

SCIENTIFIC REPORTS



OPEN

High nutrient-use efficiency during early seedling growth in diverse *Grevillea* species (Proteaceae)

Tianhua He, William M. Fowler & Casey L. Causley

Received: 26 February 2015

Accepted: 09 September 2015

Published: 26 November 2015

Several hypotheses have been proposed to explain the rich floristic diversity in regions characterised by nutrient-impooverished soils; however, none of these hypotheses have been able to explain the rapid diversification over a relatively short evolutionary time period of *Grevillea*, an Australian plant genus with 452 recognised species/subspecies and only 11 million years of evolutionary history. Here, we hypothesise that the apparent evolutionary success of *Grevillea* might have been triggered by the highly efficient use of key nutrients. The nutrient content in the seeds and nutrient-use efficiency during early seedling growth of 12 species of *Grevillea* were compared with those of 24 species of *Hakea*, a closely related genus. Compared with *Hakea*, the *Grevillea* species achieved similar growth rates (root and shoot length) during the early stages of seedling growth but contained only approximately half of the seed nutrient content. We conclude that the high nutrient-use efficiency observed in *Grevillea* might have provided a selective advantage in nutrient-poor ecosystems during evolution and that this property likely contributed to the evolutionary success in *Grevillea*.

Mediterranean climate regions, such as those in southwest Western Australia (SWA), the South African Cape region, and the Mediterranean Basin, have particularly diverse species and endemic rich flora and are considered globally significant^{1,2}. Numerous hypotheses have been proposed to explain the floristic diversity of these biodiversity hotspots³. For example, Hopper proposed that landscape age contributed to the high diversity in SWA⁴, and Lengyel *et al.* reported higher global diversification rates in myrmecochorous lineages relative to their sister groups and speculated that myrmecochory contributed to the high diversification of SWA flora⁵. However, none of the current hypotheses can explain the extreme diversity of many species-rich genera in Australia, such as *Grevillea* in the family Proteaceae.

Grevillea is a predominantly Australian genus containing 362 currently recognised species or 452 taxa, which includes subspecies, with five species present in New Guinea, New Caledonia, and Sulawesi⁶. The evolutionary time period for species accumulation has emerged as a significant explanatory variable for biodiversity^{3,4}, and this explanation is supported by fossil evidence⁷. *Grevillea* is one of the youngest genera, with only *c.* 11.2 million years of evolutionary history⁸, and it is the most species-rich genus in the Proteaceae family; thus, the greater-time-for-speciation theory does not explain the high numbers of species in this genus. However, Hopper proposed that taxa in old and climate-buffered landscapes, including *Grevillea* and other genera, such as the closely related genus *Hakea*, typically have a low extinction rate⁴. *Grevillea* species are generally dispersed by ants, which has been hypothesised to be a key evolutionary novelty that may account for the high diversification rate in this genus⁵. However, many other plant lineages in SWA, typically species that co-occur with *Grevillea* and are also dispersed by ants, have markedly fewer species within their genera. For example, ant-dispersed *Daviesia* (Fabaceae) diversified into 127 species over *c.* 26 million years, whereas ant-dispersed *Bossiaea* (Fabaceae) diversified into 80 species within *c.* 10 million years of evolutionary history³. Therefore, myrmecochory alone is unlikely to contribute to the high diversification rate in *Grevillea*. With a relatively short evolutionary history of 11.2 million years and as the third largest plant genus in Australia after *Acacia* and *Eucalyptus*, *Grevillea* probably has one of the highest net speciation rates of all large Australian plant clades.

Department of Environment and Agriculture, Curtin University, PO Box U1987, Perth, WA 6845, Australia. Correspondence and requests for materials should be addressed to T.H. (Tianhua.He@curtin.edu.au)

	<i>Hakea</i>	<i>Grevillea</i>	t-test, P value
Number of species	149 ^a	362 ^b	–
Age (million years) ^c	15.8	11	–
Seed mass (mg) ^d	30.4 (2.6–501.0)	34.9 (2.8–244.0)	–
Seed mass (mg) ^e	33.3 ± 29.2	39.5 ± 15.5	t = –0.40, P = 0.703
Seed N (%)	7.61 ± 1.36	3.53 ± 0.79	t = –1.09, P < 0.001
Seed P (%)	1.36 ± 0.38	0.63 ± 0.23	t = –1.48, P < 0.001
Germination rate	0.82 ± 0.12	0.32 ± 0.15	t = –5.07, P < 0.001
Days to first true leaf	32 ± 6	31 ± 5	t = 0.02, P = 0.942
Number of leaves	16.7 ± 5.4	8.3 ± 1.2	t = –3.57, P = 0.002
Root/shoot biomass	1.67 ± 0.71	1.34 ± 0.55	t = –1.27, P = 0.218
Total dry biomass (g)	0.54 ± 0.39	0.28 ± 0.13	t = –1.09, P = 0.093
Root length (mm)	468 ± 198	580 ± 181	t = –1.48, P = 0.152

Table 1. Seed mass and nutrient concentration (N and P), seed generation, and seedling growth in *Hakea* and *Grevillea*. ^a6; ^b6; ^c8; ^d28; ^e:species examined in this study.

Species in the family Proteaceae are highly adapted to and became diversified on the most nutrient-impoverted soils observed worldwide. Highly infertile soils have long been demonstrated to influence biological and physiological responses and speciation patterns^{4,9}. Species have developed diverse morphological and physiological adaptations to limited nutrient availability in order to successfully establish in nutrient-limited soils^{10–13}. These species typically exhibit highly efficient photosynthetic phosphorus use¹⁴.

In southwestern Australia, seed germination and successful seedling establishment predominantly occurs following winter rains after summer fires. Seed-stored nutrients, particularly nitrogen (N) and phosphorus (P), are crucial for the emergence and early growth of seedlings, and the efficient use of these nutrients is critical for the successful establishment of seedlings in severely nutrient-poor and water-limited environments. Nutrient limitation has recently been proposed as a potential driver for variation in species diversity¹⁵. Lambers *et al.* suggested that high plant species diversity on infertile soils, particularly in species-rich biomes, is associated with functional diversity for nutritional strategies, including nutrient-use efficiency¹⁶. Despite extensive research on plant species richness in relation to nutrient availability, few studies have attempted to explore the relationship between nutrient-use efficiency and speciation patterns. Here, we compare the nutrient concentration and content in seeds and nutrient-use efficiency during early seedling growth of *Grevillea* with those of the closely related genus *Hakea*, another Australian genus with 149 species that efficiently use seed nutrients during early seedling growth¹⁶. Our study aimed to provide unique insights into the potential mechanisms underlying the evolutionary success (i.e., high diversification rate) of *Grevillea*.

Results

Seed mass varies markedly in both *Grevillea* (2.8–244 mg, 116 species/subspecies surveyed) and *Hakea* (2.6–501 mg, 153 species/subspecies surveyed), and *Grevillea* has a slightly higher median seed mass than *Hakea* (24.2 vs. 18.7 mg). The average seed masses of the 30 *Grevillea* species and 29 *Hakea* species used in the present study were statistically similar (Table 1). Nutrient assays on the 30 *Grevillea* species and 29 *Hakea* species revealed significantly higher concentrations of N and P in the *Hakea* seeds than in the *Grevillea* seeds (Table 1; Table S1). The total N and P contents per seed in *Hakea* were also much higher than in *Grevillea* (N: 2.66 vs. 0.56 mg, $P = 0.002$; P: 0.47 vs. 0.17 mg, $P = 0.008$; Fig. 1).

The seeds of the 25 *Hakea* species that were tested readily germinated without requiring pre-treatment, and they had an overall germination rate of 82%. *Grevillea* seeds (29 species) pre-treated with smoke water exhibited a significantly lower germination rate of 32% (Table 1). However, for the 24 *Hakea* and 12 *Grevillea* species that yielded a sufficient number of germinants for growth comparisons, the time period to expand the first true leaves was similar, although *Grevillea* plants had significantly fewer leaves after growing for 100 days. Both *Hakea* and *Grevillea* diverted more resources to the roots than to the shoots, with a statistically similar root:shoot ratio. *Grevillea* and *Hakea* species accumulated similar amounts of dry biomass and had similar root depths in the soil after 100 days of growth (Table 1).

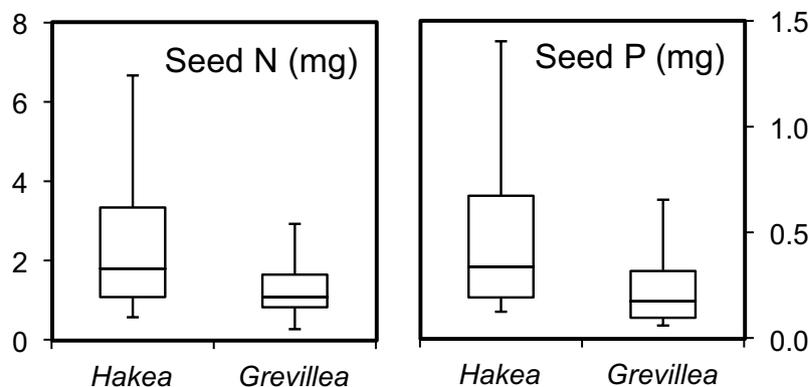


Figure 1. Comparison of nitrogen (N) and phosphorus (P) content per seed of the test species of *Hakea* and *Grevillea*. The middle lines in the box represent the median values, and the bottom and top of the box are the first and third quartiles. The ends of the whiskers are 1.5 interquartile range (IQR) above the third quartile and 1.5 IQR below the first quartile.

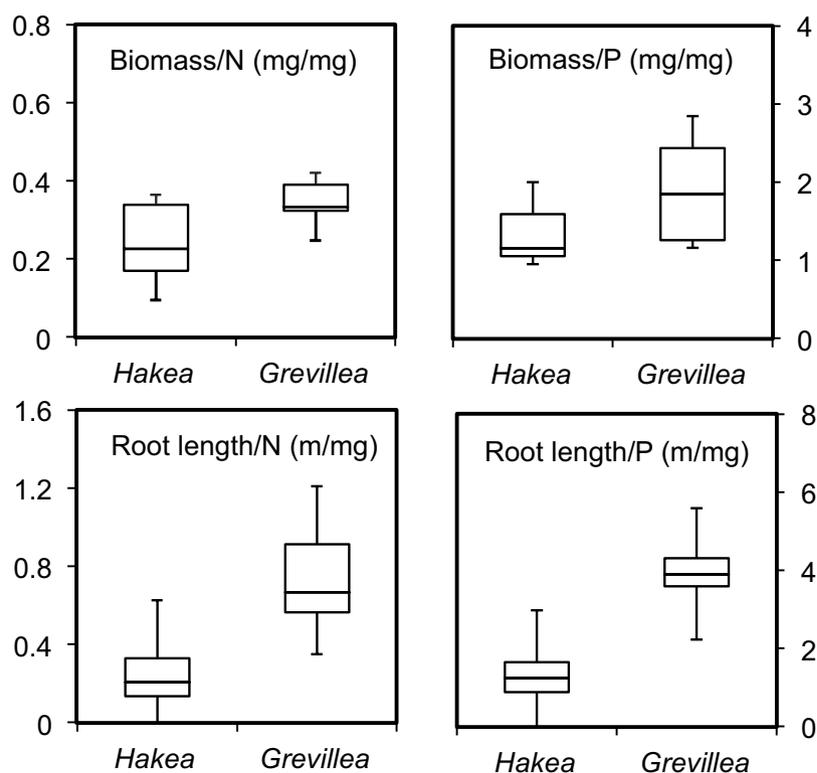


Figure 2. The nutrient-use efficiency (N and P) of the test species of *Hakea* and *Grevillea*. The middle lines in the box represent the median values, and the bottom and top of the box are the first and third quartiles. The ends of the whiskers are 1.5 interquartile range (IQR) above the third quartile and 1.5 IQR below the first quartile.

Further analysis of the seed nutrient content and seedling growth rate revealed that *Grevillea* species had significantly higher nutrient-use efficiencies (Table 1; Fig. 2). Compared with *Hakea*, for each unit of P in the seed, *Grevillea* accumulated 1.6-fold more dry biomass and had roots that descended 2.9 times deeper into the soil, whereas for each unit of N in the seed, *Grevillea* accumulated 1.4-fold more dry biomass and had taproots that descended 3.1 times deeper into the soil (Table 1, Fig. 2).

Discussion

The seed nutrient concentration in *Grevillea* reported in the present study, particularly with respect to P, was only half of that reported in previous scattered studies. For example, Hocking reported an average of $11.4 \pm 2.6 \text{ mg.g}^{-1} \text{ P}$ for ten *Grevillea* species¹⁷. The results of the present study on 30 species reported

an average of $6.3 \pm 2.3 \text{ mg.g}^{-1}$. This discrepancy is unlikely to be a consequence of methodological errors in our analysis because the 29 *Hakea* species were assayed in the same batch in the present study, and they yielded an average P content of $13.6 \pm 3.8 \text{ mg.g}^{-1}$, which is similar to the $15.1 \pm 4.2 \text{ mg.g}^{-1}$ P concentration reported for 13 species from ten previous studies (except *Hakea pycnoneura*, which was reported to have an unusually high P concentration of 36 mg.g^{-1})¹⁸. Interestingly, although the N and P concentrations in the seeds were lower in *Grevillea* than in *Hakea*, the N:P ratios in both taxa groups were similar. Notably, the lower seed N and P concentrations in *Grevillea* relative to those in *Hakea* did not reflect dilution by seed mass because the tested species in the two groups had similar seed masses.

With only half the seed nutrient content compared with species from the closely related genus *Hakea*, the species in *Grevillea* achieved similar levels of seedling growth during the early stages. Indeed, both groups of plants expanded their first true leaves within a similar time, accumulated similar amounts of dry biomass, and diverted similar proportions of resources to the shoots and roots. Consequently, the results of this analysis suggest that compared with *Hakea*, *Grevillea* species use nutrients significantly more efficiently than do *Hakea*. This efficiency represents a clear ecological advantage over *Hakea* because *Grevillea* species can accumulate twice as much dry biomass and descend their roots three times deeper into the soil for every unit of nutrient in their seeds.

Seed-stored nutrients (N and P) are crucial for the successful establishment of seedlings in nutrient-impooverished landscapes. The rapid descent of the taproots to reach more reliable soil moisture reserves prior to the onset of summer droughts is critical for the survival of seedlings, particularly in Mediterranean climates with dry summer conditions¹⁹. Using fewer nutrients, *Grevillea* species can grow roots equal in length to those of *Hakea* and survive dry Mediterranean-type summers equally as well, and these characteristics provide a selective advantage to *Grevillea* compared with other organisms that might require additional nutrients in nutrient-impooverished soils.

Seed production is inversely proportional to the per-seed reserve of nutrients, particularly P, in nutrient-impooverished soils²⁰. In the fire-prone environments of SWA, which are generally characterised by poor soil, resprouters (populations that predominantly regenerate through sprouts from the trunk or underground organs) typically produce fewer seeds than nonsprouters (populations that predominantly regenerate through seeds)²¹. However, no significant differences were observed in the P content between nonsprouting and resprouting species growing on P-impooverished soil in a survey of 41 species in SWA¹⁸. The lower nutrient requirements for seedling growth in *Grevillea* relative to *Hakea*, which has much higher nutrient (N and P) concentrations, might allow *Grevillea* species to produce more seeds regardless of the post-fire regeneration mode. Because of challenges in gauging the size of the soil seedbank, systematic surveys of *Grevillea* seed production are rare, although Pickup *et al.* reported a large seedbank for *Grevillea rivularis* at $193 \pm 73 \text{ seeds m}^{-2}$ ²². Because of its capacity to grow to 2.5 m tall and 3 m wide, this plant species has the potential to produce up to 2000 seeds per plant, which is much higher than that of most *Hakea* species²³. Levin²⁴ proposed that propagule-rich lineages are likely to have high speciation rates because the increased number of seeds increases the opportunities for ecological and geographical speciation, and this hypothesis could also be true for *Grevillea*. Interestingly, high propagule pressure has also been proposed as one of the mechanisms facilitating successful invasion²⁵.

Species in the Proteaceae family grow well in the world's most nutrient-impooverished landscapes, particularly P-impooverished soils. Apart from diverse and efficient nutrient-acquisition strategies²⁶, high nutrient-use efficiency is likely another key adaptation. Recent studies have observed high photosynthetic P-use efficiency in SWA Proteaceae species¹⁰. Lambers *et al.* reported that Proteaceae species such as *Hakea* species (*Grevillea* species were not included) from severely P-impooverished soils extensively replace phospholipids with galactolipids and sulpholipids during leaf development to achieve a high photosynthetic P-use efficiency, and proposed the “sulphur-for-phosphorus” hypothesis¹⁶. Sulpice *et al.*²⁷ observed that Proteaceae species grow with very low levels of ribosomes, particularly plastidic ribosomes, at early stages of leaf development. Many of the above studies involved *Hakea* species, although they rarely included *Grevillea* species; thus, additional research is required to test whether *Grevillea* have developed peak nutrient-use efficiency.

It can be speculated that once *Grevillea* overcome the barriers imposed by the two most limiting factors to its development, low soil nutrient availability and periodic water availability, these plants can invade most habitats. Indeed, species of *Grevillea* have been observed throughout Australia. High nutrient-use efficiency has interacted with multiple factors, including myrmecochory, a predominantly sexual reproduction strategy, and a high seed output, to allow the *Grevillea* genus to diversify into a group with 452 species/subspecies within only 11 million years.

Methods

The seeds of *Grevillea* and *Hakea* used for the nutrient analyses and germination and seedling growth comparisons were acquired from Nindethana Seed Service (Albany, Western Australia, Australia). Genus-wide seed mass information was extracted from the Seed Information Database²⁸. The nutrient analysis was conducted on seeds from 59 species (29 *Hakea* and 30 *Grevillea*, Table S1). These species cover the major intra-generic taxonomy sections and provenances; therefore, they represent a wide diversity of the respective genera. For each species, up to 20 g of dry seeds were analysed for N and P concentrations at the ChemCentre (Perth, Western Australia, Australia) using the combustion method for N and inductively coupled plasma spectrometry for P.

A total of 52 species (25 *Hakea* and 27 *Grevillea* species, Table S1) were used in the germination trials. Each species had three replicates, and there were up to 30 seeds per replicate (depending on seed size). Each replicate was contained within a 120-mm sterile Petri dish with 2 layers of sterilized Whatman Grade 1 filter paper and sealed with Parafilm. Petri dishes with *Hakea* seeds were filled with 5 mL autoclaved deionised water, whereas those with *Grevillea* seeds (because of inherent dormancy) were filled with 5 mL 10% smoke water solution (Regen 2000 Smokemaster); previous studies have suggested that *Grevillea* seed germination is enhanced by smoke stimulation²⁹. To reduce the risk of fungal growth throughout the germination experiment, the smoke water was passed through a 0.20- μ m filter. Germination was conducted in an environmental chamber at a constant 15°C with a 12 hour light/dark cycle. The germinants (seeds with radicle emergence ≥ 1 mm) were counted and recorded every two days for a period of up to 60 days. Seeds that did not germinate were assessed for viability using a cut test to visually assess the health of the embryo. The Petri dishes were randomised within the environmental chamber in each experiment.

For each of the species germinated in the above-described germination phase (12 *Grevillea* and 24 *Hakea* species, with no or minimal germination recorded in some species), ten germinants (or fewer when available) were individually planted in PVC tubes (100 cm in length and 5 cm in diameter) with a substrate of washed white sand (with no nutrients) and placed in a hoop house. The seedlings were watered (20 mL) every 2 to 3 days and grown for 100 days (from late winter to spring 2014) prior to harvesting. The measurements taken at harvest were leaf number, shoot length, root length and fresh and dry shoot and root mass. The dry biomass of the shoot and root was recorded after the samples were oven dried at 70°C for 48 hours.

The nutrient-use efficiency (N and P) was defined as the amount of biomass produced or root length gained per unit of nutrient (mg) in the seeds. The nutrient-use efficiencies of *Hakea* and *Grevillea* were compared. Comparisons of the parameter values between *Grevillea* and *Hakea* were analysed with a t-test and implemented in PAST³⁰. $P < 0.05$ was considered to represent statistical significance.

References

1. Cowling, R. M., Rundel, P. W., Lamont, B. B., Arroyo, M. K. & Arianoutsou, M. Plant diversity in mediterranean-climate regions. *Trends Ecol. Evol.* **11**, 362–366 (1996).
2. Myers, N., Mittermeier, R. A., Mittermeier, C. G., da Fonseca, G. A. B. & Kent, J. Biodiversity hotspots for conservation priorities. *Nature* **403**, 853–858 (2000).
3. Cook, L. G., Hardy, N. B. & Crisp, M. D. Three explanations for biodiversity hotspots: small range size, geographical overlap and time for species accumulation. An Australian case study. *New Phytol.* doi: 10.1111/nph.13199 (2014).
4. Hopper, S. D. OCBIL theory: towards an integrated understanding of the evolution, ecology and conservation of biodiversity on old, climatically buffered, infertile landscapes. *Plant Soil* **322**, 49–86 (2009).
5. Lengyel, S., Gove, A. D., Latimer, A. M., Majer, J. D. & Dunn, R. R. Ants sow the seeds of global diversification in flowering plants. *PLoS ONE* **4**, 5 (2009).
6. Makinson, R. O. In *Flora of Australia*, Vol. 17A (ed Makinson, O.) Ch. 1, 1–20 (CSIRO Publishing, 2000).
7. Sniderman, J. M. K., Jordan, G. J. & Cowling, R. M. Fossil evidence for a hyperdiverse sclerophyll flora under a non-Mediterranean-type climate. *Proc. Natl. Acad. Sci. USA* **110**, 3423–3428 (2013).
8. Sauquet, H. *et al.* Contrasted patterns of hyperdiversification in mediterranean hotspots. *Proc. Natl. Acad. Sci. USA* **106**, 221–225 (2009).
9. Orians, G. H. & Milewski, A. V. Ecology of Australia: the effects of nutrient-poor soils and intense fires. *Biol. Rev.* **82**, 393–423 (2007).
10. Denton, M. D., Veneklaas, E. J., Freimoser, F. M. & Lambers, H. *Banksia* species (Proteaceae) from severely phosphorus-impooverished soils exhibit extreme efficiency in the use and re-mobilization of phosphorus. *Plant Cell Environ.* **30**, 1557–1565 (2007).
11. Lambers, H. & Shane, M. W. Role of root clusters in phosphorus acquisition and increasing biological diversity in agriculture in Scale and Complexity in Plant Systems Research: Gene-Plant-Crop Relations (eds Spiertz, J. H. J., Struik, P. C., Laar, H. H.) 237–250 (Springer, 2007).
12. Lambers, H., Raven, J. A., Shaver, G. R. & Smith, S. E. Plant nutrient-acquisition strategies change with soil age. *Trends Ecol. Evol.* **23**, 95–103 (2008).
13. Sander, J. & Wardell-Johnson, G. Fine-scale patterns of species and phylogenetic turnover in a global biodiversity hotspot: implications for climate change vulnerability. *J. Veg. Sci.* **22**, 766–780 (2011).
14. Lambers, H. *et al.* Proteaceae from severely phosphorus-impooverished soils extensively replace phospholipids with galactolipids and sulfolipids during leaf development to achieve a high photosynthetic phosphorus-use efficiency. *New Phytol.* **196**, 1098–1108 (2012).
15. Venterink, O. H. Does phosphorus limitation promote species-rich plant communities? *Plant and Soil* **345**, 1–9 (2011).
16. Lambers, H., Brundrett, M. C., Raven, J. A. & Hopper, S. D. Plant mineral nutrition in ancient landscapes: high plant species diversity on infertile soils is linked to functional diversity for nutritional strategies. *Plant and Soil* **334**, 11–31 (2010).
17. Hocking, P. J. Mineral nutrient composition of leaves and fruit of selected species of *Grevillea* from south-western Australia, with special reference to *Grevillea leucopteris* Meissn. *Aust. J. Bot.* **34**, 155–164 (1986).
18. Groom, P. K. & Lamont, B. B. Phosphorus accumulation in Proteaceae seeds: a synthesis. *Plant Soil* **334**, 61–72 (2010).
19. Enright, N. J. & Lamont, B. B. Survival, growth and water relations of *Banksia* seedlings in a sand mine rehabilitation site and adjacent scrub-heath sites. *J. Appl. Ecol.* **29**, 663–671 (1992).
20. Henery, M. L. & Westoby, M. Seed mass and seed nutrient content as predictors of seed output variation between species. *Oikos* **92**, 479–490 (2001).
21. Lamont, B. B. & Wiens, D. Are seed set and speciation always low among species that resprout after fire, and why? *Evol. Ecol.* **17**, 277–292 (2003).
22. Pickup, M., McDougall, K. L. & Whelan, R. J. Fire and flood: soil-stored seed bank and germination ecology in the endangered Carrington Falls *Grevillea* (*Grevillea rivularis*, Proteaceae). *Austral Ecol.* **28**, 128–136 (2003).
23. Groom, P. K. & Lamont, B. B. In *Gondwanan heritage. Past, present and future of the Western Australian biota* (eds Hopper, S. D. *et al.*) 39–24 (Surrey Beatty and Sons, 1996).

24. Levin, D. A. Ancient dispersals, propagule pressure, and species selection in flowering Plants. *Syst. Bot.* **31**, 443–448 (2006).
25. Lockwood, J. L., Cassey, P. & Blackburn, T. M. The more you introduce the more you get: the role of colonization pressure and propagule pressure in invasion ecology. *Divers. Distrib.* **15**, 904–910 (2009).
26. Lambers, H. in *Plant life on the sandplains in Southwest Australia, a global biodiversity hotspot* (ed Lambers, H.) Ch. 10, 307–315 (UWA Publishing, 2014).
27. Sulpice, R. *et al.* Low levels of ribosomal RNA partly account for the very high photosynthetic phosphorus-use efficiency of Proteaceae species. *Plant Cell Environ.* **37**, 1276–1298 (2014).
28. Royal Botanic Gardens Kew. Seed Information Database (SID). Version 7.1. (2014) Available at: <http://data.kew.org/sid/> Date of access: 10/12/2014.
29. Morris, E. C. Germination response of seven east Australian *Grevillea* species (Proteaceae) to smoke, heat exposure and scarification. *Aust. J. Bot.* **48**, 179–189 (2000).
30. Hammer, Ø., Harper, D. A. T. & Ryan, P. D. 2001. PAST: Paleontological statistics software package for education and data analysis. *Palaentol. Electron.* **4**, 1–9 (2001).

Acknowledgements

This work was supported by Australian Research Council (DP120103389). TH acknowledges the support from a Curtin Research Fellowship. We thank Barry Price (ChemCentre, Western Australia) for assistance in seed nutrient assay, and Sophie Fox, Sh-Hoob El-Ahmirand, Thien Tran Duc for assistance in glasshouse and germination components.

Author Contributions

T.H. and C.C. designed the experiment, W.F. conducted the germination and growth experiment, T.H. and W.F. analysed the data; C.C. prepared materials for nutrient analysis. T.H. and W.F. wrote manuscript; All authors contributed to revision and reviewed the manuscript.

Additional Information

Supplementary information accompanies this paper at <http://www.nature.com/srep>

Competing financial interests: The authors declare no competing financial interests.

How to cite this article: He, T. *et al.* High nutrient-use efficiency during early seedling growth in diverse *Grevillea* species (Proteaceae). *Sci. Rep.* **5**, 17132; doi: 10.1038/srep17132 (2015).



This work is licensed under a Creative Commons Attribution 4.0 International License. The images or other third party material in this article are included in the article's Creative Commons license, unless indicated otherwise in the credit line; if the material is not included under the Creative Commons license, users will need to obtain permission from the license holder to reproduce the material. To view a copy of this license, visit <http://creativecommons.org/licenses/by/4.0/>