Soaking curve and effect of temperature on the germination of daisy seeds

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ABSTRACT

The soaking curve and the effect of temperature on the germination of daisy seeds (Chrysanthemum leucanthemum) were characterized in this study. To determine the soaking curve, four samples of 0.5 g of seeds were soaked in germitest paper moistened with distilled water and maintained in germinators at 25°C. The seeds were weighed in periods of 0, 3, 6, 9, 12, 24, 36, 48, 60, 72, 84 and 96 hours using a precision digital balance of 0.0001 g. A triphasic pattern germination curve was adjusted, allowing the determination of the beginning and duration of the phase II of the germination process. The germination test was carried out with four replications of 50 seeds disposed in “Gerbox” boxes and placed in germinators at the temperatures of 20, 25, 30 or 20-30°C. A completely randomized experimental design was used with four replications of 50 seeds. The data were submitted to the analysis of variance and the averages were compared by the Tukey test, at 5% of probability. For analysis of accumulated seed germination, regressions were adjusted based on period of experiment. The seeds presented a triphasic pattern of germination and the phases I and II lasted 12 and 48 hours, respectively. The best temperature for the germination of the seeds is 25°C. The temperature of 30°C promoted the thermoinhibition of germination and increased the dead and dormant seeds and abnormal seedlings.

Keywords: Chrysanthemum leucanthemum, thermoinhibition, floriculture, ornamental.

RESUMO

Curva de embebição e efeito da temperatura na germinação de sementes de margarida

A curva de embebição e o efeito da temperatura sobre a germinação de sementes de margarida (Chrysanthemum leucanthemum) foram caracterizados. Para determinar a curva de embebição, quatro amostras de 0,5 g de sementes foram embebidas em papel germitest umedecido com água destilada e mantidas em germinador a 25°C. As sementes foram pesadas em períodos de 0, 3, 6, 9, 12, 24, 36, 48, 60, 72, 84 e 96 horas, com auxílio de balança de precisão digital de 0,0001 g. Foi ajustada uma curva trifásica padrão de germinação, permitindo a determinação do início e a duração da fase II do processo de germinação. O teste de germinação foi realizado com quatro repetições de 50 sementes dispostas em caixas “gerbox” mantidas em germinadores nas temperaturas de 20, 25, 30 ou 20-30°C. O delineamento experimental foi inteiramente casualizado. Os dados foram submetidos à análise de variância e as médias comparadas pelo teste de Tukey a 5% de probabilidade. Para análise da germinação acumulada das sementes foi feita regressão em função dos dias de avaliação. A embebição de sementes de margarida apresentou padrão trifásico de germinação e as fases I e II duraram 12 e 48 horas, respectivamente. A melhor temperatura para a germinação das sementes de margarida foi 25°C. A temperatura de 30°C promoveu termoinibição de germinação e aumentou a porcentagem de sementes mortas e dormentes e plântulas anormais.

Palavras-chave: Chrysanthemum leucanthemum, termoinibição, floricultura, ornamental.

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observe a triphasic pattern curve of soaking process (Bewley & Black, 1994). The determination of soaking curve and the definition of change phases during germination allow adjustments of methodologies for seed osmoconditioning, which are frequently used to homogenize the germination and emergence of seedlings in the field (Rodrigues et al., 2009). Besides, the determination of soaking phases allows to achieve adjustments in several tests of seed vigor used in the analysis of the physiological quality of seeds, such as water stress, and the tetrazolium test, to discriminate the vigor of seed lots and destine them to be traded (Marcos Filho, 1999; Rodrigues et al., 2008).

Temperature may influence the percentage of germination, affecting both the speed of water absorption and the biochemical reactions (Amato et al., 2007). Temperatures within optimum ranges promote the development of normal seedlings, but low temperatures lead to lower seedlings production, and the higher temperatures cause enzymatic disorders (Marcos Filho, 1999). High temperatures cause changes in the levels of phytohormones associated to the germination of seeds, mainly abscisic acid and gibberellin, by a process called thermoinhibition of germination (Akman, 2009). Thermoinhibition is mentioned as responsible for the high percentages of dormant seeds or formation of abnormal seedlings due to lower cell elongation (Toh et al., 2008).

The Rules for Seed Analysis (RAS) present specific methodologies of temperature and dormancy breaking for germination tests of several species of the genus Chrysanthemum, but there are no established conditions for the species C. leucanthemum (Brasil, 2009). Thus, the objective of this study was to characterize the soaking curve and to define the effect of temperature on the germination of daisy seeds.

**MATERIAL AND METHODS**

The present work was carried out in a laboratory of the Federal University of Viçosa (Minas Gerais state, Brazil). Seeds of the cultivar Gigante Branca were evaluated.

Initially, seed moisture content was determined by the official method of drying in kiln at 105±3°C for 24 hours, according to the Rules for Seed Analysis (Brasil, 2009). To determine the soaking curve, four replications of 0.5 g of seeds were placed in “Gerbox” boxes over two sheets of germitest paper moistened with distilled water at the proportion of 2.5 times their weight and maintained in germinator at 25°C. In the periods of 0, 3, 6, 9, 12, 24, 36, 48, 60, 72, 84 and 96 hours, the seeds were removed from the “Gerbox” boxes and carefully dried with towel paper for the removal of the excess of water adhered to the surface of the tegument and immediately weighed in a digital balance, with the precision of 0.0001 g. The weight gain of the seeds was calculated by the formula:

\[ GP = \frac{(PF-Pi)}{Pi} \times 100 \]

Where \( GP \) = weight gain (%); \( PF \) = final weight (weight in grams at each soaking period); \( Pi \) = initial weight of the seeds in grams, before soaking.

The germination test was carried out with four replications of 50 seeds disposed in “Gerbox” boxes on two germitest paper sheets moistened with distilled water at the proportion of 2.5 times their weight and maintained in a germinator at the temperature of 20, 25, 30 or 20-30°C. All seeds were germinated at 8 hours photoperiod under white light of fluorescent lamp, submitted to alternating temperatures, the highest temperature (30°C) was released during light period and the lowest (20°C), during the night period. The number of normal seedlings was counted in the first germination counting (FGC) carried out seven days after the beginning of the test. The second germination counting (SGC) was carried out 16 days after the beginning of the test, and the number of germinated seedlings was evaluated, accordingly to the Rules for Seed Analysis (Brasil, 2009). In the second germination counting, the number of normal and abnormal seedlings, dormant seeds (non-germinated) and dead seeds were also counted.

To determine the germination speed index (GSI), the number of germinated seeds was counted daily. The seeds with primary root protrusion with 1.0 mm of length were considered germinated to calculate the GSI. The GSI was calculated according to Maguire (1962), by the expression:

\[ GSI = \frac{(G1 / N1)+ (G2 / N2) +...+ (Gn / Nn)}{n} \]

Where \( G1 \) = number of germinated seeds in the first counting; \( N1 \) = number of days elapsed before the first counting; \( G2 \) = number of germinated seeds in the second counting; \( N2 \) = number of days elapsed before the second counting; \( n \) = last counting.

We established a curve that adjusted to a triphasic pattern of the germination that allowed the determination of the beginning and duration of the phase II of the germination process. A completely randomized experimental design was used for the germination test, with four replications and 50 seeds per replication. The data were submitted to the analysis of variance and the effects of the treatments were compared by the Tukey test, at 5% of probability, with the use of the SISVAR statistical analysis software system, version 4.0. To obtain the soaking curve and the accumulated germination of seeds, polynomial models of third degree and sigmoid were fitted in function of evaluation periods, respectively.

**RESULTS AND DISCUSSIONS**

The degree of moisture content of the seeds during the performance of the experiments was 7.8%.

The soaking process starts with the contact of the seeds with the moistened substrate (Figure 1), leading to the inference that the seeds do not present restrictions to water absorption, and the increased size of the seeds is visible in the first 12 hours. A triphasic pattern is observed in the soaking of the seeds, similarly to that found in black sucupira (Bowdichia virgilioides) (Albuquerque et al., 2009).

The phase I was characterized by a high gain of the moisture content in the first 12 hours of soaking, of about 100% (Figure 1). This phase is characterized by the high absorption of water, necessary to the germination process (Guimaraes et al., 2008). Similar results...
were observed with seeds of black sucupira (*B. virgilioides*) that presented similar behavior when maintained at the temperature of 30°C. On this species the phase I of germination is the period corresponding to 12 hours of soaking (Albuquerque *et al.*, 2009). According to Bewley & Black (1994), since the phase I is a physical process and do not depend on the metabolic activity, it can occur both in viable and dead seeds provided that there is not impediment to the water flow through the tegument.

The duration of phase II was longer, about 48 hours, with reduced weight gain (Figure 1). It is characterized by the less soaking rate and is known as stationary, where the mobilization of the reserves metabolized in the phase I to the embryonic axis occurs. Similarly to what is found in daisy seeds, in soybean seeds, the beginning of the phase II occurs at around 12 hours, but the duration time was shorter, from 12 to 30 hours (Villela *et al.*, 2007).

According to Rodrigues *et al.* (2008), the behavior of the water absorption and the duration of the phases may differ among species, cultivars or lots, due to genetic differences or the physiological quality of the seed lot.

The phase III began after 48 hours, and was evidenced by the radicle protrusion (Figure 1). At this phase, the weight gain was more expressive than in the previous phase because a higher amount of water was necessary to restart the growth of the embryonary axis and formation of the vegetative structures (Pereira *et al.*, 2007; Albuquerque *et al.*, 2009).

In the package of daisy seeds was indicate the germination of 81%. But, after the test the percentage was lower, not depending of the temperatures tested (Table 1), demonstrating that there are divergences between the results issued in the commercial packages of seeds and the real germination detected in laboratory. Therefore, stricter supervision should be adopted in the trade of ornamental plant seeds, as reported by Stefanello *et al.* (2006).

The temperature of 30°C allowed

Table 1. First germination counting (FGC), second germination counting (SGC) and germination speed index (GSI) of daisy seeds submitted to different temperatures (primeira contagem de germinação (FGC), segunda contagem de germinação (SGC) e índice de velocidade de germinação (GSI) de sementes de margarida submetidas a diferentes temperaturas). Viçosa, UFV, 2009.

<table>
<thead>
<tr>
<th>Temperature (°C)</th>
<th>Second germination counting (%)</th>
<th>First germination counting (%)</th>
<th>Germination speed index</th>
</tr>
</thead>
<tbody>
<tr>
<td>20</td>
<td>70.5 a*</td>
<td>24.0 b</td>
<td>29.3 b</td>
</tr>
<tr>
<td>25</td>
<td>75.0 a</td>
<td>42.5 a</td>
<td>38.9 a</td>
</tr>
<tr>
<td>30</td>
<td>50.0 b</td>
<td>13.5 c</td>
<td>18.7 c</td>
</tr>
<tr>
<td>20-30</td>
<td>71.5 a</td>
<td>29.5 b</td>
<td>31.1 b</td>
</tr>
<tr>
<td>CV (%)</td>
<td>7.3</td>
<td>10.4</td>
<td>10.8</td>
</tr>
</tbody>
</table>

*Average values followed by same letters in the same column do not differ by the Tukey test (5%) (médias seguidas de mesma letra nas colunas não diferem entre si pelo teste de Tukey (5%)).

Table 2. Dead and dormant seeds (%), and abnormal daisy seedlings (%), according to exposure to different temperatures during germination (sementes dormentes e mortas (%) e plântulas anormais (%) de margarida sob diferentes temperaturas de germinação). Viçosa, UFV, 2009.

<table>
<thead>
<tr>
<th>Temperature (°C)</th>
<th>Dead seeds (%)</th>
<th>Abnormal seedlings (%)</th>
<th>Dormant seeds (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>20</td>
<td>4.5 b*</td>
<td>5.3 c</td>
<td>19.8 b</td>
</tr>
<tr>
<td>25</td>
<td>2.5 ab</td>
<td>7.5 bc</td>
<td>15.0 b</td>
</tr>
<tr>
<td>30</td>
<td>6.0 a</td>
<td>16.0 a</td>
<td>27.8 a</td>
</tr>
<tr>
<td>20-30</td>
<td>3.5 b</td>
<td>11.0 b</td>
<td>14.0 b</td>
</tr>
<tr>
<td>CV (%)</td>
<td>28.9</td>
<td>18.0</td>
<td>14.0</td>
</tr>
</tbody>
</table>

*Average values followed by same letter in the same column do not differ by the Tukey test (5%) (médias seguidas de mesma letra nas colunas não diferem entre si pelo teste de Tukey (5%)).

Figure 1. Soaking curve of daisy seeds at 25°C (curva de embebição de sementes de margarida a 25°C). Viçosa, UFV, 2009.
50% of germination of the seeds, which is lower than the percentages observed under the other temperatures. The Rules for Seed Analysis (RAS) recommend alternate temperatures of 20-30°C or constant of 20°C for *Chrysanthemum indicum*, *C. nivelli* and *C. parthenium*, while for the species *C. carinatum*, *C. coronarium*, *C. segetum* and *C. coccinimum*, the conditions of 20-30°C or constant of 15°C are indicated (Brasil, 2009). Considering that daisy is a species that comes from temperate regions, the temperature of 30°C may have caused thermoinhibition to the germination process. This condition stimulated the synthesis of the abscisic acid (ABA) and inhibited the synthesis of gibberellin (GA). Such regulation was considered an adaptive response of these species to abiotic stress, which results in lower percentages of germination (Yoshioka et al., 1998; Toh et al., 2008). Supraoptimal temperatures also decreased the germination percentage of *Strelitzia reginae* seeds, due to the changes in metabolic processes, therefore the most appropriate temperature for both species was 25°C (Barbosa et al., 2005).

The temperatures of 20°C and alternate of 20-30°C allowed intermediate responses that did not differ (Table 1). In this study we used seeds from the same lot. Thus, the variations caused by the different temperatures lead to infer that the highest physiological potential of daisy seeds is expressed at 25°C, since this condition allowed the achievement of a higher number of normal seedlings, in comparison to the others.

The germination speed index is much affected by the temperature. Seeds maintained at the temperature of 25°C germinated faster than those that germinated at the constant temperature of 20°C or alternate of 20-30°C. The temperature of 30°C caused the slowest germination, which can be partially explained by the thermoinhibitory effect mentioned by Toh et al. (2008). The thermoinhibitory effect was observed in seeds of many species under temperatures of 35°C, mainly affecting the speed of germination (Akman, 2009).
The lowest number of dead seeds was observed at 25°C; the highest percentage of dead seeds was achieved at the temperature of 30°C, indicating that high temperatures cause critical damage to the germination metabolism of the seeds (Table 2). The highest percentage of dormant seeds was observed at the temperature of 30°C, and it leads to infer that the germination process was thermoinhibited. The highest percentage of abnormal seedlings was also observed at the temperature of 30°C. This condition presented the highest negative effects on the performance of the seeds.

Similarly to our results, Akman (2009) observed that there were changes in the morphology of radicles and coleoptiles of lettuce seedlings when these were soaked at temperatures higher than the ideal, inhibiting their germination. Possible changes in hormone levels may have inhibited cell elongation, affecting the growth of embryo and leading to the occurrence of the highest percentage of dead seeds and abnormal seedlings characterized by the late emission of the structures essential to the development of the plant, observed at the temperature of 30°C.

According to the rules for analyzing seeds a normal plant should have all its essential structures present, developed and healthy as can be seen. When this pattern does not occur, the seedlings are classified as abnormal (Brasil, 2009). Seedlings with malformed roots, without the aerial part and with roots infected by fungus were the main abnormal characteristics observed (Figure 2). In this study we characterized seedlings with different abnormalities that can be used as reference in the evaluation of germination tests of this species.

The daisy seeds started radicle protrusion on the fourth day after the beginning of the germination test under all the temperatures tested. However, the stabilization of the germination was influenced by temperature, and it occurred on the days 14, 12, 14 and 15 under temperatures of 20, 25, 20-30 and 30°C, respectively (Figure 3).

The evolution of the germination of seeds maintained at 25°C stabilized four days before those maintained at 30°C; this difference should be considered as determinant in the production of daisy seedlings because the conditions for faster germination of seeds allow the formation of more homogeneous stands and, consequently, seedlings of better quality and more income for producers.

In summary we observed that the seeds presented a triphasic pattern of germination. The duration of the phases I and II is 12 and 48 hours, respectively. The radicle protrusion occurs 60 hours after the beginning of the soaking at 25°C. The seeds presented thermoinhibition at the temperature of 30°C. The exposure to this temperature increases the percentage of dead and dormant seeds and cause abnormality to the seedlings. The temperature of 25°C is the most appropriate for the germination of the daisy seeds.

REFERENCES


ALBUQUERQUE KS; GUIMARÃES RM; ALMEIDA IF; CLEMENTE ACS. 2009. Alterações bioquímicas durante a embebição de sementes de sucuri-preta (Brodinia virgilioides Kunth.). Revista Brasileira de Sementes 31: 12-19.

AMATO ALP; MAIA FC; MAIA MS; CAETANO LS; SIMIONI SB; CONTO L; BONINI FILHO RM. 2007. Estabelecimento de condições de luz e temperatura na germinação de sementes de amendoim forrageiro. Revista Brasileira de Sementes 29: 61-66.


CERATTI M; PAIVA PDO; SOUSA M; TAVARES TS. 2007. Comercialização de flores e plantas ornamentais no segmento varejista no município de Lavras-MG. Ciência e Agrotecnologia 31: 1212-1218.


LANDGRAF PRC; PAIVA PDO. 2009. Produção de flores cortadas no estado de Minas Gerais. Ciência e Agrotecnologia 33: 120-126.


STEFANELLO R; GARCIA DC; MENEZES NL; WRASSE CF. 2006. Influência da luz, temperatura e estresse hídrico na germinação de sementes de anis. Revista Brasileira de Agrociência 12: 45-50.

TOH S; IMAMURA A; WATANABE A; NAKABAYASHI K; OKAMOTO M; JIKUMARU Y; HANADA A; ASO Y; ISHIYAMA K; TAMURA N; IUCHI S; KOBAYASHI Y; YAMAGUCHI S; KAMIYA Y; NAMBARA E; KAWAKAMI N. 2008. KOBAYASHI M; YAMAGUCHI S; KAMIYA Y; NAMBARA E; KAWAKAMI N. 2008. High temperature-induced abscisic acid biosynthesis and its role in the inhibition of gibberellin action in Arabidopsis seeds. Plant Physiology 146: 1368-1385.
