1	A metabolic perspective of stochastic community assembly		
2			
3	Victor S. Saito ¹ , Daniel M. Perkins ² and Pavel Kratina ³		
4			
5	1- Environmental Sciences Department, Federal University of São Carlos, São Carlos/SP, Brazil.		
6	Postcode: 13565-905. victor.saito@gmail.com – corresponding author		
(Twitter: @vs_saito		
o Q	2- Life Sciences Department, University of Roenampton, London, UK. Postcoae: SW154JD. Daniel Perkins@roehampton.ac.uk. Twitter: @drDanPerkins		
10	3- School of Biological and Chemical Sciences, Queen Mary University of London, London, UK. Postcode:		
11	E1 4NS. p.kratina@qmul.ac.uk		
12			
13	Key-words: Food webs, macroecology, metabolic scaling, neutral processes, niche		
14	processes		
15			
16	Abstract: Metabolism controls the pace of life driving major ecological patterns. We		
17	propose that the scaling of metabolism with temperature influences neutral processes		
18	of community assembly by controlling population dynamics independently of species		
19	identities. This perspective provides new insights into the prevalence of niche and		
20	neutral processes through universal energetic constraints.		

21 How metabolism controls community assembly processes

22 Current synthesis in community ecology recognizes the contribution of both, 23 niche and neutral processes (see Glossary) in the assembly of ecological communities 24 [1]. The niche perspective has traditionally focused on taxonomic identity and trait 25 differences in shaping biotic interactions and environmental filtering. In contrast, the 26 random birth, death, and dispersal of organisms within trophic levels have been the key factors in purely neutral perspective [2]. The current challenge in community ecology is 27 thus to determine the factors that explain the relative contribution of niche and neutral 28 29 processes during community assembly across environmental gradients [1]. Here, we 30 address this challenge by integrating concepts from the metabolic theory of ecology [3] 31 into the niche-neutral theories. We explain how considering the universal scaling of 32 mass-specific metabolic rates (hereafter metabolic rates) with temperature casts a 33 new light on how communities are organized in nature.

Metabolism encompasses the biological processing of material and energy by 34 35 organisms via biochemical reactions. Due to an increased rate of molecular kinetics, metabolic rates increase predictably with temperature [3]. Consequently, 36 37 environmental temperature is the most important abiotic driver of metabolism that 38 propagates to all levels of biological organization [3]. The increased metabolism at 39 higher temperature governs many natural processes, including the number of 40 individuals within communities and the rates of biomass production in ecosystems [3]. Because temperature consistently changes along altitudinal, depth and latitudinal 41 42 gradients, this should generate environmental gradients of metabolic rates. Because 43 metabolism influences fundamental biological processes, we argue that it modulates the 44 importance of neutral processes during community assembly.

45 The Unified Neutral Theory of Biodiversity and Biogeography [2] assumes that 46 organisms within a trophic level can be considered as approximately equivalent in their 47 chances of birth, death and dispersal [2]. This implies that population densities within trophic levels vary largely at random and similarly among species - i.e. negative 48 density-dependency is equal among and within species and thus populations drift in 49 time [2, 4]. Investigating the ecological equivalence of individuals and species is 50 51 therefore pivotal for understanding community assembly [4] and metabolism could 52 influence neutrality in the following ways.

53 First, greater metabolic rates at higher temperatures result in decreased longevity of ectotherms [3]. These changes in longevity are linked to extrinsic and 54 55 intrinsic factors affecting population death rates [5]. Extrinsic factors are influenced by 56 species niches as the chances of death increase in unfavourable environmental conditions. Intrinsic factors instead are driven by metabolic rates due to an increased 57 58 accumulation of damage from oxidative reactions, telomere shortening and deleterious 59 mutations [5]. We hypothesize a greater proportion of intrinsic to extrinsic deaths at 60 higher temperatures (Figure 1), which could therefore reduce competitive differences 61 among species and lead to a higher competitive equivalence. This would occur because 62 intrinsic deaths are controlled by damages acting stochastically among individuals and 63 consistently among species, possibly undermining their competitive differences. 64 Consequently, populations would be under relatively weaker control of niche-based 65 processes like competitive dominance (Figure 1). Higher death rates in organisms with high metabolism have been found across a wide range of taxa indicating a strong control 66 67 of intrinsic factors [5]. However, extrinsic factors may also increase death rates as biotic 68 interactions change predictably with warming (i.e. organisms became more susceptible 69 to predation as their oxidative damages accumulate)[3, 6]. Whether increased rates of 70 total deaths are predominantly driven by intrinsic or extrinsic factors remains an area 71 of future research. In any case, increased death rates reduce population densities and 72 lead us to the second major link between individual metabolisms and neutral processes 73 of community assembly.

74 There is ample evidence that population densities decline with increasing 75 metabolic rates, especially in ectotherms [3]. This is due to the increased death rates 76 and can be explained by the greater individual energetic demands at higher 77 temperatures resulting in lower densities under a fixed supply of resources [3] and by 78 the faster biomass turnover due to shorter life cycles under these conditions [7]. At 79 lower densities, the relative importance of neutral processes is enhanced [8] (Figure 1), because the influence of demographic stochasticity during community assembly is 80 81 inversely proportional to population density [8] (Figure 1). Species with large 82 competitive differences but with low densities can have equivalent chances of extinction 83 since the effect of demographic stochasticity could overcome those of niche processes 84 [8]. In communities with high densities the impacts of demographic stochasticity would 85 be relatively weak compared to the population variation caused by niche processes

- 86 (Figure 1). Such predictable variation in neutral processes due to population density has
- 87 been suggested theoretically [8] and demonstrated empirically [9]. Given the predicted
- 88 decrease in population densities under higher temperatures [3], this should entail
- 89 consistent variation in neutral processes across temperature gradients.
- 90

91 Metabolism and the assembly of stream metacommunities

Metabolism influences community assembly in several additional ways and the 92 93 importance of individual mechanisms may differ among ecosystems. We illustrate our 94 ideas using short-lived stream insects, as the lower densities with rising temperature 95 should be less important for long-lived organisms [3, 7]. Adult insects emerge into 96 terrestrial ecosystems and recolonize the streams via oviposition, completing the life 97 cycles in months to up to few years. In tropical communities, insect densities have been found to be approximately five times lower than in high-latitude streams [9], likely due 98 to the accelerated metabolism and biomass turnover in the warmer tropics, making 99 100 communities strongly affected by demographic stochasticity [9]. In addition to the 101 general effects of increased mortality and lower densities, other mechanisms should 102 operate in these communities. For example, predation could enhance neutrality in prey 103 communities because an increased metabolic rate in predator fish generally leads to 104 more generalist and omnivorous feeding [10]. This occurs because faster metabolism 105 requires organisms to feed more often, less selectively and on high carbon content prey [10], potentially leading to a higher stochasticity in size and identity of consumed prey, 106 107 reinforcing neutrality in prey communities.

The fast biomass turnover of aquatic insects entails frequent dispersal of adults among streams; with more dispersal events for tropical insects given they have more generations per year than temperate species [7]. Since tropical communities are more neutrally assembled, dispersal and recolonization is less predictable as well [9]. At the metacommunity level, neutral processes prevail due to the frequent colonization of organisms with variable body sizes and taxonomic identities that could ultimately influence neutrality at the metacommunity level [11].

In summary, differences in metabolism should lead to predictable variation in
the relative importance of neutral processes in stream communities. This variation can
also alter the way energy flows through ecosystems, explaining food web structures that

stems from energetic constraints, such as relationships between abundance and bodymass.

120

121 Niche and neutral mass-abundance relationships

122 Size spectra have long been used to investigate relationships between body 123 mass and abundance, and understand energy allocation and transfer in ecosystems. 124 These relationships depict the frequency distribution of individual body sizes and 125 allow comparisons of communities in different environmental settings, irrespective of 126 their taxonomic composition (Figure 2). Metabolic scaling theory predicts a negative 127 power-law relationship [3] as a function of two main parameters: the transfer 128 efficiency of energy across trophic levels and the relative size of predators and prey 129 (Figure 2). We propose that the fitted parameters of the size spectrum vary with 130 temperature and the relative influence of niche and neutral processes (Figure 2) 131 providing a way to test predictions across trophic levels. First, the variation in 132 abundance explained by body mass (i.e. R² value) should be smaller at higher 133 temperatures (and under neutral community assembly) due to enhanced importance of demographic stochasticity and the frequent random dispersal of organisms, relaxing 134 135 energetic constraints [11] (Figure 2). Under these conditions, higher temporal and spatial variation in size-spectra slopes would also be expected for communities in 136 137 warmer conditions (Figure 2). Finally, the intercept should be lower in warm regions 138 because of the lower population densities and community biomass [3, 7] (Figure 2). 139 Size spectra provide an excellent tool to test these and other hypotheses (Figure 2), as 140 they directly represent energy fluxes across trophic levels.

141

142 **Towards a metabolic niche theory**

143 Whereas the mechanisms described here suggest a weaker role of niche 144 processes in community assembly under higher temperatures, variation among systems could occur. For example: i) a fast pace of life could increase interspecific differences if 145 population density is strongly constrained by carrying capacity and limiting resources 146 147 are scarcer at high temperatures (e.g. green food webs) [12]. In such conditions, a faster 148 metabolism could lead to greater importance of niche differences accelerating deterministic competitive exclusions. ii) Predators with higher metabolism could also 149 150 specialise and selectively feed on more nutritious prey, as observed in lizards [13], in

- 151 contrast to increased generalism found for fish [10]. Our key point is not to imply a
- 152 singular direction of the metabolism-stochasticity relationship, but rather to emphasize
- 153 that the metabolic perspective provides a general biological framework to
- 154 understanding variation in niche and neutral community assembly.
- Our ideas represent the first steps towards linking metabolic constraints with
 neutral processes to understand community assembly within and across trophic levels.
 Future empirical tests of this framework will be pivotal to test whether niche-neutral
 theories and the metabolic theory of ecology can be viewed as two sides of the same
- 159 coin.
- 160

161 Acknowledgements

- 162 We thank Tadeu Siqueira, Raul Costa Pereira, Luís Schiesari, two reviewers and the
- 163 editor for comments that improved our perspective. The authors gratefully
- acknowledge the following grants: a Newton Fund/FAPESP grant (#2019/05464-1) to
- 165 DP, a Royal Society grant (#NAF\R2\180791) to PK, and a FAPESP grant
- 166 (#2019/06291-3) to VSS.

167	7 References		
168			
169	1-	Shoemaker, L. G. et al. (2020). Integrating the underlying structure of	
170		stochasticity into community ecology. <i>Ecology</i> . 101, e02922.	
171	2-	Hubbell, S. P. (2001). The unified neutral theory of biodiversity and	
172		biogeography, Princeton University Press, Princeton, NJ.	
173	3-	Brown, J. H. <i>et al.</i> (2004). Toward a metabolic theory of ecology. <i>Ecology</i> . 85,	
174		1771-1789.	
175	4-	McPeek, M. A. and Siepielski, A. M. (2019). Disentangling ecologically equivalent	
176		from neutral species: The mechanisms of population regulation matter. J. Anim.	
177		<i>Ecol.</i> 88, 1755-1765.	
178	5-	McCoy, M. W. and Gillooly, J. F. (2008). Predicting natural mortality rates of	
179		plants and animals. <i>Ecol. Lett. 11</i> , 710-716.	
180	6-	Brown, J. H., Hall, C. A., & Sibly, R. M. (2018). Equal fitness paradigm explained by	
181		a trade-off between generation time and energy production rate. <i>Nature Ecol.</i>	
182		<i>Evol. 2</i> , 262-268.	
183	7-	Munch, S. B. and Salinas, S. (2009). Latitudinal variation in lifespan within	
184		species is explained by the metabolic theory of ecology. <i>PNAS.</i> 106, 13860-13864.	
185	8-	Orrock, J. L. and Watling, J. I. (2010). Local community size mediates ecological	
186		drift and competition in metacommunities. Proc. R. Soc. Lond. B. 277, 2185-2191.	
187	9-	Siqueira, T. <i>et al.</i> (2020). Community size can affect the signals of ecological drift	
188		and niche selection on biodiversity. <i>Ecology</i> . 101, e03014.	
189	10	-González-Bergonzoni, I. <i>et al.</i> (2012). Meta-analysis shows a consistent and	
190		strong latitudinal pattern in fish omnivory across ecosystems. <i>Ecosystems</i> . 15,	
191		492-503.	
192	11	-Perkins, D. M. <i>et al.</i> (2018). Bending the rules: exploitation of allochthonous	
193		resources by a top-predator modifies size-abundance scaling in stream food	
194		webs. <i>Ecol. Lett.</i> 21, 1771-1780.	
195	12	-Bernhardt, J. R. <i>et al.</i> (2018). Metabolic theory and the temperature-size rule	
196		explain the temperature dependence of population carrying capacity. Am.	
197		Nat. 192, 687-697.	
198	13	-Bestion, E. <i>et al.</i> (2019). Altered trophic interactions in warming climates:	
199		consequences for predator diet breadth and fitness. Proc. R. Soc. Lond. B. 286,	
200		2019-2227.	

201 Figure 1. The influence of temperature and metabolism on the relative importance of 202 niche and neutral processes. A) Predictions based on the universal acceleration of 203 mass-specific metabolic rates with warming due to molecular kinetics. At the 204 individual-level, mortality rates increase whereas life span decrease with increasing 205 temperature. At the population level, density decreases whereas biomass turnover 206 increases with temperature. B) At high temperatures and fast metabolic rates, 207 population densities are low due to a faster pace of life and higher mortality rate. In 208 this scenario, the relative influence of demographic stochasticity increases (blue 209 dashed brackets), whereas those of niches differences (blue solid brackets) for 210 community assembly decreases (here represented as competition between two

211 species). Given the low population densities, species in warmer environments are more

- 212 prone to random extinctions (red circle).
- 213

214

215 Figure 2. The relative influence of niche and neutral processes on local community size spectra. **A)** Tropical and temperate communities have distinct patterns of population 216 217 dynamics. In warm tropical communities, populations (individual lines) are more 218 strongly influenced by demographic stochasticity due to small population densities (red colour) compared to large population densities in cool, temperate communities 219 220 (blue lines). A higher number of generations per year also enhances the number of 221 demographic events increasing the importance of stochastic population dynamics in 222 tropical communities. Due to energetic constraints, organisms tend to be, on average, 223 smaller in warmer tropical communities [3] (indicated by the thickness of lines), even 224 though the opposite relationship exist for some taxa. The relatively higher influence of 225 neutral processes entails greater variation in rank-abundance patterns in tropical 226 communities. This is illustrated by the bar plots where species have higher abundance 227 variation from T1 to T2 in tropical than in temperate communities. **B)** Hypothetical 228 local size spectra depicting the distribution of abundance among different size classes 229 in tropical and temperate communities. In tropical communities, the higher relative 230 importance of neutral processes results in greater variation of data around the 231 regression line, with size classes with higher and lower abundances than predicted 232 based on steady-state energetic conditions. Under these conditions, higher temporal and spatial variation in size-spectra parameters are expected in tropical communities 233 234 (variation in size spectra from T1 to T2). Dashed and solid lines indicate size spectra in 235 T1 and T2, respectively.

- 236 Glossary:
- 237

238 Mass-specific metabolic rate: Demands of energy per unit of body mass per time in order to maintain 239 biological functions inherent to survival. The difference from absolute metabolic rate is important, given 240 that body size also tends to decrease with higher environmental temperatures [3], sustaining a trade-off 241 along temperature gradients. In other words, individuals demand more energy under higher 242 temperatures, but also tend to be smaller, demanding less energy per individual. Mass-specific metabolic 243 rate reflects the higher energetic expenditure per unit of body mass at higher temperatures and is 244 commonly measured (and considered in this study) as basal metabolic rate of a resting or inactive 245 organism. This basal metabolic rate is generally correlated with the mean daily metabolic rate of organisms under active periods [6].

246 247

Metabolic rate: Individual demands of energy in time to maintain biological functions inherent to
 survivorship. In heterotrophs metabolism is aerobic respiration, whereas photosynthesis is the main
 contributor to the metabolic rates in autotrophs [3].

251

252 Neutral processes: A combination of processes that can be specifically stochastic at the population level.

253 These processes include stochastic rates of birth and death, dispersal and the introduction of

evolutionary novelty via mutation and speciation. The Unified Neutral Theory of Biodiversity and
Biogeography [2] assumes that these processes are similar among species within a trophic level at a first

256 approximation.257

Niche processes: A combination of processes where species differences determine ecological outcomes.
 For example, prey differences in anti-predator behaviour can determine predation pressure, or
 differences in species tolerances can determine community composition along a gradient of salinity.

261

Size spectra: Relationship between organism body size and abundance, which commonly encompasses
 multiple trophic levels. The relationship is depicted by plotting (on double logarithmic scales) the number

of individuals within body size (or mass) classes against the mid-point of the size class. The negative slope

of the size spectrum summarises energy allocation and transfer through the food web, for which a rich

body of theory exists (e.g. [3, 6]).