Carbohydrates and secondary compounds of alpine tundra shrubs in relation to experimental warming

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Abstract
Background: It is critical to understand the sensitivity, response direction and magnitude of carbohydrates and secondary compounds to warming for predicting the structure and function of the tundra ecosystem towards future climate change.

Results: Open-top chambers (OTCs) were used to passively increase air and soil temperatures on Changbai Mountain alpine tundra. After seven years’ continuous warming (+ 1.5 °C), the vegetation coverage, nonstructural carbohydrates (soluble sugars and starch) and secondary compounds (total phenols, flavonoids and triterpenes) of leaves and roots in three dominant dwarf shrubs, Dryas octopetala var. asiatica, Rhododendron confertissimum and Vaccinium uliginosum, were investigated during the growing season. Warming did not significantly affect the concentrations of carbohydrates but decreased total phenols for the three species. Carbohydrates and secondary compounds showed significantly seasonal pattern and species-specific variation. No significant trade-off or negative relationship between carbohydrates and secondary compounds was observed. Compared to Dr. octopetala var. asiatica, V. uliginosum allocated more carbon on secondary compounds. Warming significantly increased the coverage of Dr. octopetala var. asiatica, did not change it for V. uliginosum and decreased it for Rh. confertissimum. Rh. confertissimum had significantly lower carbohydrates and invested more carbon on secondary compounds than the other two species.

Conclusions: Enhanced dominance and competitiveness of Dr. octopetala var. asiatica was companied by increased trend in carbohydrate concentrations and decreased ratio of secondary compounds to total carbon in the warming OTCs. We, therefore, predict that Dr. octopetala var. asiatica will continue to maintain dominant status, but the competition ability of V. uliginosum could gradually decrease with warming, leading to changes in species composition and community structure of the Changbai tundra ecosystem under future climate warming.

Keywords: Soluble sugars, Starch, Flavonoids, Phenols, Triterpenes, Coverage

Background
Global air temperature has been predicted to increase continuously, with the most marked increase in alpine and tundra regions [1]. During the past decades from 1950 to 2010, Changbai Mountain alpine tundra experienced significant increases in both temperature and precipitation and decreases in frost and ice days. The warming rate of growing season is 0.0239 °C y⁻¹, higher than that of the average earth surface temperature. The ice day decreased 0.2245 d y⁻¹ and the increased rate of potential evapotranspiration is 0.6312 mm y⁻¹ [2]. In addition, it is hard to see the snow on the top of the Changbai Mountain in summer. Temperature change alone, and in combination with other environmental changes,
will inevitably have considerable impacts on ecosystems [3]. Tundra vegetation on the Changbai Mountain has changed significantly over the last decades, showing that the abundance of shrubs decreased, whereas the grasses’ abundance continuously significantly increased [4]. For instance, the grasses that previously either occurred in the mountain birch forests at a lower elevation or were only occasionally observed in the tundra have extensively invaded the Changbai Mountain alpine shrub tundra, and the tundra vegetation is currently co-dominated by shrubs and six herb species (Calamagrostis angustifolia, Geranium baishanense, Ligularia jamessii, Sanguisorba parviflora, S. stipulata, and Saussurea tomentosa) [5, 6]. Jin et al. stated very recently that the Changbai Mountain shrub tundra will be replaced by a grass tundra, and which is mainly regarded as a consequence of continuous air warming [7].

Strong and sensitive reactions in phenology, photosynthesis, growth, leaf nutrients, and carbon contents and components of alpine and arctic plants to climate warming, including positive and negative responses, have been observed [8–12]. A question that remains unclear is whether species that respond negatively to warming will have disadvantages and thus must escape from the community under future climate warming. For example, it has been observed that two moss species completely disappeared, and the abundance of three dwarf shrub species decreased, but the abundance of forb and grass species did not change after four years’ warming in an alpine plant community, southwestern Norway [13]. Two previously dominant shrubs, Rhododendron chrysanthum and Vaccinium uliginosum, on the west-facing slope of Changbai Mountain alpine tundra showed different responses to climate change over the last decades. V. uliginosum significantly decreased its abundance and changed its previously even distribution pattern into a patch distribution, while Rh. chrysanthum did not [7]. Whether such decreased abundance of some species or disappearance of moss is related to physiologically negative responses to climate change, and/or whether the change in community structure can be explained by physiological competition relationships among species are poorly understood [13].

Competitive relationship among co-existing species is affected by many factors. The competitive ability of a species is influenced by available nonstructural carbohydrates (NSCs, mainly sugars and starch) and secondary metabolism and vice versa [14, 15]. Competition may affect the distribution and allocation pattern of carbon between growth and defense. NSCs and secondary compounds support plant physiological processes of vegetative and reproductive growth, maintenance, storage, and defense, hence, they may be allocated for various purposes, affecting competition or trade-offs between NSCs and secondary compounds at an individual level [16]. For example, more investment of carbon to plant growth could lower plant concentrations of carbon-based secondary compounds [17]. Growth on tundra is limited by low temperature. According to hypotheses of resource-based defence, alpine plants might invest more carbon in defence. When temperature increases, increased resource availability will prioritize growth and spend less on defence [17]. The relative growth rate of four trembling aspen was negatively related to condensed tannin content [14]. The carbon allocation to defensive compounds decreased for tomato due to competition [18]. The competition led to a decrease in starch concentration for two Larix species, but soluble sugars and total NSCs (TNC) concentrations were not directly influenced by competition [15].

Changes in environmental conditions will alter the allocation of photosynthetic fixed carbon between primary and secondary compounds and the responses might be highly species specific. Aerts et al. observed that the carbon concentration of three co-existing sub-arctic dwarf shrubs – Empetrum hermaphroditum, Andromeda polifolia and V. uliginosum responded significantly differently to experimental warming [19]. Starch reserves increased across 14 tree species along a natural temperature gradient from lowland to the alpine treeline [20]. According to a simulated warming experiment, elevated temperature did not significantly change the concentrations of soluble sugars and starch in needles, bark and wood of Larix decidua grown at the alpine treeline [21]. The carbon-based secondary compound concentrations of Tofieldia pusilla were decreased by warming, while Saussurea alpina, Carex vaginata, V. uliginosum, Selaginella selaginoides in the same growth environment did not respond to the warming treatment [17]. The tannin concentrations of Cassiope tetragona and V. vitis-idaea leaves were increased by warming [22]. We wonder, therefore, whether a synchronized change or a trade-off between NSCs for growth and the secondary compounds for defense exists in tundra plants under global warming.

We used a 7-year experimentally warming site (+ 1.5 °C in the air temperature and + 0.8 °C in the soil temperature at 10 cm depth) in the alpine tundra on the north-facing slope of Changbai Mountain to study whether changes in plant coverage are associated with changes in carbon resource availability and allocation between growth and defense. We measured NSCs (soluble sugars, starch) and carbon-based secondary compounds (total phenols, flavonoids and triterpenes) in three dominated shrub species (Dryas octopetala var. asiatica, V. uliginosum, Rh. confertissimum) which have shown marked variation in their coverage responses to warming over time.
(see Table 1). We hypothesize that (1) a decrease in the coverage of species implies negative responses of NSCs to warming, and vice versa; (2) the levels of NSCs and carbon-based secondary compounds of the three species respond to warming differently, showing a significant interaction between species and warming; and (3) there is a trade-off between NSCs and secondary compounds to warming. We aimed to assess the effects of warming on NSCs and secondary compounds in dominant shrubs on alpine tundra, to investigate the link between the change in vegetation coverage and NSCs as well as secondary compounds, and thus to understand whether the plant carbon-physiological parameters can be used as useful indicators to predict vegetation change (winner or loser) under global warming or environmental changes.

**Results**

**Responses of species coverage to warming**

Experimental warming significantly increased the coverage of *Dr. octopetala var. asiatica* (*P* < 0.01), but it highly significantly decreased (*P* < 0.001) the coverage of *Rh. confertissimum* (Table 1). The coverage of *V. uliginosum* was not markedly changed by warming (Table 1). After seven years of warming treatment, the coverage of *Rh. confertissimum* in the warming OTCs was very small, less than 5% (Table 1).

**Leaf nonstructural carbohydrates**

Warming treatment had no marked effects on leaf soluble sugars, starch and TNC concentrations, as well as the ratio of sugars to starch, but all the four traits varied over time and were significantly species-specific (Table 2). Generally, the concentrations of soluble sugars were relatively higher, and the starch concentration was lower in both *Dr. octopetala var. asiatica* and *V. uliginosum* (Fig. 1).

Starch and TNC concentrations of *Dr. octopetala var. asiatica* in the warming OTCs were 11% and 4% higher than those in the control across the whole growing season although the difference was not statistically significant. The soluble sugars of *Rh. confertissimum* in the warming decreased by 17% in July, 34% in August and 18% in September compared to the control, but starch concentration did not change with warming across the

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**Table 1** The average coverage of three shrub (*Dryas octopetala var. asiatica*, *Vaccinium uliginosum* and *Rhododendron confertissimum*) species in the warming open-top chambers and the control plots investigated before and after seven years' warming treatment (unit: %; *n* = 8). Different lowercase letters indicated significant difference in coverage among years. The statistical results of the effects of warming on coverage was shown by *P* value for each species (** *P* < 0.01, *** *P* ≤ 0.001, ns *P* > 0.05).

| Species                      | 2010     | 2015     | 2016     | 2017     | 2015     | 2016     | 2017     | 2015     | 2016     | 2017     |
|------------------------------|----------|----------|----------|----------|----------|----------|----------|----------|----------|----------|
| Dr. octopetala var. asiatica| 38 ± 2.3 | 43 ± 5.8 | 33 ± 5.8 | 32 ± 6.2 | 31 ± 3.7 | 31 ± 3.6 | 20 ± 2.1 | 24 ± 1.8 | 9 ± 0.8  | 6 ± 0.9  |
| V. uliginosum                | 29 ± 3.1 | 30 ± 2.8 | 33 ± 2.7 | 33 ± 4.1 | 29 ± 3.2 | 27 ± 3.6 | 29 ± 3.2 | 27 ± 3.6 | 6 ± 0.9  | 12 ± 2.1 |
| Rh. confertissimum           | 9 ± 0.8  | 6 ± 0.9  | <5 b     | <5 b     | 15 ± 2.0 | 19 ± 2.4 | 14 ± 2.4 | 12 ± 2.1 | ***      | ***      |

**Table 2** The statistical results of the effects of treatment (warming and control), sampling dates (July, August, September) and species (*Dryas octopetala var. asiatica*, *Rhododendron confertissimum* and *Vaccinium uliginosum*) on the soluble sugars, starch and total nonstructural carbohydrate (TNC) (sugar + starch) concentrations, the ratio of soluble sugars to starch, and secondary compounds (flavonoids, total phenols, triterpene) concentrations in leaves using three-way ANOVA.

|                          | Sugar | Starch | TNC  | Sugar/Starch | Flavonoids | Total phenols | Triterpene | F   | P  | F   | P  | F   | P  | F   | P  | F   | P  |
|--------------------------|-------|--------|------|-------------|------------|--------------|------------|-----|----|-----|----|-----|----|-----|----|-----|----|
| Treatment (T)            |       |        |      |             |            |              |            | 2.448| ns | 0.164| ns | 1.982| ns | 0.851| ns | 1.790| ns | 229.742| *** | 0.021| ns |
| Sampling dates (D)       | 38.712| ***    | 8.798| ***        | 6.042      | ***          | 38.590     | *** | 1.070| ns | 10.723| *** | 72.249| *** | 359.343| *** | 775.940| *** |
| Species (S)              | 43.192| ***    | 7.976| ***        | 39.783     | ***          | 15.228     | *** | 22.136| *** | 15.228| *** | 22.136| *** | 359.343| *** | 775.940| *** |
| T x D                    | 0.399 | ns     | 0.644| ns          | 0.916      | ns           | 0.055      | ns  | 28.124| *** | 250.941| *** | 171.625| *** | 162.755| *** | 73.310| *** |
| T x S                    | 0.553 | ns     | 1.877| ns          | 1.465      | ns           | 2.046      | ns  | 3.416| ns  | 16.275| *** | 73.310| *** | 162.755| *** | 73.310| *** |
| D x S                    | 1.345 | ns     | 7.691| ***        | 3.278      | ns           | 5.164      | **  | 0.724| ns  | 266.306| *** | 180.204| *** | 266.306| *** | 180.204| *** |
| T x D x S                | 1.171 | ns     | 0.658| ns          | 0.741      | ns           | 0.758      | ns  | 5.466| **  | 202.599| *** | 75.168| *** | 202.599| *** | 75.168| *** |

Notes: Significance ** *P* < 0.01, *** *P* ≤ 0.001, ns *P* > 0.05
whole growing season (Fig. 1). We observed that warming decreased starch concentration of *V. uliginosum* leaves by 14% in July, 12% in August and 21% in September, on average 15% decline across the whole growing season ($P = 0.003$). Therefore, the ratio of sugars to starch of *V. uliginosum* leaves in the warming was higher than that in the control, but the difference was not significant (Fig. 1).

Soluble sugars, starch and TNC in leaves of the three species showed a pronounced seasonal variation for both the warming OTCs and the controls. Soluble sugars and the ratio of sugars to starch increased with time, reaching the maximum values for the three species in September when the temperature was relatively lower (Fig. 1). The starch concentrations of *Rh. confertissimum* in the warming OTCs were relatively stable during the whole growing season (Fig. 1). We observed that warming decreased starch concentration of *V. uliginosum* leaves by 14% in July, 12% in August and 21% in September, on average 15% decline across the whole growing season ($P = 0.003$). Therefore, the ratio of sugars to starch of *V. uliginosum* leaves in the warming was higher than that in the control, but the difference was not significant (Fig. 1).
growing season (Fig. 1). The seasonal trend of starch in Dr. octopetala var. asiatica and V. uliginosum showed an opposite trend of soluble sugars, and therefore, the TNC was relatively stable with time. However, the TNC concentrations of Rh. confertissimum leaves increased with time, reflecting the seasonal pattern of soluble sugars (Fig. 1).

The concentrations of soluble sugars and TNC did not differ between Dr. octopetala var. asiatica and V. uliginosum (P > 0.05) but significantly higher than Rh. confertissimum (P = 0.007). The soluble sugars concentrations of Dr. octopetala var. asiatica and V. uliginosum were approx. 1.9 times that of Rh. confertissimum. There was no significant difference in starch concentration among the three species, ranging from 101.2 to 136.9 mg g⁻¹ when all data were pooled. The TNC concentration had a similar trend to soluble sugars (Fig. 1). The TNC concentration of Dr. octopetala var. asiatica and V. uliginosum was approx. 1.5 times that of Rh. confertissimum.

**Carbon-based secondary compounds in leaves**

Warming significantly affected the concentrations of total phenols (P < 0.001) and flavonoids of Rh. confertissimum (P = 0.026). Warming decreased the total phenols concentrations by 16% for Rh. confertissimum, 21% for Dr. octopetala var. asiatica and 7% for V. uliginosum across the growing season. On average, Rh. confertissimum in the warming OTCs had 14% higher flavonoids concentration than the controls across the whole growing season.

Total phenols and triterpenes showed pronounced seasonal fluctuation, but different species had different patterns (Fig. 2). Total phenols and triterpenes of Dr. octopetala var. asiatica grown in the warming OTCs decreased with time, but the controls had the highest...
values in August. *V. uliginosum* had relatively higher total phenols and triterpenes in September. The flavonoids were stable during the growing season for the three species.

There were significant differences in flavonoids, total phenols and triterpenes concentrations among the three species (Table 2). *Rh. confertissimum* had the lowest concentrations of flavonoids, total phenols and triterpenes compared to *Dr. octopetala var. asiatica* and *V. uliginosum* (Fig. 2). The total phenols concentration of *V. uliginosum* were 1.4 times higher than *Dr. octopetala var. asiatica* and *Rh. confertissimum* in the warming and approx. 1.2 times higher in the controls. *V. uliginosum* had 1.4 and 2.4 times higher triterpenes concentration than *Dr. octopetala var. asiatica* and *Rh. confertis - simum* across the whole growing season, with the highest difference mainly occurring in September (Table 2, Fig. 2).

**Nonstructural carbohydrates in roots**

No significant effects of warming on soluble sugars, starch and TNC concentrations and the ratio of sugars to starch in roots were observed for the three species, but there were significant differences among species (Fig. 3). *V. uliginosum* roots had 24% higher starch concentration and 12% higher TNC concentration compared to *Dr. octopetala var. asiatica* and *Rh. confertissimum*. The TNC concentration in *Dr. octopetala var. asiatica* roots was very close to *Rh. confertissimum* (Fig. 3).

**Discussion**

**Warming effects on carbohydrates and their possible effects on coverage**

Experimental warming in tundra regions often causes increased photosynthesis and growth rate [23–25]. In line with our hypothesis 1, *Dr. octopetala var. asiatica* positively responded to OTC-warming in tissue NSCs, plant coverage, photosynthesis and single leaf size [25], indicating that warming makes *Dr. octopetala var. asiatica* to fix more carbon, to grow fast, and then to occupy more space and have stronger competitiveness. The soluble sugars of *Rh. confertissimum* negatively responded to warming (Fig. 1), and decreases in the coverage of *Rh. confertissimum* have already been recorded during the experimental period (Table 1), consistent with the hypothesis 1. The coverage of *V. uliginosum* grown in the warming OTCs and the control plots did not significantly change during the whole experimental period (Table 1). Jin et al. (2019) found that *V. uliginosum* on the west-facing slope of Changbai Mountain alpine tundra showed a patch distribution deviating from the previously normal distribution [7]. Our research field was located on the north-facing
slope whose microenvironment (e.g. lower temperature) should be different from the west-facing slope (higher temperature) of Changbai Mountain alpine tundra. The starch storage of *V. uliginosum* responded negatively to the warming (Fig. 1), so we predict that the distribution of *V. uliginosum* on the north-facing slope might decrease in the future or gradually show a patch distribution like on the west-facing slope.

The responses of carbohydrates to warming in tundra shrubs were found to be species-specific in the present study which is consistent with our hypothesis 2 (Table 2). Similar results were also observed by other studies [26, 27]. Warming increased NSCs concentrations in *Himan-toria lugubris* but decreased them in *Polytrichastrum alpinum*, *Pinus sylvestris*, *Pseudotsuga menziesii* and *Picea mariana* [27–30]. No significant change in TNC was observed in *Carex vaginata*, *Sausurea alpine*, *Selaginella selaginoides*, *V. uligonosum*, *Usnea antarctica*, *U. aurantiaco-ata*, *Sanionia uncinata*, *Quercus robur* and *P. petraea* leaves in response to warming [17, 24, 26, 27]. Different species have different sensitivity to warming which can affect photosynthesis, the distribution and utilization of photosynthesize, and growth etc.

Stored carbohydrates in roots can be used by plants for defense and regrowth, or as a buffer under insufficient carbon production [31–35]. Warming did not significantly affect root carbohydrate concentrations of the three species, which is consistent with previous findings that there were non-significant responses of root carbohydrates at the end of the growing season to warming for alpine plants (*Elymus nutans*, *Euphrasia regelii* and *Swertia mussotii*) on the Tibetan Plateau [10], for grapevines in Barossa Valley of Australia [33], for *Pinus taeda* and *P. ponderosa* [36]. *V. uliginosum* roots had higher starch and TNC concentrations compared to *Dr. octopetala* var. *asiatica* and *Rh. confertissimum*, which might be related to that *V. uliginosum* is a deciduous species. Deciduous plants need more energy and nutrition for new leaves sprouting in the next spring, especially on alpine tundra regions.

The accumulation of NSCs is one of the cryoprotective mechanisms [37]. The sugar-starch system in plants adjusts the ratio of sugar to starch in response to low temperature or other stressors [38]. At high elevations, a higher sugar-starch ratio reflects that plants are subjected to lower temperatures, sometimes positively correlating with cold stress [39]. We found that the ratios of soluble sugars to starch in the warming OTCs were lower than those in the control plots for all the three species, indicating that warming can affect sugar-starch relationship. The three species grown in the warming OTCs hydrolyze less starch against freezing which might be beneficial to growth.

**Time-dependent NSC levels and their possible effects on coverage**

Marked seasonal patterns of NSCs concentrations for the three species were observed, which is in agreement with earlier findings of the seasonal NSCs fluctuation with soluble sugars concentrations reaching a higher level at the end of the growing season in various tree species [40–42]. Soluble sugars, serving as osmotic adjustment and signal substances, play an important role against cold [41, 43]. The starch concentrations peaked in July or August (the active growing season) and decreased in September, indicating that starch hydrolyzes to soluble sugars towards end-season [42]. The temporal variation in the level of NSCs also illustrates that temperature might affect the proportions of carbohydrate component.

Wintergreen *Dr. octopetala* var. *asiatica* and deciduous *V. uliginosum* are typical alpine and arctic dwarf shrubs, especially *Dr. octopetala* var. *asiatica* generally dominating community [44]. Evergreen *Rh. confertissimum* has a small distribution area compared to the two others, only in tundra regions. Generally, deciduous trees require more abundant carbohydrates for vegetative or reproductive growth before the new leaves grow [34, 45]. *V. uliginosum* had similar concentrations of TNC as *Dr. octopetala* var. *asiatica*, but *V. uliginosum* did not show the same obvious advantages in coverage, photosynthesis, leaf size and growth as *Dr. octopetala* var. *asiatica* did [25], indicating that *V. uliginosum* might allocate some carbohydrates to the growth of new organs. The roots of *V. uliginosum* had significantly higher starch and TNC concentrations than *Dr. octopetala* var. *asiatica* and *Rh. confertissimum*. Starch, different from soluble sugars, is inactive and accumulates as a storage compound. *V. uliginosum*, as a deciduous species, is assumed to store high amounts of carbohydrates over harsh winter to support leaf flush in spring [45]. The concentrations of TNC in *V. uliginosum* and *Dr. octopetala* var. *asiatica* were significantly higher than those of *Rh. confertissimum* that had significantly decreased coverage. Therefore, TNC concentrations are related to the species coverage, which is consistent with our hypothesis 1.

The NSCs responses of *Dr. octopetala* var. *asiatica* to warming support the idea that carbon allocation is a key factor for determining dominance. After five years of warming by open-top chambers in the alpine region of southwestern Norway, the carbohydrates storage of *Dr. octopetala* increased [46]. *Dr. octopetala* var. *asiatica* showed increased trend in the carbohydrate concentrations in the present study and significantly increased leaf size [25], indicating that the total carbohydrate contents (leaf biomass or size × concentration) have been stimulated. The stimulation and accumulation of carbohydrates are conducive to dominance, expansion and
improvement of competitiveness, which may further lead to changes in community composition and structure of alpine and tundra ecosystems under future climate change [44].

Responses of secondary compounds and trade-off with NSCs
Deciduous species probably have lower concentrations of secondary compounds compared to evergreen plants [47]. Higher concentrations of secondary compounds in perennial leaves could be a greater need for defending herbivorous predator due to longer life span [48]. However, we found that the deciduous V. uliginosum had the highest absolute concentrations of total secondary compounds compared to the other two species. One of the reasons is probably related to that V. uliginosum produces delicious fruits which might need more secondary compounds to defend animals, especially triterpenes.

We observed that Rh. confertissimum allocated relatively more carbon to defense than V. uliginosum and Dr. octopetala var. asiatica based on the ratio of the sum of secondary compounds to total carbon (secondary compounds + TNC). Herms and Mattson (1992) found that increased investment in secondary defense is accompanied by decreased growth, plant size and competitive ability [16]. Thus, more carbon is allocated to secondary compounds in expense of growth, dominance and competition. Warming may alter interspecific competitive relationships and community structure because of the changes in carbon allocation pattern and defense abilities [46].

Total phenols concentrations were decreased by warming in the present study, similar to the results of Holopainen et al. [49]. Phenolic compounds originate from the shikimic acid pathway which is related to the carbohydrate metabolisms [50, 51] and antioxidative potential of plants [52]. Tundra ecosystem is generally characterized by simultaneous stresses such as low temperature, high UV radiation, low nutrient availability which could make plants to produce high levels of secondary compounds or allocate more proportion carbon to the secondary compounds [46, 53]. Thus, alleviation of low temperature by warming is expected to decrease the contents of secondary compounds. The decreases in the concentrations of secondary compounds for Bistorta vivipara, Dr. octopetala, Salix reticulate, Cassiope tetragona, S. herbacea × Polaris and Tofieldia pusilla have been reported [17, 22, 46]. The decrease in total phenols concentration probably relates to the fewer carbon resources for defense substance or more carbon for growth [17]. The decreased proportion of total phenols of Dr. octopetala var. asiatica was relatively higher than the other two species (Fig. 2) which is consistent with higher NSCs and growth. Hence, Dr. octopetala var. asiatica has strong competitive potential.

Flavonoids concentrations of Rh. confertissimum were increased by warming, but OTC warming did not affect the levels of flavonoids in Dr. octopetala var. asiatica and V. uliginosum. We also found that Rh. confertissimum grown in the warming OTCs had relatively lower soluble sugars and significantly reduced coverage than the controls or the other two species. The three species in the present study changed their defense levels to some extent when experiencing continuous 7 years’ warming, consistent with our hypothesis 2 that there was significant interaction between species and warming on the total phenols and flavonoids.

Inconsistent with our hypothesis 3, no significant trade-off relationship between NSCs and secondary compounds in leaves was observed based on the correlation analysis (Figure not shown). However, the secondary compounds tended to be positively correlated with NSCs for Dr. octopetala var. asiatica while negatively correlated with NSCs for V. uliginosum and Rh. confertissimum. Further research is necessary to continue to examine if long-term warming can result in a trade-off relationship between NSCs and secondary compounds for species with decreased distribution or competitiveness.

Conclusions
We conclude that warming increased NSCs and decreased secondary compound in Dr. octopetala var. asiatica, which makes the species still maintain dominant status in the Changbai Mountain alpine tundra with climate change. The dominance of V. uliginosum could gradually decline with continuous warming. Rh. confertissimum had relatively lower carbohydrates compared to Dr. octopetala var. asiatica and V. uliginosum and increased secondary compounds investment with warming. The coverage of Rh. confertissimum in the OTCs was getting small. All these traits will weaken the competition ability of Rh. confertissimum in tundra community. Thus, we predict that Dr. octopetala var. asiatica will maintain its dominant status, V. uliginosum could gradually decrease its coverage, and Rh. confertissimum might be in danger of disappearance with air warming, leading to changes in species composition and community structure of the Changbai tundra ecosystem under future climate warming.

Materials and Methods
Study site and experimental design
The study was conducted on the north-facing slope of alpine tundra on Changbai Mountain (41°58′ - 42°42′ N; 127°67′ - 128°27′ E, 2046 m a.s.l.), northeastern China with typical characteristics of Arctic tundra [54].
Changbai Mountain alpine tundra experienced significant increases in temperature and precipitation and decreases in frost and icing days based on the data from 1950 to 2010 [2]. The mean annual temperature is -7.3 °C and mean annual precipitation is 1373 mm in this region [55]. The mean air temperature during the growing season (June to September) is 5.9 °C, and the highest mean daily temperature is less than 10 °C [2]. The majority falling as rain occurs during the short summer (July and August). A snow-free season lasts from May to September. Soils are characterized by Haplic Cambisol (Humic, Dystric). The total nitrogen, phosphorus and potassium were 3.6 ± 0.19, 1.00 ± 0.06 and 1.64 ± 0.01 mg/g, respectively, and the soil had an organic matter content of 124 ± 9.2 mg/g. The vegetation mainly consisted of Dr. octopetala var. asiatica, V. uliginosum, Rh. confertissimum, Rh. chrysanthum, Sanguisorba parviflora, S. stipulate, Calamagrostis angustifolia, etc.

Open-top chambers (OTCs) were used to increase air and soil temperature according to the criteria of the International Tundra Experiment [56]. Eight OTCs were established near the viewing platform in June 2010, about half a mile away from road, with a steep slope in the middle. The OTCs were hexagonal, 0.45 m high, had inclined sides (60°), enclosed a surface of 1.0 m², and were made of transparent polycarbonate. The OTCs were left in place year-round. Control plots were established beside each OTC with similar species composition and vegetation coverage. Air and soil temperature, air and soil humidity, radiation in the OTCs and the control plots were logged every 30 minutes during growing seasons (June to September) by Em 50 Data Collection System (Decagon, USA). On average, the OTC increased air and soil temperature at 10 cm depth by 1.5 °C and 0.8 °C (with less increase in air temperature, while decrease in soil temperature during night). The air relative humidity was not significantly changed by warming, but the soil water content was decreased by 0.05 m³ m⁻³ [25].

Three dominant species growing on Changbai Mountain tundra were selected in the present study. Dr. octopetala var. asiatica is an evergreen dwarf shrub and often forms heath communities on calcareous soils in arctic-alpine environments [46]. V. uliginosum is deciduous dwarf shrub with thin and approximate round leaves. Rh. confertissimum is an evergreen dwarf shrub with relatively thicker leaves. The specimens of the three plants can be found in the Institute of Applied Ecology, Chinese Academy of Sciences with voucher ID C.Y. Li 1962, Y.L. Zhou 1951, and P.Y. Fu 1959.

Sampling leaves and roots
Experimental area and sampling for scientific research were approved by local administrative department that supported the present project. We identified the three plant species. After all samples were dried and ground in the field station, they were stored in the laboratory in Shanghai Institute of Technology.

The leaves of Dr. octopetala var. asiatica, V. uliginosum, Rh. confertissimum were sampled from three plants in each OTC and the control plot on July 13th, August 13th and September 18th, 2017. The roots were only sampled on September 18th because of the destructive sampling. Plants were always randomly sampled away from the chamber sides.

Nonstructural carbohydrates analysis
Dried leaves and roots were ground into fine powders using a small grinder. Extraction and determination of carbohydrates from leaves and roots were based on the Anthrone method [57]. The powdered material was suspended in 80% ethanol and incubated for 30 min at 80°C for extraction of soluble sugars. The supernatant was decanted after centrifugation. Then the residual was resuspended in 80% ethanol and repeated the same procedure twice. The supernatant was in constant volume and then was quantified using anthrone as a reagent at 620 nm. The remaining pellet was kept for starch analysis. The pellet was washed two times with 4.6 mol/L HClO₄ to digest starch. The supernatant was quantified using the same method as soluble sugars. TNC were calculated by summing total soluble sugars and starch [58].

Secondary compounds analysis
Due to an insufficient number of root samples, only leaf secondary compounds were measured. Secondary compounds of total phenols, flavonoids and triterpenes were extracted by adding air dried leaf powder to 70% ethanol that was heated to reflux for two hours. The procedure was repeated two times. The filtered extract was condensed with a vacuum evaporator. Total phenols were measured at 760 nm wavelength by the Folin-Ciocalteu method [59]. Flavonoids were determined at 510 nm wavelength with Rutin as the standard solution [60]. Triterpenes were measured at 542 nm wavelength with vanillin and glacial acetic acid as the chromogenic reagent [61].

Data analysis
Twenty-four plants were respectively selected in the warming OTCs and the control for each species. The leaves of Rh. confertissimum were not enough to measure all parameters, so every four plants were mixed into one sample. MANOVA was used to test the significance of main effect factors (warming treatment, species and sampling date) and their interactions on carbohydrates and secondary compounds. Due to the overall significant effects of species on parameters studied, repeated measures ANOVAs
were used to analyze the effects of treatment (between subject), sampling date (within subject), and their interaction on parameters within each species. Multiple comparisons were used to examine the difference in levels of the parameters among the three species. Paired-samples T test was used to assess the effects of warming treatment on every parameter for each species for each sampling date. Data were analyzed statistically using SPSS 16.0 system (SPSS Inc., Chicago, IL, USA). All tests of statistical significance were conducted at a level of 0.05.

Abbreviations
OTCs: Open-top chambers; NSCs: non-structural carbohydrates; TNC: total NSCs.

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Authors’ contributions
YMZ, NY, JJJ finished lab work. ZJT and DTL collected field data and statistically analyzed the data. YMZ wrote the original draft. YMZ and XM revised the manuscript. All authors contributed to the text of the manuscript. All authors have read and approved the manuscript, and ensure that this is the case.

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Availability of data and materials
The datasets used and analyzed during the current study are available from the corresponding author on reasonable request.

Declarations

Ethics approval and consent to participate
Not applicable

Consent for publication
Not applicable

Competing interests
The authors declare that they have no competing interests.

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References
1. Deslippe JR, Simard SW. Below-ground carbon transfer among Betula nana may increase with warming in Arctic tundra. New Phytol. 2011;192:689–98.
2. Zong SW, Wu ZF, Du HB. Study on climate change in alpine tundra of the Changbai Mountain in growing season in recent 52 years. And Zone Res. 2013;30:41–9.
3. Li MH, Krauchi N, Gao XP. Global warming: can existing reserves really preserve current levels of biological diversity? J Integr Plant Biol. 2006;48:255–9.
4. Zong SW, Ju JW, Wu FZ, Qiao LL, Wang DD, Meng XJ, et al. Analysis on the process and impacts of Deynaxia angustifolia invasion on the alpine tundra, Changbai Mountain. Acta Ecol Sin. 2014;34:6837–46.
5. Jin YH, Xu JW, Liu LN, He HS, Tao Y, Zong SW, et al. Spatial distribution pattern and associations of dominant plant species in the alpine tundra of the Changbai Mountains. Sci Geogr Sinica. 2016;36:1212–8.
6. Jin YH, Zhang YJ, Xu JW, Tao Y, He HS, Guo M, et al. Comparative Assessment of Tundra Vegetation Changes Between North and Southwest Slopes of Changbai Mountains, China, in Response to Global Warming. Chin Geogr Sci. 2018;28:665–79.
7. Jin Y, Xu J, He H, Li MH, Tao Y, Zhang Y, et al. The Changbai alpine shrub tundra will be replaced by herbaceous tundra under global climate change. Plants. 2019;8:370.
8. Arft AM, Walker MD, Gurevitch J, Alatalo JM, Breit-Harte MS, Dale M, et al. Responses of tundra plants to experimental warming: meta-analysis of the International Tundra Experiment. Ecol Monogr. 1999;69:491–511.
9. Root TL, Price JT, Hall KR, Schneider SH, Rosenzweig C, Pounds JA. Fingerprints of global warming on wild animals and plants. Nature. 2003;421:57–60.
10. Shi C, Sun G, Zhang H, Xiao B, Ze B, Zhang N, et al. Effects of warming on chlorophyll degradation and carbohydrate accumulation of alpine herbaceous species during plant senescence on the Tibetan Plateau. PlosOne. 2014;9:e107874.
11. Gargallo-Garriga A, Sardans J, Pérez-Trujillo M, Oravec M, Urban O, Jentsch A, et al. Warming differentially influences the effects of drought on stoichiometry and metabolomics in shoots and roots. New Phytol. 2015;207:591–603.
12. Aspinwall MJ, Drake JE, Canpany C, Våhrammar A, Ghannoum O, Tissue DT, et al. Convergent acclimation of leaf photosynthesis and respiration to prevailing ambient temperatures under current and warmer climates in Eucalyptus tereticornis. New Phytol. 2016;204:354–67.
13. Klanderud K. Species-specific responses of an alpine plant community under simulated environmental change. J Veg Sci. 2008;19:363–72.
14. Donaldson JR, Kruger EL, Lindroth RL. Competition- and resource-mediated tradeoffs between growth and defensive chemistry in trembling aspen (Populus tremuloides). New Phytol. 2006;169:561–70.
15. Guo Q, Li J, Zhang Y, Zhang J, Lu D, Karpelainen H, et al. Species-specific competition and N fertilization regulate non-structural carbohydrate contents in two Larix species. For Ecol Manage. 2016;364:60–9.
16. Herms DA, Mattson WJ. The dilemma of plants: to grow or to defend. Quart Rev Biol. 1992;67:283–335.
17. Nybakken L, Sandvik SM, Klanderud K. Experimental warming had little effect on carbon-based secondary compounds, carbon and nitrogen in selected alpine plants and lichens. Environ Exp Bot. 2011;72:368–76.
18. Stamp N, Bradfield M, Li S, Alexander B. Effect of competition on plant allometry and defense. Am Midl Nat. 2004;151:50–64.
19. Aerts R, Callaghan TV, Dorrepaal E, Van Logtestijn RSP, Cornelissen JHC. Seasonal climate manipulations result in species-specific changes in leaf nutrient levels and isotopic composition in a sub-arctic bog. Funct Ecol. 2009;23:680–8.
20. Hoch G, Körner C. Global patterns of mobile carbon stores in trees at the high-elevation tree line. Globo Ecol Biogeogr. 2011;21:861–71.
21. Streit K, Rinne KT, Hagedorn F, Dawes MA, Saurer M, Hoch G, et al. Tracing fresh assimilates through Larix decidua exposed to elevated CO₂ and soil warming at the alpine treeline using compound-specific stable isotope analysis. New Phytol. 2013;197:838–49.
22. Hanssen AH, Jonasson S, Michelsen A, Jalkunen-Titto R. Long-term experimental warming, shading and nutrient addition affect the concentration of phenolic compounds in arctic-alpine deciduous and evergreen dwarf shrubs. Oecologia. 2006;147:1–11.
23. Wooley PA, Robinson CH, Parsons AN, Welker JM, Press MC. Environmental constraints on the growth and performance of Dryas octopetala ssp. ohtachae at a High Arctic polar semi-desert. Oecologia. 1995;104:567–78.
24. Dornemann CF. Consequences of manipulations in carbon and nitrogen supply for concentration of anti-herbivore defence compounds in Salix polaris. Ecoscience. 2003;10:312–8.
comparison of deciduous and semideciduous arboreal legumes. Sci Agric. 2006;63:40–5.
48. Aerts R. Nutrient use efficiency in evergreen and deciduous species from heatlands. Oecologia. 1999;100:391–7.
49. Holopainen JK, Virgoano V, Ghimire RP, Blande JD, Jülkenen-Titro R, Kivimäenpää M. Climate change effects on secondary compounds of forest trees in the northern hemisphere. Front Plant Sci. 2018;9:1445.
50. Seigler DS. Plant Secondary Metabolism. Dordrecht: Kluwer Academic Publishers, 1998.
51. Lindroth R. Atmospheric change, plant secondary metabolites, and ecological interactions. In: The Ecology of Plant Secondary Metabolites: From Genes to Global Processes, Iason GR, Dicke M, Hartley SE ed; 2012. p. 120–53.
52. Someya S, Yoshioka Y, Okubo K. Antioxidant compounds from bananas (Musa Cavendish). Food Chem. 2002;79:351–4.
53. Thoss V, Shevtsova A, Nilsson M-C. Environmental manipulation treatment effects on the reactivity of water-soluble phenolics in a subalpine tundra ecosystem. Plant and Soil. 2004;259:355–65.
54. Qian H. Numerical classification and ordination of plant communities in Mt Changbai. J Appl Ecol. 1990;1:254–63.
55. Liu QJ, Zhang GC, Xu QQ, Wang YD, Wang HM. Simulation of soil respiration in response to temperature under snowpacks in the Changbai Mountain, China. Chin J Plant Ecol. 2010;34:477–87.
56. Henry HGR, Molau U. Tundra Plants and Climate Change: the International Tundra Experiment (ITEX). Glob Chang Biol. 1997;3(Suppl 1):1–9.
57. Seifert S, Dayton S, Novic B, Muntwyler E. The estimation of glycosylation with the anthrone reagent. Arch Biochem. 1950;25:191–200.
58. Li MH, Hoch G, Körner C. Source/sink removal affects mobile carbohydrates in Pinus cembra at the Swiss treeline. Trees Struct Funct. 2002;16:331–7.
59. Singleton VL, Orthofer R, Lamuela-Raventós R. Analysis of total phenols and other oxidation substrates and antioxidants by means of Folin-Ciocalteu reagent. Methods Enzymol. 1999;299:152–78.
60. Wang ZZ, Lin SY, Liu JB, Wang EL, Zhang W. Study on structure identification of flavonoids in Vaccinium uliginosum L. Food Sci. 2007;28:455–7.
61. Zong W, Xia WS, Cui BL. Determination of total triterpenes in Lagerstroemia speciosa L. by thin layer chromatography-spectrophotometry. Food Sci. 2005;26:222–5.

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