

# Using fibre-optic sensing for non-invasive, continuous dendrometry of mature tree trunks

Martijn van den Ende, Eléonore Oberlé, Thierry Améglio, Robin Ardito,

Gildas Gâteblé

# ▶ To cite this version:

Martijn van den Ende, Eléonore Oberlé, Thierry Améglio, Robin Ardito, Gildas Gâteblé. Using fibreoptic sensing for non-invasive, continuous dendrometry of mature tree trunks. 2024. hal-04574205

# HAL Id: hal-04574205 https://hal.science/hal-04574205

Preprint submitted on 14 May 2024

**HAL** is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers. L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.

# Using fibre-optic sensing for non-invasive, continuous dendrometry of mature tree trunks

Martijn van den Ende<sup>1,\*</sup>, Eléonore Oberlé<sup>2</sup>, Thierry Améglio<sup>3</sup>, Robin Ardito<sup>2</sup>, Gildas Gâteblé<sup>2</sup>

1 Université Côte d'Azur, CNRS, IRD, Observatoire de la Côte d'Azur, Géoazur, France
2 INRAE, UE 1353, Unité Expérimentale Villa Thuret, Antibes Juan-les-Pins, France
3 Université Clermont Auvergne, INRAE, PIAF, F-63000 Clermont-Ferrand, France

\* Corresponding author: martijn.vandenende@oca.eu

# Abstract

Dendrometry is the main non-invasive macroscopic technique commonly used in plant physiology and ecophysysiology studies. Over the years several types of dendrometric techniques have been developed, each with their respective strengths and drawbacks. Automatic and continuous monitoring solutions are being developed, but are still limited, particularly for non-invasive monitoring of large-diameter trunks. In this study, we propose a new type of automated dendrometer based on distributed fibre-optic sensing that continuously measures the change in stem circumference, is non-invasive, and has no upper limit on the trunk diameter on which it can be installed. We perform a three-month validation experiment during which we deploy a fibre-optic cable at three localities around the trunks of two specimens of *Brachychiton*. We verify the accuracy of this new method through comparison against a conventional point-dendrometer, and we observe a consistent time lag between the various measurement locations that varies with the meteorological conditions. Finally, we discuss the feasibility of the fibre-based dendrometer in the context of existing dendrometric techniques and practical experimental considerations.

### 1 Introduction

Dendrometry plays a prominent role in plant physiology. Current techniques that measure stem diameter comprise circumferentially-averaged and local point measurements [Drew and Downes, 2009]. Out of these methods, (non-automated) dendrometer bands, tape and calipers [Liu et al., 2011, McMahon and Parker, 2015], require manual intervention for each measurement, resulting in a less than desired temporal resolution and non-uniform time sampling which preclude the use of conventional signal processing methods (such as frequency-band filtering). Moreover, since climatic variables are non-stationary and the tree response to these can be highly nonlinear [Wilmking et al., 2020], ideal measurement systems should be long-term stable and continuous in time, in order to track the response of a specimen over time. However, automated and continuously-recording sensors require signal acquisition, conditioning, and storage or telemetry systems that are adapted to the (harsh) outdoor conditions of *in-situ* studies, a feat that is not trivial to accomplish. 12

Variation in branch diameter can be monitored continuously, non-invasively and in real time using 13 continuous dendrometers [Ameglio and Cruiziat, 1992, Zweifel et al., 2000, 2006]. On a daily basis, 14 the variation in branch diameter is the result of several components, including irreversible radial 15 growth, reversible dehydration/rehydration of living cells, thermal expansion and contraction, and 16 the expansion of conducting elements due to the increase and relaxation of internal tension Daudet 17 et al., 2005, Steppe and Lemeur, 2004, Zweifel et al., 2006]. In the event of severe drought, radial 18 growth is halted and the variation in branch diameter mainly represents the fluctuation in water 19 storage in elastic tissues, including most living cells. [Andriantelomanana et al., 2024, Lamacque 20 et al., 2020, Zweifel, 2016, Zweifel et al., 2016]. Dendrometers have also given us more insight into 21 trees' resistance to freezing and the consequences of the freezing and thawing processes [Ameglio 22 et al., 2001, Charra-Vaskou et al., 2016, Charrier et al., 2017, Zweifel and Häsler, 2000]. Dendrometry 23 can be complemented with other types of instrumentation, such as stomatal conductance analysis 24 [Hernandez-Santana et al., 2023, Marchin et al., 2022], sap flow measurements [Iqbal et al., 2021, 25 Steppe and Lemeur, 2004], and water potential measurements [Cochard et al., 2001, Moriana et al., 26 2012], to provide a more holistic view of the response of the specimen. 27

However, as these techniques represent local point-measurements, they may yield biased mea-28 surements if the stem diameter variation is non-isotropic along the circumference of the stem. This 29 may arise when fluid transport is not homogeneous due to predominant positioning of branches, or 30 when wind triggers the formation of flexure wood and ovalisation of the stem [Roignant et al., 2018]. 31 Another case is given by the non-uniform mechanical response of the bark, e.g. in species with a 32 thick, scaley rhytidome (dead layers of old periderm), commonly seen in pines, or with the outer bark 33 shedding off in sheets (Platanus L. and Eucalyptus L'Hér.) [Coder, 2022]. In these and many other cases, it is desirable to measure the stem diameter variation averaged over the entire circumference, 35 which may be a more representative measure for comparison between specimens within or between 36 species. 37

Moreover, it is often impractical (or even impossible) to install instrumentation along large-38 diameter stems. Around the world, there exist many veteran tree specimens that have a particular 39 historical, cultural, or botanical interest. Local, national, and international initiatives intended to preserve this biological heritage (such as the *arbres remarguables* in France, Great trees or Champion 41 Trees in the US and UK [Roesch et al., 2024], and heritage trees in China [Jim, 2017]) have a need for 42 affordable and reliable monitoring of the health of the specimen, as well as its response to climatic 43 changes, resulting in e.g. extended hydraulic stress. The use of invasive and destructive techniques 44 on these specimens is avoided, particularly for specimens that are fragile. To some extent, automated 45 band dendrometers alleviate many of the aforementioned issues, though their accuracy has been found to be rather limited due to friction, hysteresis, and the difficulty of eliminating slack during 47 installation [Herrmann et al., 2016, Wang and Sammis, 2008]. Hence, there is a clear niche for a new 48 type of dendrometric instrumentation that can be applied in the scenarios described above. 49

Relatively recently, distributed fibre-optic sensing [Hartog, 2017] has emerged as a promising 50 technology for measuring strain (and temperature) in a distributed manner. By systematically 51

sending short pulses of laser light into an optical fibre (as commonly used in telecommunication), local strain can be inferred as a function of distance along the fibre through interferometric or 53 spectral analysis techniques. Various sensing techniques have already been adopted in a range of 54 scientific disciplines, including engineering [Hubbard et al., 2021, Liu et al., 2023], geophysics [Jousset 55 et al., 2018, Lindsey and Martin, 2021, van den Ende and Ampuero, 2021, and ocean monitoring 56 [Bouffaut et al., 2022, Rivet et al., 2021], and to a limited extent, in ecology [Ashry et al., 2020]. 57 Fibre-optic cables are flexible, robust to environmental factors like temperature and UV radiation, 58 and are relatively inexpensive. These characteristics unlock a wide range of applications and targets 59 for deployment. 60

Recognising the potential of fibre-optic sensing applied to dendrometry, we propose a novel 61 experimental set-up to continuously record the stem diameter variations of tree specimens. More 62 specifically, we conducted a 3-months long experiment measuring the radial expansion of two specimens 63 of the genus *Brachychiton* with Brillouin-based Distributed Strain and Temperature Sensing (DSTS). 64 In what follows, we describe in detail the DSTS methodology, the experimental configuration, and 65 the data processing workflow. We interpret our observations of trunk strain variations in the context 66 of the meteorological conditions encountered during the experiment, and the known differences in 67 responses to hydraulic stress between the two Brachychiton species [Reynolds et al., 2018]. We 68 conclude with an assessment of the feasibility of Brillouin-based DSTS for distributed dendrometry 69 and with perspectives for the future. 70

# 2 Methodology

#### 2.1 Description of study site and species

The data were collected from 28 July to 02 November 2023 at the Villa Thuret Botanical Gardens 73 in Antibes, South-East France (43°33'47.8"N, 7°7'30.6"E, 40 m a.s.l). The garden was established 74 in 1857 by Gustave Thuret, and is now managed by INRAE (France's National Research Institute 75 for Agriculture, Food and Environment), a public research institute. It spans an area of 3.5 ha and 76 contains a collection of 2500 plant specimens, representing 1000 species and 131 families. Pruning is 77 only done for safety reasons and acclimation to the local climate is tested by restricting artificial 78 watering to the first three years after planting [Ducatillion and Landy, 2010]. Hence, the specimens 79 present on this site are adapted to the local, coastal Mediterranean climate [Csa Köppen subtype; 80 Kottek et al., 2006]. This climate is mainly defined by drought in summer, mild winters, and 81 precipitations concentrated in autumn and spring [Daget, 1977]. The typical rainfall in Antibes 82 averages 800 mm per year, although it only reached 530 mm in 2022. The average annual temperature 83 is 15 °C, with a minimum and maximum around 0 °C and 34 °C, respectively. Locally, the clay-rich 84 soil layer extends only to shallow depths (60 cm to 80 cm), and owes its elevated pH (> 7) to an 85 important quantity of calcium [Gras, 1990]. 86

The specimens studied in this work are from the genus *Brachychiton* Schott & Endl. (Malvaceae family), which are known to tolerate drought and to grow in varied climates and soil types [Elliot and Jones, 1985]. The first, *Brachychiton discolor* F.Muell. (Lacebark tree), has a typical height of 10 m to 30 m, is deciduous before flowering, and grows in rain forests and scrubs of coastal eastern

71

Australia (Qld, NSW). The second, *Brachychiton populneus* (Schott & Endl.) R.Br. (Kurrajong tree), has a typical height of 6 m to 20 m, is evergreen but can be drought deciduous, and is found in rocky areas, riverbanks and semi-arid plains of inland SE Australia (Qld, NSW, Vic) [Elliot and Jones, 1985]. *Brachychiton populneus* is considered to be slightly more drought tolerant than *B. discolor* [Reynolds et al., 2018]. Both specimens tested at the Villa Thuret are mature, and their heights are 9.5 m, for *B. discolor* and 10 m for *B. populneus*. Their DBH are 136 cm and 115 cm respectively.

#### 2.2 Distributed Strain and Temperature Sensing

As a subcategory of fibre-optic sensing, Distributed Strain and Temperature Sensing (DSTS, also known as Brillouin Optical Time-Domain Reflectometry, B-OTDR) is ideally suited to long-term 99 stable monitoring of both fibre strain and ambient temperature. As described in detail by e.g. Hartog 100 [2017], the measurement principle of DSTS is based on spontaneous Brillouin scattering, which arises 101 from the interaction between an incident photonic wave (the probe wave), and thermally-excited 102 acoustic vibrations in the fibre lattice called phonons. Those phonons whose wavelength matches 103 that of the probe wave interfere with the photons to produce back-scattering that can be detected by 104 the instrument at one end of the fibre. The Brillouin back-scattering frequency spectrum exhibits a 105 peak with intensity I at frequency  $\nu$ , which relate to fibre strain  $\varepsilon$  and temperature T as: 106

$$\begin{bmatrix} \Delta \nu \\ \Delta I \end{bmatrix} = \begin{bmatrix} C_{\nu}^{\varepsilon} & C_{\nu}^{T} \\ C_{I}^{\varepsilon} & C_{I}^{T} \end{bmatrix} \begin{bmatrix} \Delta \varepsilon \\ \Delta T \end{bmatrix}$$
(1)

By measuring the peak frequency and intensity relative to an arbitrarily defined baseline,  $\Delta \nu$  and  $\Delta I$  can be converted into strain and temperature relative to that same baseline ( $\Delta \varepsilon$  and  $\Delta T$ , respectively). Typical values for the coefficient matrix have been established by Maughan et al. [2001] as:

$$\begin{bmatrix} C_{\nu}^{\varepsilon} & C_{\nu}^{T} \\ C_{I}^{\varepsilon} & C_{I}^{T} \end{bmatrix} = \begin{bmatrix} 0.046 \,\mathrm{MHz}\,\mathrm{m}\,\mathrm{\mu}\mathrm{m}^{-1} & 1.07 \,\mathrm{MHz}\,\mathrm{K}^{-1} \\ -8 \times 10^{-6} \,\%\,\mathrm{m}\,\mathrm{\mu}\mathrm{m}^{-1} & 0.36 \,\%\,\mathrm{K}^{-1} \end{bmatrix}$$

For a given relative spectral measurement  $(\Delta\nu, \Delta T)$ , the physical quantities  $(\Delta\varepsilon, \Delta T)$  are obtained by inverting this (constant) coefficient matrix and left-multiplying. As we have not calibrated these coefficients to account for the specific thermo-mechanical properties of our fibre-optic cable assembly, we directly utilise  $\Delta\varepsilon$  estimated during acquisition, and correct for any temperature dependencies through the use of reference cable segments (this will be described in detail in the next section).

A single DSTS measurement is distributed over the fibre, meaning that during a single acquisition 112 cycle, a Brillouin spectrum is obtained at discrete points that are evenly distributed in space (the 113 spacing of which is referred to as the gauge length L). Optionally, the gauge length can be oversampled 114 to obtain more sensing points along the fibre, although this practice does not directly lead to a 115 higher spatial resolution as two sensing points within one gauge length are statistically dependent. 116 Hence, DSTS provides measurements of  $\Delta \varepsilon$  that are averaged over a gauge length. By repeating 117 the acquisition cycle every  $\Delta t$  time units, a data set of (relative) strain is obtained with a spatial 118 resolution L and a temporal resolution  $\Delta t$ . 119



**Figure 1.** Overview of the fibre deployment. Fibre sections A (tight and loose) and B (only tight) are looped around a *Brachychiton populneus*. Fibre section C (tight and loose) is looped around a *Brachychiton discolor*, which is equipped with a PépiPIAF dendrometer installed on a branch (not in view of this picture).

#### 2.3 Experimental set-up

We performed a small-scale experiment by deploying 100 m of tight-buffered single-mode fibre (type: 121 G652.D) around the trunk of two trees; first, several winds of fibre were tightly looped around the 122 trunk of a Brachychiton populneus (location B in Fig. 1, at a height of 85cm), followed by several more 123 tight winds one metre higher up the same trunk (location A in Fig. 1, at a height of 185cm). Since 124 DSTS is simultaneously sensitive to strain and temperature, additional fibre was wound very loosely 125 around location A. Owing to the negligible coupling between this loose fibre and the trunk, this 126 section of fibre is sensitive only to temperature, and can hence be used to correct the measurements 127 of the tightly-wound sections. While Eq. (1) should (in theory) be able to separate the effects of 128 strain and temperature, we find that this separation is imperfect and that in practice the use of 129 a reference section is preferred. Finally, the fibre was looped (both tightly and loosely) around a 130 second tree (Brachychiton discolor; location C in Fig. 1, at a height of 160cm), which was equipped 131 with an independent LVDT-based micro-dendrometer (described in more detail below). 132

With the fibre in place around the trees of study, the fibre was rolled back to starting point and connected to a Febus G1-C-19002 DSTS interrogator [Clément et al., 2021]. The section of fibre

spanning between the trees and the interrogator was placed within a rubber tube for protection. The acquisition was launched with a time sampling period of 300 s and spatial resolution (gauge length) of 2 m, oversampled to obtain a spacing of 0.5 m. To geolocate specific sections of interest (rubber tube entry and exit points, tight and loose sections), tap tests were performed by manually straining the fibre at specified times, which could later be easily identified in the processed data.

As a reference instrument, we employ the PépiPIAF dendrometry system (Capt-Connect, Clermont-140 Fd, France), which features a linear variable differential transformer (LVDT) that continuously 141 monitors micro-metric variations of the diameter of branches or trunks. This instrument has previously 142 been used in various studies [e.g., Charrier et al., 2017, Daudet et al., 2005, Lamacque et al., 2020]. 143 The specific model used in the present study is the e-Pépipiaf 1.0, which is equipped with a Solartron 144 DF5.0 LVDT sensor, with a resolution of 0.5 µm, as well as a thermistor housed inside a waterproof 145 case to simultaneously record ambient temperature [Chevallier et al., 2011, Lamacque et al., 2022]. 146 The instrument is mounted on (B. discolor on a north-east facing branch with a diameter of 20 cm 147 at a height of 3.25 m above the ground. 148

After data retrieval from the instruments, the fibre segments corresponding with locations A, B, and C were extracted. The channel recordings within a given section were averaged to reduce uncorrelated noise, and the recordings of the loosely-coupled sections (which contain the apparent strain signal due to temperature changes) were subtracted from those of the tightly-coupled sections. The resulting time-series for locations A, B, and C were thus unaffected by daily temperature fluctuations. For reference, meteorological data were obtained from the Météo-France public database for a weather station at the airport of Nice (located 10 km from the study site).

#### 3 Results

Since DSTS is not yet a proven technology for dendrometry, we first consider the amplitude and 157 long-term evolution of the radial strain of B. discolor. This specimen is equipped with a PépiPIAF 158 dendrometer that has been successfully used in previous studies, and which will act as a reference 159 sensor. Even though section C of the optical fibre was not deployed directly adjacent to the PépiPIAF, 160 it is located sufficiently close to warrant a direct comparison. To facilitate this, the output of the 161 PépiPIAF LVDT sensor was converted from units of millimetres to units of microstrain through 162 division by the diameter of the branch it was mounted on (20 cm). The (equivalent) strain relative to 163 the start of the experiment is given for the PépiPIAF dendrometer and section C of the optical fibre 164 in Fig. 2b. While small deviations between the two time series can be occasionally discerned, overall 165 the agreement for both the diurnal variations and the long-term evolution is more than satisfactory. 166 Moreover, since the two sensors are not directly collocated, it is not expected to obtain perfectly 167 identical results, and so we attribute minor amplitude discrepancies to the different response of the 168 trunk and the branch at which the measurements are made. Particularly for the period after 21 169 October, corresponding to a period of net secondary growth, the trunk appears to be accumulating 170 more strain than the branch. 171

During the experiments, several brief episodes with a significant amount of precipitation occurred (indicated by the blue bars in Figs. 2-4). During the first episode starting on 2023-08-27 a modest amount of precipitation was measured (up to 40 mm over 48 hours), and a minor response of both 174

bioRxiv preprint doi: https://doi.org/10.1101/2024.05.10.593544; this version posted May 13, 2024. The copyright holder for this preprint (which was not certified by peer review) is the author/funder, who has granted bioRxiv a license to display the preprint in perpetuity. It is made available under aCC-BY 4.0 International license.



Figure 2. (a) Temperature recorded by the PépiPIAF on-board thermocouple and daily precipitation recorded by a weather station located  $\sim 10 \text{ km}$  away. (b) Strain of the fibre located at section C, and that recorded by the PépiPIAF dendrometer on a branch on the same tree (*B. discolor*). (c) Strain of the fibre located at sections A and B (*B. populneus*). All strain measurements are relative to the start of the time series, and the dates are given in local time (UTC+2).

Brachychiton specimens was recorded. The subsequent precipitation episodes were more significant, triggering a more considerable response in both specimens, dilating by as much as  $1000 \,\mu m \,m^{-1}$  (equivalent to a radial dilatancy of the trunk of approximately  $0.25 \,mm$ ).

Considering the radial strain variations in more detail, we find subtle longitudinal variations 178 in each specimen. First, when we overlay the strain recordings of the pépiPIAF dendrometer and 179 fibre section C, we observe a small but persistent time lag of the PépiPIAF with respect to the 180 fibre (Fig. 3a). This time lag can be effectively quantified by converting two narrowband-filtered 181 time series into their analytic representation and subsequently extracting the phase as the complex 182 argument. Through this analysis, we obtain the instantaneous time lag  $\Delta \tau(t)$  between the pépiPIAF 183 and the fibre recordings, as well as the instantaneous amplitude A(t) of the strain signal, which we 184 can track over time. We provide a detailed description of the analytic signal approach in Appendix I. 185

Applying this processing procedure to the PépiPIAF and fibre recordings, we estimate the time <sup>1366</sup> delay to fall in the range of 1-3 hours, typically (Fig. 3c). The corresponding instantaneous amplitude <sup>1377</sup> of the analytic signal ranges from < 100 to close to  $250 \,\mu m \,m^{-1}$ . Note that owing to the narrowband <sup>1389</sup> filtering applied (see Appendix I), this amplitude only reflects the strain associated with the diurnal <sup>1390</sup> hydration cycle. When compared to the recordings of daily precipitation, a persistent increase of the <sup>1391</sup> time lag seems to coincide with a period of drought. More strikingly, the amplitude of the diurnal <sup>1392</sup> hydration cycles correlates directly with the onset of precipitation. <sup>1392</sup>



Figure 3. Comparison between the PépiPIAF dendrometer and fibre section C (*B. discolor*). (a) Detailed view of the bandpass-filtered strain data, clearly showing the time delay between section C and the pépiPIAF. (b) Complete time series of the bandpass-filtered strain data. For reference, the time range of panel (a) is indicated by the shaded area, and the daily precipitation is given by the blue bars. The time series have the same colour coding as in panel (a). (c) The instantaneous time delay and amplitude of the strain.

Performing a similar analysis on fibre sections A and B (*B. populneus*), we observe a time lag that is systematically much smaller than that seen for *B. discolor*, usually remaining below 1 hour (Fig. 4). This is not necessarily unexpected, as fibre sections A and B are both located on the trunk, between fibre section C and the PépiPIAF dendrometer on the other specimen. Moreover, the same trends are observed as for *B. discolor*: the time lag tends to increase during periods of drought, and the instantaneous amplitude increases dramatically with each precipitation episode.

#### 4 Discussion

#### 4.1 Interpretation of hydraulic response

It is clear from our analysis that there is a direct correlation between the amplitude of the diurnal hydration cycles and the availability of water. This has been observed in previous studies [Daudet 203

200



**Figure 4.** Comparison between the fibre sections A and B (*B. populneus*). (a) Detailed view of the bandpass-filtered strain data, showing a minute time delay between sections A and B. (b) Complete time series of the bandpass-filtered strain data. For reference, the time range of panel (a) is indicated by the shaded area, and the daily precipitation is given by the blue bars. (c) The instantaneous time delay and amplitude of the strain.

et al., 2005, Fernández and Cuevas, 2010, Steppe and Lemeur, 2004], and it is well understood in terms 204 of an increased sap flow throughout the day [Zhao and Liu, 2010]. What can be seen particularly 205 well in Fig. 2c, is that a larger daily amplitude corresponds to an increase of the maximum strain 206 seen throughout a day, while the baseline diameter remains relatively constant over time, indicating 207 a lack of secondary growth. This indicates that the amplitude variations are due to an increased use 208 of water reserves in bark cells compared to a relatively constant storativity baseline. This constant 209 baseline may result from drought induced stomatal closure (corresponding with minimal diurnal 210 strain variations), in order to stabilise the water potential and avoid cavitation of the sap [Choat 211 et al., 2018]. A maximum of fluid storage, which marks the end of the periods of rehydration and 212 growth, is typically achieved around 08:00 local time. 213

In contrast to the amplitudes of diurnal dynamics, the time lag between discrete locations along the trunk is less straightforward to interpret. For both specimens we observe a systematic positive time lag, meaning that the sensor higher up on the tree (the PépiPIAF sensor for *B. discolor*, or fibre section A for *B. populneus*) exhibits a delay in the signal. Previous studies that simultaneously measured the rate of sap flow at the base of the trunk and in the crown of a specimen report that sap flow initiates near the crown, triggered by the onset of transpiration [Phillips et al., 1999, Zweifel 219 et al., 2001]. The observed time lags of the initiation of sap flow range from hours [Čermák et al., 220 2007] to minutes [Kume et al., 2008]. Hence, when considering the rate of sap flow, one would expect 221 to make the opposite observation (i.e., a negative time lag) from what is measured in the present 222 study. However, one needs to keep in mind that with both the PépiPIAF dendrometer and with 223 the fibre, one records radial strain, which is not necessarily directly linked with the rate of sap flow; 224 instead, it is the net storage of fluid in the elastic sapwood and phloem tissue that contributes directly 225 to stem diameter variations [Cermák et al., 2007]. An accumulation of sugar higher up the tree could 226 possibly induce an osmotic pressure gradient that stimulates fluid flow towards these regions, causing 227 storage at higher positions and depletion at lower positions on the tree. Nighttime transport of 228 solutes towards the base of the tree, along with rehydration, would reverse this trend, hence resulting 229 in a daily stem variation that occurs earlier in the day for lower positions. Compared with other 230 dendrometric studies (as opposed to sap flow studies), positive or zero time lags in stem diameter or 231 between sap flow and transpiration, have been occasionally reported [Ameglio and Cruiziat, 1992, 232 Simonneau et al., 1993, Steppe and Lemeur, 2004, Ueda et al., 1996]. Exactly what controls the 233 retention and release of fluids is subject of debate [Sevanto et al., 2002], and is likely highly dependent 234 on the species under investigation [Scholz et al., 2008]. We therefore cannot be conclusive about the 235 potential role of solutes in explaining the observed lag times. 236

A second candidate hypothesis is the difference in exposure to solar radiation: east facing branches 237 (which are illuminated earlier in the day) tend to have an earlier peak in sap flow, compared to west 238 facing and lower branches [Burgess and Dawson, 2008, Martin et al., 2001], and southern branches 239 can have a higher total/overall sap flow than northern branches [Steinberg et al., 1990]. In this study, 240 the PépiPIAF was mounted on a lower branch oriented to the North-East, usually in the shade of a 241 building. Since the fibre at section C measures the stem variations averaged over the circumference 242 of the trunk, one might anticipate time shifts when comparing a specific branch to the tree as a 243 whole. However, this does not explain the time delay observed between sections A and B, which are 244 both situated on the same trunk. 245

Regardless of the relation between sap flow rates and fluid storage, it is interesting to note that 246 as soon as the water availability increases (during or shortly after precipitation), the observed time 247 lag diminishes; particularly in the case of *B. populneus* this time lag approaches zero, implying 248 that the trunk essentially acts as a rigid fluid reservoir that uniformly stores and releases fluids 249 from and to other parts of the specimen. In between precipitation episodes, the time lag becomes 250 non-zero, resulting in a distributed storage and release of fluids starting at the base of the trunk and 251 propagating upward. By having several continuous measurements distributed over the length of the 252 stem, it has become feasible to study this transition from uniform to heterogeneous fluid release in 253 detail, particularly in relation to the availability of water and the state of hydraulic stress. In turn, 254 such observations can be interpreted in terms of species-dependent strategies of mitigating hydraulic 255 stress and their resilience to episodic drought [Chirino et al., 2011]. 256

Both species studied here are known to be drought resilient, with a relatively high basal increment 257 given the low transpiration measured. The natural range of *B. discolor* is slightly more mesic than 258 that of *B. populneus* [McCarthy et al., 2011], which might impact their respective hydraulic stress 259 mitigation strategies. Seedlings of *B. populneus* were found to have higher stomatal conductance 260 than B. discolor in the control setting and lower in the drought setting [McCarthy et al., 2011]. This 261 plasticity makes these seedlings better adapted to seasonal rainfall and enhances their water use 262 efficiency during droughts [Reynolds et al., 2018]. In the present study, B. populneus experiences a 263 more intense response (in terms of daily amplitude variations) to precipitation than B. discolor, which 264 suggests higher evapotranspiration and growth. Additionally, after several days without precipitation, 265 B. populneus exhibits a daily stem variation that approaches zero, potentially implying at least 266 partial stomatal closure. These results are in line with previous observations made on seedlings, and 267 support the current view on the drought resistance of these two species. 268

#### 4.2 Feasibility of DSTS and perspectives

The favourable comparison between the DSTS strain measurements and those obtained from an established instrument (the PépiPIAF dendrometer; see Fig. 2b) demonstrates that the fibre-based strain measurements reliably record the radial expansion and contraction, over time scales ranging ratio is comparable to that of the PépiPIAF LVDT sensor, facilitating detailed analyses like those presented here. Hence, DSTS offers a viable alternative to conventional dendrometric instruments, while holding the following advantages: 276

- A single instrument (the interrogator) is capable of continuously sensing up to 100 km of fibre 277 at a metric resolution, permitting the simultaneous monitoring of numerous positions on one or 278 multiple specimens. Compared to installing, maintaining, telemetering, and synchronising the 279 data of multiple instruments, the logistics of fibre-based instrumentation are greatly simplified. 280
- Fibre-optic cables are flexible yet robust to tension, resistant to heat, water, and sun exposure, 281 and they are electromagnetically inert (since no electric current runs through them). This 282 makes optical fibres highly suitable for long-term deployments in exposed (outdoor) settings. 283 Moreover, with a nominal cost of the order of 1 €/\$/£ per metre, the cost of replacement is 284 minimal, and damaged sections can be readily repaired on-site. Compared to conventional 285 electronics, fibre-optic cables require minimal protective measures for deployment in harsh 286 conditions.
- The inherently distributed nature of DSTS enables the investigation of the hydrological dynamics <sup>288</sup> in high spatio-temporal resolution. As aforementioned, follow-up studies that utilise fibre <sup>289</sup> segments located at a number of localities along the stem of a tree offer an opportunity to <sup>290</sup> elucidate fine-scale details of the storage and release of fluids. Moreover, the technique is <sup>291</sup> completely non-invasive, and does not require human intervention beyond the initial deployment. <sup>292</sup>

We therefore envision that DSTS will facilitate future long-term dendrometric monitoring programmes. Complemented with e.g. sap flow rate and hydraulic potential measurements, distributed dendrometry will lead to renewed insights into the species-specific response to climatic stressors, mitigation strategies, and long-term adaptability. 296

# 5 Conclusions

In this study we propose Distributed Strain & Temperature Sensing (DSTS) as a novel technology 298 for long-term, continuous dendrometry. To test the feasibility of DSTS for this purpose, we deployed 200 an optic fibre around the stems of two specimens of Brachychiton (B. discolor and B. populneus). 300 Brachychiton discolor was equipped with a conventional dendrometric instrument (PépiPIAF sensor) 301 that acted as a reference. From the comparison of the continuous DSTS strain recordings with those 302 derived from the conventional dendrometer, we conclude that the DSTS strain measurements are 303 both accurate and long-term stable, validating DSTS as a feasible methodology. Furthermore, the 304 spatially distributed DSTS measurements reveal intriguing lateral variations in the diurnal diametric 305 fluctuations of the stems, which we interpret to relate to the local storage and release of fluids, and 306 variations thereof driven by the availability of water. Hence, DSTS-based dendrometry has great 307 potential for complementing existing dendrometric techniques, and to deliver new insights into the 308 spatially-variable hydrological response of tree species. 309

# Appendix I: analytic signal processing

Consider a time-domain signal x(t). The corresponding analytic signal  $(\hat{x}(t))$  is obtained as  $\hat{x}(t) = 311$  $x(t) + j\mathcal{H}(x(t))$ , with  $\mathcal{H}(\cdot)$  denoting the Hilbert transform and  $j^2 \equiv -1$  being the imaginary unit. 312 This analytic representation is complex-valued, and so it can also be represented in polar form as: 313

$$\hat{x}(t) = A(t) \exp\left(-\jmath 2\pi\phi(t)\right) \tag{2}$$

with instantaneous amplitude A(t) and phase  $\phi(t)$ . Consider now two time series,  $\hat{x}_1(t)$  and  $\hat{x}_2(t)$  and that are filtered in a narrow frequency band around central frequency  $f_0$ . Multiplying  $\hat{x}_1$  with  $\hat{x}_2^*$  (\* denoting the complex conjugate), then gives: 316

$$\hat{x}_{1}\hat{x}_{2}^{*} = A_{1}(t)A_{2}(t)\exp\left(-\jmath 2\pi \left[\phi_{1}(t) - \phi_{2}(t)\right]\right)$$
  
=  $A_{1}(t)A_{2}(t)\exp\left(-\jmath 2\pi\Delta\phi(t)\right)$  (3)  
 $\approx A_{1}(t)A_{2}(t)\exp\left(-\jmath 2\pi f_{0}\Delta\tau(t)\right)$ 

Here,  $\Delta \tau(t)$  denotes the instantaneous time lag between signals  $x_1(t)$  and  $x_2(t)$ . Hence, the time lag is explicitly computed as:

$$\Delta \tau(t) = -\frac{\angle \left[\hat{x}_1 \hat{x}_2^*\right]}{2\pi f_0} \tag{4}$$

with  $\angle[\cdot]$  denoting the complex argument. Likewise, the instantaneous amplitude of x(t) is simply: 319

$$A(t) = |\hat{x}(t)| \tag{5}$$

For the analysis of the diurnal hydration cycles, we set  $f_0$  corresponding to a period of 24 hours. We then apply to the strain data a non-causal peak filter centred on  $f_0$  with a dimensionless quality factor of 3, resulting in a time series that only contains the diurnal variations. We subsequently transform the filtered data into its analytic form with Eq. (2), and estimate the time lag and

297

instantaneous amplitude with Eqs. (4) and (5), respectively. Note that the time lag inferred in this manner is independent of the time resolution of the measurement (5 minutes), and so time lags smaller than this time resolution can be correctly resolved.

# Acknowledgments

MvdE thanks A. Sladen (CNRS Géoazur) for making the DSTS instrument available for this 328 experiment, and P. Clement (Febus Optics) for in-depth discussion of the DSTS measurement 329 principle. EO, RA and GG thank the city of Antibes and especially P. Dalmasso for their financial 330 support (n° C10870) to this study. The authors thank P.-E. Lauri for his feedback on the initial draft 331 of the manuscript. For the analysis we made use of the following Python libraries: Matplotlib 3.7.2 332 [Hunter, 2007], NumPy 1.25.1 [Harris et al., 2020], Pandas 2.0.3 [Pandas Development Team, 2023], 333 SciPy 1.11.1 [Virtanen et al., 2020]. All the data and scripts needed to reproduce the figures in this 334 study are available here: https://doi.org/10.6084/m9.figshare.25773432. 335

# References

- T. Ameglio and P. Cruiziat. Daily Variations of Stem and Branch Diameter: Short Overview from a Developed Example. In T. K. Karalis, editor, *Mechanics of Swelling*, NATO ASI Series, pages 193–204, Berlin, Heidelberg, 1992. Springer. ISBN 978-3-642-84619-9. doi: 10.1007/978-3-642-84619-9\_9.
- T. Ameglio, H. Cochard, and F. W. Ewers. Stem diameter variations and cold hardiness in walnut trees. Journal of Experimental Botany, 52(364):2135–2142, 2001.
- T. Andriantelomanana, T. Améglio, S. Delzon, H. Cochard, and S. Herbette. Unpacking the point of no return under drought in poplar: insight from stem diameter variation. *New Phytologist*, 2024.
- I. Ashry, Y. Mao, Y. Al-Fehaid, A. Al-Shawaf, M. Al-Bagshi, S. Al-Brahim, T. K. Ng, and B. S. Ooi. Early detection of red palm weevil using distributed optical sensor. *Scientific reports*, 10(1):3155, 2020.
- L. Bouffaut, K. Taweesintananon, H. J. Kriesell, R. A. Rørstadbotnen, J. R. Potter, M. Landrø, S. E. Johansen, J. K. Brenne, A. Haukanes, O. Schjelderup, and F. Storvik. Eavesdropping at the Speed of Light: Distributed Acoustic Sensing of Baleen Whales in the Arctic. *Frontiers in Marine Science*, 9, 2022. ISSN 2296-7745.
- S. S. Burgess and T. E. Dawson. Using branch and basal trunk sap flow measurements to estimate whole-plant water capacitance: a caution. *Plant and Soil*, 305:5–13, 2008.
- J. Čermák, J. Kučera, W. L. Bauerle, N. Phillips, and T. M. Hinckley. Tree water storage and its diurnal dynamics related to sap flow and changes in stem volume in old-growth douglas-fir trees. *Tree physiology*, 27(2):181–198, 2007.
- K. Charra-Vaskou, E. Badel, G. Charrier, A. Ponomarenko, M. Bonhomme, L. Foucat, S. Mayr, and T. Améglio. Cavitation and water fluxes driven by ice water potential in juglans regia during freeze-thaw cycles. *Journal of Experimental Botany*, 67(3):739–750, 2016.
- G. Charrier, M. Nolf, G. Leitinger, K. Charra-Vaskou, A. Losso, U. Tappeiner, T. Améglio, and S. Mayr. Monitoring of freezing dynamics in trees: a simple phase shift causes complexity. *Plant Physiology*, 173(4):2196–2207, 2017.

- F. Chevallier, P. Balandier, J.-P. Nebout, D. Coste, B. B. Adam, and T. Ameglio. Pépipiaf: une nouvelle génération de biocapteurs pour le diagnostic stationnel en forêt. In *ECOTECHS'2011-Capteurs et Systèmes de Mesures pour les Applications Environnementales*, pages 49–52. CEMAGREF, 2011.
- E. Chirino, J. Bellot, and J. R. Sánchez. Daily sap flow rate as an indicator of drought avoidance mechanisms in five mediterranean perennial species in semi-arid southeastern spain. Trees, 25:593–606, 2011.
- B. Choat, T. J. Brodribb, C. R. Brodersen, R. A. Duursma, R. López, and B. E. Medlyn. Triggers of tree mortality under drought. *Nature*, 558(7711):531–539, 2018.
- P. Clément, R. Gabet, V. Lanticq, and Y. Jaouën. B-OTDR Solution for Independent Temperature and Strain Measurement in a Single Acquisition. *Journal of Lightwave Technology*, 39(18):6013–6020, Sept. 2021. ISSN 1558-2213. doi: 10.1109/JLT.2021.3088956.
- H. Cochard, S. Forestier, and T. Améglio. A new validation of the scholander pressure chamber technique based on stem diameter variations. *Journal of Experimental Botany*, 52(359):1361–1365, 2001.
- K. D. Coder. Tree anatomy: Periderm (bark). Warnell School of Forestry & Natural Resources, University of Georgia, 2022. Outreach Publication WSFNR-22-75C.
- P. Daget. Le bioclimat méditerranéen: caractères généraux, modes de caractérisation. Vegetatio, 34:1–20, 1977.
- F.-A. Daudet, T. Améglio, H. Cochard, O. Archilla, and A. Lacointe. Experimental analysis of the role of water and carbon in tree stem diameter variations. *Journal of Experimental Botany*, 56(409):135–144, 2005.
- D. M. Drew and G. M. Downes. The use of precision dendrometers in research on daily stem size and wood property variation: a review. *Dendrochronologia*, 27(2):159–172, 2009.
- C. Ducatillion and B.-C. Landy. L'art d'acclimater les plantes exotiques: le jardin de la Villa Thure. Editions Quae, 2010.
- R. Elliot and D. Jones. Encyclopedia of Australian Plants suitable for cultivation, Volume 2. Lothian publishing company pty. ltd., 1985.
- J. Fernández and M. Cuevas. Irrigation scheduling from stem diameter variations: a review. Agricultural and Forest Meteorology, 150(2):135–151, 2010.
- R. Gras. Etude des sols du parc de la Villa Thuret. Institut National de la Recherche Agronomique, 1990.
- C. R. Harris, K. J. Millman, S. J. Van Der Walt, R. Gommers, P. Virtanen, D. Cournapeau, E. Wieser, J. Taylor, S. Berg, N. J. Smith, R. Kern, M. Picus, S. Hoyer, M. H. Van Kerkwijk, M. Brett, A. Haldane, J. F. Del Río, M. Wiebe, P. Peterson, P. Gérard-Marchant, K. Sheppard, T. Reddy, W. Weckesser, H. Abbasi, C. Gohlke, and T. E. Oliphant. Array programming with NumPy. *Nature*, 585(7825):357–362, Sept. 2020. ISSN 0028-0836, 1476-4687. doi: 10.1038/s41586-020-2649-2.
- A. H. Hartog. An Introduction to Distributed Optical Fibre Sensors. CRC Press, Taylor & Francis Group, Boca Raton, FL, 2017. ISBN 978-1-351-64530-0.
- V. Hernandez-Santana, C. M. Rodriguez-Dominguez, J. Sebastian-Azcona, L. F. Perez-Romero, and A. Diaz-Espejo. Role of hydraulic traits in stomatal regulation of transpiration under different vapour pressure deficits across five mediterranean tree crops. *Journal of Experimental Botany*, 74(15):4597–4612, 2023.
- V. Herrmann, S. M. McMahon, M. Detto, J. A. Lutz, S. J. Davies, C.-H. Chang-Yang, and K. J. Anderson-Teixeira. Tree circumference dynamics in four forests characterized using automated dendrometer bands. *PloS one*, 11(12):e0169020, 2016.

- P. G. Hubbard, J. Xu, S. Zhang, M. Dejong, L. Luo, K. Soga, C. Papa, C. Zulberti, D. Malara, F. Fugazzotto, F. Garcia Lopez, and C. Minto. Dynamic structural health monitoring of a model wind turbine tower using distributed acoustic sensing (DAS). *Journal of Civil Structural Health Monitoring*, 11(3):833–849, July 2021. ISSN 2190-5479. doi: 10.1007/s13349-021-00483-y.
- J. D. Hunter. Matplotlib: A 2D Graphics Environment. Computing in Science & Engineering, 9(3):90–95, 2007. ISSN 1521-9615. doi: 10.1109/MCSE.2007.55.
- S. Iqbal, T. Zha, X. Jia, M. Hayat, D. Qian, C. P.-A. Bourque, Y. Tian, Y. Bai, P. Liu, R. Yang, et al. Interannual variation in sap flow response in three xeric shrub species to periodic drought. *Agricultural and Forest Meteorology*, 297:108276, 2021.
- C. Y. Jim. Urban heritage trees: Natural-cultural significance informing management and conservation. *Greening cities: forms and functions*, pages 279–305, 2017.
- P. Jousset, T. Reinsch, T. Ryberg, H. Blanck, A. Clarke, R. Aghayev, G. P. Hersir, J. Henninges, M. Weber, and C. M. Krawczyk. Dynamic strain determination using fibre-optic cables allows imaging of seismological and structural features. *Nature Communications*, 9(1):2509, July 2018. ISSN 2041-1723. doi: 10.1038/s41467-018-04860-y.
- M. Kottek, J. Grieser, C. Beck, B. Rudolf, and F. Rubel. World map of the köppen-geiger climate classification updated. *Meteorologische Zeitschrift*, 2006.
- T. Kume, H. Komatsu, K. Kuraji, and M. Suzuki. Less than 20-min time lags between transpiration and stem sap flow in emergent trees in a Bornean tropical rainforest. Agricultural and Forest Meteorology, 148(6):1181–1189, June 2008. ISSN 0168-1923. doi: 10.1016/j.agrformet.2008.02.010.
- L. Lamacque, G. Charrier, F. d. S. Farnese, B. Lemaire, T. Améglio, and S. Herbette. Drought-induced mortality: branch diameter variation reveals a point of no recovery in lavender species. *Plant physiology*, 183(4):1638–1649, 2020.
- L. Lamacque, F. Sabin, T. Améglio, S. Herbette, and G. Charrier. Detection of acoustic events in lavender for measuring xylem vulnerability to embolism and cellular damage. *Journal of Experimental Botany*, 73(11):3699–3710, 2022.
- N. J. Lindsey and E. R. Martin. Fiber-Optic Seismology. Annual Review of Earth and Planetary Sciences, 49(1):309–336, May 2021. ISSN 0084-6597. doi: 10.1146/annurev-earth-072420-065213.
- J. Liu, S. Yuan, B. Luo, B. Biondi, and H. Y. Noh. Turning Telecommunication Fiber-Optic Cables into Distributed Acoustic Sensors for Vibration-Based Bridge Health Monitoring. *Structural Control and Health Monitoring*, 2023:1–14, Apr. 2023. ISSN 1545-2263, 1545-2255. doi: 10.1155/2023/3902306.
- S. Liu, W. Bitterlich, C. J. Cieszewski, and M. J. Zasada. Comparing the use of three dendrometers for measuring diameters at breast height. *Southern journal of applied forestry*, 35(3):136–141, 2011.
- R. M. Marchin, D. Backes, A. Ossola, M. R. Leishman, M. G. Tjoelker, and D. S. Ellsworth. Extreme heat increases stomatal conductance and drought-induced mortality risk in vulnerable plant species. *Global change biology*, 28(3):1133–1146, 2022.
- T. Martin, K. Brown, J. Kučera, F. Meinzer, D. Sprugel, and T. Hinckley. Control of transpiration in a 220-year-old abies amabilis forest. *Forest Ecology and Management*, 152(1-3):211–224, 2001.
- S. M. Maughan, H. H. Kee, and T. P. Newson. Simultaneous distributed fibre temperature and strain sensor using microwave coherent detection of spontaneous Brillouin backscatter. *Measurement Science and Technology*, 12(7):834–842, July 2001. ISSN 0957-0233, 1361-6501. doi: 10.1088/0957-0233/12/7/315.
- H. R. McCarthy, D. E. Pataki, and G. D. Jenerette. Plant water-use efficiency as a metric of urban ecosystem services. *Ecological Applications*, 21(8):3115–3127, 2011.

- S. M. McMahon and G. G. Parker. A general model of intra-annual tree growth using dendrometer bands. *Ecology and Evolution*, 5(2):243–254, 2015.
- A. Moriana, D. Pérez-López, M. Prieto, M. Ramírez-Santa-Pau, and J. Pérez-Rodriguez. Midday stem water potential as a useful tool for estimating irrigation requirements in olive trees. Agricultural Water Management, 112:43–54, 2012.
- t. Pandas Development Team. Pandas-dev/pandas: Pandas. Zenodo, Nov. 2023.
- N. Phillips, R. Oren, R. Zimmermann, and S. J. Wright. Temporal patterns of water flux in trees and lianas in a Panamanian moist forest. *Trees*, 14(3):0116, 1999. ISSN 09311890. doi: 10.1007/s004680050216.
- V. A. Reynolds, L. D. Anderegg, X. Loy, J. HilleRisLambers, and M. M. Mayfield. Unexpected drought resistance strategies in seedlings of four brachychiton species. *Tree physiology*, 38(5):664–677, 2018.
- D. Rivet, B. de Cacqueray, A. Sladen, A. Roques, and G. Calbris. Preliminary assessment of ship detection and trajectory evaluation using distributed acoustic sensing on an optical fiber telecom cable. *The Journal of the Acoustical Society of America*, 149(4):2615–2627, Apr. 2021. ISSN 0001-4966. doi: 10.1121/10.0004129.
- F. A. Roesch, T. A. Schroeder, and C. A. Price. The usda forest service's forest inventory and analysis database and the national register of champion trees—a potentially symbiotic relationship. *Journal of Forestry*, page fvad058, 2024.
- J. Roignant, É. Badel, N. Leblanc-Fournier, N. Brunel-Michac, J. Ruelle, B. Moulia, and M. Decourteix. Feeling stretched or compressed? the multiple mechanosensitive responses of wood formation to bending. *Annals of Botany*, 121(6):1151–1161, 2018.
- F. C. Scholz, S. J. Bucci, G. Goldstein, F. C. Meinzer, A. C. Franco, and F. Miralles-Wilhelm. Temporal dynamics of stem expansion and contraction in savanna trees: withdrawal and recharge of stored water. *Tree Physiology*, 28(3):469–480, 2008.
- S. Sevanto, T. Vesala, M. Perämäki, and E. Nikinmaa. Time lags for xylem and stem diameter variations in a scots pine tree. Plant, Cell & Environment, 25(8):1071–1077, 2002.
- T. Simonneau, R. Habib, J.-P. Goutouly, and J.-G. Huguet. Diurnal Changes in Stem Diameter Depend Upon Variations in Water Content: Direct Evidence in Peach Trees. *Journal of Experimental Botany*, 44(3):615–621, Mar. 1993. ISSN 0022-0957. doi: 10.1093/jxb/44.3.615.
- S. L. Steinberg, M. J. McFARLAND, and J. W. WORTHINGTON. Comparison of trunk and branch sap flow with canopy transpiration in pecan. *Journal of experimental botany*, 41(6):653–659, 1990.
- K. Steppe and R. Lemeur. An experimental system for analysis of the dynamic sap-flow characteristics in young trees: results of a beech tree. *Functional Plant Biology*, 31(1):83–92, 2004.
- M. Ueda, K. Yoshikawa, and J. Okitu. Measurement of diurnal changes in stem and branch diameters using strain gauges. Journal of Forest Research, 1(3):139–142, Aug. 1996. ISSN 1610-7403. doi: 10.1007/BF02348191.
- M. P. A. van den Ende and J.-P. Ampuero. Evaluating seismic beamforming capabilities of distributed acoustic sensing arrays. Solid Earth, 12(4):915–934, Apr. 2021. ISSN 1869-9510. doi: 10.5194/se-12-915-2021.
- P. Virtanen, R. Gommers, T. E. Oliphant, M. Haberland, T. Reddy, D. Cournapeau, E. Burovski, P. Peterson, W. Weckesser, J. Bright, S. J. van der Walt, M. Brett, J. Wilson, K. J. Millman, N. Mayorov, A. R. J. Nelson, E. Jones, R. Kern, E. Larson, C. J. Carey, İ. Polat, Y. Feng, E. W. Moore, J. VanderPlas, D. Laxalde, J. Perktold, R. Cimrman, I. Henriksen, E. A. Quintero, C. R. Harris, A. M. Archibald, A. H. Ribeiro, F. Pedregosa, P. van Mulbregt, and S. . . Contributors. SciPy 1.0–Fundamental Algorithms for Scientific Computing in Python. *Nature Methods*, 17(3):261–272, Mar. 2020. ISSN 1548-7091, 1548-7105. doi: 10.1038/s41592-019-0686-2.

- J. Wang and T. Sammis. New automatic band and point dendrometers for measuring stem diameter growth. *Applied Engineering* in Agriculture, 24(6):731–742, 2008.
- M. Wilmking, M. van der Maaten-Theunissen, E. van der Maaten, T. Scharnweber, A. Buras, C. Biermann, M. Gurskaya, M. Hallinger, J. Lange, R. Shetti, et al. Global assessment of relationships between climate and tree growth. *Global Change Biology*, 26(6):3212–3220, 2020.
- W. Zhao and B. Liu. The response of sap flow in shrubs to rainfall pulses in the desert region of china. Agricultural and Forest Meteorology, 150(9):1297–1306, 2010.
- R. Zweifel. Radial stem variations-a source of tree physiological information not fully exploited yet. *Plant, Cell & Environment*, 39(2):231–232, 2016.
- R. Zweifel and R. Häsler. Frost-induced reversible shrinkage of bark of mature subalpine conifers. Agricultural and forest meteorology, 102(4):213–222, 2000.
- R. Zweifel, H. Item, and R. Häsler. Stem radius changes and their relation to stored water in stems of young norway spruce trees. Trees, 15:50–57, 2000.
- R. Zweifel, H. Item, and R. Häsler. Link between diurnal stem radius changes and tree water relations. *Tree physiology*, 21 (12-13):869–877, 2001.
- R. Zweifel, L. Zimmermann, F. Zeugin, and D. M. Newbery. Intra-annual radial growth and water relations of trees: implications towards a growth mechanism. *Journal of experimental botany*, 57(6):1445–1459, 2006.
- R. Zweifel, M. Haeni, N. Buchmann, and W. Eugster. Are trees able to grow in periods of stem shrinkage? New Phytologist, 211(3):839–849, 2016.