

# Are seed mass and seedling size and shape related to altitude? Evidence in *Gymnocalycium monvillei* (Cactaceae)

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**Abstract:** Several studies reported a negative relationship between altitude and seed mass. In cactus species, seed mass has been also related to seedling morphology (size and shape). Here we studied *Gymnocalycium monvillei* (Lem.) Pfeiff. ex Britton & Rose, a cactus species with a wide altitudinal distribution, with the main aim of analyzing how altitude affects seed mass and seedling size (height and width) and shape (globose or columnar). We collected seeds from five sites along the entire altitudinal distribution of the species in the Córdoba Mountains (sites were located between 878 and 2230 m a.s.l.), encompassing a marked climatic gradient (6 °C of mean annual temperature difference between the extreme sites). Seed mass and seedling traits were measured in the laboratory. Seedling height increased with altitude, whereas seed mass was not related to this parameter. Seedlings became more globose (reduced surface/volume ratio) with decreasing altitude. Variation in seedling shape along the altitudinal gradient may be related to the contrasting climatic conditions to which seedlings are exposed, and could account for the wide altitudinal distribution of *G. monvillei*. Our results highlight the importance of seedling traits in the species' response to climatic change.

**Key words:** altitudinal gradients, *Gymnocalycium*, seed mass, seedling morphology, Cactaceae.

**Résumé :** Plusieurs études ont rapporté une relation négative entre l'altitude et la masse des semences. Chez les cactées, la masse des semences a aussi été reliée à la morphologie des semis (taille et forme). Les auteurs ont étudié ici *Gymnocalycium monvillei* (Lem.) Pfeiff. ex Britton & Rose, une espèce de cactus ayant une large distribution altitudinale, ayant comme but principal d'analyser comment l'altitude affecte la masse des semences et la taille (hauteur et largeur) et la forme (globulaire ou columnaire) des semis. Ils ont récolté des graines à partir de cinq sites couvrant l'entière distribution altitudinale de l'espèce dans les sierras de Córdoba (les sites étaient localisés à des altitudes de 878 à 2230 m au-dessus du niveau de la mer), englobant un gradient climatique marqué (une différence de température annuelle moyenne de 6 °C entre les sites aux extrémités). La masse des semences et les traits des semis ont été mesurés en laboratoire. La hauteur des semis augmentait en fonction de l'altitude, alors que la masse des semences n'était pas reliée à ce paramètre. Les semis devenaient davantage globuleux (un ratio surface/volume réduit) en fonction de la diminution de l'altitude. La variation de la forme des semis le long du gradient altitudinal peut être reliée aux conditions climatiques contrastées auxquelles les semis sont exposés, et elles pourraient expliquer la large distribution altitudinale de *G. monvillei*. Les résultats des auteurs soulignent l'importance des traits des semis dans la réponse de l'espèce aux changements climatiques. [Traduit par la Rédaction]

**Mots-clés :** gradients altitudinaux, *Gymnocalycium*, masse des semences, morphologie des semis, Cactaceae.

## Introduction

Seed size has been related to both latitudinal and altitudinal gradients. Baker (1972) and Murray et al. (2003) found that seed size decreases with increasing altitude, whereas Moles et al. (2007) and Murray et al. (2003) found that seed size decreases with increasing latitude. These patterns have been attributed to the increasing severity of environmental conditions as temperature decreases, reduced amount of photosynthates available to seeds, or simply to changes in plant growth forms along the gradients. However, how this pattern is maintained at the intraspecific level is still not clear (Pluess et al. 2005; Guo et al. 2010). In addition, to our knowledge, no studies have analyzed inter- or intra-specific changes in seed size along altitudinal gradients in cactus species.

Seedling size is an important character determining plant survival and growth (Leishman et al. 2000), and depends strongly on seed size (Wulff 1986; Moles and Westoby 2004). In a study involving 17 cactus species, Sosa Pivatto et al. (2014) found that seed size was related not only to seedling size, but also to seedling shape: species with bigger seeds produced bigger seedlings, which were also more columnar (as adults, cactus seedlings are succulent). The authors stated that columnar seedlings would be better competitors for light, but globose seedlings would be more resistant to higher temperatures or drought events because of their lower surface/volume ratio (Felger and Lowe 1967; Cornejo and Simpson 1997). Low surface/volume ratio means that cacti have a small photosynthetic surface compared with the proportionally large respiratory demand of their water-storing parenchyma, and are

Received 31 January 2015. Accepted 11 May 2015.

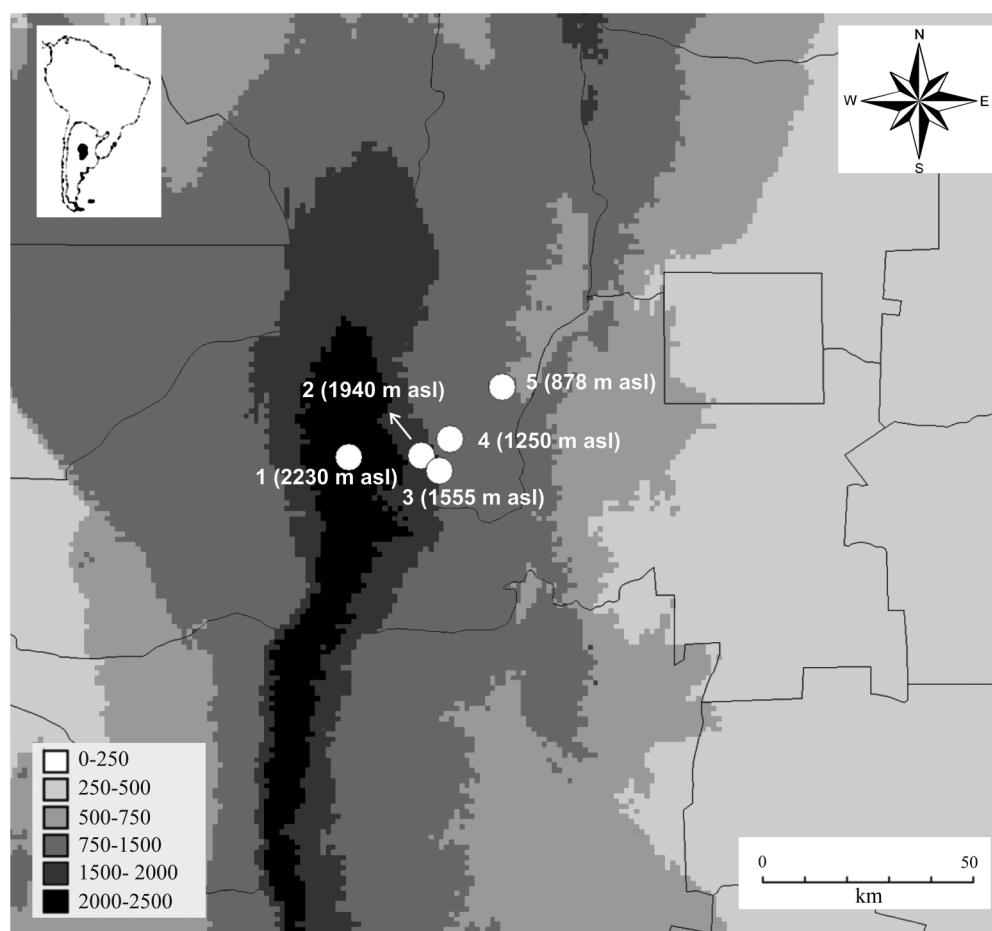
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**Fig. 1.** Location of the five collection sites of seeds of *Gymnocalycium monvillei* along an altitudinal gradient in the Córdoba Mountains, central Argentina.



therefore restricted by the amount of photosynthetically active radiation that is intercepted by the plants (Nobel 2003; Martorell et al. 2006).

*Gymnocalycium monvillei* (Lem.) Pfeiff. ex Britton & Rose is a globose cactus endemic to the Córdoba Mountains, central Argentina, and presents a very wide altitudinal distribution (Charles 2009; Gurvich et al. 2014). This gradient is related to important differences in temperature and humidity, which generate very different environmental conditions that could affect germination and establishment (García-Pérez et al. 2007). Studies of regenerative traits along gradients may help to understand how species cope with environmental conditions, and are necessary to predict the effect of climatic change on species abundances and distribution (Butler et al. 2012; Dávila et al. 2013). The aim of this study was to analyze how altitude affects seed size, seedling size, and seedling shape (columnar or globose) of the cactus species *G. monvillei* by comparing different populations along its entire altitudinal distribution.

## Materials and methods

### Study species, area characteristics, and measured variables

*Gymnocalycium monvillei* inhabits rocky outcrops from around 880 to 2200 m a.s.l. (Demaio et al. 2011; Gurvich et al. 2014). We studied five populations located along an altitudinal gradient between the localities of Cuesta Blanca ( $31^{\circ}28' S$ ,  $64^{\circ}34' W$ ) and Pampa de Achala ( $31^{\circ}41' S$ ,  $64^{\circ}50' W$ ), at 878, 1250, 1555, 1940, and 2230 m a.s.l. (Fig. 1), each population corresponding to an altitudinal class. Mean annual temperature and precipitation varied from  $16.5^{\circ}C$  and 680 mm, to  $10.3^{\circ}C$  and 790 mm, respectively,

between the lowest and highest sites of this altitudinal gradient (de Fina 1992). Vegetation varied from subtropical dry forest to cold-temperate grasslands at the extreme sites (Giorgis et al. 2011). Frost can occur all year round and snow events are common during winter at sites above 1900 m a.s.l.

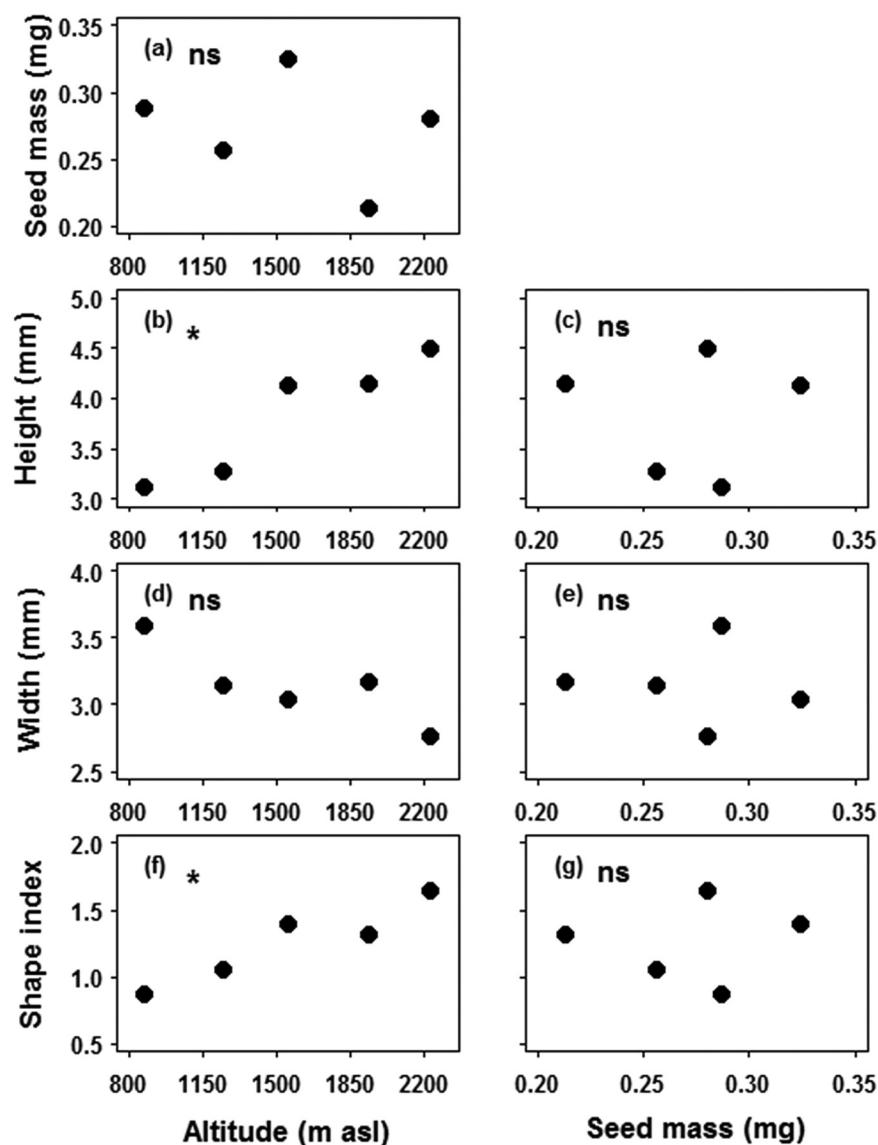
Mature fruits from 20 individuals were collected from each altitudinal class. Fruits were collected from individuals of similar sizes (around 10 cm in diameter) for all altitudinal classes. Seeds were air-dried and stored in the laboratory at ambient conditions until the start of the experiments. We estimated seed mass by determining the mass of 250 dry seeds per population with a precision balance (0.1 mg accuracy). Seeds were weighed in 25 groups of 10 seeds because of their low mass.

Seeds were germinated in Petri dishes in a germination chamber under controlled light (12 h light – 12 h dark daily photoperiod of about  $38 \mu\text{mol}\cdot\text{m}^{-2}\cdot\text{s}^{-1}$  cool white fluorescent light tubes) and temperature ( $25^{\circ}C$ ) conditions. Thirty days after germination (considered as radicle protrusion), seven seedlings per altitude class were measured, following the procedure provided in Sosa Pivatto et al. (2014). Digital photographs were taken for each seedling, and seedling height and width were measured using BIO7 software. With these data, we calculated a shape index (height/width ratio), with a value of 1 indicating spherical — globose — seedlings, and a value departing from 1 indicating that seedlings become cylindrical — the typical shape of columnar seedlings (Sosa Pivatto et al. 2014).

### Statistical analyses

We calculated the Pearson correlation coefficient to analyze the relationship between altitude, seed mass, and the different mea-

**Fig. 2.** Relationships between altitude and seed mass (a), seedling height (b), seedling width (d), and shape index (f), and between seed mass and seedling height (c), seedling width (e), and shape index (g) in *Gymnocalycium monvillei*. Pearson correlations: \*,  $p \leq 0.05$ ; ns, nonsignificant.



**Fig. 3.** Photographs of typical seedlings of *Gymnocalycium monvillei* for each altitudinal class.



sures of seedling size (height, width, and shape index), using the stats packages in R software version 2.15.1 (R Development Core Team 2012). In all cases, the data met the assumptions of normality and homoscedasticity.

## Results

Seed mass was not related to altitude or to any of the other variables (Fig. 2). Seedling height increased significantly with altitude (Fig. 2). Shape index increased significantly with altitude (i.e.,

seedlings became more columnar with increasing altitude). Seed mass was not related to any variable. Shape index was positively correlated to seedling height and negatively correlated to seedling width (correlation coefficient of 0.97 and 0.92,  $p < 0.05$ , respectively). Changes in seedling shape along the gradient are clearly observed in photographs in Fig. 3.

## Discussion

Contrary to our expectations, we did not find a relationship between seed size and altitude. Although studies performed at the interspecific level reported a clear pattern between latitude, or altitude, and seed size (Baker 1972; Murray et al. 2003; Moles et al. 2007), studies that had analyzed these patterns at the intraspecific level found contrasting results. For example, in the Swiss Alps, Pluess et al. (2005) found different seed size increases with altitude in related species but not among populations of a single species. However, in Himalayan species of the genus *Pedicularis*, Guo et al. (2010) found a decrease in seed size with altitude, both at intra- and inter-specific levels; the authors attributed those differences not to altitude per se, but to other variables, such as plant size and seed number per fruit. Although in *G. monvillei*, altitude is not a determinant of seed size, at least directly, we cannot discount the possibility that altitude in combination with other factors (e.g., humidity, soil characteristics or even biotic interactions) has an influence on this variable.

In contrast with Sosa Pivatto et al. (2014), we did not find a relationship between seed size and seedling traits. These discrepancies may be due to the lower variation in seed mass at the intraspecific level than at the interspecific level. Although seed size did not vary with altitude, seedling height and shape index showed a very clear altitude-related pattern. Seedling height increased with increasing altitude. Sosa Pivatto et al. (2014) stated that a more columnar seedling would be a better competitor for light, but less tolerant to higher temperatures or drought. Our results show that this trade-off could also operate in seedlings of similar seed sizes in response to marked differences in climate. In the present study, at higher altitudes, in a much cooler and wetter environment, seedlings were found to be columnar, which could be an advantage for light competition (Leishman et al. 2000). However, at the other extreme of the gradient, a globose morphology would make seedlings more resistant to a hotter and drier environment. In other words, factors associated with altitude (temperature and humidity) would be exerting a strong selective pressure on seedling traits, altering seedling morphology. Interestingly, low temperatures at the higher sites, where frosts occur year round, did not constrain seedling shape. Physiological, rather than morphological mechanisms are likely operating in protecting seedlings from low temperatures (Nobel 1982). For example, water loss before winter plays an important role in preventing frost damage in *Opuntia* species (Ishikawa and Gusta 1996).

Climate change is affecting organisms worldwide, and understanding species–environment relationships would help to predict species responses (Gurvich et al. 2002; Aragón-Gastélum et al. 2014). It could be accepted that species with wide distributions along climatic gradients, such as *G. monvillei*, would be less affected by climate change than narrowly distributed species (Ureta et al. 2012). However, our results suggest that *G. monvillei* populations could be negatively affected by changes in temperature if seedling morphology of the different altitudinal classes remained unchanged. In addition, dispersion in this species is mediated by ants, which are short-distance dispersers (Bregman 1988; Lengyel et al. 2010), and therefore movement of seeds among different populations is not a common occurrence. Cactus diversity across the Americas is concentrated in mountain environments (Mourelle and Ezcurra 1996; Ortega-Baes and Godínez-Alvarez 2006); therefore, understanding how species are distributed and adapted to these environments is very important to predict their responses to environmental changes.

## Acknowledgements

This work was supported by grants from Consejo Nacional de Investigaciones Científicas y Técnicas (CONICET) (PIP 11220110100873) and the Research Committee of the Cactus and Succulent Society of America. D.E. Gurvich and M.L. Las Peñas are researchers at CONICET. Two anonymous reviewers made useful comments on the manuscript and Jorgelina Brasca assisted with the English version of the manuscript.

## References

- Aragón-Gastélum, J.L., Flores, J., Yáñez-Espinoza, L., Badano, E., Ramírez-Tobías, H.M., Rodas-Ortíz, J.P., and González-Salvatierra, C. 2014. Induced climate change impairs photosynthetic performance in *Echinocactus platyacanthus*, an especially protected Mexican cactus species. *Flora*, **209**: 499–503. doi:[10.1016/j.flora.2014.06.002](https://doi.org/10.1016/j.flora.2014.06.002).
- Baker, H.G. 1972. Seed weight in relation to environmental conditions in California. *Ecology*, **53**: 997–1010. doi:[10.2307/1935413](https://doi.org/10.2307/1935413).
- Bregman, R. 1988. Forms of seed dispersal in Cactaceae. *Acta Bot. Neerl.* **37**: 395–402. doi:[10.1111/j.1438-8677.1988.tb02148.x](https://doi.org/10.1111/j.1438-8677.1988.tb02148.x).
- Butler, C.J., Wheeler, E.A., and Stuble, L.B. 2012. Distribution of the threatened lace hedgehog cactus (*Echinocereus reichenbachii*) under various climate change scenarios. *J. Torrey Bot. Soc.* **139**: 46–55. doi:[10.3159/TORREY-D-11-00049.1](https://doi.org/10.3159/TORREY-D-11-00049.1).
- Charles, G. 2009. *Gymnocalycium* in habitat and culture. Charles, Bank, Bridge, Stamford, England.
- Cornejo, D., and Simpson, B. 1997. Analysis of form and function in North American columnar cacti (tribe Pachycereeae). *Am. J. Bot.* **84**: 1482–1501. doi:[10.2307/2446610](https://doi.org/10.2307/2446610). PMID:[21708554](https://pubmed.ncbi.nlm.nih.gov/930554/).
- Dávila, P., Téllez, O., and Lira, R. 2013. Impact of climate change on the distribution of populations of an endemic Mexican columnar cactus in the Tehuacán-Cuicatlán Valley, Mexico. *Plant Biosyst.* **147**: 376–386. doi:[10.1080/11263504.2012.749955](https://doi.org/10.1080/11263504.2012.749955).
- de Fina, A.L. 1992. Aptitud agroclimática de la República Argentina. Academia Nacional de Agronomía y Veterinaria, Buenos Aires.
- Demaio, P.H., Barfuss, M.H., Kiesling, R., Till, W., and Chiapella, J.O. 2011. Molecular phylogeny of *Gymnocalycium* (Cactaceae): assessment of alternative infrageneric systems, a new subgenus, and trends in the evolution of the genus. *Am. J. Bot.* **98**: 1841–1854. doi:[10.3732/ajb.1100054](https://doi.org/10.3732/ajb.1100054). PMID:[22012926](https://pubmed.ncbi.nlm.nih.gov/22012926/).
- Felger, R.S., and Lowe, C.H. 1967. Clinal variation in the surface-volume relationships of the columnar cactus *Lophocereus schottii* in northwestern Mexico. *Ecology*, **48**: 530–536. doi:[10.2307/1936495](https://doi.org/10.2307/1936495).
- García-Pérez, J.F., Aguirre-Calderón, O.A., Estrada-Castillón, E., Flores-Rivas, J., Jiménez-Pérez, J., and Jurado, E. 2007. Germinación y establecimiento de plantas nativas del matorral tamaulipeco y una especie introducida en un gradiente de elevación. *Madera Bosques*, **13**: 99–117.
- Giorgis, M.A., Cingolani, A.M., Chiarini, F., Chiapella, J., Barboza, G., Espinar, L.A., Morero, R., Gurvich, D.E., Tecco, P.A., Subils, R., and Cabido, M. 2011. Composición florística del Bosque Chaquen serrano de la provincia de Córdoba, Argentina. *Kurtziana*, **36**: 9–43.
- Guo, H., Mazer, S.J., and Du, G. 2010. Geographic variation in seed mass within and among nine species of *Pedicularis* (Orobanchaceae): effects of elevation, plant size and seed number per fruit. *J. Ecol.* **98**: 1232–1242. doi:[10.1111/j.1365-2745.2010.01688.x](https://doi.org/10.1111/j.1365-2745.2010.01688.x).
- Gurvich, D.E., Díaz, S., Falczuk, V., Pérez-Harguindeguy, N., Cabido, M., and Thorpe, C. 2002. Foliar resistance to simulated extreme temperature events in contrasting plant functional and chorological types. *Global Change Biology*, **8**: 1139–1145. doi:[10.1046/j.1365-2486.2002.00540.x](https://doi.org/10.1046/j.1365-2486.2002.00540.x).
- Gurvich, D.E., Zeballos, S.R., and Demaio, P.H. 2014. Diversity and composition of cactus species along an altitudinal gradient in the Sierras del Norte Mountains (Córdoba, Argentina). *S. Afr. J. Bot.* **93**: 142–147. doi:[10.1016/j.sajb.2014.03.018](https://doi.org/10.1016/j.sajb.2014.03.018).
- Ishikawa, M., and Gusta, L.V. 1996. Freezing and heat tolerance of *Opuntia* cacti native to the Canadian prairie provinces. *Can. J. Bot.* **74**(12): 1890–1895. doi:[10.1139/b96-226](https://doi.org/10.1139/b96-226).
- Leishman, M.R., Wright, I.J., Moles, A.T., and Westoby, M. 2000. The evolutionary ecology of seed size. In *Seeds: the ecology of regeneration in plant communities*. Edited by M. Fenner. CABI Publishing, Wallingford. pp. 31–57.
- Lengyel, S., Gove, A.D., Latimer, A.M., Majer, J.D., and Dunn, R.R. 2010. Convergent evolution of seed dispersal by ants, and phylogeny and biogeography in flowering plants: a global survey. *Perspect. Plant Ecol. Evol. Syst.* **12**: 43–55. doi:[10.1016/j.ppees.2009.08.001](https://doi.org/10.1016/j.ppees.2009.08.001).
- Martorell, C., Vega, E., and Ezcurra, E. 2006. Morphological consequences of the trade-off between growth and reproduction in a columnar cactus (*Lophocereus schottii*). *Plant Ecol.* **183**: 125–131. doi:[10.1007/s11258-005-9011-5](https://doi.org/10.1007/s11258-005-9011-5).
- Moles, A.T., and Westoby, M. 2004. Seedling survival and seed size: a synthesis of the literature. *J. Ecol.* **92**: 372–383. doi:[10.1111/j.0022-0477.2004.00884.x](https://doi.org/10.1111/j.0022-0477.2004.00884.x).
- Moles, A.T., Ackerly, S.D., Tweddle, J.C., Dickie, J.B., Smith, R., Leishman, M.R., Mayfield, M.M., Pitman, A., Wood, J.T., and Westoby, M. 2007. Global patterns in seed size. *Global Ecol. Biogeogr.* **16**: 109–116. doi:[10.1111/j.1466-8238.2006.00259.x](https://doi.org/10.1111/j.1466-8238.2006.00259.x).
- Mourelle, C., and Ezcurra, E. 1996. Species richness of Argentine cacti: a test of biogeographic hypotheses. *J. Veg. Sci.* **7**: 667–680. doi:[10.2307/3236378](https://doi.org/10.2307/3236378).

- Murray, B.R., Brown, A.H.D., and Grace, J.P. 2003. Geographic gradients in seed size among and within perennial Australian *Glycine* species. *Aust. J. Bot.* **51**: 47–56. doi:[10.1071/BT02069](https://doi.org/10.1071/BT02069).
- Nobel, P.S. 1982. Low-temperature tolerance and cold hardening of cacti. *Ecology*, **63**: 1650–1656. doi:[10.2307/1940106](https://doi.org/10.2307/1940106).
- Nobel, P.S. 2003. Environmental biology of agaves and cacti. Cambridge University Press.
- Ortega-Baes, P., and Godínez-Alvarez, H. 2006. Global diversity and conservation priorities in the Cactaceae. *Biodivers. Conserv.* **15**: 817–827. doi:[10.1007/s10531-004-1461-x](https://doi.org/10.1007/s10531-004-1461-x).
- Pluess, A.R., Schütz, W., and Stöcklin, J. 2005. Seed weight increases with altitude in the Swiss Alps between related species but not among populations of individual species. *Oecologia*, **144**: 55–61. doi:[10.1007/s00442-005-0047-y](https://doi.org/10.1007/s00442-005-0047-y). PMID:15800741.
- R Development Core Team. 2012. R: a language and environment for statistical computing. R Foundation for Statistical Computing, Vienna, Austria. <http://www.R-project.org>.
- Sosa Pivatto, M., Funes, G., Ferreras, A.E., and Gurvich, D.E. 2014. Seed mass, germination and seedling traits for some central Argentinian cacti. *Seed Sci. Res.* **24**: 71–77. doi:[10.1017/S0960258513000366](https://doi.org/10.1017/S0960258513000366).
- Ureta, C., Martorell, C., Hortal, J., and Fornoni, J. 2012. Assessing extinction risks under the combined effects of climate change and human disturbance through the analysis of life-history plasticity. *Perspect. Plant Ecol. Evol. Syst.* **14**: 393–401. doi:[10.1016/j.ppees.2012.09.001](https://doi.org/10.1016/j.ppees.2012.09.001).
- Wulff, R.D. 1986. Seed size variation in *Desmodium paniculatum*: II. Effects on seedling growth and physiological performance. *J. Ecol.* **74**: 99–114. doi:[10.2307/2260351](https://doi.org/10.2307/2260351).