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1 Introduction

Decomposition by fungal and microbial communities is second only to photosynthesis in driving carbon cycling in the Earth's ecological communities [14]. While this significance is rarely understated, inner mechanisms and interactions within decomposition communities have been poorly understood and treated as a 'black box' in the past by scientists studying ecosystem processes [Andren1999].

Fungi are made up of hyphae: long winding filaments enmeshed together to form a thick mat of mycelium. At the frontline of battle, hyphal tips are extending forward and branching to cover new area at a velocity dictated by the hyphal extension rate, and concurrently other hyphae are dying off or merging together [3]. Like machinery, each fungus has ideal operating conditions. Fungi native to dry or arid biomes will grow efficiently in a moisture-lacking environment when other fungi will falter. Maynard et al. (2019) defines these operating conditions by assigning temperature and moisture niche-widths to 37 studied species of fungi.

1.1 Competition

When fungi come into contact with each other, competitive interactions are most often observed. Tactics studied in Boddy, L. (2000) include deploying diffusible toxins, obtaining nutrients parasitically, destruction of hyphae by intentional interference, and establishing wall-like barricades to thwart attackers [8]. Most researchers, including Maynard et al. (2019) construct a competitive ranking scale based on experimental observations to encapsulate the ramifications of these complex interactions.

Under a constant temperature regime, previous research concludes that dominant fast-growing suppress other species which negatively impacts the overall community's efficiency. However fluctuating environmental conditions change the outcome and promote complementary growth [12]. An explanation for this is presented by Lusten-

houwer et al. (2020) suggesting a spectrum between slow-growing, stress-tolerant fungi operating in a wider environmental range than fast-growing, highly competitive fungi [21].

1.2 Substrates

Determining which fungi are present in a community depends on the organic material, or 'substrate', being broken down. In the case of fungal-mediated wood decomposition, the substrate is made up of lignocellulose polysaccharides, and the fungi involved are equipped with specialized enzymes capable of splitting linkages in the polymers to harvest usable carbohydrates. Long-term wood decomposition follows a well-studied three-phase process: the first phase is dominated by fast and opportunistic fungus claiming easily-consumable soluble sugars, followed in the second phase by consistent and dependable decomposers that degrade the holocellulose constituting 65%-85% of the material [15]. Slow and highly specialized fungi trail behind in the third phase to clean up rigid lignin polymers (15%-35%) with powerful oxidative enzymes [11]. While phase progression is an important agent governing community changes on the long-term scale, this paper will focus on only the middle phase of cellulose decomposition and will not consider lignin decomposition. Fungi containing larger proportions of enzymes like acid phosphatase, which are associated with lignin decomposition, are demonstrated to strongly correlate with slower decomposition rates by Lustenhouwer et al. (2020)

1.3 Growth Model

An important step in modeling fungal communities is acknowledging separation between the growth rate and the decomposition rate of the fungus, which are influenced by different factors. Fungal growth models typically focus on the increase in overall biomass due to hyphal extension. Fungi with a greater density of hyphal tips per unit volume will grow and accrue biomass faster, as hy-

phae only grow from their tips. New tips are created when hyphae branch off, while others die or fuse together via anastomosis. Edelstein (1982) models these dynamics with the following:

$$\frac{\partial \rho}{\partial t} = n\nu - d \quad (1)$$

$$\frac{\partial n}{\partial t} = -\frac{\partial n\nu}{\partial x} + \sigma \quad (2)$$

These equations will operate standing alone, without any other influences by enzymes or growth limitations. This allows many models to treat it as a baseplate to expand upon. Edelstein (1983) and Davidson et al. (1997) do this by considering concentrations of nutrients inside the fungi and outside in the surrounding substrate. Our model will likewise follow in eliminating the assumption that fungi grow over an unlimited nutrient supply. A more contemporary approach by Du et al. (2019) describes the growth of a fungus with a 3-dimensional reaction-diffusion equation based on microscopic growth mechanisms, also focusing on tip density and extension rate but with an added term to identify the proportion of tips which are actively growing.

1.4 Environmental Influence on Hyphal Tip Extension (ν)

Environmental impacts are introduced to the decomposition model through ν , the hyphal tip elongation rate. The mechanism of elongation is relatively unknown[13]; As described in Gervais et al. (1999), "cell turgor pressure corresponds to an overpressure which allows the cell morphology, elongation, division and hence the biomass evolution" [7]. In fungi, cell turgor itself is respondent to water potential gradients rather than to active transport within the cell, making it very environmentally dependent [7]. Thus, we have reason to examine environmental impacts on fungal decomposition rates by varying ν outside of baselines for given species in set locations.

This brings us to the issue of determining ν based on environmental factors. Experimental

data from Maynard et al. 2019 shows the relationship between ν and the environmental parameters of water potential (ψ) and temperature (T) [18]. Water potential as a parameter will take into account both moisture content to fungi as well as the availability of that moisture as dictated by the soil composition.

1.5 Decomposition kinetics

Moore et al. (2015), Edelstein (1983), Moorhead et al. (2006), Parnas (1974), and Schimel et al. (2003) describe the decomposition of a certain substrate by a microbial decomposer as a function of the substrate's concentration by the Michaelis-Menton equation [Moore][4][11][2][10]. Schimel et al. (2003) and Moorhead et al. (2006) assert that decomposition levels off at a maximum rate proportional to the concentration of enzymes acting on it [10][11]. The decomposition of carbon in a substrate is described by the following equation.

$$\frac{dC}{dt} = \frac{KBC}{K_e + C} \quad (3)$$

We make the assumption that the enzyme concentration is proportional to the concentration of biomass in our system represented by r multiplied by two environmental limiting factors in $K = rS_M S_T$ [10]. Moorhead (1990) uses these limiting factors to multiply the rates of carbon uptake from microorganisms, although the model used a simpler first order, linear equation [11]. Here we use it in a similar way in limiting our maximum decay rate. The factors, S_M and S_T are computed by the following equations:

$$S_M = \alpha_2 - \lambda \log_{10}(-\psi) \quad (4)$$

$$\log_{10}(S_T) = \frac{T - 25}{10} \log_{10}(Q) \quad (5)$$

2 Model Formulation

In all models, the terms for ν , σ are taken to be constants given from experimental data. [citations].

Lin et al. (2016) takes a different approach in reviewing previous literature to apply a model that closely matches experimental data, which we will take advantage of for our model.

2.1 Growth Model

In the review of various fungal growth models described in Lin et al. (2016), a specific method of modeling branching and hyphal death was found to match experimental observations in the widest variety of settings. Based on these results we will choose to implement the following relationships, comprising dichotomous branching, tip-hypha anastomosis, and hyphal death (or YHD).

$$\sigma = \alpha_1 * n - \mu n \rho \quad (6)$$

$$d = \gamma_1 \rho \quad (7)$$

2.2 Estimating ν For Various Environments

When considering a sampling of decomposition rates in various environment types, we must determine how to estimate temperate and water potentials that ground fungi would experience. In the case of determining ν , this requires finding projected temperature and water potential.

Amongst existing biome classification models, Whitaker's scheme [1] is perhaps the simplest, being particularly trait-based. Whitaker's scheme provides a layout of biomes based on mean annual temperature and mean annual precipitation. However, modern classification of biomes has drifted away from using these traits as definitive biome identifiers [16]. These traits alone do not define all features of concern to fungal growth, such as soil composition for example. In addition, by Whitaker's scheme, we find that a given biome can have a wide range of average annual temperatures (such as arid deserts ranging from about -10 to 30 °C) [1]. Thus, we can sample a range of temperature and moisture values to output various ν and then select certain regions to profile in order to gauge potential environments where fungi can decompose and interact.

The following are the specific environments selected to represent various biomes:

Biome	Specific Environment Selected
Dessert (Arid)	Sonoran Desert, USA
Grasslands/shrublands	(Semi-Arid), central Argentina
Temperate Forest	Sal Forests, central Himalayas
Boreal Forest (Arboreal)	Pine Forests, central Himalayas
Tropical Rain Forest	Tropical Forests, Barro Colorado Island, Panama

The following are the average annual temperatures of these biomes. Note that if ranges were given, the average of the maximum and minimum is the selected value.

Biome/ Environment	Regional Average Annual Temperature [°C]	Selected Value [°C]
Dessert (Arid)	10 to 20 [Davey2007]	15
Grasslands/shrublands (Semi-Arid)	15.3 [6]	15.3
Temperate Forest	-1.42 to 26.39 [20]	12.49
Boreal Forest (Arboreal)	-1.42 to 26.39 [20]	12.49
Tropical Rain Forest	23 to 32 [19]	27.5

Although water potentials can be approximated based on predictive models, the measure is best found experimentally from soil samples [17]. Ranges for moisture potential of these environments have been found experimentally from a variety of studies is shown in the following table. Note that single water potentials are selected for sampling, as we are aiming to compare discrete environmental conditions rather than create a complete span of environmental conditions. These values are the average of the ranges or seasonal values given.

Biome/ Environment	Water Potential Range [MPa]	Selected Value [MPa]
Dessert (Arid)	-4.0 to -5.0 MPa [5]	-4.5 MPa
Grasslands/shrublands (Semi-Arid)	-1.4 to -5.0 ¹ [6]	-3.2 MPa
Temperate Forest	-0.44 (Fall), -1.19 (Winter), -0.58 (Spring), -1.42 (Early Summer), -1.81 (Summer) [9]	-1.09 MPa
Boreal Forest (Arboreal)	-0.83 (Fall), -1.20 (Winter), -0.55 (Spring), -1.61 (Early Summer), -3.36 (Summer) [9]	-1.51 MPa
Tropical Rain Forest	-1.57 MPa to 0.00 Mpa [Kupers2019]	-0.79

Given these estimates, we can have a probable example combination of temperature and moisture potential in various environments. Note that these are not wholly representative configurations, but rather examples to provide insight into how fungal species with specific traits may respond in discrete and distinct conditions likely to exist.

We can then find ν using experimental data from Maynard et al 2019. ***** (talk about continuous data technique thingy in the extra methods info sheet)

2.3 Environmental Influence on Hyphal Tip Extension structure of fungi: hyphal extension rate, branching, tips, death, etc. - proportion of enzymes relative to the phases correlates with the speed of

2.4 Decomposition Kinetics

2.5 Enzyme Implementation

2.6 Moisture

¹Measurements taken November through January at 100 cm soil depth.

3 Pre-planning intro

The Five Core Tenants: 1. growth/Decomposition
continuous model - general explanation of the

decomposition and effectiveness in a certain phase
 - substrate talk - LCI, lignin inhibition of holocellulose, common substrate compositions, how soil is different - phases talk - reasoning behind choosing a continuous model

2. interactions between different fungi with differing characteristics - competitive interactions
 * - synergy under richness and fluctuating temperatures
 * - the more differing the characteristics of the fungi, the more likely they are to compete - the competitive ranking *

3. sensitivity to environmental fluctuations
 - slower-growth stress tolerant fungi vs fast growth more niche temp/moisture fungi determining which of these two wins out depends on how much the environment is fluctuating and what subset of fungi are present
 * - fluctuating temperatures promotes growth - moisture talk * - biome talk

4. biodiversity's impact on systems overall efficiency - richness promotes growth - competition promotes efficiency
 * - dominant species that 'wipe out' the community generally harm? *

5. effect on the carbon cycle - fungi/bacteria are the only life forms which can break down lignin and efficiency break down holocellulose litter — these polysaccharides make up a huge portion of the total ground litter biomass in the earth's forests

Biodiversity's impact on the system's overall efficiency: - the greater range you have of moisture/temperature niche widths, the more tolerant your fungal community will be overall to growing in any particular environmental because at least some constituents of the community will be close to idealized in that environment. - competition rank correlates heavily with hyphal extension rate — fungal communities with species who are highly competitive will go one of two ways: 1. the environmental conditions are ideal for the smaller niches of the highly competitive isolates, and so those isolates will dominate out over the other ones 2. the environmental conditions are not ideal for the highly competitive isolates, so the slower but more robust isolates will win out

we have a spearman correlation coefficient relating production of each enzyme with de-

composition rate — enzymes primarily involved with breaking down lignin (acid phosphatase) are strongly correlated with decreased decomposition rate.

Strong correlation that increase production of acid phosphatase results in decreased competitive rank, which makes sense because on average slower-growing fungi are less competitive than faster growing ones. Competitive rank has a 0.6 spearman correlation coefficient with growth performance

Assumption: enzymes treatment of the different portions of holocellulose are the same

Implementation: each enzyme has its production rate from the data, multiply the parameter by that production rate and sum them all together to get dC/dt

Recapitulating previous literature

Prediction: The slower stress tolerant fungi will have a share of enzymes that indicate their slow decomposition rate, but a moisture and temperature width that indicate their stress tolerance to environmental fluctuations.

The faster fungi will have a share of enzymes that indicate their fast decomposition rate, but a moisture and temperature width that indicates their lack of stress tolerance to environmental fluctuations.

4 Model formulation

Maynard et. al took data on the productive rates of 8 different enzymes (5 hydrolytic and 3 oxidative to cover the spectrum of the cellulose-degradation process) and used a standard clustering approach to determine 4 representative enzymes of the entire set: cellobiohydrolase, acid phosphatase, peroxidase and phenoloxidase. The data for production of these enzymes comes from their open-source github.

We're making the assumption that the production of each enzyme is equivalent to the amount of the enzyme present in the fungus. We took the production data and normalized it to z-score val-

ues. Each z-score was turned into a portion of that enzyme with $e_x = (z - \min) / (\max - \min)$ and then we divided each portion by the sum of all enzyme portions for each enzyme (e_x) / ($e_1 + e_2 + e_3 + e_4$). This gives us a representation of the proportion of each enzyme relative to other enzymes. We made the assumption that the total mass of those four enzymes for each fungus is the same, and the fraction of enzyme mass from total mass.

We made the assumption that the enzymes are not changing over time due to C/N changes or environmental fluctuations

The time scale for our model is following the experimental results from (paper cited by question) which indicates x% log decomposition after 3 years and y% decomposition of logs after 5 years for a wide sampling of fungus

4.1 Assumptions

- Growth
 - unlimited available resources (ground litter and woody fibers)
 - neglecting effects from direct interactions between fungi
 - PROBLEM: neglecting synergistic effects on growth/metabolic efficiency
 - non-exhaustive substrates
 - not considering N addition by anthropogenic nitrogen deposition
 - LCI < 0.7 at all times because we're focusing on wood fibers and we don't want lignin block holocellulose decay in bad ways. but we could use the formula
 - LCI is not changing over time anastomosis, the formation of reconnections.
 - dynamics regarding the rate of nutrient absorption and transport of metabolites within the mycelium are ignored or simplified
- growth dynamics incurred by the formation of concentric density rings are not considered <– this doesn't need to exist because we're considering growth in a 1D tube
- only considering one configuration of growth traits out of many possible options, that configuration being: tip-hypha anastomosis, dichotomous branching, and hyphal death
- decomposition coefficients are representative of those families of enzymes
- Decomposition Kinetics
 - Elements can be stacked in any configuration without structural limitations
- Temperature/Moisture effects
 - experimental data for water potentials in specific biomes/environments are representative of the conditions of those environments
- Interactions
 - Influences from wind are neglected
 - The drones are assumed to be unobstructed by terrain
 - The drones do not experience any malfunctions
 - The earth's curve is neglected

4.2 Flight Path Sub-model

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4.3 Packing Sub-model

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4.4 Storage Location Determination

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$$L_n = \sum_{i=1}^m p_i(x_i, y_i) / \sum_{i=1}^m p_i$$

where L_n = Location number within grouping
 n being number of distinct locations (1 to 3)
 i = hospitals in the group

p = package demand of hospital i

x, y = longitude, latitude of hospital i

The output lists a discrete set of groupings of the hospitals with the corresponding locations of the storage containers for each hospital group. In figure 1, one potential combination of hospitals grouped to specific storage locations is shown.

4.5 Cost Function

I've copied the ten diff eqs from Numerical Prediction Model for Fungal Growth Coupled with Hygrothermal Transfer in Building Materials here: hygrothermal equations:

$$\frac{dH}{dT} \cdot \frac{\partial T}{\partial t} = \nabla \cdot (\lambda \nabla T) + h_v \nabla \cdot (\delta_p \nabla (\phi p_{sat})) \quad (8)$$

$$\frac{dw}{d\phi} \cdot \frac{\partial \phi}{\partial t} = \nabla \cdot (D_\phi \nabla \phi + \delta_p \nabla (\phi p_{sat})) \quad (9)$$

Chapter 15 Mathematical Modeling of Fungal Growth

$$\frac{\delta \rho}{\delta t} = nv - d(\rho) \quad (10)$$

$$\frac{\delta n}{\delta t} = \frac{\delta nv}{\delta x} + \sigma(\rho, n) \quad (11)$$

Where P represents the total medpacks delivered, $\sum t$ is an estimate of the time for all the flights to occur, and S represents the space left after packing all the drones (computed via the packing algorithm). Also, C is our cost function output, the cost. The factor of 100000 dividing S is there to adjust the units of S (it being on the order of 10^5 while $\frac{P}{\sum t}$ is on the order of 10^0) The time estimates are computed via the basic kinematic equation assuming constant speed, $\sum t = \sum \frac{d}{v_d}$ summed over all drone flights in the given plan. Here d is distance traveled in a specified flight and v_d is the max speed of the drone flying. This estimate of the time taken for a flight is assuming the drone is flying at max speed the whole way, and assuming equality of the time taken to fly to the hospital, and

the time to fly back. The assumption of perfectly sequential ordering of the flights allows us to sum the times of individual flights to get the total time.

5 Results and Analysis

5.1 Model output

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5.2 Parameter sensitivity

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5.3 Limitations/ Further Work

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