

1 **Seasonal shifts in depth to water uptake by young thinned and overstocked lodgepole pine**
2 **(*Pinus contorta*) forests under drought conditions in the Okanagan Valley, British**
3 **Columbia, Canada**

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5 Emory C. Ellis¹, Robert D. Guy², Xiaohua A. Wei³

6 ¹School of Forestry, Northern Arizona University, Flagstaff, Arizona, 86001, USA

7 ²Department of Forestry and Conservation Sciences, University of British Columbia, Vancouver, British Columbia,
8 V6T1Z4, Canada

9 ³Department of Earth, Environmental and Geographic Sciences, University of British Columbia (Okanagan
10 Campus), Kelowna, British Columbia, V1V 1V7, Canada

11 Correspondence to: Emory C. Ellis (ece58@nau.edu)

12 **Abstract:**

13 As drought and prolonged water stress become more prevalent in dry regions under climate
14 change, preserving water resources has become a focal point for maintaining forest health. Forest
15 regeneration after forest loss or disturbance can lead to over-stocked juvenile stands with high
16 water demands and low water-use efficiency. Forest thinning is a common practice with the goal
17 of improving tree health, carbon storage, and water use while decreasing stand demands in arid
18 and semi-arid regions. However, little is known about the impacts of stand density on seasonal
19 variation in depth to water uptake nor the magnitude of the effect of growing season drought
20 conditions on water availability. Existing reports are highly variable by climatic region, species,
21 and thinning intensity. In this study, stable isotope ratios of deuterium ($\delta^2\text{H}$) and oxygen ($\delta^{18}\text{O}$)
22 in water collected from soil varying depths and from branches of lodgepole pine (*Pinus contorta*)
23 under different degrees of thinning (control: 27,000 stems per ha; moderately thinned: 4,500
24 stems per ha; heavily thinned: 1,100 stems per ha) over the growing season were analyzed using
25 the MixSIAR Bayesian mixing model to calculate the relative contributions of different water
26 sources in the Okanagan Valley in the interior of British Columbia, Canada. We found that under
27 drought conditions the lodgepole pine trees shifted their depth to water uptake through the
28 growing season (June to October), to rely more heavily on older precipitation events that
29 percolated through the soil profile when shallow soil water became less accessible. Decreased
30 forest density subsequent to forest thinning did not cause a significant difference in isotopic
31 composition of branch water but did cause changes in the timing and relative proportion of water
32 utilized from different depths. Thinned lodgepole pines stands were able to maintain water
33 uptake from 35 cm below the soil profile whereas the overstocked stands relied on a larger
34 proportion of deep soil water and groundwater towards the end of the growing season. Our
35 results support other findings by indicating that although lodgepole pines are drought tolerant
36 and have dimorphic root systems, they did not shift back from deep water sources to shallow soil
37 water when soil water availability increased following precipitation events at the end of the
38 growing season.

39 **Keywords:** *Pinus contorta*; stable water isotopes; forest thinning; water-use strategies;
40 preferential water uptake; dual-isotope analysis; Bayesian isotope mixing model; soil water
41 uptake; transpiration; the interior of British Columbia

42 1. Introduction

43 As forests recover after harvesting, carbon and water demands change, and future climate
44 projections of increased drought severity will further complicate biogeochemical cycling and
45 carbon-water trade-offs (Giles-Hansen et al., 2021; Wang et al., 2019). Overpopulated
46 regenerating stands can add further stress on ecosystems; for example, light competition in dense
47 juvenile stands increases stand water demands by driving vertical growth and canopy cover (Liu
48 et al., 2011). To mitigate this stress, management strategies such as systemic thinning of high-
49 density juvenile stands have been shown to promote forest regeneration while decreasing
50 competition and providing remaining vegetation with increased light availability, rooting space,
51 nutrient access, and space for horizontal branch growth (Giuggiola et al., 2016). Over a variety
52 of forest ecosystems, reductions in stand density increase light availability, tree water use, carbon
53 storage, and water-use efficiency, an indication of improved tree health, and to decrease stand
54 water use, reducing the intensity of water stress under drought conditions (Belmonte et al., 2022;
55 Fernandes et al., 2016; Giuggiola et al., 2016; Liu et al., 2011; Manrique-Alba et al., 2020;
56 Molina & del Campo, 2012; Park et al., 2018; Sohn et al., 2012, 2016; Wang et al., 2019).
57 Because the primary goal of forest thinning is to decrease stand water use and increase
58 productivity, papers reporting the effects of this management strategy often focus on changes in
59 carbon storage, tree growth, transpiration, and water-use efficiency (Giuggiola et al., 2016;
60 Manrique-Alba et al., 2020; Park et al., 2018; Sohn et al., 2016). However, few studies have
61 reported sources of water use for vegetation water uptake and shifts in depth to water uptake in
62 association with thinning treatments in overstocked naturally regenerating forests, particularly
63 under drought conditions.

64 Quantifying stand water use is imperative to predicting the future of water availability in our
65 ecosystems. However, various studies indicate that trees do not always use the most recent
66 precipitation, and that vegetation can utilize different sources of water at different soil depths
67 depending on availability or stress (Dawson & Pate, 1996; Grossiord et al., 2017; Wang et al.,
68 2017). Many studies also report the depth of water uptake of various species and the relationship
69 between co-existing species and shared water sources (Andrews et al., 2012; Brinkmann et al.,
70 2019; Grossiord et al., 2017; Langs et al., 2020; Liu et al., 2015; Maier et al., 2019; Meinzer et
71 al., 2007; Sánchez-Pérez et al., 2008; Szymczak et al., 2020; Wang et al., 2017; Warren et al.,
72 2005). In water-limited regions such as arid and semi-arid landscapes, some species have
73 adapted to derive water from various depths over time depending on seasonal water variability,
74 indicating higher ecological plasticity and drought tolerance (Langs et al., 2020; Wang et al.,
75 2017). Understanding where in the soil profile plants obtain water, over prolonged dry periods
76 and at different stand densities, is essential in assessing the impact of forest thinning and the
77 relative importance of different seasonal water sources during shifts in water availability in arid
78 regions and under future climate conditions (Evaristo et al., 2015; Prieto et al., 2012; Sohn et al.,
79 2016). The implications of depth to water uptake and seasonal changes in water utilization, in
80 conjunction with water-use efficiency, can emphasize the importance of the timing and volume
81 of precipitation events and primary contributors to vegetation water use.

82 Stable isotope ratios can be used as powerful natural tracers to identify distinct water sources
83 such as rainfall, snow, and groundwater (Brinkmann et al., 2018; Lin & Sternberg, 1993;
84 Sprenger et al., 2017; Stumpp et al., 2018). The isotopic signature of precipitation events is
85 altered by elevation, temperature, and evaporative fractionation creating distinctive layers within

86 the soil profile (Kleine et al., 2020; Sprenger et al., 2017; Stumpp et al., 2018). More
87 specifically, soil water reflects precipitation events as they infiltrate through the soil layer with
88 the influence of evaporative fractionation until mixing with older soil water and groundwater and
89 creating individualized water isotopic signatures throughout the soil profile (Andrews & Science,
90 2009; Brinkmann et al., 2018; Dawson & Pate, 1996; Sprenger et al., 2017; Stumpp et al., 2018).
91 The isotopic composition of plant water can correspond to the water uptake depth in the soil
92 profile (Brinkmann et al., 2019; Langs et al., 2020; Meinzer et al., 2007; Stumpp et al., 2018;
93 Wang et al., 2017). Due to these unique characteristics, stable water isotopes have been used by
94 researchers to assess sources of water used by plants and their possible shifts under altered
95 environmental conditions (Evaristo et al., 2015; Flanagan & Ehleringer, 1991; Meinzer et al.,
96 2001; Stumpp et al., 2018).

97 Lodgepole pine (*Pinus contorta* Douglas) is an early successional montane conifer with a deep
98 tap root, fine roots in shallow soil layers, and an adventitious rooting system which allow this
99 species to access water throughout the soil profile (Fahey & Knight, 1986; Halter & Chanway,
100 1993). Depending on the species, root structures have two main components; namely, lateral
101 roots to exploit soil near the surface, and, in species with dimorphic root systems, sinker roots or
102 a well-developed tap root to reach deeper soil water or groundwater when surface water is
103 limited. Species with dimorphic rooting systems can access water from different depths in the
104 soil profile depending on soil moisture content and water availability, making them more
105 resilient to water scarcity or prolonged drought conditions (Dawson & Pate, 1996; Meinzer et al.,
106 2013). Wang et al. (2019) studied the short-term effects of thinning overstocked juvenile (16-
107 year-old) lodgepole pine stands in the Upper Penticton Creek Watershed, British Columbia,
108 Canada, and found a significant positive relationship between growth and water use from
109 decreased stand density and that heavily thinned treatments showed the most drought resistance.
110 Andrews et al. (2012) compared water uptake strategies between Douglas-fir (*Pseudotsuga*
111 *menziesii* (Mirb.) Franco) and lodgepole pine in southern Alberta, and found that lodgepole pines
112 are able to minimize seasonal variations in stem water potential and that tap roots are deep
113 enough to access groundwater. These findings are consistent with other literature reporting that
114 decreased stem density can improve water-use efficiency and that conifer trees can access water
115 from different depths depending on moisture availability (Meinzer et al., 2007a; Warren et al.,
116 2005). The literature therefore indicates that lodgepole pines can access water from different soil
117 layers even under extreme or prolonged drought conditions, but little is known about the shifting
118 of water use under different stand densities as a result of thinning treatments and drought
119 conditions.

120 In this study, we build on the research from Wang et al. (2019) which looked at the effects of
121 thinning on water-use efficiency during a drought and non-drought year by analyzing the stable
122 isotope ratios ($\delta^2\text{H}$ and $\delta^{18}\text{O}$) of soil and xylem water to evaluate at what depths overstocked and
123 thinned stands access water over a growing season to further our understanding of the
124 ecosystem-level impacts of thinning as a management strategy. We hypothesized that lodgepole
125 pine primarily relies on spring snowmelt, but reductions in shallow source water during the
126 growing season would drive trees to utilize deeper sources of water as the season progressed. We
127 also hypothesized that decreased stand density (thinning) would increase shallow soil
128 evaporation due to decreased canopy cover, but also decrease competitive limitations in tree
129 rooting zones so that at lower densities trees could better maintain mid-level soil water uptake.

130 Through a detailed partitioning of tree water sources, we can better understand how lodgepole
 131 pine uses water, estimate proportional dependence of lodgepole pine on specific source waters,
 132 and determine if thinning affects tree water use and uptake strategies under drought conditions.

133

134 2. Methods

135 2.1. Study site

136 The study was conducted in the Upper Penticton Creek experimental watershed (UPC) northeast
 137 of Penticton in the interior of British Columbia, Canada (49°39'34" N, 119°24',34" W). The site
 138 elevation is approximately 1675 m with steep, rocky terrain and a southern aspect (Wang et al.,
 139 2019). The luvisolic soils were formed from granite; the texture is coarse sandy-loam and is well
 140 drained with a low water holding capacity (Hope, 2011; Winkler et al., 2021; Winkler & Moore,
 141 2006). The biogeoclimatic region is the Engelmann Spruce-Subalpine Fir zone with cold, snowy
 142 conditions from November to early
 143 June and seasonal drought
 144 conditions during the summer
 145 months, June to October (Coupe et
 146 al., 1991; Wang et al., 2019). This
 147 research site was initially
 148 established as a paired watershed
 149 experiment in the early 1980s to
 150 quantify the impact of forest
 151 harvesting on water resources
 152 (Creed et al., 2014; Moore &
 153 Wondzell, 2005; Winkler et al.,
 154 2021).

155 The juvenile thinning experiment
 156 began in 2016 when 16-year-old,
 157 evenly aged, regenerating lodgepole
 158 pine stands were thinned to different
 159 densities than a control (Control - C:
 160 27,000 stem ha⁻¹, T1: 4,500 stems ha⁻¹, and T2: 1,100 stems ha⁻¹) where C represents the control
 161 stands, T1 represents the moderately thinned stands, and T2 represents the heavily thinned stands
 162 (Figure 1). The three treatments were repeated across three replicate blocks. Each block was 75
 163 m long and 25 m in width with three 20 m² plots and 5 m between treatment plots. After the
 164 initial thinning, all debris was left on site.

165 2.2. Climate and soil moisture monitoring

166 Climate stations (HOBO weather station, Onset Computer, Bourne MA, USA) were deployed
 167 across Block 1 treatments and have measured meteorological data since 2016 (ambient
 168 temperature, relative humidity (rH), wind speed, precipitation, and solar radiation) in 10-minute
 169 intervals. From these data, we calculated daily vapor pressure deficit (VPD) as well as daily and
 170 monthly potential evapotranspiration (PET) (Flint & Childs, 1991; Russell, 1960; Streck, 2003).

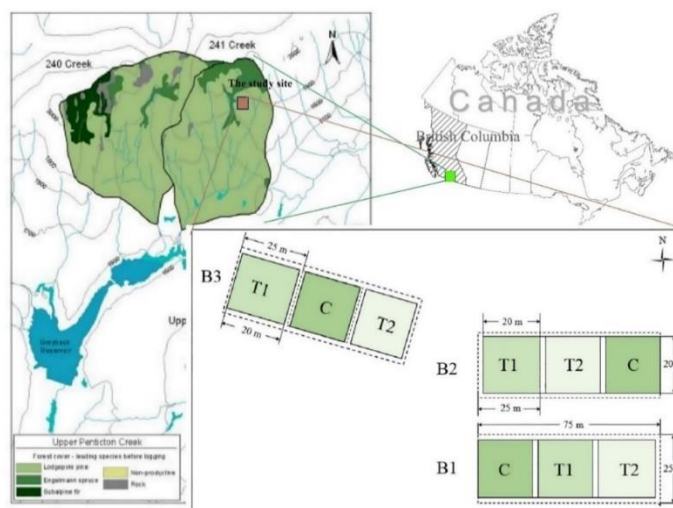


Figure 1 Watershed location and treatment plots of moderately thinned (T1), heavily thinned (T2), and the controlled (C) over-populated stands across the three replicate blocks (Wang et al., 2019)

171 Recorded historical precipitation (1997-2008) was acquired from a long-term climate station in a
 172 lodgepole pine forest in the 241 experimental watershed (climate station P7) (Moore et al.,
 173 2021).

174 Rainfall and temperature data from Block 1 were related to historical data to calculate the
 175 monthly dryness (PET/P), standardized precipitation index (SPI), and standardized precipitation
 176 evapotranspiration index (SPEI) (Table S1) (Beguería et al., 2014; Stagge et al., 2014; Wu et al.,
 177 2005). In the middle of the growing season in 2021, four soil moisture probes (HOBO TEROS
 178 11 Soil Moisture/Temp Probes) were deployed in each treatment in Block 1 to measure changes
 179 in soil moisture and temperature at 5 cm and 35 cm at 15-minute increments (n=12).

180 2.3 Sample collection

181 We sampled three trees per treatment across the three blocks and three in an adjacent mature plot
 182 south of the study site four times over the 2021 growing season in approximately six-week
 183 intervals (June 11-12, July 21-22, September 10-11, and October 7-8) around noon to capture
 184 peak transpiration time (Table 1). We used a pole pruner to cut a mid-canopy branch in the live
 185 crown. We peeled the bark off branch segments with no needle coverage to remove outer bark
 186 and phloem, placed them into 10 mL glass tubes that were then with Parafilm® wrap, covered in
 187 aluminum foil, and set in a cooler until the end of the day when they were transferred to a freezer
 188 at -18°C. During the last two sampling periods, some trees had red needles, likely an indication
 189 of dryness or higher temperatures from an early growing season heat dome that began in June.

190 *Table 1 Overview of the branch, soil, and precipitation samples collected over the four sampling periods during the*
 191 *2021 growing season and additional campaigns to collect groundwater and stream water.*

Sampling Period		1	2	3	NA	4
Sampling Date		June 11- 12	July 21- 22	September 10- 11	October 1	October 9
Sample Type	Branches	33	33	33	0	33
	5	9	9	9	0	9
	20	0	6	0	0	0
	35	9	9	9	0	9
	40	0	6	0	0	0
	60	0	6	0	0	0
	80	0	6	0	0	0
	100	0	6	0	0	0
	Rain	1	0	1	0	0
	Precipitation Snow	1	0	0	0	1
Stream	0	0	0	8	0	
Groundwater	0	0	0	6	0	

192

193 Soil samples were collected horizontally from 40 cm soil pits randomly dug within each
 194 treatment plot at 5 and 35 cm depths from the surface from June to October of 2021. Large rocks
 195 were removed from the profile. We conducted soil ribbon field tests to ensure that clay
 196 composition was less than 10% (soil ribbons were less than 20 mm in length). Soils were taken
 197 directly from the pit, then sealed in freezer seal bags and frozen until cryogenic distillation for

198 water extraction. In July, 1 m pits were dug. From the vertical pit, samples were collected in 20
199 cm increments to determine the depth of tree water access. After samples were collected, the
200 larger rocks and soils were used to fill the pits. We assumed that the isotopic signature of soil
201 water below 40 cm would be similar throughout the growing season and would be representative
202 of deep soil water. Soil samples were stored in a freezer at -18°C until cryogenically distilled.

203 Precipitation samples were collected cumulatively over individual field collection days where
204 precipitation was present (Table 1). Snow from a late spring event was collected on June 11th to
205 represent snow water isotopic composition during the sublimation and melt period of early 2021.
206 Another snow event was collected on October 11th during an active snowfall. A rain event was
207 collected on September 10th. Groundwater and stream samples were collected from the creek 241
208 watershed in early October 2021 at the beginning of the seasonal hydraulic recovery period
209 (Table 1). Groundwater was collected using a hand pump. Groundwater and stream samples
210 were collected at the end of the growing season as stream beds were dry and groundwater was
211 inaccessible during the dry period. Once the well had been pumped and cleared, 10 mL glass test
212 tubes were rinsed with ground water three times before being filled. Precipitation, groundwater
213 and stream samples were collected into 10 mL glass test tubes, sealed with Parafilm® and foil,
214 and stored in a fridge at 4°C .

215 2.4 Cryogenic extraction and isotopic analysis

216 Before extraction, branch samples remained sealed and were weighed in the glass test tubes used
217 for field collection. Branches remained in the test tubes until cryogenic distillation was complete
218 to ensure that any liquid water lost from the branch to the test tube was contained in the extract.
219 Soils samples were mixed in the Ziploc® bag, weighed, and transferred to a glass round bottom
220 flask. For stable isotope analysis, water was extracted from stem and soil samples using
221 cryogenic distillation (Orlowski et al., 2013; Percy et al., 2012). The test tube and branch
222 sample segment of the line was immersed in liquid nitrogen for 10 minutes until frozen
223 (Chillakuru, 2009). Soil sample size for extraction was roughly determined based on the
224 expected moisture of the frozen sample and soil moisture readings from continuous
225 measurements in the field. Soils were frozen for 45 minutes in a 500 mL round-bottom flask
226 using a dry-ice and 95% ethanol mixture before pumping out the air. Frozen samples were
227 pumped down to 60 mTorr, not disturbing the sample (Tsuruta et al., 2019). The vacuum-sealed
228 extraction unit was detached from the pump and transferred to a boiling water bath; the
229 extraction tube was submerged in liquid nitrogen. Branch samples were set to distill for 1 hour
230 and soil samples for 2 hours or until the tubing was clear to ensure all mobile and bound source
231 water was extracted (Orlowski et al., 2013; Tsuruta et al., 2019; Vargas et al., 2017; West et al.,
232 2006). As reviewed by Allen & Kirchner (2022), the cryogenic vacuum distillation of water from
233 plant tissues and soils can cause systematic biases in the measurements of $\delta^2\text{H}$. The degree of
234 extraction bias varies depending on species and soil type (Allen & Kirchner, 2022). In contrast,
235 bias in $\delta^{18}\text{O}$ values is close to zero (Allen & Kirchner, 2022). Reported biases in $\delta^2\text{H}$ average
236 about -6.1‰ for xylem water and -4‰ for water extracted from sandy soils, such as the soils
237 sampled here, which are of similar magnitude (Allen & Kirchner, 2022). Therefore, although we
238 used cryogenic vacuum distillation to extract water from xylem and soil media, potential
239 systematic bias introduced during the extraction process was treated as negligible as all sources
240 we identified had a difference in $\delta^2\text{H}$ greater than 4‰ (with the minimum distance being 14‰

241 between groundwater samples and deep soil water), minimizing any major effects on partitioning
 242 calculations.

243 The volume of branch water extracted ranged from 1 to 3 mL depending on the size of the
 244 branch sample. Total extracted water varied dependent on the mass of the initial sample. The
 245 volume of soil water extract ranged from 1mL to 7 mL depending on the size of the sample
 246 prepared for extraction. Samples were also weighed after extraction and compared to oven dried
 247 samples to ensure distillation was complete. Water extracted from branch and soil samples
 248 accounted for $47.9 \pm 3.2\%$ and $9 \pm 6\%$ of mean sample weight \pm standard deviation.

249 All samples were pipetted and sealed into glass vials with screw tops and shipped to the
 250 University of California Davis Stable Isotope Facility (Davis, CA, USA) for ^{18}O and ^2H analysis
 251 using headspace gas equilibration on a GasBench-II interfaced to a Delta Plus XL isotope-ratio
 252 mass spectrometer (Thermo-Finnigan, Bremen, Germany) normalized to a range of secondary
 253 reference waters calibrated against three IAEA standard waters. Precision was less than or equal
 254 to 2.0‰ for $\delta^2\text{H}$ and 0.2‰ for $\delta^{18}\text{O}$. Results were returned in the "delta" notation expressing the
 255 isotopic composition of each sample as a ratio in parts per thousand, relative to VSMOW
 256 (Vienna-Standard Mean Ocean Water) where:

$$257 \quad \delta(\text{‰}) = \left(\frac{R_{\text{Sample}}}{R_{\text{Standard}}} - 1 \right)$$

258 Sample extract was situated in an isotope biplot and compared to the global meteoric water line
 259 (GMWL) along with a local meteoric water line for the Okanagan Valley (OMWL) ($\delta^2\text{H} = 6.6$
 260 ($\delta^{18}\text{O} - 22.7$) and local evaporative line (LEL) ($\delta^2\text{H} = 5 (\delta^{18}\text{O}) - 48.4$) calculated for the
 261 Okanagan Valley by Wassenaar et al. (2011). The LEL is a linear regression that indicates the
 262 departure of water sources from the OMWL to indicate the degree of evaporative processes
 263 fractionating the isotopic composition of water sources or variance in the isotopic composition of
 264 seasonal precipitation events.

265 One extreme outlier of B1C at the 20 cm depth was removed before analysis; the high $\delta^2\text{H}$ and
 266 $\delta^{18}\text{O}$ values were likely due to contamination or incomplete cryogenic distillation. To test the
 267 variance between thinning treatments, block replicates, sampling periods, and soil depth, we first
 268 tested the assumption of normality in the subsets using the Shapiro-Wilk test and found that all
 269 subgroups were approximately normally distributed. Repeated measures ANOVAs were used to
 270 compare effects of date and treatment on $\delta^2\text{H}$ and $\delta^{18}\text{O}$ in branches, soils and groundwater to
 271 determine if changes in lodgepole pine uptake patterns occurred over time, if soil signatures
 272 varied between different depths (0-100 cm and groundwater) and densities, and if thinning
 273 juvenile stands changed seasonal shifts. All statistical analysis was conducted in R Studio
 274 (version 1.3.1073) using the appropriate tests to determine site distinctions and seasonal
 275 variability in depth to uptake (RStudio Team, 2020).

276 2.4 MixSIAR model scenarios

277 Process-based models (PBM) with a Bayesian approach include integrating other processes or
 278 existing information as priors allowing for a more informed approach than a simple linear model
 279 (Ogle et al., 2014). To accurately partition potential lodgepole pine water sources, we used the
 280 MixSIAR modeling package, a Bayesian mixing model (BMM) based on the Markov Chain

281 Monte Carlo method (MCMC) (Langs et al., 2020; Stock, 2013/2022, p. 201; Stock et al., 2018;
 282 Wang et al., 2017; Wang et al., 2019). The MixSIAR modeling package was selected over
 283 previous iterations of the dual-isotope BMM (SIAR and Simmr) and other partitioning models
 284 because of the accuracy in the analysis of covariates and the ability of the model to include
 285 source-specific uncertainties and discrimination factors (Stock et al., 2018; Wang et al., 2017).
 286 We partitioned potential water sources for five different scenarios using a combination of single
 287 and dual isotope approaches and different potential sources: scenario 1 – single isotope $\delta^{18}\text{O}$ two
 288 sources 5 cm and 35 cm depth; scenario 2 – single-isotope $\delta^2\text{H}$ two sources 5 cm and 35 cm
 289 depth; scenario 3 – dual-isotope two sources 5 cm and 35 cm depth; scenario 4 – dual isotope
 290 three sources 5 cm, 35 cm and 45-100 cm depth; scenario 5 – dual isotope three sources 5 cm,
 291 35-100 cm and groundwater; and scenario 6 – dual isotope four sources 5 cm, 35 cm, 45-100 cm
 292 and groundwater. In scenarios using deep
 293 soil water (35-100 cm depths), the isotopic
 294 composition was calculated as a weighted
 295 average between seasonally collected soil
 296 water from depth 35 and average soil
 297 water at depths collected in 202 cm
 298 intervals during the early growing season
 299 ($n=38$ per season). There were no source
 300 concentration dependencies, and the
 301 discrimination was set to zero for both
 302 isotopes in the analysis. The run length of
 303 the Markov chain Monte Carlo (MCMC)
 304 was set to ‘normal’ (chain length =
 305 100,000; burn =50,000; thin = 50; chains
 306 = 3). The Gelman-Rubin and Geweke
 307 diagnostic tests included in the model
 308 package were used to determine
 309 convergence (Gelman-Rubin score <
 310 1.01). Scenarios that did not converge
 311 were run again with a longer runtime (chain
 312 length: 300,000; burn: 200,000; thin: 100;
 313 chains = 3). No priors were used, so each
 314 water source was considered equally ($\alpha =$
 315 1).

316

317 3. Results

318 3.1. Climate and soil water content

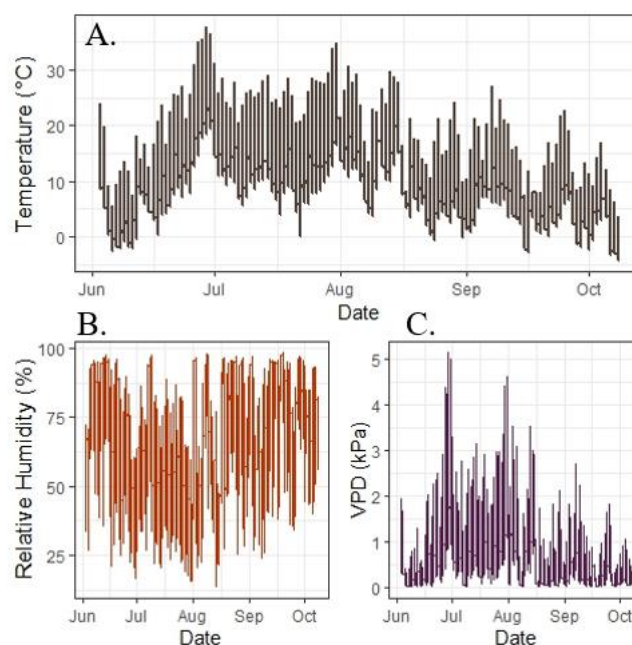


Figure 3 15-minute measurements of A. atmospheric temperature ($^{\circ}\text{C}$), B. Relative humidity (%), and C. vapor pressure deficit (VPD) (kPa).

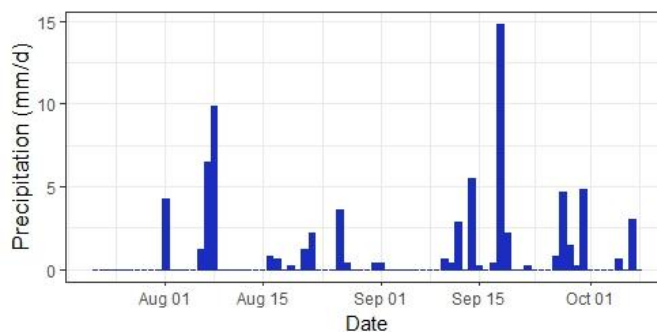


Figure 2 Rainfall (mm/d) from July 22 to October 8, 2021.

319 The ambient temperature peaked in
 320 the moderately thinned plot (T1)
 321 on June 29th with a maximum
 322 temperature of 36.3°C in an
 323 abnormally hot and dry summer
 324 (Figure 2). Relative humidity and
 325 VPD recorded in T1 showed the
 326 most variability and highest
 327 evaporative capacity during July.
 328 Atmospheric water vapor was
 329 higher in late September and
 330 October when precipitation was
 331 more frequent, and the watershed
 332 began to exhibit traits of
 333 hydrologic recovery (Figure 3).
 334 One indication of increased water
 335 availability was increased soil
 336 moisture at 5 cm and 35 cm depths
 337 and more groundwater recharge in
 338 October (Figure 4). There was
 339 17.5 mm of precipitation from
 340 September 16th to 18th that
 341 infiltrated to at least 35 cm below
 342 the soil surface along with
 343 subsequent rainfall events that likely infiltrated past the 35 cm sample depth changing the
 344 isotopic composition of deep soil water from what was measured during the deep pit sampling in
 345 July.

346 Rainfall events recorded at a nearby long-term research station between June to October from
 347 1997-2008 represented approximately 30.1% of annual precipitation (Winkler et al., 2021). Over
 348 the 2021 study period, there was 147.8 mm of rainfall, while the mean summer rainfall from
 349 1997 to 2008 was 232.5 mm, and most of the rainfall occurred in the early growing season. SPI
 350 and SPEI were significantly lower in 2021 than the mean historical range (Figure 5). Although
 351 there was precipitation and the beginning of hydraulic recovery in October, drought conditions
 352 persisted. Drought conditions of the study site reflected the drought conditions of the region as
 353 reported by Agriculture and Agri-Food Canada from June to August 2021 in moving from severe
 354 (level 2 drought) to exceptional (level 4) before recovering in September (Canada, 2014:

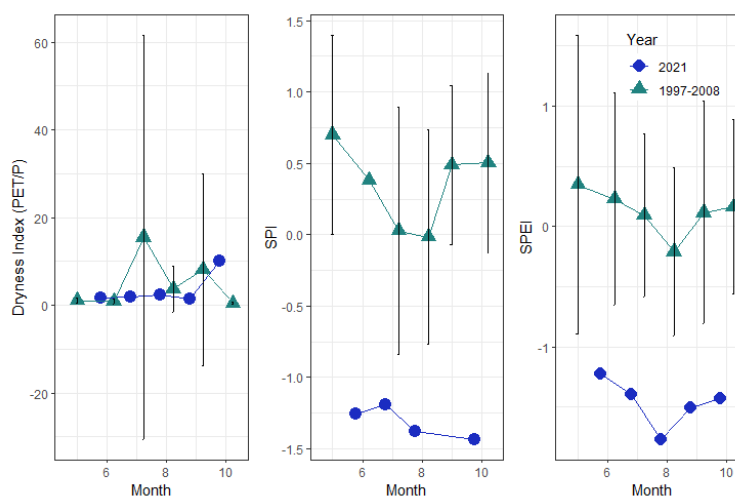


Figure 5 From left to right: dryness index (monthly PET using the Thornthwaite method divided by mean monthly precipitation) from June to October, 2021 and historic climate data from 1997 to 2008 including the including standard error for the historic climate data, standard precipitation index (SPI) with a 3-month period from June to October of 2021 and historic (1997-2008) climate data including standard error for the historic climate data, and standardized precipitation evapotranspiration index (SPEI) with a 3-month period from June to October of 2021 and historic (1997-2008) climate data including standard error for the historic climate data.

355 [https://agriculture.canada.ca/en/agricultural-zproduction/weather/canadian-drought-](https://agriculture.canada.ca/en/agricultural-zproduction/weather/canadian-drought-monitor/drought-analysis)
 356 [monitor/drought-](https://agriculture.canada.ca/en/agricultural-zproduction/weather/canadian-drought-monitor/drought-analysis)
 357 [analysis](https://agriculture.canada.ca/en/agricultural-zproduction/weather/canadian-drought-monitor/drought-analysis)).

358 3.2. Water stable isotopes

359 The biplot of sample
 360 isotopic composition
 361 shows the distribution and
 362 effect of isotopic
 363 fractionation on source
 364 water isotope ratios of
 365 samples collected during
 366 the 2021 field season.
 367 Field collected samples
 368 were compared to the
 369 Okanagan Meteoric
 370 Water Line (OMWL)
 371 (Wassenaar et al., 2011).

372 The slopes for branch and
 373 soil water were less steep
 374 than the OMWL, and the

375 intercepts more negative, indicating that evaporative fractionation contributed to the isotopic
 376 composition of these pools at the UPC (Figure 6). Soil samples seemed to follow the LEL
 377 produced by Wassenaar et al. (2011) for the region indicating similar evaporative fractionation
 378 effects. Branch water more closely following the OMWL than soils, suggesting that most
 379 samples consisted of water that was accessed from deeper in the soil profile and had infiltrated
 380 past the evaporative front. Precipitation samples collected during the field season fell along the
 381 OMWL (Wassenaar et al., 2011). The $\delta^2\text{H}$ and $\delta^{18}\text{O}$ of the June 11th rainfall event were -127.5‰
 382 and -13.03‰, respectively. The September rainfall event was much more enriched with a $\delta^2\text{H}$ of
 383 -38.4‰ and $\delta^{18}\text{O}$ of -2.89 (Figure 6). The snowfall collected on October 7th more closely
 384 resembled the lighter, colder, June precipitation event.

385 3.2.1. Soil moisture and seasonal water composition

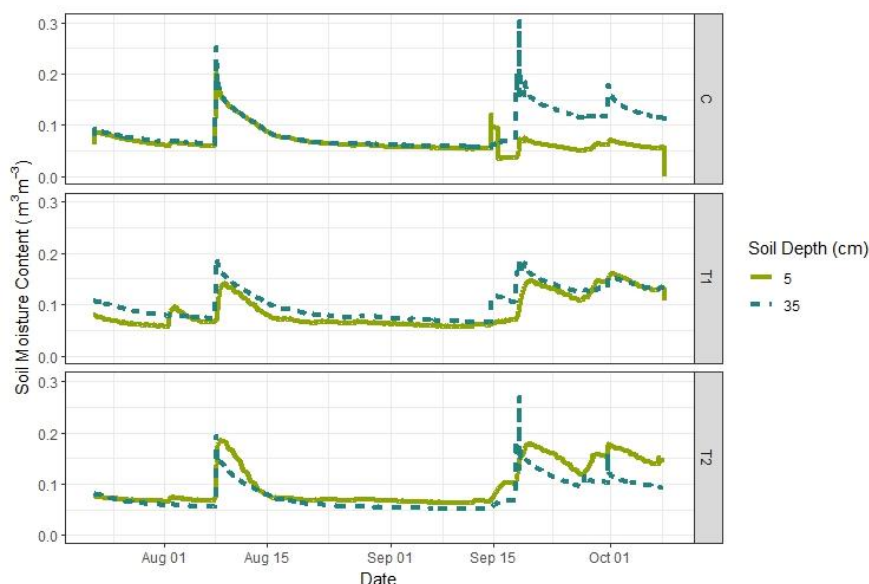


Figure 5 Average in-situ continuous measurements (15-minute interval) of soil water content (m^3/m^3) from the control, moderately thinned, and heavily thinned stands in Block 1.

386 Soil moisture probes and percent soil water content from samples collected for isotopic analysis
 387 were compared between treatments and deployment depths. Water content of soil samples was
 388 highest in June (21.5% at 5 cm and 21.6% at 35 cm) because of high snow melt and early spring
 389 precipitation, while soils were driest in September (6.32% at 5 cm and 6.19% at 35 cm).
 390 Continuous soil moisture measurements showed that soil water began to increase in mid-
 391 September as precipitation became more frequent, daily solar radiation decreased, and water
 392 percolated into deeper soil layers. There were significant differences in the continuously
 393 measured soil moisture by depths, treatments, and month, respectively (5-35 cm) (Depth: F
 394 $=3545.9$, $p < 2e-16^{***}$) (Treatment: $F=1883.3$, $p < 2e-16^{***}$) (Month: $F=3359.8$, $p < 2e-16^{***}$)
 395 (Figure 7), but soil water content of samples for isotopic analysis only varied significantly by
 396 month (August – October) ($F=22$, $p < 5.4e-9^{***}$).

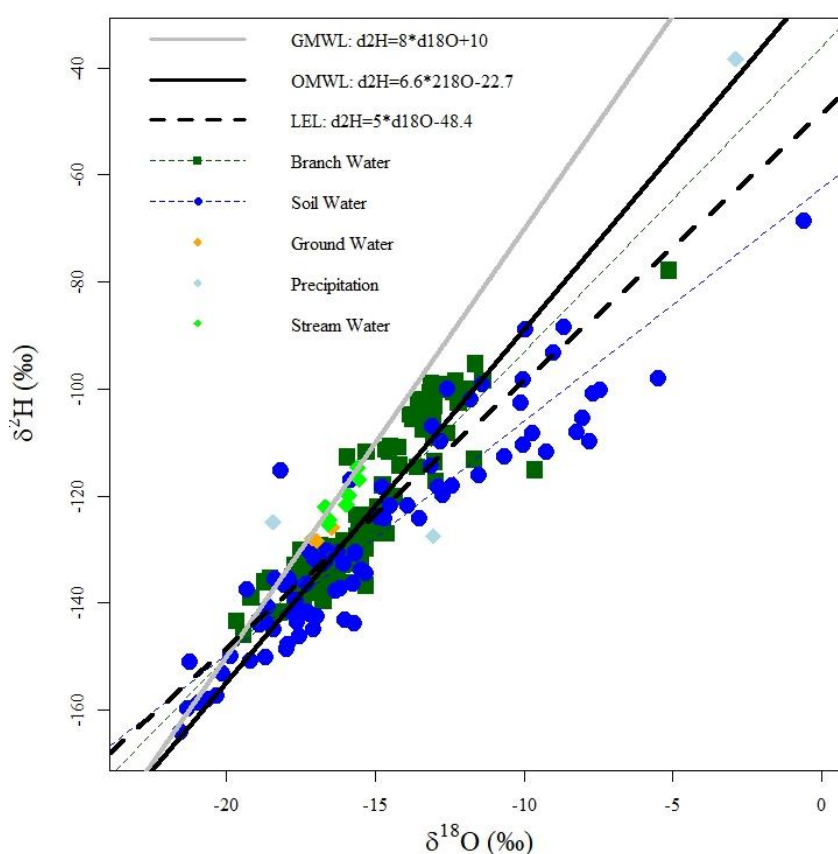


Figure 6 Biplot of $\delta^{18}\text{O}$ and $\delta^2\text{H}$ including all branch, soil, stream, groundwater, and precipitation samples collected over the 2021 study period outlined in Table 1 along with the global meteoric water line (GMWL) as well as a meteoric waterline for the Okanagan Valley (OMWL) and the Local Evaporative Line (LEL) developed by Wassenaar et al. (2011). Linear regressions are also plotted for branch water and soil water to indicate deviations in the slope and intercept from OMWL and LEL. The relationship between $\delta^{18}\text{O}$ and $\delta^2\text{H}$ in soil and branch water shows comparative ranges, but more variation among the soil water samples likely due to changes in precipitation signatures.

397 Soil isotopic results were broken into two datasets to analyze the variation in isotopic
 398 composition over time and between treatments, and then a profile of isotopic variance with depth

399 was constructed. Soil water $\delta^2\text{H}$ and $\delta^{18}\text{O}$ varied significantly by depth ($\delta^2\text{H}$: $p=2.57\text{e-}6^{***}$;
 400 $\delta^{18}\text{O}$: $p=2.45\text{e-}7^{***}$), being higher in the shallow soils than deeper in the profile (Figure 7.A.
 401 and 7.C.). $\delta^2\text{H}$ varied significantly across months ($p=2.72\text{e-}5^{**}$), but not between July and
 402 September and September and October. $\delta^{18}\text{O}$ also varied significantly across months ($p=1.5\text{e-}$
 403 5^{**}) except when directly comparing July to October and September to October. Despite
 404 treatment differences in soil moisture (Figure 4), there were no statistically significant treatment
 405 differences in the isotopic composition of soil water at either depth. In June, the mean soil water
 406 $\delta^{18}\text{O}$ at 5 cm was $-16.8\pm 2.57\text{‰}$ while the
 407 $\delta^2\text{H}$ was $-136.7\pm 13.6\text{‰}$; at 35 cm, the
 408 $\delta^{18}\text{O}$ was $-19.2\pm 1.52\text{‰}$ and $\delta^2\text{H}$ was -
 409 $149.2\pm 9.6\text{‰}$. Both $\delta^{18}\text{O}$ and $\delta^2\text{H}$
 410 increased more during the growing season
 411 at 5 cm than at 35 cm, and with more
 412 variability (Figure 7). In September, $\delta^{18}\text{O}$
 413 and $\delta^2\text{H}$ at 5 cm were -8.75‰ and -106.23
 414 and at 35 cm were -14.71‰ and -127.64
 415 respectively suggesting that soil isotopic
 416 composition nearer the soil surface
 417 follows trends in precipitation samples,
 418 being most enriched with O^{18} . By
 419 October, $\delta^{18}\text{O}$ and $\delta^2\text{H}$ at 5 cm reflected
 420 more recent precipitation events
 421 indicating that water availability in
 422 shallow soils began to increase.

423 From the isotopic soil profile, there were
 424 three significant groupings of isotopic
 425 composition ($p<0.05$): shallow soil water
 426 (5-20 cm), deep soil water (35-100 cm),
 427 and groundwater. Mean groundwater
 428 collected at the end of the growing season
 429 most closely resembled spring and fall
 430 snowfall events. The mean $\delta^{18}\text{O}$ of
 431 groundwater was $-16.82\pm 0.34\text{‰}$, which
 432 resembles that in the soil profile, but
 433 mean $\delta^2\text{H}$ was slightly higher than soil
 434 water ($n=4$). This isotope fractionation
 435 may be due to interactions with bound
 436 soil water and soils as the water infiltrates
 437 through the vadose zone, but the spread of values as potential uptake sources was greater than
 438 any predicted bias from cryogenic vacuum extraction therefor groundwater was included in the
 439 model as a isotopically distinct potential source for lodgepole pine water use (Allen & Kirchner,
 440 2022; Vargas et al., 2017).

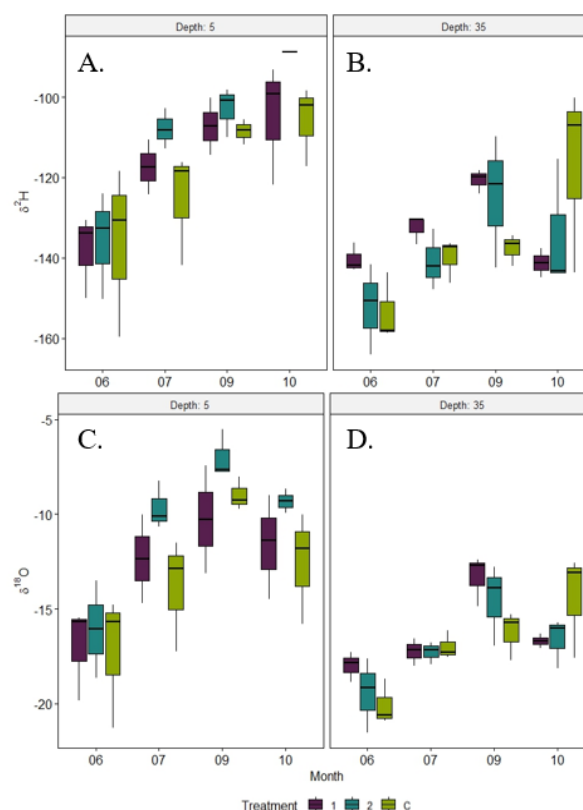


Figure 7 Boxplots of the soil water $\delta^2\text{H}$ (A. and B.) and $\delta^{18}\text{O}$ (C. and D.) at 5 (A. and C.) and 35 cm (B. and D.) depths collected four times over the growing season from each treatment and block. Mean, interquartile ranges, and standard deviation are indicated for each treatment in each month. There was a significant difference in the isotopic composition of water between months by treatment and depth indicating changes in water isotopic signature either due to evaporation or precipitation.

441 The more negative values for both $\delta^{18}\text{O}$ and $\delta^2\text{H}$ with soil depth indicate that snow melt is the
 442 main source of water to the deep unsaturated zone and that enriched summer precipitation is not
 443 infiltrating deeper soil layers (Figure 8).

444

445 3.2.2. Isotopic variability in 446 branch xylem water

447 Branch xylem for each
 448 treatment across the three
 449 blocks and the adjacent
 450 mature stand were compared
 451 for each sampling period. All
 452 treatments closely resembled
 453 the mature stand in both $\delta^{18}\text{O}$
 454 and $\delta^2\text{H}$. There were no
 455 statistically significant
 456 differences in both $\delta^{18}\text{O}$ and
 457 $\delta^2\text{H}$ of xylem water across
 458 thinning treatments; there
 459 was, however, significant
 460 variation over time ($\delta^{18}\text{O}$:
 461 $F=24.8^*$; $\delta^2\text{H}$: $F=146.6^*$).
 462 More specifically, $\delta^{18}\text{O}$ and
 463 $\delta^2\text{H}$ of xylem water varied by
 464 month for all months collected
 465 except for between June and
 466 September and July and
 467 September (Figure 9). Because the isotopic composition of xylem water showed significant
 468 change over the growing season but did not follow the same seasonal trends as soil water, the
 469 trees were likely changing their primary water source within the soil profile.

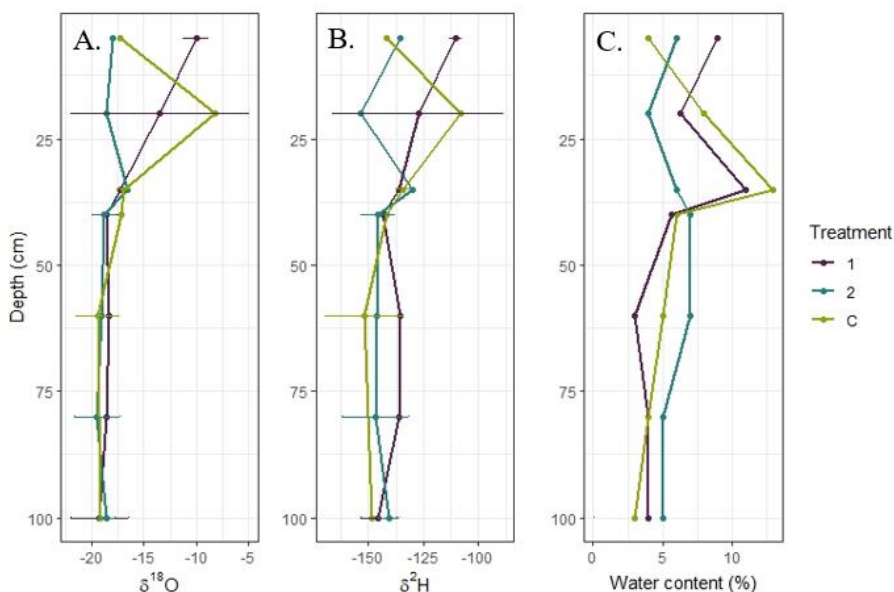


Figure 8 Vertical isotopic profiles and gravimetric soil water content from treatments in Block 2 and samples collected in mid-July where A. shows the vertical changes in $\delta^{18}\text{O}$ for each treatment, B. shows the vertical changes in $\delta^2\text{H}$ for each treatment, and C. shows the change in gravimetric water content as a percent of total soil weight.

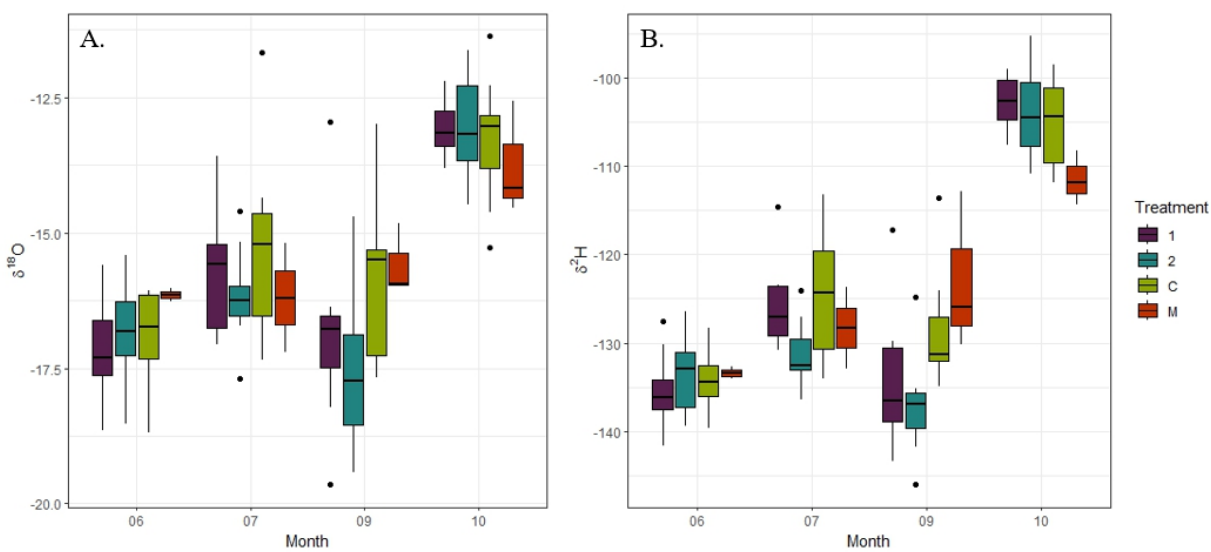


Figure 9 A boxplot showing branch mean, interquartile range, and standard deviation for A. $\delta^{18}\text{O}$ and B. $\delta^2\text{H}$ by month and treatments for the control (C), lightly thinned (T1), heavily thinned (T2), and mature (M) stands. Branch water was highest in October despite treatment effects. Mature trees were used as a reference for the isotopic composition of lodgepole pines over time but were not considered in the model of changes depth to water uptake over time. There was not statistically significant difference in $\delta^{18}\text{O}$ and $\delta^2\text{H}$ between treatments, but each treatment varied significantly by month with the highest concentration of heavy isotopes in October.

470 3.3. Partitioning xylem source water and seasonal fluxes using MixSIAR

471 Of the six scenarios considered, scenarios 1, 2 and 6 approached the Gelman-Rubin diagnostic
 472 (less than 1.05) with a runtime set to “normal” (chain length: 100,000; burn: 50,000; thin: 50;
 473 chains: 3), which indicates that they were the closest of all scenarios to reach convergence (Table
 474 S2). Out of the 6 potential scenarios, scenarios 4 (dual-isotope and 5 cm, 35 cm, and 45-100 cm
 475 soil water as sources) and 6 (dual-isotope 5 cm, 35 cm, 45-100 cm, and groundwater soil water
 476 as sources) were rerun with the run time set to “long” (chain length: 300,000; burn: 200,000;
 477 thin: 100; chains: 3) as they were hypothesized to provide the most representative results of
 478 water uptake partition from various depths. The Gelman-Rubin diagnostic for scenario 4 was
 479 120, and for scenario 6 was 17, when run for the “long” runtime, meaning scenario 6 was closer
 480 to convergence, but still greater than the convergence threshold.

481 Results of scenario 6 indicate that, in June, trees in each treatment acquired the most water from
 482 the 5 cm depth (C: 76%; T1: 77%; T2: 79%) (Figure 10). In July, shallow soil water was still the
 483 primary source for T1 and T2 at 47% and 61%, but C had 55% water from 45-100 cm deep and
 484 only 33% from 5 cm below the surface. By September, all treatments acquired less than 15% of
 485 tree water from shallow soil. Lodgepole pine water use in treatments 1 and 2 was composed of
 486 approximately 48% and 54% from around 35 cm, while 72% of water in control stand trees was
 487 from 35-100 cm. By October, although SPEI results indicate more moisture and less evaporative
 488 demand, scenario six indicated that all three treatments had most water uptake from below 45 cm
 489 in the soil profile (Figure 10). Results of the MixSIAR model support findings of branch water
 490 stable isotope trends over the growing season where the branch water started with mean $\delta^{18}\text{O}$ and
 491 $\delta^2\text{H}$ values of $-16.9 \pm 0.89\text{‰}$ and $-134.37 \pm 3.8\text{‰}$ in June, becoming slightly more enriched in July.

492 There was a shift to a source with a higher concentration of lighter isotopes in September.
 493 Branch water was most enriched with heavy isotopes in October, like shallow soil water, with
 494 mean $\delta^{18}\text{O}$ and $\delta^2\text{H}$ of $-12.9\pm 1.76\text{‰}$ and $-103.8\pm 7.0\text{‰}$, respectively. However, the MixSIAR
 495 model does not account for potential changes in the isotopic composition of water from
 496 precipitation events from mid-September to mid-October. Additionally, we did not consider
 497 extraction bias of the soil water sources nor branch water in the MixSIAR model because the
 498 previously mentioned range between distinct sources is larger than the potential change in
 499 isotopic signature during cryogenic distillation. The branch water in October was more enriched
 500 in heavy oxygen isotopes for each treatment than soil water at a depth of 35 cm and was more
 501 isotopically similar to soil water at 5 cm. Deuterium also followed a similar trend.

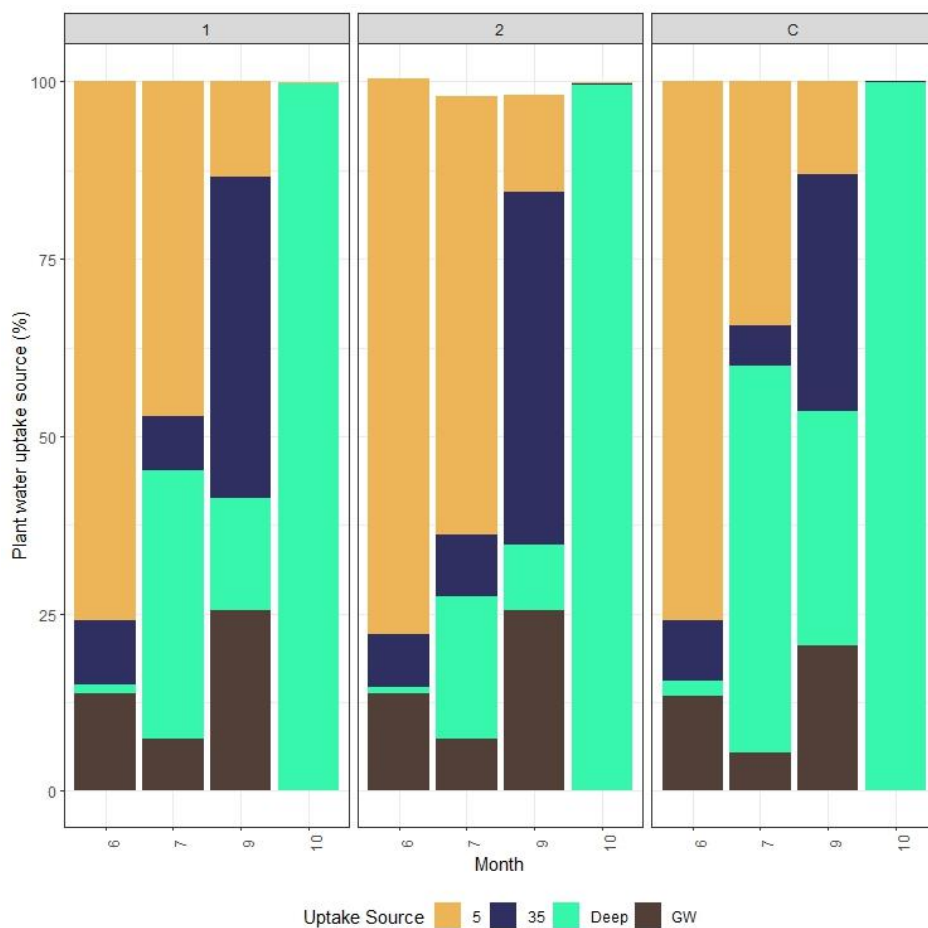


Figure 10 Stacked bar charts showing the partitioned relative contribution of different sources of water in the soil profile by the MixSIAR model of scenario 6 with long (chain length: 300,000; burn: 200,000; thin: 100; chains: 3) runtime. Scenario 6 considers both $\delta^{18}\text{O}$ and $\delta^2\text{H}$ as tracers and 5 cm, 35 cm, 45-100 cm, and groundwater as potential sources. Soil water isotopic composition at 5 cm and 35 cm changes monthly whereas the concentrations of $\delta^{18}\text{O}$ and $\delta^2\text{H}$ are held constant for the 45-100 cm and groundwater sources. Results of this model indicate that there are significant changes in the depth to water uptake of lodgepole pines between June and October of 2021 and that thinned trees can maintain a larger percentage of water uptake from shallow soil water longer than trees in the control stand.

503 4. Discussion

504 4.1. Seasonal variability in soil water

505 Deep soil water showed mixed gradient of older, more depleted, water molecules deeper in the
506 profile indicating that deep soil water mainly originates from spring snowmelt during the
507 summer months. Low intensity and less frequent summer precipitation events are evaporated out
508 of the shallow soil layers and do not infiltrate past the evaporative front to recharge the
509 unsaturated zone or groundwater. Although there was not a statistically significant difference in
510 the depth to water uptake by thinning treatments, the results of our isotopic analysis indicate that
511 there was increased evaporative enrichment, or a higher concentration of oxygen-18, in the
512 shallow soils of the heavily thinned stand compared to the oxygen-18 concentrations in the
513 moderately thinned and control stands (Figure 7.C.). The muted enrichment in ^{18}O around 35 cm
514 depth in the soil indicates a mixing of the left-over summer precipitation with older and lighter
515 water. Our results do not indicate that differences in soil exposure canopy coverage were
516 effective enough to significantly affect the isotopic composition of soil water below 5 cm in
517 depth.

518 4.2. Seasonal lodgepole pine water use

519 Literature utilizing stable water isotopic analysis to determine plant preferential water uptake in
520 arid regions indicates that vegetation can utilize precipitation despite the temporal origin
521 (Andrews et al., 2012; Brinkmann et al., 2019; Ehleringer et al., 1991). Seasonal water
522 availability depends on precipitation, soil water holding capacity and drainage, and evaporative
523 loss (Gibson & Edwards, 2002; Kleine et al., 2020; Stumpp et al., 2018). Based on the seasonal
524 shift in the isotopic composition, soil water at a depth of 5 cm was more enriched with heavier
525 isotopes over the growing season than at 35 cm due to more evaporative isotopic fractionation
526 near the soil surface and a lack of rainfall intense enough to drive precipitation deeper into the
527 soil profile before September 16, 2021 (Figure 3). The effect of evaporative enrichment of the
528 near surface soil water was most obvious in July and September in the heavily thinned stand
529 (T2). However, variability in branch isotopic composition did not follow the same trends. Our
530 results indicate that lodgepole pines access water from multiple depths in the soil profile.
531 Regardless of depth and forest density, spring snowmelt is the main source for lodgepole pines as
532 it infiltrates through the vadose zone.

533 The MixSIAR isotopic partitioning model results from each of the six scenarios indicated a
534 seasonal shift in the depth to water uptake of lodgepole pine, regardless of changes in stem
535 density, over the growing season. At the beginning of the growing season, when snow meltwater
536 is more available at shallow depths and beginning to infiltrate through the soils, lodgepole pines
537 obtain most of their water from snow melt in shallow soils with small contributions from other
538 potential sources (< 25% of June water uptake in all treatments). Then, in July, the trees in the
539 control treatment were using less shallow soil water (34.3% of plant water uptake from 5 cm
540 below the soil profile) whereas the moderately thinned and heavily thinned plots maintained a
541 greater proportion of shallow water uptake (47.1% and 61.5% respectively). The mean $\delta^{18}\text{O}$ and
542 $\delta^2\text{H}$ of branch water from each treatment in September had a higher concentration of lighter
543 stable water isotopes than in July and a larger proportion of tree water was from 35-100 cm deep
544 in the soil profile as shallow soils were dry from a lack of rainfall and surface soil evaporation.
545 By September, the control stand was more dependent on deeper soil water and groundwater with

546 only 33.4% of plant water uptake originating from 35 cm in the soil profile, whereas both
547 thinning treatments maintained more than 45% of water uptake from 35 cm in the soil profile. In
548 October, all treatments were completely dependent on deep soil water, but it is likely that the
549 isotopic profile of deep soil water sampled in July skewed the results. It is plausible that the trees
550 began to rely on shallow soil water towards the end of the growing season when soil water
551 content increased. Further research is needed with more intensive sampling of deep soil water
552 during the hydrological recharge period at the end of the growing season and beginning of
553 senescence.

554 Local monitoring close to the study site indicated that the depth to groundwater stayed at least
555 6.5 m below the surface from August through the end of the study period. The continued use of
556 deep soil water even during rewetting in late September and October suggests that the drought
557 conditions suppressed top soil water uptake, but that deeper soil was sufficiently saturated to
558 sustain root water uptake and tree function enough to limit groundwater uptake to less than 30%
559 for all treatments until the beginning of fall precipitation events recharging the saturated zone.

560 Our results indicate that lodgepole pine, like other pine species in arid regions, is flexible in its
561 ability to access deep soil water and can change its depth to water uptake over time depending on
562 water availability (Brinkmann et al., 2018; Grossiord et al., 2017; Kerhoulas et al., 2013; Kleine
563 et al., 2020; Moreno-Gutiérrez et al., 2011; Simonin et al., 2006; Sohn et al., 2014; Wang et al.,
564 2021). Our results of seasonal changes in depth to water uptake by lodgepole pine support the
565 findings of Andrews et al. (2012) on changes in lodgepole pine depth to water uptake in Alberta.
566 Tree species native to arid regions exhibit a variety of adaptations to long-term drought stress
567 and decreased water availability in the soil profile such as deep tap roots, access to the water
568 table, utilizing bound and mobile soil water, fine root mortality, and hydraulic redistribution in
569 ecosystems with low water holding capacity (Amin et al., 2020; Brinkmann et al., 2018;
570 Grossiord et al., 2017; Kerhoulas et al., 2013; Kleine et al., 2020; Langs et al., 2020; Meinzer et
571 al., 2007b; Prieto et al., 2012; Sohn et al., 2016; J. Wang et al., 2017, p. 201).

572 The literature is inconsistent across different biogeoclimatic regions and species with regards to
573 the effects of thinning on stand dynamics that influence inter-tree competition for water
574 resources or changes in depth to water uptake. (Kerhoulas et al., 2013; Moreno-Gutiérrez et al.,
575 2011; Sohn et al., 2016; Wang et al., 2021). We found no significant impact of forest thinning on
576 depth to water uptake. However, our observation of seasonal shifts in depth to water uptake
577 support results of a study on the impacts of thinning intensity on 60-year-old *Pinus halepensis*
578 Mill. in a semi-arid region of Spain which concluded that forest thinning reduced competition for
579 water resources but did not alter water uptake patterns (Moreno-Gutiérrez et al., 2011). Another
580 study on the impact of thinning *Pinus ponderosa* Dougl. on depth to water uptake concluded that
581 water was consistently more isotopically enriched in low-density stands potentially due to
582 prolonged evaporative fractionation in the soil profile, or that understory vegetation utilized
583 more shallow water sources (Kerhoulas et al., 2013). The impact of forest thinning on stand and
584 understory water use is highly variable and dependent on understory growth, canopy structure,
585 water availability, when forest thinning is implemented, and the time since stem removal
586 (Kerhoulas et al., 2013; Moreno-Gutiérrez et al., 2011; Sohn et al., 2016). More research is
587 needed to discern if lodgepole pine relies more on mobile or bound soil water, the extent of
588 lodgepole pine rooting zones, what biogeochemical factors cause seasonal shifts in water uptake,

589 and if severe seasonal drought has a lasting effect on water uptake strategies during hydrologic
590 recovery (Simonin et al., 2007; Vargas et al., 2017).

591 4.3. Impacts of the drought and implications for future climate conditions

592 The 2021 growing season was an abnormally hot and dry period for the interior of British
593 Columbia with severe to exceptional drought conditions. Wang et al. (2019) found that thinning
594 improved water-use efficiency, drought tolerance, and drought recovery by decreasing stand
595 density and improving carbon storage. Our results support the finding that lodgepole pine trees
596 can adjust to prolonged water scarcity, and over-populated stands may be more resilient than the
597 literature has initially indicated. In fact, drought conditions over the study period likely
598 intensified the change in xylem water isotopic composition over the growing season. However,
599 the scope of this study did not include pre-drought seasonal water use patterns nor the impact of
600 forest density on depth to water uptake during drought recovery. Because lodgepole pine depth
601 to water uptake changes during prolonged dry growing season conditions, the trees are more
602 reliant on winter snowpack and spring infiltration to recharge deeper source water below the
603 evaporative front. One experiment on juniper (*Juniperus monosperma* (Engelm.) Sarg.) and
604 piñon pine (*Pinus edulis* Engelm.) investigated the simultaneous stress of increased heat and
605 decreased precipitation on depth to water uptake and found that extreme temperatures and
606 decreased precipitation lead to less reversible embolism and more root death in surface soil
607 levels preventing trees from accessing shallow water sources if precipitation becomes more
608 available late in the growing season (Grossiord et al., 2017). It is becoming more imperative to
609 understand the climatic drivers of lodgepole pine water use and access as mean annual
610 temperatures continue to rise, the seasonal frequency and intensity of precipitation change, and
611 drought conditions become more severe. This study indicates that severe seasonal dryness pushes
612 lodgepole pines to rely more on snowmelt while losing function in shallow roots. Our results are
613 inconclusive in determining the depth to water uptake in September and October because of
614 limited deep soil water measurements. However, increased annual temperatures and more
615 variable precipitation patterns as a part of climate change projections are predicted to drive
616 decreases in winter snowpack and could drive lodgepole pine stands, regardless of stem density,
617 to rely on groundwater influencing water availability and depth to groundwater. These
618 projections could lead to prolonged inter-annual water scarcity along with seasonal water
619 scarcity during the late growing season.

620

621 5. Conclusions

622 Lodgepole pine, across all treatments, was able to shift access from shallow soil water at the
623 beginning of the growing season to deeper soil water as drought conditions intensified. The
624 quick-draining and sun-exposed soils of the UPC do not retain small summer precipitation
625 events, and these patterns are intensified in the shallow soil layer of the heavily thinned stand
626 because decreased canopy cover can be directly related to increased soil evaporation. As a result,
627 due to changes in water availability, lodgepole pines shift to a more readily available source in
628 the soil profile (Aranda et al., 2012; Prieto et al., 2012). Our findings support the literature that
629 lodgepole pines are a drought-tolerant species with dimorphic rooting systems allowing them to
630 access water from varying depths in the soil depending on water availability (Andrews et al.,
631 2012; Liu et al., 2011). Despite the ecological plasticity under extreme heat and low summer

632 precipitation conditions, there was no statistically significant variance in depth to water use
633 between the over-populated plots and thinned ones. Both thinned and unthinned lodgepole pine
634 stands were able to access shallow soil water during the early months (June and July), then
635 switched to deeper soil water and a larger proportion of groundwater during September.
636 Although there was not a statistically significant difference in isotopic composition of branch
637 water for the different treatments, our results indicate that decreased stem density may lead to the
638 prolonged use of soil water 35 cm below the surface during prolonged dry periods which would
639 decrease the dependency of lodgepole pine on shallow soil water and summer precipitation
640 events and rather increase the dependency on deep soil water or ground water fed by winter snow
641 accumulation and spring snowmelt.

642 Future climate projections indicate hotter growing seasons and less precipitation (Allen et al.,
643 2010). Further investigation is needed to discern how lodgepole pines, under different stand
644 densities, use water during prolonged drought and drought recovery periods (Grossiord et al.,
645 2017; Navarro-Cerrillo et al., 2019; Simonin et al., 2007; Sohn et al., 2016). From our findings,
646 stand density did not prevent lodgepole pines from accessing soil water from various depths, but
647 decreased stem density did result in lodgepole pines using soil water higher in the soil profile for
648 longer under extremely dry conditions. Lodgepole pines indicate a strong level of drought
649 tolerance and ability to access water under extreme heat conditions. If summer precipitation
650 decreases, lodgepole pine in the interior of British Columbia can access deeper soil water from
651 spring snowmelt. However, if snowpack and spring snowmelt begin to decrease, lodgepole pine
652 may need to acclimate to these hydrological shifts.

653

654 *Code and Data Availability:*

655 The codes of the data analysis and plotting are available at <https://github.com/emory->
656 [ce/LodgepolePineWaterUseStrategies2021](https://github.com/emory-ce/LodgepolePineWaterUseStrategies2021) and are available upon request (ece58@nau.edu)

657

658 *Author Contributions:*

659 EE conceived the idea as a part of their Master's research with AW, and performed the
660 extractions with RG. Analysis was primarily conducted by EE with guidance from AW and RG.
661 All authors contributed to the manuscript.

662

663 *Competing Interests:*

664 None of the authors have competing interests.

665

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671

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