# Winter Snow Depth in Arctic Alaska Results in Complex Changes in Caribou Forage Quality 

Jessica C. Richert<br>South Dakota State University

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## BY <br> 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

## 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.
A. Joshua Leffler
Advisor Date
Michele R. Dudash
Department Head Date

Dean, Graduate School
Date

## ACKNOWLEDGMENTS

I owe my sincere appreciation to a great number of people who have supported my graduate education and research. I would especially like to thank my graduate advisor, Dr. A. Joshua Leffler, for his experience, support, patience, and knowledge of all things related to R. I would also like to extend my gratitude to the other members of my committee, Dr. Jonathan Jenks and Dr. Jeffrey Welker, for their extensive knowledge, guidance, and feedback over the last few years. A special thank you goes to my fellow graduate students, Kaj Lynöe and Heidi Becker, as well as our field technicians, Monica Ague, Jeremy Buttler, Brooke Davis, and Makyla Hammer, for aiding me in collecting thousands of field samples in the middle of rain, snow, and the occasional mosquito; I could not have done it without their help. I would also like to thank Dr. Don Spalinger, Dr. Kathy Kelsey, Dr. Stine Højlund-Pedersen, and John Ferguson at University of Alaska Anchorage as well as Dr. Derek Brake and Jason Griffin in the Animal Science Department at South Dakota State University for their assistance with various lab analysis procedures and additional data collection. Finally, I would like to thank South Dakota State University, the Department of Natural Resource Management, the wonderful administrative assistants who were always happy to lend an ear and a cup of coffee, and my fellow graduate students who provided a wealth of knowledge and experience in the classroom and the field. Thank you to you all.

Funding was provided through NSF awards ARC1604249 and ARC1602440.

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# ABSTRACT <br> WINTER SNOW DEPTH IN ARCTIC ALASKA RESULTS IN COMPLEX CHANGES IN CARIBOU FORAGE QUALITY 

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 $\% \mathrm{~N}$, 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 $\% \mathrm{~N}$ 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 nearsynchronous 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 $2^{\circ} \mathrm{C}$ since the early 1900 s (Osborne et al. 2018). Temperatures are predicted to continue rising by another $2-9^{\circ} \mathrm{C}$ 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 $22^{\circ} \mathrm{C}$ (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
$13.5^{\circ} \mathrm{C}$ (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 $(\mathrm{N})$ 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 $15^{\circ} \mathrm{C}$ (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 shrub-
dominated 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 $\mathrm{CO}_{2}$ 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 ( $68^{\circ} 38^{\prime}$ N $149^{\circ} 38$ W) in the foothills of the Brooks Range, Alaska, USA. The mean annual temperature is $-8^{\circ} \mathrm{C}$, with mean summer temperatures of
$10-12^{\circ} \mathrm{C}$ and winter temperatures averaging $-20^{\circ} \mathrm{C}$ (Hobbie and Kling 2014;
Environmental Data Center Team 2019). Mean precipitation is $250-350 \mathrm{~mm}$, with $40-$ $45 \%$ falling as snow (Schimel et al. 2004). In winter, snow depths reach $30-80 \mathrm{~cm}$ 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 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 ( $3 \times 60 \mathrm{~m}$ ) was erected on the moist-acidic tussock tundra site in 1994 to artificially increase snow depth, with snow drifts reaching a maximum depth of 3 m directly behind the snow fence and declining to ambient snow depths ( $0.5-1 \mathrm{~m}$ ) 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 ( $0.5-1 \mathrm{~m}$ 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 $70^{\circ} \mathrm{C}$ 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 $1 \mathrm{~m}^{2}$ quadrats, while biomass of all other species was harvested in three $20 \mathrm{~cm}^{2}$ quadrats nested within the larger $1 \mathrm{~m}^{2}$ plot. All samples were sorted to species, then dried in a forced-air oven at $70^{\circ} \mathrm{C}$ 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 $1-\mathrm{mm}$ 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 $100^{\circ} \mathrm{C}$ 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 $500^{\circ} \mathrm{C}$ for 5 hours to determine the total proportion of non-digestible lignin-cutin and inorganic matter. Percent N was analyzed via combustion at $1800^{\circ} \mathrm{C}$ 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 $50 \mu \mathrm{l}$ 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:

$$
\begin{align*}
& \mathrm{DMD}=\left(92.31 \mathrm{e}^{-0.0451(\mathrm{LIG}) * \mathrm{NDF})+(0.831 * \mathrm{NDS}-6.97)}\right.  \tag{1}\\
& \mathrm{DP}=-3.97+0.9283 * \mathrm{CP}-11.82 * \mathrm{PPC} \tag{2}
\end{align*}
$$

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 $\mathrm{p}<0.05$.

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 $\mathrm{PPC}=0$. 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 $39 \pm 11 \mathrm{~cm}$, while snow depth in the - snow and + snow zones was $20 \pm 3.4 \mathrm{~cm}$ and $147 \pm 18 \mathrm{~cm}$ 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 $8 \pm 1 \mathrm{~cm}$ in the shallow end of the + snow treatment, $17 \pm 6 \mathrm{~cm}$ in the - snow treatment, and $14 \pm 5 \mathrm{~cm}$ in ambient conditions. Half of the + snow treatment zone was still covered by a snow drift that was $38 \pm 11 \mathrm{~cm}$ deep. By the end of the growing season (DOY 237), active layer depths in all zones were $45-60 \mathrm{~cm}$, with the snow-covered area of the + snow treatment increasing from an initial depth of $10 \pm 1 \mathrm{~cm}$ at DOY 180 to $61 \pm 10 \mathrm{~cm}$ at DOY 237. The ambient zone depth increased to similar levels at $60 \pm 9 \mathrm{~cm}$, and the - snow treatment and shallow end of the + snow treatment reached depths of $51 \pm 10 \mathrm{~cm}$ and $49 \pm 7 \mathrm{~cm}$ respectively.

## Forage Availability

Overall biomass was not significantly different among the three snow zones, with $828.60 \pm 106.31 \mathrm{~g} / \mathrm{m}^{2}$ in the ambient plots, $748.24 \pm 35.26 \mathrm{~g} / \mathrm{m}^{2}$ in the - snow treatment plots, and $764.62 \pm 106.30 \mathrm{~g} / \mathrm{m}^{2}$ 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 $\left(\mathrm{F}_{2,15}=5.027, \mathrm{p}<0.05\right)$ higher biomass of deciduous shrubs occurring in the - snow treatment at $165.36 \pm 14.38 \mathrm{~g} / \mathrm{m}^{2}$ (compared to $135.82 \pm 14.79 \mathrm{~g} / \mathrm{m}^{2}$ in the + snow treatment). The - snow treatment also had the highest biomass of evergreen dwarf shrubs $\left(138.56 \pm 43.17 \mathrm{~g} / \mathrm{m}^{2}\right)$, while the + snow treatment had the lowest $\left(59.45 \pm 13.38 \mathrm{~g} / \mathrm{m}^{2}\right)$. Both the - snow and + snow treatments had significantly $\left(\mathrm{F}_{2,15}=11.8, \mathrm{p}<0.001\right)$ lower lichen biomass $(2.43 \pm 1.80$ and $8.51 \pm 4.10 \mathrm{~g} / \mathrm{m}^{2}$ respectively) compared to ambient $\left(50.68 \pm 12.47 \mathrm{~g} / \mathrm{m}^{2}\right)$. 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 ( $\sim 2 \%$ ) 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
( $\sim 70 \%$ ) changed little throughout time, while digestibility of $V$. vitis-idaea showed a curvilinear trend over the growing season, with a lower overall digestibility of $\sim 50-60 \%$. 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 ( $\sim 45-50 \%$ in $R$. tomentosum and $\sim 60-70 \%$ 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 $0.276 \pm 0.014 \mathrm{mg}$ BSA precipitated $/ \mathrm{mg}$ 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
$0.226 \pm 0.027 \mathrm{mg}$ BSA precipitated $/ \mathrm{mg}$ DM at the beginning to a high of $0.357 \pm 0.015$ mg BSA precipitated/mg DM by DOY 217 before declining to $0.334 \pm 0.023 \mathrm{mg}$ BSA precipitated/mg DM by senescence (Fig. 5).

Caribou need a minimum of $7-8 \mathrm{~g}$ DP/ 100 g 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 $\mathrm{g} / 100 \mathrm{~g}$ 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 $V$. vitisidaea, remaining at ca. $2 \mathrm{~g} / 100 \mathrm{~g}$ DM throughout the sample period. Digestible protein in C. bigelowii mirrored the same pattern as the deciduous shrubs, beginning the season at $15-20 \mathrm{~g} / 100 \mathrm{~g}$ 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 $1-m^{2}$ 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 $\sim 25$ 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

Table 1. Model selection results for nitrogen content of each species

| Model | logLik | $\mathrm{AlC}_{\mathrm{c}}$ | $\Delta A L C_{c}$ | $d f$ | 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 2. Model selection for dry matter digestibility of each species

| Model | logLik | AIC ${ }_{\text {c }}$ | $\Delta A L C_{c}$ | $d f$ | 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 |

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

| Model | logLik | AIC $_{\text {c }}$ | $\Delta \mathrm{AlC}_{\text {c }}$ | $d f$ | 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 |

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; $\mathbf{B}$ represents the - snow treatment zone; $\mathbf{C}$ represents the + snow treatment zone.

Figure 2. Total biomass $\left(\mathrm{g} / \mathrm{m}^{2}\right)$ for each functional group by snow treatment


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 ( $7-8 \mathrm{~g} / 100 \mathrm{~g} \mathrm{DM}$ ) 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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## APPENDIX 1



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




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




| R. tomentosum |  | NDF (g/g DM) |  |  |  |  |  | ADF (g/g DM) |  |  |  |  |  | Cellulose (g/g DM) |  |  |  |  |  | Lignin (g/g DM) |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| Year | Doy | - Snow |  | Ambient |  | + Snow |  | - Snow |  | Ambient |  | + Snow |  | - Snow |  | Ambient |  | + Snow |  | -Snow |  | Ambient |  | + Snow |  |
|  |  | 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 | 0.009 | 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 | 0.009 | 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 | 0.009 |  |  | 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 | 0.009 | 0.123 | 0.006 | 0.136 | 0.009 | 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 | 0.009 | 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 | 0.009 | 0.291 | 0.015 | 0.137 | 0.010 | 0.135 | 0.005 | 0.121 | 0.009 | 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 | 0.009 | 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 | 0.009 | 0.285 | 0.014 | 0.120 | 0.008 | 0.117 | 0.009 | 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 | 0.099 | 0.006 | 0.154 | 0.009 | 0.137 | 0.011 | 0.141 | 0.007 |


| R. tomentosum |  | Ash (g/g DM) |  |  |  |  |  | Cell Solubles (\% of DM) |  |  |  |  |  | Cell Wall (\% of DM) |  |  |  |  |  | Lig/Cut (\% of NDF) |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| Year | DOY | - Snow |  | Ambient |  | + Snow |  | - Snow |  | Ambient |  | + Snow |  | - Snow |  | Ambient |  | + Snow |  | - Snow |  | Ambient |  | + Snow |  |
|  |  | 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 | 0.009 | 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 | 0.009 | 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 | 0.009 | 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 | 0.009 | 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 | 2.81 |
|  | 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 | 1.75 |



|  |  | ～ | O． | \％ | ¢ | － | O. | $\underset{0}{7}$ | N్రీ. | －1 |  | O | O | $\stackrel{7}{0}$ | O\％ | $\stackrel{7}{0}$ | O\％ | O | ¢ | O |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | $+$ | $\stackrel{\text { IN }}{\sum_{\Sigma}^{0}}$ | $\stackrel{\underset{\sim}{0}}{ }$ | $\underset{\sim}{\sim}$ | $\begin{aligned} & \text { n } \\ & \text { O. } \end{aligned}$ | ô | 긍 | $\underset{\sim}{7}$ | $\begin{aligned} & \stackrel{\sim}{0} \\ & 0 \end{aligned}$ | $\stackrel{\circ}{\circ}$ |  | $\stackrel{\overleftarrow{\circ}}{\substack{0}}$ | $\stackrel{\overleftarrow{\circ}}{\circ}$ | $\underset{0}{\mathbf{O}}$ | $\stackrel{\text { ò }}{\circ}$ | $\stackrel{n}{7}$ | $\stackrel{n}{\square}$ | $\begin{array}{r} \square \\ \vdots \\ \hline \end{array}$ | $\stackrel{\vdots}{0}$ |  |
|  |  | 0 | $\begin{aligned} & \text { O్ర } \\ & \text { Oi } \end{aligned}$ | $\stackrel{\sim}{0}$ | $\begin{aligned} & \circ \circ \\ & \hline 0 . \end{aligned}$ | $\stackrel{\infty}{\circ}$ | $\stackrel{\text { O}}{\mathrm{O}}$ | $\stackrel{n}{0}$ | ồ | $\underset{\sim}{\tilde{O}}$ | $\stackrel{0}{0}$ | Ợ | $\stackrel{0}{\circ}$ | O. | $\stackrel{\infty}{\circ}$ | $\stackrel{\hat{0}}{0}$ | Oio | $\stackrel{\text { no }}{\substack{0 \\ \hline}}$ | $\stackrel{\circ}{\circ}$ | O |
|  |  |  | $\stackrel{\rightharpoonup}{7}$ | ন্ত̆ | ® | 항 | $\begin{aligned} & \text { 강 } \end{aligned}$ | $\underset{\sim}{0}$ | $\stackrel{\underset{\sim}{0}}{\substack{0}}$ | $\stackrel{\rightharpoonup}{0}$ | $\stackrel{\circ}{0}$ | E | O. | Õ | 荅 | 긍 | $\stackrel{\circ}{\square}$ | $\stackrel{\infty}{\circ}$ | ơ |  |
|  | $\begin{aligned} & 3 \\ & \text { 30 } \\ & \text { n } \end{aligned}$ | 0 | $\begin{aligned} & \text { ñ } \\ & \stackrel{0}{0} \end{aligned}$ | $\begin{aligned} & 0 \\ & \stackrel{0}{0} \\ & \hline \end{aligned}$ | Ñ | 亿i | $\stackrel{\infty}{0}$ | O | $\begin{aligned} & \stackrel{0}{0} \\ & \text { Ö } \end{aligned}$ | ت | $\stackrel{\infty}{\circ}$ | O | $\begin{aligned} & \tilde{0} \\ & 0 \end{aligned}$ | O | $8$ | 荔 | $\begin{aligned} & 0 . \\ & \stackrel{\rightharpoonup}{0} \\ & \hline- \end{aligned}$ | $\begin{aligned} & 0 \\ & \stackrel{y}{0} \end{aligned}$ | ô |  |
|  |  | $\begin{aligned} & \stackrel{ᄃ}{\omega} \\ & \stackrel{N}{\Sigma} \end{aligned}$ | $\begin{aligned} & \stackrel{0}{7} \\ & 0 \end{aligned}$ | $$ | $\begin{aligned} & \text { on } \\ & \hline 0 \\ & 0 \end{aligned}$ | $\underset{0}{0}$ | ت̃ | $\begin{aligned} & \text { 궁 } \end{aligned}$ | $\begin{aligned} & \underset{\sigma}{\circ} \\ & \hline \end{aligned}$ | $\begin{aligned} & \circ \\ & \stackrel{0}{0} \end{aligned}$ | No | $\begin{aligned} & \text { ® } \\ & \stackrel{0}{0} \end{aligned}$ | $\begin{aligned} & \text { O} \\ & \text { O} \\ & \hline \end{aligned}$ | 흥 | $\stackrel{0}{0}$ | $\stackrel{\infty}{\underset{\sigma}{\circ}}$ | $\stackrel{\rightharpoonup}{7}$ | $\underset{0}{7}$ | $\begin{aligned} & \text { ® } \\ & \stackrel{\circ}{\circ} \end{aligned}$ |  |
|  |  | 0 | $\begin{aligned} & \text { O} \\ & \hline 0 \\ & \hline \end{aligned}$ |  | O | O |  |  | $\begin{aligned} & \text { gे } \\ & \stackrel{0}{0} \end{aligned}$ | OO |  | $\stackrel{0}{0}$ | $\stackrel{n}{0}$ | 잉 | $\stackrel{7}{0}$ | 픙 | $\stackrel{\infty}{0}$ | $\begin{aligned} & \circ \\ & \hline 8 \\ & \hline \end{aligned}$ | oi | $\stackrel{7}{0}$ |
|  |  | $\begin{aligned} & \stackrel{\pi}{0} \\ & \stackrel{N}{\Sigma} \end{aligned}$ | $\stackrel{\substack{\pi \\ \hline}}{ }$ | $\stackrel{m}{\tilde{m}}$ | $\underset{\underset{0}{\hat{0}}}{ }$ | $\begin{aligned} & \text { تٌ } \\ & \text { O} \end{aligned}$ | $\underset{\underset{0}{0}}{\tilde{0}}$ | $\stackrel{\sim}{\underset{\sim}{0}}$ | $\begin{gathered} \text { ๙̃ } \\ 0 \end{gathered}$ | $\stackrel{\tilde{m}}{\underset{0}{0}}$ |  | $\stackrel{\substack{\mathrm{m}}}{0}$ | $\underset{\sim}{\tilde{O}}$ | $\underset{\underset{0}{7}}{\substack{0 \\ \hline}}$ | $\underset{\sim}{\tilde{O}}$ | $\underset{\sim}{\underset{\sigma}{\prime}}$ | $\underset{\substack{0 \\ \hline}}{0}$ |  | $\underset{\sim}{\tilde{O}}$ |  |
|  |  | i | $\begin{aligned} & 0 \\ & \stackrel{0}{0} \\ & \hline \end{aligned}$ | $\begin{aligned} & \stackrel{m}{0} \\ & \stackrel{0}{0} \end{aligned}$ | $\begin{aligned} & \circ \\ & \hline 0 \\ & \hline 0 \end{aligned}$ | 冾 | Ò | $\stackrel{\infty}{0}$ | $\begin{aligned} & \text { ñ } \\ & \stackrel{0}{0} \end{aligned}$ | $\stackrel{7}{0}$ | ồ | $\stackrel{\mathrm{O}}{\mathrm{o}}$ | $\begin{aligned} & \text { O} \\ & \hline 0 \\ & \hline \end{aligned}$ | $\underset{0}{7}$ | $\stackrel{0}{0}$ | $\underset{0}{7}$ | $\begin{aligned} & \text { O. } \\ & \stackrel{y}{0} \end{aligned}$ | $\begin{aligned} & \circ \\ & \hline 0 \\ & \hline \end{aligned}$ | On |  |
|  |  |  | $\stackrel{\sim}{\tilde{O}}$ | $\underset{\sim}{\text { In }}$ | $\underset{\sim}{\sim}$ | $\underset{\text { İ }}{\text { In }}$ | $\underset{\substack{\mathrm{m} \\ \hline}}{\substack{0}}$ | $\stackrel{n}{\underset{\sim}{\circ}}$ | $\stackrel{\stackrel{m}{0}}{0}$ | $\stackrel{0}{\pi}$ | $\underset{\sim}{\underset{O}{9}}$ | $\stackrel{\infty}{\underset{\sim}{0}}$ | $\stackrel{\sim}{7}$ | $\underset{\underset{0}{\hat{0}}}{\substack{1 \\ \hline}}$ | $\stackrel{\sim}{\tilde{m}}$ | $\stackrel{N}{\underset{\sim}{0}}$ |  | $\stackrel{\tilde{N}_{0}^{0}}{0}$ | $\stackrel{\stackrel{\Gamma}{0}}{0}$ |  |
|  | $\begin{aligned} & 3 \\ & 5 \\ & 5 \end{aligned}$ | 0 | O | $\begin{aligned} & 0 . \\ & \stackrel{\rightharpoonup}{0} \end{aligned}$ | $\stackrel{\rightharpoonup}{0}$ | $\begin{aligned} & \text { İ } \\ & 0 \end{aligned}$ | $\begin{aligned} & \circ \\ & \hline 0 \\ & \hline 0 \end{aligned}$ | $\begin{aligned} & \text { İ } \\ & \text { O} \end{aligned}$ | O | $\begin{aligned} & \text { O. } \\ & \stackrel{1}{0} \end{aligned}$ | $\stackrel{\infty}{\circ}$ | $\underset{0}{3}$ | O | $\stackrel{m}{0}$ | $\begin{aligned} & 0 \\ & \stackrel{y}{0} \end{aligned}$ | $\stackrel{m}{0}$ | Õ | $\begin{aligned} & \text { O} \\ & \hline 0 \\ & \hline \end{aligned}$ | $\stackrel{\infty}{\circ}$ |  |
|  |  | $\begin{aligned} & \text { 厄్ల } \\ & \text { } \end{aligned}$ | $\begin{aligned} & \stackrel{\sim}{0} \\ & \text { O } \end{aligned}$ | $\underset{\sim}{\text { In }}$ | $\stackrel{\rightharpoonup}{\underset{0}{0}}$ | $\begin{aligned} & \underset{\sim}{0} \end{aligned}$ | $\stackrel{\sim}{\underset{\sigma}{0}}$ | $\stackrel{\hat{m}}{0}$ | $\stackrel{\sim}{\tilde{\sigma}}$ | $\underset{\sim}{\underset{\sim}{J}}$ | $\underset{\sim}{\tilde{\sim}}$ | $\begin{aligned} & \stackrel{0}{7} \\ & \stackrel{0}{0} \end{aligned}$ | $\underset{\sim}{\sim}$ |  | $\stackrel{\tilde{m}}{0}$ | $\stackrel{n}{0}$ | $\stackrel{\infty}{\underset{\sim}{0}}$ | $\underset{\substack{0 \\ \hline}}{\substack{0}}$ | $\underset{\sim}{\sim}$ |  |
|  |  | 0 |  | 笑 | $\begin{aligned} & \circ \\ & \hline 0 \\ & \hline 0 \end{aligned}$ |  |  |  | $\begin{aligned} & \text { No } \\ & \text { O. } \end{aligned}$ | $\stackrel{n}{0}$ |  | $\begin{aligned} & \circ \\ & \hline 0 \\ & \hline 0 \end{aligned}$ | $\begin{aligned} & \stackrel{\infty}{0} \\ & \stackrel{0}{\circ} \end{aligned}$ | $\begin{aligned} & \text { O. } \\ & \stackrel{1}{0} \end{aligned}$ | $\begin{aligned} & \text { 응 } \\ & \text {. } \end{aligned}$ | تָ. | 응 | $\begin{aligned} & \circ \\ & \hline 8 \\ & \hline 0 \end{aligned}$ | $\underset{\sim}{\sim}$ | $\stackrel{7}{0}$ |
|  |  |  | $\underset{\substack{N \\ \text { N }}}{ }$ | $\underset{\substack{\underset{\sim}{0} \\ \hline}}{ }$ | セ̛̣ | $\stackrel{\sim}{\sim}$ | Ñ | $\underset{\substack{\text { N }}}{ }$ | $\begin{aligned} & \text { مٌ } \\ & \text { No } \end{aligned}$ | $\stackrel{\sim}{\sim}$ |  | 멍 | $\underset{\sim}{\underset{\sim}{2}}$ | $\begin{aligned} & \text { N } \\ & \text { 둥 } \end{aligned}$ | הָ | $\underset{\sim}{\mathrm{N}}$ | $\stackrel{\infty}{\underset{0}{0}}$ | $\underset{\substack{\text { d } \\ \hline}}{\text { N }}$ | ત్వి |  |
|  |  | is | 筑 | $\begin{aligned} & 0 \\ & 0.0 \\ & \hline \end{aligned}$ | O | $\begin{aligned} & \text { O} \\ & \stackrel{y}{0} \end{aligned}$ | $\underset{0}{J}$ | त̈ | $\begin{aligned} & \text { O. } \\ & \stackrel{\rightharpoonup}{0} \end{aligned}$ | O | $\begin{aligned} & \circ \\ & \hline 0 \\ & \hline 0 \end{aligned}$ | oi | O | $\stackrel{0}{0}$ | $\stackrel{m}{0}$ | $\stackrel{\hat{0}}{0}$ | ô | O | O |  |
|  |  |  | $\begin{aligned} & \text { Oid } \\ & \text { O- } \end{aligned}$ | $\underset{\sim}{\text { ơb }}$ | $\underset{\sim}{\tilde{N}}$ | ~্ત | $\underset{\substack{\infty \\ 0}}{\infty}$ |  | $\underset{\sim}{\tilde{\sim}}$ | $\underset{\substack{\mathrm{N}}}{\substack{\text { a }}}$ | Niべ | N్స్రీ | ત্તু | ̈ㅡㅇ | $\begin{aligned} & \text { ON} \\ & \text { Ó } \end{aligned}$ | ợ | O. | $\underset{\sim}{\hat{\sim}}$ | $\underset{\sim}{\sim}$ |  |
|  | $\begin{aligned} & 3 \\ & \stackrel{3}{6} \end{aligned}$ | 0 | $\begin{aligned} & \stackrel{\infty}{0} \\ & \stackrel{0}{0} \end{aligned}$ | オ্তু | $\begin{aligned} & \text { O} \\ & \text { O. } \end{aligned}$ | $\begin{aligned} & 0 \\ & 0.0 \\ & 0 . \end{aligned}$ | N్రీ. | O. | $\begin{aligned} & 0 \\ & 0.0 \\ & 0 \end{aligned}$ | ت̃ | ồ | 亏⿳亠丷⿵冂⿱丷丅犬 | $\begin{aligned} & \text { O. } \\ & \stackrel{0}{0} \end{aligned}$ | 今̀ | O. | 筞 | O | $\begin{aligned} & \infty \\ & \stackrel{\infty}{\circ} \\ & \hline \end{aligned}$ | $\begin{aligned} & \circ \\ & \hline 0 \\ & \hline \end{aligned}$ |  |
|  |  |  | $\underset{\substack{0 \\ 0}}{ }$ | $\underset{\substack{0 \\ \hline}}{ }$ | ה̃ | $\stackrel{\sim}{\mathrm{N}}$ | ホ̛ণ ָ | $\underset{\sim}{0}$ | $\stackrel{\stackrel{\sim}{0}}{0}$ | Nơ | ત্ત̃ | הָ | స்̇ | $\underset{\substack{\text { ®j }}}{ }$ | $\stackrel{\substack{0 \\ \text { di }}}{ }$ | $\underset{\substack{0 \\ \hline}}{\substack{0}}$ | N్ర | $\stackrel{\substack{0}}{\substack{2}}$ |  |  |
|  | $\begin{aligned} & 3 \\ & 0 \\ & \mathbf{5} \\ & + \end{aligned}$ | 0 |  | 荅 | $\stackrel{\otimes}{\square}$ |  |  | $\stackrel{\rightharpoonup}{\circ}$ | no | $\stackrel{n}{0}$ |  | べ． | － | $\begin{aligned} & 0 \\ & \stackrel{0}{0} \end{aligned}$ | \％ | － | $\stackrel{\text { à }}{0}$ | \％ | $\stackrel{\text { äd }}{0}$ | $\stackrel{\rightharpoonup}{\circ}$ |
|  |  |  | $\begin{aligned} & \text { un } \\ & \substack{0} \end{aligned}$ | $\begin{aligned} & 0 \\ & \substack{0 \\ \hline} \end{aligned}$ | $\stackrel{\rightharpoonup}{\mathbf{N}}$ | $\underset{\sim}{\sim}$ |  | $\begin{aligned} & \stackrel{y}{N} \\ & \text { No } \end{aligned}$ | Oion | $\underset{\text { N゙ }}{\text { N }}$ |  | $\begin{aligned} & \text { O} \\ & \text { O} \end{aligned}$ | ষ্লু | N్రె | $\begin{aligned} & \text { O} \\ & \text { O} \\ & \hline \end{aligned}$ | $\begin{aligned} & \text { to } \\ & \text { O} \\ & \hline \end{aligned}$ | $\stackrel{\substack{n \\ 0}}{0}$ | $\stackrel{\substack{\infty \\ 0 \\ 0}}{ }$ | $\underset{\sim}{\tilde{N}}$ |  |
|  |  | is | $\begin{aligned} & \text { Ö } \\ & \text { Oi } \end{aligned}$ | $\begin{aligned} & \text { ờ } \\ & \text { O } \end{aligned}$ | $\begin{aligned} & \infty \\ & \hline 0 \\ & \hline 0 \end{aligned}$ | $\begin{aligned} & \text { ñ } \\ & \stackrel{0}{0} \end{aligned}$ | $\stackrel{\sim}{\circ}$ | ö | $\begin{aligned} & \stackrel{\infty}{0} \\ & \stackrel{0}{0} \end{aligned}$ | $\begin{aligned} & \text { O} \\ & \text { O. } \end{aligned}$ | $\begin{aligned} & \text { Ơ } \\ & \text { O} \end{aligned}$ | $\underset{0}{7}$ | $\underset{0}{7}$ | $\begin{aligned} & \stackrel{\circ}{7} \\ & \hline-1 \end{aligned}$ | ત్ర | N్రీ. | $\begin{aligned} & 0 . \\ & \stackrel{y}{0} \end{aligned}$ | $\begin{aligned} & \text { Ñ } \\ & \text { On } \end{aligned}$ | $\stackrel{m}{0}$ |  |
|  |  |  | $\begin{gathered} \text { n } \\ \substack{0} \end{gathered}$ | $\stackrel{\rightharpoonup}{\infty}$ | $\begin{gathered} \stackrel{\rightharpoonup}{m} \\ \text { oi } \end{gathered}$ | $\begin{gathered} \text { m} \\ 0 \\ \hline \end{gathered}$ | $\begin{aligned} & \text { ت} \\ & \text { on } \end{aligned}$ | $\begin{aligned} & \text { n } \\ & \\ & \hline \end{aligned}$ |  | $\begin{aligned} & \infty \\ & \underset{\sim}{\infty} \\ & 0 \end{aligned}$ | -i | $\stackrel{\sim}{\infty}$ | $\begin{gathered} \text { O} \\ \text { No } \end{gathered}$ | $\underset{\substack{0 \\ 0}}{\substack{0}}$ | $\begin{gathered} \text { n } \\ \substack{0} \end{gathered}$ | $\stackrel{N}{0}$ | $\underset{\substack{-0 \\ 0}}{ }$ | $\begin{gathered} \underset{\sim}{2} \\ 0 \end{gathered}$ | $\underset{\sim}{\sim}$ |  |
|  | $\begin{aligned} & 3 \\ & 5 \\ & 5 \end{aligned}$ | 0 |  | た్రి. | $\begin{aligned} & \stackrel{\rightharpoonup}{0} \\ & 0 . \end{aligned}$ | $\begin{aligned} & 0 \\ & \stackrel{0}{0} \end{aligned}$ |  | ${ }_{0}^{\text {N̈ }}$ | O. | O | $\stackrel{8}{\circ}$ | $\begin{aligned} & \text { O} \\ & 0 \end{aligned}$ | oi | へ | $\stackrel{\square}{\square}$ | ¢ | \％ | ড | ${ }_{0}^{\circ}$ |  |
|  |  |  | $\begin{gathered} \infty \\ \\ \hline 0 \end{gathered}$ | $\begin{gathered} \text { O} \\ \\ \hline 0 \end{gathered}$ | $\begin{gathered} 0 \\ \\ 0 \end{gathered}$ | $\underset{\sim}{\sim}$ | $\stackrel{0}{0}$ | $\stackrel{\underset{N}{N}}{\substack{0}}$ | $\stackrel{\otimes}{0}$ | $\begin{aligned} & \stackrel{\circ}{0} \\ & 0 . \end{aligned}$ | $\stackrel{\text { ®id }}{0}$ | $\stackrel{\underset{m}{0}}{\stackrel{\rightharpoonup}{0}}$ | $\begin{gathered} \text { t } \\ \text { Ò } \end{gathered}$ | $\begin{gathered} \underset{m}{0} \\ 0 \end{gathered}$ | $\begin{aligned} & \infty \\ & 0 \\ & 0 \end{aligned}$ | $\begin{aligned} & \text { n } \\ & \substack{0} \end{aligned}$ | $\stackrel{\circ}{0}$ | $\stackrel{\text { O}}{\substack{0}}$ | － | $\stackrel{\stackrel{\rightharpoonup}{0}}{0}$ |
| $\begin{aligned} & \stackrel{8}{8} \\ & \stackrel{0}{c h} \\ & \stackrel{y}{3} \\ & > \\ & > \end{aligned}$ | \％ |  | $\stackrel{\bigcirc}{7}$ | F | $\underset{\sim}{\square}$ | ন | $\stackrel{\sim}{\square}$ | $\stackrel{\sim}{\sim}$ | N | $\stackrel{\text { a }}{ }$ | $\stackrel{n}{7}$ | $\stackrel{\sim}{\square}$ | $\stackrel{\infty}{\sim}$ | 尔 | 8 | $\stackrel{7}{7}$ | $\stackrel{\text { İ }}{ }$ | $\underset{\sim}{\sim}$ | $\underset{\sim}{\sim}$ | $\stackrel{\sim}{\sim}$ |
|  | $\stackrel{\text { ® }}{\text { ¢ }}$ |  | స̇̈ |  |  |  |  |  |  |  | $\stackrel{\sim}{\sim}$ |  |  |  |  |  |  |  |  |  |


| V.vitis | idea | Ash (8/g DM) |  |  |  |  |  | Cell Solubles (\% of DM) |  |  |  |  |  | Cll Wall (\%\% of DM) |  |  |  |  |  | Lig/Cut (\% of NDF) |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| vear | por | - Snow |  | Ambient |  | + Snow |  | - snow |  | Ambient |  | + Snow |  | - Snow |  | Ambient |  | + Snow |  | -snow |  | Ambient |  | + Snow |  |
| Year |  | Mean | sd | Mean | so | Mean | so | Mean | so | Mean | so | Mean | sD | Mean | sD | Mean | so | Mean | so | Mean | so | Mean | so | Mean | so |
| 2017 | 170 | 0.004 | 0.001 | 0.005 | 0.001 | 0.004 | 0.001 | 61.713 | 2.425 | 60.550 | 2.63 | 63.459 | ${ }^{3.415}$ | 38.287 | 2.42 | 39.45 | 2.631 | 36.541 | 3.415 | 30.23 | 2.433 | 30.256 | 4.659 | 34.360 | 7.298 |
|  | 177 | 0.005 | 0.001 | 0.004 | 0.001 | 0.004 | 0.001 | 61.112 | 2.85 | 856 | 5.865 | 59.681 | 5.576 | 38.8 | 2.857 | 38.14 | 5.865 | 40.319 | 5.576 | 26.6 | 3.305 | 28.9 | 2.602 | 30.469 | 5.679 |
|  | 184 | 0.012 | 0.01 | 0.004 | 0.001 | 0.003 | 0.000 | 66.361 | 1.871 | 6.596 | 0.83 | 60.93 | 18.888 | 33.639 | 1.87 | 33.04 | 0.833 | 39.066 | 18.888 | 23.42 | 5.985 | 25.47 | 1.261 | 24.57 | 2.474 |
|  | 191 | 0.003 | 0.001 | 0.003 | 0.001 | 0.003 | 0.001 | 67.750 | 1.618 | 66.674 | 1.991 | 6. 804 | 0.679 | 32.25 | 1.6 | 3.33 | 1.45 | 33.196 | 0.67 | 28.20 | 0.312 | 29.079 | 1.75 | 29.12 | 1.645 |
|  | 198 | 0.003 | 0.001 | 0.004 | 0.004 | 0.003 | 0.000 | 64.137 | 2.380 | 63.850 | 1.533 | 6.418 | 2.593 | 35.863 | 2.38 | 36.15 | 1.53 | 36.58 | 2.593 | ${ }^{31.057}$ | 3.23 | 30.429 | 1.19 | 31.900 | 3.605 |
|  | 205 | 0.004 | 0.001 | 0.004 | 0.002 | 0.002 | 0.000 | 62.861 | 1.220 | 66.510 | 1.930 | ${ }_{65.356}$ | 1.921 | 37.139 | 1.220 | 33.49 | 1.93 | 34.644 | 1.921 | 32.254 | 0.848 | 33.74 | 4.87 | 32.502 | 1.967 |
|  | 212 | 0.006 | 0.004 | 0.003 | 0.001 | 0.005 | 0.003 | 63.419 | 3.935 | 65.573 | 1.768 | 64.000 | 3.524 | 36.581 | 3.93 | 34.427 | 1.768 | 36.000 | 3.524 | 32.248 | 3.93 | 29.61 | 1.58 | 34.870 | 4.790 |
|  | 219 | 0.008 | 0.002 | 0.008 | 0.003 | 0.007 | 0.002 | 63.964 | 2.555 | ${ }^{61.683}$ | 2.993 | 65.820 | 1.482 | 36.036 | 2.555 | 38.317 | 2.993 | 34.180 | 1.482 | 29.499 | 1.065 | 33.908 | 3.478 | 29.195 | 0.659 |
| 2018 | 173 | 0.006 | 0.000 | 0.005 | 0.001 |  |  | 70.477 | 0.736 | 70.956 | 0.534 |  |  | 29.523 | 0.736 | 29. | 0.534 |  |  | 24.510 | 2.607 | 27.667 | 1.039 |  |  |
|  | 180 | 0.007 | . 001 | 0.008 | 0.003 | 0.006 | 0.001 | 1 | 2.959 | .533 | 1.068 | 69.37 | 1.230 | 31.70 | 2.959 | 28.46 | 1.068 | 30.62 | 1.230 | 26.674 | 1.068 | 27.116 | 1.725 | 27.3 | 1.167 |
|  | 187 | 0.005 | 001 | 0.006 | 0.001 | 0.005 | 0.002 | 69.56 | 0.919 | 437 | 1.138 | 6. 61 | 254 | 435 | 0.919 | 9.56 | .138 | 30.3 | 2.554 | 29.3 | 3.259 | 2.772 | 0.937 | 27.554 | 1.341 |
|  | 194 | 0.005 | 0.001 | 006 | 0.001 | 0.007 | 0.001 | 66.828 | 2.699 | 311 | 10.62 | ${ }^{70.321}$ | 1.598 | 33.172 | 2.699 | 26.68 | 0.62 | 29.6 | 1.598 | 29.4 | 2.641 | 47.073 | 39.3 | 19 | 5.425 |
|  | 200 | 0.00 | 0.002 | 0.006 | 0.002 | 0.008 | 0.002 | 66.392 | 1.068 | 66.519 | 2.128 | 9.379 | 1.961 | 33.608 | 1.068 | 3.48 | 2.128 | 30.6 | 1.961 | 30.72 | 2.377 | 28.2 | 1.311 | 28.626 | 2.560 |
|  | 210 | 0.00 | 0.001 | 0.006 | 0.001 | 0.006 | 0.001 | 99 | 4.731 | 64.722 | 2.471 | ${ }^{3} .563$ | 2.088 | 40.501 | 4.731 | 35.278 | 2.471 | 36.437 | 2.088 | 33.746 | 4.562 | 31.118 | 3.4 | 37 | 1.43 |
|  | 217 | 0.00 | 0.001 | 007 | 0.001 | 0.008 | 0.002 | 64.973 | 0.897 | 63.914 | 1.012 | 4.863 | 1.878 | 35.027 | 0.897 | 36.086 | 1.01 | 35.137 | 1.878 | 30.59 | 2.13 | 30.289 | 2.639 | 29.831 | 1.368 |
|  | 223 | 0.008 | 0.003 | 007 | . 001 | 0.009 | 0.002 | 66.006 | 2.422 | 67.054 | 1.172 | 6.489 |  | 33.994 | 2.422 | 32.946 | 1.172 | 33.511 | 0.606 | 32.495 | 1.85 | 29.640 | 1.638 | 30.265 | 1.261 |
|  | 231 | 0.008 | 0.003 | 006 | 0.002 | 0.007 | . 003 | 67.363 | 0.912 | 67.591 | 1.292 | 68.841 |  | 32.637 | 0.912 | 32.409 | 1.292 | 31.15 | 1.892 | 30.052 | 2.446 | 30.660 | 1.50 | 32.02 | 0.980 |
|  | 237 | 0.00 | 0.002 | 0.008 | 0.001 | 0.008 | 0.001 | 69.025 | 1.374 | 69.313 | 1.105 | 69.654 | 1.675 | 30.975 | 1.374 | 30.687 | 1.105 | 30.346 | 1.675 |  | 1.421 | 31.348 | 2.042 | 884 |  |


|  |  | i | $\begin{aligned} & \infty \\ & \stackrel{\infty}{\infty} \\ & \underset{i}{2} \end{aligned}$ | $\underset{\substack{\text { ¢ } \\ \underset{\sim}{c} \\ \hline}}{ }$ |  | $\begin{aligned} & \text { to } \\ & \text { Ho } \end{aligned}$ | $\stackrel{\text { N }}{\underset{\sim}{n}}$ | $\begin{aligned} & \text { Q } \\ & \underset{\sim}{n} \end{aligned}$ | $\begin{gathered} \underset{\sim}{i} \\ \underset{\sim}{2} \end{gathered}$ | $\stackrel{\rightharpoonup}{\underset{\sim}{7}}$ |  | $\begin{gathered} \underset{\sim}{\sim} \\ \hline \end{gathered}$ | $\underset{\sim}{\underset{\sim}{\sim}}$ | $\stackrel{\underset{\sim}{\infty}}{\substack{\infty}}$ | $\begin{aligned} & \text { O} \\ & \underset{\sim}{6} \end{aligned}$ | $\underset{\substack{n \\ \underset{\sim}{n}}}{ }$ | $\begin{aligned} & \stackrel{\rightharpoonup}{0} \\ & \underset{\sim}{n} \end{aligned}$ | $\begin{aligned} & \text { to } \\ & \text { Nin } \end{aligned}$ | $\xrightarrow{\text { Nิ }}$ |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  |  |  | $\begin{aligned} & \stackrel{\sim}{0} \\ & \underset{\sim}{\gamma} \end{aligned}$ |  | $\begin{aligned} & \underset{U}{\mathbb{G}} \\ & \underset{\sim}{\infty} \end{aligned}$ | $\begin{aligned} & \tilde{\sim} \\ & \text { ஷુ寸 } \end{aligned}$ | $\begin{aligned} & \vec{\sim} \\ & \underset{\sim}{\prime} \end{aligned}$ | $\begin{aligned} & \underset{\sim}{7} \\ & \text { ¢ } \end{aligned}$ | $\begin{aligned} & \stackrel{\circ}{ } \\ & \underset{\sim}{2} \end{aligned}$ |  | $\begin{aligned} & \text { ® } \\ & \text { in } \\ & \hline \end{aligned}$ | ＋ in in | $\begin{gathered} \text { Win } \\ \text { in } \end{gathered}$ | $\begin{aligned} & \text { + } \\ & \text { O} \\ & \text { in } \end{aligned}$ | $\begin{aligned} & \text { In } \\ & \text { ஷु } \end{aligned}$ | $\begin{aligned} & \overrightarrow{0} \\ & \dot{6} \end{aligned}$ | $\begin{aligned} & \underset{\sim}{\infty} \\ & \underset{\sim}{\infty} \end{aligned}$ | $\begin{aligned} & \text { N} \\ & \text { in } \end{aligned}$ | ñ |
|  |  | n | $\stackrel{\stackrel{\infty}{i}}{\underset{\sim}{i}}$ | $\underset{\substack{ \pm \underset{\sim}{+}}}{\substack{0}}$ | Ơ | $\underset{\sim}{\underset{\sim}{0}}$ | $\underset{\underset{A}{A}}{\underset{\sim}{2}}$ | $\begin{aligned} & \text { to } \\ & \hline \end{aligned}$ | ợ | $\stackrel{\stackrel{\rightharpoonup}{d}}{\underset{\sim}{2}}$ | $\begin{aligned} & \text { 管 } \end{aligned}$ | $\begin{aligned} & \text { è } \\ & \stackrel{\infty}{\infty} \end{aligned}$ | No | $\underset{\substack{\infty \\ \infty \\ \infty}}{\substack{n}}$ | $\stackrel{\infty}{\stackrel{\infty}{\sim}}$ | $\begin{gathered} \underset{\sim}{\mathrm{O}} \end{gathered}$ |  | 太ু | N | $\infty$ |
|  |  |  | $\begin{aligned} & \text { N} \\ & \text { Niv } \end{aligned}$ | $\begin{aligned} & \tilde{m} \\ & \tilde{寸} \end{aligned}$ | $\begin{aligned} & \stackrel{\rightharpoonup}{\mathrm{N}} \\ & \infty \\ & \underset{\sim}{0} \end{aligned}$ | $\begin{aligned} & \stackrel{\circ}{\tilde{m}} \\ & \underset{\sim}{\alpha} \end{aligned}$ | $\begin{aligned} & \text { ö } \\ & \text { ¢i } \end{aligned}$ |  | $\begin{aligned} & \underset{\sim}{n} \\ & \underset{\sim}{n} \end{aligned}$ | $\begin{gathered} \stackrel{\circ}{0} \\ \underset{j}{2} \end{gathered}$ | $\begin{aligned} & \dot{\sigma} \\ & \underset{i}{\prime} \end{aligned}$ | $\begin{aligned} & \underset{\sim}{J} \\ & \underset{\sim}{n} \end{aligned}$ | $\begin{gathered} \text { n} \\ \text { in } \\ \text { in } \end{gathered}$ | $\begin{aligned} & \text { ひ్} \\ & \underset{\sim}{n} \end{aligned}$ | $\begin{aligned} & \hat{0} \\ & \dot{\sim} \\ & \dot{\delta} \end{aligned}$ |  |  | $\begin{gathered} \tilde{N} \\ \underset{\sim}{\hat{\alpha}} \end{gathered}$ | $\stackrel{\circ}{\stackrel{\circ}{g}}$ | － |
|  | $\begin{aligned} & 3 \\ & \stackrel{3}{5} \end{aligned}$ | $\because$ | $\stackrel{\sim}{\underset{\sim}{\mathrm{N}}}$ | $\underset{\sim}{\underset{\sim}{\sim}}$ | $\begin{gathered} \text { ñ } \\ \underset{\sim}{n} \end{gathered}$ | $\underset{\substack{\text { N } \\ \hline}}{ }$ | $\stackrel{\infty}{\stackrel{\infty}{\top}}$ |  | $\stackrel{\stackrel{\rightharpoonup}{\mathrm{N}}}{\substack{2}}$ | $\stackrel{\sim}{\underset{\sim}{i}}$ | N̈ | $\underset{\substack{\text { そ } \\ \text { in }}}{ }$ | $\stackrel{0}{0}$ | $\underset{\sim}{\sim}$ | $\begin{aligned} & \infty \\ & \infty \\ & \infty \\ & \hline \end{aligned}$ | $\underset{\sim}{\underset{\sim}{r}}$ | 卆 | $\underset{\sim}{\underset{\sim}{\mathrm{N}}}$ | N | $\overrightarrow{7}$ |
|  |  | $\begin{aligned} & \sum_{0}^{历} \\ & \sum \end{aligned}$ | $\begin{aligned} & \underset{\sim}{m} \\ & \underset{\sim}{j} \end{aligned}$ | $\begin{aligned} & \underset{\infty}{\underset{\sim}{\sim}} \\ & \underset{y}{c} \end{aligned}$ | $\begin{aligned} & \stackrel{\circ}{\underset{\sim}{2}} \\ & \underset{\sim}{\circ} \end{aligned}$ | $\begin{aligned} & \stackrel{\rightharpoonup}{m} \\ & \underset{\sim}{\dot{q}} \end{aligned}$ | $\begin{aligned} & \infty \\ & \underset{\sim}{\infty} \\ & \dot{q} \end{aligned}$ | $\begin{aligned} & \infty \\ & \stackrel{\infty}{\circ} \\ & \underset{y}{2} \end{aligned}$ | $\stackrel{\vec{N}}{\substack{\text { 号 }}}$ | $\begin{aligned} & \dot{\otimes} \\ & \underset{\sim}{g} \end{aligned}$ | $\begin{aligned} & \text { ơ } \\ & \text { in } \end{aligned}$ | $\begin{aligned} & \text { g } \\ & \dot{\text { g }} \end{aligned}$ | $\begin{aligned} & \infty \\ & \infty \\ & \text { in } \end{aligned}$ | $\begin{aligned} & \text { t } \\ & \text { in } \\ & \dot{\alpha} \end{aligned}$ | $\underset{\substack{\underset{\sim}{\sim} \\ \text { No }}}{ }$ | $\begin{aligned} & \underset{\sim}{\underset{N}{2}} \\ & \underset{\sim}{2} \end{aligned}$ | $\begin{aligned} & \underset{\sim}{\tilde{j}} \\ & \underset{\sim}{n} \end{aligned}$ | $\begin{aligned} & \vec{\infty} \\ & \underset{\sim}{\infty} \end{aligned}$ | $\begin{aligned} & \text { ơ } \\ & \dot{g} \end{aligned}$ | $\stackrel{0}{0}$ |
|  | $\xrightarrow{3}$ | 0 |  |  | $\begin{aligned} & \stackrel{\infty}{\infty} \\ & \underset{\sim}{j} \end{aligned}$ | $\begin{aligned} & N \\ & 0 \\ & 0 \end{aligned}$ |  |  | $\underset{\underset{i}{\mathrm{~N}}}{\stackrel{\rightharpoonup}{2}}$ | $\underset{\sim}{7}$ |  | $\begin{aligned} & \text { O} \\ & \stackrel{0}{0} \\ & \hline \end{aligned}$ | $\begin{aligned} & \text { ti } \\ & \text { No } \end{aligned}$ | $\stackrel{\text { ñ }}{\underset{\sim}{\mathrm{i}}}$ | $\underset{\underset{\sim}{\mathrm{N}}}{\substack{\underset{\sim}{2}}}$ |  | $\stackrel{ }{\substack{\circ \\ \hline}}$ | $\begin{aligned} & \text { ơn } \\ & \text { No } \end{aligned}$ | $\stackrel{\text { No }}{\substack{\text { ¢ }}}$ | N |
|  |  | $\stackrel{\text { IN }}{\stackrel{N}{0}}$ | $\stackrel{7}{0}$ | $\begin{gathered} \text { 㞧 } \\ \text { ơn } \end{gathered}$ | $\begin{aligned} & \stackrel{\Omega}{\infty} \\ & \underset{\sim}{\sim} \end{aligned}$ | $\underset{\infty}{\infty}$ | $\begin{aligned} & \text { O } \\ & \text { O } \\ & \infty \end{aligned}$ | $\stackrel{\infty}{\stackrel{\infty}{N}}$ | $\stackrel{\otimes}{\infty}$ | $\begin{gathered} \infty \\ \underset{\infty}{\infty} \\ \infty \end{gathered}$ |  | $\underset{\infty}{\underset{\sim}{\underset{\sim}{*}}}$ | $\underset{\sim}{\underset{\sim}{\sim}}$ | No | $\stackrel{\sim}{\infty}$ | $\underset{\infty}{\underset{\sim}{7}}$ | $\begin{aligned} & \stackrel{\otimes}{\otimes} \\ & \infty \\ & \infty \end{aligned}$ | $\stackrel{n}{\underset{\sim}{n}}$ | $\stackrel{0}{0}$ | ¢ |
|  | cre | i | $\stackrel{\underset{\sim}{\mathrm{N}}}{\substack{2}}$ | $\begin{aligned} & \circ \\ & \\ & \end{aligned}$ | $\begin{gathered} \stackrel{\circ}{0} \\ \text { on } \end{gathered}$ | $\begin{aligned} & \text { 隻 } \\ & \hline \end{aligned}$ | $\begin{aligned} & \text { గ్ } \\ & \text { On } \end{aligned}$ | $\begin{aligned} & 0 \\ & \text { ñ } \\ & \underset{\sim}{n} \end{aligned}$ | $\stackrel{\circ}{\grave{\circ}}$ | 祭 | $\stackrel{\text { F }}{\dot{\circ}}$ | $\stackrel{0}{\stackrel{0}{0}}$ | Nin | $\underset{\sim}{\tilde{m}}$ | $\begin{aligned} & \text { ® } \\ & 0 \\ & \hline 0 \end{aligned}$ | $\underset{\underset{\sim}{\sim}}{\underset{\sim}{n}}$ | $\begin{aligned} & \infty \\ & \stackrel{\infty}{0} \\ & \hline \end{aligned}$ | $\underset{\text { Ni }}{\text { N }}$ | $\begin{aligned} & \text { in } \\ & \text { in } \end{aligned}$ | ¢ |
|  |  | $\begin{aligned} & \text { 厄్ల } \\ & \stackrel{N}{\Sigma} \end{aligned}$ | $\stackrel{\circ}{\mathscr{\sigma}}$ | N్రু |  | $\underset{\sim}{\tilde{\infty}}$ | $\begin{aligned} & \infty \\ & \stackrel{\infty}{\infty} \\ & \infty \\ & \hline \end{aligned}$ | $\stackrel{\substack{\infty \\ 0}}{\substack{0}}$ | $\underset{\substack{\infty \\ \infty}}{\vec{\infty}}$ | $\stackrel{\hat{6}}{\underset{\sim}{e}}$ | $\stackrel{\stackrel{\rightharpoonup}{\circ}}{\stackrel{1}{2}}$ | $\stackrel{ }{\stackrel{\sim}{c}}$ | $\stackrel{0}{\infty}$ | 융 | $\underset{\substack{\text { of }}}{\overrightarrow{0}}$ | $\begin{gathered} \hat{N} \\ \text { in } \end{gathered}$ | $\underset{\substack{\tilde{N} \\ \infty}}{ }$ | $\underset{\infty}{\underset{\infty}{J}}$ | $\stackrel{\text { N}}{\substack{\text { n }}}$ | స్ర̆ |
|  | $\begin{aligned} & 3 \\ & \stackrel{3}{5} \end{aligned}$ | 0 | $\begin{aligned} & \text { J } \\ & \text { UO } \end{aligned}$ | $\begin{aligned} & \text { N} \\ & \underset{\sim}{\widehat{1}} \end{aligned}$ | $\stackrel{m}{\infty}$ | $\begin{aligned} & \text { ® } \\ & \text { Mo } \\ & \hline \end{aligned}$ | $\begin{aligned} & \infty \\ & \infty \\ & \infty \\ & 0 \end{aligned}$ | $\begin{aligned} & 0 \\ & \vdots \\ & 0 \end{aligned}$ | $\begin{aligned} & \stackrel{\infty}{\bullet} \\ & \hline 0 \end{aligned}$ | $\underset{\sim}{n}$ | $\underset{\sim}{7}$ | $\begin{aligned} & \infty \\ & \infty \\ & \infty \\ & \hline \end{aligned}$ | $\begin{aligned} & \infty \\ & \infty \\ & \infty \end{aligned}$ | $\stackrel{\stackrel{\rightharpoonup}{\mathrm{O}}}{\substack{0}}$ | $\begin{aligned} & \vec{\alpha} \\ & 0 \\ & 0 \end{aligned}$ | $\begin{aligned} & \text { 乌 } \\ & \infty \\ & \hline \end{aligned}$ | $\begin{gathered} \cong \\ \\ \end{gathered}$ | 충 | ホ̇ | － |
|  |  | $\begin{aligned} & \stackrel{\Gamma}{0} \\ & \stackrel{N}{\Sigma} \end{aligned}$ |  | $\begin{aligned} & 0 \\ & \infty \\ & \infty \\ & \end{aligned}$ | $\begin{aligned} & \infty \\ & \stackrel{\infty}{O} \\ & \underset{\sim}{2} \end{aligned}$ | $\underset{\sim}{\underset{\infty}{\sim}}$ | $\underset{\infty}{\underset{\infty}{\mathrm{L}}}$ | $\underset{\infty}{\underset{\infty}{\sim}}$ | $\stackrel{\underset{\infty}{\infty}}{\stackrel{1}{\infty}}$ | $\underset{\infty}{\underset{\infty}{\infty}}$ | $\stackrel{N}{\hat{\circ}}$ | $\underset{\infty}{\underset{\infty}{\infty}}$ | 苞 | $\underset{\substack{\text { on }}}{\substack{\text { N}}}$ | $\stackrel{\circ}{i}$ | $\underset{\sim}{\tilde{\infty}}$ | $\underset{\sim}{\tilde{\infty}}$ | $\stackrel{\underset{N}{\mathrm{~N}}}{ }$ | $\stackrel{\sim}{\infty}$ | $\stackrel{7}{6}$ |
|  | ¿ |  | $\stackrel{\text { ® }}{ }$ | A | $\underset{\sim}{ \pm}$ | $\checkmark$ | $\stackrel{\sim}{\square}$ | ～ | ～ | $\stackrel{\text { a }}{ }$ | $\stackrel{\text { ® }}{ }$ | $\stackrel{\sim}{\square}$ | $\stackrel{\text {－}}{\sim}$ | － | $\stackrel{\sim}{\sim}$ | $\stackrel{\text { 가 }}{ }$ | ̇ | $\underset{\sim}{\sim}$ | $\stackrel{\rightharpoonup}{\sim}$ | $\stackrel{\sim}{\sim}$ |
|  |  |  | 讯 |  |  |  |  |  |  |  | － |  |  |  |  |  |  |  |  |  |





| E. vag | tum | NDF (g/g DM) |  |  |  |  |  | ADF (g/g DM) |  |  |  |  |  | Cellulose (g/g DM) |  |  |  |  |  | Lignin (g/g DM) |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Year | DOY | - Snow |  | Ambient |  | + Snow |  | - Snow |  | Ambient |  | + Snow |  | - Snow |  | Ambient |  | + Snow |  | -Snow |  | Ambient |  | + Snow |  |
|  |  | 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 | 0.096 | 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 | 0.009 | 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 | 0.009 | 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 | 0.690 | 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 |




