# CASSIA+ Instruction and Plans Booklet

## Documentation Author: Joanna Simms

# Contents

1	$\mathbf{W}\mathbf{h}$	hat is the CASSIA model?				
	1.1	Introduction	2			
		1.1.1 Coding plan	2			
		1.1.2 Git and GitHub resources	2			
2	Tut	orials	3			
	2.1	Function List	3			
3	CAS	SSIA	6			
	3.1	Basic Model	6			
	3.2	Sugar Model	7			
		3.2.1 Allocation of Sugar Within the Tree	8			
		3.2.2 Allocation to Mycorrhiza / Soil	9			
		- · · · · · · · · · · · · · · · · · · ·	10			
	3.3		12			
	3.4	Photosynthesis	12			
4	Res	spiration	13			
5	Soil	Functions	14			
J	5.1		15			
	0.1		15			
		G I	16			
		*	16			
		1	18			
		·	19			
	5.2		$\frac{10}{20}$			
	0.2	<u> </u>	$\frac{20}{21}$			
			$\frac{21}{21}$			
	5.3		$\frac{21}{21}$			
	0.0		$\frac{21}{21}$			
		v	$\frac{21}{23}$			
	5.4		26 26			

7	App	pexdix: Parameter Values in Code	32
	7.1	repo_p Explanation	32
	7.2	common_p Explanation	33
	7.3	ratio_p Explanation	34
	7.4	parameters_p Explanation	35
	7.5	sperling_p Explanation	41

### 1 What is the CASSIA model?

#### 1.1 Introduction

The package is available from GitHub: https://github.com/josimms/CASSIA

CASSIA model is an intra-annual growth model for an individual tree in boreal conditions. Seasonal organ level cell growth is modelled, as well as sugar and water when the appropriate settings are chosen. The main mathematical structure and equations are found in Schiestl-Aalto et al. [2015] where the science behind this model as well as the basic principle and structure are clearly explained. Further details for the individual functions can be found by writing ?function\_name in the terminal in R, although a list of functions included in this package can be found in Section 2.1. And tutorials are provided Section 2.

This document and package includes newer developments not yet published in papers such as a sugar internal allocation model and more detailed xylogenesis process. If there are questions about these sections, please get in contact with the people involved in making this package.

### 1.1.1 Coding plan

The model was originially written in R and then translated to C++. The C++ version of the model changes the running time of the model from 0.397 seconds to 0.006, so is useful for calibrations, but not all subfunctions have been translated. Currently the basic model and the sugar allocation model are the ones that are working. Xylogenesis and water functions will soon be added (email Joanna if needed quicker). In 2024-2025 a soil, mycorrhizal growth and photosynthesis model are being integrated.

All C++ functions should be accessible by an R interface.

#### 1.1.2 Git and GitHub resources

If you are unfamiliar with GitHub then these are some helpful resources. If you would like to develop the model then using GitHub would be ideal, if you just want to use the model then you just need to know how to install the model from GitHub.

- GitHub: https://docs.github.com/en
- GitHub: https://docs.github.com/en/get-started/quickstart/hello-world
- RStudio: https://education.rstudio.com/learn/beginner/
- R: https://www.codecademy.com/learn/learn-r
- C++: https://www.codecademy.com/learn/learn-c-plus-plus

# 2 Tutorials

To be published: 09.05.2025 with documentation accessible from the code to also be updated around this time.

## 2.1 Function List

Table 1: Functions included in the CASSIA package grouped by function.

Purpose	Documented
Tests	
Tests the new CPP CASSIA function against the original CASSIA outputs (Pauliina's original R version).	
TODO	
Output dataset. Creates dataframes for the CAS-SIA model outputs, this isn't used yet.	N
Input test. Check that the settings are defined in a way that makes sense for the model. As not all model settings work simultaneously.	N
Input test. Function to check the parameters are from a site that is defined in the model. If a new site should be added contact Joanna.	N
Input test. The weather data should be within allowable bounds, these bounds are checked in this function	N
Plots	
1	
	Tests the new CPP CASSIA function against the original CASSIA outputs (Pauliina's original R version). TODO  Output dataset. Creates dataframes for the CASSIA model outputs, this isn't used yet.  Input test. Check that the settings are defined in a way that makes sense for the model. As not all model settings work simultaneously.  Input test. Function to check the parameters are from a site that is defined in the model. If a new site should be added contact Joanna.  Input test. The weather data should be within allowable bounds, these bounds are checked in this function  Plots

Table 1 – continued from previous page

Function	Purpose	Documente
respiration_plot	Plot to test the cpp respiration formulation	N
xylem_plot	Plot to test the cpp xylogenesis_wrapper	N
	C++ Model Functions	
CASSIA_cpp	The current CASSIA function using the cpp code	Y
01100111-0РР	and the sugar and soil options. TODO: check	-
CASSIA_calibration	TODO?	
CASSIA_soil	TODO?	
CASSIA_yearly	TODO?	
Fungal_N_Uptake		
Microbe_Uptake		
myco_decision		
myco_growth		
mycofon_balence		
plant_decision		
Plant_N_Uptake		
uptake_C		
uptake_N		
symphony		
symphony_plus		
symphony_plus_daily		
xylogenesis_wrapper		
	R Model Functions	
PRELES_GPP	R formulation of the PRELES function with the	Y
	growth_photo_coef definition. Needs to be kept	
	for the R version of the CASSIA model	
repola	R formulation of the repola function. Needs to be	Y
	kept for the R version of the CASSIA model.	
	Weather Processing	
calculate_VPD	This calculates the VPD from the temperature	Y
	and the temperature at dew point. Needed for	-
	the ERA5 data.	
generating max values monthly	y Generates the monthly max values for the specific	N
generaling_max_varaes_month,	weather data	11
generating_mean_values_daily	Generates the daily mean values for the specific	N
generating_mean_varues_dairy	weather data	11
generating mean values month	lyGenerates the monthly mean values for the spe-	N
Source and The Control of the Contro	cific weather data	11
concreting gum values monthly	Generates the monthly sum values for the specific	N
generaling sum vames momini		4.1

Table 1 – continued from previous page

Function	Purpose	Documente
$\operatorname{downloading\_data}$	Downloading the SMEAR weather data for Hyytiälä [TODO: reference]	Y
Hyytiala_Data_Creation	,	
process_files		
process_files_co2		
$process\_weather\_data$		
nporting_hyytiala_raw_data_in	to_list	
raw_to_daily_monthly_hyytiala		
binddatatable		
ead_and_combine_weather_date	a	
reading_nc		
replace_value_DataFrame		
simplifying_names	Control of the contro	
sort_CMIP	Sorts the pre-downloaded CMIP6 data for Hyytiälä.	
	Data sets	
lata_format	The best Hyytiälä formatting	
oad_data	Function to load all of the Hyytiälä data gathered,	
	mainly for calibrations and comparisons	
byewole_2015_calibration_data weather_original		
	Parameters	
common_p	The parameters needed for the common function are saved as a RData file so can be automatically accessed by the function	
parameters_p		
ratios_p		
repo_p		
sperling_p		
	Calibration	
dhtn	An alternative to the normal distribution for the boreal calibration	N
nitrogen_uptake_calibration		
ikelyhood_nitrogen_uptake	Likelihood function for the nitrogen uptake function.	N
	Likelihood function for the soil models.	N
ikelyhood_soil_models		
ikelyhood_soil_models ikelyhood_sugar_model	Likelihood function for the sugar model.	N

Table 1 – continued from previous page

Function	Purpose	Documented
common		
daily_list		
daily_to_average		
envfac.data		
GPP_2009		
GPP_previous_sum		
$GPP\_ref$		
growth_wrapper		
Hyde_daily_original		
Hyde_weather		
Hyde_yearly_original		
initialize_parameters		
$karike\_df\_all$		
leap_years		
nitrogen		
parameters_R		
printColumnNames		
ratios		
soil_model_calibration		
$transform\_variable$		

# 3 CASSIA

### 3.1 Basic Model

### Model Lowdown: CASSIA.

Model type: Carbon based growth model

Inputs: Temperature (air, soil), soil water content, photosynthesis

**Scope:** Tree level (with categorisation and cell level growth)

Timestep: Daily

Reference: Schiestl-Aalto et al. [2015, 2019]

NOTE: Please check the references [Schiestl-Aalto et al., 2015, 2019] to understand the full structure of the CASSIA model - equations will be added here when water dependencies are added (again email Joanna if these are needed sooner). But the most important aspect is the carbon sink and source effect on seasonal growth:

## Actual Growth =Poteital Gorwth Dependant on Environmental Conditionsmin [Sugar limitation, Nitrogen Limitation]

Note that the sugar limitation is the original formulation. The nitrogen limitation is currently being added to the system.

#### 3.2 Sugar Model

CASSIA includes an organ level internal sugar model based on the logic from Sperling et al. [2019], which predicts the bloom dates of almond trees by considering that bloom happens when the sugar level drops beyond a certain threshold. The sugar level is controlled by enzymes that convert sugar to starch. The enzyme activity is affected by the amount of enzymes and temperature. Plant production of enzymes is also generated by the difference between current sugar level and the "expected" sugar level (equilibrium point).

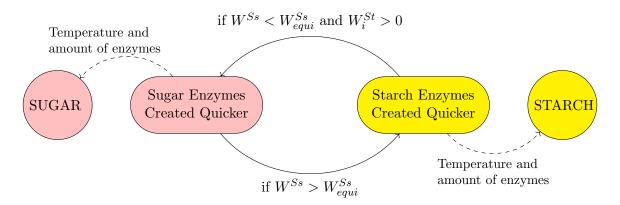


Figure 1: Figure to show the Sperling model process.

This is mathematically represented by;

$$\begin{split} W_{i}^{Ss} > W_{equi,i}^{Ss} : & (1) \\ \begin{cases} \frac{dD_{i}^{Ss}}{dt} = \lambda_{i}D_{i}^{Ss} + \delta_{i} \\ \frac{dD_{i}^{St}}{dt} = \lambda_{i}D_{i}^{St} \end{cases} \\ W_{i}^{Ss} < W_{equi,i}^{Ss} & \text{and } W_{i}^{St} > 0 : \\ \begin{cases} \frac{dD_{i}^{Ss}}{dt} = \lambda_{i}D_{i}^{Ss} \\ \frac{dD_{i}^{Sd}}{dt} = \lambda_{i}D_{i}^{St} + \delta_{i} \end{cases} \\ K_{i}^{St}(T, D^{St}) = D^{St}e^{F^{St}T} & (2) \\ K_{i}^{Ss}(T, D^{Ss}) = D^{Ss}e^{F^{Ss}T} & (3) \\ Q_{i}(W^{Ss}) = 0.004211\rho_{i}(-K_{i}^{St}(T, A_{i}^{St}) + K_{i}^{Ss}(T, A_{i}^{Ss})) & (4) \end{cases} \end{split}$$

Where W is the amount of sugar, Q is the transfer of sugar to starch, Ss is sugar and St is starch, K is the synthesis of sugar or starch dependent on temperature and amount of relevant enzymes, D is the amount of enzymes,  $\delta$ ,  $\lambda$ , F are all parameters that control the enzymatic behaviour and amount, T is temperature and  $\rho$  is the organ density. i determines the organ. Emergency supplies of sugar – when the organ's sugar concentration is less than a threshold  $L_i$  – is from the starch storage and is released by  $E(W_i^{Ss}, W_i^{St})$  seen in (5).

$$E\left(W^{Ss}, W^{St}\right) = \begin{cases} \min\left[\max\left[\frac{L_i - W_i^{Ss}}{\tau}, 0\right], W_i^{St}\right] & W_i^{St} \ge 0\\ 0 & \text{else} \end{cases}$$
 (5)

### 3.2.1 Allocation of Sugar Within the Tree

Sugar moves between the organs via a concentration-based model seen in Figure 1. Between each organ and the phloem, there is a diffusion-based relationship [Dietze et al., 2014] seen in Figure 2. The basic relationship is

$$W_{i,j}^{\text{allocation}} = \max \left[ (W_i^{Ss} + W_i^{St}) \cdot (f(W_i^{Ss} + W_i^{St}) - f(W_j^{Ss} + W_j^{St})), (W_j^{Ss} + W_j^{St}) - cB_i \right]$$
(6)

Where f is a storage term between 0 and 1 which normalizes the concentration gradient. This means that the storage limitation is the driver in the system. This is updated for each iteration, where i is the sugar origin organ, and j, is the destination organ. This is then compared to the storage capacity of each organ cB, where c is the capacity – the percentage of biomass that can be used for storage. These values are fitted with sugar organ level data from Hyytiälä. As a big debate in the field is the surplus C hypothesis of passive sugar movement compared to the active transfer, both should be considered in terms of their match for the data. Thus, when a carbohydrate conservative strategy is used we use the equation.

$$W_{i,j}^{\text{allocation}} = (W_j^{Ss} + W_j^{St}) - cB_i \tag{7}$$

More allocation strategies could be applied here, however the storage creating a gradient system is quite useful when different allocation strategies are considered in the system (3.2.2).

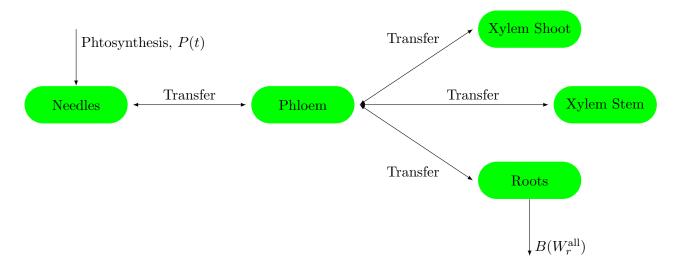


Figure 2: Figure to show the sugar transport of the model

Here  $B(W_r^{\text{all}})$  is the allocation to fungal partners or the soil which is determined in the allocation alternatives section.

G(t) is growth associated with that organ and R(t) is both growth and maintenance respiration associated with the organ. Growth and respiration are calculated to be the maximum possible growth [Schiestl-Aalto et al., 2015] and then multiplied by a scaling of the total storage of sugar and starch (f).  $\alpha$ ,  $W^{ala}$  and h are fitted parameters.

Again this f function representing the storage has been formulated in different ways to test the model behaviour. The best ones are the moment are:

$$f(W_i^{all}) = \max\left[0, \min\left[1, h_i\left(1 - \frac{1}{e^{\alpha_i\left(W_i^{all} - W_i^{ala}\right)}}\right)\right]\right]$$
(8)

$$f(W_i^{all}) = \max \left[ 0, \min \left[ 1, h_i \left( 1 - \frac{1}{e^{\alpha_i \left( W_i^{all} - W_i^{ala} \right)}} \right) \right] \right]$$

$$f(W_i^{all}) = \max \left[ 0, \frac{1}{1 + \exp(-2(\text{Sugar} + \text{Starch} - \text{Storage Capacity}))} \right]$$
(9)

#### 3.2.2 Allocation to Mycorrhiza / Soil

To understand the observed patterns we should consider a few different strategies. These strategies change the seasonality as well as the magnitude of allocation to the soil.

$$B(W_r^{all}) = \max(0.3 \cdot W_r^{all}, W_r^{all} - cB_r) \tag{10}$$

- 1. Observed: fitting the data with the sugar available considering a fixed percentage going to the mycorrhiza. This creates the desired effect in the starch. The value of this is approximately 30% of plant sugar being allocated to the soil. No nitrogen limitation.
- 2. Nitrogen feedback. However, rather than just considering the sugar leaving the system due to it fitting data could it be to harvest nitrogen? In this case consider the difference in growth, which thus effects the carbon fixed in growth and not available to be allocated below ground

- 3. Nitrogen investment. Again rather than considering this process to be passive, maybe it is active. In this case we consider the plant wanting to minimise the difference between the nitrogen and carbon limitation. Thus investing as much free carbon as it can to minimise this difference (ecoevolutionary inspired)
- 4. Strategies under development: nitrogen limitation within the model being driven by soil dynamics.

Note that the strategies above are considered due to the data available to us. Eventually the model should consider the ectomycorrizal and plant decisions more explicitly as in 5.2.

#### 3.2.3 Sugar and Starch complete process

Therefore, sugar and starch transfer is calculated via:

$$\frac{dW^{Ss}}{dt} = \begin{cases} P + Q + E - f\left[G + R\right] - W_{n,p}^{\text{allocation}} & \text{Needles} \\ Q + E - f\left[G + R\right] - W_{p,s,b,r}^{\text{allocation}} + W_{n,p}^{\text{allocation}} & \text{Phloem} \\ Q + E - f\left[G + R\right] & + W_{p,b}^{\text{allocation}} & \text{Xylem, Shoot} \\ Q + E - f\left[G + R\right] & + W_{p,s}^{\text{allocation}} & \text{Xylem, Stem} \\ Q + E - f\left[G + R\right] & + W_{p,r}^{\text{allocation}} - B(W_r^{all}) & \text{Roots} \end{cases}$$

$$\frac{dW_i^{St}}{dt} = -Q - E$$

The timing of bud burst is determined by a factor  $W_{\rm bud}^{Ss}$  which is close to the minimum possible sugar, when the total storage of sugar goes below this threshold the first day of bud growth begins. The model is run twice for each year, the first run is to find the bud date, and then the model is run again using this value for the bud date to calculate growth. These second run results are the final results of the model. The fact that the equilibrium point and the bud burst have to be defined is an obvious assumption and mean that the system isn't fully dynamic - although the point when the system would reach these hard limits is.

The symbols are defined in Table 2. When the model is calibrated the actual parameter values will be added. Both from literature and from the Bayesian calibration. When the parameters are calibrated, the uniform distribution was chosen due to a lack of prior knowledge.

Table 2: Table to define all of the symbols in the sugar model. TODO: units and values

Symbol	Function	Units (day-1)	Meaning
		•	Variable
W	Variable	kg C	Amount of of carbon in the organ indicated by the super-
			script
D	Variable	mg g-1	Amount of enzyme
T	Variable	Degrees C	Temperature, input to the model timeseries
		Proc	cess in the model
K	Process	mg g-1	Creation of sugar or starch
Q	Process	kg C	The Sperling model process amalgamated
$E(W^{Ss}, W^{St})$	Process	kg C	Emergency transfer when the sugar concentration is too low
, ,		_	if there is enough starch to send an emergency transfer of
			starch.
$f(W_i^{all})$	Multiplier	0-1	Function between 0 and 1 which scales growth based on the
			carbohydrate reserves.
$F(W_r^{all})$	Process	kg C	Allocation to mycorrhiza
R(t)	Process	kg C	Respiration is worked out as in other CASSIA papers and
,		_	includes growth and maintenance [Schiestl-Aalto et al.,
			2019]
G(t)	Process	kg C	Growth is worked out as in other CASSIA papers [Schiestl-
			Aalto et al., 2019]
			Index
Ss	Index		Sugar
St	Index		Starch
all	Index		Both sugar and starch summed
equi	Index		Equilibrium point of relevant variable derived from mea-
1			surements
n, p, s, b, r	Index		As subscripts these represent all of the organs respectively;
,1,,,,			needles, phloem, xylem shoot, xylem stem and roots.
			Parameter
λ	Parameter	1e-4	Decay rate of enzymes
δ	Parameter	25e-6	Emzymic parameter
F	Parameter		= log(Q10)/10, Q10 for sugar synthesis and 1.8 for starch.
0.004211	Parameter		Scale factor from sugar concentration in mg g <sup>-1</sup> to kg C to
			match the CASSIA units.
ρ	Parameter		density of organ
$\stackrel{r}{L}$	Parameter		Threshold for appropriate processes
$W^{ala}$	Parameter		lower bound for storage effects on growth
h	Parameter		Control of the sugar storage effects on growth
$\alpha$	Parameter		Control of the sugar storage effects on growth
$L_{r}^{M}$	Parameter	kg C	Threshold for allocation to roots
$W_{ m bud}^{Ss}$	Parameter	kg C	Threshold of sugar concentration for spring awaken / bud
bud		0 ~	burst
c	Parameter	kg C	The threshold of sugar storage in each organ fitted for each
=	_ 0.2 0.2.1.0 0.0.1		organ with the sugar data collected from Hyyitälä.
			Other
P(t)	Timeseries	kg C tree-1	Photosynthesis, timeseries input, calculated by the external
1 (6)	1 11110001100	ng C 1166-1	model or PRELES depeding on the version of the model.
			Therefore, there is no dynamic effect on the daily photo-
			synthesis input [Susiluoto et al., 2010].

#### 3.3 Growth As Investment

Currently, we have considered fluxes and allocation as the key structure. Therefore, the plant investment is in sugars, however we do see a difference in the biomass allocated as well. Currently a framework is being thought of where the objective is to investigate how the cost of different root structures affects an optimisation problem where the cost of photosynthesis is dependent on nitrogen supply.

We seek to:

Which is dependent on number of root tips, length of mycorrhized root tip and allocation of biomass to belowground infrastructure per leaf area. These traits are inspired by studies of the climate and soil fertility gradient in the hemiboreal to arctic soils [Ostonen et al., 2007], the root economic spectrum as explained by Kramer-Walter et al. [2016] and Kong et al. [2019] as well as conversations by boreal root experts.

This formulation has not been published so is not included in this documentation, but will be implemented in CASSIA when a calibration is completed. Expected Autumn 2025. If interested ask send an email to Joanna for a preview.

#### 3.4 Photosynthesis

In the original CASSIA model used SPP to calculate photosynthesis. The SPP model [TODO: reference] is first run and then the output is added into CASSIA as an input. In terms of integrated photosynthesis models PRELES [Mäkelä et al., 2008, Minunno et al., 2016] is now integrated and phydro [Joshi et al., 2022] with a nitrogen addition is being added. Preles gives ecosystem level GPP considering environmental conditions and the nitrogen multipler developed by Peltoniemi [TODO: reference] has been added to consider the entire nitrogen porcess in the model. This is explained further below. As PRELES is originally a canopy level model, the output of PRELES has been divided by the amount of trees in Hyyitälä (1010) to give an 'average tree' value. As the amount of trees changes in Hyytiälä, this could also be a variable in the future for Hyyiälä comparisons.

phydro with a nitrogen factor is currently being added to CASSIA to give an eco-evolutionary approach alternative. The nitrogen limitation added will be published in the near future and then added here.

Model Lowdown: PRELES.

**Model type:** Photosynthesis model with a water balance

**Inputs:** Climate data, fAPAR

Scope: Stand level Timestep: Daily

Reference: Mäkelä et al. [2008], Peltoniemi et al. [2015], Minunno et al. [2016]

The current PRELES structure for GPP generation is;

$$P = \beta \cdot \phi \cdot f_{APAR} \cdot f_{L} \cdot f_{S} \cdot \min(f_{D}, f_{W,P}) \cdot f_{CO_{2},P}$$
(12)

Where GPP is first defined as the maximum possible under light interception (first two terms) and then down-scaled with daily multipliers for other weather conditions such as light (L), temperature (T), VPD (D), water (W,P) and carbon dioxide (CO2). Therefore to have a nitrogen feedback effect PRELES should be modified to include a  $f_N$  multiplier. I suggest this function should be from McMURTRIE [1991]:

$$f_N(N) = \frac{\epsilon}{\epsilon_0} = \frac{\epsilon_0 \left(1 + \frac{N}{N_0}\right) - \sqrt{\left(\epsilon_0 \left(1 + \frac{N}{N_0}\right)\right)^2 - 4\theta \epsilon_0^2 \frac{N}{N_0}}}{2\theta \epsilon_0}$$
(13)

Key factors:

- To get the function to be between 0 and 1 divide  $\epsilon$  with  $\epsilon_0$  (the point at the asymptote). Currently N limits at the average needle N concentration (9.1).
- CASSIA used to generate the fAPAR as in Tian et al. [2021]
- When combining CASSIA and PRELES in this way, N is directly limiting photosynthesis, but there are no N controls in CASSIA so growth is only limited indirectly apart from diameter which directly uses GPP.

Calibration data:

- Peltoniemi
- Kainulainen and Holopainen [2002]

Table 3: McMurtrie: table of symbols

Symbol	Function in	Units	Meaning
	the system		
			Variable
$\overline{N}$	Variable	Day	Nitrogen
			Parameter
$N_0$	Parameter	mg g-1	Parameter characterising dependence of $\epsilon$ on $[N]$
$e_0$	Parameter	g MJ-1	Light utilisation coefficient at high $[N]$
θ	Parameter		Curvature parameter of relationship pf $\epsilon$ to $[N]$

# 4 Respiration

TODO: add the respiration function here from Ryhti! [Ryhti et al., 2021, 2022]

### 5 Soil Functions

The original extended model structure is shown in Figure 3. The equations are then stated afterwords according to the figure's sections. Note at the moment the Mycofon and symphony models are so altered that they don't necessarily reflect the original models.

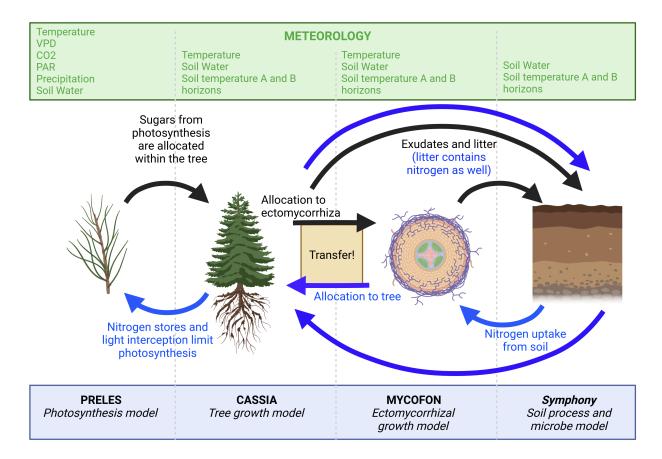


Figure 3: Figure to show the model process. Arrows show the most important inputs to the separate modules in terms of this question. Created with BioRender.com

#### **Assumption:**

- No root competition big problem in a boreal forest! Hyyitälä (the main site in this calibration / model) specifically has been shown to have intense competition [Ryhti et al., 2022]
- As in Ekblad et al. [2013] assume that the seasonality effect is produced by different parts of the model (N, C inputs and growth)
- The mycorrhiza should be reflective of one tree as there is only C input from one tree this should naturally limit the growth
- One nitrogen pool for the whole plant this nitrogen affects the photosynthetic effect, but is not specifically leaf nitrogen nor organ nitrogen for the nitroge limitation on growth

- No spacial considerations in the nitrogen patches. As mycorrhiza and plant roots are observed to create "patches" [Brandes et al., 1998], we can assume that they detect and uptake the nitrogen that is closely available to them and thus they get an average uptake of high and low nitrogen patches. The problem comes when data collecting. The value for the N:C ratio needs to be taken from lots of different locations to make sure that the average is representative of the entire area.
- All nitrogen uptake should be considered organic, nitrate and ammonium as they form a significant part of nitrogen uptake in the plant [Wallenda and Read, 1999, Näsholm et al., 2009]

#### 5.1 Uptake Functions

### 5.1.1 Nitrogen uptake

The organic uptake of N is currently controlled by considering the maximum possible uptake and then this is downscaled with environment dependent functions, taking a value between 0 and 1. These functions take the same logic, but not the same form as the PRELES equations.

$$u(N) = \frac{kN^8}{N_{limit}^8 + N^8} \tag{14}$$

$$f_T(T) = \begin{cases} \frac{T+20}{55} & T > 0\\ 0 & T \le 0 \end{cases}$$
 (15)

$$f_{SWC}(SWC) = \frac{SWC^8}{0.3^8 + SWC^8} \tag{16}$$

$$u_{actual} = f_T(T) f_{SWC}(SWC) u(N) \tag{17}$$

Where fs are the environmental modification functions and  $f \in [0, 1]$ , u() is the uptake. In equation 14, N is the concentration of a form of nitrogen,  $N_{limit}$  and k are parameters to control the uptake. Equation 15 controls the temperature response, where T is the temperature, 20 and 55 are parameters to control the uptake (non-fitted, will change with calibration). Finally the water effect is controlled by Equation 16, where SWC is soil water content and 0.3 is a parameter (will change with calibration). Equation 17 is the final nitrogen uptake effect. The parameters are dependent on the type of nitrogen.

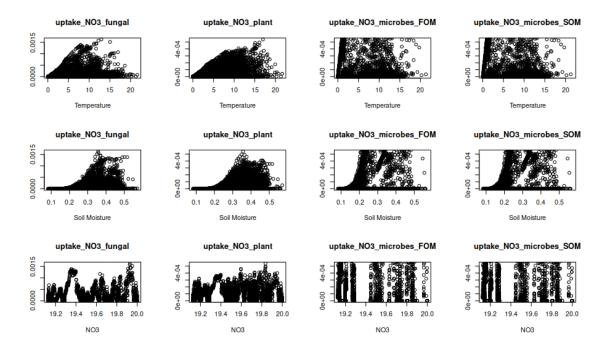


Figure 4: Figure to show the results of the uptake equation with respect to these environmental variables when the simulations are run for a few different years. Note the model is yet to be calibrated so the scales are not sensible.

#### 5.1.2Microbe Carbon Uptake

This is the carbon uptake of the mycorrhiza and other microbes in the soil. Current version of the model only has microbial uptake. This uptake is formed using the same logic as the nitrogen uptake. The carbon uptake is in the form of organic compounds, although considered in kg C in the model.

$$u(C) = \frac{kC^8}{C_{\text{limit}}^8 + C^8} \tag{18}$$

$$f_T(T) = \begin{cases} \frac{T+20}{55} & T > 0\\ 0 & T \le 0 \end{cases}$$

$$f_{SWC}(SWC) = \frac{SWC^8}{0.3^8 + SWC^8}$$
(19)

$$f_{SWC}(SWC) = \frac{SWC^8}{0.3^8 + SWC^8} \tag{20}$$

$$u_{actual} = f_T(T) f_{SWC}(SWC) u(C)$$
(21)

#### Plant N Uptake 5.1.3

NOTE: Uptake of nitrogen can also be considered as proportional to sugar exudated as well as a function of the nitrogen concentration in the soil. The following is considering a case when the soil model or at least nitrogen availability is an input.

This function is for the soil to root uptake only. This function uses the nitrogen uptake functions for each of the individual types of N compiled for the plant. The demand is calculated by the

decision function, then used to scale the maximum uptake Section 5.2.1. Although demand is currently 1 for all decision functions included until the code works better and strategies have been chosen that reflect the demand being 1 [Meyer et al., 2010, Franklin et al., 2014]. Eventually, the demand should be inversely proportional to the concentration of the desired nutrient in each organism. The nitrogen transferred to the root is a combination of all of the uptake functions, with the NO3 nitrogen uptake updated with a NH4 modifier. NH4, NO3 and FOM are the amounts in the soil. T is temperature in the soil B horizon and then SWC is the soil water content. For f(NH4), a and b are fitted parameters and the function gives a value between 0 and 1. m is the ratio of mycorrhized roots.

Original formulation:

$$f(NH4) = \frac{aNH4^8}{NH4^8 + b^8} \tag{22}$$

N to root =
$$(1-m)$$
 · demand (plant N amount) ·  $[u_{organic}(FOM, T, SWC) + u_{NH4}(NH4, T, SWC) + f(NH4)u_{NO3}(NO3, T, SWC)]$  (23)

New formulation: (to be added to code)

Allocation factor for roots = demand (plant N amount) 
$$(24)$$

N to root = 
$$(1 - m) \cdot [u_{organic}(FOM, T, SWC) + u_{NH4}(NH4, T, SWC) + f(NH4)u_{NO3}(NO3, T, SWC)]$$
 (25)

More formulations to be added!

Table 4: Uptake functions

Symbol	Function in	Units	Meaning
· ·	the system		
			Variable
$\overline{N}$	Variable	kg N	Nitrogen
NH4	Variable	kg N	NH4 pool in soil
NO3	Variable	kg N	NO3 pool in soil
FOM	Variable	kg N	FOM pool in soil
C	Variable	kg C	Carbon
T	Variable	degree C	Temperature
SWC	Variable	%	Soil water content
m	Variable	0-1	Amount of roots mycorrhized, will eventually by a moving value as in
			Equation 49
			Parameter
$N_{limit}$	Parameter	kg N	Parameter that controls when the switch in the uptake is in terms of the
			N limitations
k	Parameter		Parameter to control the nitrogen
0.3	Parameter	%	Parameter that controls when the switch in the uptake is in terms of
			SWC
20, 55	Parameter		Parameters controlling the uptake of nitrogen in terms of temperature

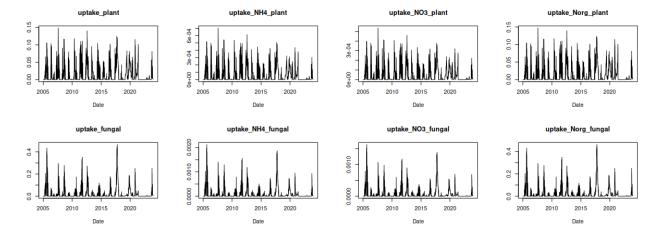


Figure 5: The uptake of nitrogen over the years 2005 to 2023 for the plant and mycorrhiza. Note the model is yet to be calibrated so the scales are not sensible. Behaviour is consistent with literature

#### **Assumptions:**

- Almost the same uptake priorities for all of the organisms (trees and mycorrhiza), similar to [Marschner et al., 1991]
- Mineral and organic uptake of N are in parallel due to the fact that when the N forms are found in a mixture the amino acid transfer doesn't decrease in proportion to the inorganic [Näsholm et al., 2009, Wallenda and Read, 1999] and as I could find relationships between  $NH_4^+$  and  $NO_3^-$  uptake [Marschner et al., 1991].
- Currently ignoring mass flow: which is bad as Oyewole et al. [2017] notes that "the results suggest that mass flow, induced by transpiration, may be a strong driver for plant nitrogen acquisition in boreal forests by delivering higher amounts of amino acids and nitrate to plant roots and mycorrhizas." The water effect on growth will be added in the next iteration of the CASSIA model so could be thought about at this stage. NOTE: Using the uptake as a reference from this paper anyway so the numbers are in the right range, but this could be a mistake if the processes are different to the ones in my model.

#### 5.1.4 Mycorrhizal N Uptake

Like the plant uptake, this function uses the nitrogen uptake functions for each of the individual types of N compiled for the mycorrhiza. The demand is calculated by the decision function, although is currently 1 for all decision functions included [Meyer et al., 2010, Franklin et al., 2014]. Then the output is the N uptake of each of the types of nitrogen calculated in parallel [Wallenda and Read, 1999, Meyer et al., 2010] and then downscaled with the demand function. Original formulation:

N to mycorrhiza = Mycorrhizal Biomass · demand (mycorhiza C amount) · 
$$[u_{organic}(FOM, T, SWC) + u_{NO3}(NO3, T, SWC)]$$
(26)

New formulation: (to be added to code)

Allocation factor for fungal = demand (mycorhiza C amount) 
$$(27)$$

N to root = Mycorrhizal Biomass 
$$\cdot [u_{organic}(FOM, T, SWC) + u_{NH4}(NH4, T, SWC) + f(NH4)u_{NO3}(NO3, T, SWC)]$$
 (28)

More formulations to be added!

Where  $u_{total}$  is the total mycorrhizal uptake.  $u_{organic}$ ,  $u_{NO3}$  and  $u_{NH4}$ , are the mycorrhizal uptake of each of the forms of nitrogen. And demand is an input to the function calculated in the ectomycorrhiza decision function 5.2.2.

### **Assumption**:

• Although in the original model there was no mycorrhizal decomposition. Decomposition is currently under development as inspired by [Lindahl et al., 2007, Baskaran et al., 2017]

### 5.1.5 Microbe Uptake

The microbe uptake holds half of the symphony model dynamics - the microbe side. Original equations from Symphony paper

Carbon Limitation 
$$=u_{\text{Norg}}C_{\text{Microbe}}$$
 (29)

Nitrogen Limitation = 
$$\frac{i(u_{\text{NH4}} + u_{\text{NO3}}) + 0.2 \cdot (\text{N:C})_{\text{Microbes, opt}} C_{\text{Microbe}}}{(\text{N:C})_{\text{Litter}} - (\text{N:C})_{\text{Microbes, opt}}}$$
(30)

Total N uptake = 
$$\max\{0, \min\{\text{Carbon Limitation}, \text{Nitrogen Limitation}\}\}\$$
 (31)

Total N uptaken =Total N uptake 
$$\cdot$$
  $\left(0.2 \cdot (\text{N:C})_{\text{Microbes, opt}} C_{\text{Microbes}} + \right)$ 

$$(N:C)_{Litter} - (N:C)_{Microbes, opt}$$
 (32)

Equations currently implemented

Total C uptaken = 
$$C_{\text{Microbes}}[u_c(\text{Relevant C compartment}) + u_c(\text{Exudes}) - (N:C)_{\text{opt}} + R(T_B)]$$
 (33)

Total 
$$N_j$$
 Uptaken =  $C_{\text{microbe}} \left( u_{N_j} - (\text{N:C})_{\text{Microbes, opt}} \right) \left( \frac{N_j \text{ Pool}}{\text{Total Nitrogen}} \right)$  (34)  
 $j \in [NH4, NO3, N_{\text{org}}]$ 

Where the uptake functions are the same as earlier, but calibrated for microbe uptake. i is immobilisation, 0.2 is a respiration parameter before the respiration is made dynamic, C, N are carbon and nitrogen respectively and (N:C) are the nitrogen carbon ratios. j represents the NH4, NO3 and organic nitrogen pools. If the SOM decomposes don't get enough nitrogen from their own pools then they can uptake more from the FOM pool, as below.

Extra FOM Uptake 
$$=u_{\text{Organic}}$$
 (35)

NOTE: the table of symbols will be added when the equations are finalised.

#### **Assumptions:**

- Carbon and nitrogen have the same units here need to be careful that they are really made equivalent in the code.
- Exudates are uptaken by the microbes and not by the mycorrhiza. This allows for a limited priming effect.
- In terms of the carbon and nitrogen limitations missing here, they should be present in the soil model namely immobilisation and growth.

#### 5.2 Decisions: transfer between organisms of N and C

Model Lowdown: Optimisation and Game Theory - currently my own model.

Model type: Decision

Inputs: Max possible transfer of C and N

**Scope:** Just the transfer amounts of C and N between the organisms

Timestep: Daily

Reference: Meyer et al. [2010], Franklin et al. [2014]

There will be many different strategies here when the model is calibrated and tested as is currently happening in 3.2.2. Plans of these include, but are not limited to:

- Original MYCOFON dynamics [Meyer et al., 2010] (Coded)
- Näsholm et al. [2013] strategies (Coded)
- The trees / mycorrhiza want to optimise the system, not themselves [Baskaran et al., 2017]
- The trees / mycorrhiza want to optimise their growth (within this C, N and C:N)
- The trees / mycorrhiza want to optimise the a growth form (within this C, N and C:N) [Valverde-Barrantes et al., 2017, Bergmann et al., 2020]
- The trees / mycorrhiza want to optimise their limiting element uptake (within this C, N and C:N)
- The C surplus theory, that C is given proportionally to the overflow from the plant.
- Combinations of these strategies will make the system Game Theoretical

Reference values come from Ingestad et al. [1986], Hobbie [2006], and behaviours from Johnson et al. [2006], Van't Padje et al. [2021], Högberg et al. [2010], Blaško et al. [2015].

#### 5.2.1 Plant Decision

Currently two strategies are coded. These are derived from Näsholm et al. [2013] and the Mycofon model [Meyer et al., 2010]. The strategy of Mycofon gives a demand of 1 and and transfer amount stated in Equation 36.

Transfer = 
$$\max\{\text{allo}_{\max}C_r, \text{allo}(C_r - (\text{root : mycorrhizal})_{\text{opt}}) - C_f\}$$
 (36)

$$allo_{\text{max}} = \begin{cases}
1 - (1 - [\exp(-50 * N_r)]^3) & \text{for } N_r < 0.01 \\
0.2 & \text{else}
\end{cases}$$

$$allo = \begin{cases}
\frac{N_{\text{allo}}}{N_r + N_{\text{allo}}} & \text{for } N_{\text{allo}} < 0.5 \\
1 & \text{else}
\end{cases}$$
(38)

$$allo = \begin{cases} \frac{N_{\text{allo}}}{N_r + N_{\text{allo}}} & \text{for } N_{\text{allo}} < 0.5\\ 1 & \text{else} \end{cases}$$
 (38)

Where  $C_r$  and  $C_f$  are the carbon in the roots and mycorrhiza respectively, (root: mycorrhizal)<sub>opt</sub> is the optimal root mycorrhizal biomass ratio,  $N_r$  is the nitrogen in the roots and  $N_{allo}$  is the nitrogen allocated by the mycorrhiza.

The strategy of Näsholm gives a demand of 1 and and a transfer shown in Equation 39

$$\max \left[ C_{\text{Allocated}}^{\text{CASSIA}}, 0.0 \right] \tag{39}$$

where CASSIA is the maximum C allocation to the mycorrhiza calculated in CASSIA.

#### 5.2.2 Mycorrhizal Decision

Currently two strategies are used. These are derived from Näsholm et al. [2013] and the Mycofon model [Meyer et al., 2010]. The strategy of Mycofon has a demand of 1 and a transfer function given as below 40.

Transfer = 
$$\max \left[ N_f \left( 1 - \frac{N_r}{C_r \cdot (N:C)_{\text{opt}}} \right), 0.0 \right]$$
 (40)

where  $N_f$ ,  $N_r$  are the N in the mycorrhiza and root respectively,  $C_r$  is the C in the root and  $(N:C)_{opt}$  is the optimal N:C ration in the root.

The strategy of Näsholm has a demand of 1 and a transfer of

$$\max\left[N_f - G_f, 0.0\right] \tag{41}$$

Where N and G are the nitrogen and growth of the mycorrhiza. NOTE: add the table of symbols when the equations are finalised

#### Mycorrhizal Code 5.3

#### Mycorrhizal Growth 5.3.1

The growth of mycorrhiza is controlled currently by the same logic as CASSIA [Schiestl-Aalto et al., 2015, where the mycorrhizal growth is first calculated as possible growth based on temperature (Equation 45). Then the Non-Structural Carbon and Nitrogen form a limitation, as in Meyer et al. [2010] and Franklin et al. [2014]. NOTE: the timing of root growth is used for mycorrhiza temporarily for testing, parameterisation will soon follow.

$$g_F(t) = \begin{cases} 0 & T_a(t) < 0\\ (1 - \exp(-\lambda M(t))(1 + \exp(-a(T_a(t) - b))^{-1} & T_a(t) \ge 0 \end{cases}$$
(42)

Where  $g_R(t)$  is the first step in calculating the possible growth.  $\lambda$  is a parameter that decreases fine root growth during water deficiency, M(t) is the soil moisture content,  $T_a$  is the temperature at soil depth a, finally a and b are fitted growth parameters.

$$f_F(t) = \begin{cases} 0 & s_i \le 0\\ \frac{1}{2} \left( \sin \left( \frac{2\pi}{s_i^c} \left( s_i(t) - \frac{s_i^c}{4} \right) \right) + 1 \right) & 0 \le s_i \le s_i^c\\ 0 & s_i \ge s_i^c \end{cases}$$

$$G_F = \frac{L0}{\text{Mycelium Lifespan}} \cdot f_F(t) \cdot g_F(t)$$

$$(43)$$

$$G_F = \frac{L0}{\text{Mycelium Lifespan}} \cdot f_F(t) \cdot g_F(t)$$
 (44)

Potential Mycelium Growth = 
$$\begin{cases} G_F & G_F > 0 \\ 0 & G_F \le 0 \end{cases}$$
 (45)

More information can be found in Schiestl-Aalto et al. [2015]! NOTE: the rest of the equations for the rest of the organs will be added to their respective sections at a later date. Now the rest of the equations not in CASSIA. The potential ectomycorrhizal growth is then limited by the stores of carbon and nitrogen.

$$f_C \approx \frac{C_{NS}^f}{C^f} \tag{46}$$

$$f_N \approx \frac{N_{NS}^f}{N^f} \tag{47}$$

$$Mycelium Growth = Potential Mycelium Growth \cdot f_C \cdot f_N$$
 (48)

#### **Assumptions:**

• Need to assume a upper limit to the storage capacities and the effect that storage would have on growth.

#### 5.3.2 Mycofon Balance

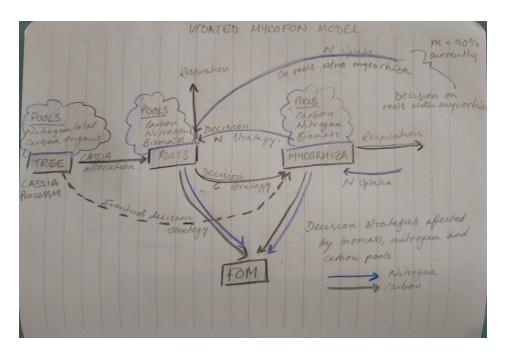


Figure 6: Flow chart for the updated MYOCFON model. Exact processes are represented by an arrow, more detailed processes are represented in the sections / flow diagrams referenced. The uptake rates etc. all have modifications by the environment, which are not written explicitly here.

### Model Lowdown: MYCOFON Inspired.

Model type: Daily N and C cycling model, with a little bit of growth

Inputs: Temperature, N and C, biomass

Scope: Tree or stand level, depends on the inputs

Timestep: Daily

**Reference:** Version inspired by: Meyer et al. [2010]

The official MYCOFON model can be gotten by contacting Hongxing He (hongxing.he@gu.se). Due to the links with the Symphony and CASSIA models I have remade the MYCOFON model in C++ for this project.

The MYCOFON model (Meyer, 2010), with updates to include the uptake properties for three types of nitrogen as well as the decision functionality. This function brings together the C or N balance as in the original MYCOFON model, the decision functions for resource transfer, mycorrhizal growth and multiple N uptake functions. The equations in this function are below.

$$m = \frac{C_f}{B_{opt}^{rf} C^r} \tag{49}$$

$$\frac{dC^r}{dt} = \text{Growth from CASSIA} - (1 - m)s^r C^r - ms^m C^r$$
 (50)

$$\frac{dC^r}{dt} = \text{Growth from CASSIA} - (1 - m)s^r C^r - ms^m C^r$$

$$\frac{dC^f}{dt} = g^f - s^{\text{mantle}} C^{\text{mantle}} - s^{\text{ERM}} C^{\text{ERM}} - R(T_{\text{B}}) C^f$$
(50)

$$\frac{dC_{\text{NonStruct}}^{r}}{dt} = C_{\text{Allocated}}^{\text{CASSIA}} - d^{r}(C^{\text{CASSIA}})$$
(52)

$$\frac{dN_{\text{NonStruct}}^r}{dt} = d^f(C^f) + u^f C^f \tag{53}$$

$$\frac{dN_{\text{NonStruct}}^{r}}{dt} = d^{f}(C^{f}) + u^{f}C^{f}$$

$$\frac{dC_{\text{NonStruct}}^{f}}{dt} = d^{r}(C^{\text{CASSIA}}) - g^{f}$$

$$\frac{dN_{\text{NonStruct}}^{f}}{dt} = u^{f}C^{f} - g^{f} - d^{f}(C^{f})$$
(53)

$$\frac{dN_{\text{NonStruct}}^f}{dt} = u^f C^f - g^f - d^f(C^f) \tag{55}$$

Table 5: Mycofon: table of symbols

			5. Mycoloff, table of symbols
Symbol	Function in	Units	Meaning
	the system		
			Variable
t	Variable	Day	Time
N	Variable	kg N	Amount of nitrogen in specified organs
С	Variable	kg C	Amount of carbon in specified organs
В	Variable	kg C	Biomass
$C_{\text{Allocated}}^{\text{CASSIA}}$	Variable	kg C	The maximum amount of sugar that could be allocated from the CASSIA
T	Variable /	Damaga C	model Terror proteins
1	,	Degrees C	Temperature
	Input time-		
	series	0.1	Marankin dan atau ati Although a famoula in annount die the dans
m	Variable	0-1	Mycorrhized roots ratio. Although a formula is presented in the docu-
			ment, 0.9 is currently being used for the testing phase.
			Index
r, f	Index		Indexes for the roots and mycorrhiza respectively
opt	Index		Refers to the optimal value of the indicated variable
m, r	Index		Mycorrhized or non-mycorrhized roots
ERM	Index		Index referring to the extra-radical mycelium
Mantle	Index		Index referring to the mantle
В	Index		B soil horizon
			Parameter
s	Parameter	kg C day-1	Turnover
			Function
d()	Function	0-1	Decision function, gives the allocated amount of carbon or nitrogen de-
			pending on the direction of the function, see Section 5.2
R(Temperature)	) Function	kg C	Respiration for indicated compartment, see Section 4
g()	Function	kg C	Growth function, for the mycorrhiza

Where N is nitrogen, C is carbon, r roots, f mycorrhiza, B biomass, opt is optimum. The uptake and decision functions are then calculated from the balance of the last iteration. Note that the

plant decision requires the N allocated, so the mycorrhizal decision is calculated first in the code. Where,  $C_{\text{Allocated}}^{\text{CASSIA}}$  is the carbon allocated from the CASSIA model, s is turnover, which is different for mycorrhized (m) and non-mycorrhized roots (r),  $T_B$  is temperature of the soil, d() is decision (including exudes and transfer to mycorrhiza) and R() is respiration, although for the roots this is calculated in CASSIA not here. Again, the growth process is currently handled in CASSIA so this is just the cumulative sum of the growth with the turnover negated. Growth for mycorrhiza is represented by g().

Note: the m value is currently 0.9, although will be changed to this formulation when the model is fully tested.

#### **Assumptions:**

• As the root:mycorrhiza surface area ratio should control the mycorrhizal uptake of C from the tree, and in boreal systems there is a high colonisation rate [Smith and Read, 2010], therefore I assume that there is an optimal relationship one, this assumption could effect the underground dynamics, both in terms of the type of N as well as the colonisation dynamics

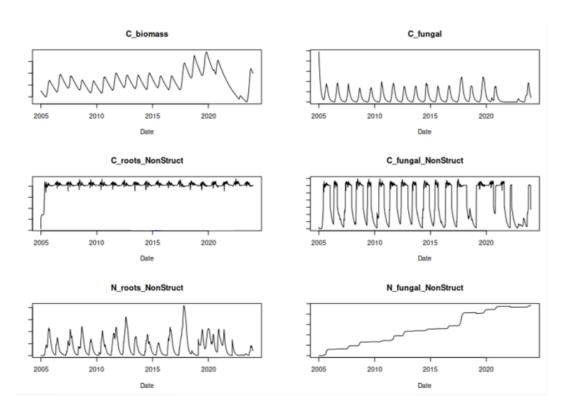


Figure 7: Figure to show the current growth in the model with the Näsholm et al. [2013] decision strategy.

### 5.4 Soil Processes: Soil Model

NOTE: Talking about replacing these equations with Ghersheen et al., however, the logic of the following equations is still valid!

Whilst the aim of the project is **to account for nitrogen and carbon transfer between trees and ectomycorrhiza in a boreal ecosystem**, to do this a soil model is needed. As the amount of nitrogen in the soil depends on the fungal competition, it doesn't make sense to study the strategic transfer between the tree and their symbionts without the other functional group competition between microbes in the soils. And also the feedback of the competitive advantage this symbiosis gives to the ectomycorrhiza in the soil.

#### Model Lowdown: Soil Model

Model type: Soil C and N, Process-Organism Model [Perveen et al., 2014]

Inputs: Litter input, initial soil compartment amounts, temperature of air and soil

Scope: Microbial compartmentalisation of soil, with enzymatic N uptake

Timestep: Daily

Reference: Perveen et al. [2014] Baskaran et al. [2017]

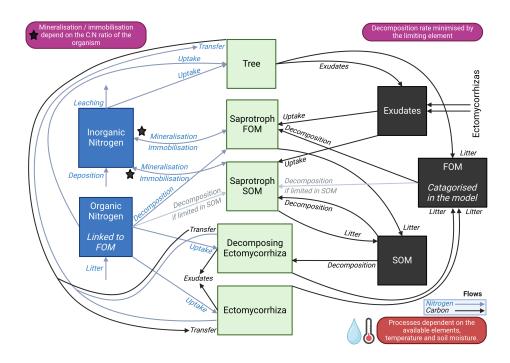


Figure 8: Temporary figure for the soil model. Figure has all of the processes that we want to be represented in the equations.

As the model is seasonal and conceptual so the model should be looked at at two levels: Short term results:

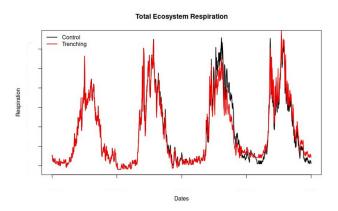
- Priming effect from trenching experiments (priming effect can be seen) [Ryhti et al., 2021, 2022]
- Nitrogen and Carbon transfer tracked at Hyyitälä (data being gathered)
- Growth patterns within the seasonal pattern [Ekblad et al., 2013] (roots come from the tree model [Ding et al., 2020] and nitrogen availability based root strategy being added)

The long term results should respond to disturbance:

- Change in fungal composition / dominance of fungal strategy Lindahl et al. [2021]
- Changes in carbon stock after disturbances reflecting the different recycling strategies of the different microbes Clemmensen et al. [2015], Sterkenburg et al. [2015]
- Nitrogen immobilisation in old growth forests Sterkenburg et al. [2015]
- Inorganic vs organic nitrogen stock changes after disturbances Lindahl et al. [2010]

This soil model was originally inspired by Perveen et al. [2014], but in order to make the model fit for boreal forests the model has ended up looking a lot like Baskaran et al. [2017] and to a lesser extent [Sulman et al., 2014]. Only with a seasonal timestep and some small differences in the equations. This is encouraging in terms of the understanding of the system, however should be carefully considered in terms of the model use. Exploring using another soil model is also in progress.

The behaviour so far:



NOTE: Equations are from an old iteration of the model so exudation and decomposing ectomycorrhiza are missing.

$$\frac{dC_{\text{SOM}}}{dt} = -u_{\text{C}}(C^{SOM})C^{\text{microbe, SOM}} + s\left(C^{\text{microbe, SOM}} + C^{\text{microbe, FOM}}\right)$$
(56)

$$C_{FOM} = C_{FOM}^{needles} + C_{FOM}^{woody} + C_{FOM}^{roots} + C_{FOM}^{mantle} + C_{FOM}^{ERM}$$

$$N_{FOM} = N_{FOM}^{needles}(N:C)^{needles} + N_{FOM}^{woody}(N:C)^{woody}$$

$$+ N_{FOM}^{roots}(N:C)^{roots} + N_{FOM}^{mantle}(N:C)^{mantle} + N_{FOM}^{ERM}(N:C)^{ERM}$$
(58)

$$+N_{FOM}^{roots}(N:C)^{roots} + N_{FOM}^{mantle}(N:C)^{mantle} + N_{FOM}^{ERM}(N:C)^{ERM}$$
 (58)

$$d_{total} = u_C(C^{FOM})C^{microbes,SOM} + u_C(C^{FOM})C^{microbes,FOM}$$
(59)

$$\frac{dC_{FOM}^{needles}}{dt} = L^{needles} - d_{total} \frac{C_{FOM}^{needles}}{C_{FOM}} \tag{60}$$

$$\frac{dC_{FOM}^{woody}}{dt} = L^{woody} - d_{total} \frac{C_{FOM}^{woody}}{C_{FOM}} \tag{61}$$

$$\frac{dC_{FOM}^{roots}}{dt} = L^{roots} - d_{total} \frac{C_{FOM}^{roots}}{C_{FOM}}$$

$$\tag{62}$$

$$d_{total} = u_C(C^{FOM})C^{microbes,SOM} + u_C(C^{FOM})C^{microbes,FOM}$$

$$\frac{dC^{needles}_{FOM}}{dt} = L^{needles} - d_{total} \frac{C^{needles}_{FOM}}{C_{FOM}}$$

$$\frac{dC^{woody}_{FOM}}{dt} = L^{woody} - d_{total} \frac{C^{woody}_{FOM}}{C_{FOM}}$$

$$\frac{dC^{roots}_{FOM}}{dt} = L^{roots} - d_{total} \frac{C^{roots}_{FOM}}{C_{FOM}}$$

$$\frac{dC^{mantle}_{FOM}}{dt} = L^{mantle} - d_{total} \frac{C^{mantle}_{FOM}}{C_{FOM}}$$

$$\frac{dC^{ERM}_{FOM}}{dt} = L^{ERM} - d_{total} \frac{C^{ERM}_{FOM}}{C_{FOM}}$$

$$(63)$$

$$\frac{dC_{FOM}^{ERM}}{dt} = L^{ERM} - d_{total} \frac{C_{FOM}^{ERM}}{C_{FOM}} \tag{64}$$

Where C and N are carbon and nitrogen respectively, (N:C) is the carbon nitrogen ratio of the indicated compartment, L is letter and the compartments are referred to explicitly. FOM is fresh organic matter and SOM is soil organic matter. Where d is the decomposition, which is the uptake of both microbe pools, which is seen in more detail in Section 5.1.2. NOTE: the uptake is only from the aggregated FOM pool rather than each of the separate FOM pools and the decomposition from each pool is currently based on the size of the pool rather than the type of material. This will be changed to reflect the type of material.

$$\frac{dNH4}{dt} = (\psi_{\text{ims}} + \psi_{\text{imf}}) \frac{NH4}{NH4 + NO3} - NH4^{p}_{used} - NH4^{f}_{used} - u_{NH4}C^{microbes,FOM} - u_{NH4}C^{microbes,SOM}$$
(65)

$$\frac{dNO3}{dt} = (\psi_{\text{ims}} + \psi_{\text{imf}}) \frac{NH4}{NH4 + NO3} - NO3_{used}^p - NO3_{used}^f - u_{NO3}C^{microbes,FOM} - u_{NO3}C^{microbes,SOM}$$
(66)

$$\frac{dN^{FOM}}{dt} = \text{Litter} - N_{used}^{FOM,p} - N_{used}^{FOM,f} - u_{Norg}C^{microbes,FOM} - u_{Norg}C^{microbes,SOM}$$
 (67)

$$\frac{dN^{SOM}}{dt} = \text{Microbe Litter} - u_{Norg}C^{microbes,SOM}$$
(68)

NOTE: in the original symphony model there is leaching and fertilisation. These should be added, but currently not in the testing phase.

Where  $NH4^p_{used}$  is the NH4 used for the plant, f represents the mycorrhizal uptake. This is an input to the function from the mycofon model. Next the microbe balance is considered.

$$\frac{dC^{\text{microbe, FOM}}}{dt} = u_{FOM}C^{\text{microbe, FOM}} - r(T_B) - s \tag{69}$$

$$\frac{dC^{\text{microbe, FOM}}}{dt} = u_{FOM}C^{\text{microbe, FOM}} - r(T_B) - s$$

$$\frac{dC^{\text{microbe, SOM}}}{dt} = u_{SOM}C^{\text{microbe, SOM}} - r(T_B) - s$$
(69)

#### Assumptions and behaviours:

- Microbes assumed to be organisms that are not in symbiosis with the tree and that form competition for the nitrogen resources from the same pools as trees or ectomycorrhiza.
- Therefore the priming effect is still included as if there is more C in the soil then the decomposition will increase to a N storage bound value. The next day the N uptake can compensate for the N that was used the previous day with the C increase. This means that over time there would be a priming effect - but it is not instantaneous.
- In the model there are no chemical processes as the idea is to look at symbionts and microbes this make sense, but could be a bad assumption.
- The respiration is not variable yet. Ryhti et al. [2022] data and models planning to be used for this.

Table 6: Soil Model: table of symbols

Symbol	Function in the system	Units	Meaning
	one system		Variable
t	Variable	Day	Time
N	Variable	kg N	Amount of nitrogen in specified organs
C	Variable	kg C	Amount of carbon in specified organs
${ m T}$	Variable	Degrees C	Temperature
NH4	Variable	kg N	The amount of NH4 in the soil
NO3	Variable	kg N	The amount of NO3 in the soil
			Index
needles	Index		Subscript indicating the needles
woody	Index		Subscript indicating the woody biomass
roots	Index		Subscript indicating the roots
ERM	Index		Subscript indicating the extra-radical mycelium
mantle	Index		Subscript indicating the mantle
FOM	Index		Subscript indicating the fresh organic matter
SOM	Index		Subscript indicating the soil organic matter
p, f	Index		Subscripts indicating the plant and fungi respectively
B	Index		Soil B horison
NH4	Index		For processes related to NH4 pools
NO3	Index		For processes related to NO3 pools
Norg	Index		For processes related to Norg pools
microbe	Index		For processes related to microbes
			Parameter
s	Parameter	kg C day-1	Turnover
			Function
u	Function	kg N	Uptake for indicated compartment, see section 5.1.2

# 6 Complete table of variables

TODO: update the following table with new equations.

Table 7: All Sections: Table of Symbols

Symbol	Function in Units the System	Meaning
		Variable
$\overline{W}$	Variable	Amount of carbon in the indicated organ
A	Variable	Amount of enzyme
T	Variable $^{\circ}\mathrm{C}$	Temperature (input to the model timeseries)
t	Variable Day	Time
N	Variable	Amount of nitrogen in specified organs
C	Variable	Amount of carbon in specified organs
B	Variable kg C	Biomass
$C_{\text{Allocated}}^{\text{CASSIA}}$	Variable	Maximum sugar allocatable from the CASSIA model
m	Variable 0–1	Mycorrhized roots ratio (0.9 used in testing phase)
$\mathrm{NH_4}$	Variable	Amount of NH <sub>4</sub> in soil
$NO_3$	Variable	Amount of $NO_3$ in soil
		Index
r, f	Index	Roots and mycorrhiza indices
opt	Index	Refers to the optimal value of the variable
m, r	Index	Mycorrhized or non-mycorrhized roots
ERM	Index	Extra-radical mycelium
Mantle	Index	Mantle tissue
needles	Index	Needles
woody	Index	Woody biomass
roots	Index	Roots
ERM	Index	Extra-radical mycelium
mantle	Index	Mantle
FOM	Index	Fresh organic matter
SOM	Index	Soil organic matter
p, f	Index	Plant and fungi
B	Index	Soil B horizon
$\mathrm{NH_4}$	Index	Related to NH <sub>4</sub> pools
$NO_3$	Index	Related to NO <sub>3</sub> pools
Norg	Index	Related to organic nitrogen pools
Microbe	Index	Related to microbial processes
		Parameter
λ	Parameter	Decay rate of enzymes
$\delta$	Parameter	Enzymatic parameter
A	Parameter	Rate of carbohydrate production

$\begin{array}{cccccccccccccccccccccccccccccccccccc$	$\operatorname{owth}$
$\begin{array}{c ccccccccccccccccccccccccccccccccccc$	$\operatorname{owth}$
$\begin{array}{cccccccccccccccccccccccccccccccccccc$	
$\begin{array}{c ccccccccccccccccccccccccccccccccccc$	ening /
$\begin{array}{c ccccccccccccccccccccccccccccccccccc$	
d()Function $0-1$ Decision function allocating C or N depending on c $u()$ FunctionUptake for specified compartment (see Section 5.1.2 $R(Temperature)$ FunctionRespiration function (see Section 4) $g()$ Functionkg CGrowth function for mycorrhiza	
u() Function Uptake for specified compartment (see Section 5.1.2 $R(Temperature)$ Function Respiration function (see Section 4) $g()$ Function kg C Growth function for mycorrhiza	
u() Function Uptake for specified compartment (see Section 5.1.2 $R(Temperature)$ Function Respiration function (see Section 4) $g()$ Function kg C Growth function for mycorrhiza	$\overline{\text{ontext}}$
g() Function kg C Growth function for mycorrhiza	
	,
Process	
K Process Creation of sugar or starch	
Q Process Sperling model amalgamated process	
$E(W^{Ss}, W^{St})$ Process Emergency transfer of starch to sugar under low sug	ar con-
ditions	
$f(W_i^{all})$ Multiplier Scaling function (0-1) for growth based on carbo	nydrate
reserves	
$B(W_r^{all})$ Process Allocation to mycorrhiza	
Other	
P(t) Timeseries Photosynthesis (input timeseries; no feedback)	
G(t) Input, state Growth based on CASSIA model [Schiestl-Aalto et a	, 2019]
variable	-
R(t) State variable Respiration including growth and maintenance [S	chiestl-
Aalto et al., 2019]	
D Data Observational data from Hyytiälä	
CASSIA $(\theta)$ Model CASSIA model with Hyytiälä-specific inputs and particles ters	

# 7 Appexdix: Parameter Values in Code

## 7.1 repo\_p Explanation

These parameters are from the paper Repola [2009] and is a multi-variable model fitted for Scots Pine and Norway Spruce in Finland.

Parameter Name	Parameter Value	Equations
b0.repo b1.repo b2.repo uk.repo eki.repo	-6.303 14.472 -3.976 0.109 0.118	This model is from Repola [2009] and not explained further here. As the model is a mixed effect model, their parameters here do not have clear biological meanings, but are fixed effect parameters.

Table 8: Explanation for the repo\_p parameters built into the package.

# ${\bf 7.2}\quad {\bf common\_p}\ {\bf Explanation}$

Parameter Name (Name in Paper	Parameter value	Units	Comments and additional references
a (a)	0.185	Degree C <sup>-1</sup>	Parameter of function g
b(b)	18.4	Degree C	Parameter of function g
$\overrightarrow{\mathrm{TR0}}$	0	G	Root temperature factor
abs_zero	273.15	K	Absolute zero temperature
b.s	4.14		Soil water potential and conductance
theetta.FC	0.62		Hölttä et al. 2009 Table 2
phi.e	$-6.8 \times 10^{-7}$	MPa	Hölttä et al. 2009 Table 2, note psi.s should be psi.e
K.sat	24.5	$\begin{array}{c} \mathrm{mol} \ \mathrm{m}^{\text{-}1} \ \mathrm{s}^{\text{-}1} \\ \mathrm{MPa}^{\text{-}1} \end{array}$	Hölttä et al 2009, Table 2
R.length	5300	${\rm m}~{\rm root}~{\rm m}^{-2}$	Index, Hölttä et al 2009
M.H2O	0.018	${\rm kg\ mol^{-1}}$	Hölttä et al 2009
r.cyl	$4.25 \times 10^{-3}$	m	Hölttä et al 2009
r.root	$3.0 \times 10^{-3}$	m	Hölttä et al 2009
ypsilon	$1 \times 10^{-14}$		To prevent dividing by zero
$ m Rg.N~(r_R^g)$	0.35	$kg C(kg C)^{-1}$	Growth respiration, share of growth (Needles)
Rg.S $(r_{\text{wood}}^g)$	0.3	$kg \overset{'}{C}(kg \overset{'}{C})^{-1}$	Growth respiration, share of growth (Wood)
Rg.R $(r_N^g)$	0.35	$kg C(kg C)^{-1}$	Growth respiration, share of growth (Fine roots)
gas.const $(R)$	8.314	J mol <sup>-1</sup> K <sup>-1</sup>	Gas constant
$M.C(M_C)$	12.01	g mol <sup>-1</sup>	Molar mass of Carbon
M.H	1.008	$g \text{ mol}^{-1}$	Molar mass of Hydrogen
M.O	16	$g \text{ mol}^{-1}$	Molar mass of Oxygen
osmotic.sugar.conc	2000000	Pa	Osmotic sugar concentration 2 MPa Hölttä et al.
m n	0.02174605		2000
m_n Uggla	1.95	_	Division to early/late wood

Table 9: Explanation for the common\_p parameters built into the package

## 7.3 ratio\_p Explanation

Parameter Name (Name in		Paramet	er value		Units	Comments and additional	
Paper)	Hyytiälä Lettosuo Väriö			Chinese Site	011100	references	
form_factor $(\varphi)$	0.6	0.6	0.55		Ratio	Lettosuo: The multiplier between a cylinder (with diam=D0, height=h0) and the total biomass of stem, coarse roots and branches	
needle_fineroot_ratio	NA	1/2.9	NA		Ratio	Depends on tree size, species and site and *	
sapwood.share	0.8	0.8	0.8		Ratio	_	
height_growth_coefficient $(\alpha_S)$	4.3	1	2.6		_	repeated value, Variö: leader shoot length / average measurement shoot length (average over years)	
diameter_growth_coefficient $(\alpha_D)$	1.6	1	0.8		_	repeated value	
height_growth_coefficient_max	5.5	1.28	NA		_	repeated value, min if growth decreases	
$height\_growth\_coefficient\_min$	3.8	0.88	NA		_	repeated value, min if growth decreases	
$diameter\_growth\_coefficient\_ma$	x 1.9	1.19	NA		_	repeated value, max if growth decreases	
$diameter\_growth\_coefficient\_min$	n 1.5	0.94	NA			repeated value, min if growth decreases	

Table 10: Explanation for the ratios\_p vector built into the package. Additional comment \* "Lettosuo: Helmisaari et al. 2006 Tree physiology -; 2.0 for VT, 3.8 for MT, 5.7 for OMT. Very nice curve for needles / fine roots vs. fine root N % leads now to  $100~\rm gC$  m-2 (roots ; 2 mm) but result lower than in Leppälammi-Kujansuu et al. (2013, Plant Soil) where they found ca 225 gC m-2 (roots ; 2 mm) in control and 300-350 in fertilized ca 225 gC m-2 (roots ; 2 mm) in control and 300-350 in fertilized (Leppälammi-Kujansuu et al. 2013, Plant Soil) "

# $7.4 \quad parameters\_p \ Explanation$

Table 11: Explanation for the parameters\_p vector built into the package  $\,$ 

Parameter Name (Name		Paramet	er value		Units	Comments				
in Paper)	Hyde	Lettosuo	Väriö	HF China	Omes	Comments				
Respiration										
Q10.N $(q_{\rm N}^{10})$	1.898	1.898	1.898		_	Needles Q10				
Rm.N $(r_N^m)$	0.00267	0.0020	0.004005		$kg C(kg C)^{-1}$	Needles R0				
Q10.S $(q_{Wood}^{10})$	1.74788	1.74788	1.74788		_	Wood Q10				
Rm.S $(r_S^m)$	5.5576e-5	5.5576e-5	8.3364e-5		$kg C(kg C)^{-1}$	Wood R0				
Q10.R $(q_{\rm R}^{10})$	2.5575	2.9662	2.0244		_	Fine roots Q10				
Rm.R $(r_R^m)$	0.00958	0.0059	0.00945		$\begin{array}{c} \mathrm{kg} \ \mathrm{C}(\mathrm{kg} \\ \mathrm{C})^{-1} \end{array}$	Fine roots R0				
Growth										
$\operatorname{sRc}(S_R^c)$	30	30	25		_	Root growth cessation				
My corrhiza										
growth.myco	0.1	0.1	0.1			Mycorrhiza growth				
root.lifetime	1.7	2.01	2		years	Root lifetime				
		Shoot	s							
НН0	10	10	10		mm	Initial shoot length				
sH0	- 1.359200388	-4.12008	-1.44		_	Shoot growth beginning				
LH	8.226401284	8.226401284	9.0			Shoot				
LH0	8.226401284	8.226401284	9.0			growth rate Initial shoot growth rate				

Table 11 continued from previous page

	Table 1	1 continued f	rom previous pa	age		
Parameter Name (Name		Parame	eter value		Units	Comments
in Paper)	Hyde	Lettosuo	Väriö	HF China	011165	Comments
sHc	14.59636279	12.8328	10.0		_	Shoot growth cessation
		Need	lles			
$\mathrm{sN0}\ (S_N^0)$	-8.37584	-3.56589	-5.6		-	Needle growth beginning
	LN parameter is	s a typo, it si	hould be LNO, be	ut will solve to		
LN	1.849493	1.849493	3.5		$\mathrm{mm}\ \mathrm{d}^{-1}$	Needle growth rate
$LN0 \ (L_{N0})$	1.849493	1.849493	3.5		mm d <sup>-1</sup>	Initial needle
$\operatorname{sNc}(S_N^c)$	5.263883	7.60671	4.327		-	growth rate Needle growth
HN0	1	1	1		mm	cessation Initial needle length
		Diam	eter			
sD0.Trad	3.724083738	-3.5	- 2.434161213		_	Diameter growth beginning
TODO: the first	LD parameter is	s a typo, it si	hould be LD0, b	ut will solve ti	his issue la	
LD	1.293443902	1.9	2.9		cells d <sup>-1</sup>	Diameter growth rate
$\mathrm{LD0}\ (L_{\mathrm{D0}})$	1.293443902	1.9	2.9		cells d <sup>-1</sup>	Initial diameter growth rate
$\operatorname{sDc} (S_D^c)$	5.077004992	5.2	4.093829285		_	Diameter growth cessation
sDc.T.count	NA	8.8	NA		_	Diameter growth cessation count
		Duration pe	arameters			

Table 11 continued from previous page

Parameter Name (Name		Parame	eter value		Units	Comments
in Paper)	Hyde	Lettosuo	Väriö	HF China	OHIOS	Comments
tau. Ee $(\tau_{\rm e}^{\rm early})$	10.68685877	5.5	10.25174759		day	Early wood cell en- largement duration
tau. El $(\tau_{\rm e}^{\rm late})$	8.789131263	4.8	4.510400352		day	Late wood cell en- largement duration
tau. We $(\tau_{\rm wa}^{\rm early})$	25.29448857	17.8	51.60724145		day	Early wood cell wall formation duration
tau. W l $(\tau_{\rm wa}^{\rm late})$	35.12148687	19.2	17.76015932		day	Late wood cell wall formation duration
		GP	P			
tau.GPP $(\tau_{\text{GPP}})$	5	5	5		_	GPP effect on daily LD
Uggla	1.95	1.8	1.95		_	Division to early/late wood
		Buc	ls			
sB0	171	171	181		day	Bud growth beginning
sBc	85	85	60		day	Bud growth cessation
LB	0.005	0.005	0.003			Bud growth rate
		Xyloge	nesis			
cell.d.ew $(d_{\text{cell}}^{\text{early}})$	35.7e-6	32.1e-6	30e-6		m	Early wood cell diameter
cell.d.lw $(d_{\text{cell}}^{\text{late}})$	24.2e-6	27.5e-6	20e-6		m	Late wood cell diameter
cell.l.ew $(l_{\text{cell}}^{\text{early}})$	2.59e-3	2.89e-3	2.59e-3		m	Early wood cell length
					Continue	d on next page

Table 11 continued from previous page

Parameter Name (Name		Paramet		0 -	Units	Comments
in Paper)	Hyde	Lettosuo	Väriö	HF China	Omes	Comments
cell.l.lw $(l_{\text{cell}}^{\text{late}})$	2.73e-3	2.97e-3	2.73e-3		m	Late wood cell length
cell.wall.density.ew	57	570	557		$ m kg~C$ $ m m^{-3}$	Early wood cell wall density
cell.wall.density.lw	57	680	557		$ m kg~C$ $ m m^{-3}$	Late wood cell wall density
wall.thickness.ew	2.61e-6	3.1e-6	2.61e-6		m	Early wood wall thickness
wall.thickness.lw	5.23e-6	3.88e-6	5.23e-6		m	Late wood wall thickness
cell.volume.growth.per.day.ev	v NA	5.49e-13	NA		$\mathrm{m}^3$ $\mathrm{day}^{-1}$	Early wood cell volume growth rate
cell.volume.growth.per.day.lw	n NA	4.62e-13	NA		$\mathrm{m}^3$ $\mathrm{day}^{-1}$	Late wood cell volume growth rate
density_tree $(\rho)$	400	400	400		${\rm kg~m^{\text{-}3}}$	Tree density
carbon_share	0.5	0.5	0.5		kg kg <sup>-1</sup>	Carbon share
D0 $(d_0)$	0.175	0.175	0.154		m	Initial diameter
h0	17.9	17.9	9.5		m	Initial height
n_age	3	5	5		years	Needle lifespan
n_lenght	34.241	13	39.2		mm	Average needle length
h_increment	309.0938	309.0938	120.00		mm	Mean height increment
SLA	13	5.5	13		$\mathrm{m^2~kg^{-1}}$	Specific leaf area
LR0 $(L_R^0)$	0.07446064	NA	0.02		kg C d <sup>-1</sup>	LR0 parameter on next page

Table 11 continued from previous page

Parameter Name (Name		Paramet	er value		Units	Comments
in Paper)	Hyde	Lettosuo	Väriö	HF China	Omos	Comments
		Repola para	meters			
b0_repo	-6.303	NA	NA		_	Repola [2009] Parameter
b1_repo	14.472	NA	NA		_	Repola [2009] Parameter
b2_repo	-3.976	NA	NA		_	Repola [2009] Parameter
		Sperling pare	ameters			
lower_bound_needles	0.02	NA	NA		kg C	Lower bound for needles
$lower\_bound\_phloem$	0.03	NA	NA		kg C	Lower bound for phloem
$lower\_bound\_roots$	0.05	NA	NA		kg C	Lower bound for roots
lower_bound_xylem_sh	0.03	NA	NA		kg C	Lower bound for shoot xylem
$lower\_bound\_xylem\_st$	0.1	NA	NA		kg C	Lower bound for stem xylem
$tau\_emergancy\_needles$	3	NA	NA		_	Emergency time constant for needles
$tau\_emergancy\_phloem$	3	NA	NA		_	Emergency time constant for phloem
tau_emergancy_roots	3	NA	NA		_	Emergency time constant for roots

Table 11 continued from previous page

Parameter Name (Name		Paramet	er value		Units	Comments
in Paper)	Hyde	Lettosuo	Väriö	HF China	Omos	Comments
tau_emergancy_xylem_sh	3	NA	NA		-	Emergency time constant for shoot xylem
$tau\_emergancy\_xylem\_st$	3	NA	NA		_	Emergency time constant for stem xylem
$lower\_bound\_W$	0.01	NA	NA		kg C	Lower bound for W
tau_emergancy	3	NA	NA		-	General emergency time constant
uk_repo	0.109	NA	0.109		_	Repola [2009] Parameter
eki_repo	0.118	NA	0.118		_	Repola [2009] Parameter
stem_no	3	NA	NA		_	Repola [2009] Parameter

# $7.5 \quad sperling\_p \ Explanation$

Table 12: Explanation for the sperling\_p vector built into the package

Parameter Name	meter Name Parameter value			Units	Comments	
Tarameter Ivanie	Hyde	Lettosuo	Flakaliden	HF China	Omes	Comments
starch0	0.3246781	0.40	0.40			Initial starch
						concentration
sugar0	0.4184208	0.35	0.35			Initial sugar
						concentration
starch.needles0	0.03	NA	NA			Initial needle starch
						concentration
starch.phloem0	0.037	NA	NA			Initial phloem starch
						concentration
starch.xylem.sh0	0.034	NA	NA			Initial shoot xylem
						starch concentration
starch.xylem.st0	0.166	NA	NA			Initial stem xylem
						starch concentration
starch.roots0	0.057	NA	NA			Initial root starch
						concentration
sugar.needles0	0.087	NA	NA			Initial needle sugar
						concentration
sugar.phloem0	0.27	NA	NA			Initial phloem sugar
						concentration
sugar.roots0	0.014	NA	NA			Initial root sugar
						concentration
sugar.xylem.sh0	0.0249	NA	NA			Initial shoot xylem
						sugar concentration
sugar.xylem.st0	0.021	NA	NA			Initial stem xylem
						sugar concentration
Wala	0.0	0.0	0.0			Wala parameter
carbon.sugar	0.4211	0.4211	NA			Carbon content in
						sugar
carbon.starch	0.4444	0.4444	NA			Carbon content in
						starch
alfa	3	3	3			Alfa parameter
tau.s	2	2	2			Tau s parameter
tau.t	2	2	2			Tau t parameter
starch00	0.3246781	0.40	0.40			Secondary initial starch concentration

Table 12 continued from previous page  $\,$ 

Parameter Name	Parameter Name Parameter value			Units	Comments
ranicoer ivanic	Hyde	Lettosuo	Flakaliden_dHF_China	011163	Comments
sugar00	0.4184208	0.35	0.35		Secondary initial sugar concentration
Q10s	3	NA	NA		Q10 for synthesis
Q10d	1.8	NA	NA		Q10 for decomposition
$\operatorname{SCb}$	0.23	NA	NA	kg C	Storage carbon baseline
sugar.level	0.41	NA	0.35	kg C	Sugar level at senescence
Ad0.needles	0.017	NA	NA		Initial needle Ad0
Ad0.phloem	0.008	NA	NA		Initial phloem Ad0
Ad0.roots	2e-04	NA	NA		Initial root Ad0
Ad0.xylem.sh	2e-04	NA	NA		Initial shoot xylem Ad0
Ad0.xylem.st	0.047	NA	NA		Initial stem xylem Ad0
lamda.needles	0.197	NA	NA		Needle lamda parameter
lamda.phloem	0.05301	NA	NA		Phloem lamda parameter
lamda.roots	0.211	NA	NA		Root lamda parameter
lamda.xylem.sh	0.00401	NA	NA		Shoot xylem lamda parameter
lamda.xylem.st	0.00401	NA	NA		Stem xylem lamda parameter
delta.needles	0.729	NA	NA		Needle delta parameter
delta.phloem	0.832	NA	NA		Phloem delta parameter
delta.roots	0.853	NA	NA		Root delta parameter
delta.xylem.sh	0.762	NA	NA		Shoot xylem delta parameter
delta.xylem.st	0.294	NA	NA		Stem xylem delta parameter
$k_{-}np$	0.3	NA	NA		k_np parameter
k_pr	0.072	NA	NA		k_pr parameter
k_pxsh	0.188	NA	NA		k_pxsh parameter
k_pxst	0.17	NA	NA		k_pxst parameter
myco.thresh	0.025	NA	NA		Mycorrhiza threshold

### References

- Preetisri Baskaran, Riitta Hyvönen, S Linnea Berglund, Karina E Clemmensen, Göran I Ågren, Björn D Lindahl, and Stefano Manzoni. Modelling the influence of ectomycorrhizal decomposition on plant nutrition and soil carbon sequestration in boreal forest ecosystems. New Phytologist, 213(3):1452–1465, 2017.
- Joana Bergmann, Alexandra Weigelt, Fons van Der Plas, Daniel C Laughlin, Thom W Kuyper, Nathaly Guerrero-Ramirez, Oscar J Valverde-Barrantes, Helge Bruelheide, Grégoire T Freschet, Colleen M Iversen, et al. The fungal collaboration gradient dominates the root economics space in plants. *Science Advances*, 6(27):eaba3756, 2020.
- Róbert Blaško, Lisbet Holm Bach, Stephanie A Yarwood, Susan E Trumbore, Peter Högberg, and Mona N Högberg. Shifts in soil microbial community structure, nitrogen cycling and the concomitant declining n availability in ageing primary boreal forest ecosystems. *Soil Biology and Biochemistry*, 91:200–211, 2015.
- Bettina Brandes, Douglas L Godbold, Arnd J Kuhn, and Georg Jentschke. Nitrogen and phosphorus acquisition by the mycelium of the ectomycorrhizal fungus paxillus involutus and its effect on host nutrition. *The New Phytologist*, 140(4):735–743, 1998.
- Karina E Clemmensen, Roger D Finlay, Anders Dahlberg, Jan Stenlid, David A Wardle, and Björn D Lindahl. Carbon sequestration is related to mycorrhizal fungal community shifts during long-term succession in boreal forests. *New Phytologist*, 205(4):1525–1536, 2015.
- Michael C Dietze, Anna Sala, Mariah S Carbone, Claudia I Czimczik, Joshua A Mantooth, Andrew D Richardson, and Rodrigo Vargas. Nonstructural carbon in woody plants. *Annual review of plant biology*, 65(1):667–687, 2014.
- Yiyang Ding, Pauliina Schiestl-Aalto, Heljä-Sisko Helmisaari, Naoki Makita, Kira Ryhti, and Liisa Kulmala. Temperature and moisture dependence of daily growth of scots pine (pinus sylvestris l.) roots in southern finland. *Tree Physiology*, 40(2):272–283, 2020.
- Alf Ekblad, Håkan Wallander, Douglas L Godbold, C Cruz, D Johnson, P Baldrian, RG Björk, Daniel Epron, B Kieliszewska-Rokicka, R Kjøller, et al. The production and turnover of extramatrical mycelium of ectomycorrhizal fungi in forest soils: role in carbon cycling. *Plant and Soil*, 366(1):1–27, 2013.
- Oskar Franklin, Torgny Näsholm, Peter Högberg, and Mona N Högberg. Forests trapped in nitrogen limitation—an ecological market perspective on ectomycorrhizal symbiosis. *New Phytologist*, 203(2):657–666, 2014.
- Samia Ghersheen, Stefano Manzoni, Marie Spohn, and Björn D Lindahl. Modelling mycelial responses to nitrogen limitation during litter decomposition. *Available at SSRN 5107467*.
- Erik A Hobbie. Carbon allocation to ectomycorrhizal fungi correlates with belowground allocation in culture studies. *Ecology*, 87(3):563–569, 2006.

- Mona N Högberg, Maria JI Briones, Sonja G Keel, Daniel B Metcalfe, Catherine Campbell, Andrew J Midwood, Barry Thornton, Vaughan Hurry, Sune Linder, Torgny Näsholm, et al. Quantification of effects of season and nitrogen supply on tree below-ground carbon transfer to ectomycorrhizal fungi and other soil organisms in a boreal pine forest. *New Phytologist*, 187(2): 485–493, 2010.
- Torsten Ingestad, Agneta S Arveby, and Monika Käfar. The influence of ectomycorrhiza on nitrogen nutrition and growth of pinus sylvestris seedlings. *Physiologia Plantarum*, 68(4): 575–582, 1986.
- Nancy Collins Johnson, Jason D Hoeksema, James D Bever, V Bala Chaudhary, Catherine Gehring, John Klironomos, Roger Koide, R Michael Miller, John Moore, Peter Moutoglis, et al. From lilliput to brobdingnag: extending models of mycorrhizal function across scales. Bioscience, 56(11):889–900, 2006.
- Jaideep Joshi, Benjamin D Stocker, Florian Hofhansl, Shuangxi Zhou, Ulf Dieckmann, and Iain Colin Prentice. Towards a unified theory of plant photosynthesis and hydraulics. *Nature Plants*, 8(11):1304–1316, 2022.
- Pirjo Kainulainen and JK Holopainen. Concentrations of secondary compounds in scots pine needles at different stages of decomposition. Soil Biology and Biochemistry, 34(1):37–42, 2002.
- Deliang Kong, Junjian Wang, Huifang Wu, Oscar J Valverde-Barrantes, Ruili Wang, Hui Zeng, Paul Kardol, Haiyan Zhang, and Yulong Feng. Nonlinearity of root trait relationships and the root economics spectrum. *Nature communications*, 10(1):2203, 2019.
- Kris R Kramer-Walter, Peter J Bellingham, Timothy R Millar, Rob D Smissen, Sarah J Richardson, and Daniel C Laughlin. Root traits are multidimensional: specific root length is independent from root tissue density and the plant economic spectrum. *Journal of Ecology*, 104 (5):1299–1310, 2016.
- Björn D Lindahl, Katarina Ihrmark, Johanna Boberg, Susan E Trumbore, Peter Högberg, Jan Stenlid, and Roger D Finlay. Spatial separation of litter decomposition and mycorrhizal nitrogen uptake in a boreal forest. *New phytologist*, 173(3):611–620, 2007.
- Björn D Lindahl, Wietse De Boer, and Roger D Finlay. Disruption of root carbon transport into forest humus stimulates fungal opportunists at the expense of mycorrhizal fungi. *The ISME journal*, 4(7):872–881, 2010.
- Björn D Lindahl, Julia Kyaschenko, Kerstin Varenius, Karina E Clemmensen, Anders Dahlberg, Erik Karltun, and Johan Stendahl. A group of ectomycorrhizal fungi restricts organic matter accumulation in boreal forest. *Ecology Letters*, 24(7):1341–1351, 2021.
- Annikki Mäkelä, Minna Pulkkinen, Pasi Kolari, Fredrik Lagergren, Paul Berbigier, Anders Lindroth, Denis Loustau, Eero Nikinmaa, Timo Vesala, and Pertti Hari. Developing an empirical model of stand gpp with the lue approach: analysis of eddy covariance data at five contrasting conifer sites in europe. *Global change biology*, 14(1):92–108, 2008.

- Horst Marschner, Max Häussling, and Eckhard George. Ammonium and nitrate uptake rates and rhizosphere ph in non-mycorrhizal roots of norway spruce [picea abies (l.) karst.]. *Trees*, 5: 14–21, 1991.
- ROSS E McMURTRIE. Relationship of forest productivity to nutrient and carbon supply—a modeling analysis. *Tree Physiology*, 9(1-2):87–99, 1991.
- Astrid Meyer, Rüdiger Grote, Andrea Polle, and Klaus Butterbach-Bahl. Simulating mycorrhiza contribution to forest c-and n cycling-the mycofon model. *Plant and soil*, 327:493–517, 2010.
- F Minunno, Mikko Peltoniemi, Samuli Launiainen, M Aurela, A Lindroth, A Lohila, I Mammarella, K Minkkinen, and A Mäkelä. Calibration and validation of a semi-empirical flux ecosystem model for coniferous forests in the boreal region. *Ecological Modelling*, 341:37–52, 2016.
- Torgny Näsholm, Knut Kielland, and Ulrika Ganeteg. Uptake of organic nitrogen by plants. *New phytologist*, 182(1):31–48, 2009.
- Torgny Näsholm, Peter Högberg, Oskar Franklin, Daniel Metcalfe, Sonja G Keel, Catherine Campbell, Vaughan Hurry, Sune Linder, and Mona N Högberg. Are ectomycorrhizal fungi alleviating or aggravating nitrogen limitation of tree growth in boreal forests? *New Phytologist*, 198(1):214–221, 2013.
- Ivika Ostonen, Krista Lõhmus, Heljä-Sisko Helmisaari, Jaak Truu, and Signe Meel. Fine root morphological adaptations in scots pine, norway spruce and silver birch along a latitudinal gradient in boreal forests. *Tree Physiology*, 27(11):1627–1634, 2007.
- Olusegun Ayodeji Oyewole, Erich Inselsbacher, Torgny Näsholm, and Sandra Jämtgård. Incorporating mass flow strongly promotes n flux rates in boreal forest soils. *Soil Biology and Biochemistry*, 114:263–269, 2017.
- Mikko Peltoniemi, Minna Pulkkinen, Mika Aurela, Jukka Pumpanen, Pasi Kolari, and Annikki Mäkelä. A semi-empirical model of boreal-forest gross primary production, evapotranspiration, and soil water-calibration and sensitivity analysis. *Luke Open Documents*, 2015.
- Nazia Perveen, Sébastien Barot, Gaël Alvarez, Katja Klumpp, Raphael Martin, Alain Rapaport, Damien Herfurth, Frédérique Louault, and Sébastien Fontaine. Priming effect and microbial diversity in ecosystem functioning and response to global change: a modeling approach using the symphony model. *Global change biology*, 20(4):1174–1190, 2014.
- Jaakko Repola. Biomass equations for scots pine and norway spruce in finland. 2009.
- Kira Ryhti, Liisa Kulmala, Jukka Pumpanen, Jarkko Isotalo, Mari Pihlatie, Heljä-Sisko Helmisaari, Jaana Leppälammi-Kujansuu, Antti-Jussi Kieloaho, Jaana Bäck, and Jussi Heinonsalo. Partitioning of forest floor co2 emissions reveals the belowground interactions between different plant groups in a scots pine stand in southern finland. *Agricultural and Forest Meteorology*, 297:108266, 2021.
- Kira Ryhti et al. Belowground carbon dynamics in scots pine stands. *Dissertationes Forestales*, 2022.

- Pauliina Schiestl-Aalto, Liisa Kulmala, Harri Mäkinen, Eero Nikinmaa, and Annikki Mäkelä. Cassia—a dynamic model for predicting intra-annual sink demand and interannual growth variation in s cots pine. *New Phytologist*, 206(2):647–659, 2015.
- Pauliina Schiestl-Aalto, Kira Ryhti, Annikki Mäkelä, Mikko Peltoniemi, Jaana Bäck, and Liisa Kulmala. Analysis of the nsc storage dynamics in tree organs reveals the allocation to belowground symbionts in the framework of whole tree carbon balance. Frontiers in Forests and Global Change, 2:17, 2019.
- Sally E Smith and David J Read. Mycorrhizal symbiosis. Academic press, 2010.
- Or Sperling, Tamir Kamai, Aude Tixier, Anna Davidson, Katherine Jarvis-Shean, Eran Raveh, Ted M DeJong, and Maciej A Zwieniecki. Predicting bloom dates by temperature mediated kinetics of carbohydrate metabolism in deciduous trees. *Agricultural and Forest Meteorology*, 276:107643, 2019.
- Erica Sterkenburg, Adam Bahr, Mikael Brandström Durling, Karina E Clemmensen, and Björn D Lindahl. Changes in fungal communities along a boreal forest soil fertility gradient. *New Phytologist*, 207(4):1145–1158, 2015.
- Benjamin N Sulman, Richard P Phillips, A Christopher Oishi, Elena Shevliakova, and Stephen W Pacala. Microbe-driven turnover offsets mineral-mediated storage of soil carbon under elevated co2. *Nature Climate Change*, 4(12):1099–1102, 2014.
- Sanna Susiluoto, Emmi Hilasvuori, and Frank Berninger. Testing the growth limitation hypothesis for subarctic scots pine. *Journal of Ecology*, 98(5):1186–1195, 2010.
- Xianglin Tian, Francesco Minunno, Pauliina Schiestl-Aalto, Jinshu Chi, Peng Zhao, Matthias Peichl, John Marshall, Torgny Näsholm, Hyungwoo Lim, Mikko Peltoniemi, et al. Disaggregating the effects of nitrogen addition on gross primary production in a boreal scots pine forest. Agricultural and Forest Meteorology, 301:108337, 2021.
- Oscar J Valverde-Barrantes, Grégoire T Freschet, Catherine Roumet, and Christopher B Blackwood. A worldview of root traits: the influence of ancestry, growth form, climate and mycorrhizal association on the functional trait variation of fine-root tissues in seed plants. *New Phytologist*, 215(4):1562–1573, 2017.
- Anouk Van't Padje, Gijsbert DA Werner, and E Toby Kiers. Mycorrhizal fungi control phosphorus value in trade symbiosis with host roots when exposed to abrupt 'crashes' and 'booms' of resource availability. *New Phytologist*, 229(5):2933–2944, 2021.
- T Wallenda and DJ Read. Kinetics of amino acid uptake by ectomycorrhizal roots. Plant, Cell & Environment, 22(2):179-187, 1999.