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Climate warming increases insect-driven seed removal of two elaiosome-bearing invasive thistle species

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**Data accessibility**

All data and code are on GitHub at <https://github.com/TrevorHD/InsectSeedRemoval> and will be made public upon acceptance for publication.

**Key Words**

Ants*, Carduus acanthoides*, *Carduus nutans*, climate change, dispersal, invasive species, myrmecochory, seed removal

**Abstract**

Ants and other insects are often a source of localised secondary dispersal for wind-dispersed plants, and thus play an important ecological role in their spatial dynamics, but there is limited information on how climate change will affect such dispersal processes. Here, we use field experiments to investigate how climate warming affects seed removal; this initiation of movement represents the first step in insect-driven secondary dispersal. We find that increased growing temperature influences seed attractiveness to insect dispersers in the invasive thistles *Carduus nutans* and *Carduus acanthoides*. Seeds from maternal plants grown at temperatures 0.6 °C higher than ambient are removed at higher rates by insect dispersers than their unwarmed counterparts. We also observe that seed elaiosomes in these two species play an important role in dispersal, as seeds without elaiosomes were significantly less likely to be removed by insects over the same time period. Significant interaction effects between elaiosome presence/absence and warming treatment were also observed, though only for *C. acanthoides*, with the boost in seed removal from warming dampened when the elaiosome was present compared to when it was absent. These findings provide evidence that climate warming may alter aspects of the dispersal process such as seed removal, and have global ramifications for dispersal in future climates since seed-bearing plants across the world may be subject to increased growing temperatures, and many these plant species bear elaiosomes and experience seed dispersal by insects.

**Introduction**

For sedentary organisms such as plants, seed dispersal is the only stage in the life cycle during which movement occurs, and is thus an essential component of plant spatial dynamics. While plant propagules are often dispersed by abiotic vectors such as wind, water, and gravity, they may also be dispersed biotically by organisms such as insects, birds, and mammals. Such dispersal of seeds by biotic vectors typically occurs as part of a larger network of parallel and serial dispersal events, both primary and secondary in nature (Figure 1). Integration of these primary and secondary pathways and the vectors responsible into a total dispersal kernel, or probability distribution of how far seeds are dispersed when accounting for all possible dispersal vectors, can help us quantify how plant populations move or expand at a variety of different spatial scales (Nathan 2007). However, significant challenges in identifying these biotic dispersers and quantifying how they disperse seeds make it challenging to fully understand their role in seed dispersal, especially for smaller seeds that cannot be easily tracked through empirical means (Rogers *et al*. 2019).

For these smaller seeds, ants and other insects are often an overlooked source of localised secondary dispersal after primary dispersal by wind, water, or even another biotic vector (Vander Wall and Longland 2004). In particular, ant-mediated seed dispersal, or myrmecochory, has been shown to be a common occurrence in many different plant species. In more than 80 plant families across the world, ant-dispersed plant species contain seeds that bear an elaiosome, a small structure on the distal end of the seed achene that facilitates removal and dispersal by ants (Edwards *et al*. 2006). The elaiosome has a high lipid content and likely serves as a reward for the ant dispersing the seed (Brew *et al*. 1989; Hughes and Westoby 1990, 1992), as ants will typically take the seeds back to their nest and consume the elaiosome, then store the seed achene in a midden (Berg 1975; Culver and Beattie 1978). This dispersal can be advantageous for a variety of different reasons, including moving propagules to areas with better germination and growing conditions as well as less competition from other plants (Handel and Beattie 1990). Removal of a seed by ants does not guarantee successful dispersal of that seed, as seeds can be consumed or destroyed in the process; nonetheless, it is a critical step in secondary dispersal because initiates one or more secondary dispersal processes that move seeds further from the parent plant (Jongejans *et al*. 2015a).

Investigations of ant-mediated seed removal can be quite challenging, however. This challenge is further complicated by climate change, where changes such as increases in temperature may affect various characteristics that determine how seeds move or are moved. For example, changes in seed production may result from elevated growing temperatures (Sato *et al*. 2006, Hedhly *et al*. 2009, Caignard *et al*. 2017) and increased CO2 levels (Edwards *et al*. 2001, Thürig *et al*. 2003, Way *et al.* 2010), thus affecting the number of seeds available for secondary dispersal. The dynamics of seed release can also be affected by increased temperatures, with some species more likely to release their seeds (Teller *et al*. 2016), also affecting the number of seeds available for secondary dispersal. Climate change may also cause shifts in the range or abundance of animal and insect species that remove seeds, as well as shifts in the traits that attract these species, with potential consequences for dispersal patterns (Ruxton and Schaefer 2012, Mokany *et al*. 2014, Sales *et al*. 2021). Even seed nutrient content has the potential to be affected by climate change and various studies, especially on agricultural crops, have shown changes in nutrient allocations and chemical makeup due to factors such as increased temperatures or elevated CO2 (Thomas *et al*. 2003, Caldwell *et al*. 2005, Singh *et al*. 2013). However, there are still gaps in knowledge regarding how climate-driven changes in seed shape, size, or nutrient allocations affect seed attractiveness to dispersers or how these dispersers move seeds. Addressing these gaps must start with better understanding the underlying dispersal mechanisms themselves (Travis *et al*. 2013).

Given the role that ants and other insects play in seed dispersal, studying how they move seeds can facilitate a better understanding of how certain plant species spread. Such an understanding starts with quantifying seed removal, as this constitutes the first step of secondary dispersal (Jongejans *et al*. 2015b). A common experimental setup to quantify seed removal by secondary dispersers consists of so-called “cafeterias” or “depots” (e.g. Xiao *et al*. 2006, Fischer and Türke 2016, Gurney *et al*. 2015, Linabury *et al*. 2019, LoPresti *et al*. 2022), where seeds are aggregated and the number of seeds removed is monitored. When paired with video recording or exclusion treatments such as cages, seed removal experiments allow researchers to observe the types of animals or insects removing seeds and approximate their contribution of rates to seed removal (Jongejans *et al*. 2015b, Griffiths *et al*. 2018). In the face of challenges associated with quantifying secondary dispersal processes and documenting the ultimate fate of removed seeds (Vander Wall *et al*. 2005a), these experiments provide tractable starting points for further inquiry into identifying secondary dispersal vectors and quantifying their contributions to the secondary dispersal.

Here, we seek to better understand secondary dispersal by investigating removal of seeds from two invasive, non-native thistle species, while noting that removal studies alone cannot distinguish ultimate seed fate (secondary dispersal or predation). We also seek to explore to what extent the elaiosome, as well as increased growing temperatures of maternal plants, affects seed removal rates and attractiveness to insect dispersers. Thus, we investigate four questions. First, at what rate are seeds removed via insect after a seed has experienced primary dispersal? Second, does the seed elaiosome significantly affect the rate of removal? Third, are seeds from maternal plants growing in increased temperatures any more or less likely to be removed by insects than their unwarmed counterparts? And fourth, does the effect of the seed elaiosome on removal rates change based on whether or not the maternal plant is warmed? These questions will allow us to better understand how climate warming may affect secondary dispersal of these important invasive species, with potential implications for the many other elaiosome-bearing plant species across the world that will be subjected to increased growing temperatures in future climates.

**Methods and materials**

***Study Species***

*Carduus nutans* L. and *Carduus acanthoides* L. are invasive thistles in the Asteraceae family that, while native to Eurasia, have spread across the world and are listed as noxious weeds across much of the United States (Skinner *et al.* 2000). These thistles are unpalatable to grazing animals (Trumble and Kok 1982), and are highly successful due to high germination rates and the large numbers of seeds they produce (Desrochers *et al*. 1988). They thrive in highly disturbed areas such as pastures, roadsides, railways, and utility corridors. Both species display monocarpic perennial reproductive behaviour where, while typically biennial, can experience increased annualism under increased growing temperatures (Keller and Shea 2021). Reproduction in both species occurs exclusively by seed and dispersal occurs primarily by wind, with dispersal facilitated by a lightweight pappus attached to the distal end of the seed achene. Seeds from both species also display nutrient-rich elaiosomes on the distal end of the achene that likely play a role in ant-driven dispersal (Weiss 1908, Pemberton and Irving 1990, Mayer *et al*. 2005). Ants and other insects have been observed moving seeds in previous seed removal experiments (Jongejans *et al*. 2015b).

***Seed collection and preparation***

*C. nutans* and *C. acanthoides* from which seeds were harvested were grown under control and ambient warming treatments as part of an experiment in Drees and Shea (*in prep.*) to estimate the effects of warming on the distribution of flower heights; that publication discusses the methods and experimental setup in greater detail, but essentials are noted here. Individuals of each species were grown in a grid-like arrangement, and a subset of individuals within each species were randomly assigned a fibreglass open-top chamber. These chambers, built to specifications of the International Tundra Experiment Manual (Molau and Mølgaard 1996), have been demonstrated to increase the ambient temperature by approximately 0.6 °C on average over a year (Zhang *et al*. 2011) without significantly affecting other possible growth factors such as soil moisture and snow depth. When plants set seed, mesh pollen bags were wrapped around the flower heads to ensure that seeds did not escape and contaminate other parts of the study area.

Once individuals completed their life cycle, they were cut down, and bagged seed heads were collected from all individuals that produced five or more viable flowers. During this process, seed heads from individuals of the same species were mixed together to randomise possible differences in seed size, shape, nutrient content, and any other factors that vary between individuals and may affect seed attractiveness to insects. The pollen bags were then removed from the seed heads, and the heads were placed in paper bags and allowed to desiccate in a dry, low-light environment at approximately 24 °C. After approximately a month of desiccation, seeds were separated from their pappi and flower heads, and were then stored in an airtight container. Seeds were then irradiated to ensure they could be freely dispersed by ants and other insects without spreading these invasive thistles and contaminating the study site. Irradiation was performed with a 1000 KR dosage of gamma radiation; this dosage was demonstrated by Jongejans *et al*. (2015b) to prevent any germination without affecting attractiveness to insects (O’Hanlon *et al.* 2019).

Once seeds were irradiated, a subset of seeds from each of the four combinations of species and warming/ambient maternal treatment were assigned to an elaoisome removal treatment. Elaiosomes were removed by holding the seed with a pair of forceps, aligning the flattest side of the seed parallel to the workbench, and then using a teasing needle to press the elaiosome against the workbench until it was severed from the seed achene. In instances where the elaiosome did not cleanly separate, the teasing needle was used to gently dig out the remainder of the elaiosome from the achene. Seeds not receiving the elaiosome removal treatment were handled with forceps in the same manner mentioned above, for approximately the same duration as the seeds whose elaiosomes were removed; this was done so that possible differences in experimental outcomes could not be attributed to differences in the way the seeds were handled. After handling, seeds were again placed in airtight containers based on treatment type.

Seed depots were then built in a controlled environment before transport to the field site, where they allowed insects to easily access and remove seeds. All depots were constructed using petri dishes with a 95 mm diameter and 15 mm edge, with black sandpaper at the bottom to mimic the appearance of soil (Jongejans *et al*. 2015b). Sandpaper was glued to the bottom of the petri dish to prevent warping or movement of the paper due to wind, moisture, sunlight exposure, or insect activity. Each depot was arrayed with 25 seeds of a given treatment combination located near the centre of the petri dish, though spread out enough so that seeds were not piled on top of each other.

A portion of the irradiated seeds were also weighed to examine whether warming of the maternal plant had any significant effects on seed mass. Because the mass of individual *C. nutans* and *C. acanthoides* seeds is so low (typically a few milligrams each), seeds were weighed in groups of 20 to ensure that measurement error due to contamination, scale oscillation, etc. would be miniscule compared to the measured mass. For each of the four combinations of species and warming treatment, 30 groups of seeds with their elaiosomes intact were weighed.

***Experimental setup***

The seed removal experiment was conducted at the Russell E. Larson Agricultural Research Farm in Rock Springs, Pennsylvania. Seed depots were arranged over an 8 x 10 grid in a small field approximately 50 m from where the seeds were collected, with individual seed depots spaced 1 m apart. Seeds within a particular seed depot represented one of eight unique combinations from three fully crossed two-level factors of species (*C. nutans* vs *C. acanthoides*), warming treatment on the maternal plant (warmed vs unwarmed, with “maternal plant warming” henceforth referred to as “warming” for brevity), and elaiosome treatment (present vs absent); each row of eight depots contained a randomised arrangement of these eight unique treatment combinations, for a total of 10 depots for each treatment combination across the 8 blocked rows. Before placing each depot, the vegetation below was trimmed and compacted so that the petri dish could easily fit in the grass canopy, reducing the chance of depot disturbance by wind and ensuring easy access for insects.

Once all seed depots were placed, each seed depot was photographed from directly above by an observer at 30-minute intervals for the first 12 hours, from 09:00 to 21:00 on 07 September 2020; after 12 hours, depots were then photographed at 24 hours, 36 hours, and 48 hours. In instances where the sandpaper bottom of the dish was disturbed, it was fixed to the bottom of the petri dish again and the remaining seeds gently placed back around the centre of the seed depot, though such instances were extremely uncommon. No precipitation was observed over the duration of the experiment, and temperatures ranged between approximately 7°C and 27°C.

***Image processing***

All images were processed manually by counting the number of seeds for each seed depot at each recorded time; this was enabled by the contrast in colour between the seeds and the black sandpaper on the seed depots. Seeds were counted as removed only if they were completely removed from the seed depot; thus, even in instances where seeds were disturbed or scattered across the depot, they were still marked as present. For each image, the number of seeds was scored twice by the same observer across independent sessions to ensure accuracy of the data.

***Statistical analyses***

All statistical analyses were conducted in R version 4.2.2 (R Development Core Team 2022). To assess significance of the warming and elaiosome removal treatments, we used the **glmer** function from the package **lme4** version 1.1-31 (Bates *et al*. 2022) to fit a generalised linear mixed-effects model to the data, using a logit link and treating the number of seeds removed at a given time as a binomial response. Since between-species comparisons were not of primary interest, models were fit separately for each of the two species to facilitate ease of model interpretation, with elaiosome and warming treatments, and their interaction, encoded as fixed effects; treatment block was encoded as a random effect. To examine significance of the treatments at different points in time, models were fit separately at the 6-, 12-, and 24-hour marks similar to analyses performed by Jongejans *et al*. (2015b), and conditioned on the number of seeds at the previous time mark. Models were not fit at the 48-hour mark due to convergence issues, as many seed depots had zero seeds remaining at this point. Survival curves from observational data across all timesteps were constructed, and two-sided Kolmogorov-Smirnov tests were used to test whether survival curves for a given combination of species and one treatment differed between levels of the other treatment. The GLMs we fit are conceptually similar to a survival model that allows for a time-varying hazard; results from fitting parametric survival models are qualitatively similar (Appendix S1).

**Results**

Over the course of the 48 hours that seed removal was documented, both photographs and direct observation suggest that insects were responsible for seed removal; no birds or mammals were documented in the marked study area or were observed removing any seeds. Throughout the entire duration of the experiment, seed removal by ants was observed and made up the vast majority of direct observations of seed removal, with the ants dragging individual seeds across the flat surface of the seed depot (Appendix S2, Figure S1) before carrying the seed over the outer walls; these observations were especially common at the 30-minute observation periods during the daytime, but also occurred at several of the observation periods after sunset. Crickets were also observed on the seed depots (Appendix S2, Figure S2), though mostly at night, where they would consume the seeds on the depot rather than carry them away and consume elsewhere. Grasshoppers frequently visited the depots during the daytime hours but were not observed consuming or removing seeds, though they consumed or attempted to remove the adhesive tape used to mark the depots. Though a variety of other small insects were observed at the depots as well, none of them were observed removing any seeds and they were not identified further.

Observed rates of seed removal from depots were high, though varied between treatment combinations. On average, 24.7% (3.6%) of seeds were removed after 6 hours, 48.4% (4.5%) after 12 hours, 84.4% (3.2%) after 24 hours, and 94.6% (1.5%) after 48 hours, where plus/minus terms indicate one standard error. Rates of removal for warmed *C. nutans* with elaiosomes were particularly high compared to the overall averages, with 55.6% (13.1%) of seeds were removed after 6 hours, 79.6% (9.2%) after 12 hours, 98.0% (2.0%) after 24 hours, and 98.4% (1.6%) after 48 hours. For both *C. nutans* and *C. acanthoides*, removal of seeds with no warming treatment and no elaiosomes lagged behind that of the other treatment combinations (Appendix S2, Figure S3); however, removal at the end of the experiment was still high, with 87.2% 5.4%) removed for *C. nutans* and 88.9% (7.1%) removed for *C. acanthoides*.

Mixed-effect model coefficient estimates from Table 1 suggest that both warming treatment and elaiosome removal impacted the proportion of seeds removed, with significant effects at most of the timesteps at which models were fit. Here, seeds from warmed maternal plants were usually more likely to have been removed than those from unwarmed maternal plants, and seeds with their elaiosomes intact were more likely to have been removed than those without their elaiosomes. This reinforces trends seen in survival curves plotted from the observed seed counts over time, where seeds with their elaiosomes intact were generally removed at a faster rate than seeds without their elaiosomes for all combinations of warming and species, except for warmed *C. acanthoides* (Figure 2). Survival curves also show that seeds from warmed maternal plants were removed faster than seeds from unwarmed maternal plants for all combinations of elaiosome presence/absence and species, except for *C. acanthoides* with intact elaiosomes (Figure 3). While survival curves for warming/elaiosome treatment combinations were generally similar between species (Appendix S2, Figure S4), explicit comparisons to demonstrate whether differences in seed removal rates between species exist were not made since such a comparison was not a focus of this investigation.

Interactions between elaiosome and warming treatments were also observed in the mixed effects models. For *C. acanthoides*, significant negative interactions at all three time points suggest that presence of the elaiosome dampens the increase in seed removal facilitated by warming – that is, seeds with elaiosomes intact experience less of an increase in removal rates under warming than seeds where the elaiosome is removed (Table 1; refer to Appendix S2, Figure S5 for visualisation of marginal effects). This dampening effect on warming can be seen in Figure 3 as well, where juxtaposed survival curves clearly indicate lower seed removal for unwarmed *C. acanthoides* when the elaiosome is absent, but not necessarily when it is present. For *C. nutans*, evidence for interactions between elaiosome and warming treatments was not as clear; while there was a significant negative interaction similar to the one above at the 6-hour marks, positive interactions were observed at the 12- and 24-hour marks but were not significant.

For each species, there were significant differences in seed mass between the warmed and unwarmed treatment groups for both *C. nutans* (d.f. = 58, *t* = 4.506, *n* = 60, *p* < 0.001) and *C. acanthoides* (d.f. = 58, *t* = 2.832, *n* = 60, *p* = 0.006). A group of 20 warmed *C. nutans* seeds weighed on average 3.4 0.7 mg more than a group of 20 unwarmed seeds, where plus/minus terms indicate one standard error, representing an approximately 5.6% increase in mass. For *C. acanthoides*, the same number of warmed seeds weighed on average 1.4 0.5 mg more compared to the unwarmed seeds, representing an approximately 4.5% increase in mass.

**Discussion**

The prevalence in the ecological literature of studies regarding seed removal and secondary dispersal indicates a growing interest in identifying and quantifying the movement of seeds by organisms such as insects, animals, and humans (Anjos *et al*. 2020a). For example, seeds of the mahaleb cherry *Prunus mahaleb* are ingested, moved, and defecated by a variety of mammals such as foxes and badgers as well as birds such as crows, thrushes, warblers, and robins (Herrera and Jordano 1981; Guitián *et al*. 1992; Jordano *et al*. 2007). Multiple dispersal pathways exist in that system, with seeds consumed directly from the tree by birds, and also consumed from the ground to undergo secondary dispersal by birds and mammals after primary dispersal via gravity; these different pathways variously drive maintenance of existing mahaleb cherry populations versus formation of new populations. The acorns of various oak trees (*Quercus* sp.) are often dispersed and cached by a variety of rodents such as mice and squirrels (Jensen and Nielsen 1986; Vander Wall 2001; Gómez *et al*. 2008). Some species can even be dispersed by carnivores when they prey on frugivores or granivores with seeds in their digestive tract (Hämäläinen *et al*. 2017). Humans are also often involved in the process of dispersing plant propagules via attachment to clothing (Wichmann *et al*. 2009; Ansong and Pickering 2014) or vehicles (Veldman and Putz 2010; Taylor *et al*. 2012), as contaminants of horticultural stock (Hodkinson and Thompson 1997), and as impurities in agricultural produce such as grains (Shimono and Konuma 2008; Michael *et al*. 2010; Wilson *et al*. 2016). Here, we provide evidence of ant-mediated seed removal and dispersal, or myrmecochory, and demonstrate the importance of the elaiosome and its interaction with the effects of climate warming, in these processes.

Our results indicate that the rate of seed removal from the ground is very high, with approximately 84.4% of all seeds removed over the course of only 24 hours. These results are similar to observed removal rates of *C. nutans* and *C. acanthoides* in a similar study by Jongejans *et al*. (2015b), who demonstrate that up to 88% of seeds are removed in 24 hours when there is no impediment to the movement of insect dispersers. These high rates of seed removal suggest that the majority of *C. nutans* and *C. acanthoides* seeds dispersed via wind also experience secondary dispersal or consumption via insects or animals. Jongejans *et al*. (2015b) also demonstrate that rates of seed removal were significantly lower when restricting ant access to seeds, indicating that ants were responsible for the majority of seed removal events; this supports our observations of a large number of ant visits to seed depots. We further demonstrate that the elaiosome plays an important role in facilitating removal by ants, as seeds of both species had significantly higher removal rates when the elaiosome was left intact. Previous studies have suggested that these elaiosomes in *C. nutans* and *C. acanthoides* are involved in myrmecochory, attracting ants and playing an important role in how they disperse seeds (Pemberton and Irving 1990, Alba-Lynn and Henk 2010); the differences in seed removal rates between elaiosome-present and elaiosome-absent seeds in our study lend further support to evidence of myrmecochory in these thistle species.

In addition to removal of the elaiosome significantly impacting seed removal rates, we find that exposing the maternal plant to increased growing temperatures also increases the rate at which its seeds are removed by insect dispersers, with seeds from warmed plants experiencing higher rates of removal. We demonstrate that seeds from warmed maternal plants have higher mass, but do not yet know if there are changes in composition or nutrient content, nor whether that increase in mass went to the achene or elaiosome. As such, the effects of increased growing temperature on factors such as seed size or nutrient content, which likely play a role in how attractive insect dispersers find seeds (Anjos *et al*. 2020b), are still not clear for *C. nutans* and *C. acanthoides*. Additional analyses will be necessary to identify the underlying mechanisms responsible for these changes.

Our study is an important first step in quantifying secondary seed dispersal of thistle seeds by ants, but important challenges remain. While we can easily observe seeds being removed from a controlled experimental area, it is much more difficult to find where and how far dispersers move seeds after removing them from the seed depots. To address this challenge, there exist a variety of techniques to assess where animal-dispersed seeds are moved, with each having its own set of strengths and weaknesses. Larger seeds can hold electronic devices and be tracked via telemetry (e.g., Pons and Pausas 2007, Hirsch *et al*. 2012), and can thus be found without visual cues, which can be particularly useful in areas with dense vegetation or when seeds are cached underground. Seeds of various sizes can be painted or dyed with a fluorescent material or pigment that makes visual location easier (e.g. Levey and Sargent 2000, Reiter *et al*. 2006, Thomson 2007), but addition of pigment to seeds can change their mass depending on the size of the seed and amount of pigment applied (Lemke *et al*. 2009), which could possibly alter the dynamics of seed removal or dispersal; such pigmentation may also alter seed scent or taste. For ant-dispersed seeds such as *C. nutans* and *C. acanthoides*, pigment may not be useful if seeds are taken underground to ant nests, where they are then no longer visible from the surface. Mapping emerging seedlings within the vicinity of a seed depot can also provide some information on secondary dispersal distances, though for invasive species such as *C. nutans* and *C. acanthoides,* there are ethical concerns about allowing dispersal of viable seeds from an experiment into the environment (Jongejans *et al*. 2015b), hence our use of gamma irradiation to render the seeds unviable. Also, this method would only provide dispersal distances for a fraction of dispersed seeds because it only examines seeds that germinate, and would present the added challenge of ensuring that a particular seedling can be traced back to its source so that dispersal distance can be accurately quantified.

Given the difficulties with tracking small seeds such as those from *C. nutans* and *C. acanthoides*, while our study sheds light on the initiation of secondary dispersal by quantifying seed removal after primary dispersal, it does not document the fate of the seeds after they have been removed. Accurately assessing seed fate after removal can be quite challenging, as removed seeds are typically exposed to some mixture of dispersal and predation; while some seeds are consumed in a destructive manner or rendered inviable post-removal, others can be left intact by ants and scatter-hoarding animals (Hulme 1998, Vander Wall *et al*. 2005a, Penn and Crist 2018). Often, caching and burial improves germination rates by protecting seeds from predation after secondary dispersal, though seeds germinating after being cached at high densities may compete as seedlings (Hulme and Kollmann 2005). However, it is rarely known what proportion of seeds are dispersed or cached unharmed versus that experiencing predation. Despite this uncertainty in seed fate, many studies have conflated seed removal with predation, and often treat removed seeds as if they are consumed without substantial evidence to support this assumption (Vander Wall *et al*. 2005b). Such an assumption may not only overestimate actual rates of seed predation, but also underestimate secondary dispersal of seeds, as seed removal may not necessarily entail destruction but could instead initiate a series of secondary dispersal events (Vander Wall *et al*. 2005b). For *C. nutans* and *C. acanthoides*, the likely case is that some proportion of removed seeds are destroyed while others are dispersed, with a fraction of the dispersed seeds successfully germinating. For this reason, we have framed our results in terms of seed removal, since our uncertainty about the proportions of removed seeds that are consumed or dispersed limits us from making any claims about post-removal seed fate. Modelling of possible outcomes to assess their potential impacts would be a fruitful way to explore the importance of different fates (Vellend *et al*. 2006).

A better understanding of seed removal rates also has implications for spatial plant population dynamics. Given that most *C. nutans* and *C. acanthoides* seeds were removed after only 48 hours, almost all seeds already dispersed by wind may travel even further provided they are not subject to predation, and current estimates of how quickly these species spread might be conservative; current estimates of spatial spread (e.g., Skarpaas and Shea 2007, Jongejans *et al*. 2011, Zhang *et al*. 2011, Teller *et al*. 2016) only account for primary dispersal by wind. Secondary dispersal may also affect the germination and recruitment of new individuals by moving seeds to locations that may be more favourable to germination than their original location, or helping them escape density-dependent mortality from predation or parent-offspring competition (Janzen 1970, Connell 1971). Ultimately, while seed removal may result in the destruction of otherwise viable seeds, it also serves as the initiation of the secondary dispersal processes and thus plays a critical role in the movement of propagules. Though further research is needed, our study is a first step to fully understanding the potential for climate warming to increase secondary seed dispersal, and the resultant rate of population infilling and spread. Given that many plant families have elaiosome-bearing species and will be subject to increased growing temperatures, potential warming-induced shifts in secondary dispersal may be seen on a global scale.

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**Author Contributions**

T.H.D and K.S. designed the field experiment. T.H.D. conducted the field experiment, collected data, performed analyses, and led writing of the initial manuscript. T.H.D. and K.S. made contributions to subsequent versions of the manuscript.

**References**

Alba-Lynn, C. & Henk, S. (2010). Potential for ants and vertebrate predators to shape seed-dispersal dynamics of the invasive thistles Cirsium arvense and Carduus nutans in their introduced range (North America). *Plant Ecology*, 210(2), 291-301.

Anjos, D. V., Leal, L. C., Jordano, P., & Del‐Claro, K. (2020a). Ants as diaspore removers of non‐myrmecochorous plants: a meta‐analysis. *Oikos*, 129(6), 775-786.

Anjos, D.V., Andersen, A.N., Carvalho, R.L., Sousa, R.M., & Del‐Claro, K. (2020b). Switching roles from antagonist to mutualist: a harvester ant as a key seed disperser of a myrmecochorous plant. *Ecological Entomology*, 45(5), 1063-1070.

Ansong, M. & Pickering, C. (2014). Weed seeds on clothing: A global review. *Journal of* *Environmental Management*, 144, 203-211.

Bates, D., Maechler, M., Bolker, B., & Walker, S. (2022). Package ‘lme4’, version 1.1-31. https://cran.r-project.org/web/packages/lme4/index.html

Berg, R.Y. (1975). Myrmecochorous plants in Australia and their dispersal by ants. *Australian Journal of Botany*, 23(3), 475-508.

Brew, C.R., O'Dowd, D.J., & Rae, I.D. (1989). Seed dispersal by ants: behaviour-releasing compounds in elaiosomes. *Oecologia*, 80(4), 490-497.

Caignard, T., Kremer, A., Firmat, C., Nicolas, M., Venner, S., & Delzon, S. (2017). Increasing spring temperatures favor oak seed production in temperate areas. *Scientific Reports*, 7(1), 1-8.

Caldwell, C.R., Britz, S.J., & Mirecki, R.M. (2005). Effect of temperature, elevated carbon dioxide, and drought during seed development on the isoflavone content of dwarf soybean [Glycine max (L.) Merrill] grown in controlled environments. *Journal of Agricultural and Food Chemistry*, 53(4), 1125-1129.

Connell, J.H. (1971). On the role of natural enemies in preventing competitive exclusion in some marine animals and in rain forest trees. *Dynamics of Populations*, 298-312.

Culver, D.C. & Beattie, A.J. (1978). Myrmecochory in Viola: dynamics of seed-ant interactions in some West Virginia species. *The Journal of Ecology*, 66(1), 53-72.

Desrochers, A.M., Bain, J.F., & Warwick, S.I. (1988). The Biology of Canadian Weeds.: 89. Carduus nutans L. and Carduus acanthoides L. *Canadian Journal of Plant Science*, 68(4), 1053-1068.

Edwards, G., Clark, H., & Newton, P. (2001). The effects of elevated CO2 on seed production and seedling recruitment in a sheep-grazed pasture. *Oecologia*, 127(3), 383-394.

Edwards, W., Dunlop, M., & Rodgerson, L. (2006). The evolution of rewards: seed dispersal, seed size and elaiosome size. *Journal of Ecology*, 94(3), 687-694.

Fischer, C. & Türke, M. (2016). Seed preferences by rodents in the agri‐environment and implications for biological weed control. *Ecology and Evolution*, 6(16), 5796-5807.

Gómez, J.M., Puerta-Piñero, C., & Schupp, E.W. (2008). Effectiveness of rodents as local seed dispersers of Holm oaks. *Oecologia*, 155(3), 529-537.

Griffiths, H.M., Ashton, L.A., Walker, A.E., Hasan, F., Evans, T.A., Eggleton, P., & Parr, C.L. (2018). Ants are the major agents of resource removal from tropical rainforests. *Journal of Animal Ecology*, 87(1), 293-300.

Guitián, J., Fuentes, M., Bermejo, T., & López, B. (1992). Spatial variation in the interactions between Prunus mahaleb and frugivorous birds. *Oikos*, 63(1), 125-130.

Gurney, C.M., Prugh, L.R., & Brashares, J.S. (2015). Restoration of native plants is reduced by rodent-caused soil disturbance and seed removal. *Rangeland Ecology & Management*, 68(4), 359-366.

Hämäläinen, A., Broadley, K., Droghini, A., Haines, J.A., Lamb, C.T., Boutin, S., & Gilbert, S. (2017). The ecological significance of secondary seed dispersal by carnivores. *Ecosphere*, 8(2), e01685.

Handel, S.N. & Beattie, A.J. (1990). Seed dispersal by ants. *Scientific American*, 263(2), 76-83B.

Hedhly, A., Hormaza, J.I., & Herrero, M. (2009). Global warming and sexual plant reproduction. *Trends in Plant Science*, 14(1), 30-36.

Herrera, C.M. & Jordano, P. (1981). Prunus mahaleb and birds: the high‐efficiency seed dispersal system of a temperate fruiting tree. *Ecological Monographs*, 51(2), 203-218.

Hirsch, B.T., Kays, R., & Jansen, P.A. (2012). A telemetric thread tag for tracking seed dispersal by scatter-hoarding rodents. *Plant Ecology*, 213(6), 933-943.

Hodkinson, D.J. & Thompson, K. (1997). Plant dispersal: the role of man. *Journal of Applied Ecology*, 34(6), 1484-1496.

Hulme, P.E. (1998). Post-dispersal seed predation: consequences for plant demography and evolution. *Perspectives in Plant Ecology, Evolution and Systematics*, 1(1), 32-46.

Hulme, P.E. & Kollmann, J. (2005). Seed predator guilds, spatial variation in post-dispersal seed predation and potential effects on plant demography: a temperate perspective. In *Seed Fate: Predation, Dispersal and Seedling Establishment*, 9-30.

Hughes, L. & Westoby, M. (1990). Removal rates of seeds adapted for dispersal by ants. *Ecology*, 71(1), 138-148.

Hughes, L. & Westoby, M. (1992). Effect of diaspore characteristics on removal of seeds adapted for dispersal by ants. *Ecology*, 73(4), 1300-1312.

Janzen, D.H. (1970). Herbivores and the number of tree species in tropical forests. *The American Naturalist*, 104(940), 501-528.

Jensen, T.S. & Nielsen, O.F. (1986). Rodents as seed dispersers in a heath-oak wood succession. *Oecologia*, 70(2), 214-221.

Jongejans, E., Shea, K., Skarpaas, O., Kelly, D., & Ellner, S.P. (2011). Importance of individual and environmental variation for invasive species spread: a spatial integral projection model. *Ecology*, 92(1), 86-97.

Jongejans, E., Skarpaas, O., Ferrari, M.J., Long, E.S., Dauer, J.T., Schwarz, C.M., Rauschert, E.S., Jabbour, R., Mortensen, D.A., Isard, S.A., Lieb, D.A., Sezen, Z., Hulting, A.G., & Shea, K. (2015a). A unifying gravity framework for dispersal. *Theoretical Ecology*, 8(2), 207-223.

Jongejans, E., Silverman, E. J., Skarpaas, O., & Shea, K. (2015b). Post-dispersal seed removal of Carduus nutans and C. acanthoides by insects and small mammals. *Ecological Research*, 30(1), 173-180.

Jordano, P., García, C., Godoy, J.A., & García-Castaño, J.L. (2007). Differential contribution of frugivores to complex seed dispersal patterns. *Proceedings of the National Academy of Sciences*, 104(9), 3278-3282.

Keller, J.A. & Shea, K. (2021). Warming and shifting phenology accelerate an invasive plant life cycle. *Ecology*, 102(1), e03219.

Lemke, A., Von Der Lippe, M., & Kowarik, I. (2009). New opportunities for an old method: using fluorescent colours to measure seed dispersal. *Journal of Applied Ecology*, 46(5), 1122-1128.

Levey, D.J. & Sargent, S. (2000). A simple method for tracking vertebrate‐dispersed seeds. *Ecology*, 81(1), 267-274.

Linabury, M.C., Turley, N.E., & Brudvig, L.A. (2019). Insects remove more seeds than mammals in first‐year prairie restorations. *Restoration Ecology*, 27(6), 1300-1306.

LoPresti, E.F., Stessman, M.E., Warren, S.E., & Toll, K. (2022). Drying conditions alter the defensive function of seed mucilage against granivores. *Journal of Ecology,* 111(2), 525-536.

Mayer, V., Ölzant, S., & Fischer, R.C. (2005). Myrmecochorous seed dispersal in temperate regions. In *Seed Fate: Predation, Dispersal and Seedling Establishment*, 175-196.

Michael, P.J., Owen, M.J., & Powles, S.B. (2010). Herbicide-resistant weed seeds contaminate grain sown in the Western Australian grainbelt. *Weed Science*, 58(4), 466-472.

Mokany, K., Prasad, S., & Westcott, D.A. (2014). Loss of frugivore seed dispersal services under climate change. *Nature Communications*, 5(1), 1-7.

Molau, U. & Mølgaard, P. (1996). International Tundra Experiment Manual. Danish Polar Centre, Copenhagen.

Nathan, R. (2007). Total dispersal kernels and the evaluation of diversity and similarity in complex dispersal systems. In *Seed Dispersal: Theory and its Application in a Changing World*, 252-276.

O'Hanlon, J.C., Hill, S.J., & Andrew, N.R. (2019). Using devitalised seeds in myrmecological research. *Austral Entomology*, 58(4), 805-809.

Pemberton, R.W. & Irving, D.W. (1990). Elaiosomes on weed seeds and the potential for myrmecochory in naturalized plants. *Weed Science*, 38(6), 615-619.

Penn, H.J. & Crist, T.O. (2018). From dispersal to predation: A global synthesis of ant–seed interactions. *Ecology and Evolution*, 8(18), 9122-9138.

Pons, J. & Pausas, J.G. (2007). Acorn dispersal estimated by radio-tracking. *Oecologia*, 153(4), 903-911.

R Development Core Team (2022). R: A language and environment for statistical computing. R Foundation for Statistical Computing, Vienna, Austria. https://www.r-project.org/

Reiter, J., Curio, E., Tacud, B., Urbina, H., & Geronimo, F. (2006). Tracking Bat‐Dispersed Seeds Using Fluorescent Pigment. *Biotropica*, 38(1), 64-68.

Rogers, H.S., Beckman, N.G., Hartig, F., Johnson, J.S., Pufal, G., Shea, K., Zurell, D., Bullock, J.M., Cantrell, R.S., Loiselle, B., & Pejchar, L. (2019). The total dispersal kernel: a review and future directions. *AoB Plants*, 11(5), plz042.

Ruxton, G.D. & Schaefer, H.M. (2012). The conservation physiology of seed dispersal. Philosophical Transactions of the Royal Society B: *Biological Sciences*, 367(1596), 1708-1718.

Sales, L.P., Kissling, W.D., Galetti, M., Naimi, B., & Pires, M.M. (2021). Climate change reshapes the eco‐evolutionary dynamics of a Neotropical seed dispersal system. *Global Ecology and Biogeography*, 30(5), 1129-1138.

Sato, S., Kamiyama, M., Iwata, T., Makita, N., Furukawa, H., & Ikeda, H. (2006). Moderate increase of mean daily temperature adversely affects fruit set of Lycopersicon esculentum by disrupting specific physiological processes in male reproductive development. *Annals of Botany*, 97(5), 731-738.

Shimono, Y. & Konuma, A. (2008). Effects of human‐mediated processes on weed species composition in internationally traded grain commodities. *Weed Research*, 48(1), 10-18.

Singh, R.P., Prasad, P.V., & Reddy, K.R. (2013). Impacts of changing climate and climate variability on seed production and seed industry. *Advances in Agronomy*, 118, 49-110.

Skarpaas, O. & Shea, K. (2007). Dispersal patterns, dispersal mechanisms, and invasion wave speeds for invasive thistles. *The American Naturalist*, 170(3), 421-430.

Skinner, K., Smith, L., & Rice, P. (2000). Using noxious weed lists to prioritize targets for developing weed management strategies. *Weed Science*, 48(5), 640-644.

Taylor, K., Brummer, T., Taper, M.L., Wing, A., & Rew, L.J. (2012). Human‐mediated long‐distance dispersal: an empirical evaluation of seed dispersal by vehicles. *Diversity and Distributions*, 18(9), 942-951.

Teller, B.J., Zhang, R., & Shea, K. (2016). Seed release in a changing climate: initiation of movement increases spread of an invasive species under simulated climate warming. *Diversity and Distributions*, 22(6), 708-716.

Thomas, J.M.G., Boote, K.J., Allen, L.H., Gallo‐Meagher, M., & Davis, J.M. (2003). Elevated temperature and carbon dioxide effects on soybean seed composition and transcript abundance. *Crop Science*, 43(4), 1548-1557.

Thomson, D.M. (2007). Do source–sink dynamics promote the spread of an invasive grass into a novel habitat?. *Ecology*, 88(12), 3126-3134.

Thürig, B., Körner, C., & Stöcklin, J. (2003). Seed production and seed quality in a calcareous grassland in elevated CO2. *Global Change Biology*, 9(6), 873-884.

Travis, J.M., Delgado, M., Bocedi, G., Baguette, M., Bartoń, K., Bonte, D., Boulangeat, I., Hodgson, J.A., Kubisch, A., Penteriani, V., Saastamoinen, M., Stevens, V.M., & Bullock, J.M. (2013). Dispersal and species’ responses to climate change. *Oikos*, 122(11), 1532-1540.

Trumble, J.T., & Kok, L.T. (1982). Integrated pest management techniques in thistle suppression in pastures of North America. *Weed Research*, 22(6), 345-359.

Vander Wall, S.B. (2001). The evolutionary ecology of nut dispersal. *The Botanical Review*, 67(1), 74-117.

Vander Wall, S.B., & Longland, W.S. (2004). Diplochory: are two seed dispersers better than one?. *Trends in Ecology & Evolution*, 19(3), 155-161.

Vander Wall, S.B., Forget, P.M., Lambert, J.E., & Hulme, P.E. (2005a). Seed fate pathways: filling the gap between parent and offspring. In *Seed Fate: Predation, Dispersal and Seedling Establishment*, 1-8.

Vander Wall, S.B., Kuhn, K.M., & Beck, M.J. (2005b). Seed removal, seed predation, and secondary dispersal. *Ecology*, 86(3), 801-806.

Veldman, J.W. & Putz, F.E. (2010). Long‐distance dispersal of invasive grasses by logging vehicles in a tropical dry forest. *Biotropica*, 42(6), 697-703.

Vellend, M., Knight, T.M., & Drake, J.M. (2006). Antagonistic effects of seed dispersal and herbivory on plant migration. *Ecology Letters*, 9(3), 319-326.

Way, D.A., Ladeau, S.L., McCarthy, H.R., Clark, J.S., Oren, R.A.M., Finzi, A.C., & Jackson, R.B. (2010). Greater seed production in elevated CO2 is not accompanied by reduced seed quality in Pinus taeda L. *Global Change Biology*, 16(3), 1046-1056.

Weiss, F.E. (1908). The dispersal of fruits and seeds by ants. *New Phytologist*, 7, 23–28

Wichmann, M.C., Alexander, M.J., Soons, M.B., Galsworthy, S., Dunne, L., Gould, R., Fairfax, C., Niggemann, M., Hails, R.S., & Bullock, J.M. (2009). Human-mediated dispersal of seeds over long distances. *Proceedings of the Royal Society B: Biological Sciences*, 276(1656), 523-532.

Wilson, C.E., Castro, K.L., Thurston, G.B., & Sissons, A. (2016). Pathway risk analysis of weed seeds in imported grain: A Canadian perspective. *NeoBiota*, 30, 49-74.

Xiao, Z., Wang, Y., Harris, M., & Zhang, Z. (2006). Spatial and temporal variation of seed predation and removal of sympatric large-seeded species in relation to innate seed traits in a subtropical forest, Southwest China. *Forest Ecology and Management*, 222(1-3), 46-54.

Zhang, R., Jongejans, E., & Shea, K. (2011). Warming increases the spread of an invasive thistle. *PLoS* *One*, 6(6), e21725.

**Figure Captions**

**Table 1.** Estimates (1 standard error), -scores, and -values for the coefficients of the mixed-effects GLM with logit link fit to the proportion of seeds removed at 6, 12, and 24 hours. The intercept (baseline) represents seeds from unwarmed maternal plants and without elaiosomes. Estimates must be transformed using the inverse logit function to yield the proportion of seeds from the previous timestep that were removed.

|  |  |  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- | --- | --- |
|  | ***C. nutans* (*n* = 40)** | | |  | ***C. acanthoides* (*n* = 39)** | | |
|  |  |  |  |  |  |  |  |
| **6 Hours** | **Estimate** | ***z*** | ***p*** |  | **Estimate** | ***z*** | ***p*** |
| Intercept | -3.394 0.515 | -6.596 | <0.001 |  | -3.827 0.451 | -7.281 | <0.001 |
| Warming | 2.273 0.321 | 7.085 | <0.001 |  | 1.965 0.334 | 5.887 | <0.001 |
| Elaiosome | 2.374 0.321 | 7.398 | <0.001 |  | 1.775 0.336 | 5.290 | <0.001 |
| Warming:Elaiosome | -0.924 0.385 | -2.399 | 0.016 |  | -2.397 0.418 | -5.733 | <0.001 |
|  |  |  |  |  |  |  |  |
| **12 Hours** |  |  |  |  |  |  |  |
| Intercept | -1.662 0.426 | -3.904 | <0.001 |  | -3.295 0.552 | -5.974 | <0.001 |
| Warming | -0.158 0.275 | -0.574 | 0.566 |  | 3.263 0.403 | 8.090 | <0.001 |
| Elaiosome | 1.670 0.263 | 6.338 | <0.001 |  | 2.980 0.385 | 7.748 | <0.001 |
| Warming:Elaiosome | 0.584 0.401 | 1.455 | 0.146 |  | -3.374 0.456 | -7.402 | <0.001 |
|  |  |  |  |  |  |  |  |
| **24 Hours** |  |  |  |  |  |  |  |
| Intercept | 0.191 0.438 | 0.436 | 0.663 |  | 1.604 0.764 | 2.100 | 0.036 |
| Warming | 0.835 0.324 | 2.576 | 0.010 |  | 1.222 0.387 | 3.158 | 0.002 |
| Elaiosome | 0.962 0.329 | 2.922 | 0.003 |  | 1.428 0.398 | 3.584 | <0.001 |
| Warming:Elaiosome | 0.363 0.659 | 0.551 | 0.582 |  | -2.401 0.608 | -3.951 | <0.001 |

Diagram

Description automatically generated

**Figure 1**



**Figure 2**



**Figure 3**