

The hidden role of multi-trophic interactions in driving diversity-productivity relationships

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This supplementary document includes:

Supplementary 1: Detailed description of simulation model

Supplementary Figures S1-12

16 **Supplementary 1: Detailed description of simulation model**

17 We used an allometric-trophic-network model to simulate the complex trophic dynamics of
18 ecosystems in a controlled environment (Schneider et al. 2016). It defines trophic interactions
19 between different species based on their body-mass ratios and utilizes a set of differential equations
20 that describes density changes of resources, producers, and animal consumers over time.

21 ***Simulating producer-resource interactions***

22 The change in biomass density P_i of primary producer species i is calculated as

$$23 \quad \frac{dP_i}{dt} = r_i G_i P_i - \sum_k A_k F_{ki} - x_i P_i,$$

24 with the first term describing resource-dependent growth, the second describing mortality due to
25 predation by animals, and the third describing metabolic demands. Both the intrinsic growth rate r_i of
26 species i , which defines its maximum possible growth rate, and the metabolic demands x_i of species i
27 scale allometrically with body mass (Enquist et al. 1998; Stephenson et al. 2014):

$$28 \quad r_i = m_i^{-0.25},$$

$$29 \quad x_i = x_p m_i^{-0.25}.$$

30 with $m_i = 10^{\mu_p}$ being the specific body mass of producer species i , where μ_p follows a uniform
31 distribution on $[0, 6]$. Metabolic demands were rescaled by $x_p = 0.138$ (Brose 2008). Growth of
32 primary producer species i was further limited by the species specific growth factor G_i , defined by
33 two limiting resource $j \in \{1, 2\}$ as

$$34 \quad G_i = \min \left(\frac{\theta_{i1}}{K_{i1} + \theta_{i1}}, \frac{\theta_{i2}}{K_{i2} + \theta_{i2}} \right),$$

35 where K_{ij} is the half-saturation density of resource j at which the resource uptake rate of primary
 36 producer i is half of its maximum, and follows a uniform distribution on $[0.1, 0.2]$. θ_{ij} is the
 37 concentration of resource j accessible by primary producer i .

38 To simulate different scenarios of resource-use dissimilarity (RUD; see methods in main text), we
 39 spread resource concentrations of each considered resource j across $C = 16$ resource compartments.
 40 The change in resource concentrations N_{jn} of resource j in resource compartment n was defined as:

$$41 \quad \frac{dN_{jn}}{dt} = D \left(\frac{S_j}{C} - N_{jn} \right) - v_j \sum_i r_i G_i P_i \frac{N_{jn}}{\theta_{ij}}.$$

42 The first term describes the rate at which resources are renewed. It is limited by the turnover rate D ,
 43 which was set to 0.25. The supply concentration S_j represents the maximum concentration of
 44 resource j . It was set to 50 and 25 for resources 1 and 2, respectively. As we defined each
 45 compartment n to be quantitatively the same, we split the supply concentration S_j equally between
 46 compartments. The second term captures the loss of resources due to primary production, which is
 47 similar to the resource-dependent growth term used to calculate the change in primary producer
 48 densities but separated for each resource compartment. The relative content of resource j in the
 49 biomass of primary producers is described as v_j and was set to 1 and 0.5 for resources 1 and 2,
 50 respectively. By keeping the ratio between S_j and v_j the same for both resources j , all resources
 51 considered can limit the growth of primary producers and consequently play a role in determining
 52 competitive advantages while contributing differently to primary production. The pool of resource j
 53 accessible by producer species i corresponds to the sum of resource concentrations in the
 54 compartments it has access to:

$$55 \quad \theta_{ij} = \sum_n \vartheta_{in} N_{jn},$$

56 with $\vartheta_{in} = 1$ if species i can access resource compartment n , $\vartheta_{in} = 0$ otherwise.

At the end of the simulations, we quantified primary production as the summed up resource uptake of both resources j , in all compartments n , and for all primary producer species i :

$$Y = \sum_i Y_i = \sum_j \sum_n v_j \sum_i r_i G_i P_i \frac{N_{jn}}{\theta_{ij}}.$$

Creating network topology and simulating animal consumers

Similar to primary producer species, each animal species k was characterized by its specific body mass $m_k = 10^{\mu_A}$ with the exponents drawn randomly from a uniform distribution on $[2, 12]$. To create a viable network topology, we calculated the probability of consumer species x to feed on an encountered resource species z as

$$L_{xz} = \left(\frac{m_x}{m_z R_{opt}} e^{1 - \frac{m_x}{m_z R_{opt}}} \right)^\gamma,$$

with the optimal consumer-resource body mass ratio $R_{opt} = 100$ and $\gamma = 2$. Low probabilities with $L_{xz} \leq 0.01$ were set to be zero. We only considered animal communities where each species had at least one resource species at maximum producer richness (i.e., 16 producer species). When lowering producer richness (see main text), animal species that lost their resource species were removed before simulations.

The change of biomass densities of species k , A_k over time, was simulated as

$$\frac{dA_k}{dt} = e_p A_k \sum_i F_{ki} + e_A A_k \sum_l F_{kl} - \sum_l A_l F_{lk} - x_k A_k,$$

with the first term describing increases due to the summed up herbivorous feeding on primary producer species i , with a conversion efficiency $e_p = 0.545$ (Lang et al. 2017). Similarly, the second term describes the summed up carnivorous feeding on animal species l , with a conversion efficiency of $e_A = 0.906$ (Lang et al. 2017). The third term captures mortality due to predation by animals l in the

78 same way as for primary producers. The last term represents metabolic demands of animal species k ,
 79 which scales allometrically with body mass (Ehnes et al. 2011) as

$$80 \quad x_k = x_A m_k^{-0.305},$$

81 with a scaling constant $x_A = 0.141$ (Ehnes et al. 2011). All trophic interactions include feeding rates

$$82 \quad F_{xz} = \frac{\omega_x b_{xz} Z_z^{1+q_{xz}}}{1 + c X_x + \omega_x \sum_{\zeta} b_{x\zeta} h_{x\zeta} Z_{\zeta}^{1+q_{x\zeta}}} \cdot \frac{1}{m_x}$$

83 as a function of the biomass densities X_x and Z_z of the consumer species x and resource species z ,
 84 respectively. Feeding rates capture the proportion of biomass of resource species z consumer species
 85 x consumes. By dividing by body mass, the per-capita feeding rate is transformed to be relative to
 86 one unit biomass. Consumers with multiple resource species have to split their feeding efforts,
 87 captured in the relative consumption rate ω_x , defined as the inverse of the number of prey species of
 88 consumer x . Time lost due to consumer interference c was drawn from a normal distribution ($\mu_c =$
 89 0.8 , $\sigma_c = 0.2$) for each food web. We used an interaction-specific, allometric Hill-exponent $1 + q_{xz}$
 90 (Kalinkat et al. 2013), which determines the functional response type of the interaction. It was
 91 calculated as:

$$92 \quad q_{xz} = \frac{q_{max} R_{xz}^2}{q_0^2 + R_{xz}^2},$$

93 with R_{xz} being the consumer-resource body mass ratio of consumer species x and resource species z .
 94 By setting $q_{max} = 1$, we assure that the functional response varies between the classic type II ($q_{xz} = 0$)
 95 and type III ($q_{xz} = 1$). At optimal consumer-resource body mass ratio $R_{opt} = 100$ we wanted q_{xz} to be at
 96 intermediate levels. Therefore, we also set $q_0 = 100$. At higher consumer-resource body mass ratios,
 97 the functional response gets closer to the classic type III, which lowers the feeding rates at low
 98 resource densities. The feeding rate was further determined by the capture coefficient:

$$99 \quad b_{xz} = b_0 m_x^{\beta_x} m_z^{\beta_z} L_{xz},$$

which describes the success rate of consumer species x to capture resource species z . It is based on the assumption that an encounter is more likely with higher movement speeds of both consumer and resource species. Since movement speed scales allometrically and based on feeding type (Hirt et al. 2017), we drew β_x and β_z from according normal distributions (carnivore: $\mu_\beta = 0.42$, $\sigma_\beta = 0.05$, omnivore: $\mu_\beta = 0.19$, $\sigma_\beta = 0.04$, herbivore: $\mu_\beta = 0.19$, $\sigma_\beta = 0.04$, primary producer: $\mu_\beta = 0$, $\sigma_\beta = 0$). Similarly, we assumed different values for b_0 based on the feeding type of the consumer (carnivore: $b_0 = 50$, omnivore: $b_0 = 100$, herbivore: $b_0 = 200$). The handling time:

$$h_{xz} = h_0 m_x^{\eta_x} m_z^{\eta_z},$$

scales with the body mass of consumer and resource species to the power of η_x ($\mu_{\eta_x} = -0.48$, $\sigma_{\eta_x} = 0.03$) and η_z ($\mu_{\eta_z} = -0.66$, $\sigma_{\eta_z} = 0.02$) respectively. The scaling constant h_0 was set to 0.4. All parameters drawn from normal distributions had to fall within the inclusive limits of $\mu \pm 3\sigma$ or be redrawn otherwise.

Simulation setup

Initial biomass densities of primary producer and animal species were randomly drawn from uniform distributions on $[0, 1]$. Resource densities were initialized for the whole resource pool with random values drawn from uniform distributions on $[S_j / 2, S_j]$, which were then evenly split between compartments. We ran simulations until $t = 150,000$. Species that reached biomass densities $< 10^{-6}$ during simulations were assumed to be extinct, and their values were set to 0.

We ran all simulations in Julia 1.2.0 (Bezanson et al. 2017) using the DifferentialEquations package (Rackauckas & Nie 2017) and utilizing a stiffness detection algorithm that automatically switched between the solvers Vern7 for non-stiff problems and Rodas4 for stiff problems.

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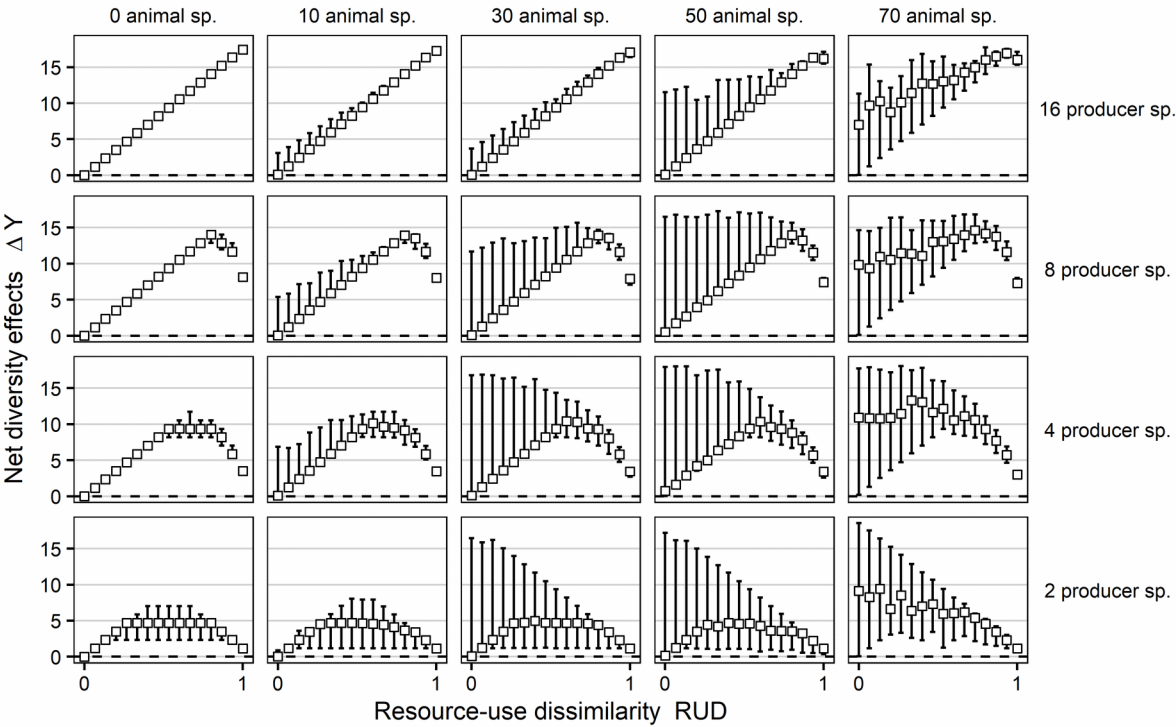
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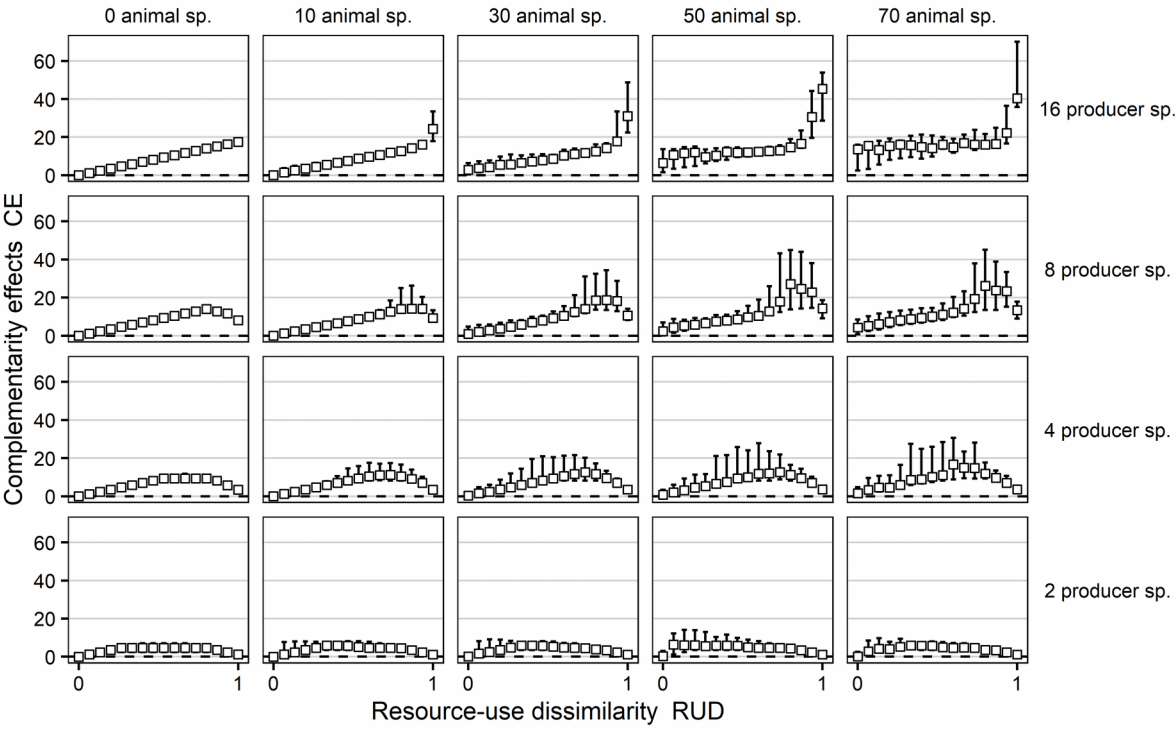
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142 **Figure S1:** Effects of resource-use dissimilarity on net diversity effects ΔY of the primary producer
 143 community. Effects are shown for different levels of producer (rows) and multi-trophic animal
 144 richness (columns). Error bars show 25th and 75th; squares show 50th percentile (i.e., median).



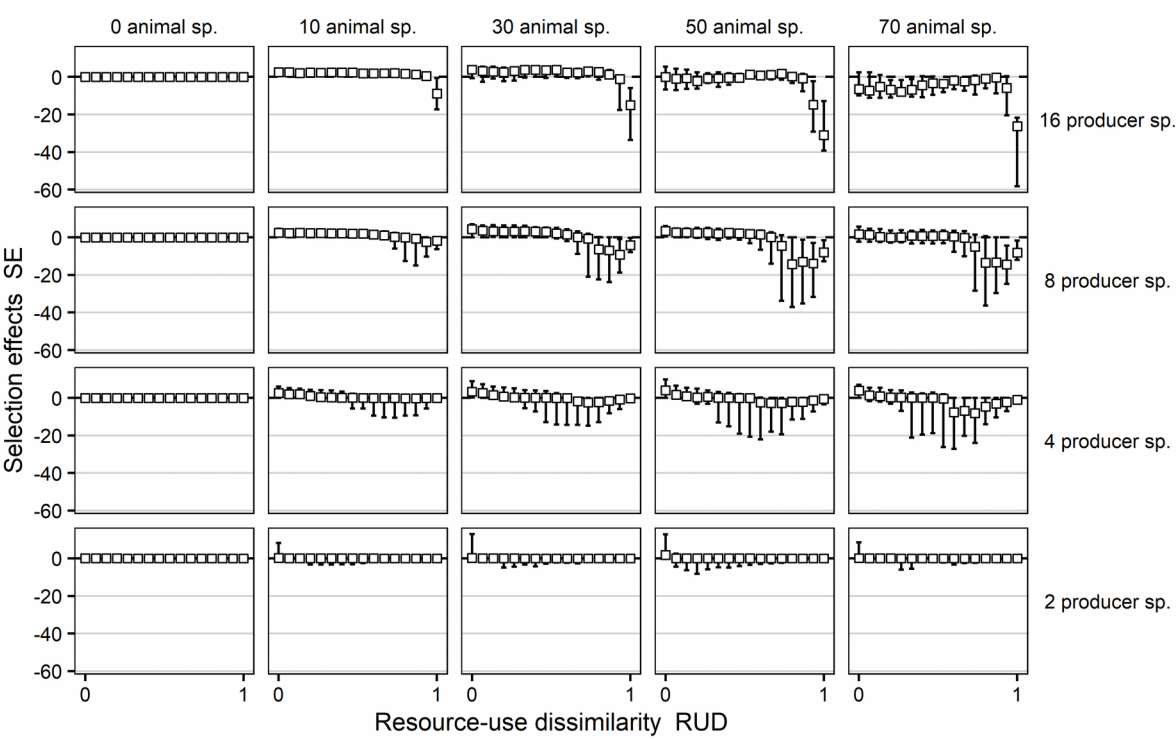
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146 **Figure S2:** Effects of resource-use dissimilarity RUD on complementarity effects CE of primary
 147 productivity. Effects are shown for different levels of producer (rows) and multi-trophic animal
 148 richness (columns). Error bars show 25th and 75th; squares show 50th percentile (i.e., median).



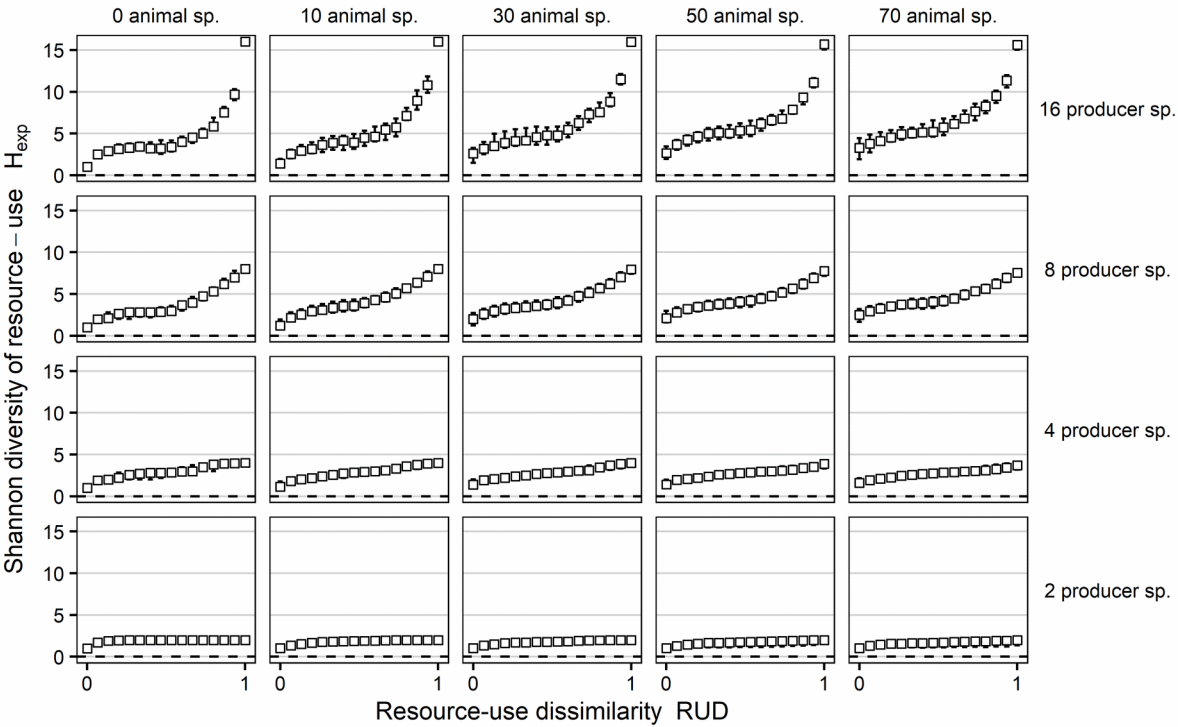
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150 **Figure S3:** Effects of resource-use dissimilarity RUD on selection effects SE of primary productivity.
 151 Effects are shown for different levels of producer (rows) and multi-trophic animal richness (columns).
 152 Error bars show 25th and 75th; squares show 50th percentile (i.e., median).



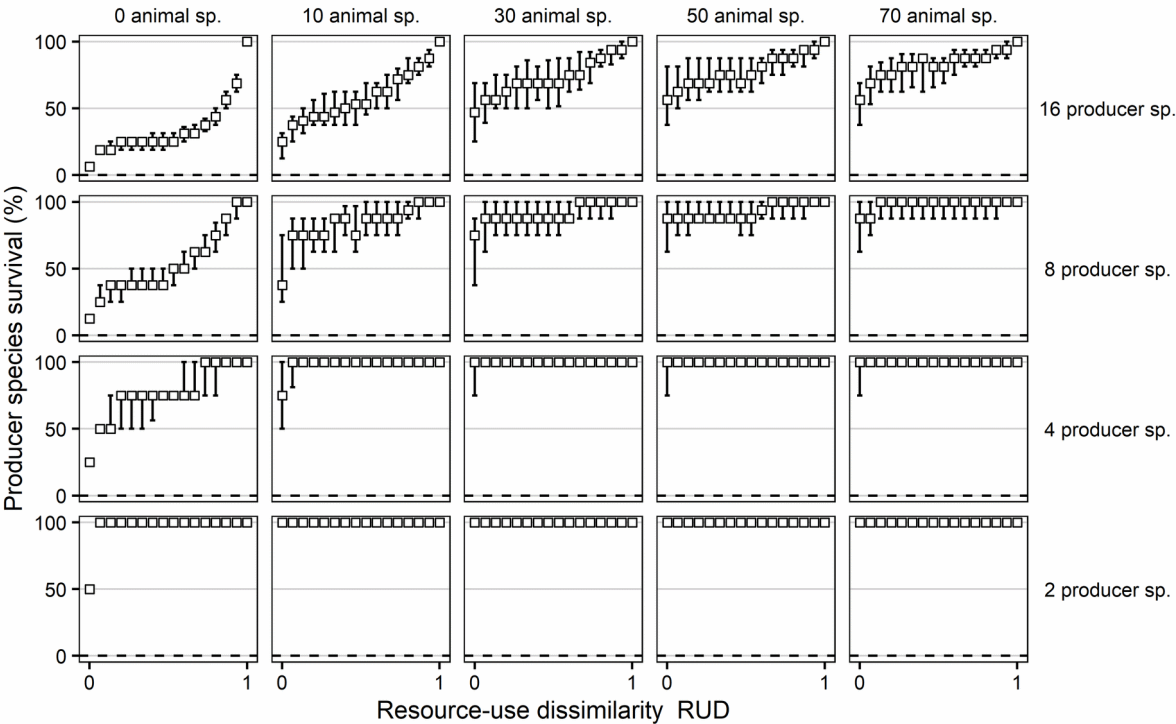
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155 **Figure S4:** Effects of resource-use dissimilarity RUD on the Shannon diversity of primary producers'
 156 resource-use H_{exp} . Effects are shown for different levels of producer (rows) and multi-trophic animal
 157 richness (columns). Error bars show 25th and 75th; squares show 50th percentile (i.e., median).



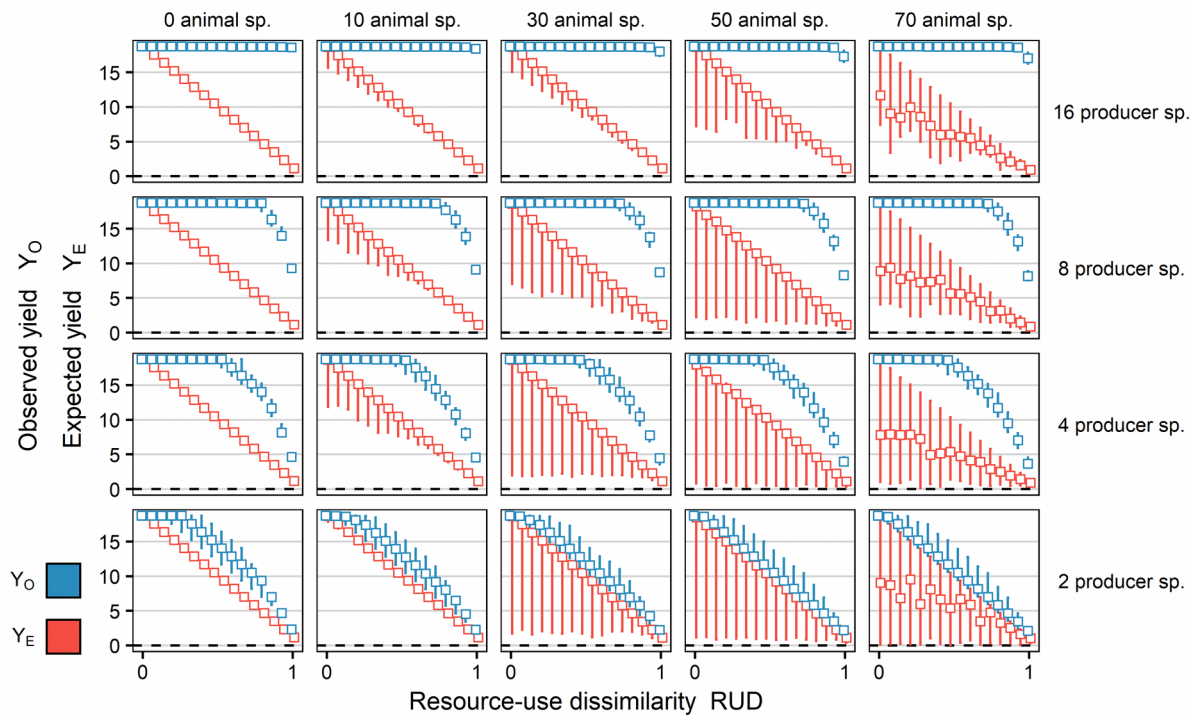
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159 **Figure S5:** Effects of resource-use dissimilarity RUD on the relative survival of the primary producer
 160 community. Effects are shown for different levels of producer (rows) and multi-trophic animal
 161 richness (columns). Error bars show 25th and 75th; squares show 50th percentile (i.e., median).



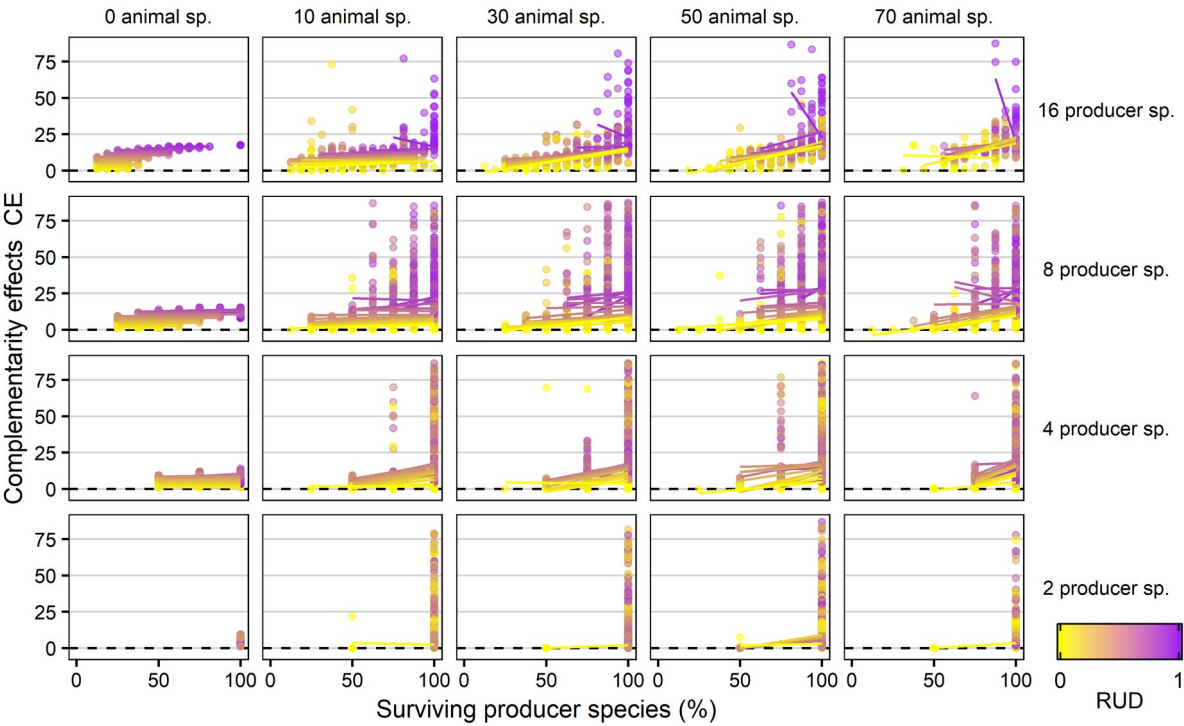
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163 **Figure S6:** Effects of resource-use dissimilarity RUD on observed yield Y_O (blue) and expected yield Y_E
 164 (red) of the primary producer community. Effects are shown for different levels of producer (rows)
 165 and multi-trophic animal richness (columns). Error bars show 25th and 75th; squares show 50th
 166 percentile (i.e., median).



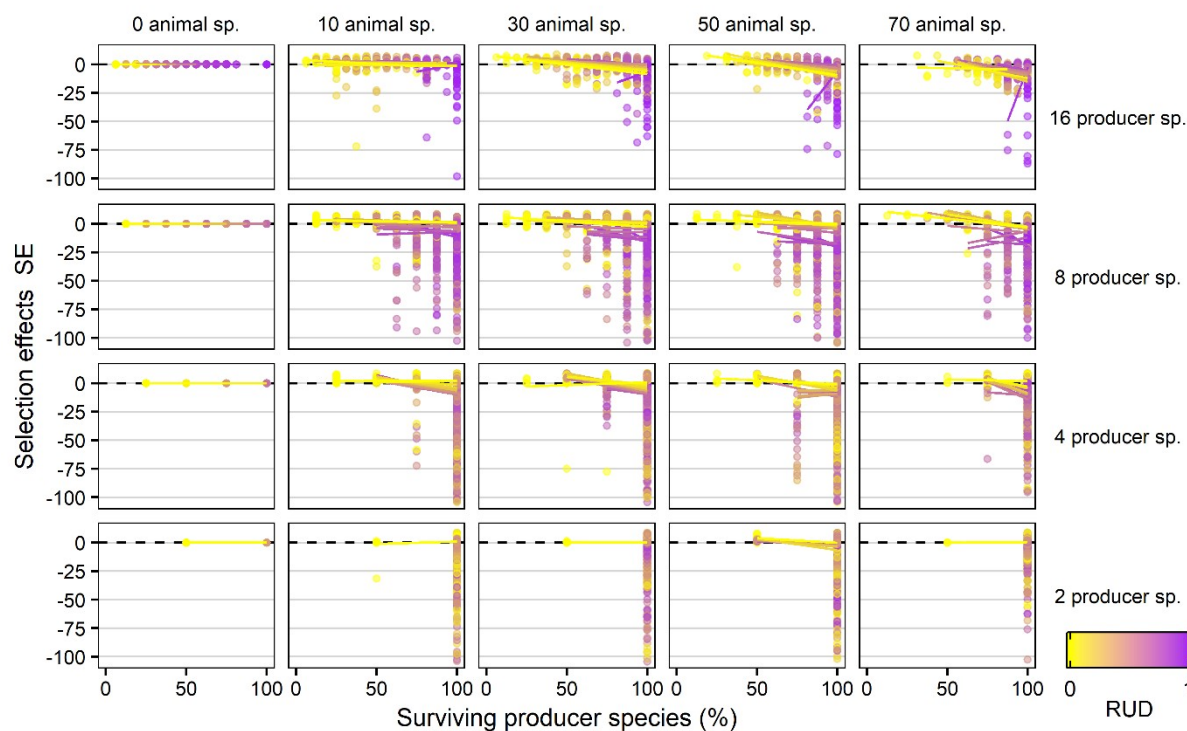
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168 **Figure S7:** Effects of relative primary producer species survival on complementarity effects CE. Effects
 169 are shown for different levels of producer (rows) and multi-trophic animal richness (columns). The
 170 level of resource-use dissimilarity (RUD) is indicated by the color (yellow: low, purple: high). To
 171 improve readability, only 95% of the simulated food webs are shown.



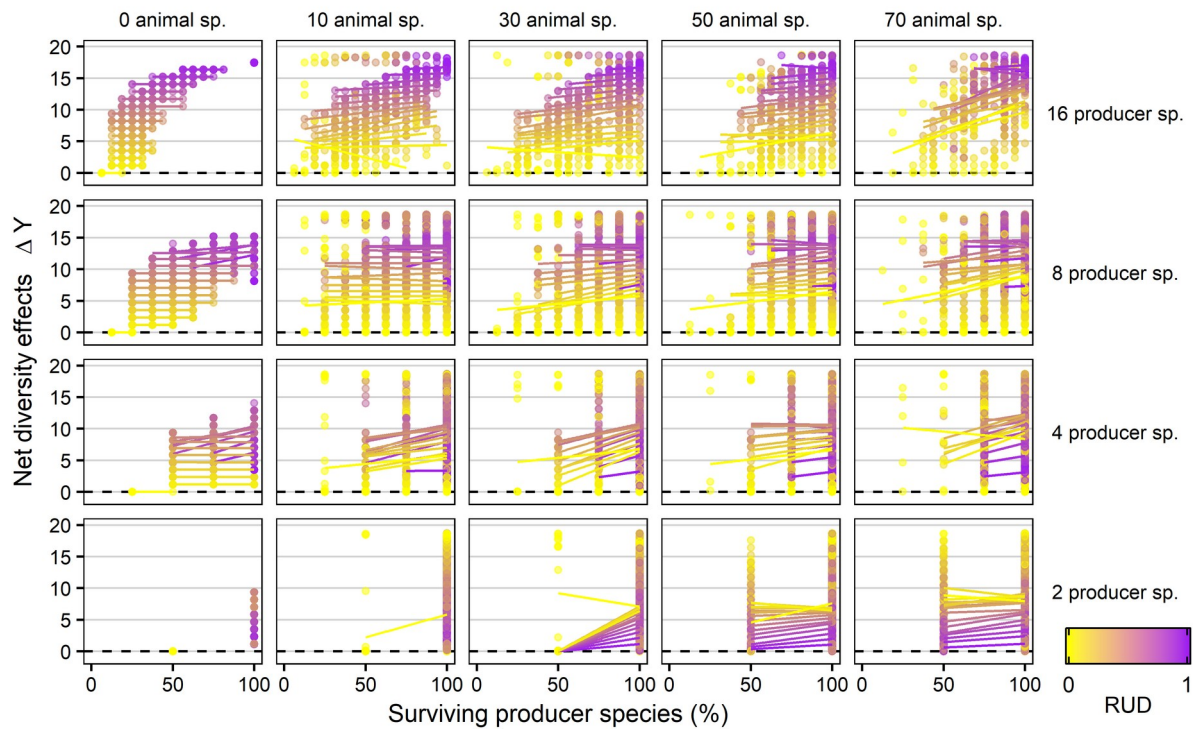
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173 **Figure S8:** Effects of relative primary producer species survival on selection effects SE. Effects are
 174 shown for different levels of producer (rows) and multi-trophic animal richness (columns). The level
 175 of resource-use dissimilarity (RUD) is indicated by the color (yellow: low, purple: high). To improve
 176 readability, only 95% of the simulated food webs are shown.



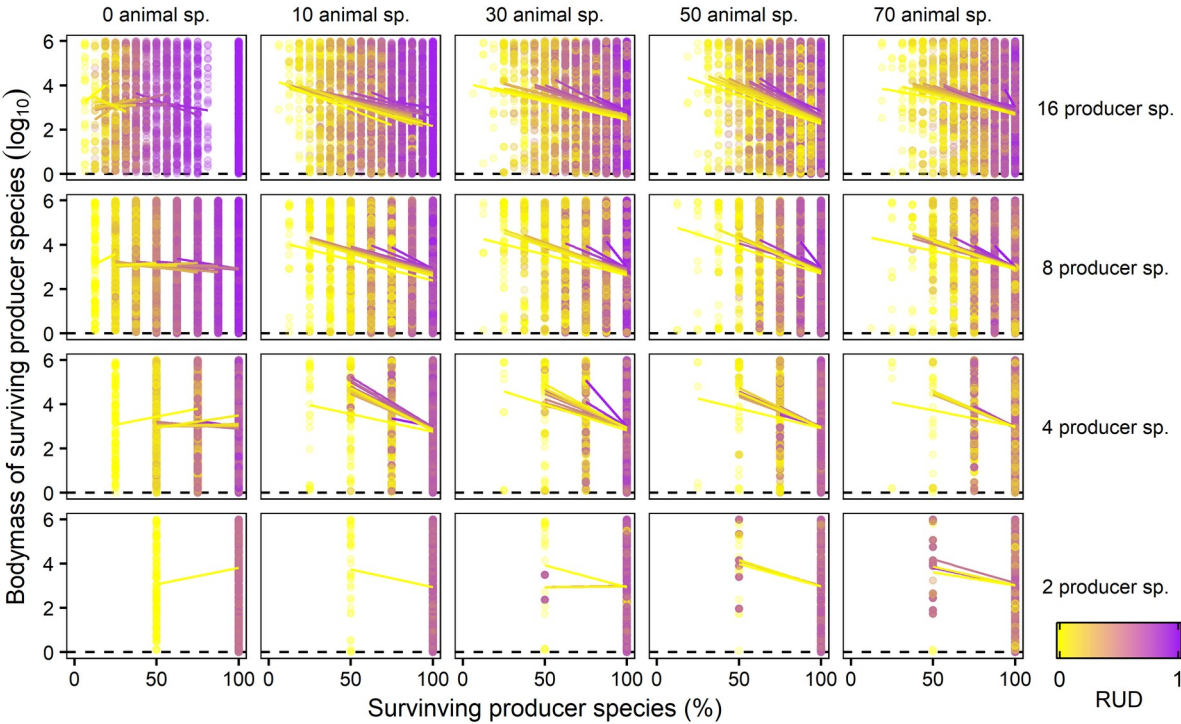
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178 **Figure S9:** Effects of relative primary producer species survival on net diversity effects ΔY . Effects are
 179 shown for different levels of producer (rows) and multi-trophic animal richness (columns). The level
 180 of resource-use dissimilarity (RUD) is indicated by the color (yellow: low, purple: high). To improve
 181 readability, only 95% of the simulated food webs are shown.



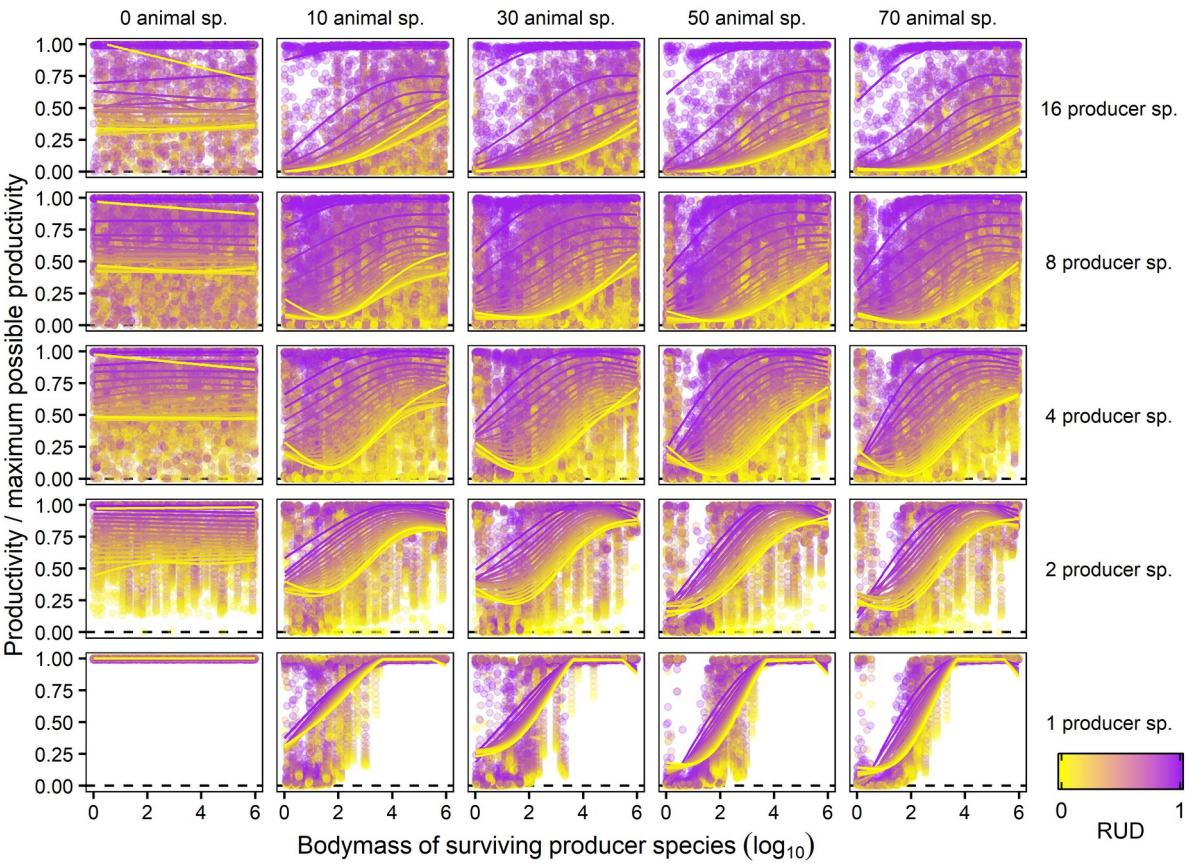
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183 **Figure S10:** Effects of relative primary producer species survival on the \log_{10} biomass of the surviving
 184 producer species. Effects are shown for different levels of producer (rows) and multi-trophic animal
 185 richness (columns). The level of resource-use dissimilarity (RUD) is indicated by the color (yellow:
 186 low, purple: high).

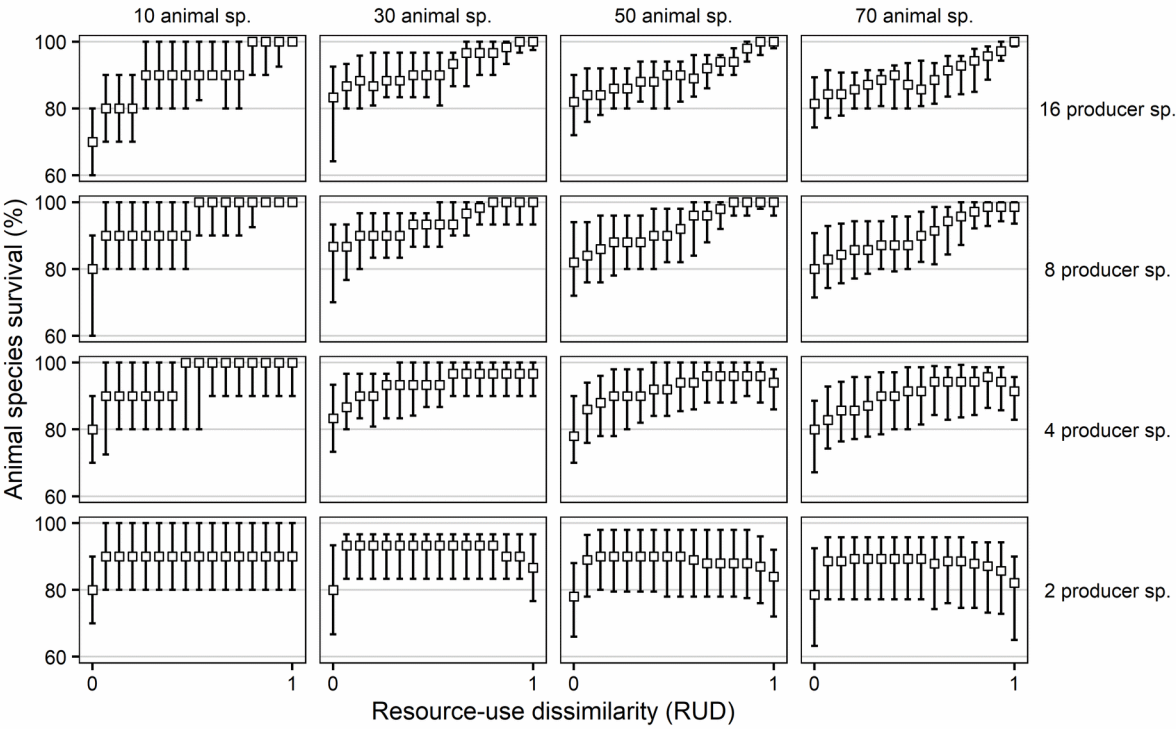


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188 **Figure S11:** Effects of \log_{10} biomass of the surviving producer species on their productivity relative to
 189 their maximum possible productivity (i.e., used resources / accessible resource). Effects are shown
 190 for different levels of producer (rows) and multi-trophic animal richness (columns). The level of
 191 resource-use dissimilarity (RUD) is indicated by the color (yellow: low, purple: high).



193 **Figure S12:** Effects of resource-use dissimilarity RUD on the relative survival of the animal
 194 community. Effects are shown for different levels of producer (rows) and multi-trophic animal
 195 richness (columns). Error bars show 25th and 75th; squares show 50th percentile (i.e., median).



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