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WINTER SNOW DEPTH IN ARCTIC ALASKA RESULTS IN COMPLEX CHANGES

IN CARIBOU FORAGE QUALITY

BY JESSICA C. RICHERT

A thesis submitted in partial fulfillment of the requirements for the Master of Science Major in Biological Sciences South Dakota State University

2019

THESIS ACCEPTANCE PAGE

Jessica C. Richert

This thesis is approved as a creditable and independent investigation by a candidate for the master's degree and is acceptable for meeting the thesis requirements for this degree. Acceptance of this does not imply that the conclusions reached by the candidate are necessarily the conclusions of the major department.

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TABLE OF CONTENTS

ABSTRACT	V
CHAPTER 1: INTRODUCTION	1
CHAPTER 2: EFFECTS OF SNOW ON FORAGE QUALITY OF TUSSOCK TUNDRA	7
Introduction	7
Materials and Methods	9
Study Site	9
Field Sampling	10
Laboratory Analyses	12
Statistical Analyses	14
Results	15
Snow and Active Layer Depth	15
Forage Availability	16
Nitrogen Content	17
Dry Matter Digestibility	18
Protein-precipitating Capacity and Digestible Protein	19
Discussion	20
Effects of Snow Depth on Forage Availability	21
Effects of Snow Depth on Forage Quality	22
Limitations	27
Conclusions	28
TABLES	29
FIGURES	32
CHAPTER 3: CONCLUSIONS	38
LITERATURE CITED	40
APPENDIX	55

WINTER SNOW DEPTH IN ARCTIC ALASKA RESULTS IN COMPLEX CHANGES IN CARIBOU FORAGE QUALITY

ABSTRACT

JESSICA C. RICHERT

2019

Caribou (*Rangifer tarandus*) rely on the short growing season for much of their annual nutrition, making them susceptible to even small changes in forage quantity and quality. Body condition in the summer and fall is linked to winter survival rates and fecundity in cows, critical factors in the robustness of caribou populations. Due to a warmer, wetter climate, snowfall is predicted to increase over Alaska's North Slope in the next several decades. Deeper snow results in higher soil temperatures, allowing microbial mineralization of nitrogen to continue throughout the winter and increasing the availability of nitrogen for plants in spring and summer; however, deeper snow can also delay the onset of spring and initial plant growth. These biophysical changes may impact the quantity, quality, and seasonality of caribou forage. I used a 20+ year snow manipulation to evaluate how a set of winter climate change scenarios may affect tussock tundra vegetation community composition and forage quality in northern Alaska. I sampled leaf tissue of six plant species (Salix pulchra, Betula nana, Rhododendron tomentosum, Vaccinium vitis-idaea, Carex bigelowii, and Eriophorum vaginatum) weekly between leaf-out and senescence in two consecutive years in areas of ambient, reduced, and added snow. Leaf tissue was analyzed for %N, dry matter digestibility, and digestible protein to quantify temporal changes in nutrition as well as differences between species and among functional groups (deciduous shrubs, evergreen dwarf shrubs, and

graminoids). Deeper snow increased leaf %N and digestible protein in the two deciduous shrubs and graminoids, but not the evergreen shrubs. Dry matter digestibility varied between species with small differences associated with divergent winter snow depths. Deeper snow also increased the duration of higher-protein forage by as much as 25 days in *S. pulchra* and 6-9 days in *B. nana* and *C. bigelowii*. Consequently, predicted increases in winter snow over the North Slope by the end of the century may enhance both summer and autumn forage quality and availability for caribou. Through multiplier effects of increased nutrition on body condition, survivorship, and fecundity, better forage conditions may improve the health and welfare of caribou in northern Alaska.

CHAPTER 1: INTRODUCTION

Caribou and reindeer (*Rangifer tarandus*) populations in the circumpolar Arctic face many challenges (Osborne et al. 2018). Forage availability, forage quality, predation, insect harassment, increasing human development, and extreme weather events have all been implicated as possible reasons for the declines of caribou and reindeer populations around the world in recent decades (Mörschel and Klein 1997, Vors and Boyce 2009, Festa-Bianchet et al. 2011, Fauchald et al. 2017). Though caribou and reindeer are highly susceptible to abiotic stochastic effects (Jefferies et al. 1994, Tyler 2010, Hansen et al. 2014) and populations are prone to decadal fluctuations in size (Gunn 2003), the near-synchronous decline of global populations is cause for concern and suggests a widespread driver like climate change may be partially responsible.

Changes in the Arctic's weather patterns and overall climate present additional challenges to caribou and reindeer populations. Over the past 60 years, the mean annual temperature has risen almost twice as fast in the Arctic as the rest of the world (ACIA 2004), with temperatures already rising by almost 2°C since the early 1900s (Osborne et al. 2018). Temperatures are predicted to continue rising by another 2-9°C by the end of the century (IPCC 2013). This increase in air temperature may worsen already existing stressors in caribou and reindeer as well as introduce new ones (Mallory and Boyce 2017). Caribou begin exhibiting signs of heat stress at temperatures above 22°C (Thompson and Barboza 2014), decreasing forage intake and reducing activity (Mörschel and Klein 1997). Insect harassment from mosquitoes and flies also increases with temperature (Mörschel and Klein 1997, Bali et al. 2013) with severe harassment affecting forage intake and large-scale movement patterns observed in air temperatures over

13.5°C (White et al. 1975). Higher temperatures also increase the frequency and risk of extreme weather events like rain on snow and icing of pastures (Rennert et al. 2009), which can lead to starvation and mortality as animals are cut off from important winter forage (Hansen et al. 2014, Mallory and Boyce 2017), though significant die-offs during such extreme events may also be due in part to density-dependent factors, that is the density of caribou or reindeer exceeds the necessary forage resources to support the herd (Tyler 2010, Hansen et al. 2019).

Warmer weather during the growing season may also decrease forage quality and contribute to phenological mismatches between migratory caribou and their food sources at a nutritionally critical part of year (Walsh et al. 1997, Fauchald et al. 2017). Warming reduces leaf-level nitrogen (N) concentrations and increases anti-herbivory compounds such as digestibility-reducing tannins (Jonasson et al. 1986, Turunen et al. 2009, Zamin et al. 2017a) while simultaneously promoting growth of deciduous shrubs like *Alnus* and *Betula spp.* These shrubs are naturally higher in anti-herbivory compounds than graminoids, potentially decreasing forage quality even as forage availability increases (Fauchald et al. 2017). Warmer temperatures may also shift the growing season earlier into the spring, potentially decoupling annual caribou physiological stages from both the timing and seasonality of critical nutrients on the landscape. While spring forage quality and availability may not be as important to capital breeders like caribou that rely more on winter body reserves for calf production and survival (Veiberg et al. 2017), phenological mismatches in the summer and fall have the potential to severely impact both individuals and populations due to the influence of even small changes in forage intake and weight

gain on growth rates and fecundity (White 1983, Cebrian et al. 2008, Proffitt et al. 2016, Gustine et al. 2017).

Effects of higher temperatures on caribou may not always be negative, however. Warmer air temperatures cause warmer springs, earlier snowmelt, and consequently, longer growing seasons (Linderholm 2006). Plant communities are already responding to these changes. Preferred summer forages like deciduous shrubs and graminoids (White et al. 1975, Thompson and McCourt 1981, Denryter et al. 2017) are thriving at the expense of less palatable evergreen shrubs and nonvascular plants (Sturm et al. 2001b, Wahren et al. 2005, Tape et al. 2006, Hobbie et al. 2017, Carlson et al. 2018), although important winter forage like lichens is also declining (Wahren et al. 2005, Hobbie et al. 2017). Asynchronous green-up of forage due to differences in snow melt-off dates between areas of shallow and deep snow in response to warmer springs may also benefit caribou by creating spatial heterogeneity of high-quality forage across the landscape and extending the length of time that such forage is available (Searle et al. 2015, Veiberg et al. 2017).

Winter precipitation patterns over the Arctic are also expected to shift along with higher air temperatures. Projections for the central Arctic region of Alaska call for an increase in winter precipitation of 13-48% by mid-century and an increase of 36-77% by the end of the century, mostly in the form of snow (Martin et al. 2009), though predictions for the entire Arctic are variable (Callaghan et al. 2011). As snow is a defining feature of Arctic ecosystems for up to nine months of the year, changes in snow cover and duration may have a greater effect on northern plant communities than a warmer growing season (Rieley et al. 1995, Jones et al. 1998, Wahren et al. 2005, Fu et al. 2014). Snow cover plays a vital role in insulating the ground from harsh winter

conditions, with even moderate increases in snow depth raising the soil surface temperature by as much as 15°C (Walker et al. 1999, Schimel et al. 2004, Pattison and Welker 2014). Higher soil temperatures allow microbial mineralization of soil nitrogen to continue throughout the winter and increases active layer thaw depths in the summer (Johansson et al. 2013) while also increasing available nitrogen pools in the soil that plants can utilize (Schimel et al. 2004, Welker et al. 2005).

Changes in winter precipitation also alter the timing of snow melt and subsequent green-up, potentially shortening or lengthening the growing season in areas. Even though the Alaskan Arctic is predicted to have more winter precipitation, the overall duration of snow cover is expected to decrease (Callaghan et al. 2011). The timing of melt-off is critical to the onset of new growth in plants as photosynthesis begins and soils thaw enough for nutrient uptake (Walsh et al. 1997, Borner et al. 2008). Deep snow accumulations may delay green-up by as much as 3-4 weeks, reducing an already short growing season and potentially reducing both productivity and overall biomass (Wipf and Rixen 2010), though this may be mitigated in some cases by a corresponding increase in photosynthesis from higher leaf N concentrations (Leffler and Welker 2013, Bosiö et al. 2014).

Moderate increases in snow depth also favor deciduous shrub growth (Wahren et al. 2005, Tape et al. 2006, Berner et al. 2018). Taller shrubs trap snow around themselves, creating deeper drifts in winter and forming a positive feedback loop wherein climate-induced increases in snowfall increases shrub growth which in turn further increases local snow depth (Sturm et al. 2001a, 2005, Jespersen et al. 2018). This loop eventually shifts vegetation communities from graminoid and ericaceous shrubdominated to those dominated by deciduous *Alnus*, *Betula*, and *Salix* species (Tape et al. 2006, 2012).

The highest nutritional demands on caribou and reindeer occur during the summer and autumn when animals are recovering from winter deprivations, cows are lactating, and calves are growing rapidly (Denryter et al. 2017, Gustine et al. 2017, Veiberg et al. 2017). Plants with higher leaf-level N concentrations, like deciduous shrubs, provide more protein to caribou and are preferentially selected during foraging (White and Trudell 1980, Denryter et al. 2017), especially early in the season when leaf N concentration is maximal (Klein 1990).

Both deciduous shrub biomass and leaf-level N increase with snow depth (Walker et al. 1999, Welker et al. 2005, Borner et al. 2008, Leffler and Welker 2013), potentially mitigating any dilution of nitrogen due to increased growth and providing caribou with an abundance of high-protein forage (Turunen et al. 2009, Zamin et al. 2017a). In addition, higher soil nitrogen can decrease the carbon-based secondary compounds, including condensed tannins and other phenolics, commonly found in arctic shrubs (De Long et al. 2016). Tannins and other phenolic compounds reduce the digestibility and available protein of plants either through binding directly to proteins in forage or by interfering with digestive enzymes in an animal's stomach (Robbins et al. 1987b, 1987a, Lambers et al. 2008). Fertilization treatments have decreased total phenolic content at multiple arctic and alpine sites (Bryant et al. 1983, Coley et al. 1985, Graglia et al. 2001, De Long et al. 2016), though effects of snow and fertilization on the actual protein-precipitating capacity of plants is less known. This study builds on decades of research from the same site, providing a continued examination of long-term trends in leaf-level nutrients after 25 years of snow manipulation as well as short-term temporal changes during the growing season. While multiple studies have examined the effects of warming, snow, and higher CO₂ on vegetation composition and leaf-level nutrients, few have specifically examined how such responses relate to caribou nutrition. This study seeks to fill in that gap and determine what effects changes in winter snow depth associated with projected climate change have on the availability and quality of forage for caribou on Alaska's North Slope.

CHAPTER 2: WINTER SNOW DEPTH IN ARCTIC ALASKA RESULTS IN COMPLEX CHANGES IN CARIBOU FORAGE QUALITY

Introduction

Due to Arctic amplification, northern latitudes are warming nearly twice as fast as the rest of the world (Martin et al. 2009), and temperatures are projected to continue rising throughout this century (IPCC 2013). Precipitation patterns are also shifting over much of the Arctic, with significant increases in winter precipitation expected over the next few decades (though model projections are variable), particularly in the form of increased snow fall (Callaghan et al. 2011). In particular, Alaska's North Slope, home to the Central Arctic Herd of caribou (*Rangifer tarandus granti*), is predicted to see a 35-70% increase in winter precipitation over portions of the herd's home range by the end of the century (Scenarios Network for Alaska and Arctic Planning 2011).

Snow cover is a defining feature of Arctic ecosystems for up to nine months of the year and changes in extent, depth, and duration may have a greater effect on northern plant communities than warming growing season temperatures (Wahren et al. 2005, Fu et al. 2014). Snow cover plays a vital role in insulating the soil from harsh winter conditions. Deeper snow results in higher soil temperatures in winter (Walker et al. 1999, Schimel et al. 2004) because snow decouples the soil from the frigid arctic air. These less cold soils lead to a deeper active layer depth in summer (Johansson et al. 2013, Pattison and Welker 2014) and facilitate microbial mineralization of organic nitrogen throughout the entire winter, increasing the soil nitrogen available for plants in the early spring (Bilbrough et al. 2000, Schimel et al. 2003, 2004, Sturm et al. 2005, Welker et al. 2005) and thus, higher leaf N all summer long. Secondary compounds are a critical attribute of tundra plants as related to herbivory and digestibility. These compounds, including tannins and other similar phenolic anti-herbivory compounds that reduce digestible protein of plants may change in response to environmental conditions (Peñuelas et al. 1997, Nybakken et al. 2013). For instance, tannin content of leaves may decrease with higher soil nutrients in Arctic and alpine sites (Bryant et al. 1983, Coley et al. 1985, Graglia et al. 2001, Schimel et al. 2004, De Long et al. 2016). There are few experimental studies that quantify whether warmer summers or changes in winter snow affect secondary compounds in tundra plants, however, an attribute that may be critical to determine as we seek to understand how climate changes will affect forage nutrition for caribou in Alaska and globally.

One of the most important facets of understanding how tundra systems will adapt as weather and climate changes is woven into the individualistic nature of plant species and functional group responses to change (Chapin III and Shaver 1985). This foundation of tundra ecology is still apparent today, as given uniform changes in environmental conditions, not all species and not all traits (i.e. flowering, leaf out, leaf-level nutrition, leaf physiology, etc) behave in a uniform manner (Arft et al. 1999). Recent changes across the Arctic are a mixture of responses with potential community and ecosystem consequences such as a strong growth response of some shrub species to warmer summers and/or deeper snow (Elmendorf et al. 2011, Tape et al. 2012). This perspective is especially important as related to caribou forage as individualistic changes in abundance in combination with changes in both magnitude and duration of higher-quality forage in spring or autumn may have large consequences for the carrying capacity of the landscape for caribou as well as the ability to support early season or prolonged nutrition during critical physiological stages of this keystone ungulate.

The primary question that this study addresses, then, is: how do various winter climate scenarios (deeper or shallower snow) affect the availability and quality of forage in tussock tundra for caribou? To address forage availability, I asked: does snow depth affect vegetation community composition and biomass? To address forage quality, I asked: does snow depth affect nutritional factors like digestibility and protein content in common tussock tundra plan species. Based on previous snow manipulation, warming, and fertilization experiments that demonstrates the effect of snow on vegetation community composition and nitrogen content, I hypothesized that: (1) enhanced growth due to warmer soil temperatures and enhanced microbial activity under deep snow would benefit deciduous shrubs over other functional groups due to deeper roots and higher phenological plasticity, leading to higher biomass; and (2) due to increased availability of nitrogen in warmer winter soils and a decrease in phenolic content in plants with experimental fertilization, measures of caribou forage quality like leaf N, digestibility, and digestible protein would be highest in areas of deeper snow, with the greatest impact on deciduous shrubs.

Materials and Methods

Study Site

This research was conducted from 2017 to 2018 in moist-acidic tussock tundra near Toolik Field Station (68°38' N 149°38W) in the foothills of the Brooks Range, Alaska, USA. The mean annual temperature is -8°C, with mean summer temperatures of Environmental Data Center Team 2019). Mean precipitation is 250-350 mm, with 40-45% falling as snow (Schimel et al. 2004). In winter, snow depths reach 30-80 cm on average, but can drift much deeper in response to winds and topography. The ground freezes to the depth of permafrost during winter, with maximum active layer depths in the

10-12°C and winter temperatures averaging -20°C (Hobbie and Kling 2014;

summer averaging 30-50 cm (Jones et al. 1998; Hobbie and Kling 2014). The site is located within the home range of the Central Arctic Herd of caribou (*Rangifer tarandus granti*) on Alaska's North Slope and is dominated by the tussock-forming sedge, *Eriophorum vaginatum*, with deciduous shrubs, evergreen dwarf shrubs, mosses, lichens, and other non-tussock forming graminoids intermixed throughout the inter-tussock areas (Whalen 2002). A wooden snow fence (3 x 60 m) was erected on the moist-acidic tussock tundra site in 1994 to artificially increase snow depth, with snow drifts reaching a maximum depth of 3m directly behind the snow fence and declining to ambient snow depths (0.5-1 m) 50-60 m from the fence (Jones et al. 1998, Walker et al. 1999, Welker et al. 2000).

Field Sampling

To test whether winter snow depth affects summer forage quality, I collected samples of six common plant species present in moist-acidic tussock tundra. These species represented three functional groups: deciduous shrubs, evergreen dwarf shrubs, and sedges. Samples of two species from each functional group were collected on a weekly basis from 19 June-7 August 2017 and 22 June-25 August 2018. The species collected were the deciduous shrubs *Salix pulchra* and *Betula nana*, the evergreen dwarf shrubs *Rhododendron tomentosum* (formerly *Ledum palustre*) and *Vaccinium vitis-idaea*,

and the sedges *Carex bigelowii* and *Eriophorum vaginatum*. Sampling methods mimicked caribou browsing, with leaves of deciduous shrubs stripped by hand and evergreen dwarf shrubs and sedges clipped at ground level. Five leaf tissue samples of each species were collected in each of three snow depth zones: + snow (1-2 m snow), snow (0.5 m snow), and ambient (0.5-1 m snow; control), for a total of 90 samples a week over a 10-12 week period throughout the growing season. After collection, all samples were dried in a forced-air oven at 70°C for 72 hours. These samples were analyzed for C and N content and dry matter digestibility. An additional five samples of *S. pulchra* and *B. nana* were collected every other week during the 2018 growing season and immediately frozen for later chemical analysis of protein-precipitating capacity (PPC), a measure of the reduction in protein digestibility in forages due to anti-herbivory compounds (Robbins et al. 1987a).

To test whether snow depth affects forage availability by altering plant community composition and abundance, I sampled biomass at three locations within each snow depth zone at peak growing season biomass of both years (13 July and 29 July, respectively) for a total of six plots per snow zone. Biomass for *S. pulchra* and *B. nana* was harvested in 1 m² quadrats, while biomass of all other species was harvested in three 20 cm² quadrats nested within the larger 1 m² plot. All samples were sorted to species, then dried in a forced-air oven at 70°C for 72 hours before being weighed to obtain the total biomass per species. Species were combined into functional groups for analysis.

In addition to forage and biomass samples, I measured snow depth during the winter of 2018-2019 and active layer depth during the summer of 2018. I measured snow depth along 12 transects perpendicular to the snow fence. Snow depths were recorded at

2-5 m intervals from the snow fence to 80 m north of the fence using a combination of a MagnaProbe (Snow-Hydro, Fairbanks, Alaska, USA) in snow to 1-m depth and an avalanche probe in areas of deeper snow. I also recorded active layer depth on a weekly basis during the 2018 growing season by inserting a probe to the freeze boundary at 5 m intervals along a 50 m transect in each snow zone. Due to late snowmelt, the + snow treatment zone was separated into two active layer depth transects, one along the shallower end of the snow drift and one along the deeper end. Measurements for these two transects began at their respective melt-off dates.

Laboratory Analyses

I quantified forage quality by measuring neutral detergent fiber (NDF), acid detergent fiber (ADF), acid detergent lignin (ADL), dry matter digestibility (DMD), percent nitrogen (N), crude protein (CP), and digestible protein (DP) in each sample (ca. 1600). Protein precipitation capacity was determined on the additional 2018 deciduous shrub samples (ca. 200 samples) collected for this purpose. Prior to chemical analysis, all samples were ground to 1-mm particle size using a Wiley mill (Thomas Scientific, Swedesboro, New Jersey, USA).

Sequential fiber analysis (determination of NDF, ADF, and ADL- See Appendix 2) was conducted on all samples using the ANKOM Technology method (ANKOM Technology 2003a, b, 2011) and an ANKOM fiber analyzer (model 200, ANKOM Technology, Macedon, New York, USA). Forages were first extracted in a neutral detergent solution with agitation at 100°C to obtain the easily digested, or solubilized, fraction (neutral detergent solubles, NDS) and insoluble fraction (NDF). NDF residues were then extracted with an acid detergent solution to yield ADF (a measure of the least digestible plant components like cellulose and lignin). The ADF residue was digested with 72% sulfuric acid to determine ADL, then ashed in a muffle furnace at 500°C for 5 hours to determine the total proportion of non-digestible lignin-cutin and inorganic matter. Percent N was analyzed via combustion at 1800°C in tin capsules using an elemental analyzer (ECS 4010, Costech Analytical Technologies Inc., Valencia, California, USA).

I assessed the protein-precipitating capacity of tannins in the deciduous shrubs, *S. pulchra* and *B. nana*, using bovine serum albumin (BSA) according to the methods developed by McArt et al. (2006). Samples were freeze-dried and ground to 1 mm particle size before the tannins were extracted in aqueous methanol using an accelerated solvent extractor (Dionex ASE-200, ASE-350, Thomas Scientific, Swedesboro, New Jersey, USA). Extracts were then serially diluted with a standard solution of BSA and an acetate buffer solution before the precipitate and 50 µl of solution was filtered into an optically clear microplate. Bradford Protein Reagent (Fisher Scientific, Pittsburgh, Pennsylvania, USA) was added, and the resulting solution was incubated at room temperature for six minutes before I read the absorbance at 595 nm on a UV-Vis microplate spectrometer (Synergy HT Multi-Mode Microplate Reader, BioTek Instruments Inc., Winooski, Vermont, USA).

DMD and DP of each sample were calculated using the following digestibility equations developed by Robbins et al. (1987) and Spalinger et al. (2010) for cervids:

$$DMD = (92.31e^{-0.0451(LIG)}*NDF) + (0.831*NDS - 6.97) \qquad eq. (1)$$

$$DP = -3.97 + 0.9283 * CP - 11.82 * PPC \qquad eq. (2)$$

where LIG is the lignin/cutin fraction calculated from sequential fiber analysis (See Appendix 2).

Statistical Analyses

Snow depth at the snow fence was calculated using a second order polynomial local area regression to interpolate between transect readings. Mean snow depth of each snow zone was then extracted from the area sampled. To examine the effect of snow depth on forage availability I ran a one-way ANOVA using the statistical computing language R (R Core Team 2018) on the biomass of each functional group and used lsmeans to determine where specific differences among the snow zones occurred. In this analysis differences among snow zones were considered significant at p < 0.05.

I examined the effect of snow depth on forage quality of each species using the nlme package (Pinheiro et al. 2018) to perform a linear mixed-effects analysis of the relationship of leaf-level N (Table 1) and dry matter digestibility (Table 2) of each species as well as the protein-precipitating capacity of the deciduous shrubs (Table 3) to the snow treatment zones. The base model included day of year as an independent variable, with plot and year as random effects to account for repeated sampling throughout the growing season and sampling in different years. Dependent variables were percent nitrogen content, percent dry matter digestibility, and protein-precipitating capacity (expressed as mg BSA precipitated/mg dry matter (DM)). This base model was compared to additional models that include additive and multiplicative interactions between snow depth treatments and day of year as well as a quadratic day of year term to account for rapid changes in the response variable during leaf expansion and senescence and slower changes during mid-summer. All proportionate data were arcsine square-root

transformed prior to analysis, and I selected the top model for each independent variable using AIC (Burnham and Anderson 2002). The top model was used to calculate predicted means and 95% confidence intervals for each variable over the sample period.

For all species except the deciduous shrubs, digestible protein was calculated directly from eq. 2 with the assumption that PPC=0. Since digestible protein is directly related to leaf N (through the crude protein variable in eq. (2)), the top N model was used with the calculated digestible protein values to obtain the predicted means and confidence intervals over the sampling period. For the shrubs, *S. pulchra* and *B. nana*, the top N and PPC models for both species (including snow zone where significant) were used to obtain daily crude protein and PPC estimates and standard errors of the estimates. I then randomly sampled from a normal distribution for each daily estimate of crude protein and PPC and calculated digestible protein using eq. (2). This procedure was repeated 1000 times for each day, producing a daily estimate and 95% confidence interval of digestible protein for both shrubs throughout the sampling period that accounts for the error of both crude protein and PPC trends.

Results

Snow and Active Layer Depth

My designated snow treatment zones matched snow depth collected in March 2019 (Fig. 1). Snow depth in the Ambient zone was 39 ± 11 cm, while snow depth in the - snow and + snow zones was 20 ± 3.4 cm and 147 ± 18 cm respectively.

Active layer depths increased rapidly throughout the growing season until DOY 217-220, when they approached maximum depth (See Appendix 1). Depths at the

beginning of the growing season (DOY 173) were only 8 ± 1 cm in the shallow end of the + snow treatment, 17 ± 6 cm in the - snow treatment, and 14 ± 5 cm in ambient conditions. Half of the + snow treatment zone was still covered by a snow drift that was 38 ± 11 cm deep. By the end of the growing season (DOY 237), active layer depths in all zones were 45-60 cm, with the snow-covered area of the + snow treatment increasing from an initial depth of 10 ± 1 cm at DOY 180 to 61 ± 10 cm at DOY 237. The ambient zone depth increased to similar levels at 60 ± 9 cm, and the - snow treatment and shallow end of the + snow treatment reached depths of 51 ± 10 cm and 49 ± 7 cm respectively.

Forage Availability

Overall biomass was not significantly different among the three snow zones, with $828.60 \pm 106.31 \text{ g/m}^2$ in the ambient plots, $748.24 \pm 35.26 \text{ g/m}^2$ in the – snow treatment plots, and $764.62 \pm 106.30 \text{ g/m}^2$ in the + snow treatment plots. Broken down by functional group though, the - snow and + snow treatments differed from ambient conditions in a few key ways. Both treatments had a higher biomass of deciduous shrubs than ambient (Fig. 2), with a statistically significant (F_{2,15} = 5.027, p < 0.05) higher biomass of deciduous shrubs occurring in the - snow treatment at $165.36 \pm 14.38 \text{ g/m}^2$ (compared to $135.82 \pm 14.79 \text{ g/m}^2$ in the + snow treatment). The - snow treatment also had the highest biomass of evergreen dwarf shrubs ($138.56 \pm 43.17 \text{ g/m}^2$), while the + snow treatment had the lowest ($59.45 \pm 13.38 \text{ g/m}^2$). Both the - snow and + snow treatments had significantly (F_{2,15} = 11.8, p < 0.001) lower lichen biomass (2.43 ± 1.80 and $8.51 \pm 4.10 \text{ g/m}^2$ respectively) compared to ambient ($50.68 \pm 12.47 \text{ g/m}^2$). For all treatments, total biomass of lichens and forbs was small compared to other functional groups, while all snow zones had similarly high biomass of both moss and graminoids. In

almost all functional groups, however, there was considerable spatial variation with each snow zone, resulting in high variation between individual plots.

Nitrogen Content

Snow depth affected N content of all sampled species except *V. vitis-idaea*, with all top models including either an additive or multiplicative interaction with snow and the quadratic day of year term (Table 1). Increased snow depth had the strongest effect on the deciduous shrub and graminoid functional groups. For both, N content was significantly greater in the + snow treatment, while there was little to no difference between the - snow treatment and ambient conditions (Fig. 3). This difference in N content among the snow zones remained relatively constant throughout the growing season. For *S. pulchra*, deep snow resulted in 11.4% greater N at the beginning of the growing season (DOY 173) compared to the ambient plots. The difference among the treatments remained relatively constant even as overall N levels declined during the season, leading to plants in the + snow plots having 90.0% higher leaf N when leaf senescence began (DOY 237). *B. nana* followed a similar pattern, although with a lesser increase in N content of 8.35% in + snow plots compared to ambient areas at the beginning of the growing season and 25.3% by the end.

Both sedges, *C. bigelowii* and *E. vaginatum*, demonstrated different patterns of N content throughout the growing season (Fig. 3). *C. bigelowii* followed much the same pattern as the shrubs, with highest N levels at the beginning of season and in the + snow treatment, declining steadily over time. While the - snow treatment and ambient plots maintained similar N levels through most of the growing season, N content of plants in ambient conditions declined sharply with the onset of senescence, resulting in 107%

higher N levels in the + snow treatment by the end of the growing season. *E. vaginatum* did not follow the same temporal pattern of N content as *C. bigelowii* or the deciduous shrubs. Instead, N content was low (~2%) over the course of the entire growing season. Plants in the + snow zone still had higher leaf N, especially near the end of the season, with a 30.4% increase over ambient.

For the evergreen dwarf shrubs, N content remained low throughout the entire growing season, with few samples ever rising above 2%. For *V. vitis-idaea*, snow depth did not have an effect on leaf N, while for *R. tomentosum* N was highest in the - snow treatment over the first few weeks of the growing season before falling below + snow levels by mid-season and rising again in the last weeks of August when N content in the + snow treatment began to decrease.

Dry Matter Digestibility

Snow depth affected dry matter digestibility less than N content for all species, with only a slight increase in digestibility of *B. nana* in the + snow treatment (Fig. 4). Even then, the increase was modest, with the highest digestibility occurring mid-season with an increase of 6.82% over ambient. By senescence, the gap among the three snow zones closed as digestibility in both the ambient and – snow plots increased slightly to match that of the + snow plots.

Few clear patterns emerged in dry matter digestibility of other sample species despite often including a snow term in the top models (Table 2). There was no effect of snow depth on the evergreen dwarf shrub, *V. vitis-idaea*, and the sedge, *C. bigelowii*, although both species showed different temporal patterns. Digestibility of *C. bigelowii*

(~70%) changed little throughout time, while digestibility of *V. vitis-idaea* showed a curvilinear trend over the growing season, with a lower overall digestibility of ~50-60%. For the deciduous shrub, *S. pulchra*, the + snow treatment showed higher digestibility at the beginning of the growing season. By mid-season, digestibility fell to the same level as the – snow treatment, both of which were below ambient levels. All three snow zones, however, differed by only a few percentage points throughout the entire growing season. Both the dwarf shrub, *R. tomentosum*, and the sedge, *E. vaginatum*, showed curvilinear trends through time as well as a slight effect of snow depth. Plants in all three snow zones exhibited similar digestibility at the beginning of the season (~45-50% in *R. tomentosum* and ~60-70% in *E. vaginatum*) only to diverge in the latter half (Fig. 4).

Protein-precipitating Capacity and Digestible Protein

Since digestible protein is directly correlated with the amount of nitrogen in a forage while also accounting for the protein loss due to anti-herbivory compounds in certain plants, digestible protein shows the same temporal patterns as the N content of each species (Fig 6). In the deciduous shrubs, however, anti-herbivory compounds like tannins bind to proteins in the plant and lower the total digestible protein an animal can obtain from them, decreasing overall forage quality (Robbins et al. 1987a). For *B. nana*, the top model for protein-precipitating capacity included snow depth (Table 3), with estimated PPC highest in the + snow treatment in the middle of the growing season at 0.276 ± 0.014 mg BSA precipitated/mg DM (Fig. 5). Both the + snow and ambient zones were similar however, and temporal variation was high, resulting in substantial overlap of confidence intervals between the two zones. For *S. pulchra*, there was no effect of snow on PPC, but PPC did increase throughout the growing season from an estimated low of

 0.226 ± 0.027 mg BSA precipitated/mg DM at the beginning to a high of 0.357 ± 0.015 mg BSA precipitated/mg DM by DOY 217 before declining to 0.334 ± 0.023 mg BSA precipitated/mg DM by senescence (Fig. 5).

Caribou need a minimum of 7-8 g DP/100g DM in their diet to maintain body condition (red line in Fig. 6) (Thompson and Barboza 2017). Even with the additional protein-precipitating capacity of anti-herbivory in the deciduous shrubs, digestible protein levels of both *S. pulchra* and *B. nana* remained above maintenance levels well into the growing season, especially in the + snow treatment (Fig. 6), with values as high as 20-25 g/100g DM. The evergreen dwarf shrubs remained a poor source of protein through the entire season, never rising above the maintenance threshold, and in the case of *V. vitis-idaea*, remaining at ca. 2 g/100g DM throughout the sample period. Digestible protein in *C. bigelowii* mirrored the same pattern as the deciduous shrubs, beginning the season at 15-20 g/100g DM and declining steadily with time until falling below maintenance levels near senescence. *E. vaginatum*, while not high in digestible protein, remained above the maintenance threshold for most of the growing season in all snow zones, and plants in the + snow treatment stayed just over the minimum protein requirement during the entire sampling period.

Discussion

My results demonstrate that increased snow depth affects both the availability and quality of forage species important to caribou on Alaska's North Slope. Deep snow areas had higher biomass of preferred functional groups over ambient snow areas as well as a decrease in the proportion of unpalatable evergreen dwarf shrubs. Deep snow also increased leaf-level N and digestible protein in both deciduous shrubs and graminoids, despite increasing the protein-precipitating capacity of anti-herbivory compounds in *B*. *nana*. While changes to forage digestibility due to snow were mixed, overall digestibility in preferred forages was high over the entire growing season. Increases in snow depth, then, may enhance both forage quantity and quality for caribou.

Effects of Snow Depth on Forage Availability

The most obvious changes in biomass among the three snow zones occurred in the deciduous shrub, evergreen dwarf shrub, and lichen functional groups, though only significantly so for deciduous shrubs and lichens. The deep snow area had the lowest evergreen shrub biomass, which is consistent with previous snow manipulation and fertilization studies (Chapin III et al. 1995, Demarco et al. 2014, Zamin et al. 2014). Many studies have also found a concomitant increase in biomass (Wipf and Rixen 2010) and/or percent cover (Wahren et al. 2005, Johansson et al. 2013, Leffler et al. 2016) of deciduous shrubs in areas with deeper snow, yet my results show the opposite, with the highest biomass of deciduous shrubs found in the reduced snow area, although both the reduced snow and deeper snow areas have higher shrub biomass than ambient conditions.

One plausible explanation for the discrepancy is that small-scale vegetation communities within the boundaries of the snow fence vary considerably, making differences due to snow depth difficult to quantify without a larger sample size than the six 1-m² plots used in this study. Another possible explanation is that several previous studies included additional treatments such as summer warming or additional N through fertilization combined with winter snow depth and the deciduous shrub response was greatest when deep winter snow interacted with these other treatments (Demarco et al. 2014, Leffler et al. 2016). Additionally, the snow depth in the deeper snow area may now

be too deep, with the snow fence amplifying natural increases in snow fall in the two decades since the fence was erected. Some studies have found decreased shrub cover in areas with deep snow (> 2 m) as opposed to areas with more moderate snow cover (1-2)m) (Wahren et al. 2005, Borner et al. 2008, Johansson et al. 2013) suggesting there is a maximum snow load a shrub-dominated community can tolerate. The number of growing degree days needed for onset of green-up at northern latitudes has a positive correlation with the amount of winter precipitation (Fu et al. 2014). With an already short growing season, snow depths that regularly delay green-up by more than two weeks may reduce productivity and eventually deplete energy reserves in plants, inducing mortality and reducing the abundance of certain species like deciduous shrubs and *Eriophorum spp*. (Walker et al. 1999, Borner et al. 2008, Wipf and Rixen 2010). In either case, the availability of important summer forage species like S. pulchra should increase with moderately more snow. The same cannot be said for important winter forages, including lichens and evergreen dwarf shrubs (Boertje 1984, Ophof et al. 2013), both of which decrease with more snow in this and other studies.

Effects of Snow Depth on Forage Quality

Additional snow increased leaf N in both the deciduous shrubs and sedges (Fig. 3), as has been found in previous studies across multiple arctic and alpine sites (Walker et al. 1999, Van der Wal et al. 2000, Welker et al. 2005, Leffler and Welker 2013). Increased snow depth had the greatest impact on N content in *S. pulchra*, but *B. nana* and the sedge *C. bigelowii* exhibited similar increases. This partially supports my hypothesis of snow having the greatest impact on the quality of deciduous shrubs due to deeper roots that can take advantage of higher soil nitrogen as well as greater phenological plasticity

in the face of environmental changes (Bret-Harte et al. 2001, Sullivan et al. 2007, Wipf and Rixen 2010). The impact of deep snow on N content of *C. bigelowii* is curious, though, as leaf N in *E. vaginatum*, another sedge, does not respond similarly to deeper snow, remaining low overall throughout the study period.

There are a few possible explanations for the difference seen in N content over the growing season between the two sedges. First, *E. vaginatum* replaces its roots annually (Chapin III 1986, Sullivan and Welker 2005) and it may not be able to take advantage of the transient increase in nutrients in the soil as early in the growing season as *C. bigelowii*, though that does not fully explain the low N levels in *E. vaginatum* throughout the entire growing season. Furthermore, *E. vaginatum* and *C. bigelowii* preferentially uptake different forms of nitrogen, with *E. vaginatum* using primarily ammonium and *C. bigelowii* using primarily nitrate (McKane et al. 2002). Deep snow increases availability of both forms of nitrogen in the soil, especially in intertussock areas (Schimel et al. 2004, Semenchuk et al. 2015) where *C. bigelowii* tends to be located (pers. obs.). Because nitrate levels are so low in arctic soils, there is more competition among plants for ammonium (McKane et al. 2002). Even as snow increases both forms of nitrogen, then, *C. bigelowii* may be better positioned to take advantage of the timing, location, and form of soil nitrogen than *E. vaginatum*.

Snow depth also influenced N concentrations in *R. tomentosum* (Fig. 3), with changes in N reflecting a phenological shift rather than a change in mean. As an evergreen, *R. tomentosum* normally produces new vegetative growth from mid-late July under ambient conditions (Murray and Miller 1982). I observed high N concentrations earlier in the growing season in the low snow area, which was snow free earliest in the

season. Leaves of *R. tomentosum* in the deep snow area exhibited similar N concentrations several weeks later, following later snow melt of the deeper snow drift. Hence, deeper snow shifted leaf production and subsequent peak leaf N in this species later in the growing season.

Interestingly, leaf N was higher for most species in the deeper snow area despite having the shallowest active layer depth over half the + snow zone (See Appendix 1). This suggests that higher nitrogen uptake by plants may rely more on snow insulating the soil enough for enhanced microbial activity during the winter than on the release of new nutrients as soils thaw deeper during the growing season (Schimel et al. 2004). While active layer depth is generally correlated with snow depth (Johansson et al. 2013), other factors such as soil moisture content, albedo and insulating properties of the vegetation types covering the soil surface cannot be discounted (Loranty et al. 2011).

Snow depth did not have as great an impact on leaf dry matter digestibility as leaf N, results similar to previous studies examining the effects of various environmental factors like shading, air temperature, and precipitation on forage quality (Lenart et al. 2002). Individual species within each functional group responded differently from one another, ranging from phenological shifts to no significant effects, and overall changes in dry matter digestibility trends, whether through time or with snow depth, usually spanned only a few percentage points. Even small changes in digestibility, however, can significantly impact dry matter intake of caribou (White 1983) and subsequent deposition of both body fat and protein (Chan-McLeod et al. 1994). As maternal winter body mass and body fat correlate strongly with animal survival and calf production (Parker et al.

2009, Proffitt et al. 2016, Veiberg et al. 2017), changes in summer forage digestibility and energy intake can propagate from the individual to the population level.

The protein-precipitating capacity of the deciduous shrubs did not respond to the winter climate change scenarios the way I initially expected. PPC of *S. pulchra* was unaffected by snow depth; however, PPC increased with snow depth in *B. nana*, albeit only slightly. Previous studies in tussock tundra in the Toolik Lake area found that concentrations of phenolics in *B. nana* increased with N fertilization treatments (Graglia et al. 2001), while other studies found a more generalized increase in carbon-based secondary compounds (including phenolics and tannins) with N fertilization (Lavola and Julkunen-Tiito 1994, De Long et al. 2016). With the increase in available soil nutrients with deep snow cover, one might expect to see lower PPC as plants use available nitrogen to shunt carbon tied up in secondary compounds into new growth rather than defense (Chapin III 1989). The results presented here show just the opposite, however.

Protein-precipitating capacity of *B. nana* was highest in the deeper snow area and lowest in the reduced snow area, although there was substantial variation among sample dates. The seasonal variation may be due in part to a phenological shift with later melt-off of the deeper snow area, as values of PPC in leaf tissue of plants in deeper snow followed roughly the same pattern as those in the ambient snow area, just offset by two weeks. One explanation for the seemingly opposite results of studies in tussock tundra surrounding Toolik Lake arises from different assays used to quantify various secondary compounds. Most studies measure total phenolic content as opposed to protein-precipitating capacity (Graglia et al. 2001, Zamin et al. 2017a), though some have measured both (De Long et al. 2016) and found a similar decrease in PPC when N is added as a fertilizer to tundra.

These fertilization responses may be sensitive to timing, however, as most measurements of both phenolic content and PPC were taken near the end of July. *B. nana* in this experiment exhibit a similar lower PPC in deeper snow areas at about the same time as prior studies; however, I observed considerable variation when analyzing trends over the entire growing season. Responses of both phenolic content and PPC to fertilization may also be specific to certain secondary compounds that use different biosynthetic pathways that may or may not compete with the synthesis of proteins necessary for plant growth (Chapin III 1989, Haukioja et al. 1998).

Despite the dampening effect of PPC on digestible protein content in the deciduous shrubs, deeper snow increased the amount of digestible protein in certain forage species (primarily through enhancing overall N content) while also increasing the length of time that digestible protein content is above the minimum maintenance levels required by caribou during the summer. The largest impact is on the value of deciduous shrubs and sedges as a protein source. For example, deep snow resulted in ~ 25 additional days of digestible protein above the maintenance threshold in *S. pulchra* compared to low snow and ambient conditions. This doubles the length of time that caribou can gain sufficient protein to recover body condition and sustain weight gain in ambient snow conditions and mimics the increased duration of protein observed along latitudinal gradients (Barboza et al. 2018). The duration of time that digestible protein in *B. nana* and C. bigelowii remained above maintenance levels was also extended, albeit more modestly at 6 and 9 days respectively. In addition, while the digestible protein content of *E. vaginatum* was lower overall than other species for most of the growing season, levels in the deeper snow area did not fall below the maintenance threshold during the sampling

period, suggesting that *E. vaginatum* may remain a useful protein source throughout much of the year (Klein 1990, Ophof et al. 2013) with deeper snow.

Limitations

There are a few limitations with this study that must be addressed. First, this study took place at a single snow fence located in moist acidic tussock tundra. There are strong regional variations in responses of tundra to warming and fertilization experiments (Wipf and Rixen 2010, Elmendorf et al. 2011), suggesting that complex interactions among climate, geology, and hydrology also determine how specific vegetation communities respond to change. However, my largest observed responses to deeper snow (i.e. higher N in S. pulchra and B. nana) are broadly consistent with several studies in different locations (Walsh et al. 1997, Schimel et al. 2004, Welker et al. 2005, Leffler and Welker 2013, Semenchuk et al. 2015). It should also be noted that the deepest snow drifts created by the snow fence in this experiment are deeper than even the most extreme predictions of increased winter precipitation in the region (Scenarios Network for Alaska and Arctic Planning 2011). More moderate increases in snow, though, may actually result in greater long-term changes to plant communities than exceptionally deep snow associated with snow fence studies (Borner et al. 2008, Wipf and Rixen 2010). For assessing forage quality, I also make a few assumptions. Since I only analyzed protein-precipitating capacity for the deciduous shrubs, I assumed the PPC for all other species was 0 when calculating digestible protein. While graminoids are low in phenolic compounds that bind protein, evergreen dwarf shrubs have high concentrations of secondary compounds that make them relatively unpalatable (Bryant et al. 1983). While the PPC of the evergreen shrubs remains unaccounted for in this study, the calculated digestible protein values are

already well below the maintenance threshold, so their value as summer forage remains unchanged. Although the results of this study may not apply to tundra ecosystems as a whole, it still provides one of the few examples of how long-term changes in snow depth affect certain tundra plants and what that means for both the quantity and quality of forage for large arctic herbivores now and in the future.

Conclusions

Because tundra ecosystems are so nutrient-limited (Bryant et al. 1983), small changes in both availability and quality of forage containing essential nutrients like protein can have outsize impacts at both the individual and population levels (White 1983). Larger quantities of high-quality plants increase forage intake by caribou, which increases the rate and amount of weight gained during the short growing season (White and Trudell 1980). Only a few kilograms of weight gain can increase the chance of conception by as much as 60% (White 1983, Proffitt et al. 2016). As capital breeders, caribou rely on body stores of fat and protein to support pregnancy and early lactation during the winter and early spring (Barboza et al. 2018). An increase in summer forage quality reduces the time needed to recover body reserves from both winter deprivations and the nutritional demand of lactation while increasing calf weight gain and growth rates (White et al. 1975, Veiberg et al. 2017). A small positive change in future forage nutrition from the increased snow predicted by climate models and observed over the past 20+ years, may potentially mitigate declines in forage quality due to higher temperatures (Jonasson et al. 1986, Turunen et al. 2009, Fauchald et al. 2017, Zamin et al. 2017a) and lead to higher survival, recruitment, and population growth rates of caribou in N Alaska.

Tables

Model	logLik	AICc	ΔAIC _c	df	weight
S. pulchra					
SNOW*DOY*DOY ²	869.63	-1707.23	0.00	15	0.87
SNOW + DOY*DOY ²	861.12	-1703.49	3.74	9	0.13
DOY*DOY ²	742.27	-1470.08	237.14	7	0.00
DOY + DOY ²	727.04	-1441.75	265.48	6	0.00
DOY	725.91	-1441.58	265.65	5	0.00
B. nana					
SNOW + DOY*DOY ²	840.98	-1663.23	0.00	9	0.87
SNOW*DOY*DOY ²	845.71	-1659.41	3.82	15	0.13
DOY*DOY ²	808.27	-1602.09	61.14	7	0.00
DOY + DOY ²	756.96	-1501.58	161.66	6	0.00
DOY	751.51	-1492.77	170.46	5	0.00
L. palustre					
SNOW*DOY*DOY ²	980.25	-1928.56	0.00	15	0.99
SNOW + DOY*DOY ²	968.63	-1918.55	10.01	9	0.01
DOY*DOY ²	962.00	-1909.55	19.01	7	0.00
$DOY + DOY^2$	654.51	-1896.69	31.87	6	0.00
DOY	950.75	-1891.26	37.30	5	0.00
V.vitis-idaea					
DOY	968.84	-1927.45	0.00	5	0.46
SNOW + DOY*DOY ²	972.45	-1926.21	1.24	9	0.25
DOY + DOY ²	968.90	-1925.47	1.98	6	0.17
DOY*DOY ²	969.30	-1924.17	3.28	7	0.09
SNOW*DOY*DOY ²	977.24	-1922.58	4.87	15	0.04
C. bigelowii					
SNOW*DOY*DOY ²	861.93	-1691.93	0.00	15	1.00
SNOW + DOY*DOY ²	843.04	-1667.37	24.56	9	0.00
DOY*DOY ²	815.11	-1615.79	76.14	7	0.00
DOY + DOY ²	795.53	-1578.74	113.19	6	0.00
DOY	793.40	-1576.57	115.36	5	0.00
E. vaginatum					
SNOW* DOY*DOY ²	861.04	-1690.19	0.00	15	0.65
SNOW + DOY*DOY ²	853.83	-1688.97	1.23	9	0.35
DOY*DOY ²	832.92	-1651.42	38.77	7	0.00
DOY + DOY ²	831.19	-1650.07	40.13	6	0.00
DOY	820.93	-1631.64	58.55	5	0.00

Table 1. Model selection results for nitrogen content of each species
Table 2. Model selection for dry matter digestibility of each species

Model	logLik	AICc	ΔΑΙϹ	df	weight
S. pulchra					
SNOW*DOY*DOY ²	710.34	-1388.64	0.00	15	1.00
SNOW + DOY*DOY ²	686.75	-1354.76	33.89	9	0.00
DOY*DOY ²	682.45	-1350.45	38.20	7	0.00
$DOY + DOY^2$	679.38	-1346.42	42.23	6	0.00
DOY	664.83	-1319.42	69.22	5	0.00
B. nana					
SNOW*DOY*DOY ²	624.33	-1216.64	0.00	15	1.00
SNOW + DOY*DOY ²	607.62	-1196.49	20.15	9	0.00
DOY + DOY ²	535.26	-1058.17	158.47	6	0.00
DOY*DOY ²	535.51	-1056.56	160.08	7	0.00
DOY	530.52	-1050.8	165.84	5	0.00
L. palustre					
SNOW*DOY*DOY ²	581.01	-1130.12	0.00	15	0.90
SNOW + DOY*DOY ²	572.15	-1125.6	4.52	9	0.09
DOY*DOY ²	566.10	-1117.76	12.36	7	0.00
DOY + DOY ²	543.41	-1074.49	55.63	6	0.00
DOY	526.78	-1043.34	86.79	5	0.00
V.vitis-idaea					
DOY*DOY ²	594.41	-1174.4	0.00	7	0.62
SNOW + DOY*DOY ²	595.81	-1172.92	1.48	9	0.30
SNOW*DOY*DOY ²	601.14	-1170.37	4.03	15	0.08
DOY	569.53	-1128.83	45.56	5	0.00
DOY + DOY ²	569.74	-1127.17	47.23	6	0.00
C. bigelowii					
DOY	291.43	-572.63	0.00	5	0.43
$DOY + DOY^2$	292.36	-572.4	0.23	6	0.39
DOY*DOY ²	292.52	-570.59	2.04	7	0.16
SNOW + DOY*DOY ²	295.56	-566.41	6.22	9	0.02
SNOW*DOY*DOY ²	297.52	-563.12	9.51	15	0.00
E. vaginatum					
SNOW*DOY*DOY ²	253.19	-474.5	0.00	15	0.50
DOY*DOY ²	244.35	-474.27	0.23	7	0.44
SNOW + DOY*DOY ²	244.51	-470.34	4.16	9	0.06
DOY	232.26	-454.3	20.21	5	0.00
$DOY + DOY^2$	232.27	-452.21	22.29	6	0.00

Model	logLik	AICc	ΔΑΙϹ	df	weight
S. pulchra					
DOY + DOY ²	83.39	-156.07	0.00	5	0.56
DOY*DOY ²	83.60	-154.20	1.87	6	0.22
DOY	80.94	-153.41	2.66	4	0.15
SNOW + DOY*DOY ²	84.46	-151.16	4.92	8	0.05
SNOW*DOY*DOY ²	91.72	-149.91	6.16	14	0.03
B. nana					
SNOW + DOY*DOY ²	127.56	-237.34	0.00	8	0.81
DOY*DOY ²	123.20	-233.40	3.94	6	0.11
DOY + DOY ²	121.17	-231.63	5.71	5	0.05
SNOW*DOY*DOY ²	131.91	-230.22	7.11	14	0.02
DOY	117.39	-226.31	11.02	4	0.00

Table 3. Model selection results for protein-precipitating capacity of S. pulchra and B. nana

Figures



Figure 1. Snow depth (cm) of sampling area behind the snow fence (located along southeast corner) in March 2019. Polygons indicate the areas sampled. **A** represents the ambient snow treatment zone; **B** represents the – snow treatment zone; **C** represents the + snow treatment zone.







Figure 3. Leaf tissue N concentration by species throughout the growing season. Points indicate individual measurements, while trend lines are the predicted median \pm 95% confidence intervals in shaded polygons. Pre-leaf emergence values for S. pulchra and B. nana are not included in the model. Note that y-axis range differs among functional groups.



Figure 4. Leaf tissue dry matter digestibility by species throughout the growing season. Points indicate individual measurements while trend lines are the predicted median \pm 95% confidence intervals in shaded polygons. Pre-leaf emergence values for *S. pulchra* and *B. nana* are not included in the model. Note that y-axis range differs among functional groups.



Figure 5. Protein-precipitating capacity by species through the growing season. Points indicate individual measurements, while trend lines are the predicted median \pm 95% confidence intervals in shaded polygons. Pre-leaf emergence values were not included in the model.



Figure 6. Leaf tissue digestible protein. Trend lines are the predicted median \pm 95% confidence intervals in shaded polygons. The maintenance level protein required by caribou (7-8 g/100g DM) is highlighted in red. Note that y-axis range differs among functional groups.

CHAPTER 3: CONCLUSIONS

Results of this research indicate that projected increases in winter precipitation over Alaska's North Slope may indirectly increase the quality of caribou forage. Deeper snow insulates the soil and allows microbial mineralization to continue throughout the winter, increasing soil nitrogen available for plant uptake in early spring. Snow depth may not have as large an impact on dry matter digestibility as leaf-level nutrients, but even the small increases seen in this study may influence forage intake, with subsequent multiplier effects on survival and fecundity. Direct changes in N availability and indirect changes in vegetation community structure, though, may have a stronger influence on overall caribou nutrition in the Arctic than species-specific changes in forage digestibility.

Caribou and reindeer populations around the Arctic face many challenges: extreme weather, predation, insect harassment, the encroachment of human development (Morschel and Klein 1997, Vors and Boyce 2009, Festa-Bianchet et al. 2011), but climate change presents a new challenge, bringing increased temperatures, shifting precipitation patterns, and altering forage availability and quality (Callaghan et al. 2011, Fauchald et al. 2017, Mallory and Boyce 2018). While warmer growing season temperatures are associated with declines in forage quality due to increased vegetative growth and nutrient dilution (Turunen et al. 2009, Fauchald et al. 2017, Zamin et al. 2017b), increases in winter snow may mitigate the magnitude of such declines. Additionally, the availability of preferred forages like deciduous shrubs is expected to continue increasing as the Arctic becomes shrubbier, though certain species, like *B. nana* (which is less preferred than *Salix spp.)*, may spread more rapidly (Bret-Harte et al. 2001, Sturm et al. 2001b, 2005, Tape et al. 2006). Arctic herbivores may adjust to the increase in available forage and potentially curb the trend towards shrubification (Gough et al. 2007, Zamin and Grogan 2013, Kaarlejärvi 2014, Kaarlejärvi et al. 2017), though perhaps only in the short term due to the strong influence of abiotic stochasticity on both plant and animal populations in such extreme environments (Jefferies et al. 1994, Loe et al. 2016). Warmer temperatures and increased shrub cover may also decrease important winter forages such as lichens and evergreen dwarf shrubs (Wahren et al. 2005, Hobbie et al. 2017), so it is difficult to say whether the net effect of changes in temperature and precipitation in the Arctic will be positive or negative for caribou. This question of the net effect, as a balance of positive feedback and feedforward, has been of interest for several decades now in the Arctic (Welker et al. 1997).

One of the major discussions as of late has been the seasonality of caribou forage as a critical component of meeting animal metabolic needs during the autumn rut and prewinter preparation and its importance to subsequent winter survival and herd fecundity (Gustine et al. 2017, Veiberg et al. 2017, Barboza et al. 2018). My data support the prediction that one of the major consequences of deeper snow in winter is that caribou available protein (CAP), delivered primarily by *Salix*, will be greatly extended through the short growing season by as much as three weeks. This higher level of CAP during a hyper-critical season may be especially important to sustaining the health and welfare of caribou in northern Alaska.

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S. pulct	hra			NDF (g,	/g DM)					ADF (g/i	g DM)					ellulose ((/g DM)					ignin (g/g	DM)		
Year	роу	- Snc	MO	Amb	ient	+ Sn	MO	- Snc	Ň	Ambi	ent	+ Sno	3	- Snov	2	Ambie	t	+ Snot	2	- Snov		Ambier	Ŧ	+ Sno	3
		Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	dean	SD	Mean	SD
2017	170	0.231	0.029	0.252	0.021	0.286	0.053	0.193	0.024	0.217	0.020	0.245	0.047	660.0	0.014	0.106	0.013	0.137	0.025	0.092	0.013 (0.107 (0.018	0.105	0.022
	177	0.164	0.005	0.160	0.006	0.144	0.007	0.120	0.005	0.118	0.003	0.110	0.005	0.061	0.003	0.060	0.002	0.058	0.001	0.056	0.004	0.055 (0.005	0.049	0.005
	184	0.195	0.004	0.214	0.011	0.190	0.016	0.137	0.004	0.143	0.007	0.135	0.011	0.075	0.001	0.076	0.003	0.072	0.008	0.060	0.004	0.065 (0.007	0.061	0.006
	191	0.208	0.016	0.211	0.006	0.186	0.019	0.135	0.006	0.143	0.005	0.124	0.013	0.074	0.006	0.079	0.004	0.066	0.007	0.058	0.004	0.063 (0.004	0.056	0.012
	198	0.220	0.006	0.221	0.010	0.224	0.023	0.142	0.003	0.136	0.004	0.139	0.013	0.076	0.002	0.073	0.002	0.076	0.008	0.065	0.003	0.061 (0.005	0.061	0.008
	205	0.230	0.004	0.228	0.006	0.251	0.015	0.144	0.002	0.144	0.003	0.150	0.008	0.076	0.005	0.073	0.006	0.073	0.016	0.067	0.005	0.070	0.006	0.075	0.012
	212	0.233	0.016	0.219	0.011	0.256	0.012	0.140	0.006	0.138	0.007	0.152	0.003	0.073	0.003	0.075	0.002	0.079	0.005	0.062) 600.c	0.056 (0.007	0.066	0.003
	219	0.261	0.007	0.235	0.015	0.261	0.014	0.160	0.008	0.142	0.007	0.155	0.011	0.080	0.002	0.079	0.003	0.077	0.006	0.073) 600.c	0.059 (0.005	0.073	0.007
2018	173	0.143	0.017	0.153	0.012			0.109	0.017	0.117	0.010			0.056	0.002	0.063	0.003			0.049	0.016	0.048 (0.007		
	180	0.165	0.020	0.172	0.010	0.123	0.004	0.125	0.016	0.135	0.008	0.096	0.004	0.064	0.004	0.067	0.005	0.058	0.001	0.055	0.012	0.064 (0.011	0.032	0.004
	187	0.192	0.019	0.165	600.0	0.155	0.018	0.134	0.014	0.120	0.008	0.118	0.014	0.072	0.002	0.024	0.097	0.064	0.004	0.059	0.013 (0.049 (0.005	0.050	0.010
	194	0.183	0.016	0.185	0.013	0.173	0.021	0.125	0.011	0.128	0.010	0.128	0.014	0.073	0.006	0.070	0.007	0.068	0.010	0.049	0.006	0.055 (0.004	0.054	0.005
	200	0.195	0.013	0.192	600.0	0.185	0.010	0.133	0.010	0.132	0.008	0.125	0.006	0.074	0.005	0.077	0.005	0.073	0.005	0.053	0.007	0.049 (0.005	0.048	0.003
	210	0.252	0.023	0.223	0.008	0.217	0.00	0.171	0.022	0.141	0.007	0.141	0.006	0.083	0.006	0.082	0.004	0.080	0.005	0.083	0.022	0.056 (0.002	0.056	0.002
	217	0.244	0.003	0.245	600.0	0.247	0.036	0.153	0.005	0.153	0.005	0.154	0.021	0.080	0.005	0.084	0.004	0.081	0.005	0.067	0.003	0.062 (0.002	0.065	0.018
	223	0.220	0.005	0.207	0.007	0.254	0.007	0.143	0.002	0.131	0.006	0.160	0.005	0.079	0.004	0.074	0.002	0.081	0.007	0.058	0.003	0.052 (0.004	0.073	0.006
	231	0.219	0.002	0.213	0.006	0.223	0.010	0.141	0.002	0.133	0.002	0.139	0.005	0.076	0.003	0.073	0.003	0.076	0.004	0.059	0.001	0.054 (0.004	0.056	0.005
	237	0.240	0.004	0.223	0.006	0.207	0.006	0.151	0.002	0.140	0.002	0.126	0.002	0.081	0.001	0.077	0.003	0.067	0.001	0.063	0.003	0.054 (0.003	0.054	0.002

APPENDIX 2: Sequential fiber analysis components by species and snow zone

			Ash (g/g DM)					ell Solubles	(% of DM)					Cell Wall (%	of DM)					Lig/Cut (%	of NDF)		
- Sno	ju o	3	Am	bient	+ Sr	wo	- Sno	Ň	Ambi	ent	+ Sno	MO	- Sno	z	Ambie	t	+ Sno	3	- Sno	3	Ambie	ant	+ Sno	z
Mean		SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD
0.003		0.001	0.003	0.001	0.003	0.001	76.859	2.910	74.776	2.093	71.408	5.251	23.141	2.910	25.224	2.093	28.592	5.251	39.798	3.091	42.460	5.377	36.707	2.060
0.003		0.001	0.003	0.001	0.002	0.002	83.569	0.513	84.020	0.635	85.594	0.746	16.431	0.513	15.980	0.635	14.406	0.746	33.971	2.238	34.117	1.632	34.173	2.031
0.003		0.000	0.002	0.000	0.002	0.000	80.496	0.422	78.558	1.108	81.007	1.560	19.504	0.422	21.442	1.108	18.993	1.560	30.505	1.477	30.188	1.699	32.277	2.864
0.003		0.001	0.001	0.001	0.002	0.001	79.210	1.573	78.901	0.640	81.403	1.893	20.790	1.573	21.099	0.640	18.597	1.893	28.064	2.012	29.725	1.457	30.141	3.571
0.002		0.001	0.001	0.000	0.001	0.000	77.989	0.621	77.931	1.043	77.636	2.267	22.011	0.621	22.069	1.043	22.364	2.267	29.325	1.575	27.753	1.515	27.389	2.955
0.002		0.001	0.002	0.000	0.001	0.000	76.970	0.442	77.209	0.567	74.922	1.523	23.030	0.442	22.791	0.567	25.078	1.523	29.105	2.455	30.595	2.267	30.038	5.039
0.005		0.002	0.007	0.002	0.007	0.003	76.679	1.619	78.075	1.062	74.376	1.174	23.321	1.619	21.925	1.062	25.624	1.174	26.675	2.390	25.422	2.753	25.745	1.274
0.007		0.002	0.005	0.002	0.004	0.003	73.946	0.701	76.470	1.461	73.937	1.368	26.054	0.701	23.530	1.461	26.063	1.368	27.941	2.916	24.917	0.718	28.102	1.291
0.004		0.001	0.006	0.001			85.689	1.744	84.698	1.245			14.311	1.744	15.302	1.245			33.626	6.443	31.160	1.941		
0.006		0.001	0.004	0.001	0.006	0.001	83.468	2.018	82.754	0.973	87.744	0.403	16.532	2.018	17.246	0.973	12.256	0.403	33.004	3.954	37.231	4.322	25.915	2.616
0.003		0.001	0.047	0.095	0.004	0.001	80.834	1.867	83.546	0.911	84.516	1.840	19.166	1.867	16.454	0.911	15.484	1.840	30.521	3.446	29.986	1.487	31.744	3.083
0.003		0.001	0.004	0.001	0.006	0.001	81.720	1.627	81.490	1.344	82.710	2.067	18.280	1.627	18.510	1.344	17.290	2.067	26.768	1.696	29.482	0.722	31.399	3.618
0.006		0.001	0.006	0.002	0.004	0.001	80.550	1.295	80.783	0.871	81.464	0.957	19.450	1.295	19.217	0.871	18.536	0.957	26.958	2.119	25.600	2.054	25.682	0.985
0.005		0.001	0.003	0.001	0.005	0.002	74.754	2.253	77.682	0.810	78.263	0.860	25.246	2.253	22.318	0.810	21.737	0.860	32.470	5.607	24.893	0.467	25.661	0.641
0.005		0.002	0.007	0.001	0.007	0.002	75.625	0.344	75.465	0.893	75.337	3.617	24.375	0.344	24.535	0.893	24.663	3.617	27.643	1.364	25.406	0.313	25.959	3.452
0.005		0.001	0.005	0.001	0.006	0.000	78.037	0.454	79.296	0.711	74.638	0.670	21.963	0.454	20.704	0.711	25.362	0.670	26.355	1.292	25.161	1.465	28.560	1.804
0.006		0.002	0.005	0.001	0.007	0.003	78.098	0.238	78.702	0.555	77.676	0.998	21.902	0.238	21.298	0.555	22.324	0.998	27.156	0.782	25.489	1.410	25.228	2.187
0.006		0.002	0.008	0.001	0.005	0.001	76.020	0.380	77.735	0.573	79.329	0.645	23.980	0.380	22.265	0.573	20.671	0.645	26.261	1.290	24.447	0.971	26.197	1.212

S. pul	chra			NDF digest	tibility (%)					NDS digest	ibility (%)		
Year	DOY	- Sn	MO	Amb	ient	us +	wot	- Snc	MO	Ambi	ient	+ Sn	MO
		Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD
2017	170	3.581	0.663	3.498	0.818	5.050	0.925	56.899	2.418	55.169	1.739	52.370	4.363
	177	3.290	0.348	3.168	0.170	2.851	0.221	62.476	0.427	62.851	0.528	64.158	0.620
	184	4.553	0.258	5.071	0.221	4.128	0.687	59.922	0.351	58.312	0.921	60.347	1.296
	191	5.447	0.816	5.101	0.276	4.415	0.534	58.853	1.307	58.597	0.532	60.676	1.573
	198	5.429	0.473	5.834	0.421	6.065	1.136	57.839	0.516	57.791	0.867	57.545	1.884
	205	5.754	0.666	5.314	0.526	660.9	1.323	56.992	0.367	57.191	0.471	55.290	1.266
	212	6.464	0.453	6.457	0.732	7.423	0.630	56.750	1.345	57.910	0.883	54.836	0.976
	219	6.857	0.840	7.056	0.327	6.768	0.160	54.479	0.583	56.577	1.214	54.472	1.137
2018	173	2.918	0.481	3.457	0.119			64.237	1.449	63.414	1.035		
	180	3.426	0.184	2.996	0.417	3.526	0.320	62.392	1.677	61.799	0.809	65.945	0.335
	187	4.458	0.281	3.925	0.117	3.398	0.174	60.203	1.552	62.457	0.757	63.263	1.529
	194	5.054	0.571	4.527	0.419	3.956	0.978	60.940	1.352	60.748	1.117	61.762	1.718
	200	5.322	0.346	5.605	0.507	5.376	0.344	59.967	1.076	60.161	0.724	60.726	0.795
	210	5.434	0.872	6.723	0.313	6.313	0.400	55.151	1.873	57.584	0.673	58.067	0.714
	217	6.478	0.414	7.204	0.343	7.007	0.367	55.875	0.286	55.742	0.742	55.635	3.006
	223	6.183	0.343	6.148	0.314	6.464	0.374	57.879	0.378	58.925	0.591	55.054	0.557
	231	5.945	0.270	6.231	0.260	6.634	0.727	57.930	0.198	58.431	0.461	57.579	0.829
	237	6.781	0.401	6.828	0.319	5.866	0.436	56.202	0.316	57.627	0.476	58.953	0.536

	MO	SD	0:030	0.010	0.006	0.013	0.020	0.012	0.009	0.016		0.011	0.008	0.013	0.003	0.015	0.015	0.018	0.008	0.007
	+ Sn	Mean	0.213	0.082	0.065	0.064	0.082	0.062	0.050	0.057		0.052	0.055	0.060	0.057	0.070	0.071	0.076	0.067	0.074
/g DM)	ient	SD	0.055	0.040	0.004	0.008	0.003	0.009	0.005	0.017	0.015	0.008	0.013	0.008	0.027	0.016	0.007	0.011	0.003	0.008
Lignin (£	Amb	Mean	0.195	0.052	0.082	0.087	0.085	0.095	0.100	0.094	0.070	0.068	0.079	0.070	0.081	0.088	0.109	0.081	0.078	0.092
	MO	SD	0.029	0.008	0.008	0.006	0.006	0.006	0.013	0.016	0.007	0.005	0.00	0.00	0.024	0.017	0.012	0.014	0.012	0.006
	-Sn	Mean	0.203	0.074	0.094	0.083	0.093	0.104	0.097	0.107	0.067	0.070	0.087	0.079	0.094	0.105	0.112	0.118	0.101	0.109
	MOL	SD	0.017	0.003	0.003	0.002	0.011	0.002	0.003	0.006		0.004	0.003	0.006	0.004	0.006	0.006	0.009	0.003	0.004
	IS +	Mean	0.139	0.058	0.062	0.056	0.064	0.059	0.056	0.058		0.058	0.059	0.061	0.066	0.072	0.067	0.065	0.066	0.068
(MD g/g)	pient	SD	0.008	0.002	0.003	0.006	0.001	0.004	0.002	0.001	0.003	0.002	0.003	0.008	0.017	0.007	0.004	600.0	0.003	0.001
Cellulose	Amt	Mean	0.105	0.048	0.068	0.068	0.065	0.069	0.069	0.066	0.061	0.061	0.063	0.064	0.065	0.075	0.077	0.066	0.067	0.072
	MO	SD	0.010	0.002	0.004	0.004	0.003	0.001	0.003	0.005	0.004	0.005	0.007	0.005	0.002	0.005	0.004	0.003	0.002	0.002
	- Sn	Mean	0.127	0.047	0.073	0.063	0.063	0.068	0.068	0.062	0.059	0.059	0.068	0.064	0.071	0.074	0.075	0.066	0.067	0.068
	wor	SD	0.044	0.010	0.008	0.016	0.029	0.011	0.013	0.017		0.013	0.009	0.019	0.007	0.020	0.020	0.025	0.008	0.006
	+St	Mean	0.359	0.143	0.128	0.121	0.148	0.123	0.112	0.122		0.117	0.117	0.125	0.128	0.147	0.145	0.147	0.139	0.149
/g DM)	pient	SD	0.055	0.004	0.005	0.011	0.004	0.010	0.006	0.016	0.018	0.009	0.014	0.015	0.044	0.021	0.010	0.017	0.007	600.0
ADF (g	Amt	Mean	0.306	0.121	0.153	0.158	0.151	0.167	0.174	0.165	0.135	0.132	0.146	0.138	0.151	0.165	0.191	0.153	0.150	0.170
	MOL	SD	0.038	0.004	0.010	0.006	0.008	0.007	0.016	0.019	0.010	0.007	0.012	0.013	0.024	0.021	0.015	0.020	0.011	0.010
	-s	Mean	0.335	0.122	0.170	0.149	0.157	0.173	0.169	0.174	0.131	0.134	0.160	0.147	0.172	0.183	0.193	0.191	0.174	0.185
	Mon	SD	0.051	0.011	0.010	0.024	0.026	0.019	0.025	0.027		0.019	0.014	0.025	0.012	0.026	0.029	0.028	0.012	0.011
	-S +	Mean	0.433	0.213	0.203	0.205	0.251	0.206	0.193	0.216		0.160	0.167	0.189	0.199	0.242	0.240	0.249	0.235	0.241
(MD B/)	bient	SD	0.051	0.006	0.010	0.014	0.011	0.025	0.015	0.015	0.023	0.011	0.017	0.015	0.062	0.035	0.011	0.028	0.015	0.014
NDF (g	Aml	Mean	0.365	0.183	0.237	0.250	0.260	0.275	0.280	0.274	0.180	0.184	0.213	0.219	0.220	0.278	0.289	0.249	0.237	0.260
	MOL	SD	0.030	0.009	0.018	0.003	0.011	0.024	0.026	0.025	0.012	0.017	0.021	0.021	0.040	0.028	0.026	0.017	0.010	0.012
	- Sr	Mean	0.414	0.199	0.274	0.249	0.286	0.293	0.290	0.301	0.186	0.197	0.244	0.250	0.263	0.299	0.308	0.308	0.278	0.295
ana	DOY		170	177	184	191	198	205	212	219	173	180	187	194	200	210	217	223	231	237
B. r	Year		2017								2018									

	Ň	SD	3.192	2.945	2.257	2.907	5.141	2.756	2.159	4.758		3.876	3.268	2.877	1.499	3.147	2.896	3.571	2.583	1.604
	+ Sno	Mean	49.167	38.327	31.877	30.802	32.398	30.027	26.083	26.144		32.370	32.949	31.573	28.694	28.876	29.383	30.123	28.436	30.538
of NDF)	ţ	SD	7.746	21.770	1.183	1.347	0.602	1.518	0.835	4.422	3.776	2.943	3.946	1.801	3.346	2.340	1.307	2.037	1.598	1.212
Lig/Cut (%	Ambie	Mean	52.781	28.669	34.714	34.919	32.689	34.634	35.948	34.102	38.306	36.774	37.126	31.702	36.012	30.965	37.596	32.510	32.909	35.378
	>	SD	4.282	2.701	2.003	2.318	1.437	1.208	1.790	3.128	1.805	3.069	2.432	1.607	5.237	3.712	1.401	2.596	2.919	1.085
	- Snov	Mean	48.871	36.972	34.412	33.523	32.526	35.396	33.286	35.469	36.211	35.755	35.606	31.493	35.318	35.069	36.384	38.242	36.345	36.880
	3	SD	5.126	1.144	0.950	2.352	2.583	1.903	2.527	2.651	 	1.894	1.363	2.492	1.212	2.568	2.878	2.779	1.230	1.114
	+ Sno	Mean	43.325	21.295	20.285	20.507	25.053	20.647	19.285	21.614		15.998	16.708	18.908	19.896	24.170	24.031	24.902	23.478	24.150
of DM)	ŧ	SD	5.060	0.578	0.965	1.379	1.060	2.493	1.509	1.482	2.320	1.066	1.722	1.508	6.207	3.531	1.111	2.800	1.471	1.358
ell Wall (%	Ambie	Mean	36.465	18.296	23.737	24.993	25.962	27.483	27.959	27.446	18.007	18.384	21.341	21.902	22.043	27.826	28.913	24.936	23.683	25.978
0	2	SD	3.038	0.874	1.837	0.320	1.058	2.412	2.589	2.518	1.159	1.657	2.051	2.069	3.960	2.800	2.614	1.685	1.003	1.202
	- Snov	Mean	41.422	19.947	27.366	24.907	28.573	29.341	29.004	30.108	18.601	19.684	24.439	25.030	26.284	29.923	30.796	30.813	27.825	29.522
	3	SD	5.126	1.144	0.950	2.352	2.583	1.903	2.527	2.651		1.894	1.363	2.492	1.212	2.568	2.878	2.779	1.230	1.114
	+ Sno	Mean	56.675	78.705	79.715	79.493	74.947	79.353	80.715	78.386		84.002	83.292	81.092	80.104	75.830	75.969	75.098	76.522	75.850
% of DM)	ŧ	SD	5.060	0.578	0.965	1.379	1.060	2.493	1.509	1.482	2.320	1.066	1.722	1.508	6.207	3.531	1.111	2.800	1.471	1.358
l Solubles (Ambie	Mean	63.535	81.704	76.263	75.007	74.038	72.517	72.041	72.554	81.993	81.616	78.659	78.098	77.957	72.174	71.087	75.064	76.317	74.022
Cel	2	SD	3.038	0.874	1.837	0.320	1.058	2.412	2.589	2.518	1.159	1.657	2.051	2.069	3.960	2.800	2.614	1.685	1.003	1.202
	- Snor	Mean	58.578	80.053	72.634	75.093	71.427	70.659	70.996	69.892	81.399	80.316	75.561	74.970	73.716	70.077	69.204	69.187	72.175	70.478
	M	SD	0.002	0.001	0.000	0.001	0.001	0.000	0.003	0.005	 	0.002	0.002	0.001	0.002	0.001	0.002	0.003	0.002	0.002
	+ Sno	Mean	0.007	0.004	0.002	0.001	0.002	0.002	0.005	0.006		0.006	0.003	0.004	0.005	0.005	0.007	0.006	0.006	0.007
(MD g/	iient	SD	0.001	0.041	0.001	0.002	0.001	0.003	0.002	0.003	0.001	0.000	0.002	0.001	0.002	0.001	0.002	0.001	0.002	0.001
Ash (g/	Amb	Mean	0.005	0.021	0.002	0.003	0.002	0.003	0.005	0.005	0.005	0.003	0.003	0.004	0.005	0.004	0.005	0.006	0.004	0.007
	MOU	SD	0.002	0.006	0.000	0.001	0.001	0.001	0.002	0.001	0.002	0.003	0.003	0.002	0.002	0.000	0.001	0.003	0.003	0.003
	<u>s</u> -	Mean	0.005	0.000	0.003	0.002	0.002	0.002	0.005	0.004	0.004	0.005	0.004	0.004	0.006	0.004	0.006	0.006	0.006	0.008
Jana	DOY		170	177	184	191	198	205	212	219	173	180	187	194	200	210	217	223	231	237
B. I	Year		2017								2018									

B. no	na			NDF diges	stibility (%)					NDS digest	tibility (%)		
Year	DOY	- Sn	wor	Amt	bient	+ Sr	wor	- Sn	MO	Amb	ient	+ Sr	MOL
		Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD
2017	170	4.252	0.677	3.151	0.653	4.382	0.742	41.708	2.524	45.827	4.205	40.127	4.260
	177	3.489	0.373	7.818	10.743	3.497	0.322	59.554	0.727	60.926	0.480	58.434	0.951
	184	5.370	0.614	4.583	0.297	4.460	0.427	53.389	1.526	56.405	0.802	59.273	0.790
	191	5.092	0.516	4.776	0.241	4.701	0.278	55.432	0.266	55.360	1.146	59.089	1.955
	198	6.088	0.354	5.490	0.311	5.403	0.928	52.386	0.879	54.555	0.881	55.311	2.147
	205	5.509	0.711	5.336	0.671	4.908	0.236	51.747	2.004	53.292	2.072	58.973	1.582
	212	5.956	0.363	5.109	0.417	5.497	0.792	52.027	2.151	52.896	1.254	60.104	2.100
	219	5.627	0.666	5.491	0.886	6.136	0.781	51.110	2.092	53.322	1.231	58.169	2.203
2018	173	3.356	0.244	2.939	0.193			60.672	0.963	61.166	1.928		
	180	3.673	0.763	3.248	0.416	3.428	0.392	59.772	1.377	60.852	0.886	62.836	1.574
	187	4.550	0.635	3.729	0.680	3.510	0.510	55.821	1.705	58.395	1.431	62.246	1.133
	194	5.581	0.439	4.834	0.205	4.184	0.324	55.330	1.719	57.929	1.253	60.417	2.071
	200	4.896	0.449	3.932	0.890	5.053	0.563	54.288	3.290	57.812	5.158	59.597	1.007
	210	5.699	0.748	6.473	0.611	6.050	0.396	51.264	2.327	53.006	2.934	56.045	2.134
	217	5.507	0.429	4.898	0.207	5.871	0.390	50.539	2.173	52.103	0.924	56.160	2.392
	223	5.074	0.366	5.323	0.679	5.896	0.468	50.524	1.400	55.408	2.327	55.437	2.309
	231	5.005	0.501	4.979	0.613	6.032	0.654	53.008	0.833	56.449	1.222	56.620	1.022
	237	5.167	0.268	4.861	0.166	5.625	0.300	51.598	0.999	54.542	1.128	56.061	0.926

R. tomen	tosum			NDF (g	(MD g/					ADF (g/i	g DM)					Cellulose (g/g DM)					Lignin (g/	g DM)		
Year	роү	- Sn	MO	Amb	ient	+ Sr	NOI	- Snc	MC	Ambi	ent	+ Sno	w	- Sno	3	Ambi	ent	+ Sno	Ň	- Snơ	ş	Ambié	ant	+ Sno	х
		Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD
2017	170	0.443	0.029	0.441	0.041	0.431	0.016	0.315	0.022	0.334	0:050	0.319	0.020	0.142	0.014	0.143	0.010	0.146	0.011	0.169	0.026	0.186	0.048	0.168	0.011
	177	0.463	0.019	0.405	0.013	0.452	0.021	0.312	0.019	0.267	0.011	0.297	0.020	0.119	0.008	0.109	0.005	0.112	0.007	0.188	0.018	0.154	0.009	0.180	0.014
	184	0.391	0.024	0.405	0.016	0.380	0.025	0.275	0.014	0.285	0.008	0.270	0.022	0.113	0.007	0.123	600.0	0.111	0.011	0.157	0.010	0.158	0.005	0.154	0.013
	191	0.381	0.014	0.365	0.007	0.364	0.012	0.273	0.012	0.264	0.006	0.263	0.009	0.112	0.009	0.113	0.004	0.114	600.0	0.157	0.008	0.148	0.005	0.145	0.009
	198	0.459	0.024	0.471	0.034	0.498	0.010	0.328	0.011	0.336	0.029	0.366	0.013	0.122	0.003	0.141	0.014	0.150	0.003	0.202	0.011	0.191	0.017	0.211	0.013
	205	0.438	0:030	0.425	0.041	0.420	0.028	0.318	0.019	0.310	0.028	0.310	0.020	0.120	0.012	0.121	0.008	0.120	0.005	0.195	0.029	0.185	0.024	0.186	0.021
	212	0.438	0.020	0.493	0.033	0.493	0.024	0.315	0.012	0.354	0.022	0.366	0.022	0.129	0.012	0.140	0.012	0.138	0.007	0.180	0.008	0.211	0.015	0.223	0.022
	219	0.476	0.027	0.453	0.039	0.447	0.007	0.345	0.017	0.331	0.034	0.327	0.007	0.144	0.014	0.147	0.013	0.139	0.007	0.195	0.006	0.173	0.028	0.179	0.010
2018	173	0.365	0.012	0.366	0.012			0.253	0.006	0.258	0.014			0.117	0.008	0.114	600.0			0.128	0.007	0.136	0.008		
	180	0.370	0.016	0.396	0.015	0.432	0.013	0.276	0.008	0.292	0.012	0.320	600.0	0.123	0.006	0.136	600.0	0.149	0.005	0.145	0.010	0.147	0.008	0.163	0.007
	187	0.376	0.034	0.410	0.021	0.400	0.022	0.282	0.026	0.294	0.010	0.286	0.020	0.123	0.012	0.129	0.007	0.127	0.010	0.154	0.019	0.158	0.005	0.152	0.013
	194	0.418	0.023	0.369	0.009	0.391	0.020	0.308	0.022	0.277	0.011	0.288	0.013	0.122	0.014	0.122	0.005	0.130	0.010	0.181	0.014	0.149	0.006	0.151	0.004
	200	0.467	0.034	0.437	0.020	0.359	0.030	0.354	0.025	0.328	0.010	0.263	0.019	0.141	0.015	0.149	0.007	0.119	600.0	0.204	0.015	0.170	0.005	0.136	0.010
	210	0.469	0.013	0.464	0.028	0.382	0.018	0.350	0.007	0.347	0.021	0.294	0.020	0.143	0.010	0.149	0.010	0.128	0.006	0.200	0.009	0.192	0.017	0.159	0.016
	217	0.442	0.016	0.433	0.014	0.379	0.023	0.324	0.011	0.319	600.0	0.291	0.015	0.137	0.010	0.135	0.005	0.121	600.0	0.178	0.011	0.175	0.009	0.162	0.007
	223	0.469	0.023	0.393	0.029	0.378	0.022	0.347	0.020	0.293	0.020	0.279	0.011	0.145	0.005	0.121	0.008	0.124	600.0	0.193	0.019	0.163	0.012	0.147	0.010
	231	0.394	0.027	0.372	0.012	0.386	0.021	0.292	0.022	0.275	600.0	0.285	0.014	0.120	0.008	0.117	600.0	0.121	0.014	0.162	0.014	0.150	0.007	0.157	0.008
	237	0.381	0.015	0.336	0.026	0:330	0.017	0.271	0.012	0.247	0.017	0.247	0.012	0.107	0.007	0.102	0.009	660.0	0.006	0.154	600.0	0.137	0.011	0.141	0.007

R. tomer	ntosum			Ash (g	/g DM)					cell Solubles	(% of DM	_			-	Cell Wall (%	of DM)				-	Lig/Cut (% o	of NDF)		
Year	DOY	- Sn	wot	Amk	bient	+ Sn	wo	- Sn	M	Amb	ent	+ Sn	MO	- Sno	3	Ambi	ant	+ Sno	ş	- Snov	z	Ambie	ŧ	+ Sno	3
		Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD
2017	170	0.004	0.001	0.005	0.001	0.004	0.001	55.710	2.931	55.925	4.059	56.872	1.585	44.290	2.931	44.075	4.059	43.128	1.585	37.987	3.949	41.820	6.595	38.974	1.267
	177	0.005	0.001	0.004	0.001	0.005	0.001	53.716	1.909	59.453	1.320	54.796	2.128	46.284	1.909	40.547	1.320	45.204	2.128	40.690	3.107	38.002	1.251	39.754	1.724
	184	0.005	0.000	0.005	0.001	0.005	0.001	60.934	2.372	59.526	1.640	62.015	2.459	39.066	2.372	40.474	1.640	37.985	2.459	40.205	1.851	39.054	2.099	40.584	1.735
	191	0.005	0.003	0.003	0.002	0.004	0.000	61.908	1.369	63.479	0.745	63.633	1.207	38.092	1.369	36.521	0.745	36.367	1.207	41.232	0.974	40.648	1.173	39.907	2.112
	198	0.004	0.001	0.004	0.001	0.005	0.001	54.148	2.398	52.853	3.406	50.238	0.959	45.852	2.398	47.147	3.406	49.762	0.959	44.167	1.555	40.446	2.105	42.488	2.336
	205	0.003	0.001	0.004	0.001	0.004	0.001	56.202	2.977	57.460	4.055	58.031	2.751	43.798	2.977	42.540	4.055	41.969	2.751	44.257	4.095	43.358	1.606	44.153	2.275
	212	0.005	0.002	0.003	0.002	0.005	0.001	56.239	2.049	50.650	3.322	50.703	2.377	43.761	2.049	49.350	3.322	49.297	2.377	41.210	1.404	42.755	2.176	45.173	2.872
	219	0.006	0.001	0.011	0.004	0.009	0.002	52.355	2.720	54.735	3.931	55.273	0.735	47.645	2.720	45.265	3.931	44.727	0.735	41.016	1.672	38.086	3.817	40.050	1.913
2018	173	0.007	0.001	0.008	0.001			63.476	1.234	63.407	1.178			36.524	1.234	36.593	1.178			35.179	2.190	37.025	1.821		
	180	0.00	0.003	0.008	0.002	0.009	0.002	62.976	1.588	60.414	1.458	56.762	1.300	37.024	1.588	39.586	1.458	43.238	1.300	39.137	1.428	37.217	1.741	37.579	0.941
	187	0.005	0.002	0.007	0.001	0.006	0.002	62.422	3.446	58.973	2.056	59.954	2.164	37.578	3.446	41.027	2.056	40.046	2.164	40.982	2.314	38.514	0.948	37.941	1.505
	194	0.005	0.001	0.006	0.000	0.007	0.002	58.171	2.290	63.090	0.919	60.870	2.011	41.829	2.290	36.910	0.919	39.130	2.011	43.174	2.637	40.264	0.870	38.765	2.226
	200	0.00	0.001	0.008	0.002	0.008	0.001	53.332	3.435	56.320	1.956	64.060	3.042	46.668	3.435	43.680	1.956	35.940	3.042	43.738	1.642	39.031	1.345	37.920	0.910
	210	0.007	0.000	0.006	0.002	0.007	0.001	53.071	1.261	53.617	2.809	61.806	1.798	46.929	1.261	46.383	2.809	38.194	1.798	42.654	2.131	41.314	2.749	41.654	2.694
	217	0.008	0.004	0.009	0.003	0.008	0.001	55.835	1.577	56.745	1.414	62.103	2.285	44.165	1.577	43.255	1.414	37.897	2.285	40.401	2.387	40.527	1.309	42.867	0.905
	223	0.00	0.002	0.009	0.001	0.009	0.001	53.144	2.324	60.703	2.860	62.208	2.216	46.856	2.324	39.297	2.860	37.792	2.216	41.190	2.310	41.564	0.754	38.925	2.314
	231	0.00	0.004	0.008	0.003	0.007	0.002	60.579	2.707	62.769	1.160	61.355	2.096	39.421	2.707	37.231	1.160	38.645	2.096	41.044	0.827	40.209	1.680	40.720	2.814
	237	0.010	0.004	0.008	0.002	0.007	0.003	61.860	1.532	66.446	2.575	66.963	1.722	38.140	1.532	33.554	2.575	33.037	1.722	40.255	1.437	40.681	1.003	42.732	1.754

				NDF digest	ibility (%)					NDS digest	ibility (%)		
R. tome	ntosum			5									
Year	РО	- Sn	MO	Amb	ient	us +	MOL	- Snc	M	Ambi	ient	+ Sne	Ň
		Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD
2017	170	7.429	1.192	6.266	1.284	6.866	0.275	39.325	2.436	39.504	3.373	40.291	1.317
	177	6.860	0.837	6.747	0.341	6.955	0.489	37.668	1.587	42.436	1.097	38.565	1.769
	184	5.906	0.694	6.460	0.894	5.633	0.534	43.666	1.971	42.496	1.363	44.564	2.044
	191	5.476	0.199	5.398	0.325	5.570	0.564	44.476	1.138	45.781	0.619	45.909	1.003
	198	5.791	0.578	7.047	0.829	6.789	0.703	38.027	1.993	36.951	2.831	34.777	0.797
	205	5.526	0.801	5.545	0.350	5.292	0.388	39.734	2.474	40.779	3.370	41.254	2.286
	212	6.314	0.596	6.661	0.909	5.955	0.628	39.765	1.703	35.121	2.761	35.164	1.976
	219	6.952	0.870	7.536	1.007	6.801	0.571	36.537	2.261	38.514	3.267	38.962	0.611
2018	173	6.931	0.751	6.375	0.533			45.778	1.026	45.721	0.979		
	180	5.853	0.323	6.841	0.621	7.334	0.357	45.363	1.319	43.234	1.212	40.199	1.080
	187	5.474	0.599	6.682	0.599	6.678	0.348	44.903	2.863	42.037	1.709	42.852	1.798
	194	5.542	0.706	5.543	0.151	6.334	0.879	41.370	1.903	45.458	0.763	43.613	1.671
	200	6.013	0.728	6.956	0.683	6.011	0.681	37.349	2.855	39.832	1.625	46.264	2.528
	210	6.356	0.700	6.681	0.884	5.405	0.541	37.132	1.048	37.586	2.334	44.390	1.494
	217	6.628	0.813	6.426	0.383	5.073	0.492	39.429	1.310	40.185	1.175	44.637	1.899
	223	6.756	0.481	5.570	0.466	6.065	0.786	37.193	1.931	43.475	2.376	44.725	1.841
	231	5.708	0.229	5.619	0.461	5.742	0.915	43.371	2.249	45.191	0.964	44.016	1.742
	237	5.737	0.393	4.949	0.439	4.457	0.502	44.435	1.273	48.247	2.140	48.676	1.431

V. vitis-ia	taea			NDF (g/	g DM)					ADF (g/£	(MU)					Cellulose (g/g DM)					Lignin (g/	(MU)		
Year	УОД	- Snc	MO	Ambi	ient	+ Sn	MO	- Snc	M	Ambie	'nt	+ Sno	3	- Snov	×	Ambie	at	+ Sno	3	- Sno	2	Ambie	ŧ	+ Sno	х
		Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD
2017	170	0.383	0.024	0.395	0.026	0.365	0.034	0.246	0.018	0.260	0.035	0.272	0.041	0.126	0.00	0.135	0.016	0.140	0.006	0.116	0.015	0.119	0.020	0.127	0.041
	177	0.389	0.029	0.381	0.059	0.403	0.056	0.233	0.024	0.239	0.033	0.262	0.045	0.124	0.010	0.124	0.013	0.133	0.008	0.104	0.016	0.111	0.023	0.125	0.042
	184	0.336	0.019	0.334	0.008	0.391	0.189	0.222	0.010	0.212	0.004	0.205	0.008	0.131	0.007	0.123	0.006	0.117	0.006	0.079	0.022	0.085	0.006	0.085	0.007
	191	0.322	0.016	0.333	0.015	0.332	0.007	0.215	0.016	0.226	0.010	0.226	0.006	0.121	0.012	0.127	0.005	0.126	0.004	0.091	0.005	0.097	0.008	0.097	0.005
	198	0.359	0.024	0.361	0.015	0.366	0.026	0.243	0.025	0.248	0.014	0.252	0:030	0.128	0.008	0.134	0.007	0.132	0.011	0.112	0.018	0.110	0.007	0.117	0.019
	205	0.371	0.012	0.335	0.019	0.346	0.019	0.261	0.014	0.232	0.012	0.243	0.016	0.137	0.012	0.115	0.018	0.128	0.008	0.120	0.004	0.113	0.015	0.113	0.011
	212	0.366	0.039	0.344	0.018	0.360	0.035	0.260	0.033	0.235	0.010	0.256	0.035	0.135	0.00	0.130	0.005	0.125	0.019	0.119	0.026	0.102	0.007	0.126	0.025
	219	0.360	0.026	0.383	0:030	0.342	0.015	0.256	0.021	0.274	0.032	0.239	0.013 (0.141	0.010	0.136	0.011	0.133	600.0	0.106	0.011	0.131	0.022	0.100	0.005
2018	173	0.295	0.007	0.290	0.005			0.201	0.007	0.205	0.009			0.123	0.008	0.119	0.007			0.072	0.008	0.080	0.003		
	180	0.317	0:030	0.285	0.011	0.306	0.012	0.227	0.017	0.203	600.0	0.220	0.006	0.136	0.011	0.118	0.007	0.130	0.003	0.085	0.009	0.077	0.005	0.084	0.004
	187	0.304	0.009	0.296	0.011	0.304	0.026	0.217	0.010	0.204	0.009	0.211	0.018	0.123	0.001	0.115	0.006	0.122	0.015	0.089	0.012	0.082	0.003	0.084	0.006
	194	0.332	0.027	0.267	0.106	0.297	0.016	0.237	0.017	0.226	0.013	0.216	0.010	0.134	0.013	0.127	0.011	0.119	0.020	0.097	0.008	0.092	0.006	0.091	0.011
	200	0.336	0.011	0.335	0.021	0.306	0.020	0.246	0.008	0.236	0.013	0.217	0.010	0.137	0.010	0.135	0.007	0.122	0.011	0.103	0.009	0.095	0.008	0.087	0.003
	210	0.405	0.047	0.353	0.025	0.364	0.021	0.301	0.043	0.259	0.017	0.271	0.021	0.157	0.013	0.143	0.011	0.149	0.012	0.138	0.034	0.110	0.017	0.115	0.011
	217	0.350	0.009	0.361	0.010	0.351	0.019	0.252	600.0	0.260	0.007	0.258	0.010	0.138	0.002	0.144	0.010	0.146	0.008	0.107	0.010	0.109	600.0	0.105	0.003
	223	0.340	0.024	0.329	0.012	0.335	0.006	0.253	0.018	0.237	0.007	0.244	0.006	0.134	0.00	0.133	0.006	0.134	0.006	0.111	0.010	0.098	0.005	0.101	0.003
	231	0.326	0.009	0.324	0.013	0.312	0.019	0.235	0.006	0.235	0.008	0.229	0.012	0.129	0.008	0.130	0.005	0.122	0.008	0.098	0.007	660.0	0.006	0.100	0.004
	237	0.310	0.014	0.307	0.011	0.303	0.017	0.224	0.009	0.224	600.0	0.221	0.011 (0.116	0.010	0.120	0.007	0.117	0.011	0.101	0.003	0.096	0.005	0.097	0.004
	MOL	SD	7.298	5.679	7.474	1.645	3.605	1.967	4.790	0.659		1.167	1.341	5.425	2.560	1.437	1.368	1.261	0.980	2.117					
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	S+	Mean	34.360	30.469	24.577	29.102	31.900	32.502	34.870	29.195		27.364	27.554	30.819	28.626	31.537	29.831	30.265	32.027	31.984					
of NDF)	ent	SD	4.659	2.602	1.261	1.756	1.195	4.877	1.582	3.478	1.039	1.725	0.937	39.349	1.311	3.458	2.639	1.638	1.500	2.042					
Lig/Cut (%	Ambi	Mean	30.256	28.940	25.447	29.079	30.429	33.744	29.611	33.908	27.667	27.116	27.772	47.073	28.278	31.118	30.289	29.640	30.660	31.348					
	3	SD	2.433	3.305	5.985	0.312	3.233	0.848	3.933	1.065	2.607	1.068	3.259	2.641	2.377	4.562	2.130	1.852	2.446	1.421					
	- Sno	Mean	30.286	26.683	23.442	28.206	31.057	32.254	32.248	29.499	24.510	26.674	29.309	29.405	30.725	33.746	30.592	32.495	30.052	32.547					
	Ň	SD	3.415	5.576	18.888	0.679	2.593	1.921	3.524	1.482		1.230	2.554	1.598	1.961	2.088	1.878	0.606	1.892	1.675					
	+ Sno	Mean	36.541	40.319	39.066	33.196	36.582	34.644	36.000	34.180		30.623	30.381	29.679	30.621	36.437	35.137	33.511	31.159	30.346					
of DM)	t	SD	2.631	5.865	0.833	1.491	1.533	1.930	1.768	2.993	0.534	1.068	1.138	10.629	2.128	2.471	1.012	1.172	1.292	1.105					
cell Wall (%	Ambie	Mean	39.450	38.144	33.404	33.326	36.150	33.490	34.427	38.317	29.044	28.467	29.563	26.689	33.481	35.278	36.086	32.946	32.409	30.687					
-		SD	2.425	2.857	1.871	1.618	2.380	1.220	3.935	2.555	0.736	2.959	0.919	2.699	1.068	4.731	0.897	2.422	0.912	1.374					
	- Snow	Mean	38.287	38.888	33.639	32.250	35.863	37.139	36.581	36.036	29.523	31.709	30.435	33.172	33.608	40.501	35.027	33.994	32.637	30.975					
		SD	3.415	5.576	8.888	0.679	2.593	1.921	3.524	1.482	 	1.230	2.554	1.598	1.961	2.088	1.878	0.606	1.892	1.675					
	+ Snov	Aean	3.459	9.681	0.934	6.804	3.418	5.356	4.000	5.820		9.377	9.619	0.321	9.379	3.563	4.863	6.489	8.841	9.654					
6 of DM)		SD	2.631 6	5.865 5	0.833 6	l.491 6	l.533 6	l.930 6	l. 768 6	2.993 6).534	l.068 6	l.138 6	.629 7	2.128 6	2.471 6	l.012 6	l.172 6	l. 292 6	1.105 6					
Solubles (9	Ambien	lean	.550	.856	.596	674	8.850	6.510	573	.683	.956	.533	.437	311 10	6.519	1.722	3.914	.054	.591	.313					
Cell		SD	.425 60	.857 61	.871 66	.618 66	380 63	.220 66	935 65	.555 61	.736 70	959 71	919 70	669 73	.068 66	.731 64	897 63	422 67	.912 67	374 69					
	- Snow	1ean	1.713 2	1.112 2	5.361 1	7.750 1	4.137 2	2.861 1	3.419 3	3.964 2	0.477 0	3.291 2	9.565 0	5.828 2	5.392 1	9.499 4	4.973 0	5.006 2	7.363 0	9.025 1					
		SD	001 6	001 6	000	.001 6	.000	.000	.003 6.	.002 6:	 7	.001 6	.002 6	.001 6	.002 6	.001 5	.002 6.	.002 6	.003 6.	.001 6					
	+ Snow	dean	0.004 0	0.004 0	0.003	0.003	0.003	0.002 0	0.005 0	0.007 0		0.006 0	0.005 0	0.007 0	0.008	0.006 0	0.008	0 600.0	0.007 0	0.008 0					
DM)	ŧ	SD	0.001	0.001	0.001	0.001 (0.004 (0.002	0.001	0.003	0.001	0.003	0.001	0.001	0.002	0.001	0.001	0.001	0.002	0.001					
Ash (g/g	Ambie	Mean	0.005	0.004	0.004	0.003	0.004	0.004	0.003	0.008	0.005	0.008	0.006	0.006	0.006	0.006	0.007	0.007	0.006	0.008					
	3	SD	0.001	0.001	0.018	0.001	0.001	0.001	0.004	0.002	0.000	0.001	0.001	0.001	0.002	0.001	0.001	0.003	0.003	0.002					
	- Sno	Mean	0.004	0.005	0.012	0.003	0.003	0.004	0.006	0.008	0.006	0.007	0.005	0.005	0.006	0.005	0.007	0.008	0.008	0.007					
idaea	роу		170	177	184	191	198	205	212	219	173	180	187	194	200	210	217	223	231	237					
V. vitis-i	Year		2017								2018														

V. vitis-	-idaea			NDF diges	tibility (%)					NDS diges	tibility (%)		
Year	РО	- Snc	M	Amb	ient		MOL	- Sn	M	Ambi	ent	+ sn	MO
		Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD
2017	170	9.021	0.624	9.476	2.154	7.301	1.608	44.313	2.015	43.347	2.187	45.764	2.838
	177	10.846	1.539	9.562	1.570	9.454	1.561	43.814	2.374	44.432	4.874	42.625	4.634
	184	11.098	3.333	9.789	0.346	14.379	13.987	48.176	1.555	48.371	0.692	43.667	15.696
	191	8.342	0.385	8.302	0.645	8.268	0.672	49.331	1.345	48.436	1.239	48.544	0.564
	198	8.177	0.838	8.468	0.559	8.056	1.086	46.328	1.978	46.090	1.274	45.731	2.155
	205	8.012	0.466	6.893	1.516	7.398	0.673	45.268	1.014	48.300	1.604	47.341	1.596
	212	7.874	0.678	8.381	0.796	6.980	1.317	45.731	3.270	47.521	1.469	46.214	2.929
	219	8.782	0.351	7.677	0.759	8.458	0.412	46.184	2.123	44.289	2.487	47.726	1.231
2018	173	9.073	1.102	7.707	0.417			51.596	0.612	51.994	0.443		
	180	8.793	0.838	7.763	0.769	8.242	0.608	49.779	2.459	52.474	0.887	50.682	1.022
	187	7.536	0.898	7.810	0.552	8.124	1.054	50.838	0.763	51.563	0.945	50.884	2.123
	194	8.207	1.360	6.150	3.453	7.032	1.765	48.564	2.243	53.951	8.833	51.467	1.328
	200	7.796	0.851	8.641	0.685	7.855	1.307	48.202	0.888	48.307	1.768	50.684	1.630
	210	8.152	0.845	8.057	1.192	8.112	0.457	42.474	3.931	46.814	2.054	45.851	1.735
	217	8.152	0.583	8.552	1.038	8.480	0.973	47.022	0.745	46.142	0.841	46.931	1.561
	223	7.262	0.727	8.014	0.727	7.915	0.550	47.881	2.013	48.752	0.974	48.282	0.504
	231	7.813	0.924	7.520	0.576	6.803	0.703	49.009	0.758	49.198	1.073	50.237	1.572
	237	6.611	0.656	6.921	0.764	6.669	1.022	50.389	1.141	50.629	0.918	50.913	1.392

C. bige	lowii			NDF (g	(MD g/					ADF (g/g	(MD					Cellulose (g/g DM)					Lignin (g/§	g DM)		
Year	ЛОД	- Sn	MO	Amb	ient	+ Sn	MO	- Snc	MC	Ambie	ut.	+ Sno	Ň	- Snov	3	Ambié	ent	+ Sno	w	- Snơ	3	Ambie	ŧ	+ Sno	3
		Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD
2017	170	0.599	0.030	0.647	0.046	0.635	0.008	0.257	0.016	0.300	0.057	0.291	0.017	0.230	0.00	0.241	0.020	0.243	0.008	0.007	0.047	0.057	0.051	0.046	0.012
	177	0.631	0.014	0.614	0.010	0.657	0.014	0.256	0.011	0.243	0.011	0.265	0.010	0.175	0.078	0.188	0.079	0.184	060.0	0.079	0.077	0.048	0.072	0.080	0.081
	184	0.682	0.019	0.704	0.033	0.701	0.019	0.277	0.003	0.302	0.027	0.284	0.011	0.247	0.005	0.246	0.043	0.238	0.025	0.028	0.005	0.054	0.043	0.044	0.021
	191	0.611	0.031	0.644	0.013	0.653	0.025	0.247	0.015	0.280	0.013	0.281	0.023	0.198	0.050	0.246	0.010	0.245	0.016	0.047	0.045	0.032	0.003	0.035	0.011
	198	0.655	0.016	0.655	0.026	0.663	0.026	0.271	0.019	0.290	0.026	0.274	0.020	0.217	0.037	0.253	0.020	0.227	0.010	0.053	0.049	0.035	0.007	0.045	0.024
	205	0.642	0.015	0.648	0.044	0.664	0.019	0.268	0.011	0.288	0.044	0.285	0.029	0.223	0.027	0.251	0.036	0.234	0.047	0.042	0.020	0.037	0.010	0.051	0.020
	212	0.601	0.008	0.635	0.021	0.628	0.015	0.236	0.010	0.266	0.023	0.250	0.018	0.187	0.051	0.231	0.024	0.217	0.020	0.046	0.048	0.032	0.003	0.028	0.007
	219	0.614	0.015	0.614	0.019	0.631	0.013	0.262	0.017	0.264	0.012	0.271	0.010	0.226	0.019	0.229	0.006	0.235	600.0	0.032	0.003	0.032	0.007	0.031	0.004
2018	173	0.607	0.018	0.586	0.015			0.262	0.011	0.240	0.009			0.232	0.030	0.214	0.020			0.025	0.028	0.022	0.024		
	180	0.638	0.018	0.620	0.009	0.617	0.016	0.262	0.019	0.255	0.008	0.256	0.006	0.242	0.014	0.235	600.0	0.233	0.010	0.016	0.005	0.015	0.008	0.017	0.011
	187	0.674	0.013	0.662	0.008	0.646	0.018	0.288	0.006	0.286	0.018	0.268	0.014	0.255	0.010	0.243	0.032	0.240	0.014	0.029	0.009	0.040	0.021	0.024	0.013
	194	0.638	0.008	0.640	0.012	0.644	0.006	0.273	0.012	0.278	0.010	0.271	0.006	0.237	0.007	0.250	0.011	0.197	0.064	0.032	0.012	0.025	0.005	0.069	0.061
	200	0.664	0.009	0.647	0.015	0.653	0.015	0.322	0.012	0.289	0.015	0.285	0.016	0.277	0.024	0.255	0.020	0.262	0.012	0.040	0.018	0.030	0.007	0.017	0.005
	210	0.673	0.036	0.639	0.032	0.639	0.014	0.286	0.021	0.279	0.015	0.277	0.012	0.252	0.019	0.235	0.020	0.244	0.012	0.029	0.009	0.040	0.028	0.028	0.002
	217	0.615	0.021	0.630	0.009	0.624	0.018	0.275	0.009	0.274	0.007	0.263	0.012	0.232	0.020	0.242	0.005	0.233	0.010	0.038	0.018	0.026	0.003	0.024	0.004
	223	0.595	0.022	0.580	0.020	0.575	0.023	0.272	0.016	0.268	0.016	0.258	0.016	0.241	0.013	0.229	0.028	0.200	0.059	0.025	0.004	0.034	0.017	0.051	0.049
	231	0.585	0.018	0.630	0.022	0.559	0.015	0.268	0.014	0.312	0.019	0.243	0.013	0.233	0.018	0.219	0.052	0.220	0.015	0.028	0.007	0.087	0.045	0.018	0.003
	237	0.616	0.021	0.630	0.052	0.558	0.008	0.296	0.025	0.324	0.042	0.256	0.012	0.218	0.074	0.284	0.034	0.218	0.029	0.072	0.056	0.034	0.011	0.030	0.029

	wo	SD	1.795	12.614	2.864	1.503	3.382	3.128	1.099	0.599		1.720	2.029	9.446	0.684	0.475	0.610	8.855	0.541	5.174
	+ Sn	Mean	7.196	12.255	6.178	5.283	6.755	7.715	4.438	4.997		2.790	3.737	10.684	2.663	4.414	3.904	8.917	3.187	5.388
of NDF)	ent	SD	6.755	11.907	5.755	0.315	0.876	1.139	0.477	1.061	3.874	1.240	3.214	0.872	1.283	4.187	0.457	3.040	7.213	1.358
Lig/Cut (%	Ambi	Mean	8.428	7.852	7.549	4.898	5.378	5.609	4.968	5.147	3.748	2.394	6.021	3.876	4.624	6.117	4.166	5.929	13.973	5.389
	3	SD	7.973	12.195	0.708	7.526	7.308	2.972	7.766	0.468	4.697	0.714	1.372	1.979	2.848	1.264	2.884	0.589	1.180	9.483
	- Sno	Mean	0.985	12.469	4.037	7.815	8.050	6.578	7.521	5.276	4.155	2.464	4.265	4.997	6.029	4.348	6.193	4.186	4.801	11.793
	Ň	SD	0.824	1.444	1.922	2.500	2.567	1.868	1.546	1.281	 	1.635	1.824	0.601	1.460	1.449	1.805	2.263	1.463	0.823
	+ Sno	Mean	63.450	65.686	70.126	65.302	66.296	66.424	62.757	63.059		61.657	64.625	64.394	65.273	63.913	62.368	57.468	55.884	55.759
% of DM)	ient	SD	4.604	0.972	3.315	1.289	2.563	4.417	2.098	1.922	1.513	0.868	0.822	1.164	1.522	3.184	0.928	1.972	2.181	5.155
Cell Wall (Amb	Mean	64.695	61.362	70.360	64.379	65.536	64.788	63.455	61.353	58.580	62.001	66.233	63.999	64.698	63.914	63.026	58.021	62.974	63.046
	wo	SD	2.983	1.420	1.946	3.055	1.612	1.539	0.761	1.454	1.848	1.774	1.291	0.763	0.926	3.561	2.077	2.243	1.775	2.101
	- Sn	Mean	59.900	63.119	68.250	61.130	65.509	64.172	60.065	61.377	60.698	63.839	67.430	63.843	66.398	67.304	61.487	59.455	58.453	61.642
	wot	SD	0.824	1.444	1.922	2.500	2.567	1.868	1.546	1.281		1.635	1.824	0.601	1.460	1.449	1.805	2.263	1.463	0.823
-	+ Sr	Mean	36.550	34.314	29.874	34.698	33.704	33.576	37.243	36.941		38.343	35.375	35.606	34.727	36.087	37.632	42.532	44.116	44.241
is (% of DM	pient	SD	4.604	0.972	3.315	1.289	2.563	4.417	2.098	1.922	1.513	0.868	0.822	1.164	1.522	3.184	0.928	1.972	2.181	5.155
Cell Soluble	Amt	Mean	35.305	38.638	29.640	35.621	34.464	35.212	36.545	38.647	41.420	37.999	33.767	36.001	35.302	36.086	36.974	41.979	37.026	36.954
	MOL	SD	2.983	1.420	1.946	3.055	1.612	1.539	0.761	1.454	1.848	1.774	1.291	0.763	0.926	3.561	2.077	2.243	1.775	2.101
	-s	Mean	40.100	36.881	31.750	38.870	34.491	35.828	39.935	38.623	39.302	36.161	32.570	36.157	33.602	32.696	38.513	40.545	41.547	38.358
	wous	SD	0.000	0.000	0.000	0.001	0.002	0.001	0.003	0.001		0.001	0.001	0.001	0.002	0.001	0.002	0.002	0.001	0.002
	+	Mean	0.002	0.002	0.002	0.001	0.002	0.001	0.005	0.004		0.006	0.004	0.005	0.006	0.004	0.005	0.007	0.005	0.007
(g/g DM)	nbient	SD	0.001	0.008	0.000	0.003	0.000	0.000	0.002	0.002	0.001	0.002	0.002	0.001	0.001	0.001	0.002	0.002	0.002	0.003
Ash (An	Mean	0.003	0.006	0.002	0.003	0.001	0.001	0.004	0.004	0.004	0.005	0.004	0.004	0.004	0.005	0.006	0.005	0.006	0.006
	Snow	SD	0.037	0.001	0.001	0.003	0.001	0.001	0.002	1 0.001	0.001	0.002	1 0.001	1 0.001	0.002	0.001	0.001	0.001	0.002	0.002
		Mean	0.020	0.002	0.002	0.002	0.001	0.002	0.004	0.004	0.005	0.005	0.004	0.004	0.005	0.004	0.005	0.006	0.007	0.005
bigelowii	r DO		7 170	17,	184	191	195	205	212	215	8 175	18(187	194	200	210	217	225	231	237
C. I	Yeâ		201								201									

C. bige	ilowii			NDF diges:	tibility (%)					NDS digest	ibility (%)		
Year	DOY	- Sr	wot	Amb	ient	+ Sr	MO	- Sne	MO	Amb	ient	+ Sn	MO
		Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD
2017	170	55.234	20.697	41.675	8.954	42.437	3.208	26.353	2.479	22.368	3.826	23.403	0.684
	177	37.000	17.122	43.586	16.326	39.120	17.709	23.678	1.180	25.138	0.808	21.545	1.200
	184	52.520	1.746	47.104	10.116	49.224	5.421	19.414	1.618	17.661	2.755	17.855	1.597
	191	41.359	11.469	47.643	0.461	47.513	2.213	25.331	2.539	22.631	1.071	21.864	2.078
	198	43.562	11.125	47.444	0.843	45.386	5.444	21.692	1.340	21.669	2.130	21.038	2.134
	205	44.290	5.177	46.390	2.080	43.740	6.865	22.803	1.278	22.292	3.671	20.931	1.552
	212	41.102	11.199	46.824	1.803	47.455	2.290	26.216	0.633	23.399	1.743	23.979	1.284
	219	44.676	1.764	44.902	1.409	46.484	1.735	25.126	1.208	25.145	1.597	23.728	1.065
2018	173	47.245	8.979	46.052	6.362			25.690	1.536	27.450	1.257		
	180	52.725	1.217	51.436	2.662	50.264	3.317	23.080	1.474	24.607	0.721	24.893	1.358
	187	51.465	3.819	47.009	6.663	50.532	4.234	20.096	1.073	21.091	0.683	22.426	1.51
	194	47.210	4.426	49.639	2.260	39.211	14.708	23.076	0.634	22.947	0.967	22.618	0.49
	200	47.029	6.157	48.592	3.740	53.440	1.470	20.953	0.769	22.366	1.265	21.888	1.21
	210	51.061	2.606	45.256	7.314	48.374	2.096	20.200	2.959	23.017	2.646	23.018	1.20
	217	43.186	5.319	48.211	0.378	48.286	1.700	25.034	1.726	23.755	0.771	24.302	1.500
	223	45.425	1.152	41.351	5.859	37.698	12.346	26.723	1.864	27.915	1.639	28.374	1.881
	231	43.517	2.920	32.436	11.286	44.697	1.792	27.555	1.475	23.798	1.813	29.690	1.216
	237	35.799	13.014	45.588	2.730	41.121	8.030	24.905	1.746	93,739	4.784	29.794	0.687

E. vagit	natum			NDF (g	(MD B/					ADF (g/.	g DM)					Cellulose (g/g DM)					Lignin (g/g	(MD)		
Year	ЛОД	us -	MO	Amb	vient	us +	MO	- Snc	MC	Ambi	ent	+ Snc	MC	- Sno	ž	Ambié	ent	+ Sno	3	- Snov	2	Ambier	ŧ	+ Sno	3
		Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD
2017	170	0.692	0.025	0.690	0.026	0.721	0.017	0.368	0.032	0.348	0.038	0.381	0.021	0.296	0.006	0.285	0.021	0.310	0.017	0.068	0.034	0.061	0.021	0.069	0.010
	177	0.663	0.012	0.679	0.022	0.682	0.007	0.298	0.018	0.307	0.012	0.310	0.006	0.264	0.018	0.271	0.009	0.271	0.014	0.031	0.017	0.033	0.008	0.036	0.015
	184	0.712	0.029	0.713	0.018	0.725	0.024	0.306	0.014	0.307	0.012	0.311	0.005	0.275	0.010	0.254	0.020	0.247	0.052	0.029	0.004	0.052	0.025	0.062	0.053
	191	0.676	0.021	0.683	0.021	0.645	0.020	0.305	0.014	0.300	0.024	0.263	0.013	0.278	0.012	0.222	0.073	0.217	0.014	0.025	0.005	0.078	0.064	0.043	0.016
	198	0.684	0.014	0.672	0.011	0.687	0.015	0.284	0.007	0.276	0.007	0.294	0.011	0.238	0.029	0.249	0.010	0.246	0.037	0.045	0.032	0.026	0.005	0.046	0.035
	205	0.720	0.033	0.711	0.013	0.700	0.008	0.313	0.018	0.304	0.013	0.314	0.006	0.243	0.024	0.218	0.082	0.217	0.097	0.069	0.034	0.085	0.075	960.0	0.094
	212	0.663	0.014	0.650	0.012	0.682	0.030	0.276	0.011	0.268	0.010	0.300	0.020	0.246	0.014	0.215	0.061	0.223	0.082	0.027	0.010	0.049	0.056	0.073	0.073
	219	0.699	0.015	0.705	0.014	0.671	0.009	0.344	0.031	0.349	0.016	0.296	0.007	0.290	0.044	0.294	0.027	0.213	0.042	0.049	0.018	0.051	0.038	0.079	0.038
2018	173	0.668	0.011	0.671	0.010			0.285	0.005	0.296	0.006			0.236	0.039	0.271	0.006			0.043	0.039	0.022	0.003		
	180	0.673	0.010	0.688	0.008	0.647	0.022	0.296	0.011	0.288	0.013	0.288	0.021	0.271	0.007	0.264	0.010	0.261	0.014	0.020	0.004	0.020	0.005	0.021	0.012
	187	0.686	0.021	0.673	0.017	0.665	0.006	0.302	0.018	0.294	0.014	0.286	0.007	0.273	0.014	0.241	0.047	0.260	600.0	0.025	0.005	0.049	0.045	0.023	0.010
	194	0.685	0.010	0.662	0.020	0.649	0.006	0.306	0.002	0.291	0.013	0.293	0.016	0.269	0.015	0.236	0.057	0.261	0.016	0.032	0.016	0.049	0.062	0.028	0.015
	200	0.701	0.034	0.682	0.011	0.677	0.007	0.306	0.020	0.304	600.0	0.303	0.011	0.262	0.006	0.273	0.011	0.271	0.023	0.039	0.015	0.026	0.004	0.028	0.018
	210	0.758	0.017	0.720	0.018	0.680	0.026	0.370	0.014	0.323	0.017	0.307	0.005	0.298	0.022	0.293	0.013	0.274	0.007	0.068	0.013	0.025	0.008	0.029	0.004
	217	0.724	0.020	0.701	0.017	0.668	0.019	0.338	0.006	0.305	0.011	0.297	0.011	0.252	0.065	0.190	0.055	0.200	0.051	0.080	0.065	0.108	0.049	0.091	0.054
	223	069.0	0.013	0.631	0.018	0.616	0.027	0.327	0.016	0.292	0.011	0.281	0.017	0.187	0.100	0.224	0.056	0.226	0.055	0.135	0.087	0.063	0.052	0.050	0.044
	231	0.660	0.022	0.622	0.022	0.594	0.019	0.300	0.009	0.282	0.009	0.273	0.006	0.220	0.036	0.228	0.040	0.238	0.027	0.075	0.039	0.050	0.041	0.025	0.017
	237	0.636	0.011	0.600	0.028	0.549	0.019	0.296	0.005	0.277	0.012	0.251	0.015	0.239	0.034	0.207	0.050	0.231	0.013	0.050	0.038	0.065	0.050	0.015	0.003

	MO	SD	1.360	2.204	7.039	2.486	4.980	13.393	11.442	5.784		1.752	1.451	2.202	2.592	0.562	7.858	7.762	2.724	0.576
	+Sn	Mean	9.532	5.222	8.415	6.733	6.622	13.663	10.972	11.785		3.190	3.463	4.255	4.176	4.237	13.451	8.346	4.121	2.710
of NDF)	ent	SD	2.720	1.080	3.308	9.469	0.827	10.841	8.939	5.209	0.354	0.652	6.636	9.244	0.565	1.065	7.023	8.483	6.388	8.194
Lig/Cut (%	Ambi	Mean	8.826	4.884	7.172	11.431	3.926	12.073	7.597	7.120	3.204	2.924	7.273	7.392	3.840	3.514	15.417	10.076	8.057	10.763
	ž	SD	4.524	2.570	0.363	0.672	4.500	4.239	1.580	2.665	5.632	0.665	0.646	2.344	1.913	1.765	9.139	12.911	5.660	5.791
	- Sno	Mean	9.730	4.620	4.110	3.671	6.446	9.410	4.016	7.047	6.392	2.970	3.624	4.692	5.476	8.924	11.105	19.757	11.295	7.870
	MO	SD	1.684	0.693	2.444	2.024	1.451	0.758	3.032	0.877		2.214	0.606	0.640	0.714	2.638	1.920	2.672	1.851	1.897
	+ Sn	Mean	72.105	68.205	72.532	64.511	68.728	70.027	68.219	67.144		64.738	66.477	64.882	67.677	67.963	66.796	61.588	59.427	54.908
% of DM)	ient	SD	2.584	2.202	1.816	2.070	1.093	1.313	1.187	1.426	0.961	0.812	1.673	2.029	1.094	1.845	1.669	1.775	2.216	2.844
Cell Wall (Amb	Mean	69.033	67.927	71.337	68.327	67.170	71.070	64.950	70.524	67.145	68.766	67.321	66.206	68.171	71.999	70.056	63.125	62.162	60.024
	wor	SD	2.521	1.181	2.923	2.111	1.356	3.343	1.395	1.513	1.135	0.959	2.124	1.017	3.439	1.714	2.032	1.255	2.160	1.102
	- Sr	Mean	69.232	66.261	71.241	67.614	68.443	71.961	66.290	69.853	66.762	67.325	68.581	68.536	70.051	75.845	72.395	68.961	65.957	63.605
	wot	SD	1.684	0.693	2.444	2.024	1.451	0.758	3.032	0.877		2.214	0.606	0.640	0.714	2.638	1.920	2.672	1.851	1.897
_	+ Sr	Mean	27.895	31.795	27.468	35.489	31.272	29.973	31.781	32.856		35.262	33.523	35.118	32.323	32.037	33.204	38.412	40.573	45.092
s (% of DM	ient	SD	2.584	2.202	1.816	2.070	1.093	1.313	1.187	1.426	0.961	0.812	1.673	2.029	1.094	1.845	1.669	1.775	2.216	2.844
ell Soluble	Amb	Mean	30.967	32.073	28.663	31.673	32.830	28.930	35.050	29.476	32.855	31.234	32.679	33.794	31.829	28.001	29.944	36.875	37.838	39.976
	MO	SD	2.521	1.181	2.923	2.111	1.356	3.343	1.395	1.513	1.135	0.959	2.124	1.017	3.439	1.714	2.032	1.255	2.160	1.102
	- Sn	Mean	30.768	33.739	28.759	32.386	31.557	28.039	33.710	30.147	33.238	32.675	31.419	31.464	29.949	24.155	27.605	31.039	34.043	36.395
	wou	SD	0.002	0.001	0.000	0.001	0.001	0.001	0.002	0.001		0.002	0.001	0.002	0.001	0.001	0.002	0.001	0.012	0.002
	s +	Mean	0.003	0.004	0.002	0.003	0.002	0.002	0.004	0.004		0.006	0.003	0.005	0.004	0.004	0.007	0.005	0.010	0.005
3/g DM)	bient	SD	0.001	0.001	0.000	0.001	0.001	0.000	0.001	0.001	0.001	0.001	0.001	0.001	0.003	0.001	0.002	0.001	0.001	0.002
Ash (Am	Mean	0.002	0.003	0.002	0.001	0.002	0.001	0.004	0.005	0.004	0.003	0.004	0.005	0.005	0.004	0.008	0.005	0.003	0.004
	MOL	SD	0.002	0.000	0.000	0.001	0.001	0.001	0.001	0.001	0.001	0.002	0.001	0.001	0.002	0.001	0.001	0.002	0.001	0.002
	-Sr	Mean	0.004	0.003	0.002	0.002	0.001	0.002	0.003	0.004	0.005	0.005	0.004	0.004	0.005	0.005	0.006	0.005	0.005	0.006
inatum	роү		170	177	184	191	198	205	212	219	173	180	187	194	200	210	217	223	231	237
E. vag	Year		2017								2018									

				NDF digest	Sibility (%)					NDS digesti	ibility (%)		
E. vagii	natum			0						D			
Year	МОД	- Sno	M	Amb	ient	+ Sn	MO	- Snc	M	Ambi	ent	÷Sn	MO
		Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD
2017	170	41.705	7.185	42.935	4.107	43.384	3.227	18.598	2.095	18.763	2.147	16.211	1.399
	177	49.940	5.701	50.313	1.818	49.926	4.548	21.067	0.982	19.683	1.829	19.452	0.576
	184	54.620	1.644	47.961	6.057	47.261	11.715	16.929	2.429	16.849	1.509	15.856	2.031
	191	52.895	1.812	40.225	14.375	44.159	4.986	19.943	1.754	19.350	1.720	22.521	1.682
	198	47.845	7.896	51.979	2.324	47.820	8.845	19.254	1.127	20.312	0.909	19.017	1.205
	205	43.833	6.356	41.248	14.743	39.013	15.761	16.331	2.778	17.071	1.091	17.937	0.630
	212	51.158	3.682	45.087	14.226	42.164	16.135	21.043	1.159	22.156	0.986	19.440	2.519
	219	47.203	5.710	48.063	9.326	37.515	10.213	18.083	1.258	17.525	1.185	20.333	0.729
2018	173	47.155	9.746	53.645	0.961			20.650	0.943	20.333	0.799		
	180	54.378	1.931	55.647	1.358	51.783	2.644	20.183	0.797	18.985	0.675	22.333	1.840
	187	53.742	0.303	46.192	11.603	52.594	3.684	19.139	1.765	20.186	1.390	20.888	0.504
	194	51.412	5.052	46.243	14.144	49.605	4.482	19.177	0.845	21.113	1.686	22.213	0.532
	200	50.493	1.886	52.921	0.801	52.022	5.730	17.918	2.858	19.480	0.909	19.890	0.594
	210	46.956	4.091	56.747	2.279	51.850	2.741	13.103	1.425	16.299	1.533	19.653	2.192
	217	43.211	15.781	33.844	12.954	35.248	12.570	15.970	1.688	17.913	1.387	20.623	1.596
	223	29.833	16.172	39.040	12.470	40.947	12.881	18.824	1.043	23.673	1.475	24.951	2.221
	231	37.430	9.345	41.074	10.297	45.748	4.757	21.319	1.795	24.474	1.842	26.747	1.539
	237	42.129	9.617	35.721	11.570	44.853	1.454	23.274	0.916	26.250	2.364	30.501	1.576