



Climate and soil factors influencing seedling recruitment of plant species used for dryland restoration

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Abstract. Land degradation affects 10–20 % of drylands globally. Intensive land use and management, large-scale disturbances such as extractive operations, and global climate change, have contributed to degradation of these systems worldwide. Restoring these damaged environments is critical to improving ecosystem services and functions, conserve biodiversity, and contribute to climate resilience, food security, and landscape sustainability. Here, we present a case study on plant species of the mining intensive semi-arid Pilbara region in Western Australia that examines the effects of climate and soil factors on the restoration of drylands. We analysed the effects of a range of rainfall and temperature scenarios and the use of alternative soil materials on seedling recruitment of key native plant species from this area. Experimental studies were conducted in controlled environment facilities where conditions simulated those found in the Pilbara. Soil from topsoil (T) stockpiles and waste materials (W) from an active mine site were mixed at different proportions (100 % T, 100 % W, and two mixes of topsoil and waste at 50 : 50 and 25 : 75 ratios) and used as growth media. Our results showed that seedling recruitment was highly dependent on soil moisture and emergence was generally higher in the topsoil, which had the highest available water content. In general, responses to the climate scenarios differed significantly among the native species which suggest that future climate scenarios of increasing drought might affect not only seedling recruitment but also diversity and structure of native plant communities. The use of waste materials from mining operations as growth media could be an alternative to the limited topsoil. However, in the early stages of plant establishment successful seedling recruitment can be challenging in the absence of water. These limitations could be overcome by using soil amendments but the cost associated to these solutions at large landscape scales needs to be assessed and proven to be economically feasible.

1 Introduction

Land degradation affects nearly 2 billion hectares of land globally, with 25 % of the total global land considered degraded to some extent (Barbero-Sierra et al., 2015; Bisaro et al., 2014; Brevik et al., 2015; Miao et al., 2015; Stan-turf et al., 2015; Torres et al., 2015; Wang et al., 2015). Restoring these damaged environments is critical to improving ecosystem services and functions, conserve biodiversity, and contribute to climate resilience, food security and landscape sustainability at the local, regional and global level

(de Moraes Sá et al., 2015; Minnemeyer et al., 2011; Perring et al., 2015; Prosdocimi et al., 2016; Roa-Fuentes et al., 2015; Zucca et al., 2015). Drylands, which include semi-arid and arid environments, are particularly vulnerable to land degradation with estimates suggesting 10–20 % of these ecosystems are degraded globally (Millennium Ecosystem Assessment, 2005; Safriel et al., 2005), and continue to be degraded across millions of hectares every year (Brauch and Spring, 2009; Wang et al., 2015; Yan and Cai, 2015). Intensive land use and management, large-scale disturbances such as ex-

tractive operations (e.g. mining), and global climate change have contributed to degradation of these systems worldwide (Anaya-Romero et al., 2011; Keesstra et al., 2016a; Kildisheva et al., 2016; Muñoz-Rojas et al., 2015).

When attempting to restore degraded, arid ecosystems challenges include limited rainfall, high temperatures, and soils with low nutrient levels and water holding capacity (Anaya-Romero et al., 2015; Muñoz Rojas et al., 2016a). Thus, despite the efforts and investments to restore these systems worldwide (Keesstra et al., 2016a), restoration of drylands has low rates of success (James et al., 2013; Sheley et al., 2011). To improve our capacity to reinstate biodiverse, viable plant communities, there is a strong need to advance our understanding of how these systems function and the effects that environmental and edaphic factors have on processes such as seedling emergence and plant growth and survival (Perring et al., 2015). For example, changes in soil water availability as a consequence of reduced rainfall and evaporation, or increases in temperature due to global warming, may affect restoration outcomes through influencing seedling recruitment (Cochrane et al., 2015; Lloret et al., 2004) or the composition and distribution of plant species (Lai et al., 2015). But the impact of environmental factors on restoration can be also compounded by unfavourable edaphic conditions (Audet et al., 2013; Thomas et al., 2015). Thus, improving soil physical and chemical properties can be decisive for successful revegetation (Machado et al., 2013), which is important in extractive industries operating in dryland environments.

During open-cut and strip mining operations, the top layer of soil is commonly removed and stockpiled before starting the extraction process and then respread before seeding the target sites for restoration (Lamb et al., 2015; Rivera et al., 2014). This topsoil is an important source of seeds, nutrients and microorganisms (Erickson et al., 2016a; Golos and Dixon, 2014; Koch, 2007; Muñoz-Rojas et al., 2016b) but its use in restoration is often limited by its scarcity and the detrimental conditions that topsoil stockpiling can have on soil functionality (Keipert et al., 2002). Waste materials produced in mining operations provide alternative substrates that are currently being used as growth media in restoration (Machado et al., 2013; Muñoz-Rojas et al., 2016b; Thomas et al., 2015). These substrates can integrate coarser materials that help to reduce slope instability and prevent erosion processes, but they are often highly deficient in organic matter which can reduce soil water retention (Shrestha and Lal, 2006). In addition, developing appropriate soil structures for restoration, for example technosoils, can be expensive and demanding in terms of time and natural resources (Rivas-Pérez et al., 2016).

Where topsoil is limiting and waste materials form the substrate for plant growth, direct seeding is the most feasible means of reinstating biodiverse plants communities, particularly at larger scales (Ceccon et al., 2016; Erickson et al., 2016a; James et al., 2011; Perring et al., 2015; Porensky et

al., 2014). However, direct seeding is inefficient in terms of the proportion of seeds that produce an established seedling; in arid ecosystems it is common for only 2–7 % of seeds to establish (Chambers, 2000; James et al., 2011; Larson et al., 2015), although the use of biochar has shown to increase these percentages (Drake et al., 2016). The early developmental life-stages of plants are usually more sensitive to environmental or edaphic constraints than the adult stages (Standish et al., 2012) and the transition from germinated seed to emerged seedling has been identified as the life-stage transition most limiting the success of direct seeding (James et al., 2011). As these first stages of plant regeneration fundamentally influence the composition of the future plant community (Jiménez-Alfaro et al., 2016), characterising abiotic factors of the edaphic environment and their effects on seeds and seedlings is necessary for developing seeding practices that can achieve the desired restoration outcomes. For example, highly erodible soils have proved to be an additional challenge for seed germination and consequently for obtaining an adequate plant cover (Bochet, 2015; Cerdà and García-Fayos, 1997, 2002; Wang et al., 2014).

With the numerous potential drivers of global change comes a wide range of potential climate change scenarios (IPCC, 2014). This hinders the incorporation of future climate predictions into restoration programs (Standish et al., 2012). In this context, more experimental studies are needed to accurately evaluate the effects of altered climatic conditions on seedling recruitment and subsequent vegetation community structure and function, all of which, in turn, are strongly linked to soil conditions (Audet et al., 2013). Such experimental approaches can be effectively addressed by manipulation of combinations of climate and soil factors under controlled conditions (Lloret et al., 2004). Here, we present a case study on plant species of the Pilbara region in the northwest of Western Australia, where we assess the effects of climate and soil factors on the restoration of semi-arid ecosystems. The Pilbara (22°03' S, 118°07' E to 23°19' S, 119°43' E) is a vast (179 000 km²) and biodiverse (~1800 plant species) semi-arid ecosystem (Erickson et al., 2016a; McKenzie et al., 2009). The region is subjected to intensive mining, particularly of iron ore, and ecological restoration following mining commonly requires reintroducing plant propagules to vastly altered growth substrates (Erickson et al. 2016a). Using five native plant species that form key elements of the vegetation of this ecosystem, the specific objectives of this study were to (i) analyse the effects of a range of climate scenarios (rainfall and temperature) and the use of alternative soil materials on seedling emergence of key native plant species from the Pilbara, (ii) determine the effects of the climate and soil scenarios on the time to emerge of these plant species, and (iii) assess the implications of these climate and soil factors on improving the restoration potential in semi-arid environments.

2 Methods

2.1 Experimental design

This study was conducted between August and December 2014 in a controlled environment room (CER) at The University of Western Australia (UWA) and a glasshouse facility at Kings Park and Botanic Garden in Perth, Western Australia. Five native plant species from five families were selected as representative of a diverse range of life-forms (e.g. perennial grass, shrub and tree components) that commonly contribute to the mature plant communities found throughout the mining intensive Pilbara region of Western Australia (Erickson et al., 2016b). These study species comprised *Acacia hilliiana* Maiden (Fabaceae), *Eucalyptus gamophylla* L'Her. (Myrtaceae), *Gossypium robinsonii* F. Muell. (Malvaceae), *Grevillea pyramidalis* R. Br. (Proteaceae) and *Triodia epactia* S. W. L. Jacobs (Poaceae). Soil materials commonly used in mine restoration operations in the Pilbara (Bateman et al., 2016; Muñoz-Rojas et al., 2016b) were collected from an active mine site in the southern part of the region and used as growth media. These materials consisted of topsoil retrieved from previously stockpiled material and an overburden waste material commonly used in landform reconstruction due to its erosive stability and physical competency.

Two experimental studies were carried out to test different climate and soil scenarios. The climate in the Pilbara region is semi-arid with mean annual rainfall ranging between 250 and 400 mm, mostly concentrated in the summer months (December to March), accounting for approximately 72 % of the total annual rainfall. This rainfall originates from sporadic summer convection thunderstorms and tropical cyclones. Mean annual temperatures range between 19.4 and 33.2 °C with average maximums over 40 °C in the summer period (Bureau of Meteorology, 2015). For the climate scenario experiment we selected a range of precipitation conditions representative of those of the Pilbara during the summer (growing season) based on the rainfall pulse duration and the amount of precipitation falling in each event (CSIRO and Bureau of Meteorology, 2007). From this selection we developed four simulated rainfall treatments, and a control (i.e. maintained at field capacity) (Table 1). Three temperature conditions were selected based on daily average temperatures in the study area (25, 30 and 35 °C). These combined rainfall ($n = 5$) and temperature ($n = 3$) treatments resulted in 15 climate scenarios and were evaluated solely in topsoil.

For the soil scenario experiment, a range of growth media blends were evaluated to assess the feasibility of using growth media mixes in restoration sites. These growth media consisted of four different blends of the soil materials collected from the mining sites: 100 % topsoil (*T*), 100 % waste (*W*), and two mixes of topsoil and waste at 50 : 50 (TW50 : 50) and 25 : 75 (TW25 : 75) ratios. Also, two watering scenarios were set up consisting of a well watered treat-

Table 1. Simulated rainfall (watering) treatments utilised in this study. Pulse durations and rainfall amounts were selected from interrogating long-term weather data accessed from the Newman Airport weather station (CSIRO and Bureau of Meteorology, 2007). Simulated rainfall treatments (R1–R4) comprised repeat daily applications of water for either 6 or 3 days and two different rainfall amounts (20 or 10 mm). The total irrigation amount of 50 or 25 mL matched the pot sizes used in this study and rainfall amount treatments required to simulate the desired simulated rain conditions.

Treatment	Control	R1	R2	R3	R4
Pulse duration (days)	–	6	6	3	3
Rainfall amount (mm)	–	20	10	20	10
Irrigation (mL)	50	50	25	50	25

ment (WW) and a water deficit treatment (WD). Both treatments were watered with 25 mL for 3 days, then 25 mL every 3rd day for WW and every 6th day for WD for a total duration of 16 days.

2.2 Experimental methods

2.2.1 Soil analyses and measurements

Topsoil and waste material from the mine site were collected and transported to the CER facilities at UWA and Kings Park in 200 L drums. To create each growth media combination, one drum of topsoil (~350 kg) and one drum of waste (~225 kg) were mixed thoroughly into the different blend proportions required (e.g. *T*, *W*, TW50 : 50 and TW25 : 75, Table 2), ensuring each blend was homogeneous once blended. From each of these growth media blends, three composited soil samples of 500 g were taken, air-dried, and sieved (2 mm mesh) for physical and chemical analysis. Soil pH and electrical conductivity (EC) were calculated in deionised water (1 : 2.5 and 1 : 5, *w/v*, respectively), with an AD 8000 microprocessor-based pH. Organic C (OC) was measured by dichromate oxidation (Walkley and Black, 1934) and total N with the Kjeldahl method (Bremner and Mulvaney, 1982). Particle size was analysed by laser diffraction using a Mastersizer 2000 (Malvern Instruments, Malvern, England) after removing the organic matter with H₂O₂. Bulk density (BD) was determined according to the method proposed by Rawls (1983).

Soil hydrological parameters (Table 2) were determined according to Conant et al. (2004) using a pressure plate device at four tensions between saturation (–0.001 kPa) and wilting point (–1500 kPa) including field capacity (–10 kPa) (Table 2). Briefly, soil samples were saturated and placed in the pressure plates and then weighed to determine moisture content after hydrostatic equilibrium was reached at each water potential.

Table 2. Soil physicochemical and hydraulic properties of growth media types (mean \pm SE, $n = 3$). EC: electrical conductivity, OC: organic C; N: total N, FC: field capacity, WP: wilting point, AWC: available water content (difference between FC and WP).

Soil	pH	EC (ms m^{-1})	OC (%)	N (%)	Clay (%)	Silt (%)	Sand (%)	Bulk density (g cc^{-1})	FC (%)	WP (%)	AWC (%)
Topsoil	7.8 \pm 0.1	46.7 \pm 0.8	0.8 \pm 0.1	0.10 \pm 0.01	4.6 \pm 0.1	24.9 \pm 0.7	70.5 \pm 0.7	1.55 \pm 0.01	28.7 \pm 0.2	8.9 \pm 0.1	19.5 \pm 0.1
TW (50 : 50)	7.6 \pm 0.1	38.5 \pm 2.5	0.4 \pm 0.1	0.03 \pm 0.01	3.1 \pm 0.1	21.9 \pm 1.7	75.0 \pm 1.6	1.57 \pm 0.01	19.1 \pm 0.4	9.1 \pm 0.1	10.0 \pm 0.3
TW (25 : 75)	7.8 \pm 0.1	38.9 \pm 2.9	0.3 \pm 0.1	0.02 \pm 0.01	2.4 \pm 0.1	12.9 \pm 0.7	84.7 \pm 0.9	1.57 \pm 0.01	17.1 \pm 0.3	87 \pm 0.2	8.1 \pm 0.2
Waste	7.3 \pm 0.1	55.7 \pm 10.7	0.1 \pm 0.1	0.01 \pm 0.01	2.1 \pm 0.3	11.8 \pm 1.0	86.0 \pm 1.0	1.57 \pm 0.01	12.4 \pm 0.4	9.1 \pm 0.2	5.4 \pm 0.2

2.2.2 Experimental set up

Seeds for each species were obtained from commercially collected seeds supplied to the mining industry for use in Pilbara restoration programs. Upon receipt at Kings Park and Botanic Garden, seeds of *Acacia hilliana*, *Eucalyptus gamophylla*, *Gossypium robinsonii*, and *Grevillea pyramidalis* were cleaned of any non-seed material (e.g. chaff in *Eucalyptus* collections) and then x-rayed to remove any empty, partially filled, or clearly non-viable seeds (Faxitron MX-20 x-ray cabinet, Tucson, Arizona, USA) following Erickson et al. (2016a). A seed was deemed filled and/or viable when the x-ray images showed no abnormalities and the image was uniform white–grey in colour. For *Triodia epactia*, a seed is dispersed in an indehiscent floret and requires removal of the floret to maximise the chances of germination (Erickson et al., 2016b). Therefore, seeds were cleaned from the covering florets structures by carefully rubbing florets on a ribbed rubber mat and separating the seed from the floret debris using vacuum separation (“Zig Zag” Selecta, Machinefabriek BV, Enkhuizen, the Netherlands). Seeds were examined under the microscope to ensure no embryo damage occurred. These cleaning processes ensured only > 95 % filled/viable material was used in each experiment and removed seed quality as a potential cause of reduced emergence.

To maximise the germination potential of each batch and accommodate seeds with primary dormancy, seed pre-treatments followed pre-treatment recommendations in Erickson et al. (2016a). Seeds of *A. hilliana* and *G. robinsonii* were treated for 1–2 min in hot water at 90 °C to break physical dormancy. Seeds of *E. gamophylla* and *G. pyramidalis* were non-dormant and did not require a pre-treatment. Once cleaned from florets, seeds of *T. epactia* were soaked for 24 h in a 1 μM concentration of karrikinolide (KAR₁; 3-methyl-2H-furo[2,3-c]pyran-2-one, synthesised following Flematti et al., 2005) and re-dried at 15 °C/15 % relative humidity for at least 2–3 days prior to sowing.

The climate scenario experiment was conducted in the CER at UWA, where temperature, CO₂ and relative humidity were controlled and monitored routinely. The CER was set to a constant 12 h day and night cycle for the duration of the experiment, where day-time temperature was the treatment temperature (25, 30 and 35 °C) and night-time temperature was set at 20 °C for all three temperature treatments. Relative

humidity was maintained at 50 % and CO₂ at 400 ppm. The soil scenario experiment was conducted in the glasshouse facilities of Kings Park and Botanic Garden where air temperatures were on average 30 °C and relative humidity \sim 50 %.

For both experiments, pots of 25 cm² surface by 12 cm height were assorted in a randomised block design and replicated 12 times. Five seeds were sown into each pot and watering regimes were imposed on day one of the experiments and applied manually using a 50 mL syringe. Volumetric soil moisture was continuously monitored across all treatments in three additional “dummy” pots. An ECHO EC-5 moisture sensor (Decagon Devices, Inc.) connected to a HOBO micro station data logger (Onset Computer Corporation, Massachusetts, USA) was inserted completely into the soil surface. Measurements of volumetric soil moisture content were recorded every 5 min for the duration of the experiment, and were later averaged for daily moisture contents (Figs. S1 and S2 in the Supplement). Air temperature was also logged in both experiments.

Seedling emergence was recorded daily in each pot for 16 days. Final emergence (%) was determined as the average emergence per pot after 16 days divided by five (the number of seeds per pot) and mean emergence time (MET) was calculated using the following equation adapted from Ellis and Roberts (1981):

$$\text{MET} = \frac{\sum Dn}{\sum n}, \quad (1)$$

where n is the number of seedlings that emerged on day D , and D is the number of days counted from the beginning of emergence.

2.2.3 Statistical analyses

Differences in seedling emergence (final proportion of emerged seedlings among climate and soil scenarios) and time to emergence among treatments were tested using analysis of variance (ANOVA). Comparisons between means were performed with the Tukey’s HSD (honestly significant difference) test ($P < 0.01$). Before ANOVA testing, the analysed variables were tested for normality and variance homogeneity using the Shapiro–Wilk and Levene tests, and data were log transformed as necessary (presented data are non-

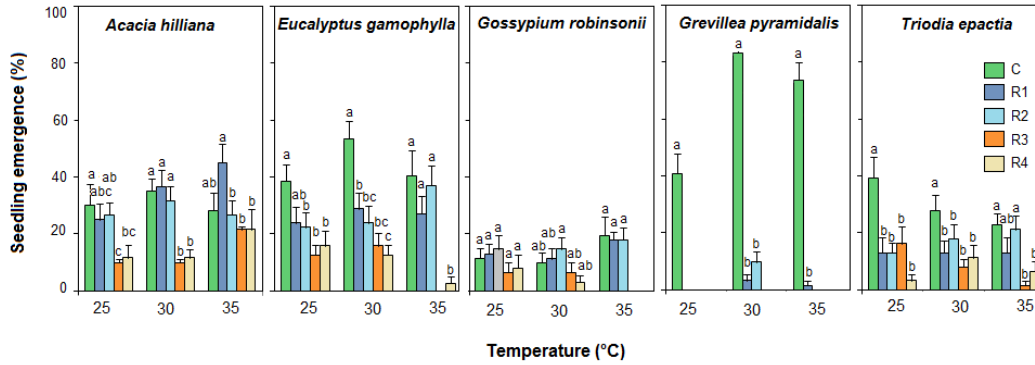


Figure 1. Total seedling emergence (% mean ± SE, $n = 12$) of Pilbara native plant species under climate scenarios (temperature and rainfall). Different letters indicate significant differences over time among watering treatments (C, R1, R2, R3 and R4) for each temperature scenario (LSD post hoc test, $P < 0.05$). Watering treatments as described in Table 1.

transformed). All analyses were performed with R statistical software version 3.1.2 (R Core Team, 2014).

3 Results and discussion

3.1 Climate effects on seedling emergence

Our results showed that seedling emergence of the Pilbara native plant species was highly dependent on soil water content in the topsoil growth media (Table 3). Total emergence varied significantly across plant species and water treatments ($P < 0.001$, Table 3; Fig. 1) and, although we did not find significant differences between temperature scenarios, interactions of temperature, water and plant species were significantly different ($P < 0.001$, Table 3).

Seedling emergence for *A. hilliana* ranged between 10 and 45 % (Fig. 1) and higher values were obtained in the control and the R1 and R2 treatments (pulse watering treatments of 10 and 20 mm daily for 6 days, respectively). The maximum number of emerged seedlings was recorded at a day temperature of 35 °C. Seedling emergence of *E. gamophylla* followed the same trend with higher emergence in the control, R1 and R2 watering treatments compared to R3 and R4 watering treatments. For this species, seedling emergence was $20.1 \pm 3.8\%$ on average and up to $40.1 \pm 6.1\%$ with available water (R1 and R2) and at 35 °C. In contrast, emergence of *G. robinsonii* was lower and differences were not significant across water and temperature treatments. Seedlings of *G. robinsonii* did not emerge at 35 °C with short initial pulses of watering (R3 and R4 watering treatments). However, maximum emergence occurred under this 35 °C temperature scenario with the 6-day pulse regime (R1 and R2). Although the maximum seedling emergence recorded for *G. pyramidalis* was higher than the other species (above 80 % in the 30 °C scenario), seedlings only emerged with continuous irrigation (control conditions); suggesting, in terms of seedling emergence, that this species has the lowest tolerance to drought.

Table 3. Effects of climate factors (temperature and water) and plant species types, and interactive effects of these factors on total emergence and mean time to emerge. Statistical significance levels: NS: not significant, *** $P < 0.001$, ** $P < 0.01$, * $P < 0.05$.

Factor	Total emergence		Mean time to emerge	
	F value	P value	F value	P value
Temperature (<i>T</i>)	2.7802	NS	15.5427	***
Water (<i>W</i>)	107.5179	***	18.0772	***
Plant species (<i>P</i>)	27.9409	***	67.2350	***
<i>T</i> × <i>P</i>	3.4951	**	3.2449	*
<i>W</i> × <i>P</i>	19.6585	***	3.8249	***
<i>T</i> × <i>W</i>	2.8951	*	0.9380	NS
<i>T</i> × <i>W</i> × <i>P</i>	3.2669	***	1.3067	NS

Patterns of seedling emergence for *T. epactia* were irregular, but in general, the seeds also proved to be dependent on higher amounts of water, and emergence generally decreased as temperature increased. Lower simulated rainfall pulse amounts seemed to be more beneficial for this species (R2 and R4).

Overall, our results showed that rainfall patterns had a large influence on seedling emergence across the five native species and suggest that seedling recruitment of these native plants may decrease in a climate scenario of increasing drought. These results are broadly consistent with other similar studies conducted in seasonally dry environments. For example, Lewandrowski (2016) found that seedling emergence of *Triodia* species decreased with water stress and high temperatures (35–40 °C). Similarly, in a study of Mediterranean shrubland of eastern Spain, Lloret et al. (2004) applied a range of warming treatments with temperature increments of 0.19–1.12 °C to analyse seedling emergence of native species. They found a moderate decrease in seedling recruitment in the warming treatments compared to the control, but differences were not statistically significant. Hogen-

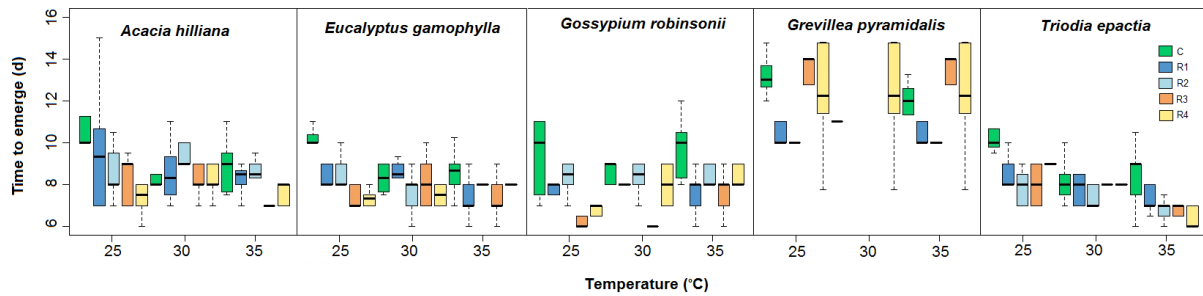


Figure 2. Mean time to emergence (days, mean \pm SE, $n = 12$) of Pilbara native plant species under climate scenarios (temperature and rainfall). Watering treatments (C, R1, R2, R3 and R4) as described in Table 1.

birk and Wein (1992) obtained larger seedling emergence at higher temperatures, but only for weedy species, suggesting that climate changes can favour weedy species over native plants. In general, the climate effect on seedling emergence seems to be more closely connected to water availability than to warming, and temperature is likely to be less of a limiting factor in the seedling emergence phase for most species (Lloret et al., 2004; Perring and Hoevendin, 2012; Woods et al., 2014).

In our study, seedling emergence responses to the watering regimes differed significantly among the five species. We found significantly decreased emergence of seedlings of *G. pyramidalis* and *G. robinsonii* under water-limited treatments, which suggest that changes in precipitation patterns can have a critical effect on the recruitment of these species. Plant species producing fewer recruits have been proposed to be more likely to disappear with drier conditions in future climate scenarios, with a consequent impact on diversity and structure of native plant communities (Lloret et al., 2004). Thus, the ability of seedlings to make use of the reduced amount of precipitation for emergence and subsequent survival will be a determinant of their distribution (Lai et al., 2015).

The mean time for emergence of the five plant species was significantly different across temperature and rainfall treatments with slightly shorter times recorded under higher temperatures, particularly in *A. hilliana* and *T. epactia* (Fig. 2); results that are in agreement with some previous studies (De Frenne et al., 2012; Richter et al., 2012). However, in the southwest of Western Australia, Cochrane et al. (2015) found that emergence of seedlings was delayed with warmer conditions, compared to control. It has been previously suggested that early emergence is a strong determinant of seedling vigour and can significantly increase plant biomass (Verdú and Traveset, 2005).

Regardless of plant species or temperature conditions, our results showed significantly higher rates of emerged seedlings with longer pulses of simulated rainfall (6 days compared to 3 days) with the same amount of accumulated water during the treatment (60 mL over the irrigation phase). Semi-arid ecosystems are particularly influenced by precipi-

tation patterns, and water availability in these environments can be highly pulsed with discrete rainfall events followed by drought periods (Miranda et al., 2001). Therefore, changes in precipitation frequency, such as rainfall pulses, can have a stronger effect than rainfall quantity in these environments (Woods et al., 2014).

Another factor that might affect plant production in global climate change scenarios is the elevated concentration of atmospheric CO₂ (IPCC, 2014). However, we have not considered this effect in this study since it is unlikely that CO₂ had any direct impact at the seedling emergence stage (Classen et al., 2010). A number of studies have previously analysed the possible impacts of CO₂ in seedling recruitment but most of them found that the response of seedlings to changes in atmospheric CO₂ are constrained by changes in precipitation patterns (Garten et al., 2008; Kardol et al., 2010).

3.2 Soil type effects on seedling emergence

Seedling emergence differed significantly between growth media types, watering treatments and plant species, but the effect of water inputs seemed to be a larger driver of emergence than growth media type ($P < 0.0001$, Table 4). With the higher soil moisture treatment (WW treatment), differences between soil materials were not significant at the $P = 0.0001$ level for *E. gamophylla*, *G. robinsonii* and *G. pyramidalis*, but emergence of *T. epactia* seedlings was significantly ($P < 0.0001$) higher in the topsoil ($56.7 \pm 7.1\%$) and the 50:50 topsoil:waste blend ($65.1 \pm 7.1\%$), as compared to the 25:75 topsoil:waste blend ($23.3 \pm 6.9\%$) and the waste ($25.1 \pm 5.6\%$) (Fig. 3). Similarly, emergence of *A. hilliana* seedlings showed a progressive decline as the amount of topsoil decreased, ranging from $58.3 \pm 6.3\%$ in the topsoil to $33.3 \pm 7.1\%$ in the waste material. In the WD scenario, seedling emergence was lower for all species with total emergence varying between $1.7 \pm 1.0\%$ in *G. pyramidalis* and $40.1 \pm 7.1\%$ in *T. epactia* in the topsoil growth media. In this water-limited scenario, seedlings of *G. pyramidalis* and *G. robinsonii* did not emerge in any growth media apart from the 100% topsoil soil type. Mean time to emergence did not differ across growth media

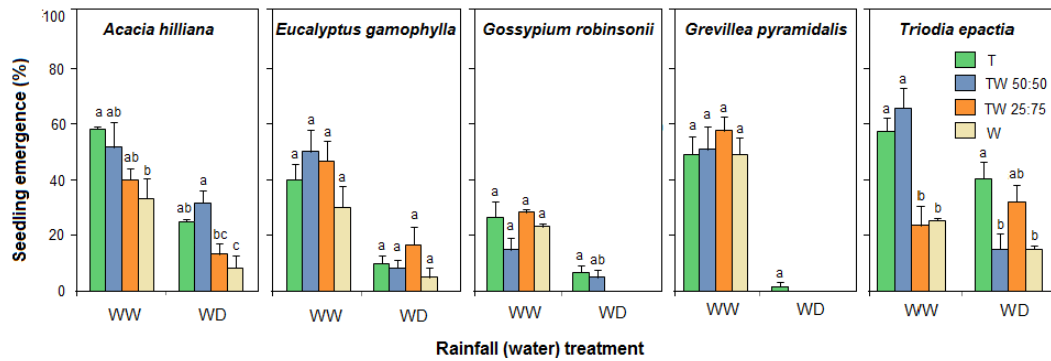


Figure 3. Total seedling emergence (%; mean ± SE, n = 12) of Pilbara native plant species for different growth media types (T: 100 % topsoil, TW 50 : 50: mix of topsoil and waste at 50 : 50 ratio, TW 25 : 75: mix of topsoil and waste at 75 : 25 ratio and W: 100 % waste) and watering treatments (WW: well watered and WD: water deficit). Different letters indicate significant differences over time among watering treatments for each temperature scenario (LSD post hoc test, P < 0.01).

Table 4. Effects of soil or growth media type, water treatments and plant species, and interactive effects of these factors on total emergence and mean time to emerge. Statistical significance levels: NS: not significant, *** P < 0.001, ** P < 0.01, * P < 0.05.

Factor	Total emergence		Mean time to emerge	
	F value	P value	F value	P value
Soil (S)	10.5853	***	0.4043	NS
Water (W)	301.1846	***	75.6453	***
Plant species (P)	19.3987	***	85.6517	***
S × P	3.07	***	0.8914	NS
W × P	12.1949	***	1.3579	NS
S × W	1.2097	NS	0.5689	NS
S × W × P	3.0291	***	1.9029	NS

types (Table 4) or in any of the interactions between growth media type, water, and plant species.

The analyses of soil physio-chemical properties showed lower contents of sand in the topsoil growth media (70.5 ± 0.7 %) consistently increased with increasing fractions of waste in the blend (Table 2). The influence of soil texture on soil water retention has been largely investigated (Saxton and Rawls, 2006) with different responses in seedling emergence (Cortina et al., 2011). Soil water holding capacity is generally higher in soils with larger clay and low sand content (Rawls et al., 2003). Higher nutrient retention in these soils rich in clay may increase seedling emergence and seedling root growth, allowing an easier extraction of water from deeper layers of the soil profile (Woodall, 2010). However, some studies showed that higher infiltration rates in soils with elevated contents of sand may increase seedling emergence allowing plants to effectively extract water following precipitation (Cortina et al., 2011).

Our study showed that seedling emergence across the five plant species was higher in the topsoil growth media which might be explained by the greater water availability as a con-

sequence of larger amounts of organic C content (Table 2). Although additional factors, such as adequate nutrient levels in the soil, can be necessary for plant establishment in degraded soils (Valdecantos et al., 2006; Brevik et al., 2015), water availability seems to be more critical at early plant life stages, particularly in semi-arid environments (Cortina et al., 2011; Miranda et al., 2001).

3.3 Implications for restoration of degraded lands

The use of growth media such as waste materials has proved to be a competent alternative to the original soil (i.e. topsoil) in restoration of degraded semi-arid areas (Machado et al., 2013; Muñoz-Rojas et al., 2006b; Rivera et al., 2014). Muñoz-Rojas et al. (2016b) showed that soil functions in a rehabilitated area of northwest Western Australia, with the use of mine waste material, can reach levels of microbial activity and organic C similar to those of topsoil once vegetation was established. However, here we show that at the early stages of plant recruitment, the use of alternative substrates depleted of organic materials can be challenging for successful seedling recruitment in the absence of water. Low contents of soil OC have been commonly associated to the loss of soil structure, which as a consequence, diminishes water holding capacity, increases bulk density, and accordingly produces soil compaction (Lal, 2004; Willaarts et al., 2016).

Overall, the results obtained in this study provide evidence that the availability of water in the soil system is a key determinant factor for increasing seedling recruitment and, therefore, optimising restoration of semi-arid lands such as the Pilbara. The application of irrigation has been proposed in restoration of semi-arid systems to control watering inputs (Bainbridge, 2002). There are several types of irrigation systems available that could effectively increase seedling recruitment, particularly in plant species most sensitive to water limitations (Padilla et al., 2009). However, there are

higher costs associated with this alternative that makes its use impractical at the landscape level (Cortina et al., 2011).

Degraded soils – frequently infertile and depleted of organic materials – can respond positively to the addition of amendments (Cortina et al., 2011; Keesstra et al., 2016b; Lozano-García et al., 2011; Valdecantos et al., 2006). Soil amendments have been commonly used in restoration to improve soil structure, restore the hydrological balance and increase the mineral nutritional capacity (Hueso-González et al., 2014; Jordán et al., 2011). Inorganic amendments (e.g. fertilisers) are usually applied to overcome plant nutritional deficiencies or physical limitations. However, the use of organic amendments such as mulch or manure has proved to increase soil water retention in soils with poor structure with a consequent increase of plant survival in mine restoration (Benigno et al., 2013). Even low doses of composted organic waste applied in degraded soils have shown to support seedling response for long periods (Fuentes et al., 2010; Yazdanpanah et al., 2016). Nevertheless, the application of organic amendments can have several implications such as competition with existing species which is compounded by the high costs of these practices at large scales in mine restoration (Cortina et al., 2011).

Since seedling establishment from seeds can be challenging in restoration (James et al., 2011), increasing seed input, or enhancing the availability of suitable micro-sites for seedling emergence through modifying the soil environment or alternatively improving the regenerative capacity of seeds represent alternative strategies for those species with limited recruitment (e.g. *G. pyramidalis* or *G. robinsonii*). Such approaches will involve new technologies for improving seed handling, processing and quality evaluation and the use of seed treatments to overcome dormancy, improve germination vigour, and increase the resilience of established seedlings (Merritt et al., 2007; Turner et al., 2013). For example, though in its infancy, seed coating procedures for native species offer promise of overcoming recruitment bottlenecks by “empowering” the seed through coating, pelleting and aggregate technologies (Madsen et al., 2014, 2016). Our results highlight the critical impact of soil water availability for seedling recruitment and the need to address this limitation, but further studies are needed to develop suitable applications and techniques in drylands restoration at a management scale. It would be useful to transfer the experiments reported here to larger-scale field trials to effectively assess applicability of the findings into restoration programs.

4 Conclusions

Seedling recruitment of the five native plants was highly dependent on soil moisture and temperature did not have a significant effect on the number of emerging seedlings. Emergence across the five plant species was higher in the top-soil growth media compared to the other soil materials, most

likely due to its larger available water content as a consequence of increased amounts of organic C. Overall, under drought scenarios total seedling emergence was below 40 % for all species and growth media types. In general, responses to the climate scenarios differed significantly among the five native species suggesting that future climate scenarios of increasing drought might affect not only seedling recruitment, but also diversity and structure of native plant communities. In particular, we found significantly decreased emergence rates in seedlings of *G. pyramidalis* and *G. robinsonii* under water-limited treatments meaning that changes in precipitation patterns may have a critical effect on the recruitment of these species. The use of growth media such as waste materials from mining operations could be an alternative to the scarce topsoil. However, at early plant stages the use of these alternative substrates that are depleted of organic materials can be challenging for successful seedling recruitment in the absence of water. These limitations could be overcome by using soil amendments but the cost associated with these solutions at large landscape scales needs to be addressed.

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