

Bio-climatic factors drive spectral vegetation changes in Greenland

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Abstract.

The terrestrial Greenland ecosystem (ice-free area) has undergone significant changes over the past decades, affecting biodiversity. Changes in near-surface air temperature and precipitation have modified the duration and conditions of snowpack during the cold season, altering ecosystem interactions and functioning. In this study, we statistically aggregated the Copernicus Arctic regional reanalysis (CARRA) and remotely sensed spectral data on green vegetation, spanning from 1991 to 2023. We use principal component analysis (PCA) to examine key sub-surface and above-surface bio-climatic factors influencing ecological and phenological processes preceding and during the thermal growing season in tundra ecosystems. Subsequently, we interpreted spatio-temporal interactions among bio-climatic factors on vegetation and investigated bio-climatic changes dependent on latitude and topographical features in Greenland. Ultimately, we described regions of ongoing changes in green vegetation distribution.

Our results show that green vegetation has responded highly to the prevailing weather patterns of the past decades, particularly along West Greenland. The PCA effectively clustered bio-climatic indicators that co-vary with summer spectral vegetation, demonstrating the potential of CARRA for biogeographic studies. The duration of the thermal growing season (GrowDays) emerged as the pivotal factor across all ecoregions (with increases up to 10 days per decade), interacting with other bio-climatic indicators to promote summer vegetation growth. The lengthening of GrowDays is explained by reduced winter precipitation associated with warming (up to 1.5°C per decade). Significant decreases in snow height occur along with earlier snowmelt (up to 20 days per decade), leading to an earlier onset of GrowDays. We find that regions with shallower snowpacks, experiencing slower snowmelt rates during the ablation period, are linked with a higher soil water content in spring; this relation not only coincides with the greenest regions in West and Southwest Greenland, but also with regions where green vegetation has recently emerged. Such processes occur prior to GrowDays and were combined with summer weather conditions that favoured warmer and clear-skies that resulted in significant summer greening. The relatively warmer and drier summer conditions experienced in the northern and interior of the studied regions evidenced surface thawing and drying. Despite these summer bio-climatic interlinks green vegetation expanded northward and upward. Green vegetation has expanded in Northeast

Greenland by 22.5% with respect to 1991–2007 period, leading to new vegetated areas. We report little to no change in the
25 length and onset of the GrowDays along the coast in Northeast Greenland, in contrast with more pronounced changes inland
and at higher elevations, hence showing an elevation-dependent response (increases up to 5 days per decade per km elevation).
Our statistical outcomes and interpretations derived from reanalysis and remote sensing data that include uncertainties, are
corroborated by in situ studies conducted in the tundra region. The bio-climatic indicators and the associated insights serve
not only as a foundation for validating bio-climatic indicators from climate models to assess future changes in vegetation,
30 but they also advocate for the inclusion of permafrost dynamics schemes. This integration will enhance the quantification of
atmosphere-vegetation-permafrost-carbon feedback loops across terrestrial Greenland amid the evolving climate.

1 Introduction

The changing climate in the past decades has profound and rapid effects on Arctic ecosystems, with regional warming in
Greenland at nearly three times the global average (Rantanen et al., 2022). This rapid warming is causing significant changes
35 in the region's climate patterns and ecosystems. Jansen et al. (2020) highlight that the current era of abrupt climate change in
the Arctic is unprecedented in the context of the past several thousand years, leading to complex and varied responses in Arctic
vegetation. Myers-Smith et al. (2020) reveal the intricacies of "Arctic greening", where increased temperatures and earlier
snowmelt drive changes in plant growth and species distribution. These changes affect ecological interactions, such as shifts in
plant community composition and alterations in soil nutrient cycling, leading to feedback mechanisms involving snow cover
40 and surface albedo. Similarly, Huang et al. (2017) discuss the rate of change in vegetation productivity across northern high
latitudes, reiterating that the response to climate change is influenced by multiple factors, including soil moisture availability,
temperature variations, and the timing and extent of snowmelt.

Over the last three decades, community plant height has increased across the Arctic (Bjorkman et al., 2018). This has largely
been caused by plant community changes, in particular due to an increase in abundance and productivity of deciduous shrub
45 species – causing “shrubification” of the tundra (e.g., Mekonnen et al. 2021; Sturm et al. 2001). A large-scale study on the
interconnection between temperature, moisture, and various key plant functional traits at 117 Arctic locations over 30 years
of warming revealed a strong relationship between temperature and particular plant traits; however, soil moisture was a strong
factor determining the strength and direction of these relationships (Bjorkman et al. 2018). Changes in plant communities are
also reliant on the availability of soil moisture (e.g., Ackerman et al. 2017; Gamm et al. 2018; Power et al. 2024). Studies
50 on dynamic tundra vegetation future scenarios have suggested that the expansion of deciduous shrubs is favored by warmer
summers, whereas graminoids are more likely to increase in wetter conditions (van der Kolk et al., 2016).

Due to high latitude and continentality, summer soil moisture availability is particularly important in certain areas of Green-
land. In these drier areas, higher temperatures and less precipitation during the summer can potentially cause desiccation and
salt accumulation at the soil surface, with a negative effect on plant growth (Zwolicki et al., 2020). Therefore, it is expected that
55 an increase in temperature will unlikely lead to a striking increase in plant growth, mainly due to the lack of precipitation. For
instance, a study on growth responses of two widespread and dominant deciduous shrub species (*Salix glauca* L. and *Betula*

nana L.) in western Greenland revealed that both species have declined in growth since the 1990s, likely due to increasing water limitation (Gamm et al., 2018). However, increasing herbivory also plays a role, such as increased moth outbreaks (Post and Pedersen, 2008), growing muskoxen populations in West (Eikelenboom et al., 2021) and East Greenland (Schmidt et al., 2015) and increases in geese populations in East Greenland (Boertmann et al., 2015). Such studies are however based on small-scale analyses and contrast with observations of increasing shrub growth in other parts of the Arctic (Metcalf et al., 2018). Also, certain inland parts of Greenland are warmer and drier than most other areas of the Arctic, and will therefore respond differently to climate change, making their spatial representativeness unclear. The critical influence of soil water availability on future changes in tundra plant communities in Greenland should not be underestimated, and may also serve as an indicator for other drier Arctic regions, which may experience similar changes in temperature and precipitation. Additionally, while certain plant communities are generally better adapted to drier conditions, and have been observed to have increased with recent warming in the colder and drier High Arctic (e.g., Heijmans et al. 2022; Opała-Owczarek et al. 2018; Weijers et al. 2017), it is likely that certain species also become decreasingly temperature- and increasingly soil moisture-dependent during summer (Weijers, 2022).

Temperature, precipitation, as well as soil water availability during the growing season are a few of the climatic indicators contributing to vegetation changes (Migala et al., 2014). However, other climatic indicators, such as snowfall, snowmelt rate and timing and frost also play an important role even before the onset of the growing season (Cooper, 2014). The projected increase in temperatures during the cold season will likely have a different impact on vegetation than warmer temperatures during the growing season (Weijers, 2022). Increased snow depth during the cold season usually causes increased plant growth in the following summer, as more snow provides insulation, less frost damage and, depending on the snowpack characteristics, increase in water availability (e.g., Lamichhane 2021; Migala et al. 2014; Wang et al. 2024). A relevant characteristic of the snowpack is that deep snow requires more energy to equalise the cold content and the liquid water holding capacity to subsequently initiate and sustain melt than shallow snowpacks (Colbeck 1976; Musselman et al. 2017). As a result, deep snow often subsists for longer periods, potentially delaying the start of the growing season, which can hinder plant growth (Schmidt et al., 2015). On the other hand, the insulation provided by deep snow has also been demonstrated to promote increased microbial decomposition, enhancing the nutrient supply for the following growing season (e.g., Cooper 2014; Pedron et al. 2023; Xu et al. 2021). The higher amount of energy input needed to melt deep snow means that it melts later but also faster, which can cause nutrient loss through increased runoff. Concurrently, meltwater from relatively shallow snow percolates the soil more efficiently during the ablation period, in contrast with fast snowmelt that quickly saturates the soil surface and runs off (Stephenson and Freeze, 1974). These slow snowmelt rates allow water to remain in the soil for extended periods, which is critical for activating soil microbe communities. These microbes then produce nutrients that are vital for vegetation growth (Glanville et al., 2012). However, if snow is limited and precipitation is falling as rain rather than snow, the resulting ice conditions can have damaging effects on the vegetation (increased branch mortality and vegetation damage, Weijers 2022) and in soil nutrient cycling. Additionally, in exceptional years like 2018, the High Arctic experienced unusually large amounts of snow, resulting in extraordinarily delayed snowmelt. This made it very difficult for plants to grow and for animals to access resources (Schmidt et al., 2019). Such conditions will strongly influence the growth of plants, and have impacts throughout

the food chain, such as for the Svalbard reindeer (Le Moullec et al., 2020) and the caribou in West Greenland (Eikelenboom et al., 2021). Overall, the amount of snow and the coupling with temperature are highly important for plant growth, and plant community composition in the Arctic, and a Greenland-focused study assessing bio-climatic changes has not yet been made.

95 Grimes et al. (2024) has recently shown that the doubling of vegetation across ice-free Greenland is linked with warming. The warming observed in Greenland over recent decades has been associated with more frequent and intense weather patterns that promote widespread clear-sky conditions and the advection of relatively warm air masses from southern latitudes along West Greenland (Barrett et al., 2020). Weather patterns can be related to indices by analysing specific atmospheric variables over time and space. For instance, the North Atlantic Oscillation is driven by surface pressure configurations in the North Atlantic
100 (Hurrell et al., 2003), and the Greenland Blocking Index measures geopotential height in the mid-troposphere over Greenland (Hanna et al., 2016). Both indices are commonly utilized in climate studies to deduce influences on various components of the climate system in Greenland and vicinity (e.g., Bjørk et al. 2018; Olafsson and Rousta 2021). Therefore, how warming impacts other interlinked bio-climatic indicators through weather patterns requires further investigation.

In order to properly assess changes in bio-climatic indicators in Greenland, it is important to consider that soil water sources
105 in the region are mainly from precipitation, snowmelt and ground thaw. The combination of hard local geology with scouring by the ice sheet has shaped the landscape, and thin soils result in less prevalent thermokarst and low water retention (Anderson 2020). Therefore, meltwater flows rather freely, eventually gathering in low-lying areas to form lakes or drains towards the sea. Therefore, tundra vegetation develops in regions adjacent to such water bodies, eventually colonizing recently drained regions (e.g., Chen et al. 2023). Due to climate warming, not only surface, but also subsurface runoff has increased in the
110 Arctic (Rawlins and Karmalkar 2024). Given the heterogeneity in soil properties and water availability sources, sub-surface bio-climatic indicators, such as volumetric soil water and, potentially, sub-surface runoff, should be considered.

In this study, we analyse 32 years (1991–2023) of remotely-sensed Normalized Difference Vegetation Index (NDVI) data to gain a deeper understanding of the spatio-temporal patterns of spectral vegetation changes across ice-free regions of Greenland, extending beyond the representativeness of point-scale studies. We examine the combined effects of bio-climatic indicators
115 ranging from sub-surface factors (such as soil water availability) to above-surface factors (such as the thermal growing season, heat stress, and frost) with summer spectral greenness. We also extend our study of bio-climatic changes beyond the summer by examining indicators from the preceding winter and spring and assessing their combined impact with summer spectral greenness. Additionally, we explore historical trends of bio-climatic indicators individually and investigate their latitudinal and topographical sensitivity. Finally, we identify regions with changes in spectral greenness and in greenness distribution.

120 2 Data

2.1 Copernicus Arctic regional reanalysis

The Copernicus Arctic regional reanalysis (CARRA) system predominantly relies on the non-hydrostatic numerical weather prediction model HARMONIE-AROME (Bengtsson et al., 2017), laterally forced by ERA5. CARRA, with a spatial resolution of 2.5 km, assimilates the same observational datasets as ERA5 (Hersbach et al., 2020), supplemented by additional station

125 data from the national meteorological services involved within the CARRA domain. This study employed the CARRA-West domain, which encompasses Greenland. For the ice-free Greenland domain, the additional station data that CARRA assimilates is sourced from the Danish Meteorological Institute and Asiaq-Greenland Survey networks. However, snow depth observations are not provided and not assimilated by CARRA. According to the CARRA Full System documentation (Schyberg et al., 2020), ice cover extent remains constant throughout the entire reanalysis period (1991-2023). The Leaf Area Index (LAI) climatol-
130 ogy in CARRA is updated based on the multi-year mean values from the Moderate Resolution Imaging Spectroradiometer (MODIS) MCD15A2H C6 (Yang et al. 2006; Yuan et al. 2011), and these have been used to update the ECOCLIMAP cover types for Greenland. ECOCLIMAP-I (Masson et al., 2003) is the global database utilized to initialize the Surface Externalisée (SURFEX, Masson et al. 2013), the soil-vegetation-atmosphere transfer scheme within CARRA. SURFEX is a multi-layer surface model that computes specific schemes dependent on the surface type (e.g., vegetation, soil, snow), allowing soil water
135 phase changes and enabling runoff over frozen and unfrozen soil. This helps to better represent areas with permafrost and ice surfaces in Greenland as they are not well described in the present version of HARMONIE-AROME. Soil properties in CARRA are derived from the Harmonized World Soil Database (Nachtergaele et al., 2010). The CryoClim project has generated a satellite-derived product of snow extent, which provides access to data collected on a daily basis from 1982 to 2015. CryoClim is a worldwide, optical snow product that utilizes the historical Advanced Very High-Resolution Radiometer - Global
140 Area Coverage (AVHRR GAC) data (Stengel et al. 2020). In the context of CARRA, the CryoClim data is ultimately used due to its comprehensive coverage for the entire period up to 2015. The data providers assure that the data for the period post-2015 have been produced and arranged in collaboration with the CryoClim developers at the Norwegian Meteorological Institute. Despite the fact that neither snow depth nor snow extent is assimilated, van der Schot et al. (2024) demonstrate in a recent study that the agreement is strong between the snow water equivalent modelled by CARRA and a snow model utilizing in
145 situ observations in both the West and East coastal regions of Greenland. They report that CARRA is capable of successfully representing snow-related indicators, with correlation coefficients exceed 0.8 and mean absolute percentage errors less than 30%.

The derived precipitation from CARRA was taken from its underlying model forecast system and is not an assimilated product. Hence, to minimize the impact of the spin up, we followed the CARRA Full System documentation (Schyberg et al.,
150 2020), which suggests combining 12 h accumulated precipitation by the difference of precipitation at lead time 18 and 6 h from forecasts initiated at 00 UTC and 12 UTC. This procedure was used for determining liquid precipitation (time integral of rain flux) and total solid precipitation (time integral of total solid precipitation flux).

2.2 NOAA Climate Data Record for Normalized Difference Vegetation Index

Phenology studies in remote sensing utilize data collected by satellite sensors, which determine the spectrum of light absorbed
155 and reflected by majorly green vegetation. Specific pigments present in plant leaves exhibit a pronounced absorption of visible light wavelengths, particularly those in the red spectrum. Conversely, the leaves exhibit a strong reflection of near-infrared (NIR) light wavelengths. While numerous vegetation indices exist, one of the most prevalent is the Normalized Difference Vegetation Index (NDVI), which uses red and near-infrared bands. NDVI serves as a measure of spectral vegetation health and

spans from -1 to 1. Biologically, NDVI values close to +1 suggest a high density of greenness and robust vegetation health, while values near zero indicate barren land or surfaces with little to no vegetation, such as rocks or sand. Negative NDVI values are typically associated with water, clouds, or snow, likely signifying areas with no vegetation presence.

The National Oceanic and Atmospheric Administration (NOAA) Climate Data Record (CDR) using the AVHRR (Vermote et al. 2018) NDVI, Version 5 (hereafter AVHRR NDVI) and NOAA CDR using the Visible Infrared Imager Radiometer Suite (VIIRS, Vermote et al. 2022) NDVI, Version 1 (hereafter VIIRS NDVI) are jointly used in this study from 1991 to 2023 on a daily basis with grid resolution of 0.05 degrees (approx. 5.5 km in latitude and around 2.5 and 0.5 km between 60 and 85 degrees North, respectively). AVHRR NDVI is available until the end of 2013, and is thereafter continued by its successor VIIRS NDVI. The surface reflectance and the associated AVHRR and VIIRS NDVI take into consideration atmospheric corrections (e.g., total column of atmospheric water vapour, ozone, and aerosol optical thickness). According to AVHRR and VIIRS technical reports, the NIR channel is centred at different wavelengths (830 nm vs. 865 nm). As there is no overlapping period available in the NOAA CDR, potential mismatches between AVHRR and VIIRS NDVI cannot be discarded. However, AVHRR NDVI uses the MODIS Land-Sea mask and its cloud mask is spectrally adjusted using 10 years of MODIS data, with 90% match accuracy over land (Franch et al. 2017). As VIIRS will eventually replace MODIS for land science, MODIS is also used to calibrate VIIRS NDVI estimates (Skakun et al. 2018).

In addition to NDVI, both products provide quality control flags. While the AVHRR NDVI flags the entire domain for latitudes above 60 degrees as polar latitudes, the VIIRS NDVI implements more stringent quality control measures, effectively flagging clouds and snow cover at polar latitudes.

2.3 Climatic oscillation indices

A variety of analytic approaches, such as principal component analysis (PCA) or *k*-means clustering, are often utilized to characterize the North Atlantic Oscillation (NAO), with input data sourced either from reanalysis or station records. Here, the NAO derived from sea-level pressure applying PCA is used. In this study, the NAO index calculated applying the leading principal component derived from sea-level pressure anomalies within the Atlantic domain (20°N–80°N, 90°W–40°E) is provided by National Center for Atmospheric Research/University Corporation for Atmospheric Research (NCAR/UCAR) (Hurrell et al., 2003). This product is posited to yield a more comprehensive representation of NAO spatial patterns compared to indices based on specific terrestrial stations. Notwithstanding, it is noteworthy to acknowledge the dynamic nature of PCA-based NAO indices, being subject to ongoing refinement with the integration of new data.

The Greenland Blocking Index (GBI) is derived from 500 hPa geopotential height over the region (60°N–80°N, 80°W–20°W), retrieved from NOAA Physical Sciences Laboratory/Earth System Research Laboratories (PSL/ESRL) (Hanna et al., 2016). Both the NAO and GBI indices originate from the NOAA National Centers for Environmental Prediction (NCEP/NCAR) reanalysis dataset (Kalnay et al. 1996). Consequently, these climatic oscillation indices have undergone seasonal standardization against the baseline period of 1950–2000.

3 Methods

3.1 Spectral greenness

Arctic regions are characterized by sparse vegetation, that typically exhibit markedly low NDVI values, often as low as 0.15 (e.g., Gandhi et al. 2015; Liu et al. 2024), with dense shrubs above 0.5 (e.g., Walker et al. 2005), and signal saturation at around 0.7 (e.g., Myers-Smith et al. 2020).

As estimates integrated through time are less likely to be influenced by temporal sampling artefacts at high latitudes than metrics based on maximum NDVI (e.g., Myers-Smith et al. 2020), we started by calculating monthly integrated NDVI. Also, since our focus is on green vegetation, only daily NDVI pixel values higher or equal to 0.15 are considered. Then, we divide the monthly integrated NDVI by the total number of monthly observations (n , see Figure S1 for the interannual variability of n) to obtain the monthly NDVI. However, before 2014 and as described in Subsection 2.2, the AVHRR algorithm was less strict in its data quality control compared to VIIRS from 2014 onward, resulting in higher n before 2014 that lowers monthly NDVI. To address temporal heterogeneities, we adjusted n from the AVHRR period with the number of monthly observations acquired during the VIIRS period. From 2014 to 2023, we identified the minimum, maximum and average number of observations for each month. Hence, using these three quantities, we generated a consistent variability range from 1991 to 2013 to recalculate monthly NDVI, considering a similar number of observations as from 2014 to 2023. This procedure assumes that the environmental conditions (i.e. snow-cover, clouds and shadow) between 1991 to 2013 are similar to those between 2014 and 2023. The maps for the average number of monthly observations and the associated standard deviation for AVHRR and VIIRS period before and after the adjustment regarding n are shown in Figures S2-S5, respectively.

Pixels with monthly NDVI equal to or greater than 0.15, representative of the area covered by green vegetation, are used to estimate the extent of green vegetation. We applied this same definition to compare how often certain pixels were classified as areas with green vegetation from 2008 to 2023 versus the earlier period from 1991 to 2007. We refer to this comparison as changes in greenness distribution. Finally, we calculated a seasonally averaged NDVI, hereafter referred to as spectral greenness and interchangeably as green vegetation.

As both gridded products in this study have different spatial resolutions, spectral greenness from NOAA NDVI is spatially interpolated to the CARRA grid with resolution of 2.5 km.

3.2 Bio-climatic factors

The set of chosen bio-climatic indicators were inspired by previous work from Aalto et al. (2023) and Rantanen et al. (2023), who proposed and investigated bioclimatic indices in Finland and across the Arctic. Our study emphasizes Greenland, considering adapted thresholds and additional climatic factors.

Table 1. Brief description of the bio-climatic indicators derived in the study.

Bio-climatic Indicator	Description	Units
T _{2m}	seasonally averaged air temperature at the height of 2 m above the surface	°C
SWE _{MAX}	mass of liquid water from melting the snow per unit area	mm w.e.
SWE _{MAX} DOY	day of the year for SWE _{MAX}	day of the year
SnowDays	snow-covered days when SWE is higher than 10 mm w.e.	days
GrowDays	days with daily T _{2m} higher than 1 °C that does not belong to SnowDays	days
DegreeDays	sum of daily T _{2m} during GrowDays	K days
Onset	first day of GrowDays	day of the year
End	last day of GrowDays	day of the year
MeltRate	mean melt rate for ablation days between SWE _{MAX} DOY and Onset of GrowDays	mm w.e. day ⁻¹
Greenness	seasonally averaged monthly NDVI, as described in Section 3.1	unitless.
Snow	seasonally accumulated mass per unit area of snow and ice particles falling on the surface	mm w.e.
RainRatio	fraction of liquid precipitation out of the total precipitation	%
RainOnSnow	RainRatio higher than 50% in SnowDays	days
Rain	seasonally accumulated mass per unit area of rain falling on the surface, when RainRatio _≥ 50%	mm w.e.
VPd	vapour pressure deficit is the difference between the amount of water vapour in the air and the amount of water vapour the air could hold when it is saturated	hPa
VSI	seasonally averaged water equivalent of volumetric soil ice content	%
VSW	seasonally averaged volumetric liquid water in the soil	%
FrostDays	SWE lower than 10 mm w.e. in spring with negative daily T _{2m}	days
DroughtDays	days with precipitation lower than 1 mm w.e. lasting for more than 10 consecutive days	days
HeatDays	days exceeding the seasonal T _{2m} climatology for the period 1991-2023 by 2 SD	days
Longitude	distance east or west of the Greenwich meridian	degrees
Latitude	distance north of the equator	degrees
Elevation	vertical elevation above sea level	m a.s.l.
Surface slope	the inclination of the surface	degrees
Surface aspect	the slope direction	degrees

220 Surface slope is transformed into sine aspect (west-east orientation) and cosine aspect (north-south orientation), given its circular orientation. Positive values in sine (cosine) aspect indicate how much the slope is facing east (north), whereas negative values indicates how much the slope is facing west (south).

From CARRA daily-averaged snow water equivalent (SWE), we derived the maximum SWE (SWE_{MAX}), the day of the year SWE_{MAX} occurs (SWE_{MAX} DOY) and snow-covered days (SnowDays) when SWE > 10 mm of water equivalent (w.e.). It is

225 important to note that SnowDays are not necessarily continuous, with sporadic snow events occurring all year-round. van der Schot et al. (2024) provide a thorough validation of CARRA SWE with in situ observations across Greenland. They report that CARRA is capable of successfully representing snow-related indicators such SWE_{MAX} and SWE_{MAX} DOY.

We calculated GrowDays by considering days that do not belong to SnowDays, with daily-averaged 2 m air temperature (T_{2m}) > 1 °C. The onset (i.e. the first) and termination (i.e. the last) day of the year of GrowDays are also derived. The indicator
230 DegreeDays is obtained by summing up T_{2m} during the previously defined GrowDays. The daily rain ratio (RainRatio) is defined as the fraction of liquid precipitation of the total precipitation. SnowDays, in combination with RainRatio higher than 50%, are used to derive days with rain-on-snow (RainOnSnow) between January and July to investigate potential snowpack warming before the thermal growing season onset. SWE_{MAX} DOY and thermal growing season onset are used to determine the length of the snow melting period. During the snow melting period, we calculated daily changes of SWE from which we
235 derived days with negative SWE changes (SWE_{melt}Days) and the mean of the negative SWE changes (MeltRate).

Certain bio-climatic indicators used in this study are based on seasonal statistics using the definition of meteorological seasons: winter includes December to February (DJF), spring from March to May (MAM), summer from June to August (JJA) and autumn from September to November (SON). Spectral greenness, T_{2m} , RainRatio, the volumetric soil water and ice (SoilWater and SoilIce) and vapour pressure deficit (VPd) are seasonally averaged, whereas precipitation, snowfall (Snow) and
240 rainfall (Rain) are seasonally accumulated.

The vapour pressure deficit in summer (VPdJJA), which is the difference between the water vapour pressure of saturated air and the actual water vapour pressure in the air, was calculated to represent continentality. Continentality in summer is expressed by high temperatures and lower humidity due to large distances to moisture sources. This lack of moisture availability contributes to lower water vapour pressure, which, when combined with high temperatures, leads to higher VPd. A high VPdJJA
245 indicates a strong drying potential in the atmosphere, which can significantly influence evaporation rates and plant water stress (e.g., Grossiord et al. 2020; Yuan et al. 2019).

DroughtDays, the number of days with precipitation lower than 1 mm w.e. lasting for more than 10 consecutive days, is seasonally aggregated in spring and summer. HeatDays are also seasonally aggregated in spring and summer, and they consist of the number of days exceeding the seasonal T_{2m} climatology for the period 1991-2023 by two standard deviations (2SD). As
250 the 32-year period is fairly normally distributed, +2SD are approximately equivalent to 97.5th percentile. FrostDays in spring are derived in the absence of snow cover, jointly with negative T_{2m} days.

Spectral greenness was compiled for summer, in order to capture the period with maximum solar radiation in Greenland and avoid snow-covered patches. Given the fact that shadow areas heavily impact reflectance, latitudes higher than 75°N are not considered due to low sun elevation. We restricted our study area to West and Northeast Greenland, as steep mountains, deep
255 fjords, expansive glaciers, and extensive ice caps inhibit the method's applicability in Southeast Greenland.

3.3 Ecoregions

Greenland extends for approximately 23 degrees of latitude, with temperature and precipitation rates varying considerably across latitudes and coasts (Westergaard-Nielsen et al., 2020). Due to the semi-permanent Icelandic Low and the steep topog-

raphy, the Southeast coast receives more precipitation than the Southwest coast (e.g., Ettema et al. 2010; Fettweis et al. 2017).
260 In general, the West and East coasts exhibit different topographic features, from a topographically complex East contrasting
with predominantly glacially eroded regions in the West (e.g., Karami et al. 2017; Anderson 2020). Nevertheless, both coasts
comprise diverse fjord systems, that often channel the wind and shield inland areas against storms. Consequently, the north-
facing slopes and the leeward side of these inland mountain systems receive reduced precipitation. Such coast-inland gradients
are therefore complex, also influencing the distribution of permafrost and freshwater systems (e.g., Westergaard-Nielsen et al.
265 2018; Abermann et al. 2019). Not only precipitation, but also temperature, tends to decrease with latitude. Other factors known
to shape the coastal climate are prominent ocean currents (e.g., East Greenland and North Atlantic current) as well as sea ice
and fjord ice conditions (e.g., Westergaard-Nielsen et al. 2020; Shahi et al. 2023).

The Arctic tundra ecosystem, including Greenland, is typically separated at around 70°N into Low Arctic and High Arctic
based on climatic and vegetation differences (Bliss et al., 1973). Greenland has also been mapped according to hydrology, soil
270 pH, percentage of water cover, floristic provinces and bioclimatic subzones (e.g., Walker et al. 2005). The former mapping
partly relies on mean July temperature thresholds and positive degree monthly temperatures to classify subzones. However, the
 T_{2m} JJA has warmed at a median rate of approx. 1°C per decade since 1991 (Fig. S6), likely shaping plant community structure
and distribution. Eythorsson et al. (2019) also shows that Köppen-Geiger classification and snow cover frequency in the Arctic
have changed and will continue to change in the Arctic this century. In order to avoid time varying metrics, we split ice-free
275 Greenland into five ecoregions (Fig. 1) based on physiogeographic features, such as adjacent seas, ocean currents and ice caps,
with direct and indirect control on heat and moisture transport.

Ecoregion 1 is the narrow coast along the Baffin Bay in Northwest Greenland, including Sigguup Nunaa, Uummannaq fjord,
Nuussuaq Peninsula and Disko Island. Disko Island is known as the transition region between High Arctic to Low Arctic, with
a smooth transition of High Arctic to Low Arctic vegetation type in between ecoregions 1 and 2. Ecoregion 2 stretches from
280 Ilulissat to the Maniitsoq Ice Cap. This ice-free part is particularly widely stretched from West to East, with climates ranging
from maritime at the coast to continental in the dry interior. Ecoregion 3 encloses mainly Southwest Greenland along the
Labrador Sea until Nunarsuit, curving from the Labrador Sea to the North Atlantic. Ecoregion 4 comprises the mountainous
and southernmost end of Greenland, facing the North Atlantic. Southeast Greenland, a very narrow coast composed of steep
slopes, is the meeting point between the relatively cold East Greenland current and the relatively warm Irminger Current,
285 leading to very foggy conditions during the warm season (e.g., Gilson et al. 2024; Laird et al. 2024). The combination of this
region's complex topography with frequent cloud cover resulted in its exclusion from the analysis. Finally, ecoregion 5 spans
from Kangertittivaq (Scoresby Sound) to the North coast of Young Sound, including Daneborg and Zackenberg. The coast of
ecoregion 5 is also commonly affected by fog conditions. However, the coastal topography usually shelters inland regions. The
Stauning Alps, the large system of mountain ranges west of Kangertittivaq, is excluded due to its very rugged and complex
290 topography, with numerous rocky peaks and active glaciers in most valleys and only minor vegetation growth.

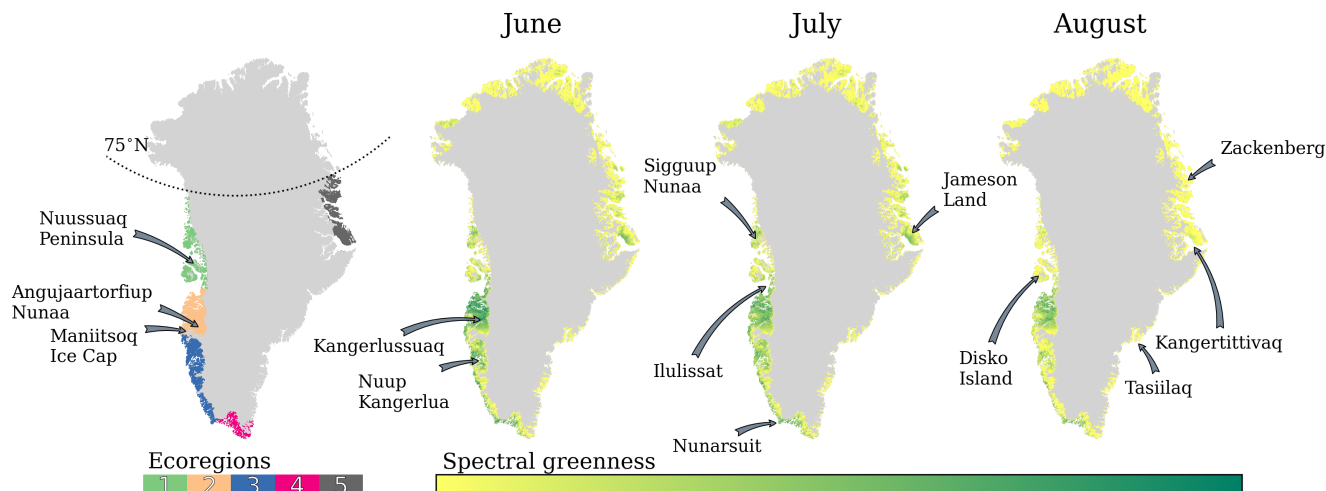


Figure 1. Ecoregions in ice-free Greenland; June, July and August averaged spectral greenness for the period 1991–2023. No scale shown in the colour bar because the aim is to illustrate spectral greenness patterns, not absolute values. Place names referenced in the study are indicated.

The onset of the thermal growing season is inherently linked with distance to coast, elevation and latitude. While distance to coast and elevation control precipitation and snow depth, latitude controls sunlight duration and near-surface air temperature. Due to less elevated and less topographically complex terrain, the thermal growing season starts earlier at the West (on average [5thpercentile, 95thpercentile], DOY: 140 [103, 172]) than at the East Coast (DOY: 171 [145, 204]). Moreover, both latitude and elevation are crucial in cooling the atmosphere, allowing snowfall to occur, which in turn marks the end of the thermal growing season. The ecoregion with the most GrowDays is ecoregion 4 (140[76, 198] days), followed by ecoregion 2 and 3 (119 [75, 145] and 121 [77, 164] days), with less than 100 GrowDays in ecoregion 1 and 5 (97 [56, 132] and 78 [47, 108]). Due to the proximity to the Atlantic cyclone track, ecoregion 4 receives the most precipitation, accumulating to SWE_{MAX} of 432 [114, 984] mm w.e. In contrast, ecoregion 2 receives about 25% of the precipitation received by ecoregion 4, with SWE_{MAX} of 143 [64, 325] mm w.e. This is due to the fact that the interior of ecoregion 2 is surrounded by high peaks in the south (e.g., Maniitsoq Ice Cap), serving as a physical barrier for poleward moisture transport.

The correlations between Greenness and North Atlantic Oscillation (NAO) index and the Greenland Blocking Index (GBI) are shown in Figure S7. NAO and GBI are well-known climate oscillations, frequently utilized to characterize the main atmospheric circulation pattern in the vicinity of Greenland. The NAO is determined by the surface pressure difference between the semi-permanent Subtropical (Azores) High and the semi-permanent Subpolar (Icelandic) Low (Hurrell et al., 2003), and its sign indicates the intensity of the North Atlantic jet stream. The phase of the NAO can explain most of the heat and moisture transported poleward, as well as temperature and precipitation anomalies in the periphery of Greenland (Björk et al., 2018). The GBI reflects the prevailing atmospheric circulation pattern, quantifying the strength of heat and moisture transport over

the Greenland region. Consequently, the GBI shows a strong correlation with near-surface variables in summer (e.g., Hanna et al. 2013; Hanna et al. 2015), such as spectral greenness.

3.4 Statistical methods

Principal Component Analysis (PCA, Pearson 1901; Lorenz 1956), often used on remotely sensed and environmental data (e.g., Mills et al. 2013; Yan and Tinker 2006), was employed to investigate the combined influence among bio-climatic indicators with summer greenness. The PCA (Pedregosa et al., 2011) solver was selected based on the input data shape. As the number of features in the input data is much less than the number of samples (geographic pixels), a classical eigenvalue decomposition on the covariance matrix was run. The classic PCA approach operates upon several assumptions, including I. linearity, which assumes that the relationships between variables can be adequately described by linear transformations; II. that there are no significant outliers in the data; III. that there is homoscedasticity, meaning that variables have equal variance. In order to overcome heteroscedasticity, we standardized all variables for each ecoregion, centering the distribution around 0 and scaling it to a standard deviation of 1. We used quartiles and the interquartile range (IQR) to filter out values beyond the upper ($Q_3 + 3 \times \text{IQR}$) and lower outer ($Q_1 - 3 \times \text{IQR}$) fence, with Q_1 and Q_3 as first and third quartile, respectively. Finally, we run a PCA for a set of bio-climatic indicators in every ecoregion between 1991 and 2023 until at least 90% of the cumulative explained variance is reached, omitting components contributing to minimal explained variance in order to accelerate the computation process.

As the classic PCA requires the variables to be linearly related, we calculated Pearson correlation coefficients to investigate bio-climatic indicators by ecoregion. However, Pearson correlation assumes that the data are stationary; that is, their statistical properties do not change over time. In order to avoid serial autocorrelation, we transform the data into non-stationary time series by linearly detrending the data before performing correlation. The calculated correlations are displayed in a correlation matrix, and bio-climatic indicators with similar correlations are sorted with hierarchical clustering. This helped to visually discern bio-climatic indicators with comparable statistical relationships and supported on the empirical reduction of indicators accounting for the relevant physical and the ecological processes on the tundra ecosystems, later used as part of the PCA. This will diminish "noise", redundancy and ultimately boost the clarity of interactions across atmosphere-biosphere-cryosphere.

Due to a change of satellite sensor from 2014 onwards, we also investigated how PCA performs interannually and whether there was a statistically significant change of the explained variance for years before and after 2014. The result is shown in Figure S8 for a set of 16 bioclimatic indicators, displaying that the two independent samples of explained variance have identical averages in all ecoregions, with a 95 % confidence level, as determined by a two-sample t-test.

We attempt a careful causal interpretation of the loading vectors from the first two principal components (PCs) of the PCA through biplots (Gabriel, 1971). Although these PCs account for most of the explained variance, their interpretation in terms of causality is limited by the nature of PCA as a descriptive statistical technique. For a cautious interpretation of the PCs, we examined not only the magnitude and direction of the loading vectors, but also trend maps of the involved bio-climatic indicators and literature on experimental studies.

We used the non-parametric Mann-Kendall (M-K) trend test (Hussain and Mahmud, 2019) to assess trend monotonicity and significance among bio-climatic indicators. However, to acknowledge autocorrelation in the greenness data, we computed the Hamed and Rao modified M-K test (Hamed and Rao, 1998), with a variance correction approach considering all significant lags
345 to improve trend analysis. The trend magnitude retrieved over decadal timescales corresponds to the Theil-Sen (T-S) estimator, a robust regression method that does not require the data to be normally distributed, hence less vulnerable to outliers than conventional methods. Under the null hypothesis that the slope is equal to zero, trends exhibiting confidence levels higher than 95% are highlighted and treated as significant trends.

4 Results

350 The green vegetation extent has progressed at different rates across summer months and ecoregions (Fig. 2). Given the colder temperatures in ecoregions 1 (first column) and 5 (last column), vegetation growth was generally not evident until July, contrasting with the more southerly located ecoregions. However, green vegetation extent was starting to increase already in June in recent years, particularly noticeable for 2019. In the southern ecoregions, the thermal growing season onset (Onset) was much earlier, and some vegetation growth began in the spring months. This is especially pronounced in ecoregion 2, due to
355 the typically shallow snow cover. By June, the vegetation was already quite developed, reaching its maximum extent in July. Ecoregion 2 comprises the largest vegetation extent, with vegetation covering approximately 80% of its area in 2015 and 2016.

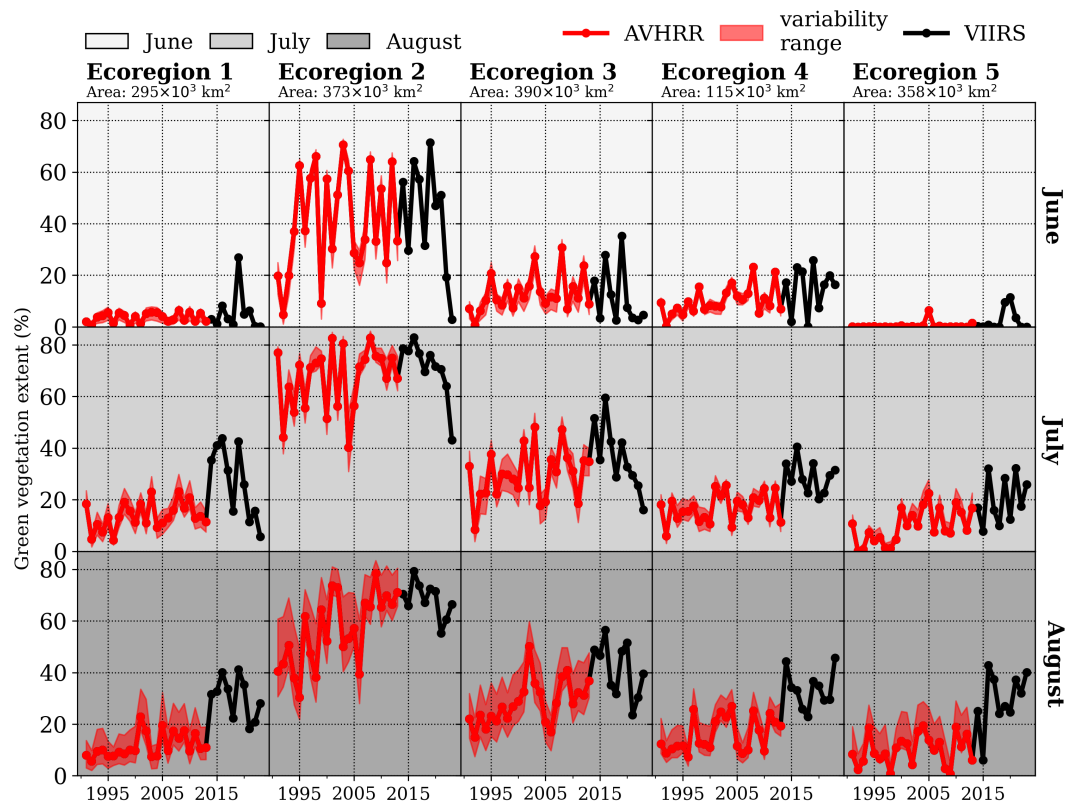


Figure 2. Development of green vegetation extent between 1991 and 2023 for June (upper row), July (middle row) and August (bottom row) across ecoregions based on AVHRR (red) and VIIRS (black). The AVHRR variability range is shaded in red, given the monthly minimum and maximum number of observations within VIIRS period (2014-2023).

It should be noted that prevailing weather patterns during summer months, like the North Atlantic Oscillation (NAO) and the Greenland Blocking Index (GBI), are highly correlated with spectral vegetation (Fig. S7). Therefore, summer weather patterns can accelerate or delay the maximum green vegetation extent given their link with temperature and precipitation. Correlations between green vegetation extent and summer GBI are investigated for three periods: AVHRR (1991-2013), VIIRS (2014-2023) and the full period (1991-2023), and are shown in Table S1. Positive and significant correlation coefficients ranging between 0.5 and 0.8 are found between ecoregion 1 and 4, generally with higher correlations for VIIRS than for AVHRR period. Green vegetation extent in ecoregion 5 is poorly correlated with the prevailing weather patterns during summer.

While the AVHRR 22-year trend evidence general expansion of green vegetation, the VIIRS 9-year trend evidence decreases, particularly in West Greenland (Table S2). However, due to high variability and small sample size, most trends in both periods are not significant. Significant long-term trends range from 2 % per decade in ecoregion 1 to approximately 6 % per decade in ecoregion 4.

4.1 Interconnectedness among bio-climatic indicators

The detrended Pearson correlation coefficients for ecoregion 2 is shown in a correlation matrix in Fig. 3. The magnitude of the statistical links vary across ecoregions, but the direction remains generally the same.

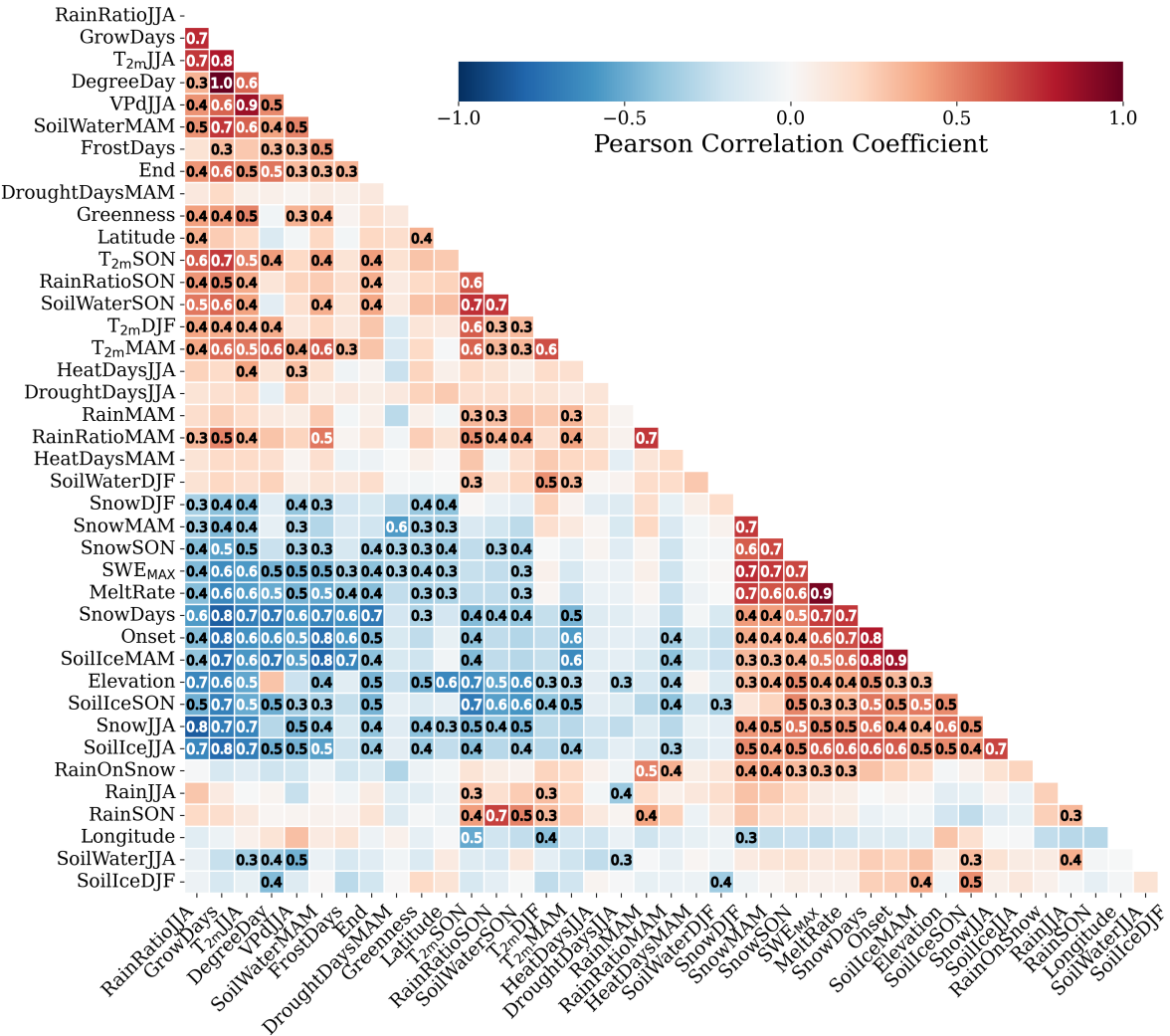


Figure 3. Correlation matrix for the bio-climatic indicators in ecoregion 2, including Elevation, Longitude and Latitude. The correlation coefficient is colour-coded and the absolute value noted for absolute correlation coefficients higher than 0.3. The abbreviations of the bio-climatic indicators are described in Section 3.2 and in Table 1.

We investigated the correlations among all the bio-climatic indicators, including physical features like elevation, latitude and longitude. These physical features are connected to climate attributes across ecoregions. For instance, the higher the elevation

and latitude, the lower the precipitation rates. It should be noted that these physical features are constant through time and were not considered when investigating the combined effect among bio-climatic indicators with greenness in the PCA.

375 A few bio-climatic indicators, such as DroughtDaysJJA, HeatDaysMAM, HeatDaysJJA, generally show correlations lower than 0.3, except DroughtDaysMAM that is negatively correlated with SnowMAM and with SWE_{MAX}. HeatDaysJJA is correlated with T_{2m} JJA, suggesting that the increased summer near-surface air temperature is associated with the increase of summer heat days. We excluded drought and heat indicators from the subsequent analysis, as greenness correlates more strongly with seasonal temperatures and precipitation amounts. Also, drought and heat variability is respectively explained by seasonal temperature and precipitation. However, it is important to highlight that increases of Heat and DroughtDays in spring and summer are reflected by increased near-surface air temperatures. RainOnSnow seems to be statistically linked with the increase of the RainMAM and RainRatioMAM, but also with SnowMAM, as RainRatioMAM represents a mix of rain and snow. In our trend analysis, it is noticeable, that RainOnSnow is increasing along East Greenland and FrostDays are locally increasing in West Greenland. However, summer spectral greenness is not statistically responding to these two bio-climatic indicators. Therefore, we removed them from further analysis. Interestingly, FrostDays is positively correlated with spring near-surface air temperature. The increase in FrostDays is also correlated with early disappearance of the snow-cover, partly related to shallower snowpacks and highly correlated with the decreasing volume of ice in the soil in spring (SoilIceMAM). SoilIce is to a large degree negatively correlated with the volume of water in the soil (SoilWater). Therefore, we decided to arbitrarily use SoilIce in winter (SoilIceDJF) and summer (SoilIceJJA) and SoilWater in spring (SoilWaterMAM) and autumn (SoilWaterSON) in the further analysis. Additionally, SnowDays and DegreeDays are not used since both are highly explained by GrowDays. While DegreeDays sum up T_{2m} during GrowDays, SnowDays is complementary of GrowDays, FrostDays and snow-free occasions with daily T_{2m} between 0 and 1 °C. Strong correlations between Rain and RainRatio are found in spring and autumn, but not in summer. Consequently, we will retain both Rain and RainRatio variables exclusively for the summer. Finally, MeltRate is removed as it is physically explained by the snowpack depth.

395 4.2 Bio-climatic indicators interlinked with greenness

PCA was used to investigate the combined influence among bio-climatic indicators with summer spectral greenness. 16 bio-climatic indicators were chosen to account for the relevant physical and the ecological processes on the tundra ecosystem, as described in the Sections 3.4 and 4.1. Figure 4 displays the combined influence of the 16 bio-climatic indicators based on the first two principal components across ecoregions. These two principal components account for most of the variability, ranging from 52% in ecoregion 2 to 65% in ecoregion 4 (Fig. S9). It takes more six to seven principal components to account for additional 30 to 40% of the explained variance.

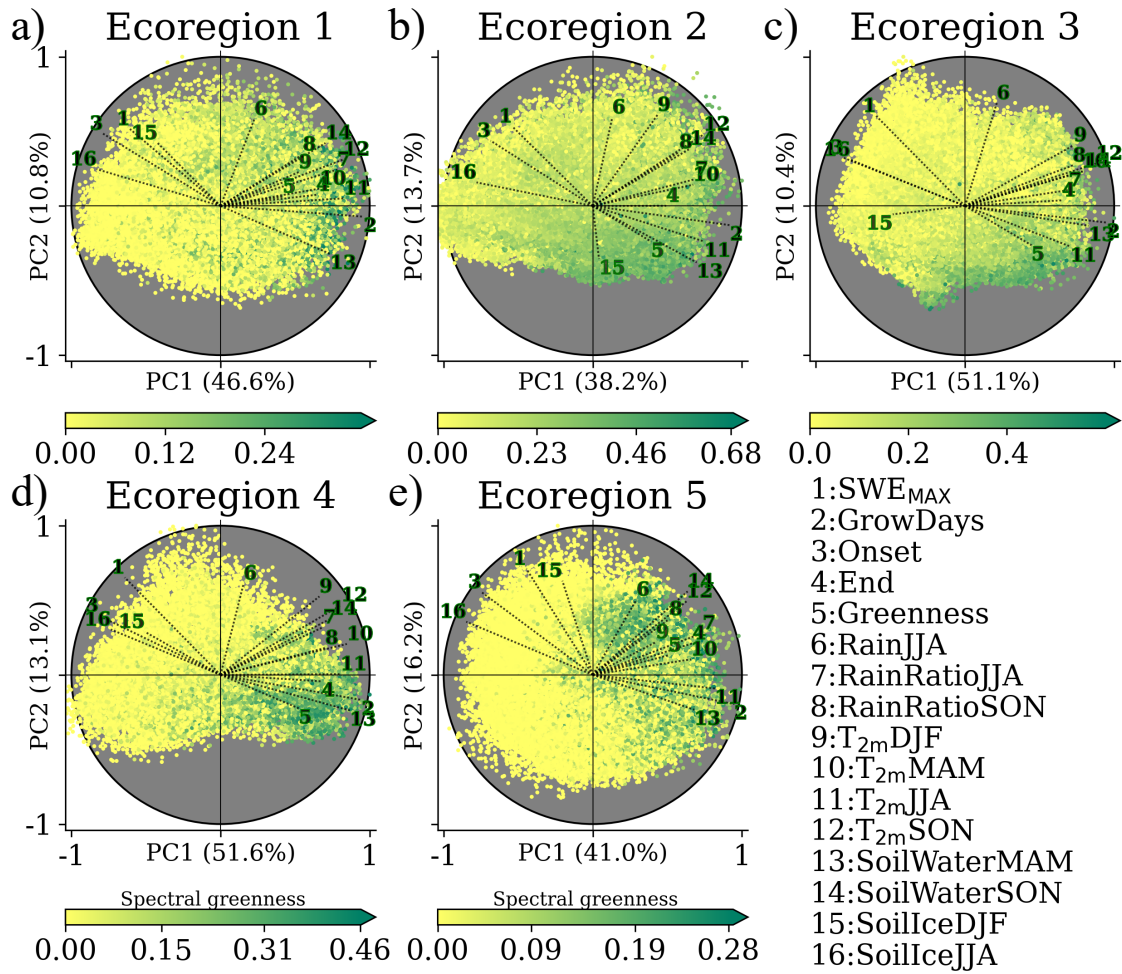


Figure 4. Biplot for scores between 1991 and 2023 for each ecoregion. The loading vectors are labelled and scaled by the maximum of each principal component. The scores are colour-coded based on the summer spectral greenness, with different scales to enhance greenness. The explained variance of the first (PC1) and second (PC2) component is labelled in the corresponding axis of the subplot. The 16 bio-climatic indicators are 1: maximum snow water equivalent (SWE_{MAX}); 2: total number of thermal growing days (GrowDays); 3 and 4: start (Onset) and termination (End) of GrowDays; 5: summer spectral greenness (Greenness); 6: rain in summer (RainJJA); 7 and 8: averaged rain ratio in summer (RainRatioJJA) and autumn (RainRatioSON); 9, 10, 11, 12: averaged 2-m air temperature in winter (T_{2m}DJF), spring (T_{2m}MAM), summer (T_{2m}JJA) and autumn (T_{2m}SON) 13 and 14: volumetric soil water in spring and (SoilWaterMAM) autumn (SoilWaterSON); 15 and 16: volumetric soil ice in winter and (SoilIceDJF) summer (SoilIceJJA). The abbreviations of the bio-climatic indicators are described in Section 3.2 and in Table 1. The spatial pattern of the averaged 1991–2023 scores for both components in every ecoregion, including their corresponding loadings, are shown in Fig. S10-S14.

According to the spatial maps of the first (PC1) and second component (PC2, Fig. S10-S14), PC1 is found to be highly controlled by the topography of the ecoregion, and is consequently related to temperature (and through that on elevation),

making GrowDays the bio-climatic indicator with the highest loading in all ecoregions, and therefore, the most significant contributor to the pattern represented by PC1. Through the analysis of the trend map for RainJJA (Fig. S15) and the spatial maps of PC2, we found that PC2 relates to precipitation and snow patterns, with SWE_{MAX} and RainJJA having the highest explanatory power.

These two principal components together largely capture Greenness distribution, as seen by scores with high summer spectral greenness often clustered in one specific quadrant of the biplot. As GrowDays is the most important loading (displayed by the longest loading vector) across ecoregions, and given its small load along PC2, GrowDays shows little dependence on precipitation and snow patterns.

In ecoregion 2, the ecoregion with the widest East-West coverage, summer greenness suggests to depend considerably on the snowpack of the preceding cold season (SWE_{MAX} loading vector opposite to Greenness loading vector). The decreasing trend of seasonal accumulated snow (SnowDJF, Fig. S16 and SnowMAM, Fig. S17) has led to SWE_{MAXDOY} to occur earlier (Fig. S18), which resulted in lower melting rates of the snowpack (MeltRate) as shown (Fig. S19). These shallow snowpacks are statistically linked to more water content in the soil in spring (SoilWaterMAM loading vector opposite to SWE_{MAX} loading vector). Additionally, the earlier snow depletion and thus earlier onset of the thermal growing season relates to enhanced spectral greenness (Onset loading vector opposite to Greenness loading vector). A wide atmospheric warming is shown by increases in T_{2mJJA} in most ecoregions (Fig. S6), which is also reflected on increases of RainRatioJJA (Fig. S20). These increases in RainRatioJJA do not seem to be linked to RainJJA between 1991 and 2023 (Fig. 4 and Fig. S15). Likewise, RainJJA do not seem to be related to higher greenness (orthogonal loading vectors in Fig. 4) across ecoregions. Also worth noting that RainJJA in the northern ecoregions is not in alignment with SoilIceJJA. In turn, the increase in T_{2mJJA} is generally aligned with less SoilIceJJA (opposed loading vectors). This is particularly evident in the northern ecoregions. The remaining ecoregions show localized increases in SoilWaterJJA, which is in contrast with the significant decreases in ecoregion 2 (Fig. S21). The same areas in ecoregion 2 show significant increases in VPdJJA (Fig. S22). Additionally, the increased T_{2mSON} is in alignment with the increase in RainRatioSON and SoilWaterSON, particularly in the southern ecoregions where the end of the thermal growing season occurs later.

We attempted to use the preceding autumn bio-climatic indicators to understand whether the start of the snow period could have played a role in the following growing season. However, the explained variance in PCA changed little (decreases of approx. 2-3% per ecoregion) and the relative importance of all loadings remained similar. Additionally, we correlated the interannual explained variance of the first two principal components with averaged climate oscillations (NAO and GBI) during the warm season (from March to September), spring and summer. We noted that the interannual change in explained variance is not significantly correlated with seasonal climate oscillations. In other words, a particular NAO or GBI phase does not boost the explained variance of the first principal components, maintaining the similar values interannually.

4.3 Coastal, latitudinal and altitudinal dependence on trends

The significant increase in length of the thermal growing season (GrowDays) across ice-free Greenland is shown in Figure 5. An evident increase in the number of GrowDays occurs in Southwest Greenland at low-laying regions below 600 meters above

sea level (asl). At a local scale, significant increase is also found at elevations above 1000 m asl, more specifically within Nuup Kangerlua (east of Nuuk) and Angujaartorfiup Nunaa (in between Maniitsoq Ice Cap and Kangerlussuaq). Such areas are in precipitation shadows, with reduced snow depths, but close to glaciers and ice caps. Along the narrow ice-free stripes in the Southeast, there is a modest increase of GrowDays (approx. 5 days per decade), at several elevations around Tasiilaq. The most pronounced increase in the number of GrowDays occurs along the coast facing the Denmark Strait.

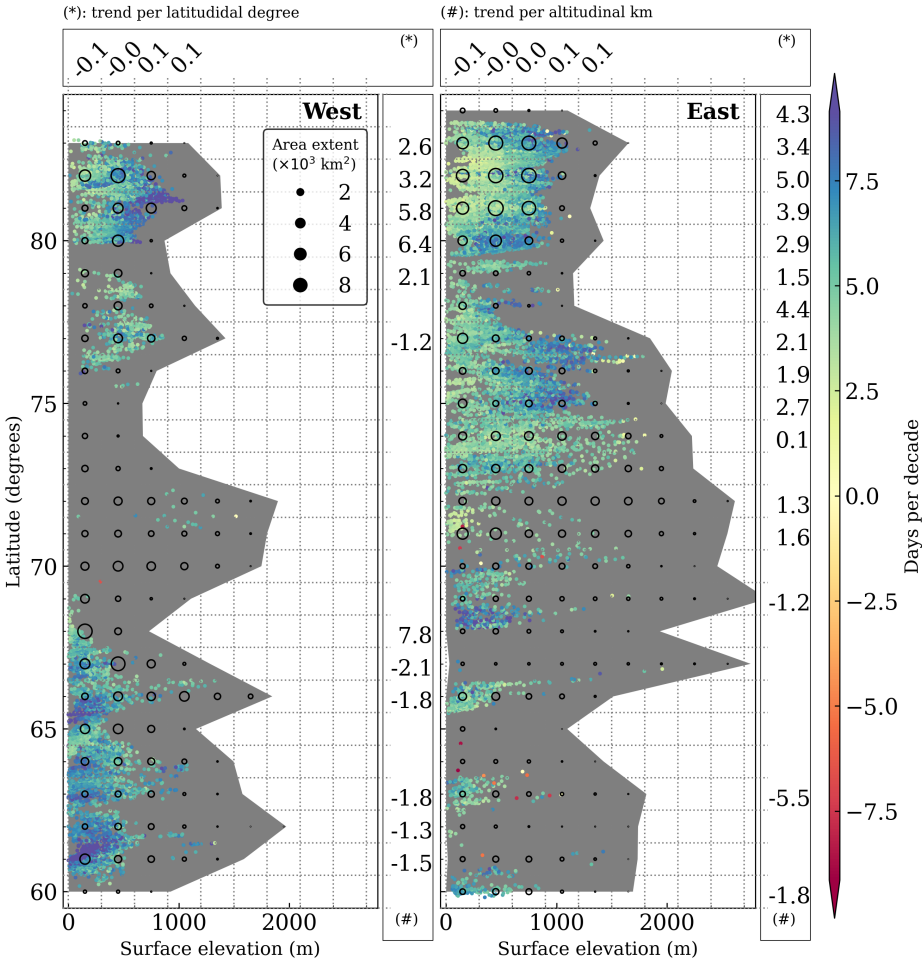


Figure 5. Significant trends for GrowDays (in days per decade) in the ice-free part of West (left panel) and East (right panel) Greenland. The trend's elevation dependency (in days per decade per altitudinal km) is binned in one degree of latitude and shown in vertical boxes marked with (#). The trend's latitudinal dependence (in days per decade per latitudinal degree) is binned every 300 m and shown in horizontal boxes marked with (*). The background grey shade displays the altitudinal extent of ice-free Greenland in the respective degree latitude, and the black circles represent the area extent by altitudinal and latitudinal bin. At least 50 pixels (approx. 312 km²) are required within each bin to compute its regression, otherwise not displayed. Trends are considered significant for confidence levels in the Mann-Kendall trend test higher than 95%, with the null hypothesis that the slope is equal to zero.

The vast and relatively flat ice-free Jameson Land (east of Ittoqqortoormiit, between 70° and 72°N), shows little evidence of GrowDays change within the past three decades. At the northernmost part of ecoregion 5 (75°N), areas at low elevations reveal the smallest increase of GrowDays in the ecoregion. This feature becomes even more pronounced in Greenland's northernmost regions, exhibiting the highest GrowDays elevation sensitivity (approx. 5 days per decade per km elevation), which is a contrasting elevation dependence in comparison with Southern Greenland. This tendency is modestly evident for the latitudinal sensitivity, mainly driven by high latitude and elevation trends: whereas GrowDays trends decrease with latitude in low-lying areas (< 300 m asl), GrowDays trends increase with latitude at higher elevations across North Greenland.

Trends for the onset of the thermal growing season resemble the trends in GrowDays, with earlier starts (approx. 8 days per decade) in southwest coastal Greenland and in the interior of Northeast Greenland. This comes as a consequence of shallower snow depths, that in combination with warming, has promoted longer snow-free periods. Thus, some areas of these ecoregions show increased trends in the number of frost days in spring.

The relationship between GrowDays and topographical features such as slope and aspect were further explored. As the surface slope is highly correlated with surface elevation, trends in GrowDays tend to significantly decrease with steepness. The dependence between GrowDays and surface aspect is rather complex, without a predominant slope orientation promoting GrowDays, in general. However, latitudes immediately south of Maniitsoq Ice Cap show increases of GrowDays in slopes with southwest orientation. On the East coast, a western slope orientation is particularly pronounced along Jameson Land, whereas northeast exposure appears favourable north of ecoregion 5. The dependence of the slope orientation for greenness changes is partly in alignment with the dependence of the slope orientation for GrowDays. Greenness trends increased in two latitudinal bands facing southeast in ecoregion 1 and 2. In Jameson Land a similar tendency for more greening is found towards southwest, while east facing slopes are preferred towards the northern part of ecoregion 5.

4.4 Greenness expansion and greening

Trends in summer spectral greenness are shown in Figure 6a. Significant greening occurs throughout Greenland, with pronounced greening across all ecoregions. Marked greening in ecoregion 1 is found in East Disko and northeast of Disko Bay. In ecoregion 2, the most pronounced greening is along the inland part of the Kangerlussuaq Fjord. The interior of Nuup Kangerlua shows the highest greening in ecoregion 3. The coastal Kujalleq municipality, facing Labrador Sea, exhibits substantial greening. There are two greening clusters in ecoregion 5, being Jameson Land in the south, and the interior of King Christian X Land, in the north. In contrast, decreases in summer greenness are shown along the Southwest coast from ecoregion 1 to ecoregion 3, and in the interior of ecoregion 2.

In order to assess which regions became greener due to spectral vegetation expansion, we detected whether a pixel meets the spectral greenness criterion annually from 1991 to 2023. Then, we evenly split the study period into two and counted the total number of summers with spectral greenness within the two sub-periods. The result (Fig. 6c) shows the number of summers that are spectrally green in the second sub-period (2008-2023) with respect to the first sub-period (1991-2007), hereafter called changes in greenness distribution. The map of changes in greenness distribution shows that a considerable part of summer trends in spectral greening result from an expansion of vegetation. With the support of such a map, we discern

that the relationship between changes in spectral greenness and its distribution are not linear. For instance, the central part of ecoregion 2 had as many summers of greenness in the second half as it had in the first half of the study period. Therefore, the changes in spectral greenness are either related to greening density of the existing vegetation or plant community change
 480 in ecoregion 2. The increased trend in spectral greenness of the remaining areas seems to be the result of spectral greenness expansion, likely due to the colonization of previously bare ground. The decreased trend in spectral greenness along the coastal southwest suggests that spectral vegetation is not as dense in the second sub-period as it used to be. Also, vegetation seems to be emerging directly adjacent to the ice-sheet.

Figure 6b combines the information of both maps by displaying significant changes in greenness as a function of latitude and
 485 latitude, colour-coded by the changes in greenness distribution. Ecoregions 3 and 4 show shrinkage in greenness at elevations lower than 500 m, while ecoregion 2 shows shrinkage up to 1000 m at certain latitudinal bands. In contrast, expansion in greenness is not only shifting upward but also northward across all ecoregions, with notorious upward advancements south of Kangerlussuaq towards Angujaartorfiup Nunaa in the west and in Jameson Land in the east.

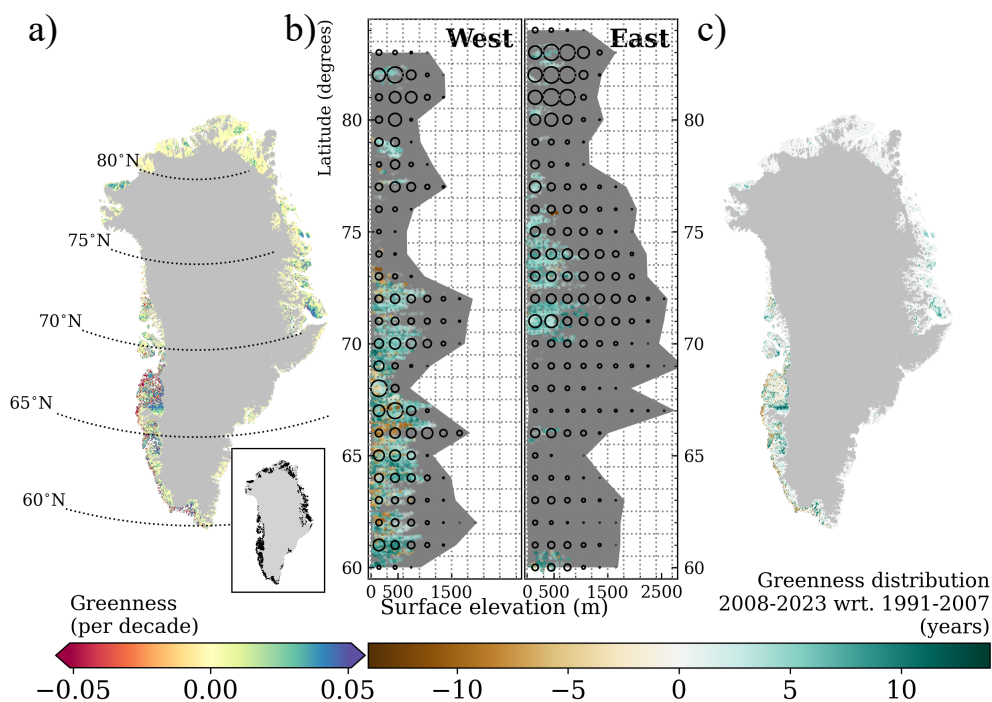


Figure 6. Trends in Greenness in Greenland (a). Significant trends are shown in the inset of (a) Trends are considered significant for confidence levels in the Mann-Kendall trend test higher than 95%, with the null hypothesis that the slope is equal to zero. Difference in summer greenness for the period 2008-2023 with respect to (wrt.) the period 1991-2007, called as changes in greenness distribution (c). Changes in greenness distribution as a function of latitude and elevation at locations with significant trends in spectral greenness in West and East ice-free Greenland (b). The background grey shade displays the altitudinal extent of ice-free Greenland in the respective degree latitude, and the black circles qualitatively represent the area extent by altitudinal and latitudinal bin.

According to Table 2, we show that ecoregion 2 experienced the highest greenness expansion at 44.2%, along with the highest greenness reduction at 33.4% between 2008-2023 compared to 1991-2007, resulting in an overall increase of 10.7% in vegetation greenness. Ecoregion 1 saw the largest increase in vegetation greenness at 22.2%, with a greenness expansion of 30.6%. Ecoregion 5 had the lowest greenness reduction at 2.7% and an overall increase in vegetation greenness of 19.8%. Ecoregions 3 and 4 also experienced increases in vegetation greenness, at 18% and 20%, respectively.

Table 2. Percentage of vegetation expansion and shrinkage, and ratio (fraction of expanded by shrank area) between 2008 and 2023 with respect to the period 1991–2007 in % of the total ecoregion area

	Ecoregion 1	Ecoregion 2	Ecoregion 3	Ecoregion 4	Ecoregion 5
Expansion	30.6	44.2	38.6	28.0	22.5
Shrinkage	8.4	33.4	20.5	7.9	2.7
Ratio	3.6	1.3	1.8	3.5	8.3

The southernmost and the northernmost ecoregions experience the highest expansion ratios, ranging from three times to eight times more expansion than shrinkage in ecoregion 4 and 5, respectively. Overall, vegetation expansion in ice-free Greenland increases two times faster than vegetation reduction between 2008-2023 compared to 1991-2007.

5 Discussion

5.1 Key findings and interpretation in the context of the current literature

Green vegetation has increased over time across Greenland, with an expansion rate of 2% per decade in ecoregion 1 to almost 5% per decade in ecoregion 4. When comparing the recent half of the time-series (2008–2023) to the distance half (1991–2007), the distribution of greenness has also changed. In ecoregion 3 and 5, greenness distributions expanded by almost two and eight times than the areas that shrank, respectively. Within the time-series, maximum green vegetation extent was observed in 2019, aligning with the end of a period of frequent, long-lasting and intense summer atmospheric blocking conditions in the vicinity of Greenland, conditions which promoted advection of relatively warm and humid air from the North Atlantic along West Greenland.

To better understand how bio-climatic indicators co-varied with spectral greenness between 1991 and 2023 across ice-free Greenland, a set of bio-climatic indicators including greenness were statistically aggregated. This was achieved by using principal component analysis (PCA) on remote-sensing data of spectral greenness and the output of a polar-adapted reanalysis. We demonstrated that the first two principal components account for most of the variability among the 16 combined bio-climatic indicators. These indicators were chosen to be of relevance to the ecological processes of tundra ecosystems. Given the fact that the chosen indicators are interlinked with supplementary indicators, we extended the interpretation to over 30 bio-climatic factors, with the support of climatology and trend maps. PCA effectively clustered bio-climatic indicators that co-vary with summer spectral greenness based on data from 1991 to 2023. Numerous indicators closely co-vary with near-surface

air temperature and topography (PC1), and less on precipitation patterns (PC2). The rank of relative importance of individual
515 bio-climatic indicators depends on ecoregion, with the number of days of the thermal growing season (GrowDays) being the
most relevant across all ecoregions, followed by soil ice during summer (SoilIceJJA) in the northern and soil water in spring
(SoilWaterMAM) in the southern ecoregions.

Our study found that areas related to green vegetation expansion in the northern ecoregions have experienced a rise in Soil-
WaterMAM along with declines in both spring soil ice content trends (SoilIceMAM) and maximum snow depth (SWE_{MAX}). In
520 Northwest Greenland, including ecoregion 1, regional exceptions of widespread increases in SWE_{MAX} with regional delays in
the onset of the thermal growing season (Onset) are found along coastal areas and not related to spectral greening. Conversely,
areas related to spectral greening are found linked with rising SoilWaterMAM, accompanied by higher spring temperatures
($T_{2m}MAM$) and earlier Onset. Despite regional trends on higher summer rainfall amounts (RainJJA, Niwano et al. 2021) in
northern Greenland, we did not find a clear link between greening and changes in RainJJA. Interestingly, trends in summer soil
525 water content (SoilWaterJJA) and soil ice content (SoilIceJJA) are both negatively related to near-surface air temperatures in
summer ($T_{2m}JJA$). This results as a consequence of surface thawing and subsequently increased evaporation caused by higher
vapour pressure deficits in these northern areas. The greening of the recently emerged vegetated areas in the northern ecore-
gions respond to different seasonal soil water contents. Greening in ecoregion 1 correlates best with SoilWaterMAM patterns,
similar to the remaining southwestern ecoregions. Conversely, ecoregion 5 is more closely connected with SoilWaterJJA, likely
530 due to a later onset of the GrowDays.

Southern ecoregions with significant decreases in SWE_{MAX} show early SWE_{MAX} day of the year (DOY) that leads to early
Onset. Despite increased amounts of fresh snow and fewer drought days in spring (winter), the decreasing trend in SWE_{MAX}
is attributed to reduced winter (spring) snowfall in West (East) Greenland.

For most ecoregions in ice-free Greenland, we find that snowpacks are becoming shallower, and consequently melt slowly,
535 but earlier in the season. This feature was mentioned by Musselman et al. (2017) and is attributed to global warming. Mussel-
man et al. (2017) explains that in Western North America regions with shallower snow are experiencing snow season contrac-
tions. Shallower snow is susceptible to snow season contraction because shallow snow requires less energy to initiate melt than
deeper snow. This earlier start of the ablation period occurs at a slower rate due to a combination of near-surface warming with
relatively low solar altitude angles. In contrast, for deep snowpacks that require more energy to initiate runoff, it is also more
540 likely for the snowmelt water to refreeze within the snowpack (Dingman, 2015). Therefore, early season slow snowmelt rates
in shallow snowpacks allow for efficient soil water percolation and subsequent water storage (Stephenson and Freeze, 1974).
The successful percolation of liquid water into soil plays a key role in tundra regions during the snow ablation period and start
of the growing season, as during this time soils are generally dry due to high drainage (Migala et al., 2014). Increased water
availability in the soil could stimulate dormant microbial communities and thus increase the decomposition of soil organic
545 matter, releasing soil nutrients (e.g., Glanville et al. 2012; Salmon et al. 2016; Xu et al. 2021). This in turn could prime the soil
for earlier and more efficient vegetation growth and colonization. The increased spring soil water content (SoilWaterMAM),
spring near-surface air temperature ($T_{2m}MAM$), and lengthening of the thermal growing season (GrowDays) indicated in our
results could therefore improve conditions for plant growth and colonization, especially in the southern ecoregions. Therefore,

it is expected that plants are more developed in early summer. Such conditions in conjunction with favourable weather patterns
550 in summer associated with increased $T_{2m}JJA$ and longer periods of solar radiation (Barrett et al., 2020), allowed for higher
greenness levels and more green vegetation distribution. The same weather patterns also brought more drought and heat days,
but apparently without an immediate negative impact on greenness.

The early onset of GrowDays has also contributed to local increases of freezing days (FrostDays) and rain on snow days
(RainOnSnow). However, the effect of FrostDays is not reflected on greenness levels. This could be related to freezing episodes
555 still during the dormant phase or rather short freezing episodes or a demonstration of certain plant community resilience (e.g.,
Gehrmann et al. 2020; Körner and Alsos 2008). Nevertheless, the early onset of GrowDays allows vegetation to be potentially
more active and responsive to solar radiation, particularly in the ecoregions in lower latitudes with longer sun exposure (Opała-
Owczarek et al., 2018).

The drier conditions in the interior of ecoregion 2 have led to substantial losses in volumetric soil water during summer
560 (SoilWaterJJA) with minimal changes in SoilIceJJA. We attribute the decrease in SoilWaterJJA to higher rates of evaporation
in the more continental areas of the ecoregion, supported by the significant increase in vapour pressure deficit (VPdJJA, Fig.
S22). The energy necessary to convert liquid water into water vapour (latent heat) cools down the soil. This is shown in our
results where the increase in $T_{2m}JJA$ and VPdJJA results in little changes in SoilIceJJA in ecoregion 2. This forces and limits
vegetation distribution towards the proximity of water bodies (Chen et al., 2023). Although there are no significant trends in
565 SWE_{MAX} in ecoregion 2, subsurface runoff from ground thaw and meltwater from nearby snow/ice bodies likely contribute to
the increase in SoilWaterMAM in the area. Episodic rainfall events in spring can also contribute to increased SoilWaterMAM.
However, no significant changes in accumulated rain nor in rain ratio in spring (RainRatioMAM) were found in the region.

We report little to no change in the length and onset of the GrowDays along the coast in Northeast Greenland. In situ
long-term measurements (e.g., Schmidt et al. 2023), state that some taxa may have reached their phenological limits despite
570 ongoing warming. Assmann et al. (2019) suggests that temperature and snowmelt explain the effects on spring phenology in
Zackenbergl, contrary to sea-ice break up in the Greenland Sea. However, the continuous southward transport of cold waters,
frequently with sea ice, through the East Greenland current likely stabilizes the onset of the GrowDays at the coast. This seems
to be corroborated given the dampened effect towards the interior and high elevations in Northeast Greenland, resulting in
elevation sensitivity.

575 Grimes et al. (2024) investigated land cover changes across Greenland by using Landsat images since the late 1980s and
found similar spatial patterns of vegetation change. For instance, they showed increases in coverage of vegetation southwest
of Kangerlussuaq. They attributed this increase of vegetation to receding lakes happening, at least, since 1995 (Law et al.,
2018). Similar to our findings, Grimes et al. (2024) detected increased vegetation cover in northeast of Kangerlussuaq. This
has been shown and modelled in other parts of the Arctic tundra (e.g., Bosson et al. 2023; Jones and Henry 2003). Specifically,
580 in ecoregion 2 and 5, green vegetation is not only expanding inland, but also upward. The interior of Greenland, less exposed to
frontal systems developing over the Atlantic and with meltwater availability, seems to be a more favourable area for vegetation
growth as shown in ecoregion 1 and 2. Increasing greenness levels were also found with a tendency for slopes facing southeast
in ecoregion 1 and 2. According to Grimes et al. (2024), the retreat of vegetation in front of the Maniitsoq Ice Cap is leading

to the exposure of bedrock. Additionally, the less dense summer vegetation in coastal ecoregion 2 and along ecoregion 3 is suggested by Grimes et al. (2024) to be related to increases in freshwater, likely due to increased river discharge. A small-scale study, north of Kangerlussuaq, reports declining growth of deciduous shrubs (Gamm et al., 2018) since the 1990s. A similar signal is seen regionally in our results. They reported that the decrease is likely due to water soil scarcity, being a markedly pronounced negative trend for SoilWaterJJA in the region. The derived spatio-temporal patterns of summer rainfall (Fig. S15) and rain ratio (Fig. S20) are also in agreement with literature (e.g., Huai et al. 2022; van der Schot et al. 2023), especially on the significant increase of the rain ratio in North and West Greenland in summer and autumn. This consistency with other studies demonstrates the potential of the Copernicus Arctic regional reanalysis (CARRA) for biogeographic studies by extending insights from experimental studies into large-scale.

The widespread summer spectral greening could be due to encroachment of vegetation on previously bare surfaces and changes in plant community composition at certain sites (Grimes et al., 2024). Spectral greenness correlates best with biophysical properties, such as leaf area index (Myers-Smith et al., 2020). Therefore, we may argue that the spectral greening is generally related to tundra shrub expansion throughout the past three decades, as early proposed by Sturm et al. (2001).

5.2 Significance and implications

Longer thermal growing seasons are shown across Greenland between 1991 and 2023. Longer thermal growing seasons, associated with higher near-surface air temperatures have in the studied period favoured green vegetation growth and expansion. However, further investigation is required to comprehend the impacts on green vegetation and ecosystem functioning in regions that have been facing freezing conditions due to reduced snow cover, earlier onset of the thermal growing season as well as in regions exposed to heat stress conditions and changes in precipitation patterns throughout the thermal growing season. Given the reportedly significant decreases in snow cover, the surface albedo is lower for longer periods, facilitating more energy absorption and enhancing surface warming. The observed changes in greenness distribution enhance the surface albedo feedback, with varying effects that extend beyond the growing season and depended on the vegetation type (e.g., Blok et al. 2011; Loranty et al. 2011).

The surplus of the surface energy budget leads to surface warming and promotes surface thawing, particularly in the northern ecoregions. However, depending on the vapour pressure deficit and the vegetation canopy, the excess of surface energy can be used for latent heat release, which in turn will cool the surface (Heijmans et al., 2022). The increase in green vegetation drives at first to greater carbon sequestration. However, if the increase in vegetation causes substantial surface thaw, the net effect could trigger the release of carbon, offsetting the compensation of carbon sequestration from vegetation (Glanville et al., 2012).

The terrestrial Arctic biosphere is an important regional source of primary biological aerosol particles (PBAPs), highly correlated with near-surface air temperature and surface vegetation. These aerosol particles were found to play an important role in cloud formation, specifically in the Arctic with low aerosol concentrations (e.g., Pereira Freitas et al. 2023; Sze et al. 2022). Therefore, the increased near-surface air temperature and changes in vegetation can significantly impact cloud properties, such as cloud phase, radiative properties, cloud lifetime, and precipitation patterns, which in turn impact the surface conditions, including the surface energy budget. Additionally, low clouds and fog are also very likely to become more frequent in certain

coastal parts of Greenland due to decreasing sea ice (Song et al., 2023). The potential warming and shading conditions were shown through an experimental study in West Greenland to reduce carbon sequestration from vegetation (Dahl et al., 2017).

620 Water droplets from fog can effectively be retained by tundra vegetation and are not accounted as a water source. This interaction between fog, vegetation and soil conditions should be better investigated particularly for coastal tundra vegetation. Such PBAPs can also be advected towards snow- and ice-covered regions, for instance the Greenland ice sheet, contributing to the surface darkening and enhancing algae growth (Feng et al. 2024) which again leads to increased melt, particularly of the ice bodies in the vicinity of densely vegetated regions.

625 Longer thermal growing season in regions with shallow soils could have also significant large-scale implications for biodiversity. Prolonged warmth may foster the proliferation of shrubs, leading to increased "shrubification" and potentially resulting in the homogenization of species compositions across these landscapes (Myers-Smith et al., 2011). In other regions with deeper and frozen soils, the active layer deepening could favour graminoids due to longer roots (van der Kolk et al., 2016). In case of permafrost degradation with deep infiltration (Liljedahl et al., 2016), graminoids would also be favoured. This ecological shifts

630 might will also affect animal communities such as birds (Boelman et al., 2015) and arthropods (Høye et al., 2018). Ultimately, the increased growing season could create favourable conditions for invasive species to establish and spread, further threatening the native biodiversity and altering the delicate balance of these unique environments (e.g., Elmendorf et al. 2012; Pearson et al. 2013).

Our study determines a set of bio-climatic indicators that have been shown relevant for spectral greenness. The statistical

635 interlink among these indicators is confirmed in experimental studies across the Arctic (e.g., Chen et al. 2023; Gamm et al. 2018; Grimes et al. 2024; Huai et al. 2022; Migala et al. 2014; Musselman et al. 2017; Opała-Owczarek et al. 2018; Schmidt et al. 2023; Stephenson and Freeze 1974; van der Schot et al. 2024), allowing the interpretation of our outcome to be expanded to large-scale, with apparent features dependent on the ecoregion and latitude. Such insights can now be used to validate whether the same bio-climatic indicators interdependence is captured during the historical period of global climate models.

640 This would guarantee more confidence in the use of these indicators for the study of future vegetation changes across Greenland under a changing climate.

5.3 Study limitations and future research directions

There are limitations in the use of NDVI for the characterization of changes in plant communities (e.g., Myers-Smith et al. 2020). This includes that NDVI is good at capturing plant communities with a high composition of shrubs (e.g., Blok et al.

645 2011), but it is not as good at detecting communities with low infrared reflectance or sparsely vegetated areas. Our methods in combination with proposed approaches as in Karami et al. (2018) and Rudd et al. (2021), who categorized tundra vegetation classes across ice-free Greenland, would allow an optimal assessment of spatio-temporal changes among plant communities. However, the high spatial resolution of optical satellite images from Landsat 8 and Sentinel 2 are only collected for approximately one decade. Another limitation regarding the NDVI analysis is that pixels associated with certain vegetation types (e.g.,

650 wet tundra) may be misinterpreted by adjacent water bodies, likely influencing spectral vegetation extent and trend magnitudes of certain areas, such as discussed for ecoregion 2. Additionally, certain low-lying strips near fjords are very narrow, potentially

causing errors in pixel reflectance calculations due to limited spatial resolution. Remote-sense NDVI products highly depend on the weather conditions in order to retrieve surface reflectance. Occasions with snow, shadows and clouds are thus assumed to be evenly distributed through time. The NDVI datasets employed in this study are sourced from two satellite products processed by NOAA, each utilizing a different type of sensor. Due to the absence of a temporal dataset overlap, the assessment of uncertainties was limited and potential for mismatches between the datasets cannot be discarded. This lack of a common calibration period raises concerns about the reliability of long-term time integrated NDVI analysis. Also, spectral greenness highly responds to the prevailing atmospheric circulation patterns. An exceptional period in frequency and intensity of anti-cyclonic activity between 2010 and 2019, promoted advection of relatively warm and humid air from the North Atlantic towards South-west Greenland (Silva et al., 2022). Such periods have favoured exceptional vegetation growth across western ecoregions as shown in our results. Despite the frequency of prevailing atmospheric circulation patterns, there is a superimposed warming signal, with less cold conditions likely promoting vegetation growth that is poorly captured due to cloudiness.

The distribution of soil nutrients are also highly influenced by topography (not only elevation, but also relief and aspect), but not entirely reflected on the vegetation changes in large-scale in our study. According to Anderson (2020), organic rich soils in Greenland generally accumulate on north facing slopes, with little to none on the south facing slopes as a result of precipitation patterns, whereas in valley bottoms and at slope breaks, thicker fen-like, organic rich deposits accumulate. Even though we have investigated how vegetation and bio-climatic indicators are changing as a function of latitude, elevation, slope and aspect potential influences due to relief and aspect apparent in our results, potentially more evident at the local scale due to less spatial heterogeneity.

Our results show that summer spectral greenness appears statistically unresponsive to changes in rain on snow days (Rain-OnSnow) and below zero temperatures (FrostDays) during spring. Future work could rather focus on the analysis of extreme events and their impacts on greenness.

Even though CARRA is able to capture spatio-temporal changes on relevant bio-climatic indicators influencing spectral greenness, most implications, such as the effect of shrub canopies on the ground conditions are likely not captured. This implies that potential feedback loops (e.g., Hallinger et al. 2010; Barrere et al. 2018), where shrub growth and expansion result in more snow trapping during winter, thereby enhancing winter soil insulation (Lamichhane, 2021), increased microbial activity (Wang et al., 2024) along with greater shading in the following summer (Blok et al., 2010) cannot be properly assessed yet. Additionally, the vegetation type was recently considered as a strong predictor of summer surface latent and sensible heat fluxes (Oehri et al., 2022). A better representation of the permafrost extent and active layer thickness together with the inclusion of dynamic tundra vegetation models within CARRA could be beneficial to deepen our knowledge on interactions among atmosphere, vegetation, carbon and nitrogen cycling, water and permafrost dynamics.

Permafrost areas will continue to likely be locations for future vegetation expansion (Chen et al., 2023), especially under the current trend of decreased summer precipitation. Moreover, permafrost thawed areas are also susceptible to fast drying (Liljedahl et al., 2016) and potentially sudden vegetation changes. Ultimately, plants can fixate along streams and small lakes as future land ice melt will continue to provide sediments and nutrients through runoff (Migala et al., 2014).

6 Conclusions

Our study aimed to better understand the long-term, large-scale interactions among various bio-climatic indicators and their collective effects with summer spectral greenness in ice-free Greenland. This study utilized remote sensing Normalized Difference Vegetation Index and bio-climatic indicators from the Copernicus Arctic regional reanalysis between 1991 and 2023.

690 Bio-climatic changes are influenced by a complex set of factors, not only centered in summer, but also dependent on winter and spring atmospheric temperatures, precipitation, solar radiation, soil properties, and soil water availability.

We conclude that regions under green vegetation expansion in ice-free Greenland are associated with reductions in winter precipitation. The resulting shallower snowpacks melt earlier in the season but slower. This slow snowmelt rate allows the ground to retain more liquid water during the ablation period. Such conditions occur prior to the start of the thermal growing season are mentioned in experimental studies to facilitate vegetation growth. Longer thermal growing seasons accompanied by prevailing summer weather patterns, with its peak in 2019, that promoted warmer and clear-sky conditions over the past decades also contributed to vegetation growth.

The spatio-temporal changes in summer greenness distribution depend on ecoregion, elevation and latitude. Overall, the bio-climatic changes during the study period led to more vegetation expansion, particularly towards the interior and northward. Ultimately, this study encourages the incorporation of dynamic tundra vegetation schemes to improve our knowledge on deeper interactions among atmosphere, vegetation, carbon and nitrogen cycling, water and permafrost dynamics, particularly for future projections.

Data availability. The Normalized Difference Vegetation Index CDR used in this study was acquired from NOAA's National Centers for Environmental Information (<http://www.ncei.noaa.gov><http://www.ncei.noaa.gov>). This CDR was originally developed by Eric Vermote and colleagues for NOAA's CDR Program.

Schyberg et al. (2020) was downloaded from the Copernicus Climate Change Service (2024). The results contain modified Copernicus Climate Change Service information 2024. Neither the European Commission nor ECMWF is responsible for any use that may be made of the Copernicus information or data it contains.

The North Atlantic Oscillation and Greenland Blocking Index data were obtained from the NCEP/CPC and the PSL/ESRL, respectively. Both climate oscillations were seasonally standardized relative to the period 1950-2000.

Author contributions. The inspiration for the paper was brought by BSW, EMB, IGA, JA and NdV, the concept and methodology was developed by TS, the original paper draft was written by TS, the data were processed and analyzed by TS, all authors contributed to the interpretation of results as well as reviewing and editing the final paper draft.

Competing interests. The authors declare that they have no conflict of interest.

715 *Acknowledgements.* The University of Graz is acknowledged for support of publication costs. Brandon S. Whitley, Elisabeth M. Biersma and Natasha de Vere have received funding from the Carlsberg Foundation. The main author would like to acknowledge the use of OpenAI's ChatGPT for assisting in the writing and editing of this manuscript. The chatbot was utilized to enhance the clarity and readability of the text. A special thanks to Inger Greve Alsos and Therese Rieckh for their valuable suggestions.

References

- 720 Aalto, J., Lehtonen, I., Pirinen, P., Aapala, K., and Heikkinen, R. K.: Bioclimate change across the protected area network of Finland, *Science of the Total Environment*, 893, 164 782, <https://doi.org/10.1016/j.scitotenv.2023.164782>, 2023.
- Abermann, J., Van As, D., Wacker, S., Langley, K., Machguth, H., and Fausto, R. S.: Strong contrast in mass and energy balance between a coastal mountain glacier and the Greenland ice sheet, *Journal of Glaciology*, 65, 263–269, <https://doi.org/10.1017/jog.2019.4>, 2019.
- Ackerman, D., Griffin, D., Hobbie, S. E., and Finlay, J. C.: Arctic shrub growth trajectories differ across soil moisture levels, *Global Change*
725 *Biology*, 23, 4294–4302, <https://doi.org/10.1111/gcb.13677>, 2017.
- Anderson, N. J.: Terrestrial ecosystems of West Greenland, *Encyclopedia of the World's Biomes*, 1, 465–479, <https://doi.org/10.1016/B978-0-12-409548-9.12486-8>, 2020.
- Assmann, J. J., Myers-Smith, I. H., Phillimore, A. B., Bjorkman, A. D., Ennos, R. E., Prevéy, J. S., Henry, G. H., Schmidt, N. M., and Hollister, R. D.: Local snow melt and temperature—but not regional sea ice—explain variation in spring phenology in coastal Arctic
730 tundra, *Global Change Biology*, 25, 2258–2274, <https://doi.org/https://doi.org/10.1111/gcb.14639>, 2019.
- Barrere, M., Domine, F., Belke-Brea, M., and Sarrazin, D.: Snowmelt events in autumn can reduce or cancel the soil warming effect of snow–vegetation interactions in the Arctic, *Journal of Climate*, 31, 9507–9518, <https://doi.org/10.1175/JCLI-D-18-0135.1>, 2018.
- Barrett, B. S., Henderson, G. R., McDonnell, E., Henry, M., and Mote, T.: Extreme Greenland blocking and high-latitude moisture transport, *Atmospheric Science Letters*, 21, e1002, <https://doi.org/10.1002/asl.1002>, 2020.
- 735 Bengtsson, L., Andrae, U., Aspelien, T., Batrak, Y., Calvo, J., de Rooy, W., Gleeson, E., Hansen-Sass, B., Homleid, M., Hortal, M., Ivarsson, K.-I., Lenderink, G., Niemelä, S., Nielsen, K. P., Onvlee, J., Rontu, L., Samuelsson, P., Muñoz, D. S., Subias, A., Tijm, S., Toll, V., Yang, X., and Køltzow, M. Ø.: The HARMONIE–AROME model configuration in the ALADIN–HIRLAM NWP system, *Monthly Weather Review*, 145, 1919–1935, <https://doi.org/10.1175/MWR-D-16-0417.1>, 2017.
- Björk, A., Aagaard, S., Lütt, A., Khan, S., Box, J., Kjeldsen, K., Larsen, N., Korsgaard, N., Cappelen, J., Colgan, W., Machguth, H., Andresen,
740 C. S., Y, P., and H, K. K.: Changes in Greenland's peripheral glaciers linked to the North Atlantic Oscillation, *Nature Climate Change*, 8, 48–52, <https://doi.org/10.1038/s41558-017-0029-1>, 2018.
- Bjorkman, A. D., Myers-Smith, I. H., Elmendorf, S. C., Normand, S., Rüger, N., Beck, P. S., Blach-Overgaard, A., Blok, D., Cornelissen, J. H. C., Forbes, B. C., et al.: Plant functional trait change across a warming tundra biome, *Nature*, 562, 57–62, <https://doi.org/10.1038/s41586-018-0563-7>, 2018.
- 745 Bliss, L. C., Courtin, G., Pattie, D., Riewe, R., Whitfield, D., and Widden, P.: Arctic tundra ecosystems, *Annual Review of Ecology and Systematics*, pp. 359–399, 1973.
- Blok, D., Heijmans, M. M., Schaepman-Strub, G., Kononov, A., Maximov, T., and Berendse, F.: Shrub expansion may reduce summer permafrost thaw in Siberian tundra, *Global Change Biology*, 16, 1296–1305, <https://doi.org/10.1111/j.1365-2486.2009.02110.x>, 2010.
- Blok, D., Schaepman-Strub, G., Bartholomeus, H., Heijmans, M. M., Maximov, T. C., and Berendse, F.: The response of Arctic vegetation
750 to the summer climate: relation between shrub cover, NDVI, surface albedo and temperature, *Environmental Research Letters*, 6, 035 502, <https://doi.org/10.1088/1748-9326/6/3/035502>, 2011.
- Boelman, N. T., Gough, L., Wingfield, J., Goetz, S., Asmus, A., Chmura, H. E., Krause, J. S., Perez, J. H., Sweet, S. K., and Guay, K. C.: Greater shrub dominance alters breeding habitat and food resources for migratory songbirds in Alaskan arctic tundra, *Global Change Biology*, 21, 1508–1520, <https://doi.org/10.1111/gcb.12761>, 2015.

- 755 Boertmann, D., Olsen, K., and Nielsen, R. D.: Geese in Northeast and North Greenland as recorded on aerial surveys in 2008 and 2009, *Dansk Ornitologisk Forenings Tidsskrift*, 109, 206–17, 2015.
- Bosson, J.-B., Huss, M., Cauvy-Fraunié, S., Clément, J.-C., Costes, G., Fischer, M., Poulenard, J., and Arthaud, F.: Future emergence of new ecosystems caused by glacial retreat, *Nature*, 620, 562–569, <https://doi.org/10.1038/s41586-023-06302-2>, 2023.
- Chen, Y., Cheng, X., Liu, A., Chen, Q., and Wang, C.: Tracking lake drainage events and drained lake basin vegetation dynamics across the
760 Arctic, *Nature Communications*, 14, 7359, <https://doi.org/10.1038/s41467-023-43207-0>, 2023.
- Colbeck, S. C.: An analysis of water flow in dry snow, *Water Resources Research*, 12, 523–527, 1976.
- Cooper, E. J.: Warmer shorter winters disrupt Arctic terrestrial ecosystems, *Annual Review of Ecology, Evolution, and Systematics*, 45, 271–295, <https://doi.org/10.1146/annurev-ecolsys-120213-091620>, 2014.
- Dahl, M. B., Priemé, A., Brejnrod, A., Brusvang, P., Lund, M., Nymand, J., Kramshøj, M., Ro-Poulsen, H., and Haugwitz, M. S.: Warming,
765 shading and a moth outbreak reduce tundra carbon sink strength dramatically by changing plant cover and soil microbial activity, *Scientific Reports*, 7, 16035, <https://doi.org/10.1038/s41598-017-16007-y>, 2017.
- Dingman, S. L.: *Physical Hydrology*, Waveland press, 2015.
- Eikelenboom, M., Higgins, R. C., John, C., Kerby, J., Forchhammer, M. C., and Post, E.: Contrasting dynamical responses of sympatric caribou and muskoxen to winter weather and earlier spring green-up in the Arctic, *Food Webs*, 27, e00196,
770 <https://doi.org/10.1016/j.fooweb.2021.e00196>, 2021.
- Elmendorf, S. C., Henry, G. H., Hollister, R. D., Björk, R. G., Boulanger-Lapointe, N., Cooper, E. J., Cornelissen, J. H., Day, T. A., Dorrepaal, E., Elumeeva, T. G., Gill, M., Gould, W. A., Harte, J., Hik, D. S., Hofgaard, A., Johnson, D. R., Johnstone, J. F., Jónsdóttir, I. S., Jorgenson, J. C., Klanderud, K., Klein, J. A., Koh, S., Kudo, G., Lara, M., Lévesque, E., Magnússon, B., May, J. L., Mercado-Díaz, J. A., Michelsen, A., Molau, U., Myers-Smith, I. H., Oberbauer, S. F., Onipchenko, V. G., Rixen, C., Schmidt, N. M., Shaver, G. R., Spasojevic, M. J.,
775 þórhallsdóttir, t. E., Tolvanen, A., Troxler, T., Tweedie, C. E., Villareal, S., Wahren, C.-H., Walker, X., Webber, P. J., Welker, J. M., and Wipf, S.: Plot-scale evidence of tundra vegetation change and links to recent summer warming, *Nature Climate Change*, 2, 453–457, <https://doi.org/10.1038/nclimate1465>, 2012.
- Ettema, J., Van den Broeke, M., Van Meijgaard, E., and Van de Berg, W.: Climate of the Greenland ice sheet using a high-resolution climate model—Part 2: Near-surface climate and energy balance, *The Cryosphere*, 4, 529–544, <https://doi.org/10.5194/tc-4-529-2010>, 2010.
- 780 Eythorsson, D., Gardarsson, S. M., Ahmad, S. K., Hossain, F., and Nijssen, B.: Arctic climate and snow cover trends—Comparing Global Circulation Models with remote sensing observations, *International Journal of Applied Earth Observation and Geoinformation*, 80, 71–81, <https://doi.org/10.1016/j.jag.2019.04.003>, 2019.
- Feng, S., Cook, J. M., Naegeli, K., Anesio, A. M., Benning, L. G., and Tranter, M.: The Impact of Bare Ice Duration and Geo-Topographical Factors on the Darkening of the Greenland Ice Sheet, *Geophysical Research Letters*, 51, e2023GL104894,
785 <https://doi.org/10.1029/2023GL104894>, 2024.
- Fettweis, X., Box, J. E., Agosta, C., Amory, C., Kittel, C., Lang, C., van As, D., Machguth, H., and Gallée, H.: Reconstructions of the 1900–2015 Greenland ice sheet surface mass balance using the regional climate MAR model, *The Cryosphere*, 11, 1015–1033, <https://doi.org/10.5194/tc-11-1015-2017>, 2017.
- Franch, B., Vermote, E. F., Roger, J.-C., Murphy, E., Becker-Reshef, I., Justice, C., Claverie, M., Nagol, J., Csizsar, I., Meyer, D., Baret, F.,
790 Masuoka, E., Wolfe, R., and Devadiga, S.: A 30+ year AVHRR land surface reflectance climate data record and its application to wheat yield monitoring, *Remote Sensing*, 9, 296, <https://doi.org/10.3390/rs9030296>, 2017.

- Gabriel, K. R.: The biplot graphic display of matrices with application to principal component analysis, *Biometrika*, 58, 453–467, <https://doi.org/10.1093/biomet/58.3.453>, 1971.
- Gamm, C. M., Sullivan, P. F., Buchwal, A., Dial, R. J., Young, A. B., Watts, D. A., Cahoon, S. M., Welker, J. M., and Post, E.: De-
795 clining growth of deciduous shrubs in the warming climate of continental western Greenland, *Journal of Ecology*, 106, 640–654, <https://doi.org/10.1111/1365-2745.12882>, 2018.
- Gandhi, G. M., Parthiban, S., Thummalu, N., and Christy, A.: Ndvi: Vegetation change detection using remote sensing and gis—A case study of Vellore District, *Procedia computer science*, 57, 1199–1210, <https://doi.org/10.1016/j.procs.2015.07.415>, 2015.
- Gehrmann, F., Lehtimäki, I.-M., Hänninen, H., and Saarinen, T.: Sub-Arctic alpine *Vaccinium vitis-idaea* exhibits resistance to strong
800 variation in snowmelt timing and frost exposure, suggesting high resilience under climatic change, *Polar Biology*, 43, 1453–1467, <https://doi.org/10.1007/s00300-020-02721-3>, 2020.
- Gilson, G. F., Jiskoot, H., Gueye, S., and van Boxel, J. H.: A climatology of Arctic fog along the coast of East Greenland, *Quarterly Journal of the Royal Meteorological Society*, 150, 706–726, 2024.
- Glanville, H. C., Hill, P. W., Maccarone, L. D., N. Golyshin, P., Murphy, D. V., and Jones, D. L.: Temperature and water controls on
805 vegetation emergence, microbial dynamics, and soil carbon and nitrogen fluxes in a high Arctic tundra ecosystem, *Functional Ecology*, 26, 1366–1380, <https://doi.org/10.1111/j.1365-2435.2012.02056.x>, 2012.
- Grimes, M., Carrivick, J. L., Smith, M. W., and Comber, A. J.: Land cover changes across Greenland dominated by a doubling of vegetation in three decades, *Scientific Reports*, 14, 3120, <https://doi.org/10.1038/s41598-024-52124-1>, 2024.
- Grossiord, C., Buckley, T. N., Cernusak, L. A., Novick, K. A., Poulter, B., Siegwolf, R. T., Sperry, J. S., and McDowell, N. G.: Plant responses
810 to rising vapor pressure deficit, *New phytologist*, 226, 1550–1566, <https://doi.org/10.1111/nph.16485>, 2020.
- Hallinger, M., Manthey, M., and Wilmking, M.: Establishing a missing link: warm summers and winter snow cover promote shrub expansion into alpine tundra in Scandinavia, *New Phytologist*, 186, 890–899, <https://doi.org/10.1111/j.1469-8137.2010.03223.x>, 2010.
- Hamed, K. H. and Rao, A. R.: A modified Mann-Kendall trend test for autocorrelated data, *Journal of hydrology*, 204, 182–196, 1998.
- Hanna, E., Jones, J. M., Cappelen, J., Mernild, S. H., Wood, L., Steffen, K., and Huybrechts, P.: The influence of North Atlantic atmospheric
815 and oceanic forcing effects on 1900–2010 Greenland summer climate and ice melt/runoff, *International Journal of Climatology*, 33, 862–880, <https://doi.org/10.1002/joc.3475>, 2013.
- Hanna, E., Cropper, T. E., Jones, P. D., Scaife, A. A., and Allan, R.: Recent seasonal asymmetric changes in the NAO (a marked summer decline and increased winter variability) and associated changes in the AO and Greenland Blocking Index, *International Journal of Climatology*, 35, 2540–2554, <https://doi.org/10.1002/joc.4157>, 2015.
- 820 Hanna, E., Cropper, T. E., Hall, R. J., and Cappelen, J.: Greenland Blocking Index 1851–2015: a regional climate change signal, *International Journal of Climatology*, 36, 4847–4861, <https://doi.org/10.1002/joc.4673>, 2016.
- Heijmans, M. M., Magnússon, R. Í., Lara, M. J., Frost, G. V., Myers-Smith, I. H., van Huissteden, J., Jorgenson, M. T., Fedorov, A. N., Epstein, H. E., Lawrence, D. M., and Limpens, J.: Tundra vegetation change and impacts on permafrost, *Nature Reviews Earth & Environment*, 3, 68–84, <https://doi.org/10.1038/s43017-021-00233-0>, 2022.
- 825 Hersbach, H., Bell, B., Berrisford, P., Hirahara, S., Horányi, A., Muñoz-Sabater, J., Nicolas, J., Peubey, C., Radu, R., Schepers, D., Simmons, A., Soci, C., Abdalla, S., Abellan, X., Balsamo, G., Bechtold, P., Biavati, G., Bidlot, J., Bonavita, M., De Chiara, G., Dahlgren, P., Dee, D., Diamantakis, M., Dragani, R., Flemming, J., Forbes, R., Fuentes, M., Geer, A., Haimberger, L., Healy, S., Hogan, R. J., Hólm, E., Janisková, M., Keeley, S., Laloyaux, P., Lopez, P., Lupu, C., Radnoti, G., de Rosnay, P., Rozum, I., Vamborg, F., Vil-

laume, S., and Thépaut, J.-N.: The ERA5 global reanalysis, *Quarterly Journal of the Royal Meteorological Society*, 146, 1999–2049, <https://doi.org/10.1002/qj.3803>, 2020.

Høye, T. T., Bowden, J. J., Hansen, O. L., Hansen, R. R., Henriksen, T. N., Niebuhr, A., and Skytte, M. G.: Elevation modulates how Arctic arthropod communities are structured along local environmental gradients, *Polar Biology*, 41, 1555–1565, <https://doi.org/10.1007/s00300-017-2204-2>, 2018.

Huai, B., van den Broeke, M. R., Reijmer, C. H., and Noël, B.: A daily 1-km resolution Greenland rainfall climatology (1958–2020) from statistical downscaling of a regional atmospheric climate model, *Journal of Geophysical Research: Atmospheres*, 127, e2022JD036688, <https://doi.org/10.1029/2022JD036688>, 2022.

Huang, M., Piao, S., Janssens, I. A., Zhu, Z., Wang, T., Wu, D., Ciais, P., Myneni, R. B., Peaucelle, M., Peng, S., Yang, H., and Peñuelas, J.: Velocity of change in vegetation productivity over northern high latitudes, *Nature Ecology & Evolution*, 1, 1649–1654, 2017.

Hurrell, J. W., Kushnir, Y., Ottersen, G., and Visbeck, M.: An overview of the North Atlantic oscillation, *Geophysical Monograph-American Geophysical Union*, 134, 1–36, <https://doi.org/10.1029/134GM01>, 2003.

Hussain, M. and Mahmud, I.: pyMannKendall: a python package for non-parametric Mann Kendall family of trend tests., *Journal of Open Source Software*, 4, 1556, <https://doi.org/10.21105/joss.01556>, 2019.

Jansen, E., Christensen, J. H., Dokken, T., Nisancioglu, K. H., Vinther, B. M., Capron, E., Guo, C., Jensen, M. F., Langen, P. L., Pedersen, R. A., Yang, S., Bentsen, M., Kjær, H. A., Sadatzki, H., Sessford, E., and Stendel, M.: Past perspectives on the present era of abrupt Arctic climate change, *Nature Climate Change*, 10, 714–721, 2020.

Jones, G. A. and Henry, G. H.: Primary plant succession on recently deglaciated terrain in the Canadian High Arctic, *Journal of Biogeography*, 30, 277–296, <https://doi.org/10.1046/j.1365-2699.2003.00818.x>, 2003.

Kalnay, E., Kanamitsu, M., Kistler, R., Collins, W., Deaven, D., Gandin, L., Iredell, M., Saha, S., White, G., Woollen, J., Zhu, Y., Chelliah, M., Ebisuzaki, W., Higgins, W., Janowiak, J., Mo, K. C., Ropelewski, C., Wang, J., Leetmaa, A., Reynolds, R., Jenne, R., and Joseph, D.: The NCEP/NCAR 40-year reanalysis project, *Bulletin of the American meteorological Society*, 77, 437–472, [https://doi.org/10.1175/1520-0477\(1996\)077<0437:TNYP>2.0.CO;2](https://doi.org/10.1175/1520-0477(1996)077<0437:TNYP>2.0.CO;2), 1996.

Karami, M., Hansen, B. U., Westergaard-Nielsen, A., Abermann, J., Lund, M., Schmidt, N. M., and Elberling, B.: Vegetation phenology gradients along the west and east coasts of Greenland from 2001 to 2015, *Ambio*, 46, 94–105, <https://doi.org/10.1007/s13280-016-0866-6>, 2017.

Karami, M., Westergaard-Nielsen, A., Normand, S., Treier, U. A., Elberling, B., and Hansen, B. U.: A phenology-based approach to the classification of Arctic tundra ecosystems in Greenland, *ISPRS Journal of Photogrammetry and Remote Sensing*, 146, 518–529, <https://doi.org/10.1016/j.isprsjprs.2018.11.005>, 2018.

Körner, C. and Alsos, I. G.: Freezing resistance in high arctic plant species of Svalbard in mid-summer, *Bauhinia*, 21, 1–8, 2008.

Laird, N. F., Crossett, C. C., Keaton, G. A., and Hopson, L. N.: Weather conditions and seasonal variability of limited surface visibility at Greenland coastal locations, *International Journal of Climatology*, 44, 393–405, <https://doi.org/10.1002/joc.8332>, 2024.

Lamichhane, J. R.: Rising risks of late-spring frosts in a changing climate, *Nature Climate Change*, 11, 554–555, <https://doi.org/10.1038/s41558-021-01090-x>, 2021.

Law, A., Nobajas, A., and Sangonzalo, R.: Heterogeneous changes in the surface area of lakes in the Kangerlussuaq area of southwestern Greenland between 1995 and 2017, *Arctic, Antarctic, and Alpine Research*, 50, S100027, <https://doi.org/10.1080/15230430.2018.1487744>, 2018.

- Le Moullec, M., Sandal, L., Grøtan, V., Buchwal, A., and Hansen, B. B.: Climate synchronises shrub growth across a high-arctic archipelago: contrasting implications of summer and winter warming, *Oikos*, 129, 1012–1027, <https://doi.org/10.1111/oik.07059>, 2020.
- Liljedahl, A. K., Boike, J., Daanen, R. P., Fedorov, A. N., Frost, G. V., Grosse, G., Hinzman, L. D., Iijma, Y., Jorgenson, J. C., Matveyeva, N., et al.: Pan-Arctic ice-wedge degradation in warming permafrost and its influence on tundra hydrology, *Nature Geoscience*, 9, 312–318, <https://doi.org/10.1038/ngeo2674>, 2016.
- Liu, Y., Wang, P., Elberling, B., and Westergaard-Nielsen, A.: Drivers of contemporary and future changes in Arctic seasonal transition dates for a tundra site in coastal Greenland, *Global Change Biology*, 30, e17118, <https://doi.org/10.1111/gcb.17118>, 2024.
- Loranty, M. M., Goetz, S. J., and Beck, P. S.: Tundra vegetation effects on pan-Arctic albedo, *Environmental Research Letters*, 6, 024014, <https://doi.org/10.1088/1748-9326/6/2/024014>, 2011.
- 875 Lorenz, E. N.: Empirical orthogonal functions and statistical weather prediction, vol. 1, Massachusetts Institute of Technology, Department of Meteorology Cambridge, 1956.
- Masson, V., Champeaux, J.-L., Chauvin, F., Meriguet, C., and Lacaze, R.: A global database of land surface parameters at 1-km resolution in meteorological and climate models, *Journal of Climate*, 16, 1261–1282, [https://doi.org/10.1175/1520-0442\(2003\)16<1261:AGDOLS>2.0.CO;2](https://doi.org/10.1175/1520-0442(2003)16<1261:AGDOLS>2.0.CO;2), 2003.
- 880 Masson, V., Le Moigne, P., Martin, E., Faroux, S., Alias, A., Alkama, R., Belamari, S., Barbu, A., Boone, A., Bouysse, F., et al.: The SURFEXv7. 2 land and ocean surface platform for coupled or offline simulation of earth surface variables and fluxes, *Geoscientific Model Development*, 6, 929–960, <https://doi.org/10.5194/gmd-6-929-2013>, 2013.
- Mekonnen, Z. A., Riley, W. J., Berner, L. T., Bouskill, N. J., Torn, M. S., Iwahana, G., Breen, A. L., Myers-Smith, I. H., Criado, M. G., Liu, Y., et al.: Arctic tundra shrubification: a review of mechanisms and impacts on ecosystem carbon balance, *Environmental Research*
- 885 *Letters*, 16, 053001, <https://doi.org/10.1088/1748-9326/abf28b>, 2021.
- Metcalfe, D. B., Hermans, T. D., Ahlstrand, J., Becker, M., Berggren, M., Björk, R. G., Björkman, M. P., Blok, D., Chaudhary, N., Chisholm, C., Classen, A. T., Hasselquist, N. J., Jonsson, M., Kristensen, J. A., Kumordzi, B. B., Lee, H., Mayor, J. R., Prevéy, J., Pantazatou, K., Rousk, J., Sponseller, R. A., Sundqvist, M. K., Tang, J., Uddling, J., Wallin, G., Zhang, W., Ahlström, A., Tenenbaum, D. E., and Abdi, A. M.: Patchy field sampling biases understanding of climate change impacts across the Arctic, *Nature Ecology & Evolution*, 2,
- 890 1443–1448, <https://doi.org/10.1038/s41559-018-0612-5>, 2018.
- Migała, K., Wojtuń, B., Szymański, W., and Muskała, P.: Soil moisture and temperature variation under different types of tundra vegetation during the growing season: A case study from the Fuglebekken catchment, SW Spitsbergen, *Catena*, 116, 10–18, <https://doi.org/10.1016/j.catena.2013.12.007>, 2014.
- Mills, R. T., Kumar, J., Hoffman, F. M., Hargrove, W. W., Spruce, J. P., and Norman, S. P.: Identification and visualization of dominant patterns and anomalies in remotely sensed vegetation phenology using a parallel tool for principal components analysis, *Procedia Computer Science*, 18, 2396–2405, <https://doi.org/10.1016/j.procs.2013.05.411>, 2013.
- 895 Musselman, K. N., Clark, M. P., Liu, C., Ikeda, K., and Rasmussen, R.: Slower snowmelt in a warmer world, *Nature Climate Change*, 7, 214–219, <https://doi.org/10.1038/nclimate3225>, 2017.
- Myers-Smith, I. H., Forbes, B. C., Wilkening, M., Hallinger, M., Lantz, T., Blok, D., Tape, K. D., Macias-Fauria, M., Sass-Klaassen, U., Lévesque, E., Boudreau, S., Ropars, P., Hermanutz, L., Trant, A., Collier, L. S., Weijers, S., Rozema, J., Rayback, S. A., Schmidt, N. M., Schaeppman-Strub, G., Wipf, S., Rixen, C., Ménard, C. B., Venn, S., Goetz, S., Andreu-Hayles, L., Elmendorf, S., Ravolainen, V., Welker, V., Grogan, P., Epstein, H. E., and Hik, D. S.: Shrub expansion in tundra ecosystems: dynamics, impacts and research priorities, *Environmental Research Letters*, 6, 045509, <https://doi.org/10.1088/1748-9326/6/4/045509>, 2011.
- 900

- Myers-Smith, I. H., Kerby, J. T., Phoenix, G. K., Bjerke, J. W., Epstein, H. E., Assmann, J. J., John, C., Andreu-Hayles, L., Angers-Blondin, S., Beck, P. S., Berner, L. T., Bhatt, U. S., Bjorkman, A. D., Blok, D., Bryn, A., Christiansen, C. T., Cornelissen, J. H. C., Cunliffe, A. M., Elmendorf, S. C., Forbes, B. C., Goetz, S. J., Hollister, R. D., de Jong, R., Loranty, M. M., Macias-Fauria, M., Maseyk, K., Normand, S., Olofsson, J., Parker, T. C., Parmentier, F.-J. W., Post, E., Schaepman-Strub, G., Stordal, F., Sullivan, P. F., Thomas, H. J. D., Tømmervik, H., Treharne, R., Tweedie, C. E., Walker, D. A., Wilmking, M., and Wipf, S.: Complexity revealed in the greening of the Arctic, *Nature Climate Change*, 10, 106–117, <https://doi.org/10.1038/s41558-019-0688-1>, 2020.
- Nachtergaele, F., van Velthuisen, H., Verelst, L., Batjes, N., Dijkshoorn, K., van Engelen, V., Fischer, G., Jones, A., and Montanarella, L.: The harmonized world soil database, in: *Proceedings of the 19th World Congress of Soil Science, Soil Solutions for a Changing World*, Brisbane, Australia, 1-6 August 2010, pp. 34–37, 2010.
- Niwano, M., Box, J., Wehrlé, A., Vandecrux, B., Colgan, W., and Cappelen, J.: Rainfall on the Greenland ice sheet: Present-day climatology from a high-resolution non-hydrostatic polar regional climate model, *Geophysical Research Letters*, 48, e2021GL092942, <https://doi.org/10.1029/2021GL092942>, 2021.
- Oehri, J., Schaepman-Strub, G., Kim, J.-S., Grysko, R., Kropp, H., Grünberg, I., Zemlianskii, V., Sonnentag, O., Euskirchen, E. S., Reji Chacko, M., Muscari, G., Blanken, P. D., Dean, J. F., di Sarra, A., Harding, R. J., Sobota, I., Kutzbach, L., Plekhanova, E., Riihimäki, A., Boike, J., Miller, N. B., Beringer, J., López-Blanco, E., Stoy, P. C., Sullivan, R. C., Kejna, M., Parmentier, F.-J. W., Gamon, J. A., Mastepanov, M., Wille, C., Jackowicz-Korczynski, M., Karger, D. N., Quinton, W. L., Putkonen, J., van As, D., Christensen, T. R., Hakuba, M. Z., Stone, R. S., Metzger, S., Vandecrux, B., Frost, G. V., Wild, M., Hansen, B., Meloni, D., Domine, F., te Beest, M., Sachs, T., Kalhori, A., Rocha, A. V., Williamson, S. N., Morris, S., Atchley, A. L., Essery, R., Runkle, B. R. K., Holl, D., Riihimäki, L. D., Iwata, H., Schuur, E. A. G., Cox, C. J., Grachev, A. A., McFadden, J. P., Fausto, R. S., Göckede, M., Ueyama, M., Pirk, N., de Boer, G., Bret-Harte, M. S., Leppäranta, M., Steffen, K., Friberg, T., Ohmura, A., Edgar, C. W., Olofsson, J., and Chambers, S. D.: Vegetation type is an important predictor of the arctic summer land surface energy budget, *Nature Communications*, 13, 6379, <https://doi.org/10.1038/s41467-022-34049-3>, 2022.
- Olafsson, H. and Rousta, I.: Influence of atmospheric patterns and North Atlantic Oscillation (NAO) on vegetation dynamics in Iceland using Remote Sensing, *European Journal of Remote Sensing*, 54, 351–363, <https://doi.org/10.1080/22797254.2021.1931462>, 2021.
- Opala-Owczarek, M., Pirożnikow, E., Owczarek, P., Szymański, W., Luks, B., Kępski, D., Szymanowski, M., Wojtuń, B., and Mięka, K.: The influence of abiotic factors on the growth of two vascular plant species (*Saxifraga oppositifolia* and *Salix polaris*) in the High Arctic, *Catena*, 163, 219–232, <https://doi.org/10.1016/j.catena.2017.12.018>, 2018.
- Pearson, K.: LIII. On lines and planes of closest fit to systems of points in space, *The London, Edinburgh, and Dublin philosophical magazine and journal of science*, 2, 559–572, 1901.
- Pearson, R. G., Phillips, S. J., Loranty, M. M., Beck, P. S., Damoulas, T., Knight, S. J., and Goetz, S. J.: Shifts in Arctic vegetation and associated feedbacks under climate change, *Nature Climate Change*, 3, 673–677, <https://doi.org/10.1038/nclimate1858>, 2013.
- Pedregosa, F., Varoquaux, G., Gramfort, A., Michel, V., Thirion, B., Grisel, O., Blondel, M., Prettenhofer, P., Weiss, R., Dubourg, V., Vanderplas, J., Passos, A., Cournapeau, D., Brucher, M., Perrot, M., and Duchesnay, E.: Scikit-learn: Machine Learning in Python, *Journal of Machine Learning Research*, 12, 2825–2830, <http://jmlr.org/papers/v12/pedregosa11a.html>, 2011.
- Pedron, S., Jespersen, R., Xu, X., Khazindar, Y., Welker, J., and Czimczik, C.: More snow accelerates legacy carbon emissions from Arctic permafrost, *AGU Advances*, 4, e2023AV000942, <https://doi.org/10.1029/2023AV000942>, 2023.

- 940 Pereira Freitas, G., Adachi, K., Conen, F., Heslin-Rees, D., Krejci, R., Tobo, Y., Yttri, K. E., and Zieger, P.: Regionally sourced bioaerosols drive high-temperature ice nucleating particles in the Arctic, *Nature Communications*, 14, 5997, <https://doi.org/10.1038/s41467-023-41696-7>, 2023.
- Post, E. and Pedersen, C.: Opposing plant community responses to warming with and without herbivores, *Proceedings of the National Academy of Sciences*, 105, 12 353–12 358, <https://doi.org/10.1073/pnas.0802421105>, 2008.
- 945 Power, C. C., Normand, S., von Arx, G., Elberling, B., Corcoran, D., Krog, A. B., Bouvin, N. K., Treier, U. A., Westergaard-Nielsen, A., Liu, Y., and L. Prendin, A.: No effect of snow on shrub xylem traits: Insights from a snow-manipulation experiment on Disko Island, Greenland, *Science of The Total Environment*, 916, 169 896, <https://doi.org/10.1016/j.scitotenv.2024.169896>, 2024.
- Rantanen, M., Karpechko, A. Y., Lipponen, A., Nordling, K., Hyvärinen, O., Ruosteenoja, K., Vihma, T., and Laaksonen, A.: The Arctic has warmed nearly four times faster than the globe since 1979, *Communications Earth & Environment*, 3, 168, 2022.
- 950 Rantanen, M., Kämäräinen, M., Niittynen, P., Phoenix, G. K., Lenoir, J., Maclean, I., Luoto, M., and Aalto, J.: Bioclimatic atlas of the terrestrial Arctic, *Scientific Data*, 10, 40, <https://doi.org/10.1038/s41597-023-01959-w>, 2023.
- Rawlins, M. A. and Karmalkar, A. V.: Regime shifts in Arctic terrestrial hydrology manifested from impacts of climate warming, *The Cryosphere*, 18, 1033–1052, <https://doi.org/10.5194/tc-18-1033-2024>, 2024.
- Rudd, D. A., Karami, M., and Fensholt, R.: Towards high-resolution land-cover classification of Greenland: A case study covering Kobbefjord, Disko and Zackenberg, *Remote Sensing*, 13, 3559, <https://doi.org/10.3390/rs13183559>, 2021.
- 955 Salmon, V. G., Soucy, P., Mauritz, M., Celis, G., Natali, S. M., Mack, M. C., and Schuur, E. A.: Nitrogen availability increases in a tundra ecosystem during five years of experimental permafrost thaw, *Global Change Biology*, 22, 1927–1941, <https://doi.org/10.1111/gcb.13204>, 2016.
- Schmidt, N. M., Pedersen, S. H., Mosbacher, J. B., and Hansen, L. H.: Long-term patterns of muskox (*Ovibos moschatus*) demographics in high arctic Greenland, *Polar Biology*, 38, 1667–1675, <https://doi.org/10.1007/s00300-015-1733-9>, 2015.
- 960 Schmidt, N. M., Reneerkens, J., Christensen, J. H., Olesen, M., and Roslin, T.: An ecosystem-wide reproductive failure with more snow in the Arctic, *PLoS Biology*, 17, e3000 392, <https://doi.org/10.1371/journal.pbio.3000392>, 2019.
- Schmidt, N. M., Kankaanpää, T., Tiisanen, M., Reneerkens, J., Versluijs, T. S., Hansen, L. H., Hansen, J., Gerlich, H. S., Høye, T. T., Cirtwill, A. R., Zhemchuzhnikov, M. K., Peña-Aguilera, P., and Roslin, T.: Little directional change in the timing of Arctic spring phenology over the past 25 years, *Current Biology*, 33, 3244–3249, <https://doi.org/10.1016/j.cub.2023.06.038>, 2023.
- 965 Schyberg, H., Yang, X., Køltzow, M., Amstrup, B., Bakketun, m., Bazile, E., Bojarova, J., Box, J. E., Dahlgren, P., Hagelin, S., Homleid, M., Horányi, A., Høyer, J., Johansson, m., Killie, M., Körnich, H., Le Moigne, P., Lindskog, M., Manninen, T., Nielsen, E. P., Nielsen, K., Olsson, E., Palmason, B., Peralta, A. C., Randriamampianina, R., Samuelsson, P., Stappers, R., Støylen, E., Thorsteinsson, S., Valkonen, T., and Wang, Z.: Arctic regional reanalysis on single levels from 1991 to present. Copernicus Climate Change Service (C3S) Climate Data Store (CDS), <https://doi.org/10.24381/cds.713858f6>, accessed on 15-12-2022, 2020.
- 970 Shahi, S., Abermann, J., Silva, T., Langley, K., Larsen, S. H., Mastepanov, M., and Schöner, W.: The importance of regional sea-ice variability for the coastal climate and near-surface temperature gradients in Northeast Greenland, *Weather and Climate Dynamics*, 4, 747–771, <https://doi.org/10.5194/wcd-4-747-2023>, 2023.
- Silva, T., Abermann, J., Noël, B., Shahi, S., van de Berg, W. J., and Schöner, W.: The impact of climate oscillations on the surface energy budget over the Greenland Ice Sheet in a changing climate, *The Cryosphere*, 16, 3375–3391, <https://doi.org/10.5194/tc-16-3375-2022>, 2022.

- Skakun, S., Justice, C. O., Vermote, E., and Roger, J.-C.: Transitioning from MODIS to VIIRS: an analysis of inter-consistency of NDVI data sets for agricultural monitoring, *International Journal of Remote Sensing*, 39, 971–992, <https://doi.org/10.1080/01431161.2017.1395970>, 2018.
- 980 Song, S., Chen, Y., Chen, X., Chen, C., Li, K.-F., Tung, K.-K., Shao, Q., Liu, Y., Wang, X., Yi, L., and Zhao, J.: Adapting to a Foggy Future Along Trans-Arctic Shipping Routes, *Geophysical Research Letters*, 50, e2022GL102395, <https://doi.org/10.1029/2022GL102395>, 2023.
- Stengel, M., Stapelberg, S., Sus, O., Finkensieper, S., Würzler, B., Philipp, D., Hollmann, R., Poulsen, C., Christensen, M., and McGarragh, G.: Cloud_cci Advanced Very High Resolution Radiometer post meridiem (AVHRR-PM) dataset version 3: 35-year climatology of global cloud and radiation properties, *Earth System Science Data*, 12, 41–60, <https://doi.org/10.5194/essd-12-41-2020>, 2020.
- 985 Stephenson, G. R. and Freeze, R. A.: Mathematical simulation of subsurface flow contributions to snowmelt runoff, Reynolds Creek Watershed, Idaho, *Water Resources Research*, 10, 284–294, 1974.
- Sturm, M., Racine, C., and Tape, K.: Increasing shrub abundance in the Arctic, *Nature*, 411, 546–547, <https://doi.org/10.1038/35079180>, 2001.
- Sze, K. C. H., Wex, H., Hartmann, M., Skov, H., Massling, A., Villanueva, D., and Stratmann, F.: Ice Nucleating Particles in Northern Greenland: annual cycles, biological contribution and parameterizations, *Atmospheric Chemistry and Physics Discussions*, 23, 1–45, <https://doi.org/10.5194/acp-23-4741-2023>, 2022.
- van der Kolk, H.-J., Heijmans, M. M., Van Huissteden, J., Pullens, J. W., and Berendse, F.: Potential Arctic tundra vegetation shifts in response to changing temperature, precipitation and permafrost thaw, *Biogeosciences*, 13, 6229–6245, <https://doi.org/10.5194/bg-13-6229-2016>, 2016.
- 995 van der Schot, J., Abermann, J., Silva, T., Jensen, C. D., Noël, B., and Schöner, W.: Precipitation trends (1958–2021) on Ammassalik island, south-east Greenland, *Frontiers in Earth Science*, 10, 1085499, <https://doi.org/10.3389/feart.2022.1085499>, 2023.
- van der Schot, J., Abermann, J., Silva, T., Rasmussen, K., Winkler, M., Langley, K., and Schöner, W.: Seasonal snow cover indicators in coastal Greenland from in situ observations, a climate model, and reanalysis, *The Cryosphere*, 18, 1033–1052, <https://doi.org/10.5194/tc-18-5803-2024>, 2024.
- 1000 Vermote, E., Justice, C., Csaszar, I., Eidenshink, J., Myneni, R., Baret, F., Masuoka, E., Wolfe, R., Claverie, M., and Program, N. C.: NOAA Climate Data Record (CDR) of Normalized Difference Vegetation Index (NDVI), Version 5, <https://doi.org/10.7289/V5ZG6QH9>, access date: 2022-05-06, 2018.
- Vermote, E., Franch, B., Roger, J.-C., Murphy, E., Becker-Reshef, I., Justice, C., Claverie, M., Nagol, J., Csaszar, I., Meyer, D., Baret, F., Masuoka, E., Wolfe, R., Devadiga, S., Villaescusa, J., and Program, N. C.: NOAA Climate Data Record (CDR) of Surface Reflectance, Version 1, <https://doi.org/10.25921/gakh-st76>, access date: 2023-07-06, 2022.
- 1005 Walker, D. A., Raynolds, M. K., Daniëls, F. J., Einarsson, E., Elvebakk, A., Gould, W. A., Katenin, A. E., Kholod, S. S., Markon, C. J., Melnikov, E. S., Moskalenko, N. G., Talbot, S. S., Yurtsev, B. A., and other members of the CAVM Team, T.: The circumpolar Arctic vegetation map, *Journal of Vegetation Science*, 16, 267–282, <https://doi.org/10.1111/j.1654-1103.2005.tb02365.x>, 2005.
- Wang, X., Li, Z., Xiao, J., Zhu, G., Tan, J., Zhang, Y., Ge, Y., and Che, T.: Snow cover duration delays spring green-up in the northern hemisphere the most for grasslands, *Agricultural and Forest Meteorology*, 355, 110130, <https://doi.org/10.1016/j.agrformet.2024.110130>, 2024.
- 1010 Weijers, S.: Declining temperature and increasing moisture sensitivity of shrub growth in the Low-Arctic erect dwarf-shrub tundra of western Greenland, *Ecology and Evolution*, 12, e9419, <https://doi.org/10.1002/ece3.9419>, 2022.

- Weijers, S., Buchwal, A., Blok, D., Löffler, J., and Elberling, B.: High Arctic summer warming tracked by increased *Cassiope tetragona* growth in the world's northernmost polar desert, *Global Change Biology*, 23, 5006–5020, <https://doi.org/10.1111/gcb.13747>, 2017.
- 1015 Westergaard-Nielsen, A., Karami, M., Hansen, B. U., Westermann, S., and Elberling, B.: Contrasting temperature trends across the ice-free part of Greenland, *Scientific Reports*, 8, 1586, <https://doi.org/10.1038/s41598-018-19992-w>, 2018.
- Westergaard-Nielsen, A., Hansen, B. U., Elberling, B., and Abermann, J.: Greenland Climates, *Encyclopedia of the World's Biomes*, 1, 465–479, <https://doi.org/10.1016/B978-0-12-409548-9.11750-6>, 2020.
- 1020 Xu, W., Prieme, A., Cooper, E. J., Mörsdorf, M. A., Semenchuk, P., Elberling, B., Grogan, P., and Ambus, P. L.: Deepened snow enhances gross nitrogen cycling among Pan-Arctic tundra soils during both winter and summer, *Soil Biology and Biochemistry*, 160, 108356, <https://doi.org/10.1016/j.soilbio.2021.108356>, 2021.
- Yan, W. and Tinker, N. A.: Biplot analysis of multi-environment trial data: Principles and applications, *Canadian journal of plant science*, 86, 623–645, <https://doi.org/10.4141/P05-169>, 2006.
- 1025 Yang, W., Tan, B., Huang, D., Rautiainen, M., Shabanov, N. V., Wang, Y., Privette, J. L., Huemmrich, K. F., Fensholt, R., Sandholt, I., Weiss, M., Ahl, D., Gower, S., Nemani, R., Knyazikhin, Y., and Myneni, R.: MODIS leaf area index products: From validation to algorithm improvement, *IEEE Transactions on Geoscience and Remote Sensing*, 44, 1885–1898, <https://doi.org/10.1109/TGRS.2006.871215>, 2006.
- Yuan, H., Dai, Y., Xiao, Z., Ji, D., and Shangguan, W.: Reprocessing the MODIS Leaf Area Index products for land surface and climate modelling, *Remote Sensing of Environment*, 115, 1171–1187, <https://doi.org/10.1016/j.rse.2011.01.001>, 2011.
- 1030 Yuan, W., Zheng, Y., Piao, S., Ciais, P., Lombardozzi, D., Wang, Y., Ryu, Y., Chen, G., Dong, W., Hu, Z., Jain, A. K., Jiang, C., Kato, E., Li, S., Lienert, S., Liu, S., Nabel, J. E., Qin, Z., Quine, T., Sitch, S., Smith, W. K., Wang, F., Wu, C., Xiao, Z., and Yang, S.: Increased atmospheric vapor pressure deficit reduces global vegetation growth, *Science advances*, 5, eaax1396, <https://doi.org/10.1126/sciadv.aax1396>, 2019.
- Zwolicki, A., Zmudczyńska-Skarbek, K., Wietrzyk-Pelka, P., and Convey, P.: High Arctic vegetation, *Encyclopedia of the World's Biomes*, 1, 465–479, <https://doi.org/10.1016/B978-0-12-409548-9.11771-3>, 2020.