

## Publication

### Explicitly accounting for needle sugar pool size crucial for predicting intraseasonal dynamics of needle carbohydrates $\delta^{18}\text{O}$ and $\delta^{13}\text{C}$

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We explore needle sugar isotopic compositions ( $\delta^{18}\text{O}$  and  $\delta^{13}\text{C}$ ) in boreal Scots pine (*Pinus sylvestris*) over two growing seasons. A leaf-level dynamic model driven by environmental conditions and based on current understanding of isotope fractionation processes was built to predict  $\delta^{18}\text{O}$  and  $\delta^{13}\text{C}$  of two hierarchical needle carbohydrate pools, accounting for the needle sugar pool size and the presence of an invariant pinitol pool. Model results agreed well with observed needle water  $\delta^{18}\text{O}$ ,  $\delta^{18}\text{O}$  and  $\delta^{13}\text{C}$  of needle water-soluble carbohydrates (sugars + pinitol), and needle sugar  $\delta^{13}\text{C}$  ( $R^2 = 0.95, 0.84, 0.60, 0.73$ , respectively). Relative humidity (RH) and intercellular to ambient  $\text{CO}_2$  concentration ratio ( $C_i/C_a$ ) were the dominant drivers of  $\delta^{18}\text{O}$  and  $\delta^{13}\text{C}$  variability, respectively. However, the variability of needle sugar  $\delta^{18}\text{O}$  and  $\delta^{13}\text{C}$  was reduced on diel and intra-seasonal timescales, compared to predictions based on instantaneous RH and  $C_i/C_a$ , due to the large needle sugar pool, which caused the signal formation period to vary seasonally from 2 d to more than 5 d. Furthermore, accounting for a temperature-sensitive biochemical  $^{18}\text{O}$ -fractionation factor and mesophyll resistance in  $^{13}\text{C}$ -discrimination were critical. Interpreting leaf-level isotopic signals requires understanding on time integration caused by mixing in the needle sugar pool.

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