Seed dormancy of *Ochradenus baccatus* (Resedaceae), a shrubby species from Arabian desert regions

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Abstract: *Ochradenus baccatus* is a perennial glycophyte growing in Middle East and it is one of the most important food sources for many animal species in desert regions. The aim of our study was to investigate the effects of seed storage, light, temperature and gibberellic acid (GA₃) on germination of *O. baccatus* seeds. We also investigated the germination characteristics of *O. baccatus* seeds under different saline concentration and their capability to recover germination once they were transferred to distilled water. Seeds were stored at room temperature (20 ± 2 °C) and at -18 °C. Germination tests were conducted at alternating temperatures of 15/25, 20/30 and 25/35 °C in either continuous darkness or photoperiod of 12-h dark/12-h light. To study the effect of GA₃ on germination of *O. baccatus* seeds, freshly-collected seeds and stored seeds were soaked for 24 h in a GA₃ water solution (1 g/L) before sowing. To assess the salinity tolerance during germination, seeds were germinated under different salinity levels (100, 200 and 400 mM NaCl). Stored seeds at room temperature and -18 °C germinate equally well at different temperature regimes and light conditions. However, freshly matured seeds were not able to germinate even when they were treated with GA₃. On the contrary, stored seeds at room temperature and -18 °C treated with GA₃ increase the final germination percentages. These results indicated that *O. baccatus* seeds have physiological dormancy and they need to be stored in order to break their dormancy. In the present study, one year of storage did not show a significant variation in germination between the two storage conditions assayed. Therefore, further research is needed to know about the maximum storage period for *O. baccatus* seeds under different storage conditions. Very few *O. baccatus* seeds (less than 5 %) germinated at the tested lowest concentration of NaCl. However, ungerminated seeds were able to germinate when salinity stress was alleviated. In conclusion, *O. baccatus* seeds have physiological dormancy, and seed storage (at room temperature and at -18 °C) for one year is effective for breaking this dormancy. In addition, *O. baccatus* seeds present ability to remain viable in saline conditions and they will be able to germinate once the salinity level decrease. Rev. Biol. Trop. 64 (3): 965-974. Epub 2016 September 01.

Key words: seed storage, seed germination, temperature, salinity, desert species, physiological dormancy.

*Ochradenus baccatus* Delile (Resedaceae) is a perennial shrub that grows on sandy and stony habitats in Middle East (Al-Fredan, 2010). The fleshy fruits of *O. baccatus* contain high water and sugar content, are consumed and dispersed by various frugivores, and is reported to be one of the most important food sources for many animal species in the Arabian desert (Bronstein et al., 2007; Spiegel & Nathan, 2007; 2011). The green stems of *O. baccatus* helps leaves in conducting photosynthesis, especially when leaves are weakened after severe drought environment (Abdulfatah, 2004). This species is widely distributed and is found in Ethiopia, Egypt, Libya, Middle East, Iran and extends into Pakistan (Al-Fredan, 2010; Abd El-Wahab, Zaghloul, Kamel, & Moustafa, 2008; Khan, Al-Qurainy, & Nadeem, 2012). The medicinal uses of *O. baccatus* have been well documented (Nawash & Al-Horani,
2011; Alqasoumi, Soliman, Awaad, & Donia, 2012); it is also useful in controlling root-knot nematodes, and can be planted as cover plant/green manure (Oka, Shuker, Tkachi, Trabelcy, & Gerchman, 2013). Furthermore, due to extreme drought tolerance ability, *O. baccatus* has the potential to be used in urban landscaping (Sudershan, Abo El-Nil, & Hussain, 2003). *O. baccatus* have also been categorized among the native plant species that could be exploited for landscaping (Global Sustainability Assessment System, 2014).

Seeds of *O. arabicus* exhibit dormancy and have hard seed coats, presence of some inhibitors, low internal hormone content or underdeveloped embryos (Nadeem, Al-Qurainy, Khan, Tarroum, & Ashraf, 2012). Contrary to this, Suleiman, Bhat, Jacob and Thomas (2011) have shown that *O. baccatus* seeds have high germination without any pretreatment. This result suggests that seeds of this species have not dormancy. In the present study, we investigated if *O. baccatus* does really display some type of seed dormancy. The inability of seeds to germinate in darkness is a trait generally associated with the ability to form a soil seed bank especially for small seeded species from arid regions (Pons, 1991; Kigel, 1995). To study the effect of light on seed germination of *O. baccatus* we tested germination in continuous darkness or 12 hr light/12 hr dark photoperiod. Besides, *O. baccatus* seeds behave like orthodox (http://data.kew.org/sid/sidsearch.html). Considering the multiple values of the species and its tolerance to seed desiccation, *ex situ* conservation is an appropriate choice to contribute to the long-term conservation of *O. baccatus* seeds. *O. baccatus* grows on dry soils and colonise arid environments where saline soils are common. Therefore, it is of great interest to study the effects of salt stress on seed germination of *O. baccatus*. It has been shown in a large number of species that an increase in salt concentration usually delays and reduces seed germination (El-Keblawy & Al-Shamsi, 2008; Guma, Padrón-Mederos, Santos-Guerra, & Reyes-Betancort, 2010). However, there is a wide range of variability in salt tolerance among species (Khan & Gulzar, 2003). Moreover, many seeds that are unable to germinate at high saline concentrations might recover the ability to germinate when salinity levels decrease (Zia & Khan, 2008). The main aims of our study were to investigate: (i) the germination characteristics of fresh and stored seeds of *O. baccatus* at different temperature and light regimes; (ii) the effect of GA<sub>3</sub> on the germination of *O. baccatus* seeds, and (iii) the salinity tolerance of *O. baccatus* seeds during germination.

**MATERIALS AND METHODS**

**Plant material:** *Ochradenus baccatus* reaches up to two meter and grows as bushes (Bronstein et al., 2007). The species produces yellow flowers, followed by whitish berry containing black seeds (Omar, Al-Mutawa, & Zaman, 2007). The species mainly flowers from December to March in Middle-East (Zohary, 1966). However, Wolfe and Burns (2001) showed that large plants of *O. baccatus* have the ability to flower continuously whereas, smaller plants are reproductive only during the winter (Wolfe & Burns, 2001). Furthermore, *O. baccatus* is a gynodioecious species, where female individuals are constant in sex expression; however, male individuals exhibit great variation in functional gender (Amer & Hassan, 2015).

**Seed collection and seed storage:** Fruits of *O. baccatus* were collected during last week of March 2013 from Shahniya Dukan Road, Doha, Qatar (25° 28’ 90” N - 51° 48’ 89” E). The climate of Qatar is hot during summer (April to September), with daytime temperatures as high as (50 °C). Precipitation is scarce (less than 152 mm annually), and mostly occurs in winter between November to March (Persian Gulf, 2012). The pulp was removed manually by mashing with hand gloves, and seeds were obtained and stored both at room temperature (20 ± 2 °C) and in a freezer at -18 °C. Freshly
collected seeds were also immediately tested for germination. However, seeds stored at room temperature as well as at -18 °C were retrieved separately after one year and tested for germination. Three replicates of 50 seeds each were used to determine the mean seed mass.

**Water uptake during seed imbition:** Water uptake capacity was determined during imbition of freshly collected seeds, seeds stored at room temperature and seeds stored at -18 °C, three replications of 50 seeds each were weighed and then reweighed after imbition in distilled water for 12 and 24 h. Percentage of water uptake (mean ± standard deviation) was calculated as the amount of water taken up relative to initial seed fresh mass (Baskin, Zackrisson, & Baskin, 2002).

**Seed germination trials:** To assess the effect of storage temperature and time on germination of *O. baccatus*, germination tests were conducted in incubators (LMS incubators, UK) set at different temperatures 15/25, 20/30 and 25/35 °C in either continuous darkness or photoperiod of 12-h dark/12-h light (the highest temperature for 12 h in light and the lowest one for 12 h in dark) (Vargas-Figueroa, Duque-Palacio, & Torres-González, 2015). Incubators were illuminated with a 25-W white fluorescent lamp. The germination was conducted in 9-cm tight-fitting Petri-dishes containing one disk of Whatman No. 1 filter paper, moistened with 10mL of distilled water. During the dark treatment, the dishes were wrapped in aluminum foil to prevent any exposure to light. Four replicates of 25 seeds each were used for each treatment. The number of germinated seeds were counted and removed every alternate day for 20 days. A seed was considered to have germinated when the emerging radicle elongated to 2 mm. Seeds incubated under dark conditions were counted after a period of 20 days.

**Effect of seed soaking in gibberellic acid on germination:** Lots of 100 freshly-collected and stored seeds were soaked for 24 h at room temperature in a gibberellic acid (Sigma- Aldrich, UK) solution (GA3, 1 g/L) before sowing. After this treatment, pre-treated seeds were tested for germination at alternating temperatures of 20/30 °C under 12-h dark/12-h light photoperiod (the highest temperature for 12 h in light and the lowest one for 12 h in dark) and continuous darkness. Untreated seeds were sown in the same conditions and used as a control.

**Effect of salinity on seed germination:** To assess the salinity tolerance during germination, seeds stored at room temperature as well as at -18 °C were germinated separately under different salinity levels (100, 200 and 400 mM NaCl). Four replicates, each of 25 seeds, were used for each treatment. Seeds were germinated in 9-cm diameter Petri dishes and two layers of Whatman No.1 filter paper, moistened with 10 mL of salt solution. Petri dishes were sealed with parafilm to minimize evaporation and incubated at 20/30°C in 12-h dark/12-h light photoperiod (the highest temperature for 12 h in light and the lowest one for 12 h in dark) and continuous darkness. Seed germination was recorded as above. Seeds sown under the same conditions but in distilled water were used as control seeds.

**Recovery percentage:** Non-germinated seeds from the 20-day NaCl incubation tests were transferred to distilled water, and incubated for additional 20 days at the same incubation temperature. Recovery percentage (RP) was calculated by the following formula: RP = (a-b/c-b)x100, where a is the number of seeds germinated in NaCl solution after a 20-d period, plus those that recovered to germination in distilled water after another 20-d period, b is the number of seeds germinated in NaCl solution after a 20-day period, and c is the total number of seeds tested (Yang, Dong, & Huang, 2010). Initial germination was recorded as (b/c)x100, and final germination as (a/c)x100. For all experiments, final germination percentage (mean value ± standard deviation) was calculated. The values of final germination percentages were arcsine square-root transformed.
and then subjected to analysis of variance (ANOVA) using SPSS (untransformed data appear in Tables). In all germination trials, the number of empty seeds in each replicate was always excluded when calculating the final germination percentage. Moreover, for seeds soaked in GA₃, mean germination time (MGT, mean value in days ± standard deviation) was calculated. This parameter was determined according to the following formula: MGT = ΣDN/ΣN; where D is the number of days counted from the date of sowing and N is the number of seeds germinated on day D (Ellis & Roberts, 1981). The effect of incubation temperatures (three levels), seed storage conditions (three levels) and light conditions (two levels) on the final germination percentage was analyzed by a three-way factorial ANOVA. Similarly, the effect of soaking in GA₃ (soaked and not soaked seeds in GA₃; two levels), light conditions (two levels) and storage conditions (three levels) on the final germination percentage, was analyzed by a three-way factorial ANOVA. The effect of soaking in GA₃ (two levels) and seed storage conditions (two levels) on MGT was analyzed by a two-way factorial ANOVA. The effect of seed incubation in NaCl solutions of different concentration (four levels) and seed storage conditions (two levels) on the initial and final germination percentage was analyzed by a two-way factorial ANOVA. Where ANOVA indicated a significant effect, a comparison of mean values was carried out through the least significant difference test at 0.05 level of probability.

RESULTS

Seed water uptake: The mean mass for a lot of 50 seeds was 42.33 ± 0.72 mg for freshly collected seeds, 44.33 ± 0.27 mg for seeds stored at room temperature, and 41.00 ± 0.82 mg for seeds stored at -18 ºC. *O. baccatus* seeds imbibed water quickly and after 12h of imbibition in distilled water. Seed mass increased by 11.93 ± 3.07 % for freshly collected seeds, 19.57 ± 2.26 % for seeds stored at room temperature, and 21.20 ± 3.69 % for seeds stored at -18 ºC. After 24h of imbibition, seed mass increase was 16.63 ± 2.14 %, 23.30 ± 2.58 %, and 22.80 ± 3.05 %, respectively. Thus, in *O. baccatus* the seed coat does not prevent absorption of water.

Seed germination trials: Light condition and incubation temperature were not significant (P > 0.05). None of the freshly matured seeds of *O. baccatus* germinated at all temperature regimes and light conditions assayed (Table 1). Germination of stored seeds ranged from 35-54 % for seeds stored at -18 ºC, and from 41 % to 56 % for after-ripening in dry storage independently of light treatment. Seed storage had a highly significant (P< 0.001) effect on seed germination of *O. baccatus*. Considering the two-way interactions, only the interaction between temperature and seed storage condition was significant (P = 0.038), while the three-way interaction was not significant (P= 0.503). The final germination percentage of fresh seeds were significantly lower (P<

<table>
<thead>
<tr>
<th>Temperature (ºC)</th>
<th>Germination (% ± SD) by storage conditions</th>
<th>RT</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Fresh seeds</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Light</td>
<td>Dark</td>
</tr>
<tr>
<td>15/25</td>
<td>0 a</td>
<td>0 a</td>
</tr>
<tr>
<td>20/30</td>
<td>0 a</td>
<td>0 a</td>
</tr>
<tr>
<td>25/35</td>
<td>0 a</td>
<td>0 a</td>
</tr>
</tbody>
</table>

Means within a row followed by the same letter are not significantly different from each other (LSD test, P > 0.05). Seeds were stored at -18 ºC or at room temperature (RT).
0.05) than that of stored seeds. There were not significant differences ($P > 0.05$) between the final germination percentages reached by seeds stored at room temperature and at -18 °C.

**Effect seed soaking in $GA_3$ on germination:** The final germination percentages of *O. baccatus* seeds soaked for 24 h in a $GA_3$ solution (1 g/L) were always higher than these of control seeds (Table 2). However, only in seeds incubated under constant darkness were found significant differences for seeds stored at room temperature and seeds stored at -18 °C. The highest germination percentage reached by stored seeds soaked in $GA_3$ was 84%. The three-way ANOVA showed a significant effect ($P < 0.001$) of soaking in $GA_3$ and seed storage conditions ($P < 0.001$) on final germination, whereas the interaction between both factors was not significant ($P = 0.163$). Light condition had no significant effect ($P = 0.274$) on seed germination. The interaction between light and seed storage condition, and between $GA_3$ and storage condition were not significant ($P = 0.057$ and $P = 0.163$, respectively). The three-way interaction was significant ($P = 0.007$). Germination speed (as expressed by MGT) was higher for seed soaked in $GA_3$ than for untreated seeds (2.73 vs. 3.21 days), but these differences were only significant ($P < 0.05$) for seeds stored at room temperature (Table 2). Seed storage condition was not significant ($P = 0.995$), whereas soaking in $GA_3$ was significant ($P = 0.029$). The interaction between seed storage condition and soaking in $GA_3$ was not significant ($P = 0.874$).

**Effect of salinity on seed germination:** Germination of *O. baccatus* seeds ranged from 0 to 3% for the different NaCl concentrations assayed (Table 3). However, the seeds were able to germinate once they were transferred to distilled water. Recovery percentages (RP) ranged from 36 to 68%. The highest RP values were reached for 100 mM NaCl (52% for seeds stored at -18 °C and 68% for seed stored at room temperature). Seed storage condition and NaCl concentration were significant ($P = 0.021$ and $P < 0.001$, respectively) for initial germination. However, the interaction between seed storage condition and seed incubation in NaCl was not significant ($P = 0.135$). Seed

### TABLE 2

Effect of soaking in a gibberellic acid solution (GA$_3$, 1 g/L) on the final germination percentages (mean values ± standard deviation) and mean germination time (MGT, mean in days ± standard deviation) of fresh and stored seeds of *Ochradenus baccatus* Delile at 20/30 °C under two light conditions (12-h light photoperiod and continuous darkness)

<table>
<thead>
<tr>
<th>Storage condition</th>
<th>Germination (% ± SD)</th>
<th>MGT (days ± SD)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Without $GA_3$</td>
<td>With $GA_3$ P</td>
</tr>
<tr>
<td>Fresh seeds</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Light</td>
<td>0 a</td>
<td>9 ± 9.12 a ns</td>
</tr>
<tr>
<td>Dark</td>
<td>0 a</td>
<td>2 ± 3.46 a ns</td>
</tr>
<tr>
<td>-18 °C</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Light</td>
<td>45 ± 9.10 b</td>
<td>59 ± 7.14 b ns</td>
</tr>
<tr>
<td>Dark</td>
<td>40 ± 12.32 b</td>
<td>84 ± 8.94 b **</td>
</tr>
<tr>
<td>RT</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Light</td>
<td>56 ± 4.90 b</td>
<td>67 ± 1.14 b ns</td>
</tr>
<tr>
<td>Dark</td>
<td>52 ± 6.32 b</td>
<td>84 ± 11.66 b *</td>
</tr>
</tbody>
</table>

Mean values in a column followed by the same letter are not significantly different from each other (LSD test, $P > 0.05$). MGT was not calculated (NC) when germination was equal to or less than 10% and for trials in continuous darkness because the number of germinated seeds was only counted at the end of germination period. $P$: For each seed storage condition and light condition, significance level between the final germination percentages reached by seeds soaked and non-soaked in $GA_3$. **$P < 0.001$; *$P < 0.05$; ns, not significant.**

Seeds were stored at -18 °C or at room temperature (RT).
storage condition and NaCl concentration were significant (P= 0.001 and P< 0.001, respectively) for final germination, and the interaction was not significant (P= 0.170).

**DISCUSSION**

Fresh matured seeds of *O. baccatus* (which have a fully developed embryo) are dormant at maturity and are unable to germinate under different temperature and light conditions. However, the seed coat of *O. baccatus* seeds (fresh as well as stored) is permeable to water as seeds imbibed water quickly after 24 h. Therefore, these seeds do not exhibit physical dormancy, according to the classification system of Baskin and Baskin (2004), where physical dormancy is defined as the result of a water-impermeable layer in the seed or fruit. In addition, *O. baccatus* seeds reached germination percentages up to 56 % and GA3 improved germination depending on temperature and light conditions.

Based on the results, we concluded that these seeds have physiological dormancy. Moreover, the breaking of seed dormancy during dry storage (after-ripening) could indicate that the level of physiological dormancy is non-deep (Baskin & Baskin, 2004). Our results are in contradiction with those obtained by Suleiman, Bhat, Jacob and Thomas (2011) for this same species. These authors found that *O. baccatus* seeds reached high germination percentages without any pretreatment. Probably, the species showed inter-population variation in seed dormancy, and/or variability in germination capacity, depending on the time of seed collection.

The presence of physiological dormancy in freshly matured seeds might allow them time for dispersal and prevents immediate germination at the time of seed release (Kucera, Cohn, & Leubner-Metzger, 2005; Finch-Savage & Leubner-Metzger, 2006). The seeds of *O. baccatus* matured in late March, when temperature increases and the chances of rainfall are very low. This suggests that seeds enter into the soil seed bank and remain dormant during unsuitable ecological conditions, when the probability of seedling survival is low (Black, Bewley, & Halmer, 2006; Soares-Oliveira, Cleiton-José, Monteiro-Ribeiro, & Rocha-Faria, 2015), as chances of rainfall during summer are minimal (Böer, 1997). Furthermore, the seeds might remain dormant in natural conditions until the winter (November to March). This strategy ensures that germination is unlikely to take place under natural environmental conditions until November (before the onset of winter rain), when conditions for seedling establishment are more favorable. These findings are supported by Wolfe and Burns (2001), who showed that seeds of *O. baccatus* produced in winter have higher germination percentage due to favorable environmental conditions for germination and seedling establishment. In the present study, *O. baccatus* seeds germination was as much as 56 % after one year of dry storage.

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**TABLE 3**

Germination percentages (mean values ± standard deviation) of *Ochradenus baccatus* Delile seeds stored at -18 °C or at room temperature (RT) after incubation in different NaCl concentrations for 20 days at 20/30 °C under 12-h light photoperiod (initial germination), and germination percentages when non-germinated seeds were incubated for another 20 days in distilled water (final germination)

<table>
<thead>
<tr>
<th>NaCl concentration (mM)</th>
<th>Initial germination</th>
<th>Final germination</th>
<th>RP</th>
<th>Initial germination</th>
<th>Final germination</th>
<th>RP</th>
</tr>
</thead>
<tbody>
<tr>
<td>-18°C</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>0 (control)</td>
<td>45 ± 9.10 b</td>
<td>45 ± 9.10 a</td>
<td>–</td>
<td>56 ± 4.90 b</td>
<td>56 ± 4.90 ab</td>
<td>–</td>
</tr>
<tr>
<td>100</td>
<td>0 a</td>
<td>52 ± 8.94 a</td>
<td>3 ± 3.32 a</td>
<td>69 ± 10.34 b</td>
<td>68 ± 11.74 b</td>
<td></td>
</tr>
<tr>
<td>200</td>
<td>0 a</td>
<td>37 ± 3.32 a</td>
<td>0 a</td>
<td>56 ± 7.48 ab</td>
<td>56 ± 7.48 ab</td>
<td></td>
</tr>
<tr>
<td>400</td>
<td>0 a</td>
<td>36 ± 5.66 a</td>
<td>0 a</td>
<td>37 ± 5.20 a</td>
<td>37 ± 5.20 a</td>
<td></td>
</tr>
<tr>
<td>RT</td>
<td></td>
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</tr>
</tbody>
</table>

Means within a column followed by the same letters are not significantly different (LSD test, P > 0.05). Control: seeds incubated for 20 days in distilled water. RP: recovery percentage.
storage (after-ripening) indicating that they are tolerant to desiccation. The poor germination by freshly harvested seeds could also be interpreted as avoiding germination in summer desert conditions.

Seed stored at room temperature and -18 °C germinated equally well at the different temperatures assayed. This wide temperature range perhaps allows them to germinate in winter (November to February) in their natural habitat (monthly average temperature between November to February vary between 22.5 to 25.9 °C - Islam, Kubo, Ohadi, & Alili, 2009). On the basis of our results, we can confirm that *O. baccatus* seeds became physiologically dormant during maturation. Therefore, seed storage (at room temperature and at -18 °C) for one year worked more effectively in breaking the dormancy of *O. baccatus* seeds. Besides, there was no significant variation in seed germination between the two storage conditions tested. It has been indicated that viability of orthodox seeds could be maintained by drying them in low moisture content at relatively lower temperature (Nahuel-Morandini, Mabel-Giamminola, & De Viana, 2013), and their longevity can be extended with reductions in both moisture content and temperature over a wide range of storage environments (http://data.kew.org/sid/sidsearch.html). Our results showed that *O. baccatus* seeds could be stored without significant loss of viability. This was further confirmed by the requirements of after ripening before their germination. However, further investigations are required in order to know the maximum seed storage period for *O. baccatus* under different storage conditions.

Germination of freshly matured seeds treated with GA$_3$ is less than 10 %. However, application of GA$_3$ treatments significantly enhanced the germination percentage of stored seeds. We assumed that stored seeds might have reached a maximum release of dormancy after storage and therefore they showed better germination compared to freshly harvested seeds. These findings further support that freshly harvested seeds of *O. baccatus* have physiological dormancy and need after-ripening to remove the dormancy, because the seeds have endogenous dormancy, that might be associated with the physiological maturity of embryos that caused lower germination in freshly matured seeds, even with the application of GA$_3$ (Schütz, Milberg, & Lamont, 2002; Gao, Jordan, & Avele, 2012). The final germination percentages of stored seeds of *O. baccatus* treated with GA$_3$ were higher than the control seeds (untreated seeds). However, seeds incubated under constant darkness showed significantly higher germination as compared to light incubated seeds. As the seeds of *O. baccatus* are neutrally photoblastic, it might be possible that the application of GA$_3$ improved the germination of both light and dark incubated seeds. However, higher germination for dark incubated seeds could indicate the biosynthesis of gibberellins (mediated through the phytochrome system) may have been inhibited by light exposure (Kato-Noguchi, 2002).

Very few seeds (less than 5 %) were able to germinate at the lowest concentration of NaCl tested. However, once transferred to the distilled water, they were able to germinate. Although the germination recovery was higher for seeds exposed to the lowest concentration of NaCl. These results indicate that although, *O. baccatus* has been categorized as a glycophyte, seeds have ability to remain viable in saline conditions and are able to germinate once the salinity level decreases by rain. Similar results have been obtained for several glycophyte species (El-Keblawy, Al-Ansari, & Al-Shamsi, 2011). This strategy might help them to form a persistent soil seed bank even in saline habitats that can avoid the vulnerability of local extinction when vegetation on the ground is removed and would be important in restoration conservation of plants (Bakker, Poschlod, Strykstra, Bekker, & Thompson, 1996). Bajji, Kinet and Lutts (2002) reported that soil seed bank has the ability to remain quiescent at a high salt level, and to germinate immediately after salinity reduction. This is common for both halophytes and other species that colonize similar environments. Results of the present study indicate that seeds of the glycophytic *O.*
Ochradenus baccatus, like those of the halophyte species, recover their germination when transferred from saline solutions to distilled water.

In conclusion, seeds stored at room temperature and at -18 °C germinate equally well under different temperature regimes and light conditions. O. baccatus seeds have physiological dormancy and need to be stored in order to break their dormancy. Considering the tolerance to seed desiccation, ex situ conservation is an appropriate choice to contribute to the long-term seed conservation of O. baccatus. Seeds of the species are able to remain viable in saline conditions and they will be able to germinate once the salinity level decreases due to rainfall. O. baccatus colonize arid environments where saline soils are common; therefore, the species could be used in plant restoration programs and in revegetation projects in arid coastal areas.

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RESUMEN

Dormición de semillas de Ochradenus baccatus (Resedaceae), una especie arbustiva de las regiones desérticas arábigas. Ochradenus baccatus es un glicófito perenne que crece en oriente Medio y es una de las fuentes de alimentación más importantes para muchas especies animales de regiones desérticas. El objetivo de nuestro estudio fue investigar los efectos del almacenamiento de semillas, luz, temperatura y ácido giberélico (GA₃) en la germinación de semillas de O. baccatus. También se ha investigado la germinación de semillas de O. baccatus bajo diferentes concentraciones salinas y su capacidad para recuperar la germinación una vez que fueron transferidas a agua destilada. Las semillas se conservaron a temperatura ambiente (20 ± 2 °C) y a -18 °C. Los ensayos de germinación se realizaron a temperaturas alternas de 15/25, 20/30 y 25/35 °C bajo oscuridad continua o fotoperíodo de 12-h oscuridad/12-h luz. Para estudiar el efecto del GA₃ en la germinación de semillas de O. baccatus, semillas recién recolectadas y semillas almacenadas se sumergieron durante 24 h en una solución acuosa de GA₃ (1 g/L) antes de la siembra. Para evaluar la tolerancia a la salinidad durante la germinación, las semillas fueron germinadas bajo diferentes niveles de salinidad (100, 200 y 400 mM ClNa). Las semillas conservadas a temperatura ambiente y a -18 °C germinaron igualmente bien en los diferentes regímenes de temperatura y condiciones de iluminación. Sin embargo, las semillas recién maduras fueron incapaces de germinar incluso cuando se trataron con GA₃. Por el contrario, las semillas almacenadas tratadas con GA₃ incrementaron los porcentajes finales de germinación. Estos resultados indican que las semillas de O. baccatus tienen dormición fisiológica y necesitan ser almacenadas para romperla. En este estudio, un año de conservación no supuso una variación significativa en la germinación entre las dos condiciones de conservación ensayadas. Por lo tanto, se precisan investigaciones adicionales para conocer cuáles son los periodos máximos de almacenamiento de semillas de O. baccatus bajo diferentes condiciones de conservación. Muy pocas semillas de O. baccatus (menos del 5 %) germinaron a la concentración más baja de ClNa. Sin embargo, las semillas no germinadas fueron capaces de germinar cuando el estrés salino fue aliviado. En conclusión, las semillas de O. baccatus tienen dormición fisiológica y el almacenamiento de las mismas (a temperatura ambiente y a -18 °C) durante un año es eficaz para romper dicha dormición. Además, las semillas de O. baccatus presentan capacidad para permanecer viables en condiciones salinas y serán capaces de germinar una vez que el nivel de salinidad disminuya.

Palabras clave: almacenamiento de semillas, germinación de semillas, temperatura, salinidad, especies de desierto, dormición fisiológica.

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