Species Preferentiality of the Pollen Tube Attractant Derived from the Synergid Cell of *Torenia fournieri*1

Tetsuya Higashiyama*, Rie Inatsugi, Sachio Sakamoto, Narie Sasaki, Toshiyuki Mori, Haruko Kuroiwa, Takashi Nakada, Hisayoshi Nozaki, Tsuneyoshi Kuroiwa, and Akihiko Nakano

Department of Biological Sciences, Graduate School of Science, University of Tokyo, Hongo, Tokyo 113–0033, Japan (T.H., R.I., S.S., T.N., H.N., A.N.); Department of Biology, Faculty of Science, Ochanomizu University, Otsuka, Tokyo 112–8610, Japan (N.S.); and Department of Life Science, College of Science, Rikkyo (St. Paul's) University, Nishi-Ikebukuro, Tokyo 171–8501, Japan (T.M., H.K., T.K.)

The synergid cell of *Torenia fournieri* attracts pollen tubes by a diffusible but yet unknown chemical attractant. Here we investigated the species difference of the attractant using five closely related species in two genera, namely *T. fournieri*, *Torenia baillonii*, *Torenia concolor*, *Lindernia (Vandellia) crustacea*, and *Lindernia micrantha*. These five species have an exserted embryo sac, and ablation experiments confirmed that their synergid cells attracted the pollen tube. When ovules of *T. fournieri* and one of the other species were cultivated together with pollen tubes of each species, pollen tubes were significantly more attracted to synergid cells of the corresponding species. The attraction was not affected by the close proximity of embryo sacs of different species. This suggests that the attractant is a species-preferential molecule that is likely synthesized in the synergid cell. The calcium ion, long considered a potential attractant, could not serve as the sole attractant in these species, because elevation of the calcium ion concentration did not affect the observed attraction. In vivo crossing experiments also showed that the attraction of the pollen tube to the embryo sac was impaired when pollen tubes of different species arrived around the embryo sac, suggesting that the species preferentiality of the attractant may serve as a reproductive barrier in the final step of directional control of the pollen tube.

The chemistry of pollen tube guidance has been studied for more than a century. Many attempts have been made to identify the guidance cues that regulate the directional growth of pollen tubes in pistils. Several molecules potentially involved in pollen tube guidance have been identified, including the guidance cue. External calcium ions were first identified as a potential attractant in the snapdragon (*Antirrhinum majus*) pistil using classic in vitro tests (Mascarenhas and Machlis, 1962). Other small molecules, including sugars (Reger et al., 1992) and nitric oxide (Prado et al., 2004), are also reported to control directionality of the pollen tube by using in vitro tests. A 9.9-kD basic protein, chemocyanin, was isolated from lily (*Lilium longiflorum*) stigma proteins by biochemical fractionation using in vitro tests of its ability to attract pollen tubes (Kim et al., 2003). A single homolog of chemocyanin may also be involved in pollen tube guidance in Arabidopsis (*Arabidopsis thaliana*), according to in vivo experiments (Dong et al., 2005). A gradient of the concentration of water in lipids (triacylglycerides) on the stigmas of tobacco (*Nicotiana tabacum*) and Arabidopsis was proposed to be critical for the initial directional growth of the pollen tube entering the stigma (Wolters-Arts et al., 1998). Transmitting tissue-specific protein in tobacco (Cheung et al., 1995; Wu et al., 1995, 2000) and stigma/stylar Cys-rich adhesin and pectin in lily (Mollet et al., 2000; Park and Lord, 2003) were identified in style tissue and shown to be necessary for the growth of the pollen tube in the style. An appropriate concentration of γ-aminobutyric acid (GABA) concentrated toward the nucellus tissue surrounding the embryo sac was proposed to aid the pollen tube in navigating toward the ovule (Palanivelu et al., 2003). In the maize (*Zea mays*) embryo sac (female gametophyte), *ZmEA1* is expressed in both the egg cell and the synergid cells, and its product appears to be secreted into the micropyle, which was shown to be required for pollen tube guidance (Márton et al., 2005). Guidance by these molecules allows pollen tubes to grow directionally toward the target embryo sacs. However, both in vivo and in vitro confirmation that these molecules are the true attractants governing pollen tube guidance are required.

The two synergid cells to either side of the egg cell are the most plausible emitters of chemoattractants and are involved in navigating the final growth of the pollen tube toward the embryo sac. In *Torenia fournieri*,
pollen tubes are directly attracted to the exserted embryo sac in vitro (Higashiyama et al., 1998). Laser ablation experiments identified the source of the diffusible signal as the two synergid cells (Higashiyama et al., 2001). Once attracted, pollen tubes of *T. fournieri* never leave the embryo sac and form narrow coils on the surface of the micropylar end of the embryo sac before entering the sac, suggesting that the pollen tube is trapped at the highest concentration of the chemotactant. In Arabidopsis, the target embryo sac itself guides the final steps of the pollen tube into the ovary (Hülskamp et al., 1995; Ray et al., 1997; Shimizu and Okada, 2000), and use of the mutant line *myb98*, which is defective in synergid cell development, revealed that the last guidance toward the micropyle is provided by the synergid cell (Kasahara et al., 2005).

Two molecules have been reported to be candidates for pollen tube attractants that are derived from the synergid cells. One is the external calcium ion. In vitro test results suggest that calcium is an attractant derived from the pistil, in particular the ovule (Mascarenhas and Machlis, 1962, 1964; Reger et al., 1992). Using various histochemical methods including calcium antimonate precipitation, proton-induced x-ray emission, and fluorescent calcium probes, high concentrations of calcium have been observed in synergid cells and the surrounding extracellular matrices in various species (Jensen, 1965; Chaubal and Reger, 1990, 1992a, 1992b, 1993; Huang and Russell, 1992; Tirlapur et al., 1993; Tian and Russell, 1997), including *T. fournieri* (Kristóf et al., 1999). The second potential attractant is *ZmEA1* (Márton et al., 2005). The *ZmEA1* gene was originally identified from a maize egg cell cDNA library and was shown to be expressed more abundantly in the synergid cell. *ZmEA1* protein fused with green fluorescent protein appeared to be secreted toward the extracellular matrix of the nucellus at the micropyle, and some antisense RNA and RNAi lines showed reduced fertility because of a defect in pollen tube guidance at the micropyle. *ZmEA1* is a member of a large family of EA1-like genes found in flowering plants (Gray-Mitsumune and Matton, 2006). *ZmEA1* is a good candidate for a synergid cell-derived chemotactant, but ability of purified *ZmEA1* to attract the pollen tube has not yet been demonstrated.

Species difference can provide insight into the chemical properties of the chemotactant, which might be a more complex compound such as *ZmEA1* rather than a low-Μ compound such as the calcium ion. Species difference could also function as a reproductive barrier at the pollen tube guidance step. To determine if the attractant derived from the synergid cell is a species-specific molecule, an in vitro system using the exserted embryo sac seems indispensable, because it enables the exclusion of other guidance steps. When the exserted embryo sac is used, the basal end of the synergid cell is directly exposed to the medium and the attraction signal spreads into the medium from the synergid cell; pollen tubes do not need to grow on the surface of the surrounding tissues of the embryo sac, which may also contribute to pollen tube guidance. By mixing ovules of two species, one can critically examine whether pollen tubes sense the attraction signal of a different species under conditions in which they can surely respond to the attraction signal of their own species. In addition, the condition of the synergid cell is easily observed in the exserted embryo sac, but the attraction can only be examined in embryo sacs with complete synergid cells (the synergid is a fragile cell; Higashiyama et al., 1998). Ablation experiments using UV lasers are also possible in the exserted embryo sac, as shown in *T. fournieri* (Higashiyama et al., 2001), and can be used to confirm the role of the synergid cell in pollen tube attraction.

In this study, to characterize the properties of the chemotactant derived from the synergid cell, we investigated the species difference of the attractant using an in vitro *T. fournieri* system and four closely related species that possess an exserted embryo sac. The attractant from the synergid cell appeared to be a species-preferential molecule and may have a role in the reproductive barrier. Finally, it was suggested that the calcium ion was not solely the synergid cell-derived attractant in *T. fournieri*.

**RESULTS**

Scrophulariaceae Species with an Exserted Embryo Sac Can Be Fertilized In Vitro in the Same Manner as *T. fournieri*

To determine if the attractant derived from the synergid cell was a species-specific molecule, we first surveyed several plant species with exserted embryo sacs. As reviewed by Maheshwari (1950), some flowering plant species exhibit exserted embryo sacs, including some species of Philipsiphus, Thesium, Galium, Utricularia, Lindernia (Vandelia), and Torenia. This study required plants with ovaries with a large number of ovules that are suitable for cultivation. Because Galium species possess only a few ovules per ovary, and excision is difficult because the ovules are embedded in the ovarian tissue, these species were unsuitable. Philadelphus species, such as *Philadelphus satsumi*, possess a large number of ovules with an exserted embryo sac, and their flowers are readily available because of their popularity as ornamental plants. However, Philadelphus ovules tended to brown and inhibit growth of pollen tubes in culture. *Utricularia* (*Utricularia* spp.), an aquatic vermivorous plant, was not used because its flowers are difficult to obtain under laboratory conditions.

The *Torenia* and *Lindernia* genera of the Scrophulariaceae contain several species with an exserted embryo sac. In plants of these genera, up to several hundred ovules were obtained from individual ovaries. These plants easily grew and bloomed in growth chambers; their pollen tubes and ovules were able to be cultivated in the same manner as those of *T. fournieri*. 

---


Copyright © 2006 American Society of Plant Biologists. All rights reserved.
Torenia plants and flowers were larger than those of Lindernia, and the Torenia genus provided the most easily usable exserted embryo sacs. All of the Torenia species tested possessed an exserted embryo sac, but Lindernia showed variations; among the four tested species, Lindernia crustacea (Vandellia crustacea) and Lindernia micrantha (Vandellia angustifolia) possessed an exserted embryo sac, but Lindernia setulosa (Vandellia setulosa) and Lindernia antipoda (Vandellia anagallis) possessed a normal embryo sac enclosed by the integument. Therefore, we chose five species in two genera for subsequent analyses: T. fournieri, Torenia baillonii, Torenia concolor, L. crustacea, and L. micrantha. Flowers and ovules of these five species are shown in Figure 1. All five species exhibit an exserted embryo sac of the Polygonum type that protrudes from the micropyle of the ovule.

Figure 2 presents the results of a phylogenetic analysis using the large subunit of Rubisco (rbcL) nucleotide sequences from three Torenia and four Lindernia species, as well as 32 other species in the Scrophulariaceae (Wolfe and dePamphilis, 1998). The genes from the Torenia and Lindernia species formed a monophyletic group in the Scrophulariaceae with high bootstrap values: 99.5% for maximum parsimony (MP) analysis and 99.8% for neighbor-joining (NJ) analysis and high posterior probability (1.0 for Bayes’ theorem). Within this monophyletic group, the five species with exserted embryo sacs subsequently used in our analyses constituted a clade with high bootstrap values (97.7% for MP and 99.3% for NJ) and high posterior probability (1.0 for Bayes’ theorem). These results suggest that the five species are closely related. Among the five species, T. baillonii appeared closest to T. fournieri, while L. micrantha appeared most divergent.

Confirmation of the Synergid Cell Origin of the Attractant in the Studied Species

Before investigating the species difference, we confirmed that the pollen tube attraction occurred in vitro in all plant species and that the source of the attraction signal was the synergid cell, as in T. fournieri (Fig. 3). Ovules and a pollinated style of each species were cultivated in a medium previously used for T. fournieri (Higashiyama et al., 1998, 2001). In proportion to the size of the pistil, both the number of ovules and the length and number of pollen tubes increased on this medium. The number of ovules in an ovary was 300 to 500 in T. fournieri and T. baillonii, more than 500 in T. concolor, about 150 in L. crustacea, and about 300 in L. micrantha. After overnight growth in culture, attraction of the pollen tube to the embryo sac was observed in all species (Fig. 3A); there were multiple pollen tubes around ovules, and some of them grew toward the micropylar end of the embryo sac to form a narrow coil, as described for T. fournieri (Higashiyama et al., 1998). Discharge of the pollen tube contents and onset of seed development were observed as frequently as in T. fournieri. When pollen tubes of these species were germinated on the medium, no attraction of pollen tubes was observed (data not shown), as also shown for T. fournieri (Higashiyama et al., 1998), probably because the pollen tubes failed to acquire the response capability.

The ratios of the complete embryo sac (possessing the egg cell, two synergid cells, and the central cell) attracting pollen tubes are shown in Figure 3B. Most complete embryo sacs of the Torenia species attracted pollen tubes (88.4% ± 6.9% in T. fournieri [n = 141], 88.0% ± 6.5% in T. baillonii [n = 161], and 91.2% ± 8.1% in T. concolor [n = 159]); the percentages refer to the ratio of embryo sacs attracting pollen tubes out of all complete embryo sacs with SD of three replications, and n refers to total number of complete embryo sacs counted. Considering the distance of attraction, a few hundred micrometers at most (Higashiyama et al., 2003), ovules that had pollen tubes in the vicinity (approximately 100 μm) were included in calculating the ratio of pollen tube attraction. The complete embryo sacs of Lindernia species also attracted pollen tubes, but at lower frequencies than Torenia embryo sacs (16.7% ± 8.0% in L. crustacea [n = 132] and 42.8% ± 11.7% in L. micrantha [n = 120]). This result was due to limitations to the numbers and lengths of pollen tubes...
in Lindernia; pollen tubes could not cover the entire area of the ovules, and there were no pollen tubes in the vicinity of many ovules. It has been proposed that the attraction of the *T. fournieri* pollen tube by the synergid cell is effective within a few hundred micrometers in this medium (Higashiyama, 2002).

We next confirmed the contribution of the synergid cell to the attraction. When the synergid cells on either side of the egg cell were ablated using a UV laser, the attraction was completely halted in all species (0%; *n* = 62 for *T. fournieri*, *n* = 72 for *T. baillonii*, *n* = 66 for *T. concolor*, *n* = 60 for *L. crustacea*, and *n* = 60 for *L. angustifolia*; Fig. 3B). We confirmed that in all tested species, the synergid cell attracts the pollen tube, as previously reported for *T. fournieri* (Higashiyama et al., 2001).

**Species Preferentiality of the Synergid Cell-Derived Pollen Tube Attractant**

To examine the species difference of the attractants derived from the synergid cells, we cultivated *T. fournieri* ovules with ovules of other plant species (Fig. 4). Pollen tubes of each plant species were then grown. Ovules of *T. fournieri* and the other species could be distinguished by appearance (size, shape, and color of chloroplasts in the integument), as shown in Figure 1. However, it was impossible to discriminate among pollen tubes of different species. Therefore, ovules were mixed together and pollen tubes of each plant species were grown in separate experiments. Mixing of ovules allowed us to check that pollen tubes were able to respond to the attractant of the same species.

Figure 4 shows the results of pollen tube attraction in the presence of ovules of two different species in a culture. When *T. fournieri* and *T. baillonii* ovules were cultivated together in the same dish, *T. fournieri* pollen tubes tended to grow toward *T. fournieri* embryo sacs (68.6% ± 15.4%; *n* = 102 [total no. of embryo sacs counted]), although the tubes also grew toward *T. baillonii* embryo sacs (34.5% ± 12.9%; *n* = 132). Similarly, *T. baillonii* pollen tubes tended to grow toward both *T. baillonii* embryo sacs (66.8% ± 20.6%; *n* = 200) and *T. fournieri* embryo sacs (54.3% ± 22.0%; *n* = 112). The difference of the attraction was significant (χ² test; *P* < 0.01). A similar tendency was observed when *T. concolor* was used in place of *T. baillonii*; *T. fournieri* pollen tubes were attracted by *T. fournieri* embryo sacs at 64.2% ± 22.7% (*n* = 96) and by *T. concolor* embryo sacs at 57.9% ± 20.5% (*n* = 112).
sacs at 36.4% ± 21.6% (n = 113), and T. concolor pollen tubes were attracted by T. concolor embryo sacs at 79.1% ± 10.8% (n = 90) and by those of T. fournieri at 35.4% ± 29.8% (n = 146; χ² test; P < 0.01). When L. crustacea was used, few T. fournieri pollen tubes grew toward L. crustacea embryo sacs (1.7% ± 3.4%; n = 134). In contrast, L. crustacea pollen tubes grew toward the embryo sacs of both T. fournieri (47.7% ± 25.8%; n = 96) and L. crustacea (13.1% ± 3.6%; n = 101); interestingly, L. crustacea pollen tubes tended to grow toward the embryo sacs of T. fournieri (t test; P < 0.05). In the most divergent combination, T. fournieri and L. micrantha, pollen tubes of each species grew primarily toward embryo sacs of the same species; T. fournieri pollen tubes were specifically attracted to T. fournieri embryo sacs at 85.0% ± 16.3% (n = 159; L. micrantha; 0% [n = 154]), and L. micrantha pollen tubes were attracted to their own embryo sacs at 30.0% ± 9.1% (n = 110) and to those of T. fournieri at 3.1% ± 6.3% (n = 90). These results indicate that the attraction signals of these plant species differ.

To determine whether different concentrations of the same attractant are responsible for the species-preferential attraction responses, we used T. fournieri and Lindernia ovules to examine whether intraspecies attraction is affected by the presence of ovules of a different species (Fig. 5). If the attractant was a species-preferential difference in concentration and not a species-preferential molecule, the resulting higher concentration of attractant should mask or erase the signal conveyed by a lower concentration of attractant. At the start of cultivation, a micromanipulator was used to move about 10 ovules of both species (10 each of Torenia and Lindernia ovules) to face the embryo sacs toward sacs of the other species. As the distance between the micropylar ends of these embryo sacs (filiform apparatus of synergid cells) was within 30 μm, they were always in the range of attraction of the T. fournieri synergid cells (Higashiyama et al., 2003; R. Inatsugi, A. Nakano, and T. Higashiyama, unpublished data). The resulting frequency of attraction was similar to that without manipulation of ovules, and the attraction was not impaired by the presence of embryo sacs of another species (Fig. 5; n = 30 in total for each column). Thus, the attractant derived from the synergid cell, at least in T. fournieri and closely related species, is likely a species-preferential molecule.

**Figure 4.** Species preferentiality of the pollen tube attraction signal derived from the synergid cell. Each column indicates the mean value with the SD of three replications. OV, Ovule; PT, pollen tube; ST, style; Tf, T. fournieri; Tb, T. baillonii; Tc, T. concolor; Lc, L. crustacea; Lm, L. micrantha.

**Figure 5.** Species preferentiality assay following micromanipulation of ovules to place embryo sacs of different genera opposite each other. A, DIC images of T. fournieri (left) and L. micrantha (right) pollen tube attraction in the presence of a heterogeneous embryo sac. Arrows indicate attracted pollen tubes. B, Frequency of pollen tube attraction following micromanipulation of ovules to place embryo sacs of different genera opposite each other. Ten sets of ovules were used in each experiment, and each column indicates the mean value with the SD of three replications. ES, Embryo sac; OV, ovule; PT, pollen tube; ST, style; Tf, T. fournieri; Lc, Lindernia crustacea; Lm, L. micrantha. Bar = 50 μm.

**Contribution of the Stigma and Style Tissues to Species Preferentiality**

Pollen tubes used in attraction studies with Torenia require a period of tube elongation within the style to gain their competence to respond to the synergid chemical signal (Higashiyama et al., 1998). To determine whether species preferentiality was involved in acquiring this competence, we next changed species of the style tissue (Fig. 6). The growth of the pollen tube decreased with increasing divergence between the pollen species and the style species. However, in all combinations tested, pollen grains germinated on the stigma and began to grow in the stylar canal of heterogeneous species. When T. fournieri pollen grains were applied to T. baillonii stigmas, emergent T. fournieri...
pollen tubes still tended to grow toward T. fournieri embryo sacs (69.0 ± 21.6%; n = 82) rather than to T. baillonii embryo sacs (33.9 ± 17.5%; n = 104). In contrast, T. baillonii pollen tubes that grew through T. fournieri styles tended to grow toward T. baillonii embryo sacs (77.2 ± 21.5%; n = 101) rather than to T. fournieri embryo sacs (63.4 ± 15.8%; n = 79). The tendency of attraction was significant (χ² test; P < 0.01). A similar pattern was observed when T. concolor was used instead of T. baillonii (χ² test; P < 0.01). These results indicate that pollen tube detection of the synergid cell attractant is conferred by pollen species, not by sporophytic factors found in the style on which the pollen tube grows.

When Lindernia species were used, Lindernia pollen tubes did not emerge from the cut ends of T. fournieri styles, as described later; however, T. fournieri pollen tubes did emerge from Lindernia styles, but the number of pollen tubes decreased (Fig. 6). The Torenia pollen tubes growing through Lindernia styles showed impaired growth, as they were considerably shorter than tubes grown in Torenia styles. These T. fournieri pollen tubes were not attracted to the synergid cells of either T. fournieri or the Lindernia species. They appeared to fail to acquire the capability to respond to the attraction signal because of the heterogeneous conditions of the stigma and style.

**Evaluation of Potential Low-Μ Calcium Ions**

The species preferentiality of the pollen tube attractant is inconsistent with the classical hypothesis of calcium ion as the attractant. The low-molecular mass external calcium ion has been thought to be the pollen tube attractant derived from the ovule, or more precisely, the synergid cell, and was thought to form a concentration gradient in the pistil. To examine whether calcium ions are the attractant derived from the T. fournieri synergid cell, we increased the concentration of calcium ions in the medium. If an appropriate calcium ion concentration gradient had already been established by the synergid cell in the medium, the additional elevation of the calcium ion concentration in the medium should disturb the signal.

Figure 7A shows the relation between the calcium ion concentration (top) and the GABA concentration (bottom) in the medium and length of pollen tubes growing semi-in vitro. The effects of GABA, a potential chemoattractant derived from ovular sporophytic tissues (Palanivelu et al., 2003), were examined in the presence of 2 mM calcium ions. Calcium ions are generally essential to pollen tube growth. In T. fournieri, calcium ions promoted pollen tube growth and were most effective at 2 mM, which was the concentration of the in vitro T. fournieri system. At calcium concentrations greater than 10 mM, pollen tube growth was considerably impaired. GABA also promoted the growth of T. fournieri pollen tubes, as reported for Arabidopsis (Palanivelu et al., 2003), but at GABA concentrations greater than 10 mM, the growth of T. fournieri pollen tubes was impaired.

Figure 7B shows the effects of increasing the calcium ion and GABA concentrations in the medium on the percentage of embryo sacs attracting pollen tubes. Even when the calcium ion and GABA concentrations were increased to 20 and 10 mM, respectively, T. fournieri pollen tubes were still attracted by the synergid cell, although pollen tube growth began to be impaired. These results indicate that calcium ions and GABA cannot be the sole attractants of pollen tubes derived from T. fournieri synergid cells.

**In Vivo Crossing Analysis**

Finally, we examined the possibility that the species preferentiality of the attractant serves as a reproductive barrier in vivo. In all crosses tested, pollen grains were germinated on heterogeneous stigma and pollen...
tubes began to grow (Fig. 8). Figure 8B shows frequencies of penetration of the embryo sac by the pollen tube in each cross. In most combinations, except for the pollination of Lindernia pollen on Torenia pistils, pollen tubes reached the ovary locules where exserted embryo sacs were present (Fig. 8A, except $Lc \times Tf$ and $Lm \times Tf$). Pollen tubes of heterologous species arrived at the ovary in the same time period as tubes of the same species but showed a decreased ratio of penetration of the embryo sac, as shown in Figure 8B. This raises the possibility that pollen tube guidance in the ovary was impaired, suggesting that the species preferentiality of the attraction signal from the synergid cell acts as a reproductive barrier. At a considerable time after pollination, such as a few days later, most embryo sacs of Torenia had received pollen tubes by interspecific crosses, and some sacs of Lindernia had received tubes by intergenic crosses using $T. fournieri$ (data not shown). Fertilization may have been delayed in these crossings.

Because no Lindernia pollen tubes were observed in Torenia ovaries, we stained pollen tubes in the Torenia style with aniline blue and found that the pollen tubes stopped growing within the style after a period of normal growth (Fig. 8A, $Lc \times Tf$ and $Lm \times Tf$). The tubes stopped growing at a particular place and formed narrow coils or zigzag growth patterns, and finally their growth appeared to be arrested. Interestingly, the tubes of $L. crustacea$ and $L. micrantha$ grew straight within the length of their own pistils, growing straight for $6.2 \pm 0.3$ mm (no. of stylar canals observed, $n = 6$) and $6.8 \pm 0.3$ mm ($n = 6$), respectively, whereas the length of pistils of $L. crustacea$ was $5.3 \pm 0.2$ mm ($n = 10$) and that of $L. micrantha$ was $6.3 \pm 0.2$ mm ($n = 10$).

**DISCUSSION**

In this study, we demonstrated that the attraction signal from the $T. fournieri$ synergid cell and closely related species is strongly species preferential. Species preferentiality and specificity in the last phases of guidance has also been implied in interspecific and intergenic crosses using Arabidopsis (Shimizu and Okada, 2000; Shimizu, 2002; Palanivelu and Preuss, 2006). However, directional growth of pollen tubes in the pistil is governed by complex multistep controls from both the female sporophyte and gametophyte...
We showed that, in *T. fournieri*, external calcium ions could not serve as the sole synergid cell-derived attractant (Fig. 7). Although calcium has been detected in the synergid cell and the neighboring extracellular matrix, as described above (Jensen, 1965; Chaubal and Reger, 1990, 1992a, 1992b, 1993; Huang and Russell, 1992; Tirlapur et al., 1993; Tian and Russell, 1997; Kristóf et al., 1999), Ca\(^{2+}\) may play a primary role in pollen tube discharge, gamete fusion, or both, but not in species-related pollen tube attraction. In an Arabidopsis mutant in which a plasma membrane Ca\(^{2+}\) pump of the pollen tube, ACA9, is disrupted, the pollen tube arrives at the embryo sac but cannot discharge its contents to the receptive synergid cell and shows overgrowth in the embryo sac (Schött et al., 2004). ACA9 appears to be involved in intercell communication between the synergid cell and the pollen tube that triggers pollen tube discharge, along with the *PERONIA* and *SIRENE* genes that are expressed in the synergid cell (Huck et al., 2003; Rotman et al., 2003). The role of Ca\(^{2+}\) in gamete fusion has been suggested in a maize in vitro fertilization system; an isolated sperm cell can fuse autonomously with an isolated egg cell in the presence of a high concentration (5 mM) of calcium (Faure et al., 1994).

GABA has also been proposed to have a role in pollen tube guidance at the ovule of Arabidopsis, as a sporophytic guidance cue (Palanivelu et al., 2003). GABA slightly promoted the growth of Torenia pollen tubes in vitro at approximately 1 mM, as in Arabidopsis. However, we observed no change in pollen tube guidance when the GABA concentration in the medium was elevated, although our results do not exclude the possibility that GABA is a sporophytic guidance cue from the ovule. It should also be noted that in Torenia, the embryo sac is directly exposed to the placenta, and guidance of the funiculus does not appear to occur.

The stigma and style tissues contribute to capacitation of the pollen tube (Higashiyama et al., 1998; Palanivelu and Preuss, 2006). Even when the stigma and style species were changed in the Torenia genus, the species preferentiality was not altered (Fig. 6). The species preferentiality of the attraction signal from the synergid cell appeared to depend on the genotype of the pollen tube and the synergid cell (or the ovule). Arabidopsis pollen tubes acquire competence when passed through the stigma and style of Arabidopsis arenosa, *Olimarabidopsis pumila*, *Capsella rubella*, and *Sysimbrium irio*, suggesting that the ability of the stigma and style tissues to promote pollen tube competence is highly conserved (Palanivelu and Preuss, 2006). Torenia pollen tubes, however, failed to be capacitated by Lindernia stigma/style tissue. The capacitation signal from stigma/style tissue might be species preferential, and these species might be diverged enough where this capacitation fails, although it should be considered that Lindernia stigma/style tissue did not support normal growth of Torenia pollen tubes in the medium.
In most in vivo crossing combinations, the arrival of the pollen tube at the embryo sac was delayed or failed, although the pollen tubes appeared to enter the ovary normally (Fig. 8). The species preferentiality of the synergid cell attractant may be involved in this defect, possibly functioning as a reproductive barrier. Delayed pollen tube arrival may result in out competition by pollen tubes of the same species. Many pre- and postfertilization reproductive barriers exist in flowering plants (Shimizu, 2002). Severe species preferentiality of the attractant derived from the synergid cell is likely to be one of these strong barriers. Lindernia pollen tubes, on the other hand, ceased growth at a particular place in the T. fournieri stylar canal after growing slightly longer than the length of own pistils. A mechanism to stop the growth of the pollen tube autonomously may exist based on the tube length. It is also interesting that L. crustacea pollen tubes cannot enter the ovary due to this mechanism, a kind of reproductive barrier, although they have an ability to respond to the synergid cell attractant of T. fournieri (Fig. 4).

CONCLUSION

The attraction signal from the synergid cells in Torenia and closely related Lindernia species was shown to be strongly species preferential. It was shown that external calcium ions could not be the sole attractant derived from the T. fournieri synergid cell. The species preferentiality of the attractant signal may also function as a reproductive barrier in the final step of guidance. Because the attractant is likely to be a molecule that rapidly diverged during evolution, molecules synthesized in the synergid cell, such as proteins and peptides, may be candidates for this attractant. Thus, analysis of genes specifically expressed in the synergid cell will provide insight into the molecular mechanisms of pollen tube attraction.

MATERIALS AND METHODS

Plant Materials and Growth Conditions

Plants of the five species used in the study, Torenia fournieri, Torenia baillonii, Torenia concolor, Lindernia crustacea, and Lindernia micrantha, were grown in a regulated chamber at 25°C with a 16-h photoperiod (approximately 150 μmol m⁻² s⁻¹). Lindernia setulosa and Lindernia antiqua were grown in the same condition and used for phylogenetic analysis. For each experiment, ovules with placenta were excised from flowers with freshly opened stigmas using a stereomicroscope. Except for T. fournieri, these plants were automatically self-pollinating; thus, they were emasculated before flowering by removing the sympetalous petals from which the stamens emerged. The pistils matured normally and were pollinated with pollen grains from other flowers. None of these plants exhibited self-incompatibility.

Phylogenetic Analysis

Total DNAs were isolated from the seven species of Torenia and Lindernia. rbcL was PCR amplified using RH1 (5'') and 1352R (3'') primers, as reported in Wolfe and dePamphilis (1997). PCR products were directly sequenced using RH1, 1352R, and a forward primer (5'--CTAGTCTGGAAAGCCCTGC-GAATCC-3') at nucleotide positions 371 to 395 on the sequence for T. fournieri (AF126842). The rbcL sequences of other 32 species of Scrophulariaceae (Wolfe and dePamphilis, 1998) were obtained from GenBank. Tobacco (Nicotiana tabacum) and Nicotiana debneyi (Solanaeaceae) were used as the outgroup of this study as described by Wolfe and dePamphilis (1998).

The rbcL sequences were aligned using ClustalX (Thompson et al., 1997), and nucleotide positions 213 to 1,533 on the full coding sequence of rbcL for tobacco were used to obtain the best alignment. Un weighted most parsimonious analyses of rbcL genes were performed, including bootstrapping (Felsenstein, 1985) based on 1,000 replications of full heuristic searches (with the tree bisection-reconnection branch-swapping algorithm), using PAUP 4.0b10. Distance matrix was calculated by applying the Jukes-Cantor method (Jukes and Cantor, 1969) in PAUP 4.0b10. NJ trees (Saitou and Nei, 1987) were constructed using PAUP 4.0b10, including bootstrap analyses (Felsenstein, 1985) based on 1,000 replications. Bayesian analyses were performed using MrBayes 3.1 (Huelsenbeck and Ronquist, 2001; Ronquist and Huelsenbeck, 2003). A general time reversible model with a proportion of invariant sites and a y-shaped distribution of rates across sites (GTR + I + G) model were deduced with hierarchical likelihood ratio tests using Mrmodeltest 2.1 (program distributed by J. Nylander, Evolutionary Biology Centre, Uppsala University). The Bayesian analysis of rbcL genes was initiated with a random starting tree and ran two runs with four chains of Markov chain Monte Carlo (MCMC) iterations simultaneously for 1,500,000 generations, keeping one tree every 100 generations. The first quarter generations (5,000 trees) were discarded as burn in, and the remaining trees were used to calculate a 50% majority rule tree and to determine the posterior probabilities for the individual branches. The 50 between two MCMC runs was below 0.01 for the MCMC iteration, indicating convergence.

Culture Conditions for in Vitro Crossing

For in vitro crosses, ovules of one or two species were excised from placenta in modified Nitsch's medium (Higashiyama et al., 1998) containing 13% (w/v) polyethylene glycol 4000, 1% (w/v) Suc, and 1.3% ultralow-gelling-temperature agarose (agarose type IX-A; Sigma; Higashiyama and Inatsugi, 2006) and cocultivated with a hand-pollinated method described previously for T. fournieri (Higashiyama et al., 1998). When pollen tubes of Torenia species were used with Torenia styles, the ovules were cultivated in an area of 6 × 6 mm² on a coverslip in glass-bottom dish so that the tubes covered the entire area of the ovules. In contrast, when pollen tubes of L. crustacea or L. micrantha species were used with Lindernia styles, the ovules were cultivated in areas of 2 × 2 mm² and 3 × 3 mm², respectively. When ovules of two different species were mixed, the numbers of ovules of each species were adjusted to approximately equal numbers and cultivated at 30°C overnight.

Microscopy

Cultures, excised ovules, and pollen tubes on the stylar canals were observed under an inverted microscope equipped with differential interference contrast (DIC) and an epifluorescence system (IX71, Olympus). Pollen tubes on the stylar canals were stained with aniline blue and observed as described (Higashiyama et al., 1997). All photographs were taken using a 3-CCD camera (C7780, Hamamatsu Photonics) attached to the microscope.

Laser Ablation

A Nd:YAG laser (355 nm; Sigma Koki) was used for laser ablation of cells as described (Higashiyama et al., 2001). The laser beam was focused at the edge of the targeted synergid cells of ovules scattered in the medium. More than 20 ovules were treated in each experiment using one dish.

Micromanipulation of Ovules

For micromanipulation of ovules, water-saturated silicone oil (KF-96, Shin-Etsu Chemical) was layered on the medium to prevent dehydration during the subsequent micromanipulation. Ovules were thrust using a glass needle produced by a glass needle puller (PC-10, Narishige) and moved using a manipulator (MMN-1, MMO-202 N, MMO-220A, Narishige) attached to the inverted microscope. About 20 ovules (each 10 ovules) were manipulated in each experiment using one dish.
Analysis of the Roles of Ca\(^{2+}\) and GABA

For the analysis of Ca\(^{2+}\) culture medium was prepared as described above but without calcium and agarose. Volumes of 500 \(\mu\)L of the liquid medium containing various concentrations of calcium were prepared in 1.5-ml tubes. Three styles, just after pollination, were cut to 10-mm lengths and placed in each tube so that only the cut end was immersed in the medium. After cultivation for 16 h at 30\(^{\circ}\)C, the lengths of the pollen tubes grown in the medium after passing the style were measured using stereomicroscopy. The relationship between the calcium concentration and the pollen tube length was examined to determine the optimal concentration of Ca\(^{2+}\) for pollen tube growth, and this concentration was included in the medium used in the in vitro crossing experiment to examine the effect of Ca\(^{2+}\) on pollen tube attraction.

For the GABA analysis, 2 \(\mu\)M Ca\(^{2+}\) was added to the medium, and the relationship between the GABA (Wako Pure Chemical Industries) concentration and the pollen tube length was examined as for calcium. The optimal concentration of GABA for pollen tube growth was then included in the medium for the in vitro crossing experiment to examine the effect of GABA on pollen tube attraction.

In Vivo Crossing

Pollen grains of each species were applied to Torenia pistils and emasculated pistils of the same species. At 1 d after pollination, ovules (and pollen tubes) were excised from the ovaries and observed in 0.12M sorbitol. When localized pistils of the same species. At 1 d after pollination, ovules (and pollen tubes) were excised from the ovaries and observed in 0.12M sorbitol. When pollen tubes were not observed in an ovary, pollen tubes on the stylar canal were observed following aniline blue staining, as described previously (Higashiyama et al., 1997).

The rbcL sequence data from this article can be found in the DDBJ/EMBL/GenBank data libraries under accession numbers as follows: T. fournieri, AB299084; T. baillonii, AB299885; T. color, AB259806; L. crustacea, AB259807; L. micrantha, AB259808; L. setulosa, AB259809; and L. antipoda, AB259810.

ACKNOWLEDGMENTS

We thank Mr. Shinee Kato (Tsuruoka, Yamagata) and Dr. Hirokazu Tsukaya (University of Tokyo; National Institute for Basic Biology, Okazaki) for providing Lindernia plant materials.

Received May 19, 2006; accepted August 14, 2006; published August 25, 2006.

LITERATURE CITED


Gray-Mitsumune M, Malton DP (2006) The egg apparatus 1 gene from maize is a member of a large gene family found in both monocots and dicots. Planta 223: 618–625


