

# Unifying occupancy-detection and local frequency scaling (Frescalo) models

Oliver L. Pescott\*

*<sup>a</sup>UK Centre for Ecology and Hydrology, Benson Lane, Crowmarsh Gifford, OX10 8BB, Oxfordshire, UK*

---

## Abstract

Frescalo’s “local frequency scaling” and classical occupancy-detection models both seek to recover true species-occurrence signals from imperfect data. In this paper, we show that the two approaches rest on the same underlying detection mathematics. Occupancy models treat each site’s repeat visits as independent detection trials and separately estimate occupancy probability and per-visit detectability. Frescalo, by contrast, pools data across ecologically defined neighbourhoods and infers a single combined detection rate and a temporal “time-factor” to capture trends. We demonstrate that the Bernoulli-trial formulation of occupancy-detection converges to Frescalo’s Poisson-process framework, with occupancy and detectability collapsing into a single rate parameter. This equivalence clarifies how Frescalo’s neighbourhood and time corrections function as a coarser-scale analogue of repeat-visit models. By casting Frescalo in occupancy modelling terms, we hope to promote further investigation into the adoption of occupancy-model diagnostics, extensions and covariate tests within Frescalo analyses, improving transparency and rigour when working with opportunistic biodiversity data.

*Keywords:* occupancy models, sampling effort, effort correction, citizen science, unstructured data, Frescalo

---

## 1. Introduction

Occupancy-detection models [9] and the Frescalo “local frequency scaling” method [6] both aim to correct raw biological records (i.e. species occurrence) data for imperfect sampling. Classical occupancy models do this at the scale of repeated visits to individual sites, explicitly estimating true presence probabilities ( $\psi$ ) and detectability ( $p$ ) via a hierarchical likelihood. Frescalo was designed to work at larger spatio-temporal scales, exploiting emergent patterns of relative frequency in “neighbourhoods” to derive Poisson-process-based scaling factors ( $\alpha$ ) and species’ relative “time factors” indexing true fluctuations in site occupancy. Given that many datasets lack repeat-visit structure, and/or may exhibit variation in the detection process that is unmodellable due to a

---

\*Corresponding author

*Email address:* [olipes@ceh.ac.uk](mailto:olipes@ceh.ac.uk) (Oliver L. Pescott)  
*Preprint submitted to Elsevier*

18 lack of knowledge of its determinants [11], understanding how Frescalo recovers  
 19 effort-adjusted trends from aggregated data can broaden the toolkit of ecologists.

20 Although the two models can appear quite different, Pescott et al. [11]  
 21 informally suggested that Frescalo could be seen as a type of occupancy-detection  
 22 model “where an adjustment for overlooked species is made in relation to spatial  
 23 rather than temporal replication, whilst simultaneously adjusting for variable  
 24 regional effort”. We here show that this suggestion can be formalised due to  
 25 the two model types’ reliance on the same core mathematics of Bernoulli versus  
 26 Poisson detections. Below we (1) recall each framework, (2) write down their  
 27 key equations, and (3) algebraically map one onto the other, demonstrating that  
 28 Frescalo time trends are based on an implicit occupancy-detection model whose  
 29 “visits” and “occupancy” are folded into a single site/species discoverability rate  
 30 parameter  $\lambda$  and standardised neighbourhood effort index.

## 31 **2. Occupancy-detection models**

### 32 *2.1. Basic single-season model*

33 Following MacKenzie et al. [9], at each site  $i$  for species  $j$  assume a latent  
 34 occupancy indicator

$$z_{ij} \sim \text{Bernoulli}(\psi_{ij}).$$

35 Conditional on presence,  $v$  total survey visits indexed by  $k$  produce

$$y_{ij1}, \dots, y_{ijv} \mid z_{ij} = 1 \sim \text{Bernoulli}(p_{ij})$$

36 where  $p$  is detectability. If  $z_{ij} = 0$  (i.e. species absent), then all  $y_{ijk} = 0$ .  
 37 Marginalising out  $z_{ij}$ , it is well-known that the probability of at least one  
 38 detection across  $v$  visits is

$$P(\max_k y_{ijk} = 1) = \psi_{ij}[1 - (1 - p_{ij})^v].$$

39 Thus the model simultaneously estimates

$$\psi_{ij} = \text{Pr}(\text{occupied}), \quad p_{ij} = \text{Pr}(\text{detect}|\text{occupied}),$$

40 and inference proceeds via the full likelihood over all sites and detection histories.

## 41 **3. Frequency scaling using local occupancy (Frescalo)**

### 42 *3.1. Neighbourhood frequencies*

43 Frescalo [6] pools presence-only data across a neighbourhood around target  
 44 site  $i$ . We denote the observed proportion of neighbourhood sites in which  
 45 species  $j$  was recorded by  $f_{ij}$  (in practice this frequency may relate to a weighted  
 46 neighbourhood as per Hill [6], but this detail is not crucial for what follows).  
 47 Under a Poisson-process model of species discovery with rate  $\lambda_{ij}$  and unknown  
 48 total neighbourhood-level sampling effort  $s_{i(N)}$ , one has

$$f_{ij} = 1 - \exp(-\lambda_{ij}s_{i(N)}).$$

49 Subsequently, a frequency-weighted neighbourhood index

$$\phi_i = \frac{\sum_j f_{ij}^2}{\sum_j f_{ij}}$$

50 is then “standardised” to a target value  $\Phi$  by solving for a site-specific effort  
51 multiplier  $\alpha_i$  such that

$$\phi_i(\alpha_i) = \frac{\sum_j [1 - (1 - f_{ij})^{\alpha_i}]^2}{\sum_j [1 - (1 - f_{ij})^{\alpha_i}]} = \Phi.$$

52 Mathematically,  $\Phi$  is chosen so that every neighbourhood’s weighted-mean  
53 frequency  $\phi_i = \sum_j f_{ij}^2 / \sum_j f_{ij}$  equals  $\Phi$ . Hill [6] showed that  $\phi_i$  is the ratio of  
54 the mean species richness to the ‘effective number of common species’ (often  
55 called  $N_2$ , the reciprocal of Simpson’s index; Hill [5]), which means that  $\phi_i$   
56 isolates sampling intensity from true differences in richness and evenness. By  
57 fixing  $\phi_i = \Phi$ , we therefore align all neighbourhoods to the same effort scale  
58 without erasing real ecological differences.

59 This process yields the standardised neighbourhood frequencies

$$\tilde{f}_{ij} = 1 - (1 - f_{ij})^{\alpha_i}$$

60 which are independent of time (i.e. they are calculated with respect to the  
61 entire time period under consideration, rather than any subdivisions of this used  
62 for trend calculations), and serve as a proxy for the “true” discoverability- or  
63 effort-standardised neighbourhood species rank-frequency curve.

### 64 3.2. Temporal correction

65 Within each time period  $t$ , one chooses a set of “benchmark” species [8] and  
66 computes the proportion recorded per site and time period (Hill’s  $s_{it}$ ) as an index  
67 of site-level recording effort. (Note that there are potentially many ways to choose  
68 ones’ neighbourhood benchmarks, but Hill [6] proposed a fixed proportion  $R^*$  of  
69 the standardised species rank-frequency curve after an additional normalisation  
70 step involving the division of species’ ranks by the expected species count  $\sum_j \tilde{f}_{ij}$ ;  
71 however, the precise method of choosing benchmarks does not affect what follows).  
72 For each species  $j$  in period  $t$ , Hill then defines a Poisson-link intensity

$$Q_{ijt} = -\ln[1 - s_{it}\tilde{f}_{ij}],$$

73 The modelled “discovery” probability is then

$$P_{ijt}(x_{jt}) = 1 - \exp(-Q_{ijt}x_{jt}).$$

74 Hill [6] estimates the time-factor  $x_{jt}$  by matching the *total* modelled to *total*  
75 observed presences  $y_{ijt}$ :

$$\sum_i y_{ijt} = \sum_i P_{ijt}(x_{jt}).$$

76 In practice one iterates  $x_{jt}$  in the exact Poisson form above until those  
77 sums coincide (e.g. see the *R* code of Pescott [10]). The difference between  
78 the (summed) observed presences  $y_{ijt}$  and the model's baseline expectation  
79 after standardising time-independent neighbourhood effort  $\alpha_i$  and adjusting for  
80 site/time specific effort  $s_{it}$  is therefore captured by the time factor  $x_{jt}$ . Frescalo  
81 can thus deliver detection-corrected trends from unstructured data when its core  
82 assumptions are met.

#### 83 4. Bridging the gap

##### 84 4.1. Static occupancy and detection

85 We can compare the static (i.e. single season) single-species occupancy-  
86 detection model probability of at least one detection in  $v$  visits

$$\psi[1 - (1 - p)^v]$$

87 with the Poisson-process discovery rate (conditional on the all-time frequency  
88 curve) used in Frescalo

$$1 - e^{-\lambda s_{i(N)}}.$$

89 For small  $pv$ ,  $(1 - p)^v \approx e^{-pv}$ , hence  $\psi[1 - \exp(-pv)] \approx 1 - \exp(-\psi pv)$ ; now  
90 identifying  $\lambda = \psi p$  and  $v = s_{i(N)}$  recovers the approximate Frescalo detection  
91 probability  $1 - \exp(-\lambda s_{i(N)})$ . For any value of  $p$ , exact equivalence can be found  
92 by solving

$$1 - e^{-\lambda v} = \psi[1 - (1 - p)^v]$$

93 for

$$\lambda = -\frac{1}{v} \ln[1 - \psi(1 - (1 - p)^v)],$$

94 but this only reduces to  $\psi p$  in the limit  $pv \rightarrow 0$ . Frescalo's  $\lambda$  therefore combines  
95 occupancy and detectability into one Poisson rate, approximating the occupancy-  
96 detection model probability of at least one detection across  $v$  visits. Occupancy  $\psi$   
97 and per-visit detectability  $p$  collapse into  $\lambda_{ij} = \psi_{ij} p_{ij}$  and the unobserved number  
98 of site visits  $v$  in the Frescalo context becomes the continuous neighbourhood  
99 effort index  $s_{i(N)}$ , standardised across neighbourhoods via  $\alpha_i$ . Frescalo can  
100 therefore be interpreted as an occupancy-detection analogue at the neighbourhood  
101 scale: it replaces the two-parameters ( $\psi$ ,  $p$ ) and discrete  $v$  with a Poisson rate  $\lambda$   
102 and a continuous effort-multiplier  $\alpha$  equalising variable survey effort  $s_i$  (inferred  
103 by the neighbourhood level  $s_{i(N)}$ ) across sites.

104 A key step in recognising the equivalent elements of these models is to  
105 appreciate that Frescalo applies its discoverability standardisation at a large scale:  
106 not only is the adjustment done with respect to the multi-site neighbourhood and  
107 across all species, but it is also calculated across all time periods in the analysis.  
108 The standardised neighbourhood frequencies  $f_{ij}$  and the species rank-frequency  
109 curve they form is estimated once, independently of time, before temporal change  
110 is examined.

111 *4.2. Time trend interpretation*

112 A time trend in occupancy derived from a classical occupancy-detection  
113 model is modelled simply by letting  $\psi_{ij}$  vary linearly or non-linearly over time,  
114 conditional on both model-specific [16] and other standard survey sampling  
115 assumptions [3] being reasonable. Frescalo, by contrast, posits a single time-  
116 independent set of discoverability-adjusted baseline frequencies  $\tilde{f}_{ij}$ , and then uses  
117 benchmarks and the site/period effort index  $s_{it}$  to compute expected frequencies  
118 under an assumption of stasis, subsequently letting the time factors  $x_{jt}$  absorb  
119 any residual differences as true ecological change.

120 This underscores a key difference in how effort-adjustment processes func-  
121 tion in each model type. Occupancy-detection models assume that true site  
122 occupancies, and so trends in these, are directly recoverable from visit-level  
123 information. Frescalo assumes that fine-scale visit data is generally unavailable  
124 and/or uninformative for all or part of the time series of interest, and so models  
125 species' discoverability at a much larger scale. The main aim of this adjustment  
126 is to ensure a common scale across which neighbourhoods, and therefore sites,  
127 can be compared: without the harmonisation of effort across neighbourhoods,  
128 the time factors estimated for each site for a species would not be comparable,  
129 making average time factors and trends in these meaningless.

130 Another fundamental difference is the meaning of the site occupancy values  
131 produced. As noted,  $\psi_i$  has the simple meaning of predicted site occupancy under  
132 the classical model (notwithstanding debates around usage versus occupancy  
133 when these types of models are applied at different scales; [14]). The Frescalo  
134 time factor  $x_{jt}$  is, however, defined by the benchmark average, and values  $>1$  or  
135  $<1$  indicate that a species is at a higher or lower average frequency relative to the  
136 common species where it occurs, rather than in absolute occupancy probability.  
137 This may be an important limitation to inferring effort via observable recording  
138 outcomes, rather than having knowledge of those factors that directly map onto  
139 effort, such as the actual number of visits and covariates that are known to explain  
140 an important portion of observed variance in species' visit-level detectability  
141 [6, 15, 7].

142 One way around this issue is the observation of Bijlsma [1] that site occupancy  
143 probabilities can actually be derived from Frescalo via the combination of the  
144 standardised species' frequencies  $\tilde{f}_{ij}$  and the time-factors  $x_{jt}$ , and this has been  
145 exploited in at least one published analysis [4]. However, this requires a note of  
146 caution: whilst sensitivity analyses published in Hill [6] suggested that the trends  
147 in time-factors estimated by Frescalo can be relatively insensitive to the choice of  
148  $R^*$ , the benchmark threshold (variation in this parameter changing the intercept  
149 of estimated trends but not their slope), the same is not true of back-calculated  
150 site occupancy probabilities. Because the relationship between time-factors and  
151 species frequencies is non-linear, the shifts in time trend intercept seen using  
152 different values of  $R^*$  will not translate into the same proportional changes  
153 in predicted site occupancies over time (Pescott, pers. obs.) This may be  
154 particularly important when these trends are used to classify species' into risk  
155 categories, as for example happens in Red Listing exercises [e.g. 12].

156 **5. Conclusions**

157 Unstructured species occurrence data are too valuable to ignore, especially  
158 for historical periods where no information about the visit-level data collection  
159 process survives [11]. Hill’s “frequency scaling using local occupancy” or Frescalo  
160 method allows the careful analyst to infer a large-scale detectability or effort  
161 metric that can subsequently be used to place neighbourhoods on a common  
162 footing for the estimation of distribution trends. The large-scale formulation of  
163 this approach not only allows for the potential inclusion of more data sources  
164 (e.g. records extracted from Atlases or museums), but may also act to reduce  
165 the actual error in species’ trends intrinsically [2, 13].

166 By demonstrating how Frescalo collapses the classical occupancy-detection  
167 model’s  $\psi$  and  $p$  into  $\lambda$ , and how it infers visit-related effort via an emergent  
168 community-level mean rate, the approach performs an occupancy-detection-  
169 type correction even when explicit or informative temporal repeat-visit data  
170 are lacking. By highlighting this link, we hope to promote the development of  
171 additional diagnostics, extensions and more rigorous uncertainty quantification  
172 for the frequency scaling using local occupancy method.

173 **References**

- 174 [1] Bijlsma, R.J., 2013. The Estimation of Species Richness of Dutch Bryophytes  
175 between 1900 and 2011. Technical Report 15. Dutch Bryological and Licheno-  
176 logical Society. Gouda, the Netherlands.
- 177 [2] Boyd, R.J., Bowler, D.E., Isaac, N.J.B., Pescott, O.L., 2024. On the  
178 trade-off between accuracy and spatial resolution when estimating species  
179 occupancy from geographically biased samples. *Ecological Modelling* 493,  
180 110739. doi:10.1016/j.ecolmodel.2024.110739.
- 181 [3] Boyd, R.J., Powney, G.D., Pescott, O.L., 2023. We need to talk about  
182 nonprobability samples. *Trends in Ecology & Evolution* doi:10.1016/j.  
183 tree.2023.01.001.
- 184 [4] Eichenberg, D., Bowler, D.E., Bonn, A., Bruelheide, H., Grescho, V., Harter,  
185 D., Jandt, U., May, R., Winter, M., Jansen, F., 2021. Widespread decline in  
186 Central European plant diversity across six decades. *Global Change Biology*  
187 27, 1097–1110. doi:10.1111/gcb.15447.
- 188 [5] Hill, M.O., 1973. Diversity and evenness: A unifying notation and its  
189 consequences. *Ecology* 54, 427–432.
- 190 [6] Hill, M.O., 2012. Local frequency as a key to interpreting species occurrence  
191 data when recording effort is not known. *Methods in Ecology and Evolution*  
192 3, 195–205.

- 193 [7] Kéry, M., Royle, J.A., Schmid, H., Schaub, M., Volet, B., Häfliger, G.,  
194 Zbinden, N., 2010. Site-Occupancy Distribution Modeling to Correct  
195 Population-Trend Estimates Derived from Opportunistic Observations. *Con-*  
196 *servation Biology* 24, 1388–1397. doi:10.1111/j.1523-1739.2010.01479.  
197 **x**.
- 198 [8] Latour, J., van Swaay, C., 1992. Dagvlinders als indicatoren voor de  
199 regionale milieukwaliteit. *De Levende Natuur* 93, 19–22.
- 200 [9] MacKenzie, D.I., Nichols, J.D., Lachman, G.B., Droege, S., Royle, J.A.,  
201 Langtimm, C.A., 2002. Estimating Site Occupancy Rates When Detection  
202 Probabilities Are Less Than One. *Ecology* 83, 2248–2255. doi:10.1890/  
203 0012-9658(2002)083[2248:ESORWD]2.0.CO;2.
- 204 [10] Pescott, O., 2025. An R translation of Hill’s fortran code for Frescalo.  
205 doi:10.5281/zenodo.15305437.
- 206 [11] Pescott, O.L., Humphrey, T.A., Stroh, P.A., Walker, K.J., 2019. Temporal  
207 changes in distributions and the species atlas: How can British and Irish  
208 plant data shoulder the inferential burden? *British & Irish Botany* 1,  
209 250–282. doi:10.33928/bib.2019.01.250.
- 210 [12] Stroh, P., Leach, S., August, T., Walker, K., Pearman, D., Rumsey, F.,  
211 Harrower, C., Fay, M., Martin, J., Pankhurst, T., Preston, C., Taylor, I.,  
212 2014. A Vascular Plant Red List for England. *Botanical Society of Britain*  
213 *and Ireland*, Bristol.
- 214 [13] Stroh, P., Walker, K., Humphrey, T., Pescott, O., Burkmar, R. (Eds.), 2023.  
215 *Plant Atlas 2020. Mapping Changes in the Distribution of the British and*  
216 *Irish Flora*. *Botanical Society of Britain and Ireland & Princeton University*  
217 *Press*, Princeton.
- 218 [14] Valente, J.J., Jirinec, V., Leu, M., 2024. Thinking beyond the clo-  
219 sure assumption: Designing surveys for estimating biological truth with  
220 occupancy models. *Methods in Ecology and Evolution* 15, 2289–2300.  
221 doi:10.1111/2041-210X.14439.
- 222 [15] van Strien, A.J., van Swaay, C.A., Termaat, T., 2013. Opportunistic citizen  
223 science data of animal species produce reliable estimates of distribution  
224 trends if analysed with occupancy models. *Journal of Applied Ecology* 50,  
225 1450–1458. doi:10.1111/1365-2664.12158.
- 226 [16] Warton, D.I., Stoklosa, J., Guillera-Aroita, G., MacKenzie, D.I., Welsh,  
227 A.H., 2017. Graphical diagnostics for occupancy models with imperfect  
228 detection. *Methods in Ecology and Evolution* 8, 408–419. doi:10.1111/2041-  
229 210X.12761.