LUKE HEATON - NICK JONES - MARK FRICKER

FUNGAL T2M -SOFTWARE MANUAL

PLANT SCIENCES, OXFORD FIRST EDITION

Copyright © 2019 Luke Heaton - Nick Jones - Mark Fricker

All rights reserved.

Redistribution of this manual and the associated software and use in source and binary forms, with or without modification, are permitted provided that the following conditions are met:

- Redistributions of source code must retain the above copyright notice, this list of conditions and the following disclaimer.
- Redistributions in binary form must reproduce the above copyright notice, this list of conditions and the following disclaimer in the documentation and/or other materials provided with the distribution.
- Neither the name of Plant Sciences, University of Oxford nor the names of its contributors may be used to endorse or promote products derived from this software without specific prior written permission.

This software is provided by the copyright holders and contributors "as is" and any express or implied warranties, including, but not limited to, the implied warranties of merchantability and fitness for a particular purpose are disclaimed. In no event shall Mark Fricker be liable for any direct, indirect, incidental, special, exemplary, or consequential damages (including, but not limited to, procurement of substitute goods or services; loss of use, data, or profits; or business interruption) however caused and on any theory of liability, whether in contract, strict liability, or tort (including negligence or otherwise) arising in any way out of the use of this software, even if advised of the possibility of such damage.

PUBLISHED BY PLANT SCIENCES, OXFORD FIRST EDITION

First printing, November 2019

Current version, January 2020

Contents

1	Inst	allation	5
	1.1	Overview	5
	1.2	Download all files from the github repository	5
	1.3	Installation of the MATLAB app	6
	1.4	Installation of the stand-alone program	6
	1.5	Installation of additional program files needed	8
2	Rur	nning the simulation using the GUI	9
	2.1	Introduction	9
	2.2	Environmental parameters	11
	2.3	Organism parameters	11
	2.4	Running the simulation	11
	2.5	Display	12
	2.6	Performance of different classes of organism on specific resources	14
	2.7	Inclusion of user defined resources	15
3	Con	nmand Line Version	17
	3.1	Overview	17
	3.2	Running the simulation for varying resource quality and recalcitrance	•
	3.3	Running the simulation for a specific resource quality in varying abundance	

4 LUKE HEATON - NICK JONES - MARK FRICKER

Acknowledgements

This work was supported by a Visiting Fellowship at The Institute of Advanced Studies in Durham, The Leverhulme Trust (RPG-2015-437) and The Human Frontier Science Program (RGP0053/2012).

The export_fig package was written by Oliver Woodford and Yair Altman and downloaded from: https://uk.mathworks.com/matlabcentral/fileexchange/23629-export-fig

The colorcet.m package was written by Peter Kovesi and downloaded from: https://peterkovesi.com/projects/colourmaps/

Peter Kovesi. Good Colour Maps: How to Design Them. arXiv:1509.03700 [cs.GR] 2015 https://arxiv.org/abs/1509.03700

Installation

1.1 Overview

The code to run the simulation is provided as open source under a GNU General Public License v3.0 from:

https://github.com/markfricker/Fungal_multicellularity

The code is provided in three formats:

- A set of script files and functions that will run in MATLAB® on any platform;
- A MATLAB® app with a GUI interface that can be installed in MATLAB on any platform (requires release 2019b or later);
- A compiled standalone version that can be installed and run on Windows 64 bit platforms;

1.2 Download all files from the github repository

On the home page of the github repository, click on the green Clone or download button (Figure 1.1) and then the blue Download ZIP button.

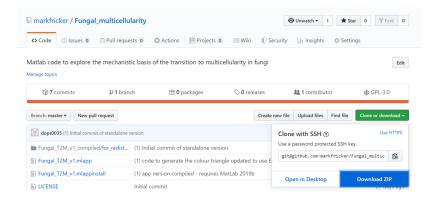


Figure 1.1: Download the software in a zip file from the github repository

Once the file has downloaded, extract all the files. If you intend to run the script versions, this folder needs to be on the MATLAB path.

1.3 Installation of the MATLAB app

The MATLAB[®] app installer file, .mlappinstall, contains everything necessary to install and run an app within the MATLAB[®] environment, including the source code, supporting data, information (such as product dependencies), and the app icon (Figure 1.2).

Double-clicking on the .mlappinstall file should launch MATLAB if it is not already running, and install the app in the app toolbar (Figure 1.3). The program can be run, by clicking on the icon in the toolbar.

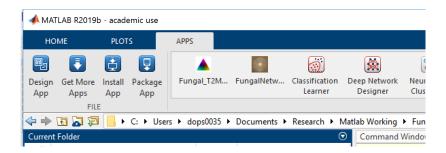




Figure 1.2: The program icon

Figure 1.3: Location of the Fungal_T2M app in the APPS menu within MATLAB

The app version of the program also requires the MATLAB Image Processing Toolbox TM to be installed.

1.4 Installation of the stand-alone program

The software has been tested on Windows 10, and requires a minimum screen resolution of 1600 x 900. In addition, an appropriate version of the MATLAB Compiler Runtime (MCR) is required to install the set of shared libraries that enables execution of the compiled MATLAB application. The MCR should automatically download from the MathWorks website when the program is installed for the first time. Alternatively MCR can be downloaded from the MathWorks Website:

http://www.mathworks.com/products/compiler/mcr.

To install the MCR and standalone package, double-click the compiled MATLAB self-extracting *.exe file located in the subfolder:

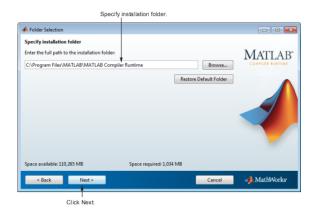
\Fungal_T2M_v1_compiled\for_redistribution

This extracts the MATLAB Runtime Installer from the archive, along with all the files that make up the deployed MATLAB environment. Once all the files have been extracted, the MATLAB Runtime Installer starts automatically. When the MATLAB Runtime Installer starts, it displays the following dialog box. Read the information and then click **Next** to proceed with the installation.

Specify the folder in which you want to install the MATLAB runtime in the Folder Selection dialog box and click **Next**. It is recommended to keep the default settings as this ensures the path to other program files is set automatically.



Note: On Windows systems, you can have multiple versions of the MATLAB runtime on your computer, but only one installation for any particular version. If you already have an existing installation, the MATLAB runtime Installer does not display the Folder Selection dialog box because you can only overwrite the existing installation in the same folder.



Confirm your choices and click Install. The MATLAB Runtime Installer starts copying files into the installation folder



Click Finish to exit the installer.



MATLAB Runtime Installer Readme File: A readme.txt file is included with the MATLAB Runtime Installer. This file, visible when the MATLAB Runtime Installer is expanded, provides more detailed information about the installer and the switches that can be used with it.

1.5 Installation of additional program files needed

A number of additional files needed to run the full suite of programs may also be installed at the same time as the main program. The latest version of Java needs to be installed, and is available from:

http://www.java.com/en/

Output of images at full resolution uses *export_fig.m* originally written by Oliver Woodford (2008-2014) and now maintained by Yair Altman (2015-). When exporting to vector format (PDF or EPS) this function requires that ghostscript is installed on your system. Ghostscript can be downloaded from:

http://www.ghostscript.com.

When exporting images to eps and pdf formats, <code>export_fig</code> additionally requires pdftops, from the Xpdf suite of functions. This is included in the xpdf tools package and can be downloaded from:

https://www.xpdfreader.com/download.html

Running the simulation using the GUI

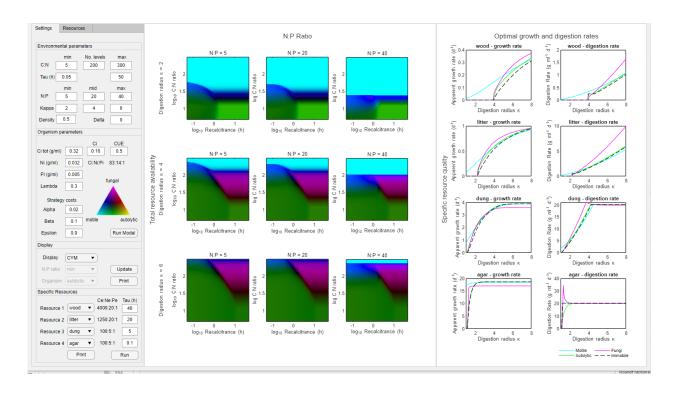


Figure 2.1: The GUI interface for the fungal multicellularity simulation

2.1 Introduction

The MATLAB app version and the standalone version both provide the same interface to run the simulation (Figure 2.1). The set of left-hand panels allow the user to change the parameters controlling the external resource environment and the internal organism parameters, including the cost terms for each of the different strategies. The definition of the parameters is given in Table 2.1.

Tunable Environmental Parameters	Symbol and Units	Interpretation	
Turumeters	$C_{\text{E}}, N_{\text{E}}, P_{\text{E}}$	Grams of C, N and P per millilitre of substrate,	
Supply of C, N and P		set by the C:N and N:P ratio of the resource,	
		and the resource density in g ml ⁻¹	
	au hours	Time required for an exoenzyme to supply a	
Recalcitrance		mass of C, N or P equal to the total mass re-	
		quired to synthesise the exoenzyme in question	
Polative digestion radius	К	Cylindrical cells and hyphae digest resource	
Relative digestion radius	K.	over a distance times the cell radius	
		Ratio of C that has to be digested to release	
Resource accessibility	δ	each N, to reflect that N is embedded within	
		C-rich polymers	

Organism Parameters			
Core demand for C, N and P	$C_{I} = 0.33 \text{ g ml}^{-1}$ $N_{I} = 0.032 \text{ g ml}^{-1}$ $P_{I} = 0.005 \text{ g ml}^{-1}$	Core demand for C, N and P per unit volume of any organism	
Motility cost	$\alpha = 0.02$	Mass of C and N required to synthesise motile apparatus, relative to core demand for C and N	
Vesicle transport cost	$\beta=0.1$	Mass of C and N in vesicles, relative to the total C and N in exoenzymes and the fungal core	
Recycling efficiency	$\epsilon = 0.5$	Fraction of C, N and P that is recouped by autolysis	
Maximal rate of resource use	$\lambda = 0.3 \text{ g ml}^{-1} \text{ hour}^{-1}$	Maximal mass of resource any cell can use per unit time and volume	

Model Variables			
Relative density of exoen-	~	Mass of C and N used for exoenzymes relative	
zymes	X	to the core demand for C and N	
Relative density of	$x_{\rm C}, x_{\rm N}, x_{\rm P}$	Mass of C and N used for C, N or P digesting	
C, N or P		exoenzymes relative to the core demand for C	
C, N of F		and N	

Functions of <i>x</i>		
Time to exhaust local	T hours	Time until one of C, N or P is locally exhausted
resource		
Time to exhaust C, N or	$T_{\rm C}$, $T_{\rm N}$, $T_{\rm P}$ hours	Time until relevant element is locally exhausted
P		Time until relevant element is locally exhausted
Specific growth rate of	$\eta \text{ hour}^{-1}$	Rate of synthesis of cell biomass, per unit of
growing cell	η πουτ	cell biomass, in the growing cells
Ammanant amazyth mata	μ hour ⁻¹	Volumetric rate of colonisation, per unit vol-
Apparent growth rate		ume colonised
	Ω	Maximum number of daughter cells and their
Total limiting supply		exoenzymes that can be synthesised, given the
		local supply of resource
Data of macaumaa uga	$\Gamma \mathrm{g} \mathrm{ml}^{-1} \mathrm{hour}^{-1}$	Total mass of resource used per unit time and
Rate of resource use		volume, in the growing cells

Table 2.1: Definition and values of modelling parameters

Environmental parameters

The C:N ratio of the resource ranges from the minimum value (min), typically around 5, to the maximum value (max), with the number of intermediate levels set by levels (Figure 2.2).

This forms the y-axis for each of the resultant plots (on a log_{10} scale). The x-axis is set by the recalcitrance (τ) , and ranges from min to max with the same number of intermediate levels. Results are plotted for three values of the N:P ratio, set by min, mid and max and three values for the overall resource availability, given by the radius of digestion (κ), again set by *min*, *mid* and *max*.

The overall resource density in g ml⁻¹ is used to convert the C:N and C:P ratios into grams ml-1 of Ce, Ne, and Pe

The value of the additional accessibility parameter δ , reflects the fact that some fraction of the available C must be digested in order to access N, irrespective of the category of organism. For example, if the C:N ratio of the substrate is 200:1, imposing a value of $\delta = 0.1$ forces organisms to digest at least 20 C for every N they acquire.

Settings Resources Environmental parameters No. levels max C:N 5 200 300 Tau (h) 0.05 50 mid max 5 20 40 Kappa 2 4 6 Density 0.5 Delta 0

Figure 2.2: Controls for the environmental parameters that define the resource quantity and auality

2.3 Organism parameters

The internal C, N and P required by all organisms are set by C_i , N_i and P_i , respectively, in g ml⁻¹. These values are also displayed as the molar C:N:P ratio in the adjacent text box (Figure 2.3). The default values are typical of fungal and microbial cells, although there is considerable variation in both the absolute amounts and the relative ratios.

The total amount of C required (C_i tot) also includes that used in respiration. This is given as a proportion that depends on the carbon use efficiency (CUE), defined as the fraction of growth to total assimilation. The default value of CUE is set as 50%, but values can be much lower than this1.

The maximum rate that resource that can be used (by any organism) is set by Lambda, with a default value of g ml⁻¹ h⁻¹. This would equate to a doubling time of just under 1h for the default resource density, and represents an upper limit unlikely to be achieved for organisms in the wild.

In addition, autolytic, motile and fungal cells have a single organism-specific parameter that helps to define their different strategies. Thus, Alpha is the additional cost of being motile, Beta is the cost associated with internal transport, whilst Epsilon is the fraction of an autolytic cell that can be recycled.

Running the simulation 2.4

The Run Model button will iterate through the external resource environmental parameters for each organism, and returns pseudocolour coded maps showing which class of organism is predicted to grow most rapidly under each set of conditions (Figure 2.4),

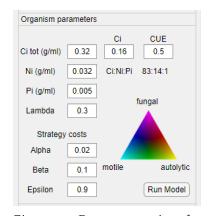


Figure 2.3: Parameter settings for the biological organisms

according to the inset colour triangle (Figure 2.5).

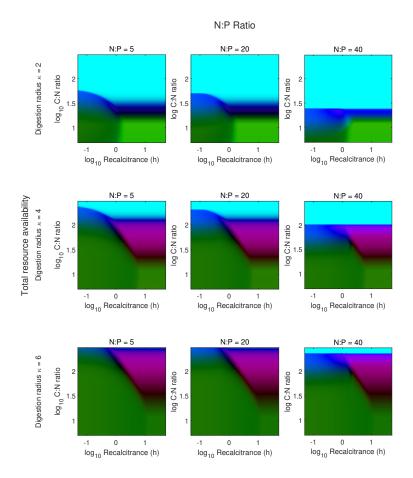


Figure 2.4: Colour-coded maps showing the relative performance of each type of organism across a wide set of resource quantity and quality.

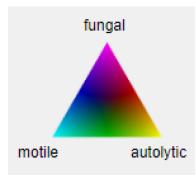


Figure 2.5: Legend colour code

Thus regions of parameter space where motile cells dominate are coloured cyan, regions where fungi dominate are magenta and regions where autolytic cells dominate are yellow. In many regions, two or more organisms may fare equally well, and there are tyically extensive regions where motile and fungal organisms both thrive (blue), or motile and autolytic are equivalent (yellow).

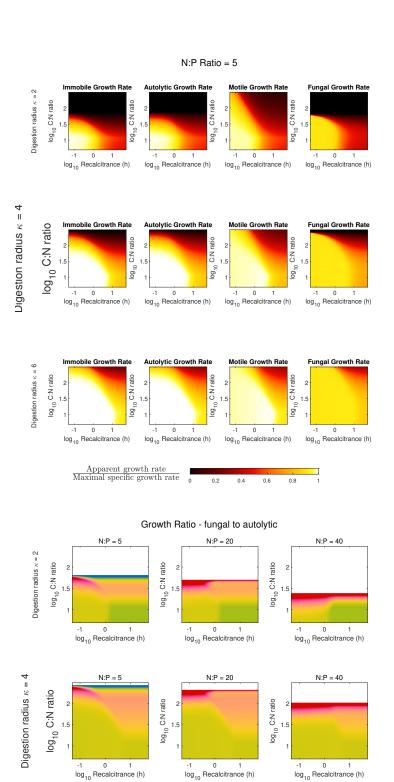
2.5 Display

The cyan-magenta-yellow maps show the relative performance of each class of organism compared to the others. In the *Display* panel (Figure 2.6), the *Display* drop down menu can be used to show the growth rates compared to the fastest growing cell by selecting the *growth* option (Figure 2.7). This provides plots for each organism for each digestion radius, whilst the *N:P ratio* has to be selected from the corresponding drop down menu.

Alternatively, the relative growth rate of a fungal organism can be compared with one of the other classes of organism to compare directly the magnitude of the hyphal advantage (Figure 2.8). The **Print** button saves a copy of the display as a png or pdf file.



Figure 2.6: Display options



log₁₀ Recalcitrance (h)

N:P = 20

-1 0 1 log₁₀ Recalcitrance (h)

 $\frac{10}{12}$ $\frac{10}{10}$

N:P = 5

-1 0 1 log₁₀ Recalcitrance (h)

 $\frac{\text{Hyphal growth rate}}{\text{Unicell growth rate}}$

log₁₀ C:N ratio

Digestion radius $\kappa=6$

log₁₀ C:N ratio

log₁₀ Recalcitrance (h)

N:P = 40

-1 0 1 log₁₀ Recalcitrance (h)

log₁₀ C:N ratio

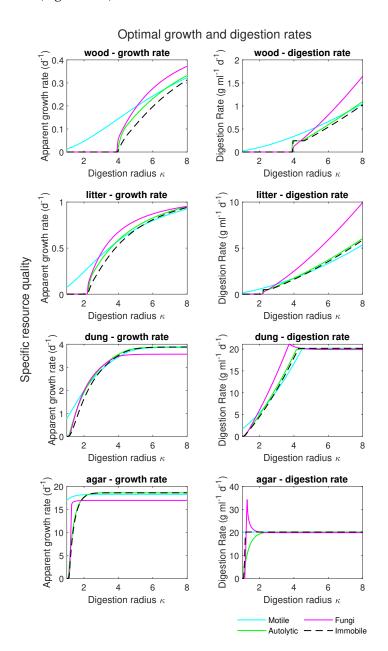
Figure 2.7: Colour-coded maps showing the relative growth of each type of organism compared to the maximum growth of any type of organism.

Figure 2.8: Colour-coded maps showing the relative growth of a hyphal organism to a specific class of unicellular organism.

2.6 Performance of different classes of organism on specific resources

The colour-coded maps display results for a very extensive range of resource environments. The *Specific resources* panel (Figure 2.9) provides options to plot the performance for four specific resources, with the defaults set to wood (C:N:P 4000:20:1, τ = 40h), leaf litter (C:N:P 1250:20:1, τ = 20), dung (C:N:P 100:5:1, τ = 5h) and agar (C:N:P 100:5:1, τ = 0.1h).

The Run button runs the simulation for these specific resource configurations using the other parameters defined for the full model (Figure 2.10)



Whilst the values for C:N:P are widely reported in the literature, the values for τ are not defined experimentally for most substrates.

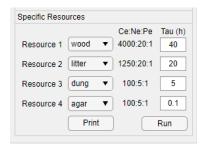


Figure 2.9: Display options

Figure 2.10: Colour-coded maps showing the relative growth of a hyphal organism to a specific class of unicellular organism.

Thus, there is the option to vary τ for each resource using the adjacent edit box. For direct comparison, the same resource can be selected in all four dropdown menus and the simulation run with different values of τ . Increasing the value of tau significantly decreases the growth rate for all organisms, but in general, changing tau only has a small effect on the relative performance of different classes of organisms.

Inclusion of user defined resources 2.7

Given that there are many different resource possibilities other than the four defaults used, there is an option to define more substrates using the Resources tab at the top of the control panels (Figure 2.11)

The Table includes the four default resource types. Additional resources can be included using the Add button, which adds an additional row to the table. Likewise the Remove button will delete the currently selected entry. The modified table can be saved using the Save button, and re-loaded using the Load button. The Update button adds the new set of resources to the Resource drop down menus in the **Specific Resource** panel in the main **Settings** tab. The effect of changing resources can then be explored using the Run button.

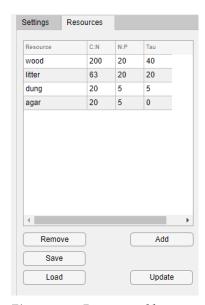


Figure 2.11: Resource table to allow addition of specific resources with user-defined C:N:P ratios and recalcitrance (Tau)

Command Line Version

3.1 Overview

The set of MATLAB® scripts that can be used to run the different components of the simulation are given in Table 3.1. These should work in any release of MATLAB® on any platform. They are the same functions as called by the GUI and standalone versions of the program.

Script Name	Description
vary_CNP_and_tau.m	The main script to run the simulation and generate colour-coded maps of relative performance
vary_kappa.m	Script to generate the growth rate and investment in digestion for each class of organism in a specific resource environment

Function Name	Description
find_best_immobile.m	Function to determine numerically the best growth rate (μ) for a given investment in digestive enzymes (x) using the parameter values for immobile cells. Calls the function $find_mu_given_x_immobile.m$ which implements equations 1-6 of the Supplementary Information.
find_best_autolytic.m	Function to determine numerically the best growth rate (μ) for a given investment in digestive enzymes (x) using the parameter values for autolytic cells. Calls the function $find_mu_given_x_autolytic.m$ which implements equations 7-10 of the Supplementary Information.
find_best_motile.m	Function to determine numerically the best growth rate (μ) for a given investment in digestive enzymes (x) using the parameter values for motile cells. Calls the function $find_mu_given_x_motile.m$ which implements equations 11-13 of the Supplementary Information.
find_best_fungi.m	Function to determine numerically the best growth rate (μ) for a given investment in digestive enzymes (x) using the parameter values for fungal cells. Calls the function $find_mu_given_U_fungi.m$ which implements equations 14-17 of the Supplementary Information.

find_best_cell.m	Function to determine numerically the best growth rate (μ) for a given
	investment in digestive enzymes (x) using the parameter values for a
	cell growing at the maximum possible rate without substrate depletion
	which implements equations 1-6 of the Supplementary Information.

Display functions	Description
generate_CYM_merge.m	Function to generate the Cyan-Magenta-Yellow maps of relative growth rate. If the fastest growing colony has grown by a factor of n, and a slower growing colony has grown by a factor of m, the channels for the colonies are coloured accordingly with intensity 1 and m/n respectively. Higher n increases the apparent difference in colour, making a lighter CYM merge.
heatmap_comparison.m	Generates colour-coded ratio map of the relative growth of fungal cells and autolytic cells in the standard set of resource environments. Calls <i>colorcet.m</i> to define a perceptually uniform ratio colourmap ('R3') and <i>applycolourmap.m</i> to apply it to the ratio growth image. See SI Figure 3.
heatmap_growth_inhibition.m	Generates heatmap showing the relative growth of each class of organism compared to the maximum growth possible with unlimited resource. Calls <i>colorcet.m</i> to define a perceptually uniform ratio colourmap ('L3') and <i>applycolourmap.m</i> to apply it to the relative growth inhibition image. See SI Figure 4.

Table 3.1: MATLAB scripts and functions

3.2 Running the simulation for varying resource quality and recalcitrance

Open the vary_CNP_and_tau.m script in the MATLAB® Editor. The first part of the code sets the resource and organism parameters (Listing 3.1), which can be modified manually to explore different environmental resources or impose different costs on the strategy used by each organism. For initial trials, the res1 parameter can be set to a much lower value (20) to reduce the computation time, whist still titrating the range of C:N and recalcitrance values (Figure 3.2).

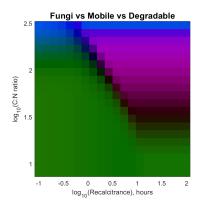


Figure 3.1: Coarse output for rapid prototyping

Listing 3.1: Setting parameter values at the start of the vary_CNP_and_tau code

```
%% set up parameters. These can be manually edited.
  % resl is the number of different values for the C:N ratio and recalcitrance
 % that are tried
_{4} res1 = 200;
```

^{6 %} xres determines the number of different values for x that are tried in finding

```
7 % the optimal solution. Final value found is accurate to xres^2.
s 	ext{ xres} = 200;
_{10} % res is the number of iterations used in finding best solution for a given
_{11} % environment and a given rate of sythesis.
12 \text{ sres} = 60;
% N_to_P set the N:P ratio of the resource.
N_{to} = 20;
  % kappa is the radius of digestion, normalised to the radous of the cells/hypha,
17
18 % and sets the total amount of resource available.
19 kappa = 6;
_{21} % C_to_N_min and C_to_N_max define the limits of the C:N ratio in the resource.
_{22} % The number of intermediate steps is controlled by resl. The C:N ratio is
23 % plotted on the y-axis of the resultnt map.
24 C_to_N_min = 8;
C_{5} C_{5} = 300;
_{27} % tau_min and tau_max define the limits of the resouce recalcitrance, where tau
{}_{28} % is the time in hours for hydrolases to digest their own mass . the number of
_{29} % intermediate steps is controlled by res1. The recalcitrance is plotted on the
30 % x-axis of the resultnt map.
31 tau_min = 0.1;
32 tau_max = 100;
_{34} % Ci, Ni and Pi are the mass of carbon, nitrogen and phosphorous, respectively,
35 % needed per unit volume of growth, in g per ml.
_{36} Ci = 0.33;
_{37} Ni = 0.032;
_{38} Pi = 0.005;
_{40} % dry weight in grams per ml of the substrate.
_{41} density = 0.5;
_{43} % epsilon is the efficiency of recycling for senscent autolytic cells.
_{44} epsilon = 0.5;
_{46} % alpha is the additional mass of machinery needed for cell mobility, relative
_{
m 47} % to the mass of essential metabolic machinery.
_{48} alpha = 0.02;
_{50} % beta is the mass of material in vesicles, relative to the mass of the rest of
_{51} % the fungus, including hydrolases.
<sub>52</sub> beta = 0.1;
54 % lambda is the maximum rate of resource use per unit volume, in g per ml per
55 % hour.
_{56} lambda = 0.3;
_{58} % _{\Delta} is the ratio of C that has to be digested to release each N, to reflect
59 % that N is embedded within C-rich polymers.
60 \quad \Delta = 0;
```

The output is an array of the relative growth (Mu) and the amount of digestive enzymes released (x) for each class of organism. The relative growth can be compared using the three different display scripts:

- generate_CYM_merge.m gives a three-way comparison using a cyan-magenta-yellow colour-code for motile, fungal and autolytic cells, respectively.
- *heatmap_growth_inhibition.m* compares the relative growth of each organism against a cell with no limitation on resource availability

such that the colony continues to grow at the maximum rate under the conditions selected.

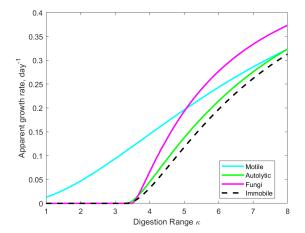
- heatmap_comparison.m compares the rate of fungal growth explicitly with autolytic cells to provide a more quantitative visual indication of the hyphal advantage in different resource environments.
- Running the simulation for a specific resource quality in varying abundance

The second main script (Listing 3.2) requires the user to define the resource quality (C:N:P ratios and recalcitrance), but then calculates the relative growth rate for each class of organism and the amount invested in digestive enzymes to give the optimal strategy. As with the vary_CNP_and_tau script, the first section has to be edited manually to set up different parameter configurations. The output is automatically generated as a plot of growth rate against resource abundance (varying kappa) and investment in digestive enzymes.

Listing 3.2: Setting parameter values at the start of the vary_kappa code

```
1 %% set up parameters. These can be manually edited.
2 % resl is the number of different values for kappa that we try
_{3} res1 = 200;
5 % xres is the number of iterations used in finding best solution for
6 % a given environment and a given rate of sythesis
_{7} xres = 200;
9 % sres is the number of iterations used in finding best solution for
10 % a given environment and a given rate of sythesis
sres = 60;
_{13} % k\_\text{min} and k\_\text{max} are the minimum and maximum values for the resource
% acquisition length that we try
_{15} k_min = 1;
_{16} k_max = 8;
_{18} % Ci, Ni and Pi are the mass of carbon, nitrogen and phosphorous, respectively,
19 % needed per unit volume of growth, in g per ml.
20 Ci = 0.33;
Ni = 0.032;
Pi = 0.005;
24 % C_to_P and N_to_P set the C:P and N:P ratio of the resource, respectively.
C_{to} = 2000;
_{26} N_to_P = 10;
{}_{28} % tau is the time in hours for hydrolases to digest their own mass and reflects
29 % the recalcitrance of the resource.
32 % dry weight in grams per ml of the substrate
_{33} density = 0.5;
  % \Delta  is the minimal fraction of C that must be digested in order to
_{36} % digest the available N
_{37} \Delta = 0;
```

```
% epsilon is the efficiency of recycling for autolytic cells
   epsilon = 0.5;
41
  % alpha is the mass of machinery needed for cell mobility, relative
42
  % to the mass of essential machinery
43
_{44} alpha = 0.02;
45
  % beta is the mass of material in vesicles, relative to the mass of
46
  % the rest of the fungus, including hydrolases
47
  beta = 0.1;
48
  % lambda is the maximum rate of resource use per unit volume,
50
  % in g per ml per hour
_{52} lambda = 0.3;
```



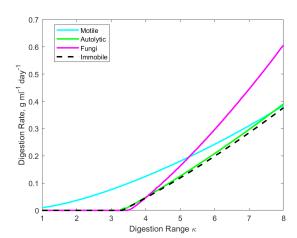


Figure 3.2: Growth rate and digestion rate for a specific resource quality and quantity