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Atmospheric drivers of storage water use in Scots pine

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BGD

4, 615–650, 2007

Drivers of storage water use in Scots pine

H. Verbeeck et al.

Title Page

Abstract

Introduction

Conclusions

References

Tables

Figures

◀

▶

◀

▶

Back

Close

Full Screen / Esc

Printer-friendly Version

Interactive Discussion

EGU

Abstract

In this study we determined the microclimatic drivers of storage water use in Scots pine (*Pinus sylvestris* L.) growing in a temperate climate. The storage water use was modeled using the ANAFORE model, integrating a dynamic water flow and – storage model with a process-based transpiration model. The model was calibrated and validated with sap flow measurements for the growing season of 2000 (26 May–18 October).

Because there was no severe soil drought during the study period, we were able to study atmospheric effects. Incoming radiation was the main driver of storage water use. The general trends of sap flow and storage water use are similar, and follow more or less the pattern of incoming radiation. Nevertheless, considerable differences in the day-to-day pattern of sap flow and storage water use were observed, mainly driven by vapour pressure deficit (VPD). During dry atmospheric conditions (high VPD) storage water use was reduced. This reduction was disproportionately higher than the reduction in measured sap flow. Our results suggest that the trees did not rely more on storage water during periods of atmospheric drought, without severe soil drought.

A third important factor was the tree water deficit. When storage compartments were depleted beyond a threshold, storage water use was limited due to the low water potential in the storage compartments. The maximum relative contribution of storage water to daily transpiration was also constrained by an increasing tree water deficit.

1 Introduction

The water balance of a tree depends on the transpirational loss, uptake from the soil and internal storage of water. Storage water use or refilling may induce a considerable time lag between transpiration and sap flow near the base of the stem (Goldstein et al., 1998). Stored water plays a biologically significant role in trees (Čermák et al., 2007). It is accessible to the individual tree only, unlike soil water, which may be taken up by other plants or lost through evaporation or drainage. Water released from storage in

BGD

4, 615–650, 2007

Drivers of storage water use in Scots pine

H. Verbeeck et al.

Title Page

Abstract

Introduction

Conclusions

References

Tables

Figures

◀

▶

◀

▶

Back

Close

Full Screen / Esc

Printer-friendly Version

Interactive Discussion

EGU

the stem serves as buffer to preclude low xylem water potentials that might otherwise be caused by peaks of transpiration. Therefore, the xylem water potential is smoothed along the flow path, which may help avoid xylem cavitation. Trees can use stored water to allow continued photosynthesis when transpiration exceeds soil water uptake (Zweifel et al., 2001). Despite the importance of water storage in regulating the water budget of trees, limited information exists on the contribution of internal water storage to their total daily water consumption (Meinzer et al., 2001).

Several authors found that the contribution of internal water to transpiration is not a constant in the water budget of trees (Loustau et al., 1996; Köstner et al., 1998; Zweifel et al., 2005; Verbeeck et al., 2007), but changes dynamically from day to day. To determine the complete water balance of a forest ecosystem and its dynamics, it is therefore important to know which factors influence the use of stored water.

This paper investigates the effect of the atmospheric microclimate on the use of stored water of Scots pine for the growing season of 2000 (26 May–18 October). Since there was no severe soil drought during the study period, we were able to investigate the effect of atmospheric drought. Low soil water availability and high evaporative demand are often treated together as aspects of drought stress. However, their effects on plants may differ: the first is a measure of how much water is available in the source, while the second (high evaporative demand) reflects the sink effect of the atmosphere (Bhaskar and Ackerly, 2006). In contrast to low soil water availability, the influence of high evaporative demand or atmospheric drought on hydraulic traits is less well understood, in part due to the difficulty of isolating this factor (Zweifel et al., 2005).

Several experimental studies mentioned by Čermák et al. (2007), calculate storage water use from sap flow measurements at different heights. In these studies, sap flow measurements at the top of the canopy are used as a measure for the transpiration at leaf level. Over the last couple of years, several models that take into account water storage have been developed to model the non-steady-state flow of water through a tree (e.g. Zweifel et al., 2005; Steppe et al., 2006), using transpiration as an input value.

Nevertheless, as suggested by Chuang et al. (2006), a stomatal model can easily

BGD

4, 615–650, 2007

**Drivers of storage
water use in Scots
pine**

H. Verbeeck et al.

Title Page

Abstract

Introduction

Conclusions

References

Tables

Figures

◀

▶

◀

▶

Back

Close

Full Screen / Esc

Printer-friendly Version

Interactive Discussion

EGU

be coupled with a model of the hydraulic system of a tree to provide independent time series of transpiration. This has been done by Lhomme et al. (2001) using an empirical Jarvis-type stomatal model and a big leaf transpiration model, and by Verbeeck et al. (2007) using the process-based ANAFORE model.

5 In this paper, the ANAFORE model was used to model sap flow and transpiration for various Scots pine (*Pinus sylvestris* L.) trees in Brasschaat, Belgium. This model integrates a water flow and storage submodel in a process-based transpiration model (companion paper: Verbeeck et al., 2007). The goal of this study was to determine the drivers of day-to-day and diurnal variation of storage water use.

10 2 Materials and methods

2.1 Site description

The forest under investigation is “De Inslag”, a mixed patchy coniferous/deciduous forest located in Brasschaat (51°18′33″ N, 4°31′14″ E), in the Belgian Campine region, about 20 km north-east of Antwerp. The site is part of the European Carboeurope-IP network and is a level-II observation plot (ICP-II forests) for intensive monitoring of forest ecosystems (EC-UN/ECE, 1996), managed by the Ministry of the Flemish Community (AMINAL). The landscape is a coastal plain, almost flat (slope <0.3%) and at a mean elevation of 16 m. The climate is temperate maritime with a mean annual temperature of 9.8°C and 750 mm of annual precipitation (30 years average). The site is located in an area with high nitrogen deposition (30–40 kg ha⁻¹ year⁻¹; Neiryneck et al., 2002).

This relatively small (150 ha) forest consists of many patches of different coniferous and deciduous species, with a variety of understorey species. This paper focuses on one particular Scots pine (*Pinus sylvestris* L.) stand, described in detail by Curiel Yuste et al. (2005). The even-aged Scots pine stand was planted in 1929. In 1997, tree density was 556 trees ha⁻¹. In November 1999, a thinning was performed reducing

BGD

4, 615–650, 2007

Drivers of storage water use in Scots pine

H. Verbeeck et al.

Title Page

Abstract

Introduction

Conclusions

References

Tables

Figures

◀

▶

◀

▶

Back

Close

Full Screen / Esc

Printer-friendly Version

Interactive Discussion

EGU

the stem number to 377 trees ha⁻¹. In winter 1999-2000, the mean DBH (Diameter at Breast Height) was 29.4 cm, the mean height 21.4 m.

The soil is loamy sand, moderately wet, with a distinct humus and iron B-horizon (Baeyens et al., 1993) and is classified as an Umbric Regosol (FAO classification, Roskams et al., 1997). Below the sandy layer, at a depth of 1 to 3 m, lies a clay layer, making that the site has poor drainage, and groundwater depth is usually between 1.2 and 1.5 m (Baeyens et al., 1993). The soil is typically moist, but rarely saturated because of the high hydraulic conductivity of the upper sandy layers. More details about the physical and chemical properties of the soil can be found in Janssens et al. (1999) and Neiryneck et al. (2002).

2.2 Sap flow measurements

The sap flow data used in this study were obtained during a measurement campaign during the growing season of 2000. From 26 May until 18 October 2000, sap flow was measured on 14 sample trees (two groups of 7 trees). The main attributes of the sampled trees are given in Table 1. Leaf area of the sample trees was determined according to the method explained in Čermák et al. (1998), based on destructive measurements on a series of sample trees from the same stand. Sapwood area was estimated from the radial sap flow profiles: the border between sapwood and heartwood was found where sap flow reached zero.

The sap flow in the sample trees was measured with the heat field deformation (HFD) method (Nadezhdina et al., 1998, 2004, 2006; Nadezhdina and Čermák, 2000; Čermák et al., 2004). This method is based on the observed changes of an artificial heat field around a linear heater in a tree stem, which depends on the sap flow rate and the xylem tissue properties. Deformation of the heat field generated by the linear heater in a certain tangential section of the stem is characterized by the ratio of temperature gradients measured around the heater in the axial and tangential directions. This ratio represents the basis for sap flow calculation.

BGD

4, 615–650, 2007

Drivers of storage water use in Scots pine

H. Verbeeck et al.

Title Page

Abstract

Introduction

Conclusions

References

Tables

Figures

◀

▶

◀

▶

Back

Close

Full Screen / Esc

Printer-friendly Version

Interactive Discussion

EGU

Two types of sensors were used for the sap flow measurements in the sample trees: small, single-point sensors for routine sap flow measurements during the entire growing season and long, multi-point sensors for the determination of the variable flow pattern along the xylem radius in the tree stem. Each sap flow sensor consisted of two sets of differential thermocouples and a linear heater. Needles of the multi-point sensor contained six thermocouples, 10 or 16 mm apart. Four multi-point sensors were used, resulting in 24 measurement points. By moving the needles along the radius at half the distance between the thermocouples (5 or 8 mm), 24 additional measuring points were obtained. The conducting system of stems at breast height was therefore characterized by a total of 48 measuring points in each sample tree. The radial pattern was used to scale-up the flow from routine single-point measurements to the whole tree level, according to the procedure previously described by Nadezhdina et al. (2002). The sap flow was measured every minute and recorded as means over 15 min intervals by data loggers (EMS-12, Unilog & Environmental Measuring Systems Inc., Brno, Czech Republic).

2.3 Supporting measurements

Meteorological parameters, i.e., global radiation (pyranometer, Kipp and Zonen CM6B, the Netherlands), net radiation (REBS 07, Seattle, WA, USA), photosynthetically active radiation (PAR, JYP-1000 sensor, SDEC, Tours, France), air temperature and relative humidity (Didcot Instrument Co Ltd, Abingdon, United Kingdom DTS-5A), wind-speed (Didcot DWR-250G) and precipitation (Didcot DRG-51) were continuously measured at the top of a tower (40 m) located within the Scots pine stand, and recorded half-hourly on a data logger (Campbell CR10, CSI, Logan, UT, USA). More details about the instruments and methods (e.g., gapfilling) used at the site can be found in Overloop and Meiresonne (1999), Kowalski et al. (2000) and Carrara et al. (2003).

Soil moisture was measured twice a week at two locations in the vicinity of the two groups of 7 Scots pine trees with a series of TDR-sensors (Time Domain Reflectometry), placed every 25 cm down to a depth of 175 cm (cabletester: Tektonix 1502B,

BGD

4, 615–650, 2007

Drivers of storage water use in Scots pine

H. Verbeeck et al.

Title Page

Abstract

Introduction

Conclusions

References

Tables

Figures

◀

▶

◀

▶

Back

Close

Full Screen / Esc

Printer-friendly Version

Interactive Discussion

EGU

Redmond, USA). Volumetric soil water content measurements were linearly interpolated to obtain estimates for the entire measurement period. Soil water potential was calculated from the volumetric soil water content using the power function model of van Genuchten (1980). Parameters for the soil water retention curve were determined at the site by Meiresonne et al. (2003).

2.4 Model description

The ANAFORE model (ANALysis of FORest Ecosystems) is a mechanistic tree and forest stand model. To model water fluxes, it integrates a dynamic water flow model in a process based transpiration model as described in a companion paper by Verbeeck et al. (2007). Stomatal aperture, sap flow, photosynthesis (Farquhar et al., 1980) and transpiration (Penman, 1948; Monteith, 1965) are simulated on a half-hourly basis. The most relevant equations to model water fluxes are described below.

The stomatal conductance model is derived from the formula of Dewar (2002), and described in Verbeeck et al. (2007):

$$g_s = \frac{[a_1(A_n + R_d) + b]}{c_i \left(1 + \frac{VPD^2}{D_0}\right)} \exp\{-[ABA]\beta \exp(\delta\psi_{leaf})\} \quad (1)$$

where g_s is the stomatal conductance to CO_2 ($\text{mol CO}_2 \text{ m}^{-2} \text{ s}^{-1}$), A_n is the net leaf CO_2 assimilation rate ($\text{mol CO}_2 \text{ m}^{-2} \text{ s}^{-1}$), R_d is the dark respiration rate ($\text{mol CO}_2 \text{ m}^{-2} \text{ s}^{-1}$), c_i is the intercellular CO_2 concentration ($\text{mol CO}_2 \text{ mol}^{-1} \text{ air}$), VPD is the atmospheric vapour pressure deficit (kPa), a_1 (dimensionless), b ($\text{mol CO}_2 \text{ m}^{-2} \text{ s}^{-1}$) and D_0 (kPa) are empirical constants which are parameter combinations, [ABA] is the leaf xylem concentration of abscisic acid (ABA) (mol ABA m^{-3}), ψ_{leaf} is the leaf water potential (MPa), β is the basal sensitivity of stomata to ABA ($\text{m}^3 \text{ mol}^{-1} \text{ ABA}$) and δ (MPa^{-1}) is a factor relating stomatal ABA sensitivity to ψ_{leaf} . According to Dewar (2002), the leaf

BGD

4, 615–650, 2007

Drivers of storage water use in Scots pine

H. Verbeeck et al.

Title Page

Abstract

Introduction

Conclusions

References

Tables

Figures

◀

▶

◀

▶

Back

Close

Full Screen / Esc

Printer-friendly Version

Interactive Discussion

EGU

xylem concentration of ABA is calculated as follows:

$$[ABA] = \frac{-\lambda_{\text{root}}\psi_{\text{root}} - \lambda_{\text{leaf}}\psi_{\text{leaf}}}{V_w(E + a)} \quad (2)$$

where ψ_{root} and ψ_{leaf} are the root and leaf water potentials (MPa), λ_{root} en λ_{leaf} are the root and leaf ABA synthesis coefficients, V_w is the molal volume of H_2O , E is the transpiration rate and a is the ABA sequestration rate. Stomatal conductance (Eq. 1) is coupled with the photosynthesis model of Farquhar et al. (1980) via a system of equations. This system is solved according to Op de Beeck et al. (2007).

The actual transpiration at leaf level is calculated according to the Penman-Monteith approach (Penman, 1948; Monteith, 1965):

$$\lambda E = \frac{s(R_n - G) + \rho_a c_p g_b (\text{VPD})}{s + \gamma \left(\frac{g_b}{g_w} \right)} \quad (3)$$

where E is the canopy transpiration ($\text{kg m}^{-2}\text{s}^{-1}$), λ the latent heat of vaporization (J kg^{-1}), s the slope of the saturated water vapour pressure curve ($\text{Pa } ^\circ\text{C}^{-1}$), R_n the net radiation above the canopy layer (W m^{-2}), G the sensible heat storage (W m^{-2}), ρ_a the density of the air (kg m^{-3}), c_p the specific heat of dry air ($\text{J kg}^{-1} ^\circ\text{C}^{-1}$), g_b the boundary layer conductance for H_2O (m s^{-1}), VPD the vapour pressure deficit (Pa), γ the psychrometric constant ($\text{Pa } ^\circ\text{C}^{-1}$), and g_w the total conductance for H_2O (m s^{-1}) between the evaporating site and the bulk air (i.e., the series sum of the stomatal and boundary layer conductances) (Jones, 1992). The sunlit and shaded leaf area of each canopy layer is used to scale up from transpiration at leaf level to the entire tree.

The sap flow submodel used in ANAFORE is based on the electrical analogon approach and is a simplified version of the water transport submodel of the RCGro model (Steppe et al., 2006). A diagram of the model is given in Fig. 1. This simplified dynamic flow and storage model uses three flow resistances and one capacitance. Transpiration of an individual tree and soil water potential are the inputs to calculate sap flow and

BGD

4, 615–650, 2007

Drivers of storage water use in Scots pine

H. Verbeeck et al.

Title Page

Abstract

Introduction

Conclusions

References

Tables

Figures

◀

▶

◀

▶

Back

Close

Full Screen / Esc

Printer-friendly Version

Interactive Discussion

EGU

water storage. The vertical water transport in the xylem can be described by Ohm's law:

$$E = \frac{\psi_{\text{stem}} - \psi_{\text{leaf}}}{R_X} \quad (4)$$

$$F_{\text{tree}} = \frac{\psi_{\text{soil}} - \psi_{\text{stem}}}{R_X} \quad (5)$$

5 where E is the transpiration rate (mg s^{-1}), F_{tree} is the sap flow in the stem (mg s^{-1}), ψ_{soil} is the soil water potential (MPa), ψ_{stem} is the water potential in the stem xylem and R_X is the xylem flow resistance (MPa s mg^{-1}). Both xylem flow resistances are assumed equal. Beside vertical water transport, internally stored water can contribute to the daily transpiration loss because of the hydraulic connection between the xylem and storage pools. This contribution depends on the magnitude of the hydraulic exchange resistance, R_S (MPa s mg^{-1}) that must be overcome when water flows from the storage compartments to the xylem to equilibrate the imbalance between water supply and demand during the day (Lhomme et al., 2001). Water flow to or from the storage pools f_{storage} (mg s^{-1}) can be expressed as:

$$15 \quad f_{\text{storage}} = \frac{\psi_{\text{stem}} - \psi_{\text{storage}}}{R_S} \quad (6)$$

where ψ_{storage} is the water potential (MPa) in the storage compartment. Values of f_{storage} are negative when water is withdrawn from the storage pool. Flow rates out of the xylem compartments can be derived from Kirchhoff's electric current law (Fig. 1):

$$E = F_{\text{tree}} - f_{\text{storage}} \quad (7)$$

20 The capacitance C (mg MPa^{-1}) of the storage tissue can be defined as the ratio between the change in amount of water in the storage tissue and the change in water potential of the tissue (e.g., Hunt, 1991; Phillips et al., 2003).

$$C = \frac{\Delta W}{\Delta \psi_{\text{storage}}} \quad (8)$$

Drivers of storage water use in Scots pine

H. Verbeeck et al.

Title Page

Abstract

Introduction

Conclusions

References

Tables

Figures

◀

▶

◀

▶

Back

Close

Full Screen / Esc

Printer-friendly Version

Interactive Discussion

where W is the water content of the tree (mg), corresponding to the total charge on the capacitor of an electrical circuit. The capacitance is considered here as a constant parameter per tree (Steppe et al., 2006).

The flow rate to or from the storage pool f_{storage} described in Eq. (7) is equivalent to the change in water content of the storage pool:

$$f_{\text{storage}} = \frac{dW}{dt} \quad (9)$$

Our sap flow model combines the above set of algebraic and differential Eqs. (4)–(9). As initial conditions the water potential of the storage tissue is considered to be zero when the water content is maximal. The maximum water content (W_{max} , kg) of the 14 pine trees (Table 2) was estimated by taking 40% of the stem volume (Steppe et al., 2004). From the difference between W_{max} and the minimum of W during the entire studied period, the maximum available tree water content (AW_{max} , kg) could be calculated. The tree water deficit (WD , %) can be defined as the actual depleted fraction of W_{max} .

The entire set of equations of the stomatal, photosynthesis, transpiration and sap flow models is solved numerically for each time step, by an iteration procedure that converges to an equilibrium leaf water potential using the “golden section search” procedure (Press et al., 1996). The leaf water potential of the previous time step is used as an initial value to start the procedure.

As stated in Eq. (7), the storage water flow can be calculated from the difference between sap flow (measured or modeled) and modeled transpiration. Positive values of storage water flow indicate that water storage is being recharged. Negative values indicate that storage water is being used to fulfill the transpiration needs of the leaves. Each day, the use of storage water can be estimated from the curve of the storage water flow as the integrated area below the zero line. For the entire studied period the maximum daily storage water use (STO_{max} , kg day⁻¹) and the maximum contribution of storage water to daily transpiration (%) were calculated for each individual tree.

Drivers of storage water use in Scots pineH. Verbeeck et al.

Title Page

Abstract

Introduction

Conclusions

References

Tables

Figures

◀

▶

◀

▶

Back

Close

Full Screen / Esc

Printer-friendly Version

Interactive Discussion

2.5 Model parameterization and validation

In a companion paper, the model was parameterized and validated using sap flow measurements and storage estimates for the 14 pine trees (Verbeeck et al., 2007). Parameter values (used in the equations to calculate stomatal conductance, sap flow, photosynthesis) are based on measurements at the Brasschaat site or on general parameters from the literature (Sampson et al., 2006; Verbeeck et al., 2007). A subset of three parameters (a_1 , C and R_X) was selected to be calibrated with half-hourly sap flow data (Verbeeck et al., 2007). The validation of the model revealed that a calibration using only the first two weeks of sap flow data yielded parameter values that were valid for the rest of the growing season (145 days of data) (Verbeeck et al., 2007). Nevertheless, for the present study we re-calibrated the model every two weeks of the growing season. The two-weekly parameterization yielded a slightly better fit to the sap flow measurements, but no clear trends (seasonal or climatic) could be found in the parameters. Therefore, the results of this parameterization are not presented or discussed further in this paper.

2.6 Site conditions

For the analysis in this study we selected six weeks (May 2–July 7, 2000) of data that contained sunny days with high sap flow rates as well as cloudy days with lower sap flow. Moreover, the 3rd and 4th week had almost no precipitation and the 4th week showed a peak in temperature and VPD (Fig. 2).

Although the soil water potential reached the lowest value of the growing season (June 23) due to a fortnight without precipitation, there was never severe soil water stress for the trees. Soil water potential dropped only once below field capacity (-0.03 MPa) in the top soil layer (25 cm). The deeper layers, which can be easily reached by the roots of the pines, were always above field capacity. A regression showed that soil water content could only explain less than 4% of the variability in the daily sap flow measurements. This could be expected because transpiration of woody species

BGD

4, 615–650, 2007

Drivers of storage water use in Scots pine

H. Verbeeck et al.

Title Page

Abstract

Introduction

Conclusions

References

Tables

Figures

◀

▶

◀

▶

Back

Close

Full Screen / Esc

Printer-friendly Version

Interactive Discussion

EGU

remains unaffected until the soil water fraction has decreased to a level between 23% and 32% of field capacity (Sinclair et al., 2005). However, soil water fraction is not the only factor influencing water availability for tree. Several other factors can play a role (e.g. level of the water table, hydraulic lift, . . .). Nevertheless, the absence of severe soil water stress for the pine trees during the study period gave the opportunity to investigate the effects of atmospheric drought.

2.7 Selection of typical days to analyze microclimate effects

Based on the daily measured sap flow and the modeled contribution of storage water to daily transpiration of all pine trees (example in Fig. 3 for pine #23) we selected four typical days for a detailed analysis of the storage water use: 4 June and 9 June before the period of high VPD, 20 June during the period of high VPD and 24 June shortly after that period. These four days were paired as such that both pairs had two days with equal measured daily sap flow, but with contrasting storage water use. The first pair (4 June and 24 June) contained the days with a moderate sap flow and entirely different climatic conditions. 4 June had high incoming radiation, temperature and VPD. In contrast, on 24 June incoming radiation, temperature and VPD were low. The second pair (9 June and 20 June) was chosen to examine the effect of high VPD. Both days had similar high incoming radiation, but the temperature and VPD of 20 June were much higher as compared to 9 June.

3 Results and discussion

3.1 Variability of storage water use

The pine trees showed a considerable day-to-day variation in the relative contribution of storage water use to daily transpiration (Fig. 3). The variation in absolute daily use of stored water was even larger (data not shown). In a companion paper (Verbeeck et

Drivers of storage water use in Scots pine

H. Verbeeck et al.

Title Page

Abstract

Introduction

Conclusions

References

Tables

Figures

◀

▶

◀

▶

Back

Close

Full Screen / Esc

Printer-friendly Version

Interactive Discussion

al., 2007) we already showed that the general seasonal trend of daily storage water use is similar to that of the daily transpiration or sap flow. In contrast, Fig. 3 showed different patterns in measured sap flow and relative contributions of stored water to daily transpiration. Days with an equal amount of sap flow had very different amounts of storage water use. Comparing 4 June with 24 June for all trees shows that, except for trees #13 and #16, all trees had a similar amount of sap flow on both days, but all trees showed a much lower storage water use on 24 June (Fig. 4). The absolute amount of storage water use varied from tree to tree and could be linked to the dimensions of the trees, as previously described (Verbeeck et al., 2007).

Maximum use of stored water over the growing season varied among trees from 4.6 to 14.6 kg day⁻¹ (Table 1). This corresponds well to the average value of 14 kg that Köstner et al. (1998) found for easily available water in *Picea abies* trees. The maximum contribution of the storage water to daily transpiration varied between 28 and 45%. These values fall within the wide range of values that can be found in the literature: ranging from 1 up to 50% (Čermák et al., 2007) or even up to 65% (Zweifel et al., 2001).

3.2 Radiation is the main driver of storage water use

When plotting half-hourly sap flow and transpiration against incoming radiation (Fig. 5), it became clear that radiation is an important factor influencing the difference between these two flows, and thus the use of water from the storage compartments. With higher radiation the increase in sap flow is less steep than the increase in transpiration due to the buffering effect of storage water. Storage water use was high at high rates of incoming radiation and storage water recharge took place when incoming radiation was low (Fig. 6a). On the other hand, when the storage water flow was plotted against VPD (Fig. 6b) or temperature (data not shown) the effect of these factors on storage water use was less pronounced than the effect of incoming radiation.

Two days with similar measured sap flow but with totally different climatic conditions were compared (Fig. 7). 4 June was warm and sunny and 24 June was cool and cloudy

Drivers of storage water use in Scots pine

H. Verbeeck et al.

Title Page

Abstract

Introduction

Conclusions

References

Tables

Figures

◀

▶

◀

▶

Back

Close

Full Screen / Esc

Printer-friendly Version

Interactive Discussion

with an incoming radiation of 19.1 and 8.2 MJ m² day⁻¹, respectively. These differences in microclimate and thus in transpiration caused a large difference in storage water use (e.g. 13.3 and 1.0 kg day⁻¹, respectively for pine tree #23). Although modeled stomatal conductance was high during the whole day on 24 June, low radiation and VPD kept the transpiration rates small. In this extreme case, the transpiration during the cloudy day (24 June) was higher than the sap flow only during one short period around midday, which means that storage water was used only during this short period and that during the rest of the day storage recharge was taking place.

3.3 Effects of vapour pressure deficit

To explain the residuals in the relationship between the storage water flow and incoming radiation, we compared two days (9 June and 20 June) with a similar measured sap flow and equal incoming radiation (29.3 and 28.6 MJ m² day⁻¹, respectively), but different VPD (peak values of 1.6 and 2.8 kPa, respectively). This comparison showed that atmospheric drought is a second important driver of storage water use (Fig. 8). It is clear that on 20 June (high VPD) stomatal closure in the afternoon was much stronger, which reduced transpiration and thus use of storage water. Furthermore, the refilling of the storage pool was higher on 20 June. The response of stomates to VPD has been reported by several authors (e.g. Monteith, 1995; Martin et al., 1997; Irvine et al., 1998; Oren et al., 1999a; Gao et al., 2002; Motzer et al., 2005). It is recognized as one of the most important sources of variation in stomatal conductance, but the mechanism of the response is still under debate (Bunce, 1997, 2006; Whitehead, 1998). The modeled stomatal conductance values of this study are consistent with other data for Scots pine (Stewart, 1988; Irvine et al., 1998).

The VPD effect is even better illustrated by the typical hysteresis patterns (Fig. 9). The hysteresis is caused by time shifts between the diurnal patterns of radiation, VPD, temperature, transpiration and sap flow. Analysis of these patterns gives a better insight into the interactions between climate and the diurnal patterns of water flow in the

BGD

4, 615–650, 2007

Drivers of storage water use in Scots pine

H. Verbeeck et al.

Title Page

Abstract

Introduction

Conclusions

References

Tables

Figures

◀

▶

◀

▶

Back

Close

Full Screen / Esc

Printer-friendly Version

Interactive Discussion

EGU

tree.

The relation between water flow and VPD shows a clockwise hysteresis. On days with moderate VPDs, there is a typical plateau at VPDs close to 1.5 kPa (Fig. 9iii), while on days with extreme high VPD (Fig. 9iv) transpiration even declined at VPDs above 1.5 kPa. On 9 June (Fig. 9iii) transpiration increased until 14 h but on 20 June (Fig. 9iv) transpiration started decreasing at 10 h due to the high VPD values that were already reached. Due to the buffering effect of the storage water, this decrease was not observed in the sap flow measurements.

The widths of the hysteresis loops between transpiration and VPD were larger than the widths of the loop of sap flow versus VPD (Fig. 8). This typical pattern of clockwise hysteresis has been found by several authors (Meinzer et al., 1997, 1999; Ewers et al., 2005; Motzer et al., 2005).

Plotting sap flow versus solar radiation typically produces a counter-clockwise hysteresis (Meinzer et al., 1997; Ewers et al., 2005). We also observed this on days with moderate VPD. However in case of high VPD, we found a clockwise hysteresis (Fig. 9ii). The decrease of transpiration starting around 10 h in the morning caused this different pattern. In contrast, during the sunny day with low VPD (Fig. 9i), transpiration increased until 14 h. This illustrates the high impact of VPD on the diurnal pattern of water flow within a tree.

The large discrepancy between sap flow at the stem base and transpiration at the leaf level indicates that sap flow measurements should not be used to calculate diurnal patterns of canopy conductance by inverting the Penman-Monteith equation without taking the possible time-lag into account. Sap flow can only be used as a measure for diurnal transpiration if the measurements are corrected for the time lag with transpiration. A commonly used method to account for the time-lag is to artificially shift the sap flow time series to maximize correlation with VPD or radiation (Oren et al., 1999b; Ford et al., 2005).

Contrary to the expectation that trees would rely more on their storage water in periods of atmospheric drought, we found in this study that the storage water use did not

BGD

4, 615–650, 2007

Drivers of storage water use in Scots pine

H. Verbeeck et al.

Title Page

Abstract

Introduction

Conclusions

References

Tables

Figures

◀

▶

◀

▶

Back

Close

Full Screen / Esc

Printer-friendly Version

Interactive Discussion

EGU

increase with atmospheric drought. However, in conditions of high soil water availability, storage water use was mainly driven by transpiration. Therefore, the hypothesis of Phillips et al. (2003) that storage water use increases with drought, only holds for soil water stress and not for atmospheric drought. According to our results, atmospheric drought did not induce an extra use of storage water. The use of storage water is not an extra advantage for trees in overcoming atmospheric drought in well-watered soils. Nonetheless, storage water contributes substantially to daily transpiration, irrespective of the climatic conditions. These results confirm that stored water plays an important role not only during periods of drought, but whenever water transport occurs within the tree (Zweifel et al., 2001). These last mentioned authors found that even in well-watered soils the tree water reserves were depleted every morning.

3.4 Constraints by tree water content deficit

Due to the use and recharge of stored water, the internal tree water content (W) fluctuated daily. The daily minimum tree water content was clearly determined by daily solar radiation and by daily maximum VPD (e.g. for pine #23 in Fig. 10). The total observed amount of stored water available for transpiration (AW_{\max}) varied among trees from 2 to 64 kg. The available storage water ranged between 2% and 36% of the total tree water content (Table 2).

Besides the diurnal fluctuations of tree water content, there was also a considerable day-to-day variation in mean tree water content, due to the net daily storage water use or recharge. When the relative contribution of storage water use to transpiration was plotted against daily mean values of the water deficit in the tree (WD) (Fig. 11), it becomes clear that the tree water content put a constraint on the maximum possible contribution to the daily transpiration. In other words: the increase of transpiration was higher than the increase of storage water use with tree water deficit.

On the other hand, the absolute amount of storage water use continues to increase with increasing tree water deficit, until a maximum is reached (Fig. 12). Beyond this threshold of tree water deficit ($WD_{\text{threshold}}$) a further depletion of the storage pool is

BGD

4, 615–650, 2007

Drivers of storage water use in Scots pine

H. Verbeeck et al.

Title Page

Abstract

Introduction

Conclusions

References

Tables

Figures

◀

▶

◀

▶

Back

Close

Full Screen / Esc

Printer-friendly Version

Interactive Discussion

EGU

more difficult due to the low water potential in the storage compartment, resulting in a lower storage water use. The threshold for tree water deficit varied considerably among the 14 individual trees, ranging from 1% to 21% (Table 2).

4 Conclusions

Relations between water fluxes, storage water use and climatic drivers were analysed in this study. As expected, solar radiation is the main driver of transpiration, and thus also of sap flow and storage water use. Nevertheless the day-to-day patterns of measured sap flow and storage water use were quite different. VPD was the main driver of these differences. In conditions of high soil water availability and atmospheric drought, trees did not rely more on their storage water. In contrast, when VPD was high, closing stomata caused low transpiration and low storage water use.

A third important factor was the tree water deficit. When storage compartments were depleted beyond a threshold, storage water use became limited due to low water potential in the storage compartments. Also the maximum relative contribution of storage water to daily transpiration was constrained by an increasing tree water deficit.

The next step in the development of the model will be the analysis of effects of soil water stress on storage water use. Maintenance of considerable transpiration rates after soil drying often depends on the availability of water in deeper soil layers, but is also hypothesized to be a result from contributions of storage water to transpiration (Sinclair et al., 2005). Future analysis with the ANAFORE model might enable us to test this hypothesis.

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BGD

4, 615–650, 2007

Drivers of storage water use in Scots pine

H. Verbeeck et al.

Title Page

Abstract

Introduction

Conclusions

References

Tables

Figures

◀

▶

◀

▶

Back

Close

Full Screen / Esc

Printer-friendly Version

Interactive Discussion

EGU

References

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BGD

4, 615–650, 2007

Drivers of storage water use in Scots pine

H. Verbeeck et al.

Title Page

Abstract

Introduction

Conclusions

References

Tables

Figures

◀

▶

◀

▶

Back

Close

Full Screen / Esc

Printer-friendly Version

Interactive Discussion

EGU

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BGD

4, 615–650, 2007

**Drivers of storage
water use in Scots
pine**

H. Verbeeck et al.

Title Page

Abstract

Introduction

Conclusions

References

Tables

Figures

◀

▶

◀

▶

Back

Close

Full Screen / Esc

Printer-friendly Version

Interactive Discussion

EGU

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BGD

4, 615–650, 2007

Drivers of storage water use in Scots pine

H. Verbeeck et al.

Title Page

Abstract

Introduction

Conclusions

References

Tables

Figures

◀

▶

◀

▶

Back

Close

Full Screen / Esc

Printer-friendly Version

Interactive Discussion

EGU

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BGD

4, 615–650, 2007

Drivers of storage water use in Scots pine

H. Verbeeck et al.

Title Page

Abstract

Introduction

Conclusions

References

Tables

Figures

◀

▶

◀

▶

Back

Close

Full Screen / Esc

Printer-friendly Version

Interactive Discussion

EGU

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BGD

4, 615–650, 2007

**Drivers of storage
water use in Scots
pine**

H. Verbeeck et al.

Title Page

Abstract

Introduction

Conclusions

References

Tables

Figures

◀

▶

◀

▶

Back

Close

Full Screen / Esc

Printer-friendly Version

Interactive Discussion

EGU

Drivers of storage water use in Scots pine

H. Verbeeck et al.

Table 1. Main attributes and maximum storage water use of the 14 sampled Scots pine (*Pinus sylvestris* L.) trees in the year 2000 at the experimental plot in Brasschaat, Belgium. All values are reported for individual trees. Diameter at breast height (DBH), height (H), sapwood area (SA), leaf area (LA), leaf-to-sapwood area (LA/SA), maximum daily storage water use (STO_{max}), maximum contribution of storage water to daily transpiration (CON_{max}). STO_{max} and CON_{max} are maximum values for each tree over the entire studied period (May 27–October 18, 2000).

Tree #	DBH (cm)	H (m)	SA (cm ²)	LA (m ²)	LA/SA (m ² cm ⁻²)	STO _{max} (kg day ⁻¹)	CON _{max} (%)
11	36.0	18.2	564	90.5	0.16	11.9	36.7
12	32.5	18.1	494	75.1	0.15	12.9	35.2
13	31.2	19.0	511	69.3	0.14	14.6	44.2
14	29.9	16.3	428	65.3	0.15	13.4	43.1
15	27.9	15.2	393	56.7	0.14	10.3	39.7
16	26.7	15.9	320	53.7	0.17	8.8	39.0
17	30.6	15.5	430	66.1	0.15	13.1	44.3
21	34.4	23.1	563	82.7	0.15	12.8	41.5
22	32.2	22.0	424	74.3	0.18	11.2	43.1
23	29.6	16.7	425	63.3	0.15	7.7	30.2
24	28.2	19.2	423	58.6	0.14	5.3	30.1
25	26.7	19.9	420	53.3	0.13	6.0	32.0
26	23.7	16.0	297	41.9	0.14	4.6	28.4
27	27.7	18.4	434	57.1	0.13	11.7	44.6

Title Page

Abstract Introduction

Conclusions References

Tables Figures

◀ ▶

◀ ▶

Back Close

Full Screen / Esc

Printer-friendly Version

Interactive Discussion

Drivers of storage water use in Scots pine

H. Verbeeck et al.

Table 2. Water content and availability of the 14 sampled Scots pine (*Pinus sylvestris* L.) trees in the year 2000 at the experimental plot in Brasschaat, Belgium. All values are reported for individual trees. Diameter at breast height (DBH), maximum tree water content (W_{\max}), maximum available tree water content (AW_{\max}) ($= W_{\max}$ – minimum W over the entire studied period), tree water content deficit threshold (WD).

Tree #	DBH (cm)	W_{\max} (kg)	AW_{\max} (kg)	AW_{\max} (% of W_{\max})	$WD_{\text{threshold}}$ (% of W_{\max})
11	36.0	271	63	23.2	14.8
12	32.5	214	44	20.6	13.8
13	31.2	213	64	30.0	10.9
14	29.9	171	54	31.6	18.2
15	27.9	146	52	35.6	20.7
16	26.7	114	32	28.1	15.8
17	30.6	175	56	32.0	11.7
21	34.4	258	53	20.5	7.7
22	32.2	182	42	23.1	8.4
23	29.6	168	30	18.9	9.9
24	28.2	159	18	11.3	6.6
25	26.7	150	25	16.7	9.6
26	23.7	94	2	2.1	1.1
27	27.7	160	57	35.6	20.6

Title Page

Abstract Introduction

Conclusions References

Tables Figures

◀ ▶

◀ ▶

Back Close

Full Screen / Esc

Printer-friendly Version

Interactive Discussion

Drivers of storage water use in Scots pine

H. Verbeeck et al.

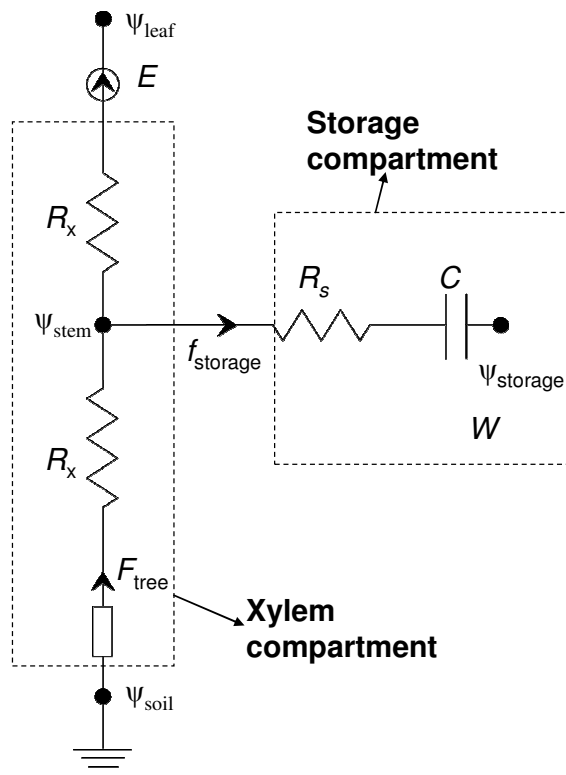


Fig. 1. Electrical analogon diagram of the water storage model. Xylem and storage compartments are shown. ψ_{soil} : soil water potential, ψ_{stem} : water potential of the xylem in the stem, ψ_{leaf} : leaf water potential, $\psi_{storage}$: water potential in the storage compartment, $f_{storage}$: water exchange between the xylem and the storage compartment, F_{tree} : water flow in the stem, R_s : exchange resistance between the xylem and the storage compartment, R_x : flow resistance of the xylem, C : capacitance of the storage compartment, E : transpiration rate, W : water content of the storage compartment.

Title Page

Abstract

Introduction

Conclusions

References

Tables

Figures

◀

▶

◀

▶

Back

Close

Full Screen / Esc

Printer-friendly Version

Interactive Discussion

Drivers of storage water use in Scots pine

H. Verbeeck et al.

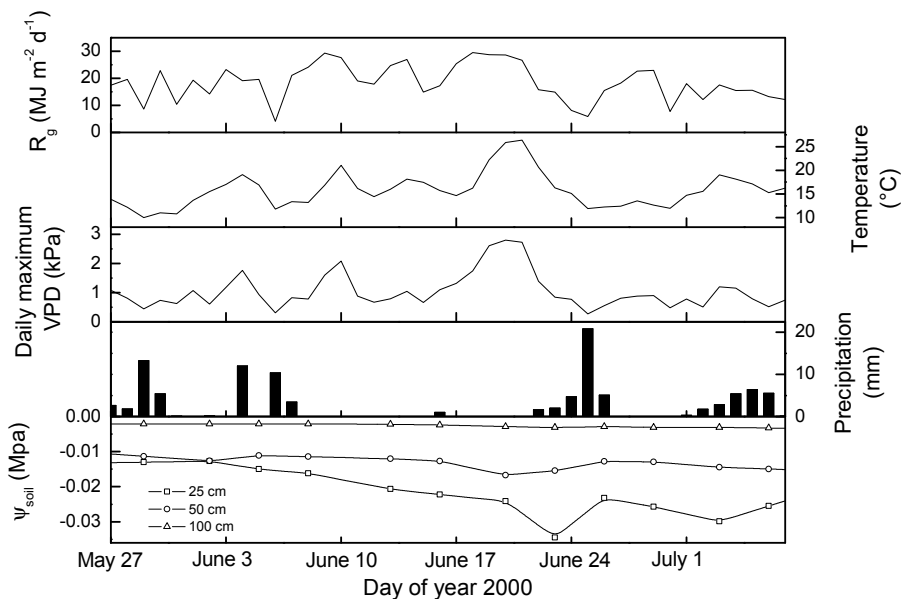


Fig. 2. Climatic conditions during the studied period (27 May–7 July 2000): daily incoming solar radiation (R_g), mean daily temperature, daily maximum vapour pressure deficit (VPD), precipitation and soil water potential (ψ_{soil}).

Title Page

Abstract

Introduction

Conclusions

References

Tables

Figures

◀

▶

◀

▶

Back

Close

Full Screen / Esc

Printer-friendly Version

Interactive Discussion

Drivers of storage water use in Scots pine

H. Verbeeck et al.

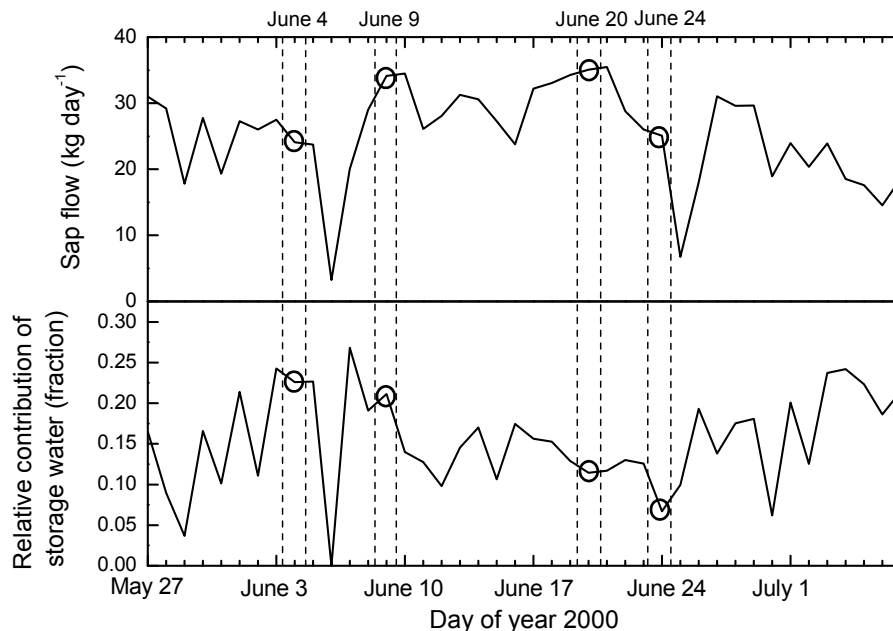


Fig. 3. Day to day variation of the measured sap flow of pine #23 and the relative contribution of storage water use to daily transpiration. Four typical days were selected (4 June, 9 June, 20 June, 24 June).

Title Page

Abstract

Introduction

Conclusions

References

Tables

Figures

◀

▶

◀

▶

Back

Close

Full Screen / Esc

Printer-friendly Version

Interactive Discussion

Drivers of storage water use in Scots pine

H. Verbeeck et al.

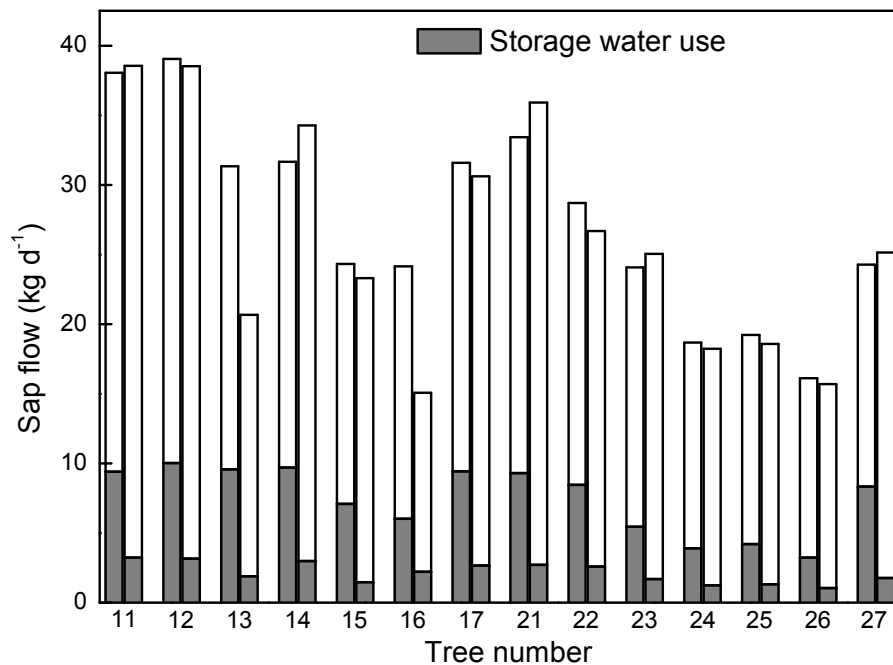


Fig. 4. Comparison of measured sap flow and storage water use between 4 June and 24 June for all 14 pine trees.

Title Page

Abstract Introduction

Conclusions References

Tables Figures

◀ ▶

◀ ▶

Back Close

Full Screen / Esc

Printer-friendly Version

Interactive Discussion

Drivers of storage water use in Scots pine

H. Verbeeck et al.

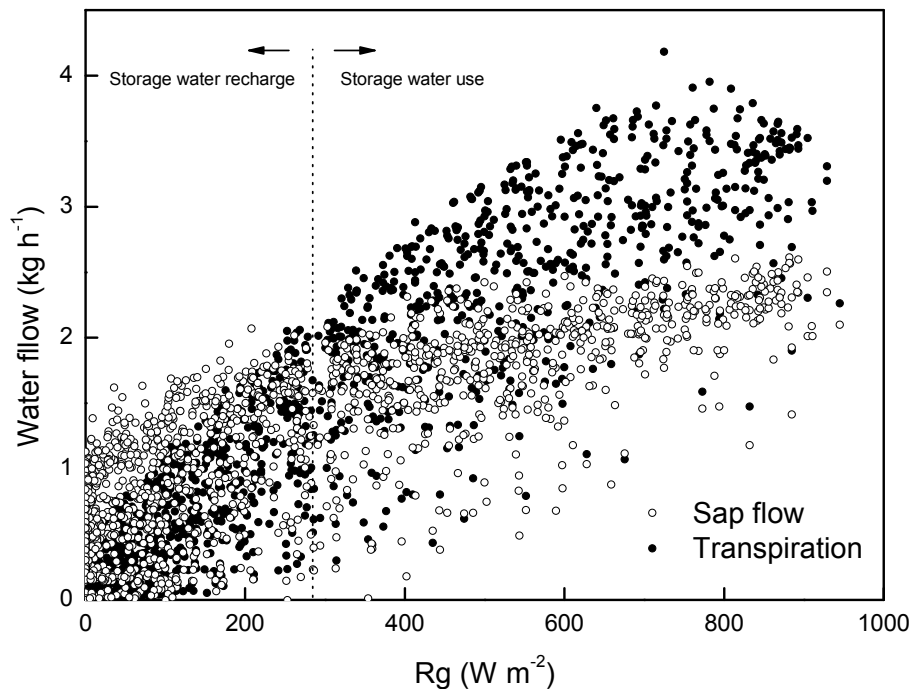


Fig. 5. Half-hourly values of measured sap flow (\circ) and modeled transpiration (\bullet) versus incoming solar radiation (R_g) for pine #23 (27 May–18 October 2000).

[Title Page](#)[Abstract](#)[Introduction](#)[Conclusions](#)[References](#)[Tables](#)[Figures](#)[◀](#)[▶](#)[◀](#)[▶](#)[Back](#)[Close](#)[Full Screen / Esc](#)[Printer-friendly Version](#)[Interactive Discussion](#)

Drivers of storage
water use in Scots
pine

H. Verbeeck et al.

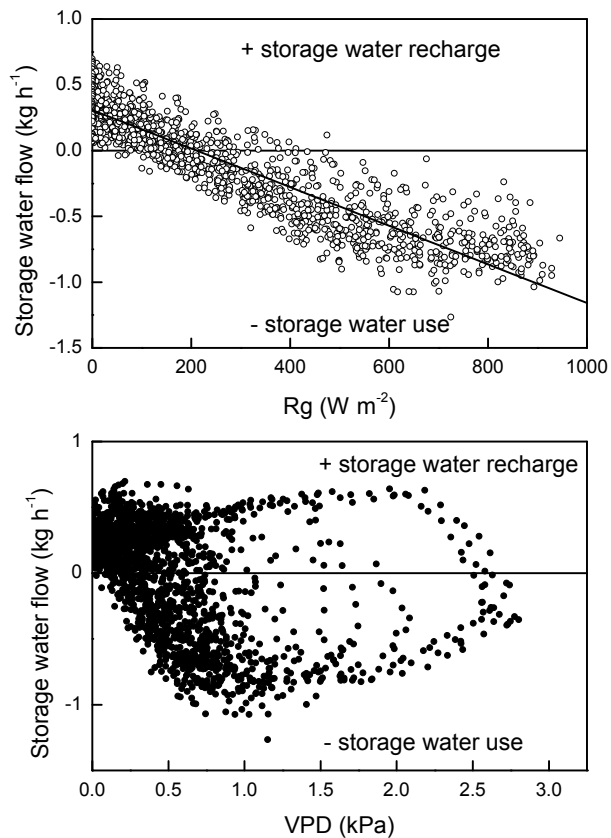


Fig. 6. Half-hourly values of modeled storage water flow versus (top) incoming solar radiation (R_g) and (bottom panel) vapour pressure deficit (VPD), for pine #23 (27 May–18 October 2000).

[Title Page](#)[Abstract](#)[Introduction](#)[Conclusions](#)[References](#)[Tables](#)[Figures](#)[◀](#)[▶](#)[◀](#)[▶](#)[Back](#)[Close](#)[Full Screen / Esc](#)[Printer-friendly Version](#)[Interactive Discussion](#)

Drivers of storage water use in Scots pine

H. Verbeeck et al.

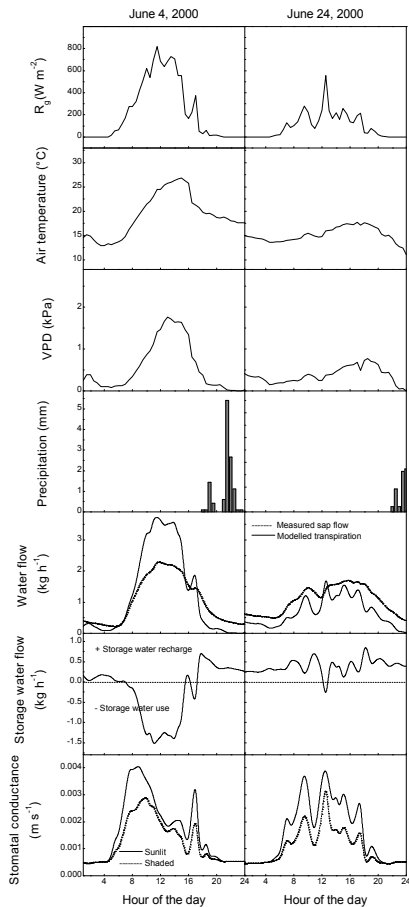


Fig. 7. Comparison of diurnal patterns during two days (4 June and 24 June 2000): climate, water flow and modeled stomatal conductance (g_s) of pine #23. (R_g = incoming solar radiation; VPD = vapour pressure deficit).

Title Page

Abstract

Introduction

Conclusions

References

Tables

Figures

◀

▶

◀

▶

Back

Close

Full Screen / Esc

Printer-friendly Version

Interactive Discussion

Drivers of storage water use in Scots pine

H. Verbeeck et al.

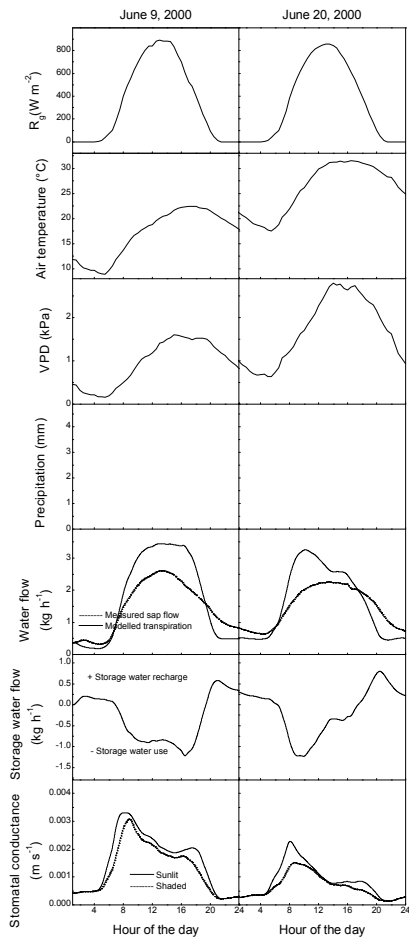


Fig. 8. Comparison of diurnal patterns during two days (9 June and 20 June 2000): climate, water flow and modeled stomatal conductance (g_s) of pine #23. (R_g = incoming solar radiation; VPD = vapour pressure deficit).

Title Page

Abstract

Introduction

Conclusions

References

Tables

Figures

◀

▶

◀

▶

Back

Close

Full Screen / Esc

Printer-friendly Version

Interactive Discussion

Drivers of storage water use in Scots pine

H. Verbeeck et al.

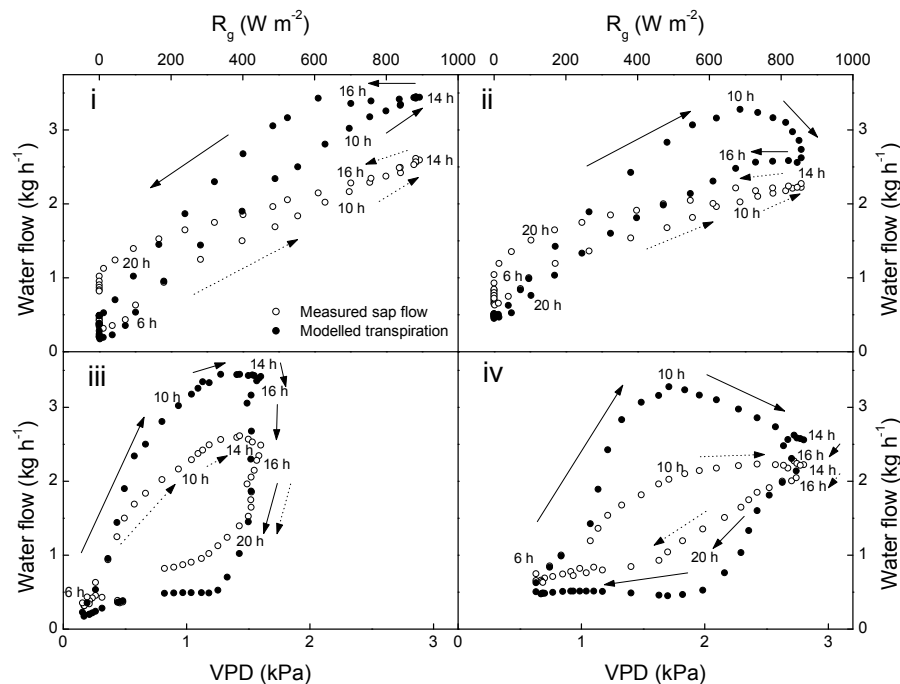


Fig. 9. Hysteresis patterns of sap flow and transpiration versus solar radiation (R_g) (i and ii) and versus vapour pressure deficit (VPD) (iii and iv) for pine #23 on 9 June and 20 June (2000), respectively.

Title Page

Abstract

Introduction

Conclusions

References

Tables

Figures

◀

▶

◀

▶

Back

Close

Full Screen / Esc

Printer-friendly Version

Interactive Discussion

Drivers of storage water use in Scots pine

H. Verbeeck et al.

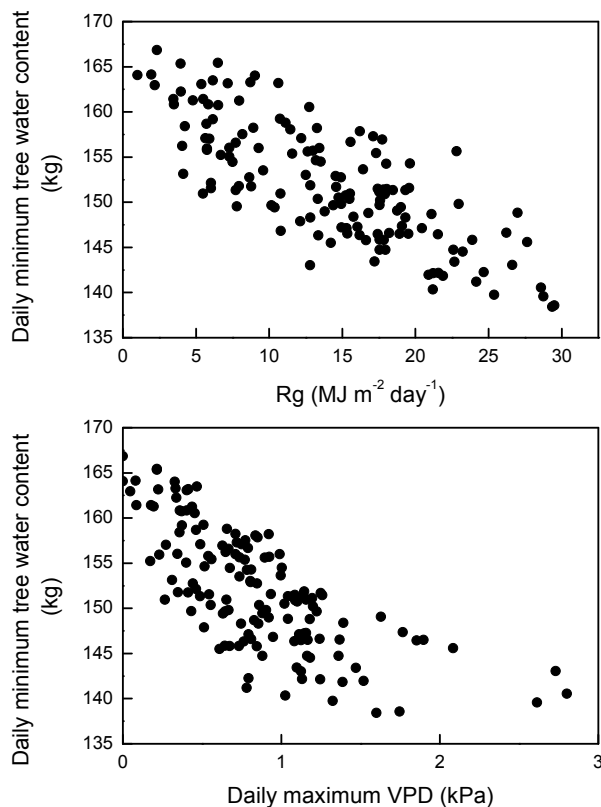


Fig. 10. Daily minimum values of tree water content plotted versus incoming solar radiation (R_g) (top) and versus daily maximum vapour pressure deficit (VPD) (bottom panel). Data from pine #23 for the period 27 May–18 October 2000.

Title Page

Abstract

Introduction

Conclusions

References

Tables

Figures

◀

▶

◀

▶

Back

Close

Full Screen / Esc

Printer-friendly Version

Interactive Discussion

Drivers of storage water use in Scots pine

H. Verbeeck et al.

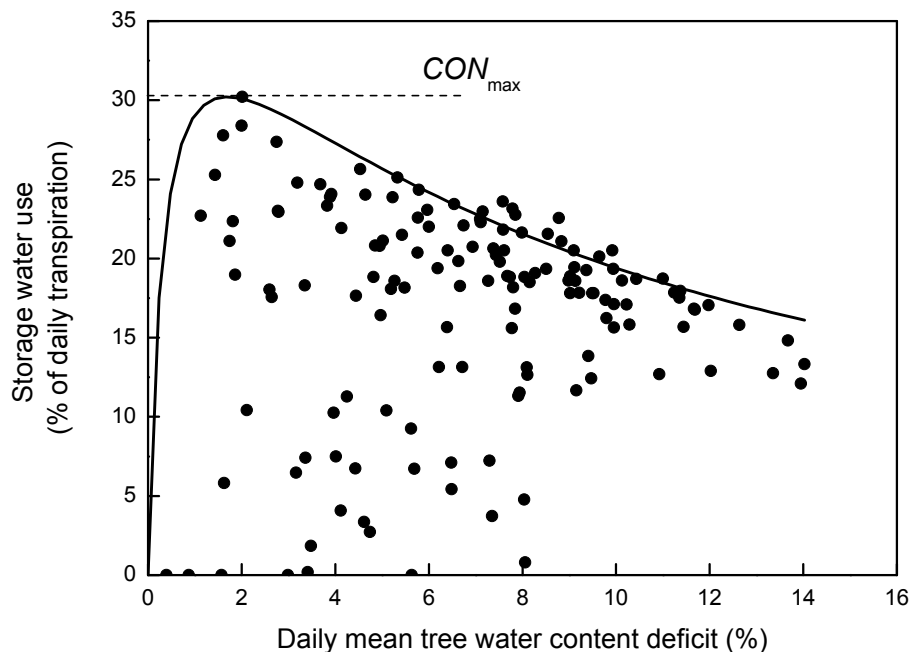


Fig. 11. Contribution of storage water use to daily transpiration versus mean daily tree water deficit (%). The data were divided in categories per 0.5 % tree water deficit. The curve was fitted through the maxima of each category. Data from pine #23 for the period 27 May–18 October, 2000. CON_{max} = maximum contribution of storage water to transpiration.

Title Page

Abstract

Introduction

Conclusions

References

Tables

Figures

◀

▶

◀

▶

Back

Close

Full Screen / Esc

Printer-friendly Version

Interactive Discussion

Drivers of storage water use in Scots pine

H. Verbeeck et al.

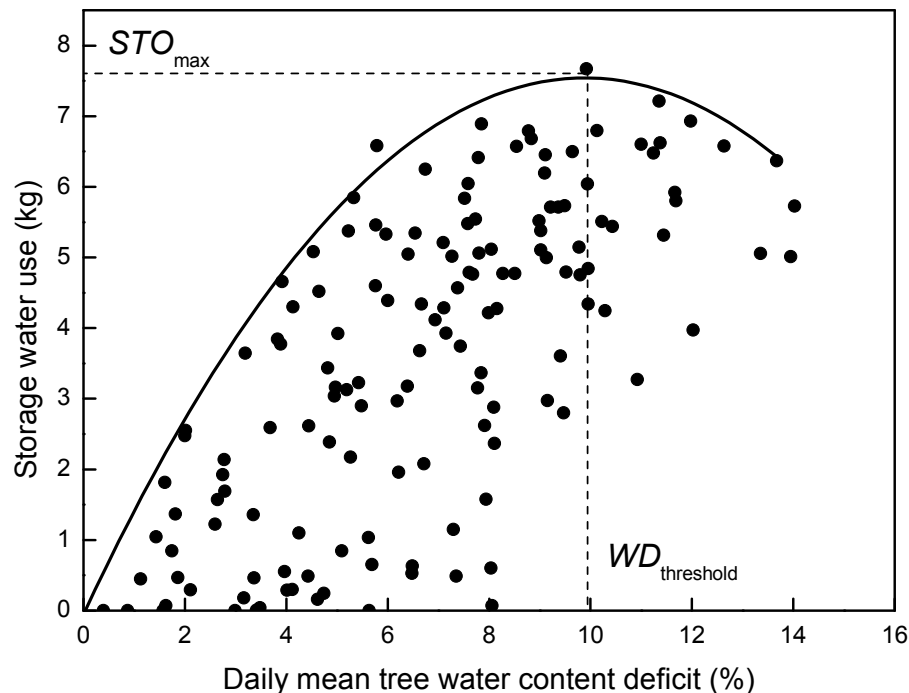


Fig. 12. Storage water use (kg) versus mean daily tree water deficit (%). The data were divided in categories per 0.5 % tree water deficit. The curve was fitted through the maxima of each category. Data from pine #23 for the period 27 May–18 October 2000. STO_{max} = maximum daily storage water use; $WD_{threshold}$ = tree water content deficit threshold.

Title Page

Abstract

Introduction

Conclusions

References

Tables

Figures

◀

▶

◀

▶

Back

Close

Full Screen / Esc

Printer-friendly Version

Interactive Discussion