

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

Oliver L. Pescott*

^aUK 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 unified Poisson-process discovery rate and a temporal “time-factor” to capture trends. We demonstrate that the Bernoulli-trial formulation of occupancy-detection can be linked to Frescalo’s Poisson framework, with occupancy and detectability represented by a single rate parameter (which approximates the product ψp when overall sampling intensity is low). This connection clarifies how Frescalo’s neighbourhood-scale 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 less-structured biodiversity data.

Keywords: occupancy models, sampling effort, effort correction, citizen science, unstructured data, Hill numbers

1. Introduction

Occupancy-detection models [13] and the Frescalo “local frequency scaling” method [10] 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 (ψ) 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 (α) and species’ relative “time-factors” (x) 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 only poorly explained by

*Corresponding author
Preprint submitted to Elsevier
Email address: olipes@ceh.ac.uk (Oliver L. Pescott)

18 available covariates [15], understanding how Frescalo recovers effort-adjusted
 19 trends from aggregated data can broaden the toolkit of ecologists.

20 Whilst the place of occupancy-detection models in the quantitative ecologist’s
 21 armoury is well-established (e.g. MacKenzie et al. [13] has almost 6000 citations
 22 according to Google Scholar, May 2025, ~260 per year since 2002, a figure that
 23 is almost certainly a large underestimate of actual applications), Frescalo has
 24 only seen occasional use by comparison (143 citations, around 11 per year since
 25 2012). This may be due partly to the broader application of occupancy models,
 26 covering both small-scale monitoring and applications to less structured data at
 27 coarser scales [e.g. 20], but, even so, the scope for the use of Frescalo to derive
 28 time-trends and other metrics from unstructured data is likely to be larger than
 29 currently realised: within the outputs that have utilised the method feature
 30 a number of national species distribution Atlases [3, 18, 1], Red Lists [17, 7]
 31 and national biodiversity “status” reports [6]. Arguably then, an increase in
 32 the familiarity of ecologists with the approach would lead to even more such
 33 successful applications.

34 Although the two model types can appear quite different, Pescott et al. [15]
 35 informally suggested that Frescalo could be seen as a type of occupancy-detection
 36 model “where an adjustment for overlooked species is made in relation to spatial
 37 rather than temporal replication, whilst simultaneously adjusting for variable
 38 regional effort”. We here show that this suggestion can be formalised due to
 39 the two model types’ reliance on the same core mathematics of Bernoulli versus
 40 Poisson detections [cf. 16]. Below we (1) recall each framework, (2) write down
 41 their key equations, and (3) algebraically map one onto the other, demonstrating
 42 that Frescalo time-trends are based on an implicit occupancy-detection model
 43 whose “visits” and “occupancy” are folded into a single site/species discoverability
 44 rate parameter λ and standardised neighbourhood effort index $s_{(N)}$.

45 2. Occupancy-detection models

46 2.1. Basic single-season model

47 Following MacKenzie et al. [13], at each site i for species j assume a latent
 48 occupancy indicator

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

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

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

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

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

53 Thus the model simultaneously estimates

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

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

55 3. Frequency scaling using local occupancy (Frescalo)

56 3.1. Neighbourhood frequencies

57 Frescalo [10] pools presence-only data across a neighbourhood around target
 58 site i . We denote the observed proportion of neighbourhood sites in which
 59 species j was recorded by f_{ij} (in practice this frequency may relate to a weighted
 60 neighbourhood as per Hill [10], but this detail is not crucial for what follows).
 61 Under a Poisson-process model of species discovery with rate λ_{ij} and unknown
 62 total neighbourhood-level sampling effort $s_{i(N)}$, one has

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

63 Subsequently, a frequency-weighted neighbourhood index

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

64 is then “standardised” to a target value Φ by solving for a site-specific effort
 65 multiplier α_i such that

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

66 Mathematically, Φ is chosen so that every neighbourhood’s weighted-mean
 67 frequency $\phi_i = \sum_j f_{ij}^2 / \sum_j f_{ij}$ equals Φ . Hill [10] showed that ϕ_i is equivalent to
 68 the ratio of the neighbourhood’s mean species richness to the ‘effective number
 69 of common species’ (often called N_2 , the reciprocal of Simpson’s index; Hill
 70 [9]), which means that ϕ_i isolates neighbourhood sampling intensity from true
 71 differences in richness and evenness. By fixing $\phi_i = \Phi$, we therefore align all
 72 neighbourhoods to the same effort scale without erasing real ecological differences.

73 This process yields the standardised neighbourhood frequencies

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

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

78 3.2. Temporal correction

79 Within each time period t , one chooses a set of local “benchmark” species [12]
 80 and computes the proportion recorded per site and time period (Hill’s s_{it}) as an
 81 index of site-level recording effort. (Note that there are potentially many ways
 82 to choose ones’ site benchmarks, but Hill [10] proposed a fixed proportion R^* of
 83 the standardised neighbourhood species rank-frequency curve after an additional
 84 normalisation step involving the division of species’ ranks by the expected species
 85 count $\sum_j \tilde{f}_{ij}$; however, the precise method of choosing benchmarks does not

86 affect what follows). For each species j in period t , Hill then defines a Poisson-link
 87 intensity

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

88 The modelled “discovery” probability is then

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

89 Hill [10] estimates the time-factor x_{jt} by matching the total modelled to total
 90 observed presences y_{ijt} :

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

91 In practice one can iterate x_{jt} in the exact Poisson form above until those sums
 92 coincide (e.g. see the R code of Pescott [14]), although analytical solutions are
 93 also possible (J.M. Yearsley, pers. comm.) The difference between the (summed)
 94 observed presences y_{ijt} and the model’s baseline expectation after standardising
 95 time-independent neighbourhood effort α_i and adjusting for site/time specific
 96 effort s_{it} is therefore captured by the time factor x_{jt} . Frescalo can thus deliver
 97 detection-corrected trends from unstructured data when its core assumptions
 98 are met.

99 4. Bridging the gap

100 4.1. Static occupancy and detection

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

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

103 with the Poisson-process discovery rate (conditional on species’ presence in the
 104 all-time frequency curve) used in Frescalo

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

105 For modest values of the product pv , one may use the standard Taylor-series
 106 fact

$$(1 - p)^v \approx e^{-pv} \quad \text{for } pv \ll 1,$$

107 which turns $\psi[1 - (1 - p)^v]$ into $\psi[1 - e^{-pv}]$. On the other hand, setting $\lambda = \psi p$
 108 and $v = s_{i(N)}$, Frescalo’s Poisson form $1 - e^{-\psi pv}$ expands in exactly the same
 109 way: to first order both are $\psi(pv)$ with only $O((pv)^2)$ differences (i.e. only
 110 quadratic and higher terms differ). We therefore recover Frescalo’s $1 - e^{-\lambda s_{i(N)}}$
 111 approximately whenever pv is small.

For larger pv , the neglected second-order terms no longer agree, so the approximation is lost. However, one can always recover the exact Poisson rate by solving:

$$1 - e^{-\lambda v} = \psi[1 - (1 - p)^v] \implies \lambda = -\frac{1}{v} \ln[1 - \psi(1 - (1 - p)^v)], \quad (2)$$

but that formula reduces to $\lambda = \psi p$ only in the limit $pv \rightarrow 0$.

Frescalo's Poisson rate λ is therefore exactly the function of occupancy, detectability and visit count that makes the first part of equation (2) true [cf. 16]. Whilst in Frescalo we never observe v directly, we infer it via the continuous neighbourhood effort index $s_{i(N)}$, standardised across all neighbourhoods by the spatial scaler α_i . Frescalo can therefore be interpreted as an occupancy-detection analogue at the neighbourhood scale: it replaces the two-parameters (ψ , p) and discrete v with a Poisson rate λ and a continuous effort-multiplier (α_i) equalising variable survey effort s_i (inferred by the neighbourhood level $s_{i(N)}$) across sites.

Underpinning all of this is the assumption that, within any neighbourhood, the sequence of species/site detections behaves like a multi-species Poisson point-process. It is the assumption which justifies the log-link given in equation (1) above, and the Poisson moment relations that let us quantify and standardise sampling effort via ϕ_i . When sampling intensifies (so that pv is no longer small), higher-order Poisson moments become more important and the simple $\psi p \approx \lambda$ approximation breaks down.

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

4.2. Time-trend interpretation

A time-trend in occupancy derived from a classical occupancy-detection model is modelled simply by letting ψ_{ij} vary linearly or non-linearly over time, conditional on both model-specific [21, 22] and other standard survey sampling assumptions [5] being reasonable. Frescalo, by contrast, posits a single time-independent set of discoverability-adjusted baseline frequencies \tilde{f}_{ij} , and then uses benchmarks and the site/period effort index s_{it} to compute standardised frequencies under an assumption of stasis, subsequently letting the time-factors x_{jt} absorb any residual differences as true ecological change.

This underscores a key difference in how effort-adjustment processes function in each model type. Occupancy-detection models assume that true site occupancies, and so trends in these, are directly recoverable from visit-level information; Frescalo assumes that fine-scale visit data is generally unavailable and/or uninformative for all or part of the time series of interest, and so models species' discoverability at a much larger scale. The main aim of this adjustment

153 is to ensure a common scale across which neighbourhoods, and therefore sites,
 154 can be compared: without the harmonisation of effort across neighbourhoods,
 155 the time-factors estimated for each site for a species would not be comparable,
 156 making average time-factors and trends in these meaningless.

157 Another fundamental difference is the meaning of the site occupancy values
 158 produced. As noted, ψ_{ij} has the simple meaning of predicted species' site
 159 occupancy under the classical model (notwithstanding debates around usage
 160 versus occupancy when these types of models are applied at different scales;
 161 [19]). The Frescalo time-factor x_{jt} is, however, defined relative to the benchmark
 162 average, and values >1 or <1 indicate that a species is at a higher or lower
 163 average frequency relative to the common species where it occurs, rather than in
 164 absolute occupancy probability. This may be an important limitation to inferring
 165 effort via observable recording outcomes, as opposed to having knowledge of
 166 those factors that directly map onto effort, such as the actual number of visits
 167 and covariates that are known to explain an important portion of observed
 168 variance in species' visit-level detectability [10, 20, 11].

169 One way around this issue is the observation of Bijlsma [2] that site occupancy
 170 probabilities can be back-calculated from Frescalo via the combination of the
 171 standardised species' frequencies \hat{f}_{ij} , the species' time-factors x_{jt} , and by setting
 172 $s_{it} = 1$ across all sites and time periods (i.e. constant effort), and this has
 173 been exploited in at least one published analysis [8]. However, this requires a
 174 note of caution: whilst sensitivity analyses published in Hill [10] suggest that
 175 trends in time-factors estimated by Frescalo can be relatively insensitive to the
 176 choice of R^* , the benchmark threshold (variation in this parameter changing
 177 the intercept of estimated trends but not their slope), the same is not true of
 178 back-calculated site occupancy probabilities (P_{ijt} in Frescalo terms). Because
 179 the relationship between time-factors and species' frequencies is non-linear, the
 180 shifts in time-trend intercept seen using different values of R^* will not translate
 181 into the same proportional changes in predicted site occupancies over time. This
 182 may be particularly important when these trends are used to classify species'
 183 into risk categories, as for example happens in Red Listing exercises [e.g. 17].

184 5. Conclusions

185 Unstructured species occurrence data are too valuable to ignore, especially
 186 for historical periods where no information about the visit-level data collection
 187 process survives [15]. Hill's "frequency scaling using local occupancy" or Frescalo
 188 method allows the careful analyst to infer a large-scale detectability metric that
 189 can subsequently be used to place neighbourhoods on a common footing for the
 190 estimation of distribution trends. The large-scale formulation of this approach
 191 not only allows for the potential inclusion of more data sources (e.g. records
 192 extracted from Atlases or museums), but may also act to reduce the actual error
 193 in species' trends intrinsically [4, 18].

194 By demonstrating how Frescalo represents the classical occupancy-detection
 195 model's ψ and p with λ , and how it infers visit-related effort via an emergent

community-level mean rate ϕ , the approach performs an occupancy-detection-type correction even when explicit or informative temporal repeat-visit data are lacking. By highlighting this link, we hope to promote the development of additional diagnostics, extensions and more rigorous uncertainty quantification for the frequency scaling using local occupancy method.

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