Effects of duration and conditions of storage on germination of seeds of Pedicularis sceptrum-carolinum (Orobanchaceae)

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Abstract

Choosing optimum conditions for plants of the Pedicularis genus to productively germinate and undergo the initial stages of development is currently a relevant problem in the search of solutions to successfully grow these taxa. For the experiments, seeds of Pedicularis sceptrum-carolinum L. (Lamiales, Orobanchaceae) were collected in the first decade of September in the vicinity of Clashinskoe Lake. The study of specifics of germination of seeds and the initial stages of the development of P. sceptrum-carolinum was carried out in controlled laboratory conditions in a climate chamber with illumination (1,200–1,500 lux, photoperiod of 9/15, temperature of 23–25 °C). After a month-long storage of seeds in their fruit capsules in the laboratory conditions, the greatest germination (83.3–93.3%) was achieved after their subsequent dry maintenance (taken out of the fruits) in a refrigerator at the temperature of +2...+3 °C for 3 or 6 months. Lower values of final germination were obtained after maintaining dry seeds at the temperature of −24...−28 °C for 3 months. Increasing periods of such storage up to six months led to decrease in the final germination and energy of germination. After-ripening lasting different periods provided lower values of the two most important parameters — final germination and energy of germination, even in cases of quite long periods of dry storage in the laboratory. The initial stages of the development of plants from seeds of P. sceptrum-carolinum, which had undergone 3-month stratification in a refrigerator, were studied during the period of 2.0 (2.5) months in different conditions: Petri dishes on moistened filter paper, and in glass vessels with settled tap water, in soil in a plastic container (pure groups of sowed seeds) and also in soil sown together with seeds of Avena sativa L., with seeds put singly into a plastic block of 9 cassettes. The study revealed morphological differences in plants that had developed over the two-months growth, in each variant of the experiment. We recorded fragmentary development of haustorial hairs on the lateral roots of the plants in the pure sown group and also the haustoriums in the group sown together with common oat. We achieved no further development and the plants died. The plants grew for a longer period (2.5 months) in the pure sown groups, which then died as well. The study we performed may be a basis for preparing successful introduction and cultivation of P. sceptrum-carolinum, which would be an important source of preservation of a species that raises concerns on account of the rapid decreases in its populations, narrowing of its range and rare occurrence.

Keywords: rare species; hemiparasites; generative reproduction; germination; Avena sativa; morphological differences.
on this subject are available in an article about biology and morphological-anatomical peculiarities of this species (Petrova & Pavlenko, 2017). At the same time, seeds of most species of Pedicularis L. genus germinate slowly and unevenly and have a low percentage of germination (McDonough, 1970; Jensen, 2004; Li et al., 2007; Kirmizi et al., 2010). All this complicates successful cultivation of these taxa (Li et al., 2007). Further details regarding the germination requirements of seeds of this genus (Ren & Guan, 2008). Furthermore, taking into account the high mortality rate of seedlings of various species of Pedicularis (Petrů, 2005), the initial stages of the development of these plants are of special interest. Likewise, growth of these plants from seeds is quite difficult in laboratory conditions (Petrova & Pavlenko, 2017; Kirillova, 2018).

Therefore, the objectives of our study were assessing the influence of duration and conditions of storage, after-ripening, dry stratification and low minus temperatures on germination of seeds of P. sceptrum-carolinum and analysis of the initial stages of the development of this species in different experimental conditions.

Materials and methods

Peculiarities of germination were studied on seeds collected in the first decade of September of 2017 in natural conditions of Chashnitskoe Lake (Yaroslavl Oblast, Rostov district, 56.936668° N, 39.374617° E). The plants grew along the edge of a waterlogged drainage ditch (of a former sand quarry) located along the south bank of the lake (Fig. 1) in a community with Marchantia sp., Equisetum hyemale L., E. scirpoides Michx., E. palustre L., Phragmites australis (Cav.) Trin. ex Steud., Salix myrinifolia Salisb. and young plants of Populus tremula L. Vegetation was sparse; soil was large-grained sand.

![Fig. 1. Growing location of Pedicularis sceptrum-carolinum (a) and appearance of the plant (b)](image)

From the moment of collection up to the start of the experiment, the seeds were in the fruits – round dark brown capsules (about 1.0 cm in diameter) kept for one month in the laboratory conditions. Then, the seeds were taken out of the capsules and divided into three equal parts: one part of them was kept dry in Petri dishes on filter paper in laboratory conditions, the second one – in a refrigerator (temperature of +2.5...+3.0 °C), and the third was put into a freezing chamber (the temperature of –24...–28 °C).

In the first experiment, seeds of P. sceptrum-carolinum were subjected to germination after a month, right after they had been taken out of the capsules – 01.10.2017; in the second, after maintenance in different conditions for 3 months – 01.01.2018; in the third – after storage for 6 months – 01.04.2018.

Peculiarities of germination of seeds of P. sceptrum-carolinum were studied in controlled laboratory conditions in a climate chamber with illumination (luminostat, illuminance of 1,200–1,500 lux, photoperiod of 9/15, and 23–25 °C temperature). Seeds (20 to each) were put in Petri dishes on filter paper moistened with settled tap water (pH = 8.2). The experiments were replicated three times; the experiments lasted for 60 days. Throughout the experiment, moisture on the filter paper was maintained at the same level.

Germination of the seeds was made according to the earlier described method (Belyakov & Lapirov, 2015). We determined the following main parameters of germination (Shipley & Parent, 1991): lag time (L) – time in days between the beginning of the experiment and start of germination; final germination (Gfin or G) – percentage of germinated seeds at the end of the experiment, which corresponds to the term “laboratory germinating capacity” in the domestic literature (Nikolaeva et al., 1999); energy of germination (E) – percentage of seeds germinated in 7 days; period of germination (P) – number of days during which the seeds germinated.

To study the initial stages of the development, seeds of P. sceptrum-carolinum, after three month stratification in the refrigerator, were put into the climate chamber with illumination (in the same controlled conditions, see above) for monitoring further development of plants in various conditions: experiment 1 – in Petri dishes on moistened filter paper (20 seeds in each); experiment 2 – in glass vessels (of 300 mL capacity) with settled tap water (50 seeds in each); experiment 3 – in soil in the plastic container (17.0 × 12.0 × 4.5 cm) (100 seeds); experiment 4 – in soil in a plastic container (17.0 × 12.0 × 4.5 cm) together with seeds of Avena sativa L. (100 seeds of each species). Common oat was chosen randomly as a host plant for monitoring possible development of the haustoria on the roots of moor-king lousewort; experiment 5 – in soil, one seed in each plastic block consisting of 9 cassettes (size of each equaling 6.0 × 5.0 × 5.5 cm).

Experiments 1 and 2 were replicated three times, experiments 3–5 – once, the duration of the experiment was 2.5 months. In experiments 3–5,
the seeds were put in soil at the depth of 1.0 cm. In the experiments, we used universal, completely ready-made nutritive soil Terra vita (Nord Palp, Russia, 2018) prepared for growing seedlings. The soil was moistened when needed.

In each variant of the experiment, after 2.0 (2.5) months, we selected 5–10 plants. During the monitoring, we performed in-detail morphological analysis of separate elements of above-ground and underground spheres of the plants. We recorded the appearance of the embryonic root, development of cotyledon leaves, development of adventitious and lateral roots, emergence of the initial and subsequent true leaves, development of haustoria and haustorial thickenings. To measure the main morphometric parameters of the plants, we used binocular microscopes MB-10 (LZOS, Russia, 2010) and MSP-2 (LOMO-Microsystems, Russia, 2009) that have micrometric scales.

All obtained results were presented as mean value ± standard deviation (x ± SD). The data were analyzed in Statgraphics Centurion XVI (StatPoint Technologies, Inc., USA, 2010). Significance of the differences between the values (P < 0.05) was determined using ANOVA in PAST (Hammer et al., 2001; Paleontological Statistics Software Package, Norway, 1999–2019). To determine significant differences, we used the Tukey test (with additions by Copenhaver & Holland (1988) and Bonferroni correction).

Results

In the central part of European Russia, blooming of P. sceptrum-carolinum occurs in July, and the fruits ripen in late July–September. The fruits are round dark brown capsules of around 1.0–1.4 cm in diameter, gathered in whorls comprising 3–4 capsules. According to our data, the average number of the capsules in the inflorescences of this plant is 16.6 ± 6.2. The number of seeds in the capsule of P. sceptrum-carolinum varies, on average equaling 94.8 ± 53.7. Large amounts of seeds are in the capsules located in the middle part of the inflorescence (150.2 ± 45.5 seeds). A total of 73.3 ± 39.7 seeds ripen in the capsules located in the lower part of the inflorescence, and 42.5 ± 25.5 in the upper part, and oftentimes the fruits have no time to form the ovary. Weight of 100 seeds is 0.024 ± 0.003 g. Seeds are elongated, light or dark brown, about 2.0–2.5 mm long, 1.0 mm wide. Our observations indicate that the dissemination of P. sceptrum-carolinum is stretched out in time, for when drying, the shoots that bear the capsules with remaining seeds do not fall over, and in conditions of intense wind, the remaining seeds continue to fall out of the hatched capsules. Analyzing the main parameters of germination of the seeds (Table 1), we should note that the highest values of two most important parameters – final germination and energy of germination – were recorded after cold stratification of dry seeds in the refrigerator. Low enough values of these parameters were obtained when storing dry seeds in laboratory conditions, whereas 6 months storage in the refrigerator led to 3-fold decrease compared with the three month period.

The general pattern of the development of plants out of seeds and initial stages of this process are presented below.

As our observations demonstrated, germination of P. sceptrum-carolinum is above-ground, hypocotylary. The first to emerge after a short period of absorbing water and swelling of seed (5–7 days) was the main root, which with growth was for some time under the protection of the external reticular sheath. Then, 2–3 days later, it comes through the reticular sheath. The main root of the seedling is white with sharpened end and indumentia of thin root hairs (sometimes they may be absent). Then, it was followed by the strong light green hypocotyl, separated from the main root by crown-like indumentum of thin semi-translucent hairs (on the root neck), the epicotyl was expressed poorly (Fig. 2).

![Fig. 2. Emergence of the main root and hypocotyl (a) from the seed coating (well noticeable indumenta of thin root hairs on the main root and root neck, b) 1 – hypocotyl, 2 – root neck, 3 – the main root)](image)

At the same time, the main root begins to bend arch-like and quickly grows in length. On the 10–11th days after sowing of seeds, two cotyledons are released from the seed coating, which gradually, on 15–16 days stretch spatially. The cotyledons are green, elongated, with entire margins, on short petioles. At the initial stage of the development, in P. sceptrum-carolinum, they may be rounded, elongated, heart-shaped, more often oval-ovoid. The first assimilating leaf emerges on the 17–18th, the second – 22–24th days, and then the plant starts developing 1–2 lateral roots. The roots were short, filament-like, located spatially horizontally under a 90 degree angle in relation to the main root.

Table 1

| Parameters               | Seeds were kept in fruit capsules in laboratory conditions for 1 month | Seeds were stored in laboratory conditions for 3 months | Seeds were kept in the fridge for 3 months | Seeds were kept in freezing chamber for 3 months | Seeds were kept in laboratory conditions for 6 months | Seeds were kept in the fridge for 6 months | Seeds were kept in freezing chamber for 6 months |
|-------------------------|------------------------------------------------------------------------|------------------------------------------------------|-------------------------------------------|-------------------------------------------------|---------------------------------------------------|-------------------------------------------|------------------------------------------------|
| L, 24 h                 | 6.33 ± 2.44               | 6.33 ± 2.44              | 4.67 ± 0.89                                | 4.33 ± 0.44                               | 8.50 ± 0.50                                      | 8.00 ± 0.00                                 | 5.67 ± 1.56                                      |
| P, 24 h                 | 9.00 ± 5.33               | 5.33 ± 2.90              | 12.67 ± 3.56                               | 13.33 ± 2.22                               | 12.50 ± 7.50                                      | 8.00 ± 0.00                                 | 7.00 ± 3.33                                      |
| G, %                    | 18.33 ± 7.86              | 13.33 ± 11.11            | 88.33 ± 22.22                              | 70.00 ± 13.33                              | 93.33 ± 4.44                                      | 25.00 ± 3.33                                 | 20.00 ± 3.33                                      |
| E, %                    | 13.33 ± 4.44              | 11.67 ± 8.89             | 80.00 ± 3.33                               | 58.33 ± 5.56                               | 6.67 ± 5.56                                      | 83.33 ± 4.44                                 | 20.00 ± 3.33                                      |

Note: different letters indicate values that statistically significantly differ from one another within one line of the table according to the results of comparison by the Tukey test with Bonferroni correction at P < 0.05.
In each of 5 variants of our experiments on growing *P. sceptrum-carolinum* from seeds in different experimental conditions in the course of two months, the plants that had grown had distinctive traits (Table 2). Therefore, during that period, the distinctive feature of the plants developing in Petri dishes (variant 1) was weak development of assimilating leaves, the sizes of which did not exceed 0.1 cm (Fig. 3, Table 2). Plants on the water surface (variant 2) were observed to have an extremely long hypocotyl, 2–8 times longer than in the other variants of the experiment and absence of lateral roots. During the development of plants in the groups of seeds sown in plastic containers (variant 3), despite the overall amount of assimilating leaves being the same as in the previous variants, the number of lateral roots was 4(8) times greater than in the rest of the experimental plants (Table 2). At the same time, on the lateral roots (variant 3), a fragmental development of haustorial hairs began, visually similar to the typical root hairs. Contrary to this, in variant 4, we observed development of lateral haustoria of rounded shape (around 0.5 mm in diameter) on the lateral roots or in the basal part of the main root (Fig. 4).

We should note that on the roots of some plants, three haustoria were developing at the same time, which were at different distances from one another. At the same time, haustoria of *P. sceptrum-carolinum*, in most cases, were located on the roots of *Avena sativa* near the tillering node, which led to gradual inhibition, and then death of the common oat. In seeds sown into soil singularly (variant 5), the main root practically completely rotted. On its short region, there usually remained one and more rarely up to three lateral roots. The development of short haustorial hairs on them was seen in singular cases. Furthermore, on a thickened part of the leaf petioles became reddish.

In seeds sown into soil individually (variant 5), the main root practically completely rotted. On its short region, there usually remained one and more rarely up to three lateral roots. The development of short haustorial hairs on them was seen in singular cases. Furthermore, on a thickened part of the leaf petioles became reddish.

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Despite the fact that the plants had morphological differences in each particular variant of the experiment (Table 2), and also differences in the rates of the process of growth and development in each particular case, for the same time interval – two months, they all were characterized by formation of two cotyledons, 1–2 weakly developed long petiole leaves, root system comprising the main and adventitious roots of the first order (at the initial stage of the development, the adventitious roots were absent in some cases). The exception was variant 5, where the number of developed green assimilating leaves in the plants was significantly higher than such in all the remaining variants of the experiments, the cotyledons died, and along with the lateral, there developed a small number of adventitious roots.

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**Table 2**
The main morphometric parameters of *Pedicularis sceptrum-carolinum* plants grown from seeds in different conditions after two months of growth (x ± SD, n = 5–10)

| Main morphometric parameters of plant | Experiment 1: Plants grown in Petri dishes on moistened filter paper | Experiment 2: Plants grown in glass vessels of 300 mL capacity with settled tap water | Experiment 3: Plants grown in soil in plastic container | Experiment 4: Plants grown in soil in plastic container together with seeds of *Avena sativa* L. | Experiment 5: Plants grown in soil, one in plastic block of 9 cassettes |
|--------------------------------------|---------------------------------------------------------------|---------------------------------------------------------------|---------------------------------------------------------------|---------------------------------------------------------------|---------------------------------------------------------------|
| **Total number of assimilating leaves** | 1.10 ± 0.85 | 2.00 ± 0.00 | 2.00 ± 0.33 | 1.33 ± 0.44 | 4.75 ± 0.90 ** |
| **Total number of lateral roots** | 2.00 ± 0.00 | 1.60 ± 0.20 | 1.53 ± 0.27 | 0.85 ± 0.25 | 3.25 ± 1.25 (3.02 ± 0.93) ** |
| **Length of hypocotyl, in cm** | 0.55 ± 0.30 | 0.50 ± 0.20 | 0.50 ± 0.20 | 0.22 ± 0.04 | – |
| **Length of the main root, in cm** | 1.46 ± 0.44 | 2.10 ± 0.63 | 1.20 ± 0.40 | 0.30 ± 0.06*** | – |
| **Maximum length of lateral roots, in cm** | 0.35 ± 0.10 | 1.53 ± 0.27 | 0.85 ± 0.25 | 0.25 ± 0.03 | – |
| **Length of cotyledon, in cm** | 0.27 ± 0.04 | 0.45 ± 0.04 | 0.34 ± 0.06 | – | – |
| **Width of cotyledon, in cm** | 0.20 ± 0.02 | 0.35 ± 0.03 | 0.25 ± 0.03 | – | – |
| **Length of petiole of the cotyledon, in cm** | 0.56 ± 0.06 | 0.36 ± 0.04 | 0.33 ± 0.02 | – | – |
| **Length of the 1st leaf, in cm** | 0.53 ± 0.10 | 0.53 ± 0.03 | 0.30 ± 0.10 | 0.52 ± 0.12 | – |
| **Length of the petiole of the 1st leaf, cm** | 0.25 ± 0.03 | 0.13 ± 0.02 | 0.25 ± 0.03 | 0.57 ± 0.14 | – |
| **Length of the 2nd leaf, cm** | 0.11 ± 0.03 | 0.46 ± 0.20 | 0.30 ± 0.10 | 0.54 ± 0.13 | – |
| **Length of the petiole of the 2nd leaf, cm** | 0.90 ± 0.04 | 1.00 ± 0.32 | 0.25 ± 0.14 | 0.52 ± 0.04 | – |
| **Length of the 3rd leaf, cm** | – | – | – | – | – |
| **Length of the petiole of the 3rd leaf, cm** | – | – | – | – | – |
| **Length of the 4th leaf, cm** | – | – | – | – | – |
| **Length of the petiole of the 4th leaf, cm** | – | – | – | – | – |

**Note:** *–* organ is absent, **–** sizes of the leaves are extremely small and do not exceed 0.1 cm, ***–** in round parentheses (in the corresponding lines), there is number and sizes of the adventitious roots, ***– residue of the main root.

*Regul. Mech. Biosyst.,* 2021, 12(2)
Unfortunately, we could not have observed the further development of the plants due to their gradual death. Plants that survived the longest (2.5 months) were the plants of moor-king lousewort in pure groups of sown seeds. Moor-king lousewort plants at that time were represented by rosette shoot with 4 (6) opposite long petiole leaves with enlarged base and rounded leaf laminae (0.2–1.0 cm in diameter), divided into 5–7 rounded parts in wave-like pattern. On the laminae and petioles, there were indumenta of scattered glandular hairs. The cotyledons usually remained. The main root was well differentiated, vertical, quite strong, white, reaching on average 1.5–2.0 (2.5) cm length. The number of lateral roots increased 1.5-fold (up to 11–12), the vast majority of them ransnaffed (up to order II), and singular ones locally bore haustorial hairs (Fig. 5). Despite such pattern of the development, with no visually seen threat to the plants and constant care for them, all of the plants ultimately died.

![Fig. 5. Appearance of Pedicularis sceptrum-carolinum plant from groups of sown plants after 2.5 months growth: 1–6, signs the same as in Fig. 3; 7 – haustorial hairs](image)

**Discussion**

Seeds of most species of the *Pedicularis* L. genus are unable to germinate right after the harvest due to being in the state of physiological dormancy (Li et al., 2007). Among various methods of overcoming this type of dormancy, one of the common ones is after-ripening – the period (usually for several months) of dry storage of freshly harvested ripe seeds at room temperature (Probert, 2000; Finch-Savage & Leubner-Metzger, 2006). Unfortunately, the use of this procedure for seeds of *P. sceptrum-carolinum* provided quite low values of the two most important parameters – final germination and energy of germination even in the conditions of long storage in laboratory conditions (Table 1). At the same time, regardless of whether the fruits contained seeds or not (in different periods of dry storage in the laboratory conditions), no significant differences between the corresponding parameters were observed. The same low percentage of germination (12.0 ± 5.7% and lower) as in our case was observed by the Chinese scientists (Ren & Guan, 2008) in the experiments with seeds of three species of louseworts (*Pedicularis rex* C. B. Clarke ex Maxim., *P. rhinanoides* Schrenk, and *P. longiflora* Rudolph), stored dry for one month in laboratory conditions.

Usually, to increase germination of seeds of representatives of the *Pedicularis* genus, cold stratification is used, sometimes combined with other types of treatments (gibberellic acid, scarification, light, darkness) (Kaye et al., 1997; Li et al., 2007; Ren & Guan, 2008; Serap et al., 2010; Belaeva et al., 2017). Therefore, for example, in freshly harvested seeds of *P. sceptrum-carolinum* picked up in late August of 2013–2014 in the vicinity of the Polar Alpine Botanical Garden (Kirovsk, Murmansk Oblast) in the foothills of Kukis Mountain along the road near waterlogged forest and in *Sphagnum* wetland and then subjected to humid cold stratification in a refrigerator (temperature + 4°C), germination varied 16% (2013) to 60% (2014) (Petrova & Pavlenko, 2017). We found no research on the roles of dry stratification and low minus temperatures for increasing germination of seeds of this species in the literature sources.

The high parameters of final germination and energy of germination which we obtained after storing dry seeds in a refrigerator for three and six months (Table 1) indicates the effectiveness of such treatment. Different parameters were seen for seeds stored for the same time in a freezing chamber. A significant effect (though significantly lower than after storage in refrigerator) was also seen three months after storage in a freezing chamber (Table 1), which makes it significantly more different from the one obtained after longer maintenance in the same conditions. Increase in duration of maintenance in low minus temperatures likely causes the seeds to enter the state of secondary dormancy, causing rapid decrease in the main parameters of germination.

Research on the conditions needed for germination revealed that the germination of this species is adapted to the environmental conditions, for it is regulated by environmental factors (Van Assche et al., 2002). Knowledge of the conditions needed for germination of local species, especially rare species, is crucial for protection and restoration of biological diversity (Cerabolini et al., 2004). Furthermore, for threatened species, preservation of the components of biological diversity outside their natural habitats (ex situ preservation), growing plants from seeds, is considered a vital and cheap method (Serap et al., 2010). Taking into account that vegetative reproduction of *P. sceptrum-carolinum* is possible, but rarely takes place, and may occur only on small scales of one mother shoot, generative reproduction (as a system of reproduction and distribution by seeds) is essential for this species. At the same time, one needs to take into account that seeds of this plant have no morphological adaptations to spreading across large areas, but the fruits and shoots are sometimes eaten by animals that may be seed carriers across large distances (Wroblewska, 2013). Difficulties with process of growing moor-king lousewort from seeds were seen earlier, because the species is unable to develop on its own for a long time at early stages of the development, even when added to reliably determined hosts (Petrova & Pavlenko, 2017). As possible explanations of unsuccessful attempts to grow this plant from seeds, these scientists named damage to the root system of reliable host plants after transfer from the natural environment to the artificial, their nonadaptiveness to new conditions, and also presence of a particular specialization of *P. sceptrum-carolinum* to host plants. In our case, further gradual death of plants in all variants of the experiment after two-months growth, except pure groups of sown seeds (variant 3), was likely related not only to high mortality among seedlings (Petrů, 2005). As some researchers indicate, seeds of the representatives of *Pedicularis* genus germinate regardless of host plants, and seedlings for their successful development at different stages require functional haustorial connections to the roots of host plants (Ter Borg, 1985; Rúžica, 1999 – quoted according to Petrů, 2005). The reason for failure of growing *P. sceptrum-carolinum* together with common oat is likely the certain selectivity of host plants, which is typical for this species (Petrova & Pavlenko, 2017). Furthermore, death of plants in our experiment is likely related to deficiency of nutrients, intake of which is extremely sensitive for the growth of *Pedicularis* genus in the absence of host plants (Li et al., 2013). It is especially relevant for nitrogen and phosphorus, especially because the latter element limits the growth of roots of hemiparasitic plants (Li et al., 2013). Therefore, it is fair to say that more experiments are needed to seek the most productive conditions for growing *P. sceptrum-carolinum*. Successful experiments on growing some species of lousewort (*P. rex* and *P. tricolor*) in a greenhouse, conducted by the group of Chinese scientists (Li et al., 2013), confirm that growing hemiparasitic plants is quite possible.
Conclusions

Taking into account the vulnerability of populations of P. sceptrum-carolinum and decrease in the number of locations of its growth, search for effective methods of its equal and fast germination with high germination percentage is essential for cultivating this plant and making it cultivable. Our studies demonstrated that maximum value of the final germination may be achieved after keeping dry seeds of this plant in a refrigerator for 3 or 6 months. Knowledge of the conditions needed to grow the seeds of this plant would make it possible to predict the course of this process in conditions of land amelioration and drainage reclamation of wetlands, in cases of cold spells returning in early spring (low plus temperatures) or after minus temperatures in winter. Furthermore, the conditions needed for effective germination of seeds of P. sceptrum-carolinum which we found in our study will be useful for the future ex situ storage of this species.

Attempts to grow plants from seeds indicated that their development in experimental conditions is possible during 2.0–2.5 months. Longer period of germination and to cultivation likely requires thorough selection of host plants, as well as storage conditions, taking into account complete provision of the plants at the early development stages with a set of nutrients needed for their growth and development.

The work was carried out within the framework of the State Assignment of the Ministry of Education and Science of the Russian Federation No. 12105100099-5.

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