

# Comparing tree species-specific water uptake strategies via continuous in-situ monitoring of stable water isotopes



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# Introduction

## Research Questions:

- Where do trees get their water from and how long does it take from water uptake to transpiration?
- Do different tree species have different water use strategies to access soil water resources?

# Approach:

- To understand how soil water is used by different tree species, we setup a semi-controlled outdoor pot experiment with three different tree species.
- We took advantage of stable water isotope techniques by tracing

# plant water uptake through isotopically labelled irrigation water.

We followed the isotopic composition of soil and xylem water in high temporal resolution via in-situ isotope probes, which were directly attached to an isotope analyzer.

# **Expected Implications:**

These questions are

fundamental in order to estimate how trees will be affected by a change in available soil water resources under predicted climate change.

# Experimental setup

- Three 20 year-old trees (4-6 m high): pinus, alnus, quercus
- Planted into clayey loamy substrate (Fig. 1)

## **Equipment:**

- In-situ porous-membrane isotope probes in the soils and tree xylem
- Soil moisture and temperature probes
- Soil matrix potential probes
- Sapflow sensors
- Portable photosynthesis system (LI-COR)
- Climate station

# Comparison with destructive samples:

Soils (2 depths) and tree xylem (two heights) were sampled destructively four times for the vapour equilibration technique (Fig. 3)

# Isotope labeling:

Quercus

 Deuterated water was applied as irrigation water (three campaigns) (Fig. 2, dashed line)

# Alnus incana Quercus suber Rainout-shelter rainout-shelter rainout-shelter valve box valve box in-situ isotope probes soil moisture sensor Picarro isotope analyzer (water + soil)

Fig. 1: Overview of experimental setup and measurement equipment.

# Conclusions

- In-situ isotope measurements in tree xylem and soil compared reasonably well to isotope values gained by destructive sampling but yield much higher temporal resolution.
- → In-situ isotope measurements are a powerful tool to trace ecohydrological fluxes.
- Very heterogeneous behavior in tree specifies water uptake patterns under controlled conditions.
- Species showed distinguishably different sapflow dynamics, but quick responses to the isotope label uptake

# <u>Implications & Outlook:</u>

- Integrating information on species-specific event-water use and root-water uptake dynamics will deliver first hints on how trees may funnel water towards their active root zones.
- This will become important under future climatic conditions in terms of development of adaptation strategies for sustainable forest management.
- Additional labeling experiments in 2020.

# Results & Discussion

# Tree-specific label water uptake (Tab.1): Alnus:

Distinct response in xylem and soil (30cm)
 δ<sup>2</sup>H values after 1 day for both labelings. Soil isotopic signature more responsive in 30cm than in 15cm soil depth.

#### <u>Pinus:</u>

Soil 15cm

Soil 30cm

- Stem 15cm

Stem 150cm

Stem 270cm

Destructive Measurements

Labeling

Sap Flow

10cm

- 150cm

Soil Moisture

15cm root bale

- 30cm root bale

15cm south

15cm north

30cm south

Water Potential

15cm root bale

15cm south

15cm north

- 30cm north

- Distinct response in xylem δ<sup>2</sup>H values
   (15cm) after 1 day; stronger effect for label 1
- Despite highest sapflow rates, isotopic label difference was lower than for other species
- Similar isotopic dynamic but higher δ<sup>2</sup>H values in 15cm than in 30cm soil depth.

### Quercus:

More moderate response of increasing xylem  $\delta^2$ H signatures (low sapflow rates), but quick response in the soil.

## In-situ vs. destructive sampling (Fig. 3):

 Destructive samples show a wider isotopic spread but were mostly consistent with insitu measurements.

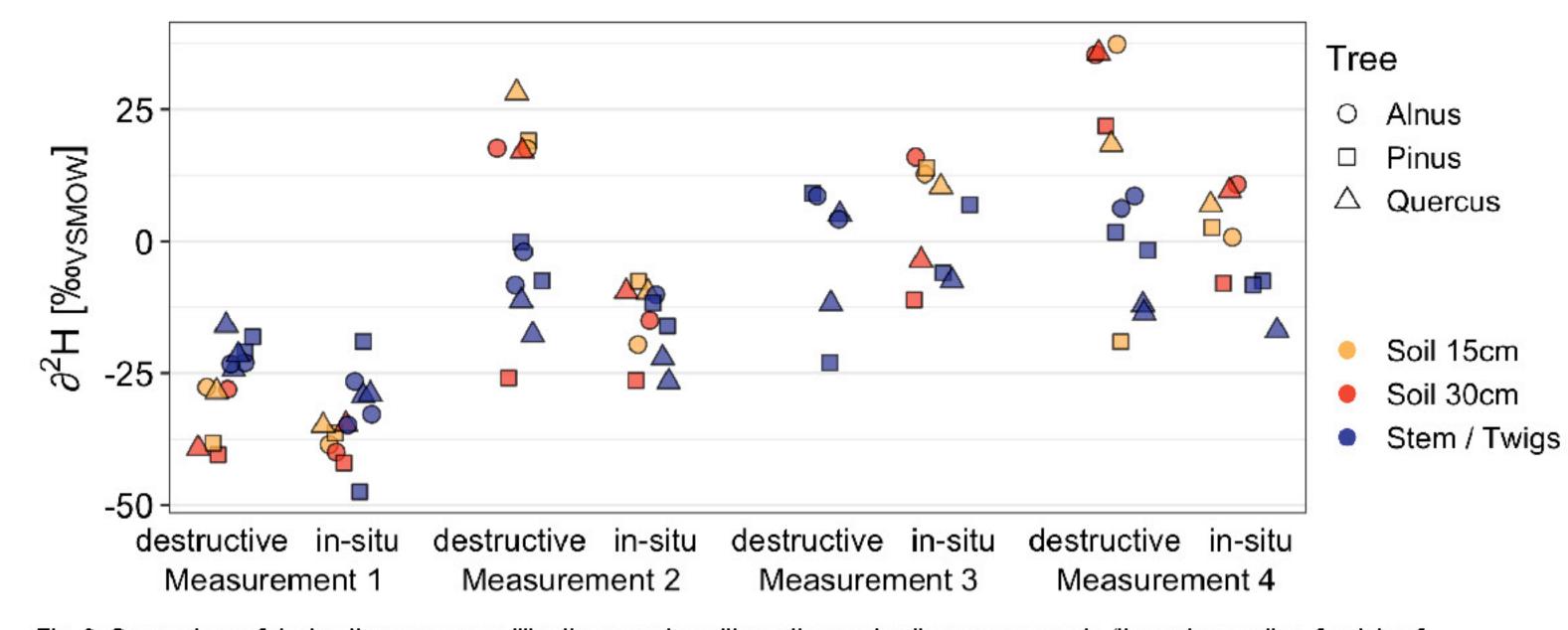


Fig. 3: Comparison of destructive vapour equilibration samples with continuous in-situ measurements (three-day median, for date of measurements see Fig. 2, grey bars).

Tab. 1: In-situ measurement response after label 1 (a&b) and 2. \* significant differences (p<0.05) between  $\delta^2$ H values three days before and after labeling (ANOVA), -- no visible change in  $\delta^2$ H values within one week after labeling, ++ visible change, but no delay time, due to five days measurement failure.

		Alnus				Pinus				Quercus			
		Soil		Stem		Soil		Stem		Soil		Stem	
Label 1		15cm	30cm	<b>1</b> 5cm	150cm	15cm	30cm	15cm	150cm	15cm	30cm	15cm	150cm
	delay [days]	6	< 1	< 1	> 6	< 1		< 1	ca. 6	< 1	< 1		
	difference [ð²H ‰]	+ 20	+ 40 *	+ 40 *	+ 20	+ 20 *		+ 20 *	+ 20	+ 20 *	+ 20 *		
7													
Label	delay [days]	++	< 1	< 1		< 1	++	++	++	< 1	< 1		
	difference [ð²H ‰]	+ 30	+ 35	+ 35		+ 20	+ 20	+ 20	+ 20	+ 30 *	+ 30 *		

# Fig. 2: Variation of $\delta^2$ H, sapflow, soil moisture, water potential and climatic parameters (solar radiation (TSI), vapour pressure deficit (VPD) and air temperature ( $T_{air}$ )) over the course of the experiment.