means across 259 260 the whole data set.

261

262 Additionally, we matched all drone and Sentinel-2 scenes, as well as all drone and Landsat 8 scenes that were less than two days apart. We resampled the red and near-infrared drone 263 bands to the relevant Sentinel-2 / Landsat 8 grids and calculated the NDVI. We restricted the 264 analysis to Landsat 8 pixels fully contained within the study plots and reprojected the drone 265 data from UTM 7N to UTM 8N using a bilinear reprojection where the Landsat 8 scenes 266 were provided in this projection. Finally, we tested the predictive relationship between the 267 resampled drone and satellite NDVI pixel-pairs for a random subsample of Sentinel pixels 268 (10% of total, n = 700) and all available Landsat 8 pixels (n = 198) with Bayesian linear 269 models (Table S8 and S9 for Sentinel-2, S10 and S11 for Landsat 8) using the MCMCglmm 270 271 v.2.29 package (93).

272

We used the 'resample' function of the R raster package v. 3.0-12 (94) for resampling from finer to coarser resolutions. The function first aggregates the smaller grid to the largest clean divisor of the larger grid using the mean and then, if required, resamples to the larger grid using bilinear interpolation. We also tested an alternative resampling approach by first

277 resampling to a common resolution and grid of 0.5 m and then aggregating by mean, but 278 found no qualitative differences in our results (Figure S2). Further details about software and 279 package versions used for raster manipulations and visualisations can be found in the 280 Supplementary Materials.

281

282

#### 283 Spatial autocorrelation

284

285 To assess the spatial autocorrelation of variation in tundra greenness within the eight plots, 286 we sampled variograms and fitted variogram models using the gstat v. 2.0-5 package (95,96). First, we pre-thinned the acquired drone-data by randomly sampling 5% of the ca. 4 287 million pixels of each orthomosaic. We then sampled variograms for all plots at the peak of 288 289 the 2017 season (26 and 28 July) and fitted variogram models, letting the gstat algorithm 290 select the best fit amongst spherical, exponential and Matern models. The only exception was Area 3 for which the closest available complete set of flights was on the 18th July 2017. 292 To test conformity of the variograms across the season, we repeated the analysis for the 293 surveys from the 26 June and 9 August 2017 for Area 1 and 2. No change in the variogram patterns were observed across the 2017 season and we therefore assume that our analysis 294 is representative for the 2016 season also. All variograms were sampled with a bin width of 295 0.15 m from 0 to 15 m and a bin width of 3 m from 0 to 45 m. 296

297

#### 298 Grain size and phenology

299

We tested the influence of grain-size on observations of tundra greenness phenology by fitting simplified growing season curves to the raster stacks for each plot and season. We first resampled the drone bands for all time-points to grids with grain sizes of 0.5, 1, 5, 10, 20 and 33.33 m. We then calculated the NDVI and fitted simple quadratic models to each pixel in the growing season stacks ( $y = ax^2 + bx + c$ , where x is the day of year and y the pixel NDVI, a the quadratic coefficient, b the linear coefficient and c the constant term). We found a strong negative correlation between the quadratic and linear coefficients of the models (Figure S6), we selected only the quadratic coefficient for further analysis. Further details on model choice and analysis can be found in the method section of the Supplementary Materials.

310

#### 311 Ground validation

312

To test the correspondence between our ground-based phenology measurements and the drone observations, we derived time-series of the plot mean standardised longest leaf length for all species (using a z-score – centred data with a standard deviation of 1) and the drone-greenness for each 2 m x 2 m ground-based monitoring plot. See supplementary methods for details on how the leaf measurements were standardised. The drone-based plot mean NDVI values were then matched with the plot mean standardised longest leaf length values from the closest ground-based survey date (Table S7). We then calculated the Spearman's rank correlation between mean NDVI and mean longest standardised leaf length for each plot and season. We replicated the analysis using Sentinel-2 data where available (see Supplementary Materials). Finally, we also conducted a by-species version of 323 the analysis using the by-species mean of the absolute longest leaf measurements for each 324 2 m x 2 m plot rather than the mean based on the standardised longest leaf measurements.

325

### 326 Results

327

## 328 Landscape greenness corresponded among sensors

329

Landscape greenness corresponded well among drone, Sentinel-2, Landsat 8 and MODIS across both the 2016 and 2017 growing seasons. Growing season curves of the plot mean NDVI were similar (Figure 1) and the seven-day block plot mean NDVI values were highly correlated across sensors (Spearman's  $\rho > 0.59-0.98$ , Table S12). However, we observed a positive offset between drone and satellite plot-mean NDVI for the seven-day block means between 0.027 (Landsat 8) and 0.073 (Sentinel-2B) absolute NDVI that was consistent across satellite platforms (Table S13). The Landsat 8 offset of 0.027 fell within the range of the estimated error associated (±0.03) with the drone-derived plot mean NDVI for the study plots determined previously with the same survey method (81).

339

340 Resampled drone pixels (10 m and 30 m) and the corresponding spatially co-located Sentinel-2 and Landsat 8 pixels were highly correlated (marginal  $R^2 = 0.69$  and marginal  $R^2$ 341 342 = 0.58 respectively, see Figure 2 and Table S8 and S10). We found that vegetation type, the 343 time-difference between satellite scene and drone data acquisition, and the specific Sentinel 344 platform (Sentinel-2A / Sentinel-2B) influenced the relationship between Sentinel-2 pixel 345 NDVI and drone-derived NDVI (marginal  $R^2 = 0.87$  see Table S9). While the Sentinel platform (Sentinel-2A / Sentinel-2B) had the strongest impact on the intercept and slope of 346 347 the linear model, vegetation type and time-difference mainly influenced the slope, with 348 time-difference having the smallest effect on slope and intercept overall (Table S9). In 349 contrast, we only detected a statistically meaningful difference in days between satellite and 350 drone scene acquisition for the Landsat 8 - drone pixel model (marginal  $R^2 = 0.70$ ); vegetation type did not have a statistically meaningful influence on this relationship (Table 351 S11). 352

353

## 354 Spatial variation in landscape greenness peaked at approx. 0.5 m

355

356 Spatial variability in the NDVI values associated with distance peaked at ranges below 0.5 meter (mean range 0.44 m) during the peak-season of 2017 (26-28 July), and little additional 357 autocorrelation structure in the NDVI was found between pixel pairs for distances of up to 45 358 m (Figure 3). This pattern was consistent across vegetation types in seven out of our eight 359 plots (Figure 3, S3 and S4). The only exception is the Dryas-vetch plot in Area 3, which 360 showed the same patterns for distance below 10 m, but thereafter spatial variation steadily 361 increased (Figure S4). Peak variability (sill) in NDVI decreased as the growing season 362 progressed (Figure S5), and varied with vegetation type (Figure 3, S3, and S4). Unexplained 363 variability (nugget) was consistently low across all Areas (Figure 3, S3, and S4). 364

365

366

### 367 Seasonal-variation was lost when aggregating to medium grain sizes

We observed a notable loss in the amount of seasonal variation in tundra greenness when aggregating grain size from ultra-fine-grain drone to medium-grain satellite data. The loss was particularly pronounced at grain-sizes above 10 m – the grain size of Sentinel-2 MSI pixels (46.2 - 63.9%) (Figure 4). The variation in the quadratic coefficient of the simple growing season curves (Figure 4b and S6) decayed logarithmically with grain size (Figure 4a), while no change occurred in the mean tendency of the coefficient (Figure S7). The quadratic and linear coefficients of the growing season curves were strongly correlated (Spearman's  $\rho$  = -0.999), thus the same pattern holds true for the linear component of the curve fit (Figure S6).

378

# 379 Drone-derived spectral greenness correlated well with leaf measurements

380

381 Drone-derived spectral greenness correlated well (mean  $\rho = 0.70$ ) with ground-based measurements of cross-season phenology for graminoids and deciduous plants (Figure 5). 382 The Spearman's correlation coefficient of the plot mean standardised leaf length and the 383 mean drone-derived NDVI values in the ground-based phenology plots (mean  $\rho = 0.70$ , 384 Table S14 and Figure 5) matched the by-species analysis based on absolute leaf lengths 385 (mean  $\rho = 0.68$ , Table S15 and Figure S9). The graminoids and deciduous shrub species 386 followed this mean tendency well across all time-series, while the partially-evergreen D. 387 integrifolia showed mixed responses between plots and years (mean  $\rho = 0.22$ , Figure S9). 388 The drone-based greenness time-series of the 2 m x 2 m ground-phenology plots highlight 389 390 fine-scale differences in phenology such as the continuous greening of tussocks that was visible at the tussock sedge tundra plot in Area 2 (Figure 5c). Sentinel-2 greenness showed 391 slightly weaker correlations (mean  $\rho = 0.58$ , Figure S10) with the plot mean standardised 392 leaf length, but no combined Sentinel-2 and ground-based phenology time-series of 393 sufficient length were available for 2016 and peak-season observations in 2017 were limited. 394 395

## 396 Discussion

397

398 Our analysis of cross-scale time-series of landscape greenness on Qikiqtaruk highlights four main findings: 1) Measures of mean tendency in landscape greenness were consistent 399 400 across sensors, but drone-derived NDVI values were lower than those from Sentinel-2, Landsat 8 and MODIS products (Figures 1 and 2). 2) The majority of variation in landscape 401 greenness was contained at scales of around half-a-metre, and is thus not captured by 402 medium-grain satellites such as Sentinel-2 (Figure 3). 3) When aggregating growing season 403 404 curves from ultra-fine-grain drone to medium-grain satellite pixel sizes, a notable amount 405 (46.2 - 63.9%) of variation in greenness phenology was lost (Figure 4). 4) Drone-based 406 measures of landscape greenness correlated well with ground-based measurements of leaf length (Figure 5). Taken together, our results highlight that drone platforms and compact 407 multispectral sensors can capture key ecological processes such as vegetation phenology 408 and bridge the existing scale gap between satellite and ground-based monitoring in tundra 409 ecosystems. 410

411

412 The correspondence between drone and satellite-derived NDVI has yet to be 413 comprehensively tested across Arctic sites (13,14). Siewert and Olofson (14) also 414 demonstrate cross-sensor agreement between drone- and satellite-derived NDVI from Arctic 415 Sweden. While similar or higher levels of cross-sensor agreement have been observed in 416 other natural and agricultural systems (14,97,98), some non-Arctic studies showed mixed or 417 poor agreement (99–101). Continued efforts in replicating these studies at different study 418 sites and systems is much needed to comprehensively evaluate cross-sensor 419 correspondence in Arctic tundra systems and beyond.

420

We observed a small but consistent offset between drone- and satellite-derived NDVI that 421 warrants further investigation. A similar offset has been detected in rice fields in Italy (100) 422 and with spectroradiometer readings in ecologically similar tundra in Alaska (77), but see 423 Siewert and Olofson (14) for a lack of offset in the more heterogeneous tundra of Arctic 424 Sweden. Both technical and ecological factors could explain the offset. We were not able to 425 conduct spectroradiometer readings coinciding with our drone surveys for on-the-ground 426 427 comparisons. Technical reasons for the observed offset may include: atmospheric effects, differences in viewing geometries, sensor properties (e.g. band widths) and signal 428 processing (e.g. radiometric calibration) between drones and satellites, but also among 429 different drone studies. Ecologically, the variation in land cover (especially the 430 presence/absence of non-vegetative surfaces) or topography within a landscape may 431 correspondence between vegetation greenness across scales due to 432 influence non-linearities in how aggregating different patch sizes and cover types influence measures 433 of NDVI (12,102). The high homogeneity of the survey plots on Qikiqtaruk likely contributes 434 to the strong correlation between drone- and satellite-derived NDVI that we have observed. 435 Yet, in our drone data fine-grain patterns of higher and lower NDVI within the landscape 436 including bare-ground patches and areas of more productive vegetation in wetter parts of the 437 landscape were evident (Figures 1-3). Heterogeneity in land cover and landscape-level 438 variation in NDVI at different spatial scales from vegetation patterning to tundra patterns in 439 topographically variable terrain could be differentially influencing correspondence in NDVI in 440 drone and satellite data across scales. Further research is needed to evaluate cross-sensor 441 and cross-scale correspondence in NDVI and other vegetation indices across Arctic tundra 442 443 systems.

444

445 We found that a plateau of spatial variation in tundra greenness occurred around 0.5 m, approximately the same width as biological and environmental patterning at this site. The E. 446 vaginatum sedges that dominate the tussock sedge vegetation type typically have diameters 447 of  $\sim 0.1 - 0.5$  m (Figure 3b) (103). The tussock sedge vegetation type is underlain by 448 ice-wedge polygons that when thawed create bands of wetter or drier plant communities with 449 widths of ~ 0.5 m - 3.0 m (104). Dryas-vetch vegetation is often found on gentle sloping 450 uplands where active layer disturbances such as cryoturbation and solifluction create 451 452 characteristic bare-ground patches perpendicular to the slope (85) with dimensions of  $\sim 0.3$ m - 0.5 m width and ~ 0.3 - 1.0 m length (Figure 3b). We expect that spatial variation would 453 increase with distances beyond the one-hectare extents of our plots as more topographic 454 diverse terrain is encountered and vegetation type transitions are reached. Topography is a 455 key proxy for many processes that structure heterogeneity in tundra vegetation (105–107) 456 and the plots were selected for little topographic variation to allow us to isolate specific 457 458 effects of land cover on scaling of greenness patterns from topography. The plot with the highest elevational range (Area 3 - Dryas-vetch tundra: 8.7 m) showed a small but steady 459 460 increase in spatial variation in distance classes above 10 m (Figure S4). Our findings

461 illustrate that on Qikiqtaruk, grain sizes of 0.5 m or less are required to capture key spatial 462 variation in vegetation greenness.

463

464 In our study, ecological information was lost when upscaling from ultra-fine-grain (~ 0.05 m) drone to moderate grain (~ 10 – 30m) satellite resolutions. Even the most recent generation 465 of freely-available multispectral satellite products can be limited in their ability to capture 466 fine-grain ecological processes of tundra vegetation change (13). Information transfer during 467 upscaling leads to the loss of more information in tundra ecosystems compared to other 468 biomes (14,108) as land cover and vegetation structure are fragmented at finer scales (109). 469 However, exactly how spatial aggregation influences the loss in observed ecological 470 variability across the diversity of Arctic landscapes remains poorly quantified (11). Yet, this 471 variability is critical to understanding climate-driven changes in vegetation phenology 472 473 (35,36,88), plant-pollinator interactions (110), and trophic interactions (111). With fine-grain observations, we were able to detect a subtle decrease in the magnitude of the spatial 474 variability in landscape-level phenology as the growing season progressed (Figure S5), while 475 aggregation to moderate satellite grains obscured both the magnitude and timing of 476 phenological heterogeneity (Figure 4). Thus, time-series of fine-grain remotely-sensed 477 observations will be critical for answering key research questions about tundra ecosystem 478 functioning in a warming Arctic (112). 479

480

Our results indicate that drone-based greenness time-series captured variation in plot-level 481 leaf-growth of deciduous tundra plant species. We demonstrate how drones can be used to 482 measure variation in tundra plant phenology of metre-scale patches at landscape extents. 483 Drones have been successfully used to monitor phenology of individual plants (trees) in 484 temperate forest ecosystems (113-115), and our study indicates that individual plant-level 485 phenology monitoring of sub-decimeter variability from drones could also be carried out in 486 tundra ecosystems. Future studies that quantify plant growth or phenology events such as 487 leaf emergence and flowering across the landscape could provide key information on 488 resource availability for plant-consumer interactions (110,111). Our findings also highlight 489 490 known limitations of NDVI to track phenology in evergreens or other non-deciduous taxa (D. integrifolia, Figure S9), suggesting that tests of alternative vegetation index - plant growth 491 relationships (115) are needed to capture within-growing-season variation in plant metabolic 492 activity of tundra evergreen and moss species. Combining drone-based time-series with 493 observations from phenocams, satellite and ground-based study plots has the potential to 494 revolutionise our understanding of landscape-scale phenology (13) by moving beyond the 495 previously small samples of individuals monitored in the Arctic tundra (36,37,39,116). 496 497

The collection of multispectral drone time-series in Arctic ecosystems has limitations and challenges. Recent studies have discussed challenges with radiometric consistency and repeatability when using compact multispectral drone sensors (81,117,118). Due to logistical constraints, we were not able to always conduct surveys under optimal conditions due to sun angle or cloud cover nor as frequently as planned due to wind or precipitation (Table S2), which likely introduced bias and/or noise into our drone data (e.g., Figure 4b). Access limitations meant that we could not capture spring and autumn on Qikiqtaruk. As an early-generation multispectral drone sensor, the Parrot Sequoia was tailored for deriving the NDVI, which despite being the legacy workhorse of tundra remote-sensing has limitations 507 (11,13). In particular, NDVI can be confounded by moisture and surface water (11,73,119), 508 complicating interpretation in wet tundra and particularly in fine-grain size data. However, the 509 rapid technological development of drones and sensors, as well as further consolidation and 510 standardisation of methods (120), will allow for pan-Arctic syntheses of fine-grain data to 511 resolve the uncertainty and complexity of Arctic greening patterns trends (13,14,81) (see 512 also the High Latitude Drone Ecology Network - <u>https://arcticdrones.org/</u>).

513

Our study demonstrates that drones can fill the scale-gap between satellite and field studies 514 515 in the observation of terrestrial Arctic vegetation change. Rather than investigating and 516 explaining patterns at scales pre-defined by satellite datasets or field-based networks, researchers can use drones to identify scale-domains that are most closely associated with 517 ecological processes of interest. Field ecologists can now use scaling theory provided by the 518 519 remote sensing community (74,121–124) at scales and temporal intervals that will allow for 520 hypothesis testing about what mechanisms are driving landscape-level ecological change. Drone imagery will also allow the remote sensing community to track the effects of sub-pixel 521 522 heterogeneity on satellite products down to the grain of individual plants and communities 523 (14) that have been long studied by field-based monitoring networks, like the International 524 Tundra Experiment (75). Only by improving our understanding of how ecologically important 525 information is captured across grain sizes can we reduce uncertainties in the medium- and coarse-grain satellite observation that feed into Earth system models and shape their 526 predictions (4,8). Fine-scale remote sensing from drones and aircraft provides key tools for 527 disentangling the drivers behind the greening of the Arctic (14,79,112). 528

#### 529

### 530 Conclusions

531

Novel remote-sensing technologies such as drones now allow us to study ecological 532 variation in landscapes continuously across scales. Fine-grain ecological observation is of 533 particular importance where variation in plant growth happens at very small spatial scales 534 535 such as in tundra ecosystems (13,71). Our finding of a peak in spatial variation found at 536 distances of ~0.5 m in the plots on Qikiqtaruk shows the grain size at which phenological information within the plant communities is best captured at this site. We demonstrate that 537 key ecological information is lost when observing the tundra at even decimeter or coarser 538 539 scales, such as those of medium grain satellites (~ 10 – 30m). Despite the methodological 540 challenges of collecting multispectral drone imagery in remote environments (81), our time-series of vegetation greenness correlated well with ground-based measurements of leaf 541 growth in the validation plots. Drones now enable cross-scale studies that fill scale gaps 542 between satellite and ground-based observations facilitating the identification of key drivers 543 of vegetation change to inform projections of climate change impacts and feedbacks in the 544 tundra biome. 545

546

### 547 Acknowledgements

548

549 We would like to thank the Team Shrub field crews of the 2016 and 2017 field seasons for 550 their hard work and effort invested in collecting the data presented in this research, this 551 includes Will Palmer, Santeri Lehtonen, Callum Tyler, Sandra Angers-Blondin and Haydn 552 Thomas. Furthermore, we would like to thank Tom Wade and Simon Gibson-Poole from the 553 University of Edinburgh Airborne GeoSciences Facility, as well as Chris McLellan and 554 Andrew Gray from the NERC Field Spectroscopy Facility for their support in our drone 555 endeavours. We also want to express our gratitude to Ally Phillimore, Ed Midchard and Toke 556 Høye for providing feedback on earlier versions of this manuscript.

557

We thank the Herschel Island - Qikiqtaruk Territorial Park Team and Yukon Government for providing logistical support for our field research on Qikiqtaruk including: Richard Gordon, Cameron Eckert and the park rangers Edward McLeod, Sam McLeod, Ricky Joe, Paden Lennie and Shane Goosen. We thank the research group of Hugues Lantuit at the Alfred Wegener Institute and the Aurora Research Institute for logistical support. Research permits include Yukon Researcher and Explorer permits (16-48S&E and 17-42S&E) and Yukon Parks Research permits (RE-Inu-02-16 and 17-RE-HI-02). All airborne activities were licensed under the Transport Canada special flight operations certificates ATS 16-17-00008441 RDIMS 11956834 (2016) and ATS 16-17-00072213 RDIMS 12929481 567 (2017).

568

569 Funding for this research was provided by NERC through the ShrubTundra standard grant 570 (NE/M016323/1), a NERC E3 Doctoral Training Partnership PhD studentship for Jakob 571 Assmann (NE/L002558/1), a research grant from the National Geographic Society 572 (CP-061R-17), a Parrot Climate Innovation Grant, the Aarhus University Research 573 Foundation, and the European Union's Horizon 2020 research and innovation programme 574 under the Marie Skłodowska-Curie grant agreement (754513) for Jeffrey Kerby, a NERC 575 support case for use of the NERC Field Spectroscopy Facility (738.1115), equipment Ioans 576 from the University of Edinburgh Airborne GeoSciences Facility and the NERC Geophysical 577 Equipment Facility (GEF 1063 and 1069).

578

579 Finally, we would like to thank the Inuvialuit people for the opportunity to conduct research in 580 the Inuvialuit Settlement Region.

581

## 582 Author Contributions

583

JJA and IMS conceived the study with input from JTK and AMC. JJA carried out data processing and analysis. JJA and IMS led the drone and ground-validation field work in 2016. AMC led the drone field surveys with input from JTK and GD led the ground-validation for 2017 with input from JTK. JJA, IMS and JTK wrote the manuscript with input from AMC and GD. IMS supervised and acquired funding for the research.

589

## 590 Data availability

591

592 All processed drone and Sentinel imagery is available via a data repository on Zenodo 593 (embargoed till publication of this manuscript).

594 Should the reviewers wish to access the data prior publication, a mirror of the Zenodo 595 repository can be accessed via this confidential link:

596

598 All code used to conduct the analysis, produce figures ands as well as summary data 599 outputs and MODIS pixel values can be found on this GitHub repository (already openly 600 available):

601 <u>https://github.com/jakobjassmann/qhi\_phen\_ts</u>

## 603 References

- Levin SA. The Problem of Pattern and Scale in Ecology: The Robert H. MacArthur
   Award Lecture. Ecology. 1992 Dec 1;73(6):1943–67.
- Anderson CB. Biodiversity monitoring, earth observations and the ecology of scale.
   Ecol Lett. 2018 Oct 1;21(10):1572–85.
- 609 3. Estes L, Elsen PR, Treuer T, Ahmed L, Caylor K, Chang J, et al. The spatial and 610 temporal domains of modern ecology. Nat Ecol Evol. 2018 May;2(5):819.
- 613 28;310(5748):657–60.
  614 5. Loranty MM, Goetz SJ. Shrub expansion and climate feedbacks in Arctic tundra.
- 615 Environ Res Lett. 2012 Mar 1;7(1):011005.
- 616 6. Pearson RG, Phillips SJ, Loranty MM, Beck PSA, Damoulas T, Knight SJ, et al. Shifts
  617 in Arctic vegetation and associated feedbacks under climate change. Nat Clim Change.
  618 2013 Jul;3(7):673–7.
- Richardson AD, Keenan TF, Migliavacca M, Ryu Y, Sonnentag O, Toomey M. Climate
  change, phenology, and phenological control of vegetation feedbacks to the climate
  system. Agric For Meteorol. 2013 Feb 15;169:156–73.
- Ernakovich JG, Hopping KA, Berdanier AB, Simpson RT, Kachergis EJ, Steltzer H, et
   al. Predicted responses of arctic and alpine ecosystems to altered seasonality under
   climate change. Glob Change Biol [Internet]. 2014 Mar 1 [cited 2014 Apr 9]; Available
   from: http://onlinelibrary.wiley.com/doi/10.1111/gcb.12568/abstract
- 626 9. Diepstraten RAE, Jessen TD, Fauvelle CMD, Musiani MM. Does climate change and 627 plant phenology research neglect the Arctic tundra? Ecosphere. 2018;9(9):e02362.
- Metcalfe DB, Hermans TDG, Ahlstrand J, Becker M, Berggren M, Björk RG, et al.
  Patchy field sampling biases understanding of climate change impacts across the
  Arctic. Nat Ecol Evol. 2018 Sep;2(9):1443.
- Beamish A, Raynolds MK, Epstein H, Frost GV, Macander MJ, Bergstedt H, et al.
   Recent trends and remaining challenges for optical remote sensing of Arctic tundra
   vegetation: A review and outlook. Remote Sens Environ. 2020 Sep 1;246:111872.
- Stoy PC, Williams M, Disney M, Prieto-Blanco A, Huntley B, Baxter R, et al. Upscaling
   as ecological information transfer: a simple framework with application to Arctic
   ecosystem carbon exchange. Landsc Ecol. 2009 Aug 1:24(7):971–86.
- 637 13. Myers-Smith IH, Kerby JT, Phoenix GK, Bjerke JW, Epstein HE, Assmann JJ, et al.
- Complexity revealed in the greening of the Arctic. Nat Clim Change. 2020
   Feb;10(2):106–17.
- 640 14. Siewert MB, Olofsson J. Scale-dependency of Arctic ecosystem properties revealed by
  641 UAV. Environ Res Lett [Internet]. 2020 Jul 2 [cited 2020 Jul 6]; Available from:
  642 https://iopscience.iop.org/article/10.1088/1748-9326/aba20b
- 643 15. Myneni RB, Keeling CD, Tucker CJ, Asrar G, Nemani RR. Increased plant growth in the
- 644 northern high latitudes from 1981 to 1991. Nature. 1997 Apr;386(6626):698–702.
- Tucker CJ, Slayback DA, Pinzon JE, Los SO, Myneni RB, Taylor MG. Higher northern
  latitude normalized difference vegetation index and growing season trends from 1982
  to 1999. Int J Biometeorol. 2001 Nov 1;45(4):184–90.
- 648 17. Bhatt US, Walker DA, Raynolds MK, Comiso JC, Epstein HE, Jia GJ, et al. Circumpolar
  649 arctic tundra vegetation change is linked to sea ice decline. Earth Interact. 2010
  650 Aug;14(8):1–20.
- 651 18. Guay KC, Beck PSA, Berner LT, Goetz SJ, Baccini A, Buermann W. Vegetation 652 productivity patterns at high northern latitudes: a multi-sensor satellite data
- assessment. Glob Change Biol. 2014;20(10):3147–3158.

19. Zhu Z, Piao S, Myneni RB, Huang M, Zeng Z, Canadell JG, et al. Greening of the Earth 654 and its drivers. Nat Clim Change. 2016 Aug;6(8):791-5. 655 656 **20**. Keenan TF, Riley WJ. Greening of the land surface in the world's cold regions consistent with recent warming. Nat Clim Change. 2018 Sep;8(9):825-8. 657 658 21. Zeng H, Jia G, Epstein H. Recent changes in phenology over the northern high latitudes detected from multi-satellite data. Environ Res Lett. 2011;6(4):045508. 659 660 22. Zeng H, Jia G, Forbes BC. Shifts in Arctic phenology in response to climate and anthropogenic factors as detected from multiple satellite time series. Environ Res Lett. 661 662 2013;8(3):035036. 23. Zhao J, Zhang H, Zhang Z, Guo X, Li X, Chen C. Spatial and Temporal Changes in 663 664 Vegetation Phenology at Middle and High Latitudes of the Northern Hemisphere over the Past Three Decades. Remote Sens. 2015 Aug;7(8):10973-95. 665 666 24. Potter C, Alexander O. Changes in Vegetation Phenology and Productivity in Alaska 667 Over the Past Two Decades. Remote Sens. 2020 Jan;12(10):1546. Tucker CJ. Red and photographic infrared linear combinations for monitoring 25. 668 vegetation. Remote Sens Environ. 1979 May 1;8(2):127-50. 669 670 **26**. Myneni RB, Tucker CJ, Asrar G, Keeling CD. Interannual variations in satellite-sensed vegetation index data from 1981 to 1991. J Geophys Res Atmospheres. 671 672 1998;103(D6):6145-60. 673 27. Elmendorf SC, Henry GHR, Hollister RD, Björk RG, Boulanger-Lapointe N, Cooper EJ, 674 et al. Plot-scale evidence of tundra vegetation change and links to recent summer warming. Nat Clim Change. 2012 Apr 8;2:453-7. 675 676 **28**. Tape KD, Sturm M, Racine CH. The evidence for shrub expansion in Northern Alaska 677 and the Pan-Arctic. Glob Change Biol. 2006 Apr;12(4):686-702. Myers-Smith IH, Forbes BC, Wilmking M, Hallinger M, Lantz T, Blok D, et al. Shrub 678 29. 679 expansion in tundra ecosystems: dynamics, impacts and research priorities. Environ Res Lett. 2011 Dec 1;6(4):045509. 680 30. Tape K, Hallinger M, Welker J, Ruess R. Landscape heterogeneity of shrub expansion 681 in Arctic Alaska. Ecosystems. 2012;15(5):711–24. 682 31. García Criado M, Myers-Smith IH, Bjorkman AD, Lehmann CER, Stevens N. Woody 683 plant encroachment intensifies under climate change across tundra and savanna 684 biomes. Glob Ecol Biogeogr. 2020; (in press). 685 Bjorkman AD, Myers-Smith IH, Elmendorf SC, Normand S, Rüger N, Beck PSA, et al. 32. 686 Plant functional trait change across a warming tundra biome. Nature. 2018 687 Oct;562(7725):57-62. 688 33. Høye TT, Post E, Meltofte H, Schmidt NM, Forchhammer MC. Rapid advancement of 689 spring in the High Arctic. Curr Biol. 2007 Jun 19;17(12):R449–51. 690 34. Kerby JT, Post E. Advancing plant phenology and reduced herbivore production in a 691 692 terrestrial system associated with sea ice decline. Nat Commun [Internet]. 2013 Oct 1 693 [cited 2014 Oct 23];4. Available from: 694 http://www.nature.com/ncomms/2013/131001/ncomms3514/full/ncomms3514.html?me 695 ssage-global=remove 35. Post E, Kerby J, Pedersen C, Steltzer H. Highly individualistic rates of plant 696 697 phenological advance associated with arctic sea ice dynamics. Biol Lett. 2016 Dec 1;12(12):20160332. 698 Assmann JJ, Myers-Smith IH, Phillimore AB, Bjorkman AD, Ennos RE, Prevéy JS, et 36. 699 700 al. Local snow melt and temperature—but not regional sea ice—explain variation in 701 spring phenology in coastal Arctic tundra. Glob Change Biol. 2019;25(7):2258–74. 702 37. Oberbauer SF, Elmendorf SC, Troxler TG, Hollister RD, Rocha AV, Bret-Harte MS, et al. Phenological response of tundra plants to background climate variation tested using 703 the International Tundra Experiment. Philos Trans R Soc B Biol Sci [Internet]. 2013 Aug 704

705 19 [cited 2013 Aug 20];368(1624). Available from: http://rstb.royalsocietypublishing.org/content/368/1624/20120481 706 707 38. Bjorkman AD, Elmendorf SC, Beamish AL, Vellend M, Henry GHR. Contrasting effects of warming and increased snowfall on Arctic tundra plant phenology over the past two 708 decades. Glob Change Biol. 2015 Dec 1;21(12):4651-61. 709 710 39. Prevéy JS, Rixen C, Rüger N, Høye TT, Bjorkman AD, Myers-Smith IH, et al. Warming 711 shortens flowering seasons of tundra plant communities. Nat Ecol Evol. 2019 712 Jan;3(1):45. 713 40. Jia GJ, Epstein HE, Walker DA. Spatial heterogeneity of tundra vegetation response to 714 recent temperature changes. Glob Change Biol. 2006 Jan;12(1):42-55. Raynolds MK, Comiso JC, Walker DA, Verbyla D. Relationship between 715 **41**. satellite-derived land surface temperatures, arctic vegetation types, and NDVI. Remote 716 717 Sens Environ. 2008 Apr 15;112(4):1884-94. Jia GJ, Epstein HE, Walker DA. Vegetation greening in the Canadian Arctic related to 718 **42**. decadal warming. J Environ Monit. 2009;11(12):2231. 719 720 43. Bhatt US, Walker DA, Raynolds MK, Bieniek PA, Epstein HE, Comiso JC, et al. Recent declines in warming and vegetation greening trends over Pan-Arctic tundra. Remote 721 722 Sens. 2013 Aug 29;5(9):4229-54. 723 44. Xu L, Myneni RB, Chapin Iii FS, Callaghan TV, Pinzon JE, Tucker CJ, et al. 724 Temperature and vegetation seasonality diminishment over northern lands. Nat Clim 725 Change [Internet]. 2013 Mar 10 [cited 2013 Mar 12]; Available from: http://www.nature.com/nclimate/journal/vaop/ncurrent/full/nclimate1836.html 726 727 45. Piao S, Nan H, Huntingford C, Ciais P, Friedlingstein P, Sitch S, et al. Evidence for a 728 weakening relationship between interannual temperature variability and northern vegetation activity. Nat Commun. 2014 Oct 16;5:5018. 729 730 46. Vickers H, Høgda KA, Solbø S, Karlsen SR, Tømmervik H, Aanes R, et al. Changes in 731 greening in the high Arctic: insights from a 30 year AVHRR max NDVI dataset for Svalbard. Environ Res Lett. 2016;11(10):105004. 732 733 **47**. Walker DA, Leibman MO, Epstein HE, Forbes BC, Bhatt US, Raynolds MK, et al. 734 Spatial and temporal patterns of greenness on the Yamal Peninsula, Russia: 735 interactions of ecological and social factors affecting the Arctic normalized difference vegetation index. Environ Res Lett. 2009 Oct;4(4):045004. 736 737 48. Macias-Fauria M, Forbes BC, Zetterberg P, Kumpula T. Eurasian Arctic greening 738 reveals teleconnections and the potential for structurally novel ecosystems. Nat Clim Change. 2012;2:613-618. 739 740 49. Post E. Erosion of community diversity and stability by herbivore removal under warming. Proc R Soc B Biol Sci. 2013 Apr 22;280(1757):20122722. 741 Bhatt US, Walker DA, Walsh JE, Carmack EC, Frey KE, Meier WN, et al. Implications 742 **50**. 743 of Arctic sea ice decline for the Earth system. Annu Rev Environ Resour. 2014;39:57-89. 744 745 51. Fauchald P, Park T, Tømmervik H, Myneni R, Hausner VH. Arctic greening from warming promotes declines in caribou populations. Sci Adv. 2017 Apr 1;3(4):e1601365. 746 747 **52**. Elmendorf SC, Henry GHR, Hollister RD, Fosaa AM, Gould WA, Hermanutz L, et al. 748 Experiment, monitoring, and gradient methods used to infer climate change effects on plant communities yield consistent patterns. Proc Natl Acad Sci. 2015 Jan 749 13;112(2):448-52. 750 53. Pattison RR, Jorgenson JC, Raynolds MK, Welker JM. Trends in NDVI and tundra 751 752 community composition in the Arctic of NE Alaska between 1984 and 2009. 753 Ecosystems. 2015 Mar 19;18(4):707-19. Piao S, Tan J, Chen A, Fu YH, Ciais P, Liu Q, et al. Leaf onset in the northern 754 54. hemisphere triggered by daytime temperature. Nat Commun. 2015 Apr 23;6:6911. 755

- 756 55. Liu Q, Fu YH, Zhu Z, Liu Y, Liu Z, Huang M, et al. Delayed autumn phenology in the Northern Hemisphere is related to change in both climate and spring phenology. Glob 757 758 Change Biol. 2016 Nov 1;22(11):3702-11. Semenchuk PR, Gillespie MAK, Rumpf SB, Baggesen N, Elberling B, Cooper EJ. High 759 **56**. Arctic plant phenology is determined by snowmelt patterns but duration of phenological 760 periods is fixed: an example of periodicity. Environ Res Lett. 2016;11(12):125006. 761 Prevéy J, Vellend M, Rüger N, Hollister RD, Bjorkman AD, Myers-Smith IH, et al. 762 **57**. Greater temperature sensitivity of plant phenology at colder sites: implications for 763 convergence across northern latitudes. Glob Change Biol [Internet]. 2017 Feb 1 [cited 764 765 2017 Feb 6]; Available from: http://onlinelibrary.wiley.com.ezproxy.is.ed.ac.uk/doi/10.1111/gcb.13619/abstract 766 Jia GJ, Epstein HE, Walker DA. Greening of arctic Alaska, 1981–2001. Geophys Res 767 58. Lett. 2003 Oct 29;30(20):HLS 3-1. 768 Fraser RH, Olthof I, Carrière M, Deschamps A, Pouliot D. Detecting long-term changes 769 **59**. to vegetation in northern Canada using the Landsat satellite image archive. Environ 770 771 Res Lett. 2011 Oct 1;6(4):045502. Ju J, Masek JG. The vegetation greenness trend in Canada and US Alaska from 772 60. 773 1984–2012 Landsat data. Remote Sens Environ. 2016 Apr: 176:1–16. Raynolds MK, Walker DA, Verbyla D, Munger CA. Patterns of Change within a Tundra 774 61. 775 Landscape: 22-year Landsat NDVI Trends in an Area of the Northern Foothills of the 776 Brooks Range, Alaska. Arct Antarct Alp Res. 2013 May 1;45(2):249-60. 777 62. Miles VV, Esau I. Spatial heterogeneity of greening and browning between and within bioclimatic zones in northern West Siberia. Environ Res Lett. 2016;11(11):115002. 778 779 63. Lara MJ, Nitze I, Grosse G, Martin P, McGuire AD. Reduced arctic tundra productivity linked with landform and climate change interactions. Sci Rep. 2018 Feb 5;8(1):2345. 780 781 64. Thompson JA, Koenig LS. Vegetation phenology in Greenland and links to cryospheric 782 change. Ann Glaciol. 2018 Dec;59(77):59-68. Phoenix GK, Bjerke JW. Arctic browning: extreme events and trends reversing arctic 783 65. 784 greening. Glob Change Biol. 2016 Sep 1;22(9):2960-2. 785 66. National Academies of Sciences E. Understanding Northern Latitude Vegetation 786 Greening and Browning: Proceedings of a Workshop [Internet]. 2019 [cited 2019 Oct 787 21]. Available from: 788 https://www.nap.edu/catalog/25423/understanding-northern-latitude-vegetation-greenin 789 g-and-browning-proceedings-of-a 790 67. Frost GV, Bhatt US, Epstein HE, Walker DA, Raynolds MK, Berner LT, et al. Arctic 791 Report Card: Update for 2019 - Tundra Greenness [Internet]. Arctic Program. 2019 792 [cited 2019 Feb 23]. Available from: 793 http://www.arctic.noaa.gov/Report-Card/Report-Card-2015/ArtMID/5037/ArticleID/221/T 794 undra-Greenness Blok D, Schaepman-Strub G, Bartholomeus H, Heijmans MMPD, Maximov TC, 795 68. Berendse F. The response of Arctic vegetation to the summer climate: relation between 796 shrub cover, NDVI, surface albedo and temperature. Environ Res Lett. 2011 Jul 797 798 1:6:035502. 799 69. Raynolds MK, Walker DA, Epstein HE, Pinzon JE, Tucker CJ. A new estimate of tundra-biome phytomass from trans-Arctic field data and AVHRR NDVI. Remote Sens 800 Lett. 2012 Sep 1;3(5):403-11. 801 Berner LT, Jantz P, Tape KD, Goetz SJ. Tundra plant above-ground biomass and shrub 802 70. 803 dominance mapped across the North Slope of Alaska. Environ Res Lett. 2018 804 Feb;13(3):035002. Cunliffe AM, Assmann JJ, Daskalova G, Kerby JT, Myers-Smith IH. Aboveground 805 71.
- biomass corresponds strongly with drone-derived canopy height but weakly with

807 greenness (NDVI) in a shrub tundra landscape. Environ Res Lett [Internet]. 2020 [cited 2020 Jul 10]; Available from: http://iopscience.iop.org/10.1088/1748-9326/aba470 808 809 72. Stow D. Remote sensing of vegetation and land-cover change in arctic tundra ecosystems. Remote Sens Environ. 2004 Feb;89(3):281-308. 810 Bartsch A, Widhalm B, Leibman M, Ermokhina K, Kumpula T, Skarin A, et al. Feasibility 73. 811 of tundra vegetation height retrieval from Sentinel-1 and Sentinel-2 data. Remote Sens 812 813 Environ. 2020 Feb 1;237:111515. 814 **74**. Woodcock CE, Strahler AH. The factor of scale in remote sensing. Remote Sens 815 Environ. 1987 Apr 1;21(3):311-32. Henry GHR, Molau U. Tundra plants and climate change: the International Tundra 75. 816 Experiment (ITEX). Glob Change Biol. 1997 Dec 1;3(S1):1-9. 817 818 76. Beck PSA, Jönsson P, Høgda K-A, Karlsen SR, Eklundh L, Skidmore AK. A ground-validated NDVI dataset for monitoring vegetation dynamics and mapping 819 820 phenology in Fennoscandia and the Kola peninsula. Int J Remote Sens. 2007 Oct 821 10;28(19):4311-30. 77. Gamon JA, Huemmrich KF, Stone RS, Tweedie CE. Spatial and temporal variation in 822 primary productivity (NDVI) of coastal Alaskan tundra: Decreased vegetation growth 823 following earlier snowmelt. Remote Sens Environ. 2013 Feb 15;129:144–53. 824 78. Kerby JT. Phenology in a changing Arctic: Linking trophic interactions across scales. 825 2015 Sep 14 [cited 2018 Oct 19]; Available from: 826 https://etda.libraries.psu.edu/catalog/26992 827 79. Anderson K, Gaston KJ. Lightweight unmanned aerial vehicles will revolutionize spatial 828 ecology. Front Ecol Environ. 2013 Mar 18;11(3):138-46. 829 830 80. Klosterman S, Richardson AD. Observing Spring and Fall Phenology in a Deciduous Forest with Aerial Drone Imagery. Sensors. 2017 Dec;17(12):2852. 831 832 **81**. Assmann JJ, Kerby JT, Cunliffe AM, Myers-Smith IH. Vegetation monitoring using multispectral sensors — best practices and lessons learned from high latitudes. J 833 Unmanned Veh Syst. 2018 Dec 5;7(1):54-75. 834 82. Klosterman S, Hufkens K, Richardson AD. Later springs green-up faster: the relation 835 between onset and completion of green-up in deciduous forests of North America. Int J 836 Biometeorol. 2018 Sep 1;62(9):1645-55. 837 Myers-Smith IH, Hik DS. Shrub canopies influence soil temperatures but not nutrient 838 83. 839 dynamics: An experimental test of tundra snow-shrub interactions. Ecol Evol. 2013 Oct 1;3(11):3683-700. 840 84. Smith C, Kennedy C, Hargrave A, McKenna K. Soil and Vegetation of Herschel Island. 841 Whitehorse, Yukon Territory, Canada: Agriculture Canada; 1989. Report No.: No 1. 842 Obu J, Lantuit H, Myers-Smith I, Heim B, Wolter J, Fritz M. Effect of Terrain 843 85. Characteristics on Soil Organic Carbon and Total Nitrogen Stocks in Soils of Herschel 844 845 Island, Western Canadian Arctic. Permafr Periglac Process. 2015 Jan 1;28(1):92–107. Myers-Smith IH, Hik DS, Kennedy C, Cooley D, Johnstone JF, Kenney AJ, et al. 86. 846 Expansion of Canopy-Forming Willows Over the Twentieth Century on Herschel Island, 847 Yukon Territory, Canada. AMBIO. 2011 Sep;40(6):610-23. 848 849 87. Raynolds MK, Walker DA, Balser A, Bay C, Campbell M, Cherosov MM, et al. A raster 850 version of the Circumpolar Arctic Vegetation Map (CAVM). Remote Sens Environ. 2019 Oct 1;232:111297. 851 852 88. Myers-Smith IH, Grabowski MM, Thomas HJD, Angers-Blondin S, Daskalova GN, 853 Bjorkman AD, et al. Eighteen years of ecological monitoring reveals multiple lines of 854 evidence for tundra vegetation change. Ecol Monogr. 2019;89(2):e01351. 855 89. Didan K. MOD13Q1 MODIS/Terra Vegetation Indices 16-Day L3 Global 250m SIN Grid V006 [Internet]. NASA EOSDIS Land Processes DAAC; 2015 [cited 2018 Oct 22]. 856 Available from: https://doi.org/10.5067/MODIS/MOD13Q1.006 857

Engine: Planetary-scale geospatial analysis for everyone. Remote Sens Environ. 2017 859 Dec 1;202:18-27. 860 Mueller-Wilm U. Sen2Cor Software Release Note: Ref.: 861 **91**. S2-PDGS-MPC-L2A-SRN-V2.4.0 [Internet]. 2017 [cited 2018 Oct 22]. Available from: 862 863 http://step.esa.int/thirdparties/sen2cor/2.4.0/Sen2Cor 240 Documenation PDF/S2-PD 864 GS-MPC-L2A-SRN-V2.4.0.pdf Molau U, Mølgaard P. International Tundra Experiment Manual [Internet]. 1996. 865 92. Available from: https://www.gvsu.edu/itex/library-8.htm 866 Hadfield JD. MCMC Methods for Multi-Response Generalized Linear Mixed Models: 867 93. The MCMCgImm R Package. J Stat Softw [Internet]. 2010 [cited 2018 Mar 16];33(2). 868 Available from: http://www.jstatsoft.org/v33/i02/ 869 870 **94**. Hijmans RJ. raster: Geographic Data Analysis and Modeling [Internet]. 2020. Available 871 from: https://CRAN.R-project.org/package=raster Pebesma EJ. Multivariable geostatistics in S: the gstat package. Comput Geosci. 872 95. 2004;30:683-91. 873 874 96. Gräler B, Pebesma E, Heuvelink G. Spatio-Temporal Interpolation using gstat. R J. 875 2016;8(1):204-18. 876 **97**. Fernández-Guisuraga JM, Sanz-Ablanedo E, Suárez-Seoane S, Calvo L. Using Unmanned Aerial Vehicles in Postfire Vegetation Survey Campaigns through Large and 877 878 Heterogeneous Areas: Opportunities and Challenges. Sensors. 2018 Feb;18(2):586. 98. Fawcett D, Panigada C, Tagliabue G, Boschetti M, Celesti M, Evdokimov A, et al. 879 880 Multi-Scale Evaluation of Drone-Based Multispectral Surface Reflectance and Vegetation Indices in Operational Conditions. Remote Sens. 2020 Jan;12(3):514. 881 Matese A, Toscano P, Di Gennaro SF, Genesio L, Vaccari FP, Primicerio J, et al. 882 99. 883 Intercomparison of UAV, Aircraft and Satellite Remote Sensing Platforms for Precision Viticulture. Remote Sens. 2015 Mar;7(3):2971–90. 884 100. Franzini M, Ronchetti G, Sona G, Casella V. Geometric and Radiometric Consistency 885 886 of Parrot Sequoia Multispectral Imagery for Precision Agriculture Applications. Appl Sci. 887 2019 Jan;9(24):5314. 101. Khaliq A, Comba L, Biglia A, Ricauda Aimonino D, Chiaberge M, Gay P. Comparison of 888 Satellite and UAV-Based Multispectral Imagery for Vineyard Variability Assessment. 889 890 Remote Sens. 2019 Jan;11(4):436. 102. Riihimäki H, Luoto M, Heiskanen J. Estimating fractional cover of tundra vegetation at 891 892 multiple scales using unmanned aerial systems and optical satellite data. Remote Sens 893 Environ. 2019 Apr 1;224:119-32. 103. Mark AF, Fetcher N, Shaver GR, III FSC. Estimated Ages of Mature Tussocks of 894 895 Eriophorum Vaginatum along A Latitudinal Gradient in Central Alaska, U.S.A. Arct Alp 896 Res. 1985 Feb 1;17(1):1-5. 104. Fritz M, Wolter J, Rudaya N, Palagushkina O, Nazarova L, Obu J, et al. Holocene 897 ice-wedge polygon development in northern Yukon permafrost peatlands (Canada). 898 899 Quat Sci Rev. 2016 Sep 1;147:279–97. 105. Billings WD, Bliss LC. An Alpine Snowbank Environment and Its Effects on Vegetation, 900 901 Plant Development, and Productivity. Ecology. 1959;40(3):388-97. 106. Suvanto S, roux PCL, Luoto M. Arctic-alpine vegetation biomass is driven by 902 903 fine-scale abiotic heterogeneity. Geogr Ann Ser Phys Geogr. 2014 Dec 904 1;96(4):549-60. 905 107. Riihimäki H, Heiskanen J, Luoto M. The effect of topography on arctic-alpine 906 aboveground biomass and NDVI patterns. Int J Appl Earth Obs Geoinformation. 2017 Apr;56:44-53. 907 908 108. Stoy PC, Williams M, Spadavecchia L, Bell RA, Prieto-Blanco A, Evans JG, et al. Using

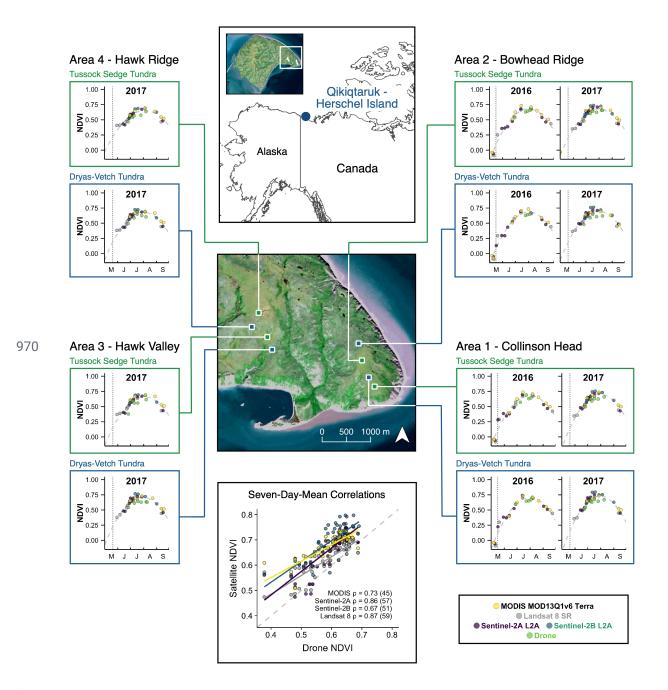
Gorelick N, Hancher M, Dixon M, Ilyushchenko S, Thau D, Moore R. Google Earth

90.

909 Information Theory to Determine Optimum Pixel Size and Shape for Ecological Studies: Aggregating Land Surface Characteristics in Arctic Ecosystems. Ecosystems. 2009 Jun 910 1;12(4):574-89. 911 912 109. Virtanen T, Ek M. The fragmented nature of tundra landscape. Int J Appl Earth Obs Geoinformation. 2014 Apr 1;27:4-12. 913 914 110. Høye TT, Post E, Schmidt NM, Trøjelsgaard K, Forchhammer MC. Shorter flowering 915 seasons and declining abundance of flower visitors in a warmer Arctic. Nat Clim Change. 2013 Aug;3(8):759-63. 916 111. Armstrong JB, Takimoto G, Schindler DE, Hayes MM, Kauffman MJ. Resource waves: 917 phenological diversity enhances foraging opportunities for mobile consumers. Ecology. 918 2016;97(5):1099-112. 919 112. Miller CE, Griffith PC, Goetz SJ, Hoy EE, Pinto N, McCubbin IB, et al. An overview of 920 921 ABoVE airborne campaign data acquisitions and science opportunities. Environ Res 922 Lett. 2019 Jul;14(8):080201. 923 113. Klosterman S, Melaas E, Wang JA, Martinez A, Frederick S, O'Keefe J, et al. 924 Fine-scale perspectives on landscape phenology from unmanned aerial vehicle (UAV) 925 photography. Agric For Meteorol. 2018 Jan 15:248:397-407. 926 114. Berra EF, Gaulton R, Barr S. Assessing spring phenology of a temperate woodland: A 927 multiscale comparison of ground, unmanned aerial vehicle and Landsat satellite observations. Remote Sens Environ. 2019 Mar 15;223:229-42. 928 929 115. D'Odorico P, Besik A, Wong CYS, Isabel N, Ensminger I. High-throughput drone-based 930 remote sensing reliably tracks phenology in thousands of conifer seedlings. New Phytol. 2020;226(6):1667-81. 931 932 116. Prevéy J, Vellend M, Rüger N, Hollister RD, Bjorkman AD, Myers-Smith IH, et al. Greater temperature sensitivity of plant phenology at colder sites: implications for 933 934 convergence across northern latitudes. Glob Change Biol. 2017;23(7):2660-71. 935 117. Fawcett D, Anderson K. Investigating impacts of calibration methodology and irradiance variations on lightweight drone-based sensor derived surface reflectance 936 937 products. In: Remote Sensing for Agriculture, Ecosystems, and Hydrology XXI 938 [Internet]. International Society for Optics and Photonics; 2019 [cited 2020 Jul 11]. p. 939 111490D. Available from: 940 https://www.spiedigitallibrary.org/conference-proceedings-of-spie/11149/111490D/Invest 941 igating-impacts-of-calibration-methodology-and-irradiance-variations-on-lightweight/10. 942 1117/12.2533106.short 118. Huang S, Tang L, Hupy JP, Wang Y, Shao G. A commentary review on the use of 943 944 normalized difference vegetation index (NDVI) in the era of popular remote sensing. J 945 For Res [Internet]. 2020 May 31 [cited 2020 Jul 11]; Available from: https://doi.org/10.1007/s11676-020-01155-1 946 947 119. Zhang Q, Yao T, Huemmrich KF, Middleton EM, Lyapustin A, Wang Y. Evaluating impacts of snow, surface water, soil and vegetation on empirical vegetation and snow 948 indices for the Utgiagvik tundra ecosystem in Alaska with the LVS3 model. Remote 949 Sens Environ. 2020 Apr 1;240:111677. 950 120. Aasen H, Honkavaara E, Lucieer A, Zarco-Tejada PJ. Quantitative Remote Sensing at 951 952 Ultra-High Resolution with UAV Spectroscopy: A Review of Sensor Technology, 953 Measurement Procedures, and Data Correction Workflows. Remote Sens. 2018 954 Jul;10(7):1091. 955 121. Strahler AH, Woodcock CE, Smith JA. On the nature of models in remote sensing. 956 Remote Sens Environ. 1986 Oct 1;20(2):121–39. 122. Chen JM. Spatial Scaling of a Remotely Sensed Surface Parameter by Contexture. 957 Remote Sens Environ. 1999 Jul 1;69(1):30-42. 958

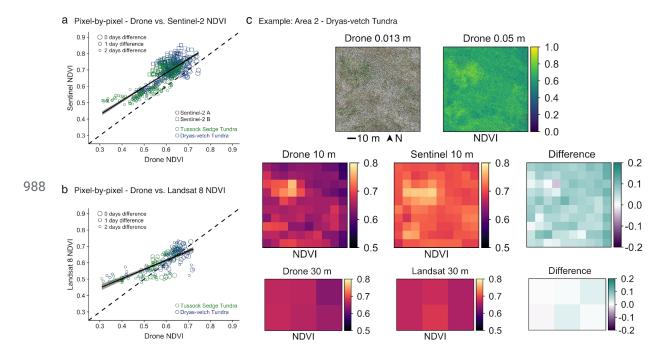
959 123. Chen JM, Chen X, Ju W. Effects of vegetation heterogeneity and surface topography

- 960 on spatial scaling of net primary productivity. Biogeosciences. 2013 Jul
- 961 18;10(7):4879–96.
- 962 124. Garrigues S, Allard D, Baret F, Weiss M. Influence of landscape spatial heterogeneity
   963 on the non-linear estimation of leaf area index from moderate spatial resolution remote
- sensing data. Remote Sens Environ. 2006 Dec 30;105(4):286–98.
- 965 125. Becker RA, Wilks AR, Brownrigg R, Minka TP, Deckmyn A. maps: Draw Geographical
   966 Maps [Internet]. 2018. Available from: https://CRAN.R-project.org/package=maps
- 967 126. Becker RA, Wilks AR, Brownrigg R. mapdata: Extra Map Databases [Internet]. 2018.
- 968 Available from: https://CRAN.R-project.org/package=mapdata

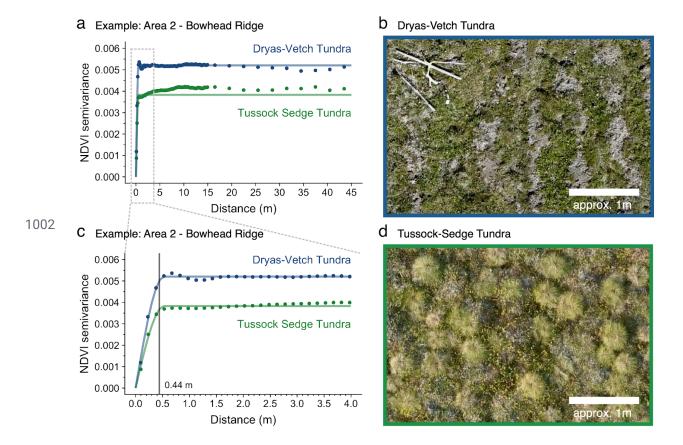


971 Figure 1: Drone-data captured the temporal variation in satellite data across vegetation communities, areas and 972 years. This figure showcases variation in mean landscape greenness (NDVI) across the eight one-hectare 973 sampling plots on Qikiqtaruk as derived from drone orthomosaics and the MODIS Vegetation Index 974 (MOD13Q1.v006 Terra), Landsat 8 Level 2 and Sentinel-2 Level-2A products. Vertical dotted grey lines represent 975 the average snow-melt at long-term monitoring plots close to Area 3 - Hawk Valley for the given year (88). 976 Dashed grey lines represent simple quadratic phenology curves (NDVI ~ a  $x^2$  + b x + c, where x is the day of 977 year, a the quadratic coefficient, b the linear coefficient and c the y-axis intercept) fitted to all data points pooled 978 across sensors. The lower central panel demonstrates the close correspondence between seven-day mean 979 values from drone and satellite NDVI, albeit with a positive offset for all satellite sensors. For this panel, drone 980 NDVI values were spatially aggregated by mean to the one-hectare plots and temporally aggregated by mean in 981 consecutive seven-day blocks starting on the first of May in both growing seasons (2016 and 2017) where data 982 was available. Matching seven-day block pairs between drone and satellite platforms were then identified and 983 plotted as shown. Spearman's rank correlation as well as mean differences (offsets) in NDVI amongst all platform 984 combinations can be found in Tables S12 and S13 respectively. The grey dashed line in this panel represents the

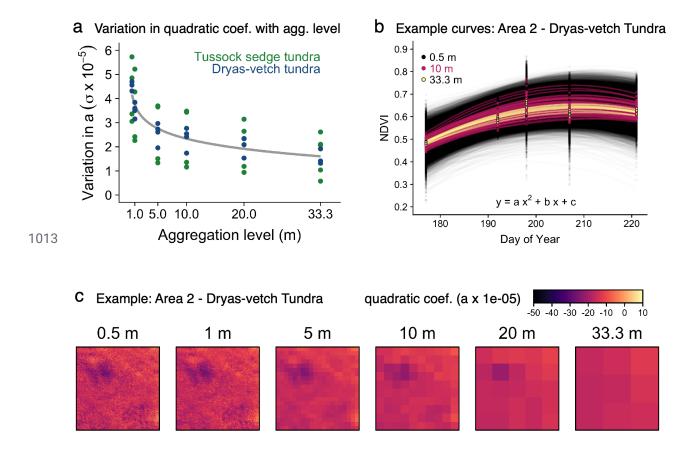
- 985 one-to-one line. Map sources: North America (125,126) in latitude and longitude on the WGS84 reference
- 986 ellipsoid and Qikiqtaruk, Copernicus Sentinel-2 true colour image July 2017 in UTM 7N based on the WGS84
- 987 reference ellipsoid.



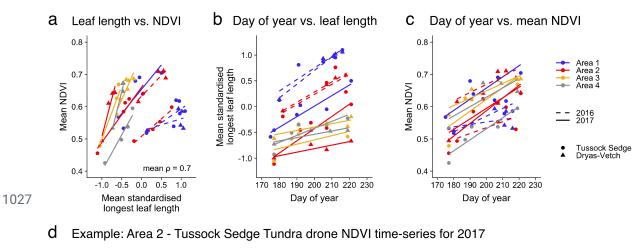
989 Figure 2: Drone-data better captured spatial heterogeneity in NDVI relative to Sentinel-2 MSI and Landsat 8 OLI 990 in pixel-by-pixel comparisons. a) Pixel-by-pixel correlations between 10 m aggregated drone NDVI and native 10 991 m Sentinel-2 NDVI for a random sample of pixels (10% of total pixels, n = 700) across all drone-sentinel image 992 pairs for the 2017 growing season that were a maximum of two days apart. No drone-sentinel image pairs were 993 available for the 2016 season that fitted the latter criterium. The black line represents a simple linear model 994 describing the relationship, see Table S8 for details. b) Pixel-by-pixel correlations between 30 m aggregated 995 drone NDVI and native 30 m Landsat NDVI for the total number of available pixels (n = 198) across all 996 drone-sentinel image pairs for the 2016 and 2017 growing season. The black line represents a simple linear 997 model describing the relationship, see Table S10 for details. c) Example visualisations from the Dryas-vetch 998 tundra at Area 2 - Bowhead Ridge for the 17 July 2017 showing ultra-fine-grain 0.013 m true colour RGB 999 imagery, 0.05 m native-scale drone NDVI, 10 m resampled drone NDVI, 10 m native Sentinel-2 NDVI, the 1000 absolute difference between resampled drone and Sentinel-2 NDVI, 30 m resampled drone NDVI, 30 m native 1001 Landsat 8 NDVI and difference between resampled drone and Landsat 8 NDVI.

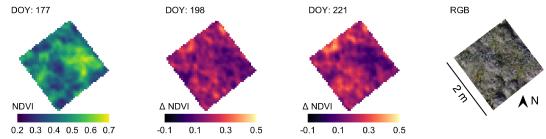


**Figure 3:** Spatial variation of vegetation greenness peaked at distances of ~0.5 m in both studied vegetation types, with little or no increase in the spatial dependence of greenness at distances above ~0.5 m. Figure shows example variograms. Overall spatial variation in greenness is higher in the Dryas-Vetch Tundra when compared to the Tussock-Sedge Tundra (a and c). Left panels: variograms for the Dryas-vetch and tussock sedge tundra plots in Area 2 for distances up to 5 m (a) and 45 m (c) at peak season in 2017. The light grey dotted lines in 1008 panel (a) indicate the subset of the distance range depicted in panel (c). The dark grey line in (c) indicates the wariogram models of both vegetation types from Areas 1, 2, and 4 during peak-season (26 and 28 July) in 2017 (see also Figure S1). Right panels: Dryas-vetch tundra with bare ground 1011 patches caused by cryoturbation and solifluction (c) and tussocks sedge tundra with distinctive patterns of 1012 tussocks interspersed by patches of willows and herbs (d).



**1014 Figure 4:** Fine-scale variation representing key ecological heterogeneity in tundra phenology was lost when 1015 aggregating from ultra-fine-grain drone to medium-grain satellite pixel sizes. We observed a logarithmic decay in 1016 variation (standard deviation) in the quadratic coefficient of simple growing season curves fitted to the eight 1017 vegetation plots in the 2017 season when aggregating the drone data across grain sizes (a). To provide an 1018 example of the underlying raw data, we visualised the pixel-by-pixel curves fitted to the time-series of pixels from 1019 the dryas-vetch tundra plot in Area 2 for a subset of three grain sizes (b). Here, each point represents a pixel 1020 NDVI value at a given day of year and grain size (indicated by colour). The transparent lines represent the simple 1021 quadratic curves fitted to each pixel across the time-series, again the colour of the line indicates the pixel's 1022 associated grain-size. See also Figure S8, which shows a random sample of nine curves for all grain sizes from 1023 the same study plot. Furthermore, to provide an example of the spatial distribution of the quadratic coefficient and 1024 how it changes across grain sizes, we plotted the respective rasters for Area 2 dryas-vetch tundra in panel (c). 1025 Similar patterns are found across all areas (a).





**Figure 5:** Time-series of ground-based plot mean standardised longest leaf lengths correlated well with 1029 drone-derived mean NDVI on Qikiqtaruk. a) Correlations between the mean standardised longest leaf length for 1030 all individuals across all monitored species and the drone-derived NDVI in the 2 m x 2 m ground-phenology plot 1031 for each area, vegetation types and year combination. The time-series of mean standardised longest leaf length 1032 (b) and drone NDVI (c) corresponding to the values in (a). Lines represent least-square regressions to illustrate 1033 the relationships for each area, vegetation type and year combination. A species-by-species version using 1034 absolute mean longest leaf length for each plot can be found in Figure S7. (d) As an example, we illustrate the 1035 drone-based NDVI observations by showing the start, midpoint and end of the timeseries for the 2 m x 2 m 1036 ground-validation plot in the tussock sedge tundra of Area 2 in 2017. The first time-point in (c) represents the 1037 greenness in the plot at the beginning of the time-series, the two subsequent plots show the relative difference in 1038 greenness to this first observation at the given day of year (DOY), and the final plot shows a true-colour image of 1039 the plot taken by drone on the 17 July 2017 (DOY 198).