



## Research article

# UV-screening and springtime recovery of photosynthetic capacity in leaves of *Vaccinium vitis-idaea* above and below the snow pack<sup>☆</sup>



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## ARTICLE INFO

## Keywords:

spring dehardening  
Flavonoids  
Anthocyanins  
Understorey  
Spectral quality  
Photoprotection  
Arctic browning

## ABSTRACT

Evergreen plants in boreal biomes undergo seasonal hardening and dehardening adjusting their photosynthetic capacity and photoprotection; acclimating to seasonal changes in temperature and irradiance. Leaf epidermal ultraviolet (UV)-screening by flavonols responds to solar radiation, perceived in part through increased ultraviolet-B (UV-B) radiation, and is a candidate trait to provide cross-photoprotection. At Hyytiälä Forestry Station, central Finland, we examined whether the accumulation of flavonols was higher in leaves of *Vaccinium vitis-idaea* L. growing above the snowpack compared with those below the snowpack. We found that leaves exposed to colder temperatures and higher solar radiation towards the top of hummocks suffered greater photoinhibition than those at the base of hummocks. Epidermal UV-screening was highest in upper-hummock leaves, particularly during winter when lower leaves were beneath the snowpack. There was also a negative relationship between indices of flavonols and anthocyanins across all leaves suggesting fine-tuning of flavonoid composition for screening vs. antioxidant activity in response to temperature and irradiance. However, the positive correlation between the maximum quantum yield of photosystem II photochemistry ( $F_v/F_m$ ) and flavonol accumulation in upper hummock leaves during dehardening did not confer on them any greater cross-protection than would be expected from the general relationship of  $F_v/F_m$  with temperature and irradiance (throughout the hummocks). Irrespective of timing of snow-melt, photosynthesis fully recovered in all leaves, suggesting that *V. vitis-idaea* has the potential to exploit the continuing trend for longer growing seasons in central Finland without incurring significant impairment from reduced duration of snow cover.

## 1. Introduction

An expected outcome of ongoing human-induced climate change is a shift in seasonal precipitation and temperature affecting growing season length (Harsch and HilleRisLambers, 2016; IPCC, 2014). This has contributed to a reduction in annual snow-cover in the Arctic by as much as 5 days each decade since 1979 (Derksen et al., 2016). At high northern latitudes, less of the winter precipitation is expected to fall as snow and more as rain (Mellander et al., 2007; Rasmus et al., 2004) and temperatures favourable for growth are forecast to occur earlier in the spring (Henry, 2008; Jylhä et al., 2008). In Finland, mean annual

temperature is expected to increase by 4–11 °C and precipitation (and hence cloudiness) by 10–60% by the end of 21st century, with most increases occurring during the winter (Finland's Seventh National Communication under the United Nations Framework Convention on Climate Change, 2017). The benefit of this change for plant growth and photosynthesis is uncertain because warmer mean temperatures together with reduced precipitation as snow may lead a smaller snow pack that melts earlier (Kreyling et al., 2012). The absence of snow cover will imply greater temperature fluctuations close to the ground, exposing plants to colder minimum, and warmer maximum, temperatures when they are starting to dehardening at the end of winter (Blume-

<sup>☆</sup> This article is part of the Special Issue “Interactive effects of UV-B radiation in a complex environment” published at the journal Plant Physiology and Biochemistry 134, 2019.

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<https://doi.org/10.1016/j.plaphy.2018.09.003>

Received 8 May 2018; Received in revised form 3 September 2018; Accepted 4 September 2018

Available online 08 September 2018

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Werry et al., 2016; Preece and Phoenix, 2013).

Plants possess several photoprotective mechanisms which mitigate abiotic stresses under harsh conditions: (1) when reaction centres are open photochemical quenching ( $q_P$ ) entails activation of the specific enzymes and metabolites involved in charge separation; (2) excess light energy is dissipated through xanthophyll-cycle pigments (carotenoids) zeaxanthin and antheraxanthin in a process regulated by the pH the thylakoid lumen (Adams et al., 2004; Murchie and Lawson, 2013); (3) additional antioxidants such as phenolic compounds are accumulated to quench reactive oxygen species (ROS) (Agati et al., 2013; Hideg et al., 2013; Salter et al., 2017). It is reasonable to assume that the pressure on these mechanisms is increased when a plant is subject to both fluctuating temperatures and irradiances above the snowpack rather than the more stable conditions offered under snow (Loik et al., 2004; Preece and Phoenix, 2013). Low temperatures dampen physiological processes meaning that even moderate irradiances cause photoinhibition through photoinactivation of the  $Mn_4CaO_5$  cluster or inhibition of the photosystem II (PSII) electron acceptors (Neuner et al., 1999; Saarenin et al., 2011). Sunlight can penetrate the snowpack, where temperature fluctuations are small, potentially allowing some photosynthesis at low irradiances (Saarenin et al., 2016). Some studies have found the relative transmittance of different regions of the spectrum by snow to be quite even (Gorton and Vogelmann, 2003) while others have reported spectrally-differential attenuation through the snowpack (Curl Jr et al., 1972; Richardson and Salisbury, 1977). Although solar irradiance, particularly ultraviolet-B (UV-B: 280–315 nm) radiation, is low in winter, the ratio of ultraviolet (UV-B plus UV-A: 280–400 nm) to photosynthetically active radiation (PAR: 400–700 nm) in forest understorey shade is enriched (Jansen et al., 2017; Hartikainen et al., 2018).

Flavonoids such as flavonols and anthocyanins are the most-common class of phenolic compounds produced in response to UV-radiation and blue light (Searles et al., 2001; Sullivan et al., 2007; Taulavuori et al., 2016). Flavonols accumulate in the leaf adaxial epidermis absorbing UV-radiation so screening the photosynthetic apparatus in the mesophyll below (Barnes et al., 2017; Caldwell et al., 1983; Nybakken et al., 2004). Although their absorption spectrum is predominately in the UV region, it has been proposed that flavonoids provide cross-protection to high solar irradiance in general (Jansen et al., 2017), and a positive correlation is sometimes reported between flavonoid accumulation and the yield of PSII (Robson et al., 2014; Wargent et al., 2011). This relationship may be attributable to protection against PSII photodamage or enhanced ROS scavenging conferred by flavonoids (reviewed by Takahashi and Badger, 2011). The accumulation of flavonoids in response to UV-radiation is moderated by temperature, being less pronounced in warm than in cool temperatures (Bilger et al., 2007; Coffey et al., 2017; Petridis et al., 2016) where flavonoids can be important in mitigating against oxidative stress (Lütz et al., 2005) and can increase frost hardiness (Chalker-Scott and Scott, 2004; Schulz et al., 2016). Hence, the accumulation of flavonoids could provide a feasible strategy whereby evergreen plants ameliorate photoinhibition, imposed by increasing irradiance and temperature fluctuations, as they dehardening over the spring.

Boreal evergreen and wintergreen plants adjust light harvesting complexes to optimise quenching processes at the expense of the light harvesting and photosynthetic capacity of their leaves during winter when light and temperature conditions are unfavourable (Adams et al., 2004). Subsequently, they undergo a recovery period in preparation for more favourable growth conditions in late spring and summer (Muller et al., 2005). During this period of dehardening, full functionality returns to the photosynthetic apparatus of leaves and this can be measured using chlorophyll fluorescence as an increase in the yield of PSII (Murchie and Lawson, 2013; Saarenin et al., 2011). Monitoring leaf-level chlorophyll fluorescence of evergreen trees (Ensminger et al., 2004; Porcar-Castell et al., 2008) and field-layer plants (Starr and Oberbauer, 2003) through spring in the boreal forest, has revealed that recovery of  $F_v/F_m$  is modulated by temperature and light.

Here, we aimed to compare leaf epidermal flavonoids (flavonols and anthocyanins), chlorophyll content and the PSII yield at different positions on hummocks relative to the snowpack, through a seasonal time-course of measurements of evergreen-shrub *Vaccinium vitis-idaea* L. During the spring dehardening period, the over-wintering leaves of this species are adapted to take advantage of favourable early-spring conditions by photosynthesizing when still under the snowpack (Lundell et al., 2008; Saarenin et al., 2011; Starr and Oberbauer, 2003). Our hypothesis was that leaves at the top of the hummock above the snowpack during winter and early spring would suffer greater photoinhibition because they encounter fluctuating temperatures and higher irradiance than those at the base of hummocks receiving lower irradiance, of both PAR and UV radiation, and more stable temperatures remaining closer to 0 °C under the snow. However, we also expected higher accumulation of epidermal UV-absorbing compounds (flavonols and anthocyanins) in leaves above the snow, and for this to partially ameliorate the photoinhibition they suffered relative to those at equivalent temperatures beneath the snowpack.

## 2. Methods

### 2.1. Study site and species

The study was performed under an open *Pinus sylvestris* L. canopy, basal area 23 m<sup>2</sup>ha<sup>-1</sup>, on level ground at Hyytiälä Forestry Station of the University of Helsinki in central Finland (61.84741°N 24.29477°E 181 m asl). The hummock-forming evergreen shrub *V. vitis-idaea* covers vast areas of peatland and boreal forest at high latitudes in the Northern Hemisphere (Karlsson, 1985; Lundell et al., 2008). At our site, the understorey comprises almost exclusively *V. vitis-idaea* in hummocks of up to 75 cm in height intermingled with *Vaccinium myrtillus* L. Four hummocks, positioned at least a metre from the nearest pine tree, were selected at random within a 10-by-10-m area of understorey. These hummocks were divided vertically into three height categories on 14-02-2017: the “upper” hummock being the uppermost shoots, most exposed to the wind and sun, the “middle” hummock being the outer shoots on the south side of the hummock, somewhat sheltered from the wind but just below the snow pack through the winter, and the “lower” hummock being shoots at the very base of the hummock on the south side, below the snow pack during the whole winter and early spring.

The snow depth relative to the lower, middle and upper hummocks, and the leaf phenology of the *Vaccinium* species, was monitored throughout the study concomitantly with leaf traits, and these data were compared against a continuous monitoring by Phenocam time-lapse images nearby in the stand (Kolari et al., 2017). By 19-06-2017, plants of both *V. vitis-idaea* and *V. myrtillus* produced new leaves. From this date onwards, until our final measurement date, 08-11-2017, a set of measurements was also made on these new *V. vitis-idaea* leaves (unfurling in June 2017) in same way as those on leaves produced in June 2016. This allowed us to obtain data for leaves from the time of their unfurling through their first and second years of growth. To allow this continuum of leaf age to be visualised in the figures we have combined these two set of leaf traits in a single time series; with 2017 leaves plotted from their unfurling until 08-11-2017 (day 312 of the year, leaf age 152 days) and 2016 leaves from 14-02-2017 onwards (day 45 of the year, leaf age 250 days).

### 2.2. Characterisation of the forest canopy by hemispherical photograph

Plant area index (PAI)(m<sup>2</sup>m<sup>-2</sup>), was estimated at each of the four hummocks by taking hemispherical photographs (at 40 cm height from the ground) through a Sigma 4.5 mm f2.8 EX DC HSM circular Fisheye lens (Sigma Corporation of America, Ronkonkoma, NY) on a Nikon D7000 camera (Nikon corporation, Tokyo, Japan) levelled on a tripod. Measurements were done on a completely overcast day in mid-July (20-07-2017) to ensure homogeneous grey sky and a good contrast between

the sky and the canopy (Fig S1). All photos were analysed following the procedure detailed in Hartikainen et al. (2018).

### 2.3. Measurement of spectral irradiance at the study site

The solar spectral irradiance was measured close to solar noon, directly to the south of each of the four hummocks on 07-03-2017 and 19-06-2017 in sunflecks and canopy shade on completely clear-sky days. These data were compared with corresponding measurements in an open field next to the forest stand earlier on the same days. Spectral irradiance recordings were made using a Maya 2000 Pro (Ocean Optics, Dunedin, FL, USA) array spectrometer. The cosine diffuser (Bentham Instruments Ltd. Reading UK) was levelled on the tripod 40-cm high and attached with a fibre-optic cable to the spectrometer. Measurements were corrected for dark-noise and stray light in the UV range ( $< 400$  nm) using a protocol detailed by Hartikainen et al. (2018). The spectrometer had been calibrated against a Bentham scanning spectroradiometer for outdoor measurements of solar spectral irradiance, over the wavelength range 290–900 nm. Pre-processing of the data was performed using functions from the R packages Ooacquire and Photobiology in R version 3.4.3 (R Core Team, 2018).

### 2.4. Optical measurements of *V. vitis-idaea* leaf pigments and photosynthetic capacity

On each measurement date, starting 14-02-2017, then 07-03-2017, and at 2-week intervals subsequently until 08-07-2017, with an additional final measurement date of 08-11-2017, leaf optical properties and photosynthetic capacity were measured. Ten leaves (remaining attached to the plants) were measured *in situ* from each height (upper, middle, lower) on four hummocks. Sun-exposed leaves of equivalent size were measured (or leaves which would have been sun exposed had it not been cloudy on overcast days), all from different shoots of each *V. vitis-idaea* hummock, avoiding the three distal leaves of a shoot. Lower and middle hummock leaves were, when necessary, excavated from beneath the snow to be measured and subsequently the snow was replaced over these leaves. Leaves that were damp from dew, rain or melting snow were quickly but gently dried with a tissue prior to measurement.

Dualex Scientific<sup>+</sup> (Force-A, University Paris-Sud, Orsy, France) was used to optically assess the epidermal flavonol concentration (epidermal absorption in the UV-A region at 375 nm) and anthocyanin concentration (absorption at 515 nm), from both the adaxial and abaxial sides of each leaf, as well as the leaf chlorophyll concentration (assessed optically by transmittance in the red at 650 nm vs. near-infrared at 710 nm, Cerovic et al., 2012). We recorded the maximum quantum yield of PSII photochemistry ( $F_v/F_m$ ; where  $F_v$  [variable fluorescence] is the maximal fluorescence,  $F_m$ , minus the minimal fluorescence,  $F_0$ , of dark-adapted leaves, as defined by Murchie and Lawson, 2013) measured with a mini-PAM (Heinz Walz GmbH, Effeltrich, Germany) following 30 min of dark-adaptation with darkening clips. According to Bokhorst et al. (2010), 30 min in darkness should be sufficient for full dark adaptation of *V. vitis-idaea* leaves year 'round. We also recorded the effective quantum yield of PSII photochemistry ( $\phi_{PSII}$  calculated as  $F_q'/F_m'$ ; where  $F_q'$  is the maximal fluorescence,  $F_m'$ , minus the steady-state fluorescence,  $F'$ , of leaves in ambient sunlight, as defined by Murchie and Lawson, 2013) of leaves in the sun and shade in the mid-to-late morning. Where possible, the same ten leaves were used for chlorophyll fluorescence and Dualex measurements on any given day, but it was not feasible to measure the same ten leaves from one measurement date to another, so from day to day leaves meeting the above criteria were selected at random.

### 2.5. Harvest of leaves for trait measurements

On 08-07-2017, leaves of age 29 days and 394 days were harvested.

From each age cohort 8–10 leaves were sampled from each of the four hummocks at each of the three heights (2 epidermis  $\times$  10 leaves  $\times$  3 heights  $\times$  4 hummocks). Fresh leaves were placed in zip-lock plastic bags, weighed and digitally photographed against an area reference on the day of sampling, then dried at 40 °C for 9 days in a drying oven with forced ventilation before reweighing to obtain their dry weight. Leaf areas were calculated from the photographs using ImageJ (US National Institute of Health, Bethesda, Maryland, USA: following Wang, 2017) and used to calculate the leaf mass area (LMA  $\text{mg mm}^{-2}$ ). The leaf area covering the optical window of the Dualex Scientific+ was 47.7  $\text{mm}^2$  (Area =  $\pi r^2$ ,  $r = 3.90$  mm), used with LMA, to calculate per area and per mass values of pigment concentrations on the harvest dates.

### 2.6. Environmental monitoring of temperature during the experiment

At the time of each measurement of leaf optical properties, the temperature of the leaves of *V. vitis-idaea* at each height on all four hummocks was measured at close range with a high precision infra-red thermometer with special macro optics (LS laser-sight Optris GmbH, Berlin, Germany). Measurements were also recorded at the tree base and at 1 m 50 cm height on the trunks (emissivity  $c$  0.98) of the four nearest trees to the hummocks at opposite compass points (North and South at midday) in the sun and shade, and on the snow surface (emissivity of tree trunks and snow  $c$  0.98).

Meteorological data for the experimental period were obtained from the SMEAR II ICOS Hyytiälä station through the AVAA data research portal (<https://avaa.tdata.fi/>). Temperature in the understorey was recorded every minute with a PT100 sensor (Pentronic, Västervik, Sweden) mounted on a mast at 4 m height and soil surface temperature was recorded every minute with a KTY 81-110 sensor (NXP Philips, Amsterdam, Netherlands). Understorey photosynthetically active radiation (PAR) was measured with an array of four Li190SZ PAR sensors (LI-COR, Lincoln, NE) at 0.6 m height next to the hummocks. These time series of meteorological data were used to calculate averages over a range of time periods preceding each measurement (1 h, 1 day, 2 days, and 2 weeks) and the relationship between temperature and PAR over these periods and leaf trait data was plotted. Of these four time integrals the best fit to the trait data was retained.

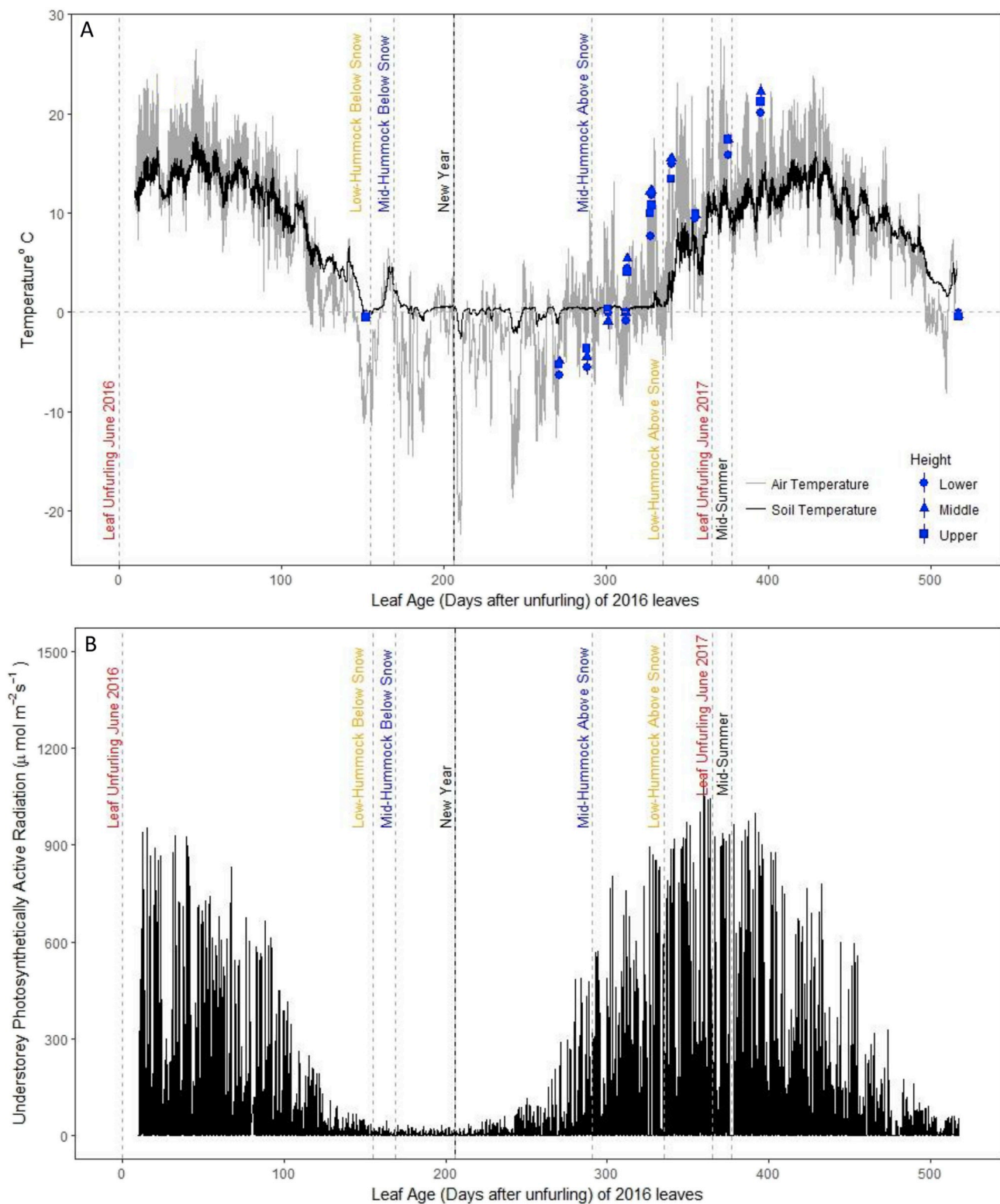
### 2.7. Data analysis

Differences due to hummock height among optically-assessed leaf traits were tested using a two-way ANOVA with hummock height, and leaf age (days) as fixed factors, or a three-way-ANOVA for the responses of flavonols and anthocyanins which were measured in both the adaxial and abaxial epidermis of each leaf (with leaf side as a nested third fixed factor). One measurement date (Leaf Age: 250 days), when only the adaxial epidermis was measured with the Dualex, was excluded from the three-way ANOVA. The equivalent single-factor ANOVA was used to test the effect of hummock height on traits measured only once, e.g. from harvested leaves at the end of the experiment. The four hummocks were the experimental units of replication and height was nested within each hummock. When significance differences were produced by the ANOVA, pairwise differences were identified between the responses across three levels of hummock height on either leaf side on each measurement date (function `glht` in the package `Multcomp`). Holm's adjustment was used to account for multiple pairwise comparisons. The relationship of  $F_v/F_m$  with temperature and PAR was assessed using linear regression (function `lm`()). All the analyses were performed using R version 3.4.3 (R Core Team, 2018).

## 3. Results

### 3.1. Forest cover, snow depth and temperature

The snow depth was  $c.$  40 cm at the first winter measurement (14-



**Fig. 1.** A. Time series of air and soil temperature with mean leaf temperature on dates of leaf trait measurements. B. Daily time course of understorey photosynthetically active radiation (PAR) at 0.6 m height, at Hyytiälä Forestry Station; from the unfurling of leaves in June 2016 (followed until November 2017), and including photosynthetic recovery period (June 2017 to November 2017) and subsequent new leaf unfurling in June 2017.

02-2017), meaning that the upper shoots of hummocks (upper hummocks) were above the snow, the middle of the hummocks were just below the top of the snowpack, and lower shoots of hummocks (lower hummocks) were entirely beneath the snowpack. From 14-02-2017 onwards, the snow depth didn't increase from this maximum and

remained above the middle-hummock shoots until 27-03-2017 and above the lower-hummock shoots until snow melt was complete on 10-05-2017.

Soil temperatures under the snow were relatively stable, starting to fluctuate only when snow melt was almost complete and the lower



**Table 1**

Maximum and Minimum Air and Soil Temperatures, and maximum PAR (PPFD) on the days of chlorophyll fluorescence and leaf pigments measurements. Freeze-thaw cycles are calculated for the period between measurements (usually 2 weeks, including 14-02-2017) and represent the number of times that the temperature passed above and returned below freezing during this period. Tree trunk temperatures were recorded at breast height on the completed shaded north side of the tree during winter and spring.

Date of Measurement	DOY	Leaf Age Days	Max-Air Temperature°C	Min-Air Temperature°C	Max-Soil Temperature°C	Min-Soil Temperature°C	Max-PAR $\mu\text{mol m}^{-2}\text{s}^{-1}$	Freeze-thaw cycles	Snow Temperature°C	Tree Trunk Temperature (North)°C
14-02-2017	45	250	5.3	−1.3	0.5	0.3	112.9	34	−	−
07-03-2017	66	271	−0.4	−10.4	0.0	−1.2	289.5	8	−7.9	−6.6
24-03-2017	83	288	1.8	−4.4	0.4	0.2	474.7	14	−10.0	−4.2
06-04-2017	96	301	8.6	−0.3	0.5	0.4	764.4	15	−3.5	0.8
18-04-2017	108	313	3.7	−7.3	0.3	−0.2	677.0	−	−2.6	−1.4
19-04-2017	109	314	5.5	−5.9	0.4	−0.1	519.8	14	−2.7	2.2
02-05-2017	122	327	12.2	0.2	0.6	−0.4	828.9	−	−3.0	8.7
03-05-2017	123	328	10.9	0.6	0.8	0.5	868.8	17	−2.8	8.1
16-05-2017	136	341	13.5	−2.7	3.4	1.1	919.6	−	−	7.7
17-05-2017	137	342	10.4	1.6	3.7	2.1	271.1	16	−	6.6
30-05-2017	150	355	13.6	0.1	7.6	4.7	290.0	−	−	9.8
19-06-2017	170	375	20.9	9.7	11.9	9.9	932.3	3	−	−
08-07-2017	189	394/29	20.3	7.2	12.7	9.2	836.8	1	−	−
09-07-2017	190	395/30	21.9	9.9	13.9	10.3	899.6	−	−	−
08-11-2017	312	517/152	5.2	2.7	4.3	0.4	51.7	34	−	−

hummock leaves above the snow (Fig. 1; Table 1). From the time series of snow depth and air temperature data, compared with IR-thermometer measurements of snow-pack temperature (Fig. 1), we were able to estimate that the upper-hummock leaves were subject to 111 freeze-thaw cycles, the middle-hummock leaves 86 and lower-hummock leaves 38 such cycles, during the period between New Year and mid-summer 2017 which encompasses full dehardening (Table 1).

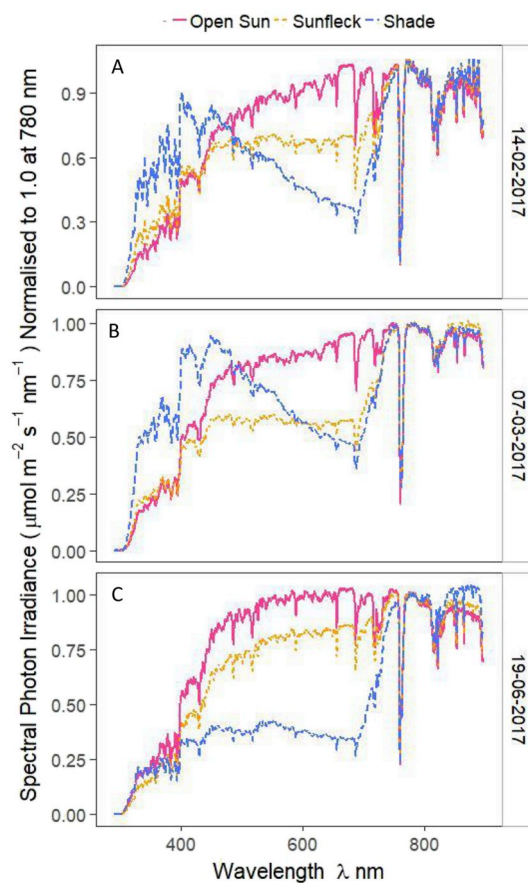
### 3.2. Spectral irradiance at the study site

The solar spectral irradiance was measured on three dates during the winter and spring (14-02-2017, 07-03-2017 and 19-06-2017). On all three dates, the normalised spectra show that PAR was depleted in the under-canopy sunflecks and shade compared with the open field, and the relative contribution of UV radiation was similar in the sunflecks and open sun but enriched in the shade (Fig. 2 and S2, Table S1). The spectral composition of the sunflecks varied with sunfleck size as well as time of year, and the relative contribution to shade by needles and branches also affected the shape of these spectra. The Plant Area Index (PAI) and sunfleck duration on 20-07-2017 were  $2.14 \pm 0.05 \text{ m}^2 \text{ m}^{-2}$  and  $169.6 \pm 11.3 \text{ min day}^{-1}$  respectively, and was similar among the four hummocks (Table S2).

The reflectance (up to 90%) and transmittance of sunlight by the snowpack modifies amount and composition of radiation received by leaves above and below the snowpack at different hummocks heights. We found the snowpack to transmit about 10% of incident PAR irradiance to a depth of 10 cm, and a similar or slightly higher percentage of UV radiation to reach the same depth (Robson unpublished data) in line with comparable measurements elsewhere (Richardson and Salisbury, 1977; Gorton and Vogelmann, 2003; Perovich, 2007).

### 3.3. Seasonal changes in leaf pigments at different heights on the *V. vitis-idaea* hummocks

Leaf flavonoid concentration (i.e. both flavonols and anthocyanins), measured optically and presented as an index of absorption on a per leaf-area basis, was most similar directly after leaf unfolding both among hummocks and between leaf sides. As the leaves matured, their flavonol concentration increased in the adaxial epidermis and initially decreased in the abaxial epidermis between the first and second measurement of new leaves during the summer (Leaf Age 10–29 days,



**Fig. 2.** Solar spectral Irradiance at midday measured at the experimental site under a pine canopy in a sunfleck and canopy shade, and in an adjacent open field, in clear sky conditions during the measurement period on (A) 14-02-2017, (B) 07-03-2017 and (C) 19-06-2017. Spectral photon irradiance is normalised to 1 at 780 nm (raw data at given in Fig. S2 and broken down in Table S1). (For interpretation of the references to colour in this figure legend, the reader is referred to the Web version of this article.)

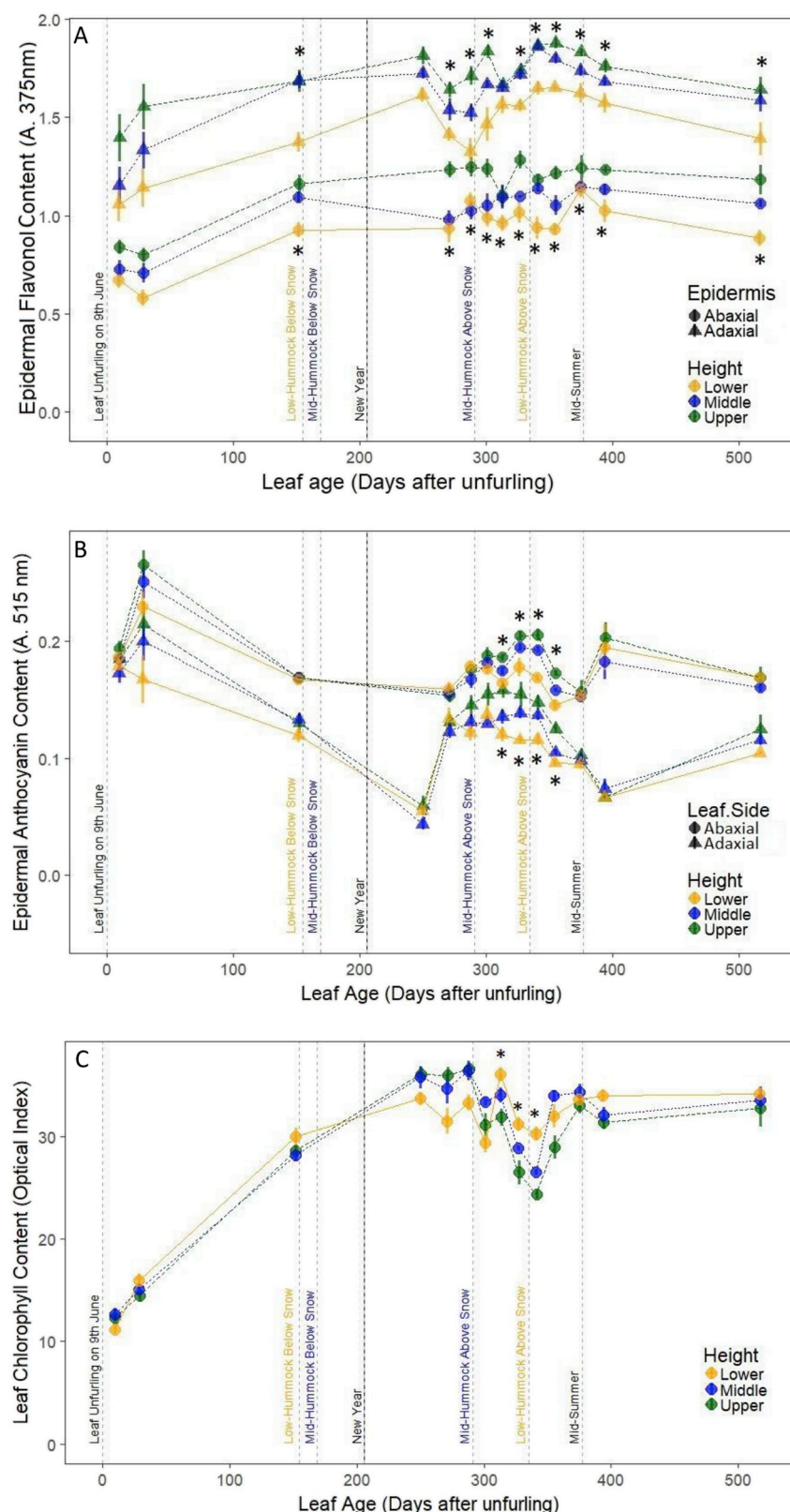


Fig. 3. Indices of (A) epidermal flavonols estimated from epidermal absorption at 375 nm; (B) epidermal anthocyanins, estimated from epidermal absorption at 515 nm, and (C) leaf chlorophyll estimated from the relative leaf transmission at 680–710 nm, on a leaf area basis. The period covered is from leaf emergence until leaves of *Vaccinium vitis-idaea* were one-and-a-half years old. Mean  $\pm$  1 SE of the 4 hummocks ( $n = 4$ ) are plotted for the adaxial (upper side) and abaxial (lower side) leaf epidermis, at the 3 heights on the hummock, and the period under the snowpack marked with vertical lines. \* indicates a significant difference with height, ANOVA summary Table S3.

Fig. 3A). During this period, UV-screening by flavonols was greater in the upper-, and middle-, hummock leaves than the lower leaves. Accumulation of flavonols continued until the start of winter in the adaxial-, and to a lesser extent the abaxial-, epidermis, and only started to definitely decline with age following the spring and summer of their second year (Leaf Age 271 days; Fig. 3A). Leaves of *V. vitis-idaea* can

remain productive for 5 years or more, but after their first growing season maximum photosynthesis declines sharply (Karlsson, 1985).

Adaxial epidermal flavonol concentration in *V. vitis-idaea* leaves transiently decreased during winter in lower- and mid-hummock leaves when they were beneath the snowpack, but started to increase again from 24-03-2017 (Leaf Age 288 days) coinciding with temperatures

rising above freezing. In sequence, the upper, then middle, then lower hummock leaves' adaxial flavonol concentration increased around the time that the middle hummocks leaves emerged from the snowpack (Fig. 3A, from 27-03-2017, Leaf Age 291 days). This phase of flavonol accumulation continued until the end of the snow melt period (16-05-2017 Leaf Age 341 days), after which flavonol concentration was stable and eventually started to decline from 19-06-2017 Leaf Age 375 days (Fig. 3A). Flavonol accumulation in the adaxial epidermis during spring was greater than that in the abaxial epidermis, leading the adaxial-to-abaxial flavonol ratio to increase until 08-07-2017, Leaf Age 394 days, but it returned to the equivalent of pre-spring ratios later in the year (08-11-2017, Leaf Age 517; Fig. 3A and S3A).

Leaf anthocyanin concentration is typically highest in young leaves, and higher in the abaxial than the adaxial epidermis, declining once leaves are mature (Landi et al., 2015): our results were consistent with this pattern until late winter, when anthocyanin accumulation rebounded (07-03-2017, Leaf Age 271 days, Fig. 3B). Initially, there were no differences in anthocyanin concentration with respect to height on the hummocks, and this remained the case until 24-03-2017 (Leaf Age 313 days) after the middle-hummock had emerged but while the lower hummock was still beneath the snow pack. During this period (24-03-2017 to 18-04-2017), adaxial epidermal anthocyanins continued to accumulate in the upper hummock leaves (Fig. 3B) and the difference in adaxial epidermal anthocyanin concentration with height was maintained, with higher values in upper hummock leaves compared with the others (Fig. 3B). In the abaxial epidermis, a corresponding difference with height occurred later during the 6-week spring dehardening period (19-04-2017 to 30-05-2017: Leaf Age 313–355 days), persisting until abaxial anthocyanin concentration was maximal in the upper and mid hummock leaves between (17/30-05-2017, Leaf Age 341–355 days). By this time, the abaxial epidermal anthocyanins had already started to decline in the lower-hummock (Fig. 3B). By mid-summer, both adaxial and abaxial epidermal anthocyanins had converged to similar values across the three height classes (19-06-2017: Leaf Age 375) (Fig. 3B). The ratios of adaxial-to-abaxial epidermal anthocyanins all declined throughout the spring, but differed among the three heights, being highest in the upper-hummock leaves and lowest in the lower-hummock leaves (Fig. S3B).

Chlorophyll concentration per leaf area increased in young leaves until 14-02-2017 (Leaf Age 250 days), by which time it was lower in the lower-hummock leaves than the middle-and-upper-hummock leaves (Fig. 3C). This difference persisted until the upper and middle hummocks were above the snowpack but the lower hummock beneath the snowpack (18-04-2017, Leaf Age 313 days), at which time chlorophyll concentration of the lower-hummock leaves increased beyond that of the exposed upper and middle hummock leaves where it declined (Fig. 3C). Lower-hummock leaves retained the highest chlorophyll concentration from then until midsummer by which time chlorophyll values at the three heights converged again (19-06-2017, Leaf Age 394 days) (Fig. 3C).

The ratio leaf flavonols vs. anthocyanins (sum of both epidermis) was higher in the upper hummocks than the lower hummocks through the winter (Fig. 4A). In the adaxial epidermis, the rate of flavonol accumulation was faster than that of anthocyanins during the dehardening period so this ratio rose, particularly in the lower hummock leaves (due to increasing flavonol accumulation as described above - Fig. 4A). There was negative relationship between flavonols and anthocyanins in the adaxial epidermis when data are plotted for all leaves and all dates (Fig. 4B). This relationship may be confounded by leaf age but nevertheless persisted for the adaxial epidermis when the relationships were calculated separately for each measurement date (Table S5).

### 3.4. Seasonal changes in photosynthetic yield of $PS_{II}$ in leaves at different heights on the *V. vitis-idaea* hummocks

The  $F_v/F_m$  of leaves across the three heights on the hummocks attained their maximal values of about 0.82 on reaching maturity on their second measurement following unfurling (09-07-2017, Leaf Age 30 days) and  $F_v/F_m$  remained high prior to snow cover in the winter (Fig. 5A). During the winter and spring,  $F_v/F_m$  was depressed most in the upper hummocks, then middle hummocks and least depressed in the lower hummocks (Fig. 5A). This ranking of  $F_v/F_m$  according to height on the hummocks was retained throughout the spring dehardening period (until 30-05-2017: Leaf Age 355 days), but by midsummer one-year old leaves at all three heights had attained similar maximal  $F_v/F_m$  again (19-06-2017, Leaf Age 375 days) (Fig. 5A).

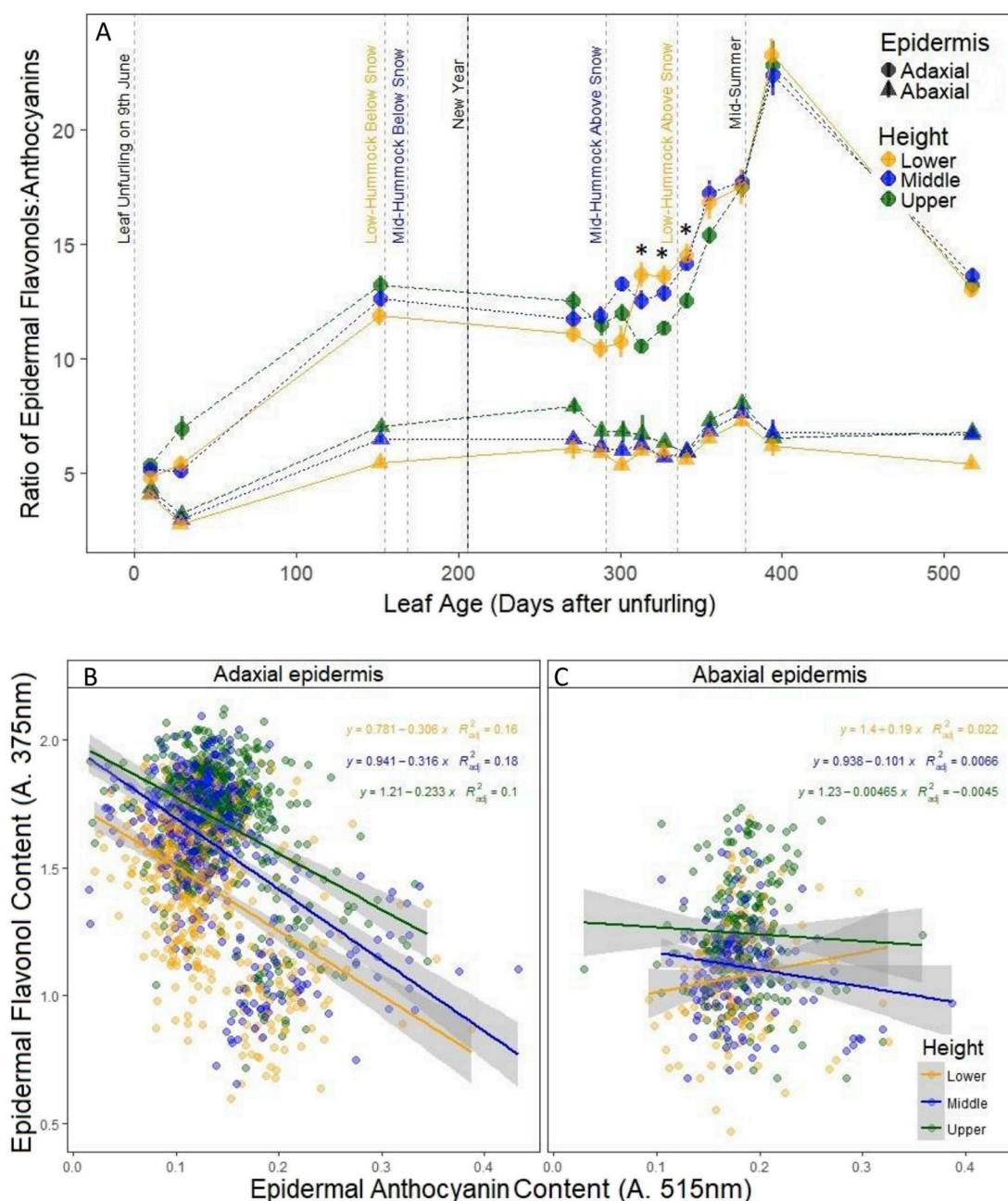
The  $\phi PS_{II}$ , measured *in situ* under ambient sunlight, varied with the solar irradiance, i.e. on clear-sky measurement days  $\phi PS_{II}$  was very low during the dehardening period (Fig. 5B). At sub-zero temperatures (e.g.  $-5.5^\circ\text{C}$  on 07-03-2017, Leaf Age 313 days) with clear-sky irradiance during early spring, the  $\phi PS_{II}$  was lowest and approached zero. Though bear in mind that it was necessary to remove the snow from covered leaves during winter to take the measurement, exposing them briefly to higher PAR than they would otherwise have received and thus probably caused greater photoinhibition than when under the snow. Nevertheless, in early spring on days 83 and 96 (leaf age 288 and 301 days),  $\phi PS_{II}$  of almost 0.2 was reached in the lower hummock leaves (Fig. 5B). Both measures of photosynthetic capacity ( $F_v/F_m$  and  $\phi PS_{II}$ ) responded to day-to-day differences in environmental conditions, as evident from differences between  $F_v/F_m$  values from consecutive daily measurements made on three occasions during the spring dehardening period (Fig. 5B, Leaf Age 313–314, 327–328, 341–342 days).

### 3.5. The relationship between *V. vitis-idaea* photosynthetic yield, leaf flavonoids and environmental conditions

We tested the relationship between temperature and  $F_v/F_m$  for leaves from the upper, middle, and lower hummocks, identifying their position relative to the snowpack prior to each  $F_v/F_m$  measurement. The air temperature ( $T_{aw}$ ) or soil temperature ( $T_{sw}$ ) at the site were positively correlated with the average  $F_v/F_m$  of leaves above or below the snowpack respectively, and in order to make a direct comparison with other studies of *V. vitis-idaea* (Bokhorst et al., 2010; Saarinen et al., 2016), we tested a range of different time periods to find the closest relationship. Of these time periods, the temperature over the 2 weeks prior to each  $F_v/F_m$  measurement gave the best fit; for leaves above the snowpack vs.  $T_{aw}$  (adj.  $R^2 = 0.91$ , Table S4), and leaves below the snowpack vs.  $T_{sw}$  (adj.  $R^2 = 0.81$ , Table S4). These relationships with temperature were stronger than  $F_v/F_m$  vs. PAR on the day of measurement for leaves above the snowpack (adj.  $R^2 = 0.74$ , Table S4).

Individual leaf measurements of  $F_v/F_m$  were also positively correlated with leaf temperature (Fig. 6A) and PAR incident on the leaf (Fig. 6B) at the time of measurements, and likewise with adaxial-epidermal flavonol concentration on the day of measurement (Fig. 6C). However, these relationships were strongly conditioned by measurement day and held only during the period from 24-03-2017 to 17-05-2017: Leaf Age 281–342 days (Fig. 6). The relationships broke down once spring recovery of  $F_v/F_m$  was almost complete (30-05-2017 Leaf Age 355 days – not shown) and in winter when leaf temperatures were consistently around  $0^\circ\text{C}$ . During dehardening, all three relationships were strongest in the upper hummocks and remained strong in the middle hummocks, where  $F_v/F_m$  recovery coincided with an increase in adaxial epidermal flavonols during this time period, but were but much less strong in the lower hummocks (Fig. 6C). The same relationship was not evident for anthocyanins in either epidermis (not shown).





**Fig. 4.** (A) Time course of the relationship between estimated epidermal flavonols and anthocyanins from Fig. 3A and B. Scatterplots and linear regression of (B) adaxial and (C) abaxial epidermal flavonols and anthocyanins, for all measurement dates together but the three hummock heights presented separately for comparison. \* indicates a significant difference with height, ANOVA summary Table S3.

## 4. Discussion

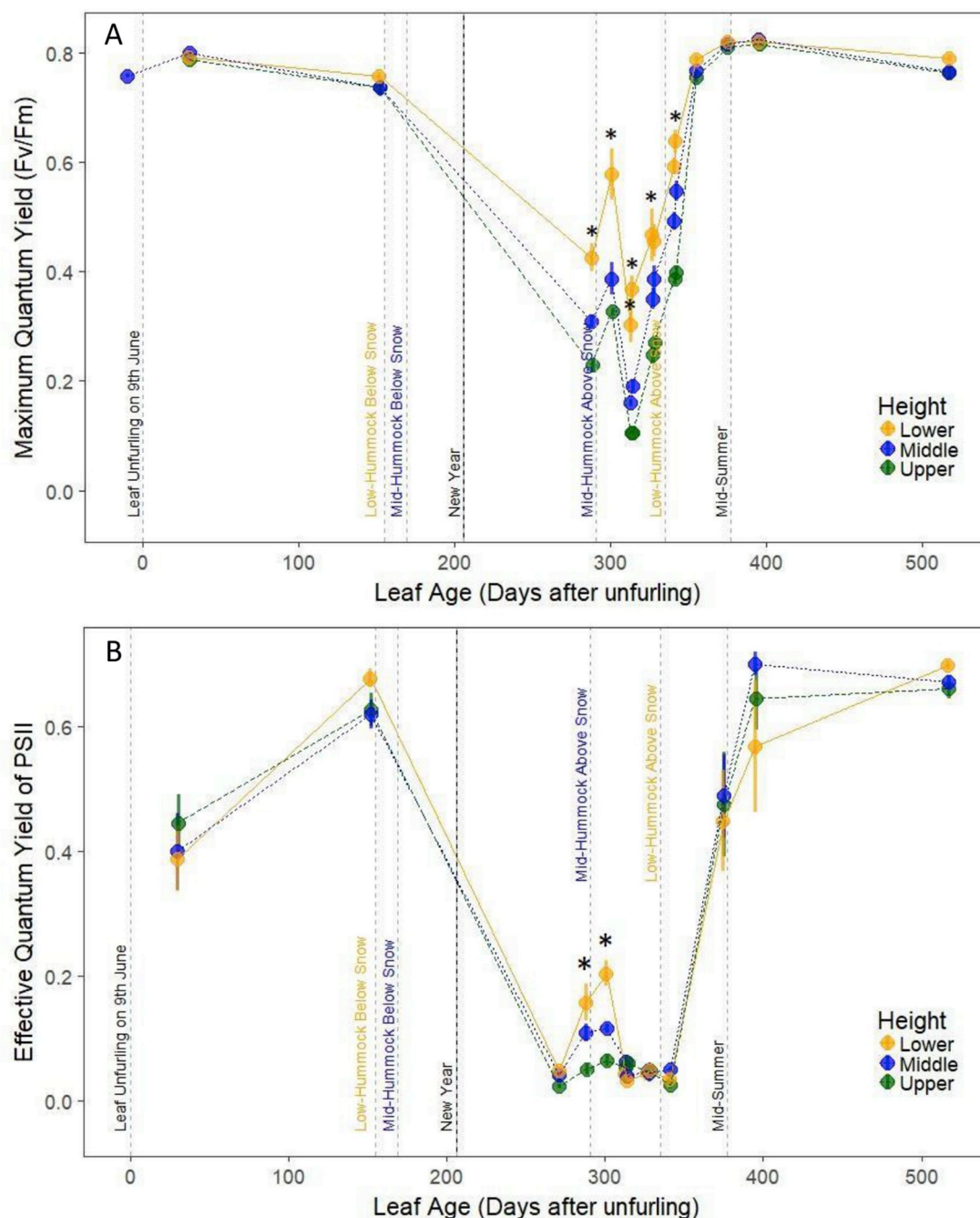
### 4.1. Consistent pattern of increasing flavonoids with hummock height throughout the year

Differences in exposure to harsh weather conditions among adjacent plants or shoots may be particularly pronounced in the understorey of open canopies typical of the boreal forest which unevenly intercept solar radiation (both incoming and reflected or emitted radiation) and snow (Hedstrom and Pomeroy, 1998). The undulating topography created by hummock-forming plants, adds to the heterogeneity of irradiance, temperature, and snow cover (Essery et al., 2008; Pomeroy et al., 2002; Sturm et al., 2001). Accordingly, a large proportion of their leaves can remain above the snowpack over winter where they are subject to greater fluctuations in temperature and irradiance than those

beneath the snowpack. In our study, this was the case for the upper leaves of *V. vitis-idaea* hummocks, which accumulated more epidermal UV-screening (attributable to flavonols) than lower hummock leaves. Using a Dualex to optically detect UV-screening allowed us to repeatedly and non-destructively follow changes over time on the same plant but not to gather biochemical information on flavonoid composition.

The results of studies based on optical properties of the leaf epidermis *in vivo* and measurements of whole-leaf extracts can be difficult to reconcile because the two methods are not equivalent. In *V. vitis-idaea*, flavonoids can be found in the trichomes and bound to the epidermal cell walls (Semerdjieva et al., 2003a, 2003b) where they will affect UV-screening, whereas they are spread throughout the leaf including chloroplasts in the mesophyll and cuticular waxes in species like *V. myrillus* (Semerdjieva et al., 2003b). Despite a certain lack of

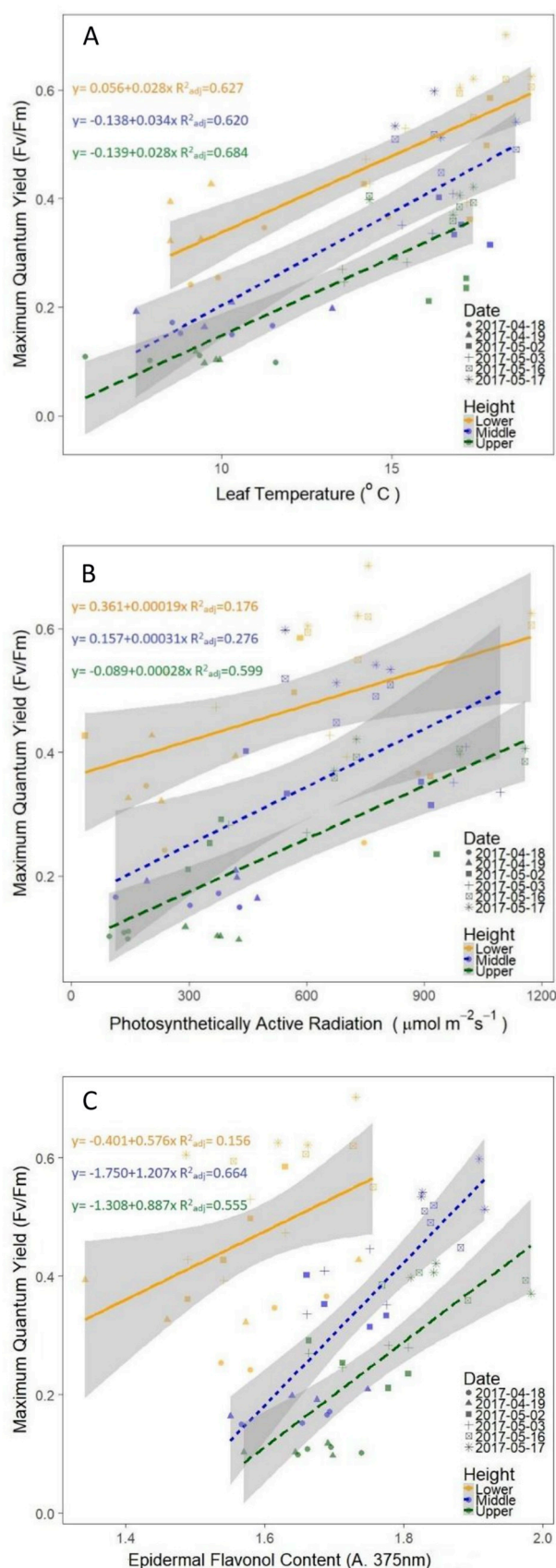




**Fig. 5.** Time course of (A) maximum quantum yield of photosystem II ( $F_v/F_m$ ) and (B) effective quantum yield of photosystem II ( $\phi_{PSII}$ ) over the period from leaf emergence until leaves of *V. vitis-idaea* were 517 days old. Mean  $\pm$  1 SE of the 4 hummocks ( $n = 4$ ) are plotted, and the period under the snowpack marked with vertical lines. \* indicates a significant difference with height, ANOVA summary Table S3. The fluorescence parameters used for these calculations are given in Fig. S4.

equivalency, it is worth noting that the relative differences we found in UV-screening within *V. vitis-idaea* hummocks, among leaves differing only in the time spent above and below the snowpack (Fig. 1), were larger than those produced in the same species by a 25% UV supplementation (Semerdjieva et al., 2003b). A strong negative correlation between accumulation of leaf flavonoids and increasing temperature can obscure the effects of UV-B radiation (Coffey et al., 2017), and consequently, the differences in temperature across hummock height related to differences in seasonal snow cover may partially explain the vertical gradient in flavonols accumulation we report. It has not yet been elucidated whether both temperature and UV-B radiation may regulate flavonoid accumulation via the UVR8 pathway, but this would constitute an interesting future avenue of research.

The response of *V. vitis-idaea* leaves to UV radiation at high latitudes has already been extensively studied in the context of northern-hemisphere stratospheric ozone depletion in the late 1990s and 2000s (e.g. Johansson et al., 1995; Phoenix et al., 2001; Semerdjieva et al., 2003a). The leaves of evergreen species emerge from snow cover in spring when seasonal UV-B exposure can be periodically high, but a meta-analysis generally found evergreen species like *V. vitis-idaea* to be resilient against such seasonal increases in UV radiation (Newsham and Robinson, 2009). The modest flavonoid induction caused by increased UV-B radiation could help protect from photodamage to  $PS_{II}$  and improve antioxidant capacity during dehardening at the start of the growing season when high irradiance and cold temperatures in unison exacerbate photoinhibition (Takahashi and Badger, 2011).



**Fig. 6.** The relationship between  $F_v/F_m$  and (A) leaf temperature, (B) PAR and (C) epidermal flavonols at the time of measurement measured on six occasions. Each point is the mean of 10 measurements of each of 4 hummocks. These relationships only hold during the spring dehardening period (17-04-2017 to 18-05-2017) when the temperature is above freezing but leaves have not yet recovered to maximal  $F_v/F_m$  of 0.82.

Most studies from the 1990s-early-2000s have presented these data on a per mass basis which is not as easily interpretable as leaf-area based measures in terms of screening. Differences in LMA among the one-year old leaves in our experiment caused the effect of hummock height on flavonols and anthocyanins per-leaf-area to be reduced when considered on a per-mass basis (Table 2). Likewise, the effect of UV-B radiation on flavonol accumulation is sometimes smaller on a per-leaf-mass than per-leaf-area basis in understory shrubs, because UV-B exposure causes mature leaves to be thicker and tougher (Newsham and Robinson, 2009; Semerdjieva et al., 2003a).

The increase in the ratio of adaxial to abaxial epidermal flavonols in the lower hummock leaves (Fig. S3A) during snow melt suggests that accumulation of adaxial flavonols could aid in photoprotection in addition to amelioration of cold temperature stress as antioxidants (Chalker-Scott and Scott, 2004). Comparison of the UV screening on either side of a leaf show this relationship to be highly species specific and environmentally dependent: sometimes UV screening by the adaxial and abaxial epidermises are strongly positively correlated (Barthod et al., 2007; Siipola et al., 2014) sometimes this relationship is weak (Kolb and Pfundel, 2005; Morales et al., 2011), as was the case in our study. The accumulation of epidermal flavonoids is commonly attributed to exposure to UV-B radiation, or high irradiance of UV-A radiation and blue light (Barnes et al., 2017), although flavonoid accumulation is also known to increase in response to cold temperatures (Agati et al., 2013). Temporal differences in flavonoid (both flavonol and anthocyanin) accumulation and their localization in the planophile leaves of *V. vitis-idaea* might reflect differences in the allocation of resources based on differences between the adaxial and abaxial epidermis in their relative exposure to cold and high irradiance.

Over the whole dataset, there was an inverse correlation between the anthocyanin and flavonol concentrations in the adaxial epidermis, even though during spring dehardening both tended to increase (Fig. 4). A similar pattern showing a trade-off between these two phenolic compounds was reported in *Arabidopsis thaliana* under controlled conditions, where the ratio of flavonoids to anthocyanins was higher when they received either blue light or UV-A radiation (Brelsford et al., 2018). Anthocyanins are typically found in greater amounts in young leaves and have a specific role as antioxidants involved in cold and high-light tolerance (Havaux and Kloppstech, 2001; Landi et al., 2015). Although anthocyanins only infrequently respond to elevated UV-B radiation (Searles et al., 2001), both flavonols and anthocyanins are phenolic compounds of the flavonoid pathway and it is possible that regulation of this pathway shifts during spring to emphasize the photoprotective function of flavonols driven by seasonally-increasing irradiance of blue light and UV radiation at the expense of anthocyanins whose antioxidant function would be more important during winter (Agati et al., 2013).

#### 4.2. Prolonged snow cover accelerates recovery of photosynthetic capacity during spring dehardening

The timing of spring dehardening in our study was better correlated with temperature than irradiance, and the relationship with temperature was maintained over 2 weeks prior to measurement (Fig. 6), in agreement with Karlsson (1985). Several studies have found a consistent relationship between the ambient temperature and  $F_v/F_m$  of *V. vitis-idaea* leaves across a range of latitudes (Bokhorst et al., 2010; Saarinen et al., 2016; Table S6 and S7) and in *Pinus sylvestris* at our

**Table 2**

Comparison of leaf pigment measurements and leaf morphological traits on per area and per mass basis. Final harvest of one-year old leaves (leaf age 394 days) and young leaves (leaf age 29 days) leaves of *V. vitis-idaea* in July 2017. Mean of 4 hummocks  $\pm$  1 SE. Percentage changes of upper and middle hummocks from lower hummock values are given in parentheses. ANOVA results given in Table S5.

Leaf Trait	Young leaves (Leaf Age 29 days old)			One-year old leaves (Leaf Age 394 days old)		
	Lower Hummock	Middle Hummock	Upper Hummock	Lower Hummock	Middle Hummock	Upper Hummock
<b>Morphological Trait</b>						
Leaf Area (mm <sup>2</sup> )	240 $\pm$ 9	240 $\pm$ 15 (0%)	246 $\pm$ 12.3 (+3%)	174 $\pm$ 11	206 $\pm$ 27 (+18%)	188 $\pm$ 20.6 (+8%)
Leaf Dry Matter Content (mg)	14.5 $\pm$ 1.9	13.6 $\pm$ 1.5 (−6%)	13.7 $\pm$ 1.6 (−5%)	28.9 $\pm$ 2.4	35.8 $\pm$ 4.8 (+24%)	34.9 $\pm$ 3.9 (+21%)
Leaf Mass Area (mg mm <sup>−2</sup> )	0.060 $\pm$ 0.006	0.056 $\pm$ 0.003 (−6%)	0.055 $\pm$ 0.005 (−8%)	0.166 $\pm$ 0.005	0.174 $\pm$ 0.01 (+5%)	0.187 $\pm$ 0.01 (+12%)
<b>Dualex (per area)</b>						
Chlorophyll index per leaf area (Dualex A mm <sup>−2</sup> )	78.6 $\pm$ 5.8	74.6 $\pm$ 6.4 (−5%)	77.8 $\pm$ 4.8 (−1%)	122.6 $\pm$ 5.6	136.8 $\pm$ 23.4 (+12%)	121.7 $\pm$ 17 (−1%)
Epidermal flavonol index per leaf area (Dualex A mm <sup>−2</sup> )	5.6 $\pm$ 0.7	6.8 $\pm$ 0.4 (+22%)	8.0 $\pm$ 0.7 (+43%)	5.7 $\pm$ 0.1	7.2 $\pm$ 0.9 (+25%)	6.8 $\pm$ 0.9 (+19%)
Epidermal anthocyanin index per leaf area (Dualex A mm <sup>−2</sup> )	0.89 $\pm$ 0.01	1.03 $\pm$ 0.07 (+16%)	1.07 $\pm$ 0.08 (+20%)	0.26 $\pm$ 0.03	0.36 $\pm$ 0.03 (+39%)	0.34 $\pm$ 0.02 (+29%)
<b>Dualex (per mass)</b>						
Chlorophyll index per leaf mass (Dualex A mg <sup>−2</sup> )	1346 $\pm$ 162	1322 $\pm$ 52 (−2%)	1445 $\pm$ 173 (+7%)	739 $\pm$ 14	798 $\pm$ 154 (+8%)	657 $\pm$ 99 (−11%)
Epidermal flavonol index per leaf mass (Dualex A mg <sup>−2</sup> )	93 $\pm$ 6	120 $\pm$ 4 (+30%)	146 $\pm$ 13 (+57%)	35 $\pm$ 1	42 $\pm$ 6 (+20%)	37 $\pm$ 5 (+6%)
Epidermal anthocyanin index per leaf mass (Dualex A mg <sup>−2</sup> )	15.2 $\pm$ 1.3	18.3 $\pm$ 0.6 (+20%)	19.4 $\pm$ 0.3 (+27%)	1.6 $\pm$ 0.2	2.1 $\pm$ 0.3 (+34%)	1.8 $\pm$ 0.1 (+15%)

Hyttiälä site (Porcar-Castell, 2011). Saarinen et al. (2016) report that  $F_v/F_m$  in leaves of *V. vitis-idaea* plants was depressed by snow removal or even when covered by only a thin layer of snow. In our study, upper-hummock leaves had the lowest  $F_v/F_m$  and mid-hummock leaves, which were also only thinly snow-covered, had lower  $F_v/F_m$  than lower-hummock leaves. Prior to their emergence from under the snow, the  $F_v/F_m$  of lower-hummock leaves was already increasing. It is conceivable this recovery was aided by the low to moderate irradiance that penetrates the upper cm's of the snowpack, potentially activating photoprotective mechanisms during snowmelt (Richardson and Salisbury, 1977). This could partially explain why the difference in  $F_v/F_m$  was maintained between leaves at different heights on the hummocks for several weeks following snow melt rather than the lower leaves exhibiting a cold-shock response upon exposure.

Large fluctuations in temperature and sustained winter warming can both prematurely activate dehardening creating the potential to bring forward the growing season but also increasing the risk of damage if cold winter temperatures return (Bokhorst et al., 2008). Photo-inhibition of exposed shoots was high in our study, and leaf chlorophyll concentration dropped in the upper and middle hummock leaves during the spring dehardening period but remained stable in the lower hummock leaves (surpassing that in the upper hummocks) following emergence from under the snowpack. The regulation of leaf chlorophyll may be a means of acclimation, to reduce photon capture under high irradiance to balance the limited carboxylation capacity at low temperature in the upper and middle hummocks when photoprotective mechanisms were surpassed.

Winter- and spring-time depression of  $F_v/F_m$  and  $\Phi PS_{II}$  was largely attributable to a decrease in maximal fluorescence ( $F_m$  and  $F_m'$  respectively) rather than an increase in  $F_0$  or  $F_0'$  (Fig. S5). This suggests that the reduction in photosynthetic capacity is the result of decreased light harvesting by  $PS_{II}$  lowering the  $F_m$  and  $F_m'$ . Such a result has previously been attributed to smaller antenna size or sustained NPQ associated with decreased epoxidation of xanthophylls or  $PS_{II}$  reaction centre quenching under cold temperatures (Ivanov et al., 2008), rather than damage or disassembly of the  $PS_{II}$  core which would be reflected in higher  $F_0$  or  $F_0'$  (Öquist and Huner, 2003). The positive correlation between epidermal flavonols and  $F_v/F_m$  found in the upper and middle, but not lower, hummock leaves during dehardening suggests that these leaves are acclimating to the more-exposed conditions. This is consistent with some mechanistic studies which report flavonoids to determine of cold acclimation and frost tolerance of *Arabidopsis thaliana*

(Havaux and Kloppstech, 2001; Schulz et al., 2016). However, knowledge of the contribution of xanthophyll cycle pigments at different hummock heights, which was not quantified in this study, would allow us to better describe this response (Adams et al., 2004).

#### 4.3. Ecological consequences of changing snow cover for *V. vitis-idaea*

In our study in central Finland, all leaves achieved full functionality following dehardening. Likewise, Saarinen et al. (2016), found that exposed leaves of *V. vitis-idaea*, subjected to experimental removal of the snow pack during winter, fully recovered  $PS_{II}$  capacity during the spring in southern Finland, albeit later than those from which snow was not removed. This result is consistent with other Arctic shrub species, e.g. *Rhododendron ferrugineum* L., whose leaves above the snow in winter have transiently reduced  $F_v/F_m$  compared with those that are covered by snow (Neuner et al., 1999). These findings suggest that *V. vitis-idaea* could benefit from the expected warmer winters and earlier snow melt in Finland, and much of the rest of its range (Rasmus et al., 2004; Henry, 2008; Jylhä et al., 2008), by dehardening earlier and contributing more in the future to photosynthetic carbon gain in the understorey (Gehrmann et al., 2018; Larsen et al., 2007). However, this does not exclude the possibility that environmental changes involving reduced duration or size of snowpack can also be detrimental to understorey plants. Further north than our study, across a range of habitats in the Arctic where winters are colder than in central Finland, a reduced snow pack can cause sufficient cold stress to damage the leaves and shoots of shrubs including *V. vitis-idaea* (Preece and Phoenix, 2013). Reduced snow cover can also lead to colder soils and prolonged freezing temperatures which can be a problem over a wider geographical range across Europe causing root mortality and affecting biogeochemical cycling (Groffman et al., 2001; Helama et al., 2011).

The link between increasing temperature and irradiance, and  $F_v/F_m$  recovery in *V. vitis-idaea* during dehardening reinforces established knowledge of this and other species (Ensminger et al., 2004; Havaux and Kloppstech, 2001; Muller et al., 2005; Neuner et al., 1999; Porcar-Castell et al., 2008; Saarinen et al., 2016). At our site in central Finland, winter temperatures are relatively mild ( $-7.37 \pm 1.15$  °C; 10-year mean  $\pm$  1 SD January temperature, 2008–2018), and are expected to increase by 4–11 °C by the end of the century (Jylhä et al., 2008; Finland's Seventh National Communication under the United Nations Framework Convention on Climate Change, 2017). Winter warming is forecast to increase productivity at the stand level (Porcar-Castell et al.,



2008). However, the expected 30–40% reduction in snow cover (Jylhä et al., 2008; Finland's Seventh National Communication under the United Nations Framework Convention on Climate Change, 2017) may lead to increased soil freezing or freeze thaw cycles which may partially negate the increase in productivity (Taulavuori et al., 2011). New estimates of ecosystem carbon assimilation aim to include the contribution of understorey vegetation in addition to that by tree species (Nikopienius et al., 2015; Starr and Oberbauer, 2003). To estimate understorey photosynthesis during spring a better understanding of how it is affected by uneven snow cover under patchy canopies will be needed (Blume-Werry et al., 2016; Kreyling, 2010).

## 5. Conclusions

In our study, there was a positive correlative relationship during the winter and spring between UV-screening by epidermal flavonols and  $F_v/F_m$  in leaves that were largely above the snowpack. More flavonols and anthocyanins accumulated in these upper-hummock leaves compared to leaves below the snowpack. The cross-protection provided by epidermal flavonols to cold and high light is likely to be regulated by solar radiation and temperature, but further research is needed to disentangle the roles of these two cues. Recent evidence that leaves can rapidly adjust their UV-screening to acclimate to light conditions (Barnes et al., 2017) suggests that a high-degree of fine tuning of these responses is possible. This assertion is supported by our finding that flavonoids (flavonols and anthocyanins) accumulate in the upper hummocks even in cold conditions when metabolic activity is low. Leaves with a higher index of epidermal flavonols tended to have a lower epidermal anthocyanin index, and a compositional shift in this ratio towards flavonols during spring suggests fine-tuning of acclimation in response to increasing irradiance and a less-cold temperatures. Finally, on hummocks at our site in central Finland, leaves that remained under snow cover longer surpassed exposed leaves in their  $F_v/F_m$  directly following snowmelt despite having lower epidermal flavonols and anthocyanins but all leaves eventually reached a similar maximum photosynthetic capacity.

## Declaration of authorship

TMR SMH conceived & designed the experiments. TS, SN, SMH, MP TMR, did the majority of the field work and sampling, TS, MP, AS, PJA TMR analysed the data, JAPC JA provided site-specific help, and access to meteorological data integral to the experiment, TS, PJA, TMR wrote the manuscript, and all authors provided editorial input.

## Funding

This study was funded by the Academy of Finland (grant number # 304519) to TMR. Grants from Finnish National Agency for Education (EDUFI fellowship program) and the Tiina and Antti Herlin Foundation (project number 20180642) funded TS.

## Acknowledgments

Craig C Brelsford and Marieke Trasser helped to do the field work. We thank Timo Saarinen and Stef Bokhorst, for making their raw data available for our comparisons of  $F_v/F_m$ , light and temperature.

## Appendix A. Supplementary data

Supplementary data related to this article can be found at <https://doi.org/10.1016/j.plaphy.2018.09.003>.

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