**Mortality and recruitment in a woodland undergoing collapse**

**Introduction**

* Mortality and recruitment are vital for understanding forest dynamics and recent studies have suggested alteration of both

An understanding of what influences tree mortality and recruitment is vital to understanding forest dynamics. As a result of changes in fire regimes, increases in water deficit and novel pests and pathogens numerous recent studies have shown increases in tree mortality and reductions in recruitment (van Mantgem *et al.* 2009; Peng *et al.* 2011; Mcintyre *et al.* 2014). In forests where individual species recruitment has not kept pace with mortality, such as in the Western USA, this can lead to alteration of forest composition (Mcintyre *et al.* 2014). Where mortality is particularly high, as seen in Eastern Amazonian forests where repeated fires have taken place, this can result in shifts to relatively treeless states (Barlow & Peres 2008; Brando *et al.* 2014). Any such transitions from forest to a non-forest ecosystem would inevitably result in large emissions of carbon as well as potentially negative impacts on other ecosystem services and conservation priorities.

* Identification of what drives tree recruitment and mortality could aid production of indicators of resilience

While numerous studies have details of increases in mortality over recent decades relatively few identify what factors determine the likelihood of tree death when a transition is occurring. Coomes *et al.* (2003) suggest that in the event of disturbance larger trees are likely to be more prone to death than small or intermediate trees and this appears to be true in many cases where transitions to non-forest states are occurring (Barlow *et al.* 2002; Mcintyre *et al.* 2014). Given that large trees produce large amounts of seeds (Greene & Johnson 1994) and play a vital role in maintaining the microclimatic conditions of forests by limiting the light reaching the forest floor their loss may have important consequences for recruitment. In addition to tree size it has been widely seen that slow growing trees tend to have increased risk of mortality. This observation may be particularly useful in the context of forests at risk of transitions to non-forest states since it may serve as an early warning indicator of low resilience (Camarero *et al.* 2015).

* In addition there is a potential in systems that positive feedbacks can drive system transitions

Positive feedbacks are important for maintaining ecosystems in their state. For example, in forests the presence of a canopy maintains relatively moist, dark conditions on the forest floor which results in relatively low ground flora biomass. However, unfavourable positive feedbacks ecosystems can cause transition from one state to another such as when forest canopies are disturbed leading to reduced seedling recruitment and invasion by grasses and shrubs in Mediterranean climates (Acácio *et al.* 2007). Positive feedback can also occur between different drivers such as when degradation and fragmentation increase fire risk in forest (Benchimol & Peres 2015). However, feedbacks such as these can occur at multiple different scales and thus can be difficult to identify (Reyer *et al.* 2015).

* Though there are numerous studies on forests in a state of transition to relatively treeless states few investigate the processes that affect individual trees to cause such changes.

**Statement of aims**

* To test the influence of tree size, growth rate, competition and proximity to dead trees on determining tree mortality.
* To test the factors what influenced seedling and sapling density to enable an understanding of recruitment
* Test the influence of tree size and competition on growth rate
* Use parameter estimates from these models along with estimates from the literature to produce an individual based model to look at the potential for regime shift in the system, in particular:
  + Hysteresis – does the system recover along the same trajectory it declined over

**Hypotheses**

* Recruitment did not keep pace with mortality for any species during any of the survey periods
* Mortality likelihood is u-shaped in relation to tree size
* Slow growing trees are more likely to die
* Proximity to trees that have died is more likely to result in tree death
* Under current conditions the woodland will show increased loss of tree cover. In situations where recruitment is maximized dramatic losses will be averted.

**Methods**

**Site history and characteristics**

The site used in this study, Denny Wood, is located in the New Forest National Park in Southern England (Lat: 50o 51.5’ N and 1o 32.5’ W). Detailed site descriptions are provided by Mountford *et al*. (1999), and are briefly summarized here. The site is situated on gentle slopes (1–3°), primarily on clay-rich brown earth soils (pH 4.5–5.0 at 10 cm depth) that are prone to winter waterlogging and summer drying, with localised areas of strongly acid (pH 3.5–4.5) podsols developed on sandier substrates. Woodland vegetation is dominated by old-growth beech (*Fagus sylvatica*)withfrequent pedunculate oak (*Quercus* robur) and birch (*Betula pendula, B. pubescens*), and an understory primarily of holly (*Ilex aquifolium*). In open areas the ground vegetation is mostly comprised of *Agrostis*-dominated grassland or stands of bracken (*Pteridium aquilinum*); *Rubus fruticosus* agg., *Juncus effusus* and *Molinia caerulea* are also locally abundant. There are large populations of deer, ponies and cattle int eh New Forest, resulting in high herbivore pressure. The portion of the site which we use for this study is likely to have experienced high browsing pressure since the 1960s (Mountford & Peterken 2003).

**Data collection**

Measurements were conducted in one 20 m-wide 1 km long transect, which was originally established in the 1950s. The transect was subdivided into contiguous 51 20 x 20 m (0.04 ha) subplot and surveyed in 1964, 1984, 1988, 1996 and 2014. Details of earlier measurements are presented by Mountford et al. (1999) and Mountford and Peterken (2003) and those for 2014 in Martin et al. (2015). In each survey, the location and species name of all woody stems >1.3 m in height were recorded, their diameter at breast height measured using diameter tapes, and their status assessed as either alive or dead. Each stem >1.3m height was given a unique ID number to allow changes in individual trees to be tracked.

In 2014 a number of variables were not recorded in previous surveys were also collected: seedling density, canopy openness, soil types and, proxies of herbivore pressure. The density of tree seedlings of all species present in 10 x 10 m plots located in the centre of the 20 x 20 m plots was recorded. In addition in 2014 the canopy openness of plots was assessed using a concave spherical densitometer in all four corners and the centre of 20 x 20 m plots and calculating the mean of these values. In 2014 the soil types in each plot were assessed by sampling from within each 20 x 20m plot and then processing these samples in the lab. Herbivore pressure was assessed by counting the amount and condition of dung for both ponies and deer in each plot.

**Statistical analysis**

**Individual tree mortality**

Statistical models of individual tree mortality were developed using logistic mixed effects models, which describe the probability that a tree dies in a given period of time. For this model we tested largely individual-level predictors of mortality. Models were developed in a four-step process similar to the workflow of Chao et al. (2008). In step 1 we prepared predictors classified into 3 groups: (i) tree size, (ii) tree growth, (iii) proximity to dead trees. Tree size represented diameter based measurements prior to death, specifically diameter at breast height, basal area and relative size. Tree growth was based on measurements of diameter at two separate time periods prior to death and represented both relative and absolute growth rates. Proximity to dead trees related to the prior to death distance of an individual tree to a dead tree or the abundance of dead trees in a 10m buffer.

In step 2 we selected we selected the best predictor for each group by choosing the univariate logistic mixed effect models which had the lowest AICc (Burnham & Anderson 2002; Chao *et al.* 2008). This step reduces intercorrelation of variables which can lead to difficulty in interpreting effects. In step 3 a full multivariate model was developed using these selected variables. In step 4 model averaging was used to produce parameter estimates using the most parsimonious models. Each of these steps is described in more detail below.

During data processing tree status (alive or dead) was coded at census time . Only trees with diameter measurements at both and prior to time were included in analyses. Censuses on the transect were undertaken 5 times from 1964-2014 with a mean (± SD) census interval period of 12.5 ± 6.7 years (range 4-20 years). Since trees used in mortality models required censuses at both and prior to time we selected three non-overlapping census periods: 1984-1988, 1988-1996 and 1996-2014 (mean census period 10 ± 5.9 years). To correct for the variation in census interval we used a complementary log-log link with an offset equal to the census interval. Plot ID was used as a random effect to account for repeated sampling of the same plots.

Tree size metrics used were (i) DBH (cm): diameter at height of 1.3m of tree at time ; (ii) BA (m2): basal area of tree at time ; (iii) DBH (cm) and DBH2 (m2) – used to indicate possible u-shaped mortality curve (Holzwarth *et al.* 2013) and; (iv) The relative size of tree at time given with the equation:

For the biggest tree in the transect is close to 1 for the smallest it is close to 0 (Monserud & Sterba 1999).

Tree growth rate metrics used were: (i) Annual diameter growth rate (mm year-1) calculated as:

where and are the *DBH* values for a tree at time and respectively; (ii) Basal area growth rate (cm2 year-1) calculated as:

where and are the *DBH* values for a tree at time and respectively; (iii) relative DBH growth rate (% DBH increase year-1) calculated as:

and; (iv) relative BA growth rate (% BA increase year-1) calculated as:

Metrics of proximity to dead trees were: (i) *deadSD*=stem density of dead trees at time within 10 metres of the target tree; (ii) *deadSD45*=stem density of dead trees ≥45cm DBH at time of death at time within 10 metres of the target tree; (iii) *deadDist*=distance to nearest dead tree (m) at time ; (iv) *deadDist45*=distance to nearest dead tree ≥45cm DBH at time of death (m) at time . Our rationale for focussing on dead trees ≥45cm DBH was that this represents the upper quartile of tree diameters in the woodland and we hypothesised that the death of larger trees plays a stronger role in altering local microclimate conditions.

As well as individual based variables that may affect tree mortality we assessed the importance of soil properties as well, given that finer soils may be prone to water logging in the winter, which impedes root growth and consequently reduces the ability of trees to take advantage of water deeper in the soil in dry summers.

* Poisson mixed models were used to assess how density of saplings and seedlings changed over time and with density of mature trees

**Individual based models of recruitment and mortality**

* An individual based model built in netlogo was parameterised using values from these analyses to investigate the potential changes in the woodland

**Results**

**Discussion**

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