Appendix B. Distinctiveness values in the special case of even abundances for all species.

In the special case of even abundances for all species, Rao's quadratic diversity is

$$
Q \text{eq} = \sum_{i=1}^{N} \sum_{j=1}^{N} \frac{1}{N} \frac{1}{N} d_{ij}
$$

the associated rarity and originality indices become, respectively,

$$
\rho \text{eq} = \frac{N-1}{N}
$$

and

$$
O \text{eq}_j = \sum_{i=1}^N \frac{1}{N} d_{ij}
$$

yielding to the following associated distinctiveness index:

$$
D\mathrm{eq}_j = \sum_{i \neq j}^{N} \frac{1}{N - 1} d_{ij}
$$

The FD-distinctiveness values are given in Tables B.1 and B.2 for the parametric extensions of the quadratic diversity.

Table B.1 Functional distinctiveness indices associated with parametric extensions of Rao's quadratic diversity in the special case of even abundances for all species. Distinctiveness is expressed in terms of dissimilarities ($d_{ij} \ge 0$) or similarities ($s_{ij} = 1 - d_{ij}$, in which case $0 \le d_{ij} \le 1$) between species; Notations are identical as those in the main text).

* case of all species similar ($s_{ij} = 1 \forall i, j$, or all species positioned on the same tip in the phylogenetic tree) excluded.

Table B.2 Phylogenetic distinctiveness indices associated with parametric extensions of Rao's quadratic diversity in the special case of even abundances for all species. Distinctiveness is expressed in terms of the branch lengths (*Lb*) of a phylogenetic tree with species as tips, the number of species descending from branch *b* (*N_b*), *C*(*j*, root) the set of branches from j to the root of the tree, $H_j = \sum_{b \in C(j, \text{root})} L_b$, *h_j* the length of the terminal branch that connects species *j* to the rest of the tree and *H* a value such that $H > \max_i H_i$ (Notations are identical as those in the main text).

	First parametric extension	Third parametric extension
	(phylogenetic approach)	(phylogenetic approach)
	$\alpha \Delta \text{eq}_{j} = \frac{N^{\alpha-1} - (\sum_{b \in C(j, \text{root})} \frac{L_b}{H} N_b)^{\alpha-1}}{N^{\alpha-1} - 1}$	$\alpha \Delta \text{eq}_{j}^{*} = \sum_{b \in C(j, \text{root})} L_b \left(\frac{N^{\alpha-1} - N_b^{\alpha-1}}{N^{\alpha-1} - 1} \right)$
$\alpha \rightarrow -\infty$	$^{\alpha}$ Δ eq _j $\frac{}{\alpha \rightarrow -\infty}$ 0	$\alpha \Delta \mathrm{eq}^*_{i} \longrightarrow h_{i}$
$\alpha = 0$	${}^{0}\Delta$ eq _j = $\frac{1/\left(\sum_{b \in C(j, \text{root})} \frac{L_b}{H} \frac{N_b}{N}\right) - 1}{N - 1}$	$^{0}\Delta$ eq [*] _j = $\sum_{b \in C(j, \text{root})} L_b \left(\frac{\frac{N}{N_b} - 1}{N - 1} \right)$
$\alpha \rightarrow 1$	$\alpha \Delta \text{eq}_{i} \longrightarrow 1 - \log_{N} \left(\sum_{b \in C(i,\text{root})} \frac{L_b}{H} N_b \right)$	$\alpha \Delta$ eq [*] _j $\xrightarrow{\alpha \rightarrow 1}$ $\sum_{b \in C(j, \text{root})} L_b (1 - \log_N N_b)$
$\alpha = 2^*$	² Δ eq _j = $\frac{N - \sum_{b \in C(j, \text{root})} \frac{L_b}{H} N_b}{N - 1}$	² Δ eq [*] _j = $\sum_{b \in C(j, \text{root})} L_b \left(\frac{N - N_b}{N - 1} \right)$
	$=\frac{H-H_j}{H}+\sum_{i\neq j}^{N}\frac{1}{N-1}\frac{H-H_j+\delta_{ij}}{H}$	$=\sum_{i\neq j}^N\frac{1}{N-1}\delta_{ij}$
$\alpha \rightarrow +\infty$ **	$\alpha \Delta$ eq _i $\frac{\alpha}{\alpha \rightarrow +\infty}$ 1	$\alpha \Delta$ eq [*] _i $\xrightarrow{\alpha \rightarrow +\infty}$ H _j

 $*$ δ_{ij} , as in the main text, is the sum of branch lengths on the shortest path from tip *j* to its most recent common ancestor with species *i*. For ultrametric trees H_j is constant over j. If $H = Hj$, then ${}^2 \Delta \text{eq}_j = \sum_{i \neq j}^{N} \frac{1}{N-1} \frac{\delta_{ij}}{\max_j}$ $j = \sum_{i \neq j} \frac{1}{N-1} \frac{1}{\max_i \delta_{ij}}$ δ Δ eq_j = $\sum_{i \neq j}^{N} \frac{1}{N-1} \frac{\delta_{ij}}{\max_{i} \delta_{ii}}$.

** case of all species similar (i.e. all species positioned on the same tip in the phylogenetic tree) excluded.

Proofs for the limits:

If species are not all similar, because $\sum_{i=1}^{N}$ *N* $\sum_{i=1}^{N} s_{ij}$ < *N* , then when *α*→+∞

$$
N^{\alpha-1} - \left(\sum_{i=1}^{N} S_{ij}\right)^{\alpha-1} \sim N^{\alpha-1}
$$
 and $N^{\alpha-1} - 1 \sim N^{\alpha-1}$

yielding

$$
\frac{N^{\alpha-1} - \left(\sum_{i=1}^{N} S_{ij}\right)^{\alpha-1}}{N^{\alpha-1} - 1} \xrightarrow{\alpha \to +\infty} 1
$$

Similarly, because $N_b < N$,

$$
\frac{N^{\alpha-1}-N_b^{\alpha-1}}{N^{\alpha-1}-1}\longrightarrow 1
$$

yielding,

$$
{}^{\alpha}\Delta eq_j \longrightarrow {}_{\alpha \to +\infty} H_j = \sum\nolimits_{b \in C(j, \text{root})} L_b
$$

The same reasoning yields

$$
\frac{\left(N^{\alpha-1}-c^{\alpha-1}\right)}{\left(N^{\alpha-1}-1\right)} \longrightarrow 1
$$

and thus to

$$
{}^{\alpha}Deq_j^* \longrightarrow \sum_{c=1}^{N-1} u_{c|j} = \sum_{c=1}^{N-1} \left(d_{c|j,j} - d_{c-1|j,j} \right) = d_{N-1|j,j} - d_{0|j,j} = \max_{i \neq j} d_{ij}
$$

Note that in all cases if species are all similar then their distinctiveness is null for all values of *α*.

$$
\frac{1-\left(p_j\right)^{\alpha-1}}{\alpha-1} \longrightarrow \ln\left(1/p_j\right)
$$
 (Patil and Taillie 1982)

and

and
\n
$$
\frac{1 - \left(\sum_{i=1}^{N} p_i s_{ij}\right)^{\alpha - 1}}{\alpha - 1} \longrightarrow \ln\left(1 / \sum_{i=1}^{N} p_i s_{ij}\right)
$$
 (Ricotta and Szeidl 2006)

yields

yields
\n
$$
{}^{1}Deq = \frac{\ln\left(N/\sum_{i=1}^{N} S_{ij}\right)}{\ln\left(N\right)} = 1 - \log_{N}\left(\sum_{i=1}^{N} S_{ij}\right)
$$

$$
^{\alpha}\Delta\mathrm{eq}_{j}=\sum\nolimits_{b\in C\left(j,\mathrm{root}\right)}L_{b}\left(\frac{N^{\alpha-1}-N_{b}^{\alpha-1}}{N^{\alpha-1}-1}\right)
$$

5 For the phylogenetic tree, we reasonably consider that all branches support more than one species except the terminal branches. In this summation for the equation of ${}^{\alpha}\Delta$ eq_j, N_b = 1 thus only when *b* is the terminal branch that connects species *j* to the rest of the tree. For all other branches, $N_b > 1$. To highlight this point, ["] Δ eq_{*j*} can be rewritten as

$$
^{\alpha}\Delta\mathrm{eq}_{j}=h_{j}+\sum\nolimits_{b\in C\left(A_{j},\text{root}\right) }L_{b}\left(\frac{N^{\alpha-1}-N_{b}^{\alpha-1}}{N^{\alpha-1}-1}\right)
$$

where A_j is the most recent ancestor of species *j*. When α tends to $-\infty$, both $N^{\alpha-1}$ and $N_b^{\alpha-1}$ tends to 0 while $N^{\alpha-1}-1$ tends to -1. This yields

$$
\sum\nolimits_{b \in C\left(A_j, \mathrm{root} \right)} L_b \left(\frac{N^{\alpha - 1} - N_b^{\alpha - 1}}{N^{\alpha - 1} - 1} \right) \xrightarrow{\alpha \to -\infty} 0
$$

and thus $\alpha \Delta \text{eq}_j \longrightarrow h_j$.

and thus "
$$
\triangle
$$
eq_j $\frac{}{\alpha \to \infty} h_j$.
Knowing that ${}^{\alpha}Deq_j^* = \frac{\sum_{c=1}^{N-1} u_{c|j} (N^{\alpha-1} - c^{\alpha-1})}{(N^{\alpha-1} - 1)} = u_{1j} + \frac{\sum_{c=2}^{N-1} u_{c|j} (N^{\alpha-1} - c^{\alpha-1})}{(N^{\alpha-1} - 1)}$

$$
\frac{\sum_{c=2}^{N-1} u_{c|j} \left(N^{\alpha-1} - c^{\alpha-1}\right)}{\left(N^{\alpha-1} - 1\right)} \xrightarrow{\alpha \to -\infty} 0
$$

and

$$
u_{1,j}=\min_{i\neq j}d_{ij},
$$

$$
{}^{\alpha}Deq_j^* = \underbrace{-\longrightarrow \min_{\alpha \to -\infty} d_{ij}}_{i \neq j} d_{ij}.
$$

The same reasoning can be applied to ${}^{\alpha}Deq_i = \frac{N^{\alpha-1} - (\sum_{i=1}^{N} s_{ij})^{\alpha-1}}{N^{\alpha-1} - N^{\alpha-1}}$ yield 1 1 $eq_j = \frac{1}{N^{\alpha-1}}$ 1 *N* $i=1$ ³*ij j* $N^{\alpha-1} - \left(\sum_{i=1}^{N} s_i\right)$ *D N* $a-1$ $\left(\nabla^N\right)^{\alpha}$ α α $-1\left(\sum_{\alpha}^{N}\right)^{\alpha-1}$ = \overline{a} \overline{a} $=$ \overline{a} \sum yielding $^{\alpha}$ *D*eq_{*j*} $\frac{\alpha}{\alpha \rightarrow -\infty}$ </sub> 0.

$$
\text{ }^{\alpha}\Delta \text{eq}_j=\sum\nolimits_{b\in C\left(j,\text{root}\right)}L_b\left(\frac{1-\left(\frac{N_b}{N}\right)^{\alpha-1}}{1-\displaystyle\frac{1}{N^{\alpha-1}}}\right)
$$

Because $N_b < N$,

1 $1 - \left(\frac{N_b}{N}\right)^{\alpha-1} \longrightarrow \ln\left(\frac{N_b}{N}\right)$ $\left(\frac{R}{N}\right)$ \longrightarrow $\left(\frac{N}{N}\right)$ α α - $-\left(\frac{N_b}{N}\right)^{\alpha-1} \longrightarrow \ln\left(\frac{N_b}{N}\right)$ $1 - \frac{1}{N^{\alpha-1}} \longrightarrow \ln\left(\frac{1}{N}\right)$ $\frac{1}{N^{\alpha-1}} \longrightarrow \ln\left(\frac{1}{N}\right)$ $-\frac{1}{N^{\alpha-1}}\longrightarrow \ln\left(\frac{1}{N}\right)$

yielding

yielding
\n
$$
\alpha \Delta \text{eq}_{j} \longrightarrow \sum_{b \in C(j,\text{root})} L_b \left(1 - \log_N N_b\right)
$$

As
$$
c < N
$$
, the same reasoning yields $^{\alpha}Deq_j^* \xrightarrow[\alpha \to 1]{} \sum_{c=1}^{N-1} u_{c|j} (1 - \log_N(c)).$

References:

Patil, G.P., Taillie, C., 1982. Diversity as a concept and its measurement. Journal of the American Statistical Association 77, 548–561. https://doi.org/10.2307/2287709

Ricotta, C., Szeidl, L., 2006. Towards a unifying approach to diversity measures: Bridging the gap between the Shannon entropy and Rao's quadratic index. Theoretical Population Biology 70, 237–243. https://doi.org/10.1016/j.tpb.2006.06.003