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Effects of heat on the germination of sclerophyllous forest species in the highlands of Madagascar.

Swanni T. Alvarado a, b, , Elise Buisson a, Harison Rabarison b, Charlotte Rajeriarison b, Chris Birkinshaw c and Porter P. Lowry II d, e

Corresponding Author: Swanni T. Alvarado (e-mail: swanni_ta@yahoo.es )

a Institut Méditerranéen de Biodiversité et d’Ecologie (IMBE), Université d’Avignon et des Pays de Vaucluse, UMR CNRS IRD Aix Marseille Université, IUT, AGROPARC BP 61207, F-84911 Avignon cedex 09, France. Phone: +33 (0)4 90 84 38 58; Fax: +33 (0)4 90 84 38 07

b Département de Biologie et Ecologie Végétales, Faculté des Sciences, Université d’Antananarivo, Madagascar. BP 906 - Antananarivo 101, Madagascar. Phone: +261 32 02 236 77

c Missouri Botanical Garden, Madagascar Research and Conservation Program, BP 3391, d Missouri Botanical Garden, Africa and Madagascar Department, P.O. Box 299. St. Louis, MO 63166-0299. USA. Phone: + 1-314-577-9453; Fax: + 1-314-577-9596

d Missouri Botanical Garden, Africa and Madagascar Department, P.O. Box 299. St. Louis, MO 63166-0299. USA. Phone: +1-314-577-9453; Fax: +1-314-577-9596


Present address: Departamento de Botânica, Laboratório de Fenologia, Plant Phenology and Seed Dispersal Group, Instituto de Biociências, Universidade Estadual Paulista (UNESP), CP 199, Rio Claro, São Paulo, Brazil
Abstract
The effects of fire on germination have been extensively studied in many ecosystems. Several studies have shown that plant species in ecosystems frequently exposed to fire can survive through two main mechanisms: vegetative regeneration (re-sprouts) and recruitment of new individuals from a fire-resistant seed bank. In Africa, an increase in temperature can break seed dormancy and stimulate germination of some herbaceous and woody species. In Madagascar, the once widespread highland ecosystems dominated by woody species are now highly fragmented and dominated by anthropic grasslands and fields, with a significantly reduced area occupied by sclerophyllous forests referred to as "tapia woodlands". Six species of this endemic vegetation type were studied: *Abrahamia ibityensis* (Anacardiaceae), *Aphloia theiformis* (Aphloiacae), *Carissa edulis* (Apocynaceae), *Pentachlaena latifolia* (Sarcolenaecae), *Uapaca bojeri* (Phyllanthaceae) and *Vaccinium secundiflorum* (Ericaceae). Germination tests were conducted 1) by soaking seeds in water for 24 hours (imbibition) or 2) by exposing the seeds to dry heat. Four different temperatures (40°C, 60°C, 80°C and 120°C), where seeds were exposed for 10, 30, 60 and 90 minutes. To simulate hotter-faster fires, two higher temperatures (100°C and 120°C) were also evaluated by exposing seeds to dry heat for 5 minutes. The results did not reveal any significant effect of water 24-hour imbibition on germination. For most species germination decreased with increasing temperature of treatment using dry heat. *Uapaca bojeri* did not germinate under any treatment. Further studies on the biological and ecological characteristics of tapia woodland species in response to fire are needed to help guide conservation, management and restoration activities focusing on this vegetation type.

Key words: Tapia woodland, Ibity mountain, conservation, *Abrahamia ibityensis*, *Aphloia theiformis*, *Carissa edulis*, *Pentachlaena latifolia*, *Uapaca bojeri*, *Vaccinium secundiflorum*. 
1. Introduction

Fire is a natural disturbance element in many ecosystems, such as savannas and sclerophyllous woodlands in Africa (Kikula 1986; Campbell 1996; Kull 2000) and South America (Cerrado: Simon et al. 2009), and several Mediterranean-like ecosystems (Pausas et al. 1999, 2004; McCoy et al. 2002; Syphard et al. 2006; Schaffhauser et al. 2012). Fire determines the physiognomy and structure of vegetation and often influences species diversity (Bond and Keeley 2005). In fire-prone ecosystems, plant species can survive by using two main strategies: 1) resprouting, in which plants regenerate their above-ground biomass by growing from dormant buds on branches, trunks and underground; and 2) seedlings, which enable plants to recruit new individuals from a fire-resistant seed bank (Keeley and Zedler 1978; Pausas et al. 2004; Paula and Pausas 2008). Species are thus classified as ‘resprouters’ or ‘seeders’, according to the main mechanism involved in regeneration after fire, although both may be used. Other characteristics related to recruitment in fire-prone environments include post-fire seed dispersal and fire-induced flowering (Keeley and Fotheringham 1998).

The effects of fire on germination have been extensively studied in many ecosystems associated with this form of disturbance (Keeley 1987; Brown 1993; Mucunguzi and Oryem-Origa 1996; Gasshaw and Michelsen 2002; Syphard et al. 2006; Chou et al. 2012). A large number of studies have concluded that temperature increase and smoke from burning, or their interaction, can break seed dormancy (Brown 1993; Keeley and Fotheringham 1998; Dayamba et al. 2010) and can stimulate germination of herbaceous and woody species (Keeley 1987; Mucunguzi and Oryem-Origa 1996; Keeley and Fotheringham 1998; Chou et al. 2012). These phenomena have been described for several families (e.g. Fabaceae, Rhamnaceae, Convolvulaceae, Malvaceae, Cistaceae, and Sterculiaceae) that have physically dormant fire-tolerant seeds (Baskin and Baskin 2000). In some species, germination is favored only when soil heating is low and time of heat exposure is short (Mucunguzi and Oryem-Origa 1996; Gasshaw and Michelsen 2002). Other studies have shown that although fire increases the germination of some species, it may also inhibit germination, increase seed mortality and reduce seedling establishment (Moreno and Oechel 1991; Dayamba et al. 2008).

In the highlands of Madagascar, spatial and temporal mosaics of forests, savannas, woodlands and shrublands are the result of fire regimes that existed long before the arrival of humans ca. 2,000 years ago (Burney 1987; Dewar and Burney 1994). Currently, the highland ecosystems dominated by woody species are highly fragmented and dominated by anthropic grasslands and fields, with a significantly reduced area occupied by sclerophyllous forests referred to as "tapia woodlands" (Koechlin et al. 1974; Cornet and Guillaumet 1976), which have some similarities to the African "miombo" (Kull 2002). Tapia woodlands are dominated by the tapia tree Uapaca bojeri (Phyllanthaceae) associated with other woody species belonging to families endemic to Madagascar, such as Sarcoalaenaceae and Asteropeiaceae (Lowry II et al. 1997). The woody species present in tapia woodlands are fire tolerant: adult individuals typically have thick, spongy bark and weakly flammable leaves, and can re-sprout after fire (Campbell 1996).

Previous studies have, however, shown that fire can be a significant threat to this distinctive vegetation type (Kull 2002; Birkinshaw et al. 2006) because it causes changes in structure and composition (Alvarado et al. 2014a) and reduces natural regeneration by killing seedlings (Perrier de la Bâthie 1921; Gade 1996).

The biological characteristics (e.g. seed dispersal, germination, phenology, etc.) of the species present in tapia woodlands, most of which are endemic to Madagascar, are still poorly known. Fruit production and dispersal of tapia woodland species occur mainly during the rainy season,
between November and May (Alvarado et al. 2014b), a pattern that is also observed in some
other tropical ecosystems, such as the Cerrado (Batalha and Mantovani 2000; Weiser and
Godoy 2001) and the Atlantic forests (Morellato et al. 2000) of Brazil. Germination thus also
potentially occurs in the rainy season, when soil moisture is higher, which increases seed
hydration.

Understanding the biological and ecological characteristics of tapia woodland species in
response to fire is important for the conservation, management and restoration of this vegetation
type, which is currently only partly protected within the Ibity and Itremo New Protected Areas
and Isalo National Park. Restoration of degraded areas and the reinforcement of wild
populations of threatened and/or ecologically important plant species requires the use of seeds
and/or seedlings, and this must be based on sound knowledge of the processes involved and
how best to manage storage and germination. The study of environmental drivers like fire is
important to get an understanding of how this disturbance affects seed germination and in
consequence how fire has the potential to affect reproduction rates/success. This information is
important to better manage vegetation. In order to help address this need, an ex-situ study was
thus performed on the main woody species of tapia woodlands to determine the effects of
imbibition and fire on seed germination. Our goals were to determine whether 1) precipitation
and imbibition alone stimulate their germination; and 2) if the seeds of these species can
survive after being affected by different scenarios of fire, and thus whether fire could represent
a barrier to natural regeneration in this vegetation type.

2. Methods

2.1. Seed material

Six species were studied: Abrahamia ibityensis (H. Perrier) Randrian. & Lowry, ined.
(Anacardiaceae), Aphloia theiformis (Vahl) Benn. (Aphloiaceae), Carissa edulis (Forssk.) Vahl
(Apocynaceae), Pentachlaena latifolia H. Perrier (Sarcolaenaceae), Uapaca bojeri Baill.
(Phyllanthaceae) and Vaccinium secundiflorum Hook. (Ericaceae). The choice of species was
defined based on two considerations: 1) the selected species were the dominant woody elements in the
canopy and mid-storey (trees and shrubs with a DBH > 1cm): U. bojeri, represents 82% of the
trees counted on Ibity (Alvarado et al. 2014a) and the other species are common in the lower
canopy and mid-storey; and 2) these species are endemics to Madagascar and are potentially
threatened by changes in fire regimes (Alvarado et al. 2014b). It is thus important to study their
biology to increase knowledge within the framework of future conservation, restoration or
reinforcement programs. Seeds used in this study were collected on the Ibity Massif, a New
Protected Area, located in the Antananarivo province, in Madagascar’s Central Highlands,
200km south of the capital city, Antananarivo, and 25km south of the town Antsirabe (47°07’S,
20°01’E). The area’s climate, classified as Cwb according to the Köppen system (Peel et al.
2007), is characterized by a well-defined dry season (June-October) and wet season (November
to May). Average annual rainfall reaches 1583mm and average temperature is 17.5°C, with a
mean maximum temperature of 27.0 °C and a mean minimum temperature of 9.1 °C in the wet
season and a mean maximum temperature of 27.3 °C and a mean minimum temperature of
5.5°C in the dry season (based on data from 1961 to 1990; Meteorology Service of
Ampandrianomby). Seeds were harvested randomly from 5-10 mother-plants of each species
during the wet season (November-December) in 2010 and 2011, according to their availability;
seeds were stored in a dry place at the laboratory; germination trials were run during the
following 1-2 months.

2.2. Germination trials
Germination experiments, conducted using a series of treatments (Table 1), were performed at the Laboratory of Plant Physiology at the University of Antananarivo. One set of trials was done without pre-germination treatment on a set of “control” seeds, and a second set was performed to determine water requirements for germination (imbibitions) by soaking seeds in water for 24 hours prior to the germination tests. Using a third set of seeds, the effect of pre-germination dry heat was tested on four of the six species studied (Aphloia theiformis, Carissa edulis, Uapaca bojeri and Vaccinium secundiflorum) to simulate increased temperature produced by fires. Four different temperatures (40°C, 60°C, 80°C and 120°C) were used, following the protocol of Munyanziza and Msanga (1996). For the other two species (Abrahamia ibityensis and Pentachlaena latifolia), we did not have enough seeds to run the dry heat tests. Almost 50% of seeds of these two species germinated in the paper bags before to start the trial. Seeds were heated in a mufla oven with an external thermometer to verify the required temperatures; another thermometer was placed inside the oven to double-check the temperature. Seeds were introduced into the mufla after five minutes of constant temperature.

For each temperature, seeds were separated in paper bags and exposed in Petri dish for 10, 30, 60 and 90 minutes in an oven. To simulate hotter-faster fires, the effects of two higher temperatures (100°C and 120°C) were also evaluated by exposing seeds to dry heat for 5 minutes. Seeds of all species were heated together for a same treatment, but replicates were heated separately in order to avoid pseudoreplication in the experimental design (Morrison and Morris 2000). Treatments were designed to cover the range of conditions that seeds could potentially encounter on the ground or in open sites during wild fires (Keeley 1987).

After treatment, seeds were sown on 90 × 15mm Petri dishes covered with Whatman 90mm diameter filter paper, and placed in a growth chamber at a constant temperature of 25°C and a 12 hour light regime, following the protocol of Baskin & Baskin (2000). The numbers of replicates (=petri dishes) and of seeds used for each species depended on their availability and on seed size (15, 25, 30 or 100 seeds per dish, Table 2).

The seeds were monitored every 2 days for 8 weeks. Germination was recorded when a 2 mm seed radicle had emerged. The effect of each treatment on seed germination was measured as the percentage of germination (PG) and mean germination time (MGT) (Munyanziza and Msanga 1996; Chou et al. 2012), calculated using the following formulas:

\[ PG(\%) = \frac{\sum n_i}{N} \times 100 \]

and

\[ MGT = \frac{\sum (t_i \times n_i)}{\sum n_i} \]

where \( n_i \) is the number of germinated seeds, \( N \) is the total number of seeds, and \( t_i \) is the day when \( n_i \) seeds had germinated.

### 2.3. Statistical analyses

The effects of treatments on germination were examined using generalized linear models with a binomial error structure (Zuur et al. 2009). Separate analyses were performed for each species. The response variable was the percent of germinated seeds after 8 weeks. First, the inhibition treatment and the control were used as factors and compared with one another. Then dry heat treatments were compared with the control. We used Akaike’s Information Criterion to select the best model(s) (Akaike 1973; Burnham and Anderson 2002).

To assess the difference in MGT between the pre-germination imbibition treatment and the control, t-tests were performed for each species. Two-way ANOVAs were conducted to determine the effects of temperature (40°C, 60°C, 80°C, 100°C and 120°C) and exposure time (5, 10, 30, 60 and 90 minutes) on MGT. For all ANOVA, MGT values were treated as variables.
and temperature and exposure time as factors. The normal distribution of data values was verified with the Shapiro-Wilk test (<40 data) or the Lilliefors test (>40 data) and square root transformations were performed as necessary. The homogeneity of variances was confirmed with Bartlett’s tests. When two-way ANOVA were significant (P<0.05), post-hoc Tukey tests were performed. When the normal distribution of data was not confirmed, Kruskal-Wallis tests, followed by Wilcoxon tests, adjusted with the Bonferroni correction, were performed. All tests were performed using R software (The R Foundation for Statistical Computing, version 2.15.1, (R Development Core Team 2012).

3. Results
3.1. Germination
The germination treatment by imbibition had no significant effect on any of the five species studied (Table 3). Germination percentages were similar between the control and the imbibition treatment: they are relatively high for Abrahamia ibityensis (77.9±7.5%) and Carissa edulis (97.6±1.1%), intermediate for Aphloia theiformis (42.0±1.8%) and Pentachlaena latifolia (43.9±3.74%), and comparatively low for Vaccinium secundiflorum (26.2±3.47%). No germination was observed for Uapaca bojeri.

Only one species, Carissa edulis, had a significant difference in MGT between the two treatments: the imbibition treatment lowered MGT by one day (Table 3). Pentachlaena latifolia germinated in 3.98±0.26 days (Table 3), whereas Vaccinium secundiflorum germinated in 19.7±0.63 days (Table 3). MGT is intermediate for the other three species: Abrahamia ibityensis (5.9±0.71 days), Aphloia theiformis (11.3±0.37 days) and Carissa edulis (6.9±0.20 days).

3.2. Effect of dry heat on seed germination
Temperature had a significant effect on PG and MGT for all tested species except Uapaca bojeri which did not germinate under any conditions. PG for Aphloia theiformis was significantly different between treatments (P(CHi) < 0.001; Figure 1a). For this species, maximum PG was obtained from control seeds (43.3±2.8%) which was not significantly different from seeds heated at 40°C or 80°C; the PG was lower for seeds heated at 120°C for 10 minutes (6.7±1.1%) and significantly the lowest for seeds heated at 120°C for 30, 60, 90 minutes (0±0%, Table 4, Figure 1a). Heating increased MGT, which was maximum when seeds were treated at 120°C (27.7±1.6 days; F(Temperature)=49.9, P<0.001, Table 4). Carissa edulis PG were significantly different between treatments (P(CHi) < 0.001, Figure 1b). For this species, maximum PG was obtained from control seeds and those heated at 40°C for 30 minutes (96±1.95% and 97.54±1.6%, respectively, Table 4, Figure 1b) and was not significantly different from seeds heated at 40°C for 10, 60 and 90 minutes or 60°C for 10, 30, 60 minutes; PG was intermediate for seeds heated at 60°C for 90 minutes (73.6±4.1%) or 100 and 120°C for 5 minutes (81.6 ± 3.9 and 75.2 ± 4.08 respectively) and significantly the lowest for seeds heated at 80 and 120°C for more than 10 minutes (4.8±1.96%, Table 4, Figure 1b); the MGT varied from 5.7 days to 7.9 days, depending on temperature (Kruskal-Wallis chi²(Temperature)=38.23, P < 0.001, Table 4). Finally, for Vaccinium secundiflorum, PG was significantly different between treatments (P(CHi) < 0.001, Figure 1c). The maximum PG for V. secundiflorum was obtained from control seeds (26.2±3.5%, Table 4, Figure 1c) and was not significantly different from seeds heated at 40°C for 10 or 90 minutes, at 80°C and at 120°C for 10 and 30 minutes; the PG was significantly the lowest for seeds heated at 60°C for 60 and 90 minutes, at 100°C for 5 minutes and at 120°C for 60 and 90 minutes (Table 4, Figure 1c). At 80°C, PG was higher (between 17.2±0.23 and 20.2±0.79%) than at 60°C but did not reach the level obtained in the control. The MGT for this species varied depending on the temperature at which the seeds were heated and the time of exposure to heat (F(Temperature×Time)= 5.52, P <
0.001, Table 4). The highest value of MGT was observed when seeds were heated at 120°C for 5 minutes and at 60°C for 60 minutes (34.5±1.2 and 32.93±1.6 days, respectively), and the lowest when they were preheated at 120°C for 120 minutes (2.7±2.7 days respectively).

### 4. Discussion

#### 4.1. Germination

Two of the six species studied (Carissa edulis and Abrahania ibityensis) exhibited percentages of germination (PG) over 84% under the control treatment, indicating that their seeds are mainly not dormant. The other four species examined had PG values under 45.5%. The water imbibition treatment did not significantly improve germination success in the species tested, but only resulted in a slight decrease in the mean germination time (MGT) of one species, C. edulis. The results show that none of the tapia woodland species considered in our study increase the percent of germination by imbibition alone. However, more germination tests including other combinations of factors (e.g. different incubation temperatures, ripening or stratification prior to germination, etc.) are needed to increase knowledge on seed physiology and ecology, and to confirm whether there is any kind of dormancy. In tropical dry woodlands and savannas, establishment from seeds has been reported as a rare event (Sarmiento and Monasterio 1983) and the seed ecology of germination for woody species is variable: some species exhibit seed dormancy (Brown 1993; Mucunguzi and Oryem-Origa 1996; Dayamba et al. 2008), some have non-dormant seeds, and others are recalcitrant (Oliveira and Valio 1992).

A phenology study performed on selected woody tapia woodland species (Alvarado et al. 2014b) shows that seed dispersal takes place during the rainy season for five of the species studied: Abrahania ibityensis, Aphloia theiformis, C. edulis, Uapaca bojeri and Vaccinium secundiflorum. Seed germination of these and other woody species is potentially linked to rainfall because precipitation brings an increase in soil moisture, which promotes germination (Oliveira and Silva 1993; Baskin and Baskin 2000). This finding makes it difficult to interpret the results of the imbibition tests; we expected germination to be improved by water imbibition, as shown by Choinski and Tuohy (1991) for two African trees, Acacia tortilis (Forssk.) Hayne and Acacia karroo Hayne.

Concerning the dominant species of tapia woodlands, Uapaca bojeri, we did not observe any germination in our laboratory tests, regardless of the treatment. Kull et al. (2005) reported that U. bojeri seeds are recalcitrant, losing their viability within a few months after dispersal, a finding that was confirmed by Randrianavosoa et al. (2011). They also found that the percent of germination of U. bojeri in laboratory experiments was between 45% and 95%, when seed water content was 20-30%, regardless of whether lowered water content was obtained after 10-15 days of drying in the shade or 26 days storage at a relative humidity of 60-84%. On-site experiments in local soil, at the Ibity nursery, showed 17.8% germination after three months, with germination beginning six weeks after seeds were sowed (Alvarado et al. 2010). This could explain the absence of germination U. bojeri after eight weeks of observation in laboratory conditions. Moreover, since the viability of the seeds was not tested, it is not possible to conclude whether this result is due to the fact that the seeds were dormant or dead.

With the exception of Abrahania ibityensis and Carissa edulis, we found that germination of the studied species was less than 50%. In A. ibityensis and C. edulis, PG values in the field nursery were lower (A. ibityensis 35.7% and C. edulis 22.4%, Alvarado et al. 2010) than in the laboratory (A. ibityensis 84.3% and C. edulis 96%). In Aphloia theiformis, PG in the nursery (26.9%; Alvarado et al. 2010) was just over half that in the laboratory (42.0%).
4.2. Effect of dry heat on seed germination

High temperatures failed to induce germination in the three species studied, suggesting that they have a germination strategy different from that of typical species found in most fire-adapted ecosystems, where fire at low intensities stimulates germination and at high intensities increases seed mortality (Keeley 1987; Moreno and Oechel 1991; Mucunguzi and Oryem-Origa 1996). Although the seeds of Aphloia theiformis, Carissa edulis and Vaccinium secundiflorum can tolerate moderate temperatures (60°C – 80°C depending of the species), they are not resistant to fire, as a reduction of PG was observed after both short or long exposure to high temperatures. Stimulation of germination by fire is common among species of Fabaceae (Portlock et al. 1990; Tarrega et al. 1992; Bradstock and Auld 1995; Mucunguzi and Oryem-Origa 1996; Auld and Denham 2006), Combretaceae (Dayamba et al. 2010) and Ericaceae, which do not have the ability to re-sprout after fire, and are thus totally dependent on the seed bank for reestablishment after burning (Keeley 1987). These species exhibit physical or physiological dormancy and germination is frequently stimulated by smoke or a combination of heat and smoke (Auld and Ooi 2008). However, we found that germination of the ericaceous species V. secundiflorum was not stimulated by increased temperature, a result contrary to the findings reported by Gonzalez-Rabanal & Casal (1995) for Mediterranean species, who noted that germination in the Ericaceae they studied was stimulated by heat shock, however, a wide range of responses can be detected in members of this family. Even if V. secundiflorum germination does not increase with high temperatures as expected, it is relevant to note that this species is at least fire tolerant. For the other two species examined in our study, exposure to moderate heat (<60°C for A. theiformis and <80°C for C. edulis), even for long periods, did not affect germination, whereas at temperatures >100°C, their seeds were sensitive to the time of exposure, which is similar to the response of certain woody species in Australia and Burkina Faso (Auld and O’Connell 1991; Zida et al. 2005). Fire temperature has been measured in several tropical ecosystems, ranging from 47°C (50 cm above the ground surface) to 537.5°C (at the surface) in Brazilian savannas (Fidelis et al. 2010), from 98-458°C in Australian savannas (Morgan 1999), and up to 150°C in Australian woodlands (Bradstock and Auld 1995). While some studies suggest that temperatures above 60°C as being fatal to plant tissues (Whelan 1995), but exposure to dry heat at 60°C and 100°C renders seeds of some species more permeable, thus promoting germination (Baskin and Baskin 2000).

Although tapia woodlands are a vegetation formation in which fire is a natural disturbance (Kull 2000, 2002), germination was not stimulated by dry heat in any of the species we studied. They showed varying levels of tolerance to different temperatures and duration of exposure to heat. Carissa edulis was the most sensitive, surviving in temperatures up to 60°C, regardless of the time of exposure to dry heat without reducing seed viability, but suffering significant reduction in PG (to less than 20%) at 80°C, although exposure for 5 minutes to 120°C did not kill seeds and only slightly reduced PG. Aphloia theiformis was less sensitive to dry heat. At 40°C, PG remained close to that observed in the control, but exposure to 40°C for 90 minutes and to 80°C, regardless of exposure time, PG decreased from 10-30%, and at 120°C all seeds died. In this species, higher temperatures reduced germination and increased MGT. Vaccinium secundiflorum had the most variable behavior faced with dry heat of the three species studied: high temperatures (80°C and 120°C) had a less negative impact on germination than low temperatures (40°C and 60°C).

5. Conclusion

Seed germination varied among the woody species we studied, but germination was low in untreated samples for 4 species out of 6 (exceptions: Abrahamia ibityensis and Carissa edulis) and decreased with increasingly hot shock treatment using dry heat for most species. Contrary
to our expectation, water imbibition did not stimulate seed germination. The reduced percentage of germination with increasing temperature suggests that high temperatures reduced viability. According with the literature and as suggested by our results, at least one of the species (*Uapaca bojeri*) has recalcitrant seeds and it is quite possible that some of the other species have similar (or at least intermediate) seeds. However, in order to conclude if seeds are dormant or recalcitrant, viability tests should be carry out to confirm if the lack of germination in the controls is due to seeds being dormant or dead. On Ibity Mountain, where the current fire regime has been modified by human activities, fires had a variable impact on seed germination in these species. Our results provide some useful insights, but further studies of the seed ecology (e.g. the role of smoke, the possible role of seed scarifications, etc.) of these and other species occurring in tapia woodlands are required in order to develop a better understanding of the relationship between fire and germination, and of the impacts of the current human-caused fire regime found on Ibity Mountain. Insights from additional studies would also be valuable for developing improved conservation and management strategies for the native species of tapia woodlands, especially with regard to efforts aimed at reintroducing species to areas where they have been extirpated and reinforcing populations that have been impacted in this localized and threatened ecosystem.

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References


Table 1: Summary of treatments applied to the seeds of six tapia woodland species (*Abrahamia ibityensis*, *Aphloia theiformis*, *Carissa edulis*, *Pentachlaena latifolia* *Uapaca bojeri* and *Vaccinium secundiflorum*). For additional information on the seeds, see some photos on Electronic appendix 1.

<table>
<thead>
<tr>
<th>Treatment</th>
<th>Treatment details</th>
<th>Species tested</th>
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<td>Control</td>
<td>No pre-germination treatment</td>
<td>All species</td>
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<td>Imbibition</td>
<td>Seeds soaked in water for 24 hours prior to germination</td>
<td>All species except <em>V. secundiflorum</em></td>
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<td>Dry heat</td>
<td>40°C 10, 30, 60 or 90 minutes</td>
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<td></td>
<td>60°C 10, 30, 60 or 90 minutes</td>
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<tr>
<td></td>
<td>80°C 10, 30, 60 or 90 minutes</td>
<td>All species except <em>A. ibityensis</em> and <em>P. latifolia</em></td>
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<tr>
<td></td>
<td>120°C 10, 30, 60 or 90 minutes</td>
<td>All species except <em>A. ibityensis</em> and <em>P. latifolia</em></td>
</tr>
<tr>
<td>Fast burnt</td>
<td>100°C 5 minutes</td>
<td><em>C. edulis</em>, and <em>V. secundiflorum</em></td>
</tr>
<tr>
<td></td>
<td>120°C 5 minutes</td>
<td><em>C. edulis</em>, and <em>V. secundiflorum</em></td>
</tr>
</tbody>
</table>
Table 2: Studied species and details of the experimental setup. Number of replicates, seeds/dish, total number of treatments and total number of seeds are indicated for each species.

<table>
<thead>
<tr>
<th>Species</th>
<th>Family</th>
<th>Life-form/habit</th>
<th>Geographic distribution</th>
<th>Replicates (Petri dishes)</th>
<th>Seeds / dish</th>
<th>Total treatments</th>
<th>Total seeds</th>
</tr>
</thead>
<tbody>
<tr>
<td>Abrahamia ibityensis</td>
<td>Anacardiaceae</td>
<td>Shrub</td>
<td>Endemic to Madagascar (Ibity and Itremo massifs only)</td>
<td>7</td>
<td>10</td>
<td>2</td>
<td>140</td>
</tr>
<tr>
<td>Aphloia theiformis</td>
<td>Aphloiacae</td>
<td>Shrub or tree</td>
<td>Madagascar, Comoros, Mascarenes, Seychelles, Africa</td>
<td>5</td>
<td>30</td>
<td>14</td>
<td>2100</td>
</tr>
<tr>
<td>Carissa edulis</td>
<td>Apocynaceae</td>
<td>Shrub</td>
<td>Madagascar, Africa, Indian Ocean islands, Asia (India)</td>
<td>5</td>
<td>25</td>
<td>20</td>
<td>2500</td>
</tr>
<tr>
<td>Pentachlaena latifolia</td>
<td>Sarcolaeaceae</td>
<td>Shrub or tree</td>
<td>Endemic to Madagascar (Ibity only)</td>
<td>4</td>
<td>15</td>
<td>2</td>
<td>120</td>
</tr>
<tr>
<td>Uapaca bojeri</td>
<td>Phyllantheae</td>
<td>Tree</td>
<td>Endemic to Madagascar (widespread)</td>
<td>6</td>
<td>10</td>
<td>20</td>
<td>1200</td>
</tr>
<tr>
<td>Vaccinium secundiflorum</td>
<td>Ericaceae</td>
<td>Shrub</td>
<td>Endemic to Madagascar (widespread)</td>
<td>3</td>
<td>100</td>
<td>19</td>
<td>5700</td>
</tr>
</tbody>
</table>
Table 3: Average percent of germination (PG) and mean germination time (MGT) of six woody species from tapia woodland (Madagascar) tested under laboratory conditions for water imbibition before the start of germination tests. Germination was monitored for 8 weeks in a culture chamber at a constant temperature of 25°C and a 12 hours light regime. No water imbibition tests were carried out for *Vaccinium secundiflorum* (NT = not tested).

<table>
<thead>
<tr>
<th>Species</th>
<th>PG (%)</th>
<th>MGT (days)</th>
<th>Significance (PG)</th>
<th>Significance (MGT)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Mean ± se</td>
<td>Mean ± se</td>
<td>P(&gt;Chi)</td>
<td>t</td>
</tr>
<tr>
<td><strong>Abrahamia ibityensis</strong></td>
<td>71.4±14.4</td>
<td>84.3±4.8</td>
<td>0.07</td>
<td>7.0±1.21</td>
</tr>
<tr>
<td><strong>Pentachlaena latifolia</strong></td>
<td>42.3±4.7</td>
<td>45.5±6.8</td>
<td>0.71</td>
<td>3.8±0.52</td>
</tr>
<tr>
<td><strong>Aphloia theiformis</strong></td>
<td>40.7±2.4</td>
<td>43.3±2.8</td>
<td>0.64</td>
<td>10.8±0.56</td>
</tr>
<tr>
<td><strong>Carissa edulis</strong></td>
<td>99.2±0.8</td>
<td>96±1.95</td>
<td>0.08</td>
<td>6.3±0.13</td>
</tr>
<tr>
<td><strong>Uapaca bojeri</strong></td>
<td>0</td>
<td>0</td>
<td>-</td>
<td>0</td>
</tr>
<tr>
<td><strong>Vaccinium secundiflorum</strong></td>
<td>NT</td>
<td>26.2±3.5</td>
<td>-</td>
<td>NT</td>
</tr>
<tr>
<td>Temperature</td>
<td>Time</td>
<td>Aphloia theiformis</td>
<td>Carissa edulis</td>
<td>Vaccinium secundiflorum</td>
</tr>
<tr>
<td>-------------</td>
<td>------</td>
<td>-------------------</td>
<td>----------------</td>
<td>------------------------</td>
</tr>
<tr>
<td></td>
<td></td>
<td>PG Mean ± SE</td>
<td>MGT Mean ± SE</td>
<td>PG Mean ± SE</td>
</tr>
<tr>
<td>Control</td>
<td></td>
<td>43.3 ± 2.8</td>
<td>11.8 ± 0.4</td>
<td>96 ± 1.95</td>
</tr>
<tr>
<td>40</td>
<td>10</td>
<td>38.7 ± 2.7</td>
<td>11.6 ± 0.9</td>
<td>95.3 ± 1.9</td>
</tr>
<tr>
<td>40</td>
<td>30</td>
<td>39.9 ± 2.1</td>
<td>13.9 ± 1.0</td>
<td>97.5 ± 1.6</td>
</tr>
<tr>
<td>40</td>
<td>60</td>
<td>43.3 ± 3.0</td>
<td>13.3 ± 0.9</td>
<td>94.3 ± 1.0</td>
</tr>
<tr>
<td>40</td>
<td>90</td>
<td>28.7 ± 3.1</td>
<td>13.7 ± 0.7</td>
<td>91.2 ± 3.0</td>
</tr>
<tr>
<td>60</td>
<td>10</td>
<td>NT</td>
<td>NT</td>
<td>91.2 ± 4.6</td>
</tr>
<tr>
<td>60</td>
<td>30</td>
<td>NT</td>
<td>NT</td>
<td>88.8 ± 3.9</td>
</tr>
<tr>
<td>60</td>
<td>60</td>
<td>NT</td>
<td>NT</td>
<td>83.2 ± 5.0</td>
</tr>
<tr>
<td>60</td>
<td>90</td>
<td>NT</td>
<td>NT</td>
<td>73.6 ± 4.1</td>
</tr>
<tr>
<td>80</td>
<td>10</td>
<td>26.7 ± 2.4</td>
<td>14.8 ± 0.8</td>
<td>6.4 ± 2.0</td>
</tr>
<tr>
<td>80</td>
<td>30</td>
<td>17.3 ± 3.2</td>
<td>15.3 ± 0.9</td>
<td>8.8 ± 1.5</td>
</tr>
<tr>
<td>80</td>
<td>60</td>
<td>14.0 ± 3.1</td>
<td>16.6 ± 0.6</td>
<td>10.4 ± 2.7</td>
</tr>
<tr>
<td>80</td>
<td>90</td>
<td>17.3 ± 4.3</td>
<td>17.6 ± 2.2</td>
<td>8.8 ± 1.5</td>
</tr>
<tr>
<td>120</td>
<td>10</td>
<td>6.7 ± 1.1</td>
<td>27.7 ± 1.6</td>
<td>4.8 ± 2.0</td>
</tr>
<tr>
<td>120</td>
<td>30</td>
<td>0 ± 0</td>
<td>NC</td>
<td>17.6 ± 3.5</td>
</tr>
<tr>
<td>120</td>
<td>60</td>
<td>0 ± 0</td>
<td>NC</td>
<td>20.8 ± 3.2</td>
</tr>
<tr>
<td>120</td>
<td>90</td>
<td>0 ± 0</td>
<td>NC</td>
<td>21.6 ± 3.7</td>
</tr>
<tr>
<td>100</td>
<td>5</td>
<td>NT</td>
<td>NT</td>
<td>81.6 ± 3.9</td>
</tr>
<tr>
<td>120</td>
<td>5</td>
<td>NT</td>
<td>NT</td>
<td>75.2 ± 4.08</td>
</tr>
</tbody>
</table>

Table 4: Average percent of germination (PG) and mean germination time (MGT) of three woody species from tapia woodland (Madagascar) tested under laboratory conditions for control and dry heat treatments. Germination was monitored for 8 weeks in a culture chamber at a constant temperature of 25°C and a 12 hours light regime. No germination was observed in *Uapaca bojeri* (NT = not tested). NC = not calculated. GLM results for PG were significant for the interaction Temperature x Time for all species.

<table>
<thead>
<tr>
<th>Statistic result</th>
<th>P(&gt;Chi) Temperature x Time &lt; 0.001</th>
<th>F&lt;sub&gt;Temperature&lt;/sub&gt;=49.9, P&lt;sub&gt;Time&lt;/sub&gt;&lt;0.001</th>
<th>P(&gt;Chi) Temperature x Time &lt; 0.001</th>
<th>Kruskal-Wallis chi² Temperature = 38.23, p-value&lt;0.001</th>
<th>P(&gt;Chi) Temperature x Time &lt; 0.001</th>
<th>F&lt;sub&gt;Temperature x Time&lt;/sub&gt; = 5.52, p-value&lt;0.001</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Statistic result</td>
<td></td>
<td>Statistic result</td>
<td></td>
<td>Statistic result</td>
<td></td>
</tr>
</tbody>
</table>
Figure 1: Percent of germination of the three (out of five) species studied using various treatments to test the effect of dry heat pre-germination. Four temperatures (40°C, 60°C, 80°C and 120°C) were evaluated; for each temperature, seeds were exposed to heat for 10, 30, 60 and 90 minutes. To simulate a hot, fast-burning fire, seeds were exposed to dry heat at two temperatures (100°C and 120°C) for 5 minutes. a) Aphloia theiformis, b) Carissa edulis, c) Vaccinium secundiflorum. White bars (control), light grey bars (seeds heated at 40°C), grey bars (seeds heated at 60°C), dark grey bars (seeds heated at 80°C),
black bars (seeds heated at 120°C), barred bars (seeds heated at 100°C for 5 minutes). Post-hoc results are showed by letters on the bar (a, b, c, etc.).