

Genome-wide analysis of RopGEF gene family to identify genes contributing to pollen tube growth in rice (*Oryza sativa*)

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Abstract

Background: In plants, the key roles played by RopGEF-mediated ROP signaling in diverse processes, including polar tip growth, have been identified. Despite their important roles in reproduction, a comprehensive analysis of RopGEF members has not yet been performed in rice (*Oryza sativa*). To determine whether RopGEF regulators are involved in rice pollen tube growth, we performed genome-wide analysis of this family in rice. **Results:** Phylogenomic and meta-expression analysis of eleven RopGEFs in rice showed that four genes were preferentially expressed in mature pollen. These four genes contain the plant-specific Rop nucleotide exchanger (PRONE) domain and possible phosphorylated residues, suggesting a conserved role in polar tip growth with *Arabidopsis*. In subcellular localization analysis of the four RopGEFs through tobacco (*Nicotiana benthamiana*) infiltration was identified in various cellular organelles, indicating their diverse roles during pollen germination in rice. Moreover, these candidate genes possessing unique cis-acting elements in their promoters compared with the other genes. **Conclusions:** In this study, four RopGEF genes were identified as pollen-specific gene in eleven members of rice, and the expression pattern, promoter analysis, and evolutionary relationship of the RopGEF family were studied compared with *Arabidopsis*. Our study indicated that four RopGEF genes might function during pollen germination in distinct subcellular localization. Our study could provide valuable information on the functional study of RopGEF in rice.

Background

Rho-type GTPases of plants (ROP), also known as RACs, are a plant-specific subfamily of Rho small GTP-binding proteins [1] and participate in diverse signal transduction processes, including disease resistance, pollen tube growth, root hair development, reactive oxygen species (ROS) production, cell wall patterning, and hormone responses [2]. Similar to Rho family proteins in other eukaryotic cells, ROPs regulate endocytosis and exocytosis through cytoskeleton organization and intracellular kinase cascades through activation of NADPH oxidase, processes that are key for cell polar growth in plants [3-6]. Activation and inactivation of ROP, by change in its conformation following GTP binding and hydrolysis (GDP-binding form), functions as a molecular switch in signal transduction [6, 7]. Conversion from the GDP and GTP form of Rho GTPases is catalyzed by guanine nucleotide exchange factors (GEFs). Plant-specific RopGEFs spatiotemporally regulate the activity of ROPs/Racs through the catalytic plant-specific ROP nucleotide exchanger (PRONE) domain [8].

Although few studies of *RopGEFs* have been carried out in other plant species, recent studies in the model plant *Arabidopsis* support their roles in various plant development processes as well as in defense. *Arabidopsis* has 14 *RopGEFs* in its genome, with a high degree of sequence similarity [8, 9]. Functional studies showed that AtRopGEF1 and AtRopGEF4 specifically regulate ROP11 in abscisic acid (ABA)-mediated stomatal closure [10], and that AtRopGEF1 plays a role in controlling lateral root growth [11] and polar auxin transport to achieve cell polarity during early plant development [12]. Among them, seven members are specifically or highly expressed in pollen tubes and function redundantly [8, 13]. Pollen-specific RopGEFs have conserved C-termini, which function by auto-inhibition of the PRONE domain [13].

Recent findings suggest that pollen-specific receptor-like kinases (PRKs) transduce ROP signaling via phosphorylation of RopGEFs at the C-terminus to regulate polar pollen tube growth [13-17]. When some ligands are received by receptor-like kinases (RLK), RopGEFs are activated and convert GDP to GTP, which binds to ROP [18]. AtPRK2 promotes ROP1 via phosphorylation of RopGEFs in the control of polarized pollen tube growth [16]. Interaction of AtPRK6 with AtRopGEF8 and AtRopGEF12 plays essential roles in polarized growth of pollen tubes [17]. Another RLK, FERONIA, belonging to the *Catharanthus* receptor-like kinase (CrRLK) family, was shown to function as an upstream regulator of RopGEFs, mediating auxin efflux on root hair growth in Arabidopsis [19-21].

In rice (*Oryza sativa*), among the first 12 reported genes, 11 *OsRopGEFs* encoded full-length proteins [22]. *OsRopGEFs* were shown to play a role in the activation of *OsRac1* in the disease-resistance response [23]. *OsRac1* negatively regulates cell death and innate immunity by production of ROS [24, 25]. It was shown that *OsRac1* activates plant NADPH oxidase, known as OsRbohB (respiratory burst oxidase homolog), by direct interaction [26]. In addition, *OsRopGEF10* activates the development of small cuticular papillae on leaf surfaces [27], whereas *OsRopGEF7B* is involved in regulating floral organ development [28]. Compared with Arabidopsis, the molecular function of rice RopGEF members in response to plant developmental cues is poorly understood.

To reveal whether roles in polar tip growth in rice exist for the conserved RopGEFs, we investigated in the present study the phylogenetic relationship of rice and Arabidopsis RopGEF proteins, analyzing the PRONE domain and characterizing the amino acid residues. We identified four pollen-preferred *OsRopGEF* genes through meta-expression analysis using our pollen/anther database, followed by confirmation by qRT-PCR. In addition, we performed subcellular localization of each RopGEF through tobacco infiltration and *cis*-element analysis of the four pollen-preferred *OsRopGEF* genes, revealing a conserved element. Based on our findings, we propose a model for the *RopGEF* role in rice pollen tube growth.

Results

Expression profiling analysis of *RopGEF* genes

To identify the pollen-preferred genes of rice, we examined a publicly available Affymetrix rice microarray data [29]. Expression patterns conserved between *indica* and *japonica* rice varieties at various developmental stages of anthers and pollen, and anatomical meta-expression data from our established database, Rice Anther Pollen Expression Database (RAPED) (<http://ricephylogenomics-khu.org/RAPED/home.php>), were analyzed. We then constructed a phylogenetic tree by aligning the protein sequences of all *OsRopGEFs*. Meta-expression data in the context of the phylogenetic tree were combined to check the function of *OsRopGEFs* with respect to anther and pollen development, compared with other tissues/organs (Fig. 1A). As a result, the rice *RopGEF* gene family can be divided into two subfamilies, one group to which *OsRopGEF2, 3, 4, 6, and 8* belong, and the other to which the remaining six genes belong, namely *OsRopGEF1, 5, 7, 9, 10, and 11*.

In addition, we identified that *OsRopGEF2*, *OsRopGEF3*, *OsRopGEF6*, and *OsRopGEF8* showed selective expression at the stage of late pollen development, i.e., at stages of tricellular pollen grains, mature pollen (MP), and germinated pollen (GP). These four genes were used for further analyzes. Since *OsRopGEF3* was also highly expressed in root hairs, we predicted that this gene could affect tip growth (Additional file 1). Genes *OsRopGEF4* and *OsRopGEF7* showed some expression in the early stages of anther development, although their expression levels decreased during the later stages. Furthermore, because expression is also detected in tissues other than anther and pollen, these two genes cannot be considered to be pollen-preferred expressed genes. In addition, *OsRopGEF5* showed weak expression in *japonica* rice anthers and other tissues, showing no preferred expression in anthers and pollen.

Next, to verify the meta-expression data, we performed quantitative real-time PCR (qPCR) using ten tissues: shoot, root, leaf blade, panicle, flower, seed, tetrad microspore, young microspore, vacuoles, and MP (Fig. 1B). By sampling anthers at various developmental stages, from pollen mother cell to MP and GP, we tried to determine at which stage the target genes were being expressed. As a result, *OsRopGEF2*, *OsRopGEF3*, *OsRopGEF6*, and *OsRopGEF8* were shown to be highly expressed in MP. *OsRopGEF6* showed the highest level of expression in MP, and *OsRopGEF2* and *OsRopGEF8* showed similar expression levels in MP. These native pollen-preferred *OsRopGEFs* were generally not expressed in other tissues (shoot, root, leaf blade, panicle, flower, seed, tetrad, microspore, vacuolated pollen stage). Therefore, these four genes may play key roles in pollen tube growth and other processes that occur after pollen maturation.

Comparative analysis of the conserved domains in RopGEF genomic and protein sequences between rice and Arabidopsis

Genes *OsRopGEF2*, *OsRopGEF3*, *OsRopGEF6*, and *OsRopGEF8* encode proteins with similar amino acid sequences. *RopGEFs* are considered to be a novel gene family with a unique structure containing the PRONE catalytic domain that is exclusively found in the plant *RopGEF* gene family [9]. The PRONE domain is known to be required for RopGEFs to convert GDP to GTP. The PRONE domain of RopGEFs is highly conserved in all *OsRopGEFs*. More interestingly, the pollen-preferred *OsRopGEFs* were found to retain the C-terminus region after the PRONE domain (Fig. 2), whereas the pollen-non-preferred *OsRopGEF1*, *OsRopGEF9*, and *OsRopGEF11* lack this C-terminus.

For the comparative analysis of *RopGEF* gene families, we collected the protein sequences of 14 *AtRopGEFs* and 11 *OsRopGEFs* and constructed a Neighbor-joining tree (Fig. 2). Four *OsRopGEF* genes showing pollen-preferred expression were clustered together with *AtRopGEF8* and *9*, which also exhibited pollen-preferred expression. In the Arabidopsis *RopGEF* family, the expression patterns revealed by qPCR were not consistent with those from previous reports: *AtRopGEF1*, *8*, *9*, *12*, *14* were detected in pollen tissues [8] and *AtRopGEF8*, *9*, *10*, *11*, and *13* showed pollen-preferred expression, respectively, and, of these, the latter data corresponded more closely with the transcriptome data in Genevestigator (Additional file 2). In the phylogenetic tree, rice members with pollen-preferred expression were closely clustered with those in Arabidopsis, and the C-terminal amino acid sequences were more similar than those for other

members in rice. The invariant serine residue (S510, numbered as in AtRopGEF12) within the C-terminus, which is important for C-terminal inhibition [13] was also located in pollen-preferred *RopGEF* members in rice (Additional file 3), supporting the hypothesis that phosphorylation-regulated GEF activity was conserved.

Protein structure

Protein sequence analysis and hydropathy plot profiling revealed that AtRopGEF and OsRopGEF, both with pollen-preferred expression, had mostly similar amino acid sequences but differed at the N- and C-termini (Fig. 3). The analyzed rice RopGEF 3-D models exhibit a butterfly-shaped three-dimensional structure in which two PRONE protomers dimerize via their N-terminal groups (Fig. 4). Each protomer consisted of two subdomains, with the subdomain 1 containing the WW-loop, which is a common characteristic in Arabidopsis [7] (Table 1). To more precisely characterize the 3-D model structures, we compared the pollen-preferred rice RopGEF 3-D models with those from Arabidopsis. The subdomains differed in the number of alpha helices and the number of residues in the WW-loop between the OsRopGEFs and the AtRopGEFs. Rice and Arabidopsis RopGEF 3-D models displayed trends in the number of α -helices from 14 to 17, and in the number of residues in the WW-loop from 25 to 50.

OsRopGEFs contained 14 to 15 alpha helices, and 25 to 40 residues in the WW-loop, whereas AtRopGEFs contained 14 to 17 alpha helices, and 38 to 50 residues in the WW-loop. The main difference between OsRopGEFs and the reported AtRopGEF [30] is in the WW-loop, where AtRopGEF displayed α -helices or beta-strands in the WW-loop, whereas OsRopGEFs did not contain α -helices or beta-strands.

Subcellular localization of four RopGEF proteins

The RopGEF is known to regulate ROP signaling in the plasma membrane (PM). Arabidopsis RopGEF8, RopGEF9, and RopGEF14 were shown to be localized to the apical PM of the pollen tube, and RopGEF1 to the entire PM. On the other hand, RopGEF12 was barely detected in the PM, being present in the cytosol of the pollen tube [8], suggesting that the different localizations of RopGEF in pollen tubes represented differences with respect to multiple regulatory ROP signaling actions. Therefore, we examined the subcellular localization of OsRopGEF members to check whether their spatial locations were the same or different. We introduced GFP fusion proteins controlled by the CaMV 35S promoter into the epidermal cells of tobacco leaves. As shown in Fig. 5, OsRopGEF3 and OsRopGEF6 were specifically localized into the PM, consistent with the common RopGEF location, while the GFP signals of OsRopGEF2 and OsRopGEF8 were predominantly found in the cytosol as well as in the PM. The GFP signal of OsRopGEF2, located in the cytosol, was associated with the endoplasmic reticulum (ER), rather than the nucleus, while the OsRopGEF8 signal was strongly associated with the nucleus. The results suggest that the four pollen-preferred members of OsRopGEF could carry out unique functions during pollen germination, as reflected by their distinct subcellular locations.

Promoter analysis of pollen-preferred genes

Up to now, the information about the roles of transcription factors and regulatory motifs in pollen germination in rice has been unclear. Based on the microarray data and the qPCR results, we performed a promoter analysis to identify the *cis*-elements conserved in the promoters of *OsRopGEFs* associated with pollen-preferred expression.

We searched the known *cis*-regulatory elements (CREs) in the promoter based on existing research (Additional file 4). Major pollen-preferred CREs such as POLLEN1LELAT52 (AGAAA), PB Core (CCAC), and GTGANTG10 (GTGA) [31] were identified. In the case of POLLEN1LELAT52, 12 CREs existed in the *OsRopGEF3* promoter and seven CREs existed in the *OsRopGEF8* promoter. In the case of GTGANTG10, 11 CREs existed in the *OsRopGEF8* promoter. In the case of the PB Core, there were five copies in the *OsRopGEF2* promoter, and four copies in each of the *OsRopGEF6* and *OsRopGEF8* promoters. However, in the case of the remaining *OsRopGEF* family genes, there were also many pollen-preferred CREs present in the promoters of genes which exhibited little or no expression in pollen. On average, there were 4.3 copies of POLLEN1LELAT52, 6.4 copies of GTGANTG10, and 1.6 copies of the PB Core in the promoters of the seven *OsRopGEFs* compared with the four highly pollen-preferred genes. We calculated the p-values to determine how many of the four highly pollen-expressed *OsRopGEF* genes had significantly more CREs relative to the rest of the *OsRopGEF* genes, but all did not exceed 0.01. Based on these results, we assume that known pollen CREs do not regulate the expression of the four pollen-preferred *OsRopGEFs*.

We then searched to find CREs that existed in the promoters of only pollen-preferred *OsRopGEFs*. Firstly, we searched for motifs that were common to the promoters of the four pollen-preferred genes, using MEME (Fig. 6). We downloaded the 2000-base pair upstream sequences of the four *OsRopGEF* genes and found ten conserved elements. Among them, three CREs were absent from the promoters of the other, pollen-non-preferred *OsRopGEF* genes. These three CREs were present at -2000 to -1500 bp from the transcription start sites of *OsRopGEF3* and *OsRopGEF6*, and mainly existed within -1000 bp of the transcription start sites of *OsRopGEF2* and *OsRopGEF8*. All CREs were present once in the promoter of each gene, except for the first CRE (presented as a dark blue box; Fig. 6), which was present twice in the promoter of *OsRopGEF2*. We predicted that these three CREs will be involved in pollen expression. Next, using TOMTOM, we revealed that these elements were present in transcription factor genomic sequences such as MYB and bHLH, indicating that the four *OsRopGEFs* could be regulated in pollen by transcription factors such as MYB, and bHLH, for example.

Discussion

In animal and plant cells, a wide range of key cellular processes that require the establishment of cell polarity are governed by Rho GTPases or ROPs [6, 32, 33]. ROPs transduce intracellular and extracellular stimuli in a spatially and temporally regulated manner, resulting in localized regulation of intracellular responses [6]. The number of ROP proteins varies among plant species: seven in rice, nine in maize, and 11 in Arabidopsis. In Arabidopsis, three *ROPs* (*ROP1*, *ROP3*, and *ROP5*) are highly expressed in pollen and

are functionally redundant in the regulation of pollen tube growth. AtROP1 is localized to the apical region of the pollen tube and is dynamically regulated by endocytosis during pollen tube growth [34, 35]. Disruption of *AtROP1* by overexpression of the constitutively active form resulted in swollen tubes whereas disturbance of AtROP1 activation resulted in short pollen tubes [36, 37], indicating that the control of ROP activity level is important for pollen tube shape. By contrast, there was no pollen-preferred ROP member in rice, as analyzed by the heatmap of tissue expression (Additional file 5a). Instead, *OsRac6*, a phylogenetically close member of *AtROP1/3/5* (Additional file 5b and c), was expressed in various tissues including mature anthers and pollen, suggesting that control of ROP activity through control of one member, *OsRac6*, might be important in rice pollen germination.

To achieve regulation of ROPs, plants might have evolved specific regulators such as RopGEFs [9]. As a result of comparative expression analysis, we found that four and five genes were highly expressed in pollen among 11 rice and 14 Arabidopsis *RopGEF* family members, respectively. This suggests that the ROP regulatory mechanism by RopGEF in pollen might be conserved between rice and Arabidopsis. Quadruple mutants of *AtRopGEF1*, *AtRopGEF9*, *AtRopGEF12*, and *AtRopGEF14* showed reduced pollen tube elongation, though this effect was not exhibited by any single gene mutation [16]. Due to functional redundancy, single mutants showed no defects in pollen germination and tube growth. Knock-out mutants of *OsRopGEF7B* were ubiquitously expressed in different tissues, including pollen, causing abnormal development of floral organs but not altered with respect to pollen formation or pollen germination [28]. Instead, *OsRopGEF2*, *OsRopGEF3*, *OsRopGEF6*, and *OsRopGEF8* were preferentially expressed in pollen, strongly suggesting their significant roles in pollen germination and tube growth. Therefore, in future studies, we would expect to see defects during pollen tube growth by generating knock-out mutants for multiple *OsRopGEF* genes which exhibited pollen-preferred expression. The pollen-preferred expression pattern of the four genes might be explained by the existence of the three *cis*-acting elements absent from promoters of the other seven, pollen-non-preferred members. Since these *cis*-acting elements also exist in promoters of regulatory genes such as *MYB* and *bHLH*, we expect that either MYB or bHLH would be the potential transcription factors regulating the expression of the *RopGEFs*. However, results from the MEME-suite are required to demonstrate the binding site of the *cis*-acting elements in promoters of the transcription factors, through further analysis such as the use of the yeast-one hybrid system.

Despite of variable isoforms, gene sequences and protein structures are well conserved throughout AtRopGEFs and OsRopGEFs, especially with respect to the PRONE domain, indicating the conserved activity of the RopGEFs. The differences in the number of alpha helices in the RopGEF 3-D structure may be involved in stabilizing the WW-loop conformation [38]. The 3-D structures of OsRopGEF2 and 6 display the same length of the WW-loop (40 residues) compared with the reported AtRopGEF8 [30], although OsRopGEF3 and 8 exhibit shorter lengths, 34 and 25 residues, respectively (Table 1). The WW-loop is known to be variable in its length and sequence among different members of the RopGEF family [8, 9, 30, 38]. We found that the phosphorylatable serine residue [13] was also conserved in the pollen-preferred OsRopGEFs as in the AtRopGEFs. Further investigations will be required to determine how the four pollen-preferred OsRopGEFs are regulated by interacting partners such as the pollen-preferred RLKs [17, 39].

Our report of differential distribution of OsRopGEFs in tobacco epidermal cells indicates that they could regulate different processes of pollen germination. In Arabidopsis, it was observed that RopGEFs interacted differentially with the GTP- and GDP-bound forms of ROP1 and were localized in different subcellular locations [8]. Deletion of the C-terminal domain in AtRopGEF12 abolished the membrane association when it was expressed in tobacco pollen [8]. RopGEFs probably activate ROPs in the PM, and their spatial distribution reflects the sites of ROP activation [6]. It was suggested that the apical location of ROP1 in turn feed-forward regulates exocytosis of RLK and RopGEFs [5]. Another study reported that the phosphorylation of the PRONE domain of RopGEFs by AGC kinase was critical for the localization of RopGEFs to the apical PM, and subsequently for ROP activation and pollen tube polarity [40]. Cellular ROP signaling is also involved in root hair formation, another polar tip-growing plant cell type. Recently, functional analysis revealed that root-hair-preferred AtRopGEF3 functions in root hair initiation by AtROP2 polarization, while another one, AtRopGEF4, regulates subsequent root hair growth [41]. Therefore, it is possible that RopGEF members can be regulated differently by kinases to achieve different localizations and activations, and subsequently function in different phases of polar cell initiation and growth. Further studies of the dynamic distribution of OsRopGEF by other signals or interacting partners would provide a better understanding of the molecular network necessary for the process of rice pollen germination.

Based on our results and other reports, we were able to draw a RopGEF model in a pollen cell (Fig. 7). *RopGEF* is expressed in pollen due to the action of transcription factors such as bHLH and MYB. Pollen-preferentially expressed RopGEFs could be regulated for the activity through RLK toward serine residues. When activated, it can be present in various positions such as the cytosol, PM, and ER. The PRONE domain of the RopGEF protein converts the GDP attached to ROP2 to GTP. This process activates ROP and regulates the other proteins, such as RBOH, to affect pollen tip growth or pollen germination. To further test this model, a binding assay between transcription factors and the *RopGEF* promoter, and observation of the changes in the expression of subpopulations of *RopGEF* or *ROP* mutant plants will be required.

Conclusion

Our study identified four genes that are highly expressed in pollen among RopGEF genes in rice, and confirmed their conserved domain in protein sequence, similarity in protein secondary and tertiary structure, and identified each subcellular localization. We also found a novel cis-regulatory element that is expected to affect pollen expression by promoter analysis and present RopGEF model in the pollen cell. Our study could provide valuable information on the functional study of RopGEF in rice.

Methods

Multiple sequence alignment and phylogenetic tree construction

To perform a phylogenetic analysis of RopGEF and ROP/Rac in rice and *A. thaliana*, we collected the protein sequences with locus ID from the Rice Genome Annotation Project

(<http://rice.plantbiology.msu.edu/>), the National Center for Biotechnology Information (NCBI, <https://www.ncbi.nlm.nih.gov/>) and the Phytozome platform (<https://phytozome.jgi.doe.gov/pz/portal.html>) (Additional file 1). Multiple amino acid sequences were aligned using ClustalW [42]. The phylogenetic analysis was performed using MEGA 7.0.26 under maximum likelihood and neighbor-joining methods [43].

Meta-expression analysis

We used a publicly available rice Affymetrix microarray data set prepared from anthers and pollen from the NCBI GEO [44] to identify late-pollen-preferred genes. To examine these data, we used the Affy package encoded by R language to normalize the signal intensity and then transformed them into \log_2 values. The normalized data, with averaged Affymetrix anatomical meta-expression data, were then used for further investigations, e.g., heatmap construction, and identification of the late-pollen-preferred genes [45].

RNA extraction, cDNA construction, and real-time PCR

Tissues, including pollen of rice (*O. sativa* cv. Dongjin) grown in paddy fields, were frozen in liquid nitrogen and ground with a Tissue-Lyser II (Qiagen; Hilden, Germany). RNA was extracted with TRIzol buffer, and cDNAs were synthesized using SuPrimeScript RT premix from GeNet Bio [46]. For tissue-specific expression by qPCR, we used control primer sets for rice ubiquitin 5 (*OsUbi5*, LOC_Os01g22490). All of the qRT-PCR primers we have used in our experiments are listed here (Additional file 2). The PCR cycle conditions used were 95°C for 30 s, 57°C for 30 s, and 72°C for 1 min 30 s for 22–38 cycles. For real-time PCR (qPCR), we used cycling conditions of 95°C for 15 s, 57°C for 30 s, and 72°C for 60 s using Roter-Gene Q instrument system (Qiagen, Hilden, Germany). We used 2x Prime Q-Mastermix (GeNet Bio) which contains SYBR Green1 for qPCR buffer. To determine the significant expression changes of four *RopGEF* genes in different tissues, we performed qPCR analysis with three independent biological replicates. Relative transcript levels and fold change were calculated by the $2^{-\Delta C t}$ and $2^{-\Delta \Delta C t}$ methods, respectively [47].

Cis-acting elements analysis

To analyze the pollen-specific promoter regions of four *OsRopGEF* loci, we extracted 2-kb upstream sequences from the start codon of 11 *OsRopGEF* loci from EMBL (https://plants.ensembl.org/Oryza_sativa/Info/Index) and Gramene (http://ensembl.gramene.org/Oryza_indica/Info/Index). To find the *cis*-elements in the promoter sequences of the *OsRopGEF* genes, the 2-kb upstream sequence was scanned using the PLACE (Plant *cis*-acting regulatory DNA elements) database [48]. To discover novel motifs, which exist only in pollen-specific RopGEF genes, we performed Multiple Em for Motif Elicitation (MEME)-suite searches with those

sequences in the FASTA format via the Web server hosted by the University of Queensland (<http://meme-suite.org/tools/meme>) [49]. First, we analyzed the 2-kb promoter sequence of all 11 *RopGEF* genes at once. Next, the 2-kb promoter sequences of the four pollen-specific *RopGEF* genes were analyzed, and we compared the two data sets using MEME. Among the motifs discovered, we chose those that were found only in the promoter sequence of pollen-specific *RopGEF* genes and searched for some motifs in databases of known motifs, using TOMTOM. The search conditions were “motif length 5 to 10bp,” “indicating only the motifs commonly found in the input data,” and “checking both strands.”

Subcellular localization analysis

The coding sequences (CDSs) of four *RopGEF* genes were amplified from mature anther cDNA and cloned into pGreen vector fused with C-terminal green fluorescence protein (GFP). All of the cloning primers we have used in our experiments are listed here (Additional file 2). The constructs were transfected into *A. tumefaciens* strain GV3101 and used for *Nicotiana benthamiana* infiltration as described by [50]. Two to three days after infiltration, GFP fluorescence was observed with a confocal laser scanning microscope (Zeiss LSM 510, Jena, Germany) with spectral settings of 500–530 nm for emission and 488 nm for excitation.

Protein structure analysis

The three-dimensional (3-D) structural models were computed on a SWISS-MODEL Workspace in a fully automated mode [51] using *RopGEF* amino acid sequences as templates. The obtained 3-D structures were visualized using the University of California, San Francisco (UCSF) Chimera 1.10 program [52].

Additional Files

Additional file 1: Figure S1. Meta-expression analysis and genome-wide identification of every *OsRopGEF*. *OsRopGEF* expression was observed in all tissues except pollen. Yellow color in the heatmap indicates high level of expression; dark blue, low level of expression. Numeric values indicate the average of the normalized log₂ intensity of microarray data.

Additional file 2: Figure S2. Meta-expression analysis of entire *AtRopGEF* genes. The heatmap was prepared using the Genevestigator. We chose five representative tissues, including pollen. It revealed that five *Arabidopsis* *RopGEFs* were highly expressed in pollen. The dark red color of the heatmap indicated the highest expression; white color, lowest expression.

Additional file 3: Figure S3. Protein sequence alignment domain analysis and conserved phosphorylated amino acid residues of C-termini of *RopGEF* genes. Every *OsRopGEF* and *AtRopGEF* protein sequence was collected and aligned, followed by the PRONE domain (C1, C2, C3) and the WW-motif. At the end part of the sequence, we found some conserved regions. According to previous studies, S510 in the C-terminus of *AtRopGEF12* is involved in the C-terminal inhibition of GEF activity. As in *AtRopGEF12*, the

serine residue is conserved in each of the OsRopGEF2, OsRopGEF3, and OsRopGEF8 genes but not in OsRopGEF6. However, K (Lysine) can also be phosphorylated.

Additional file 4: Figure S4. Promoter analysis of each OsRopGEF gene. Specific-CREs known to affect expression in pollen were identified using PLACE database, namely three pollen-CREs and the TATA box. The number above the yellow bar shows the upstream position of the promoter base pair when taken at +1 of ATG. We analyzed up to the upstream 2,000-base pair. The p-values indicate how significantly the four genes exhibiting high expression in pollen differed from the other seven genes.

Additional file 6: Table S1. Accession numbers of RopGEF genes and RAC/ROP genes.

Additional file 7: Table S2. OsRopGEF isogene-specific primers for qPCR and pGREEN cloning.

Abbreviations

CRE: cis-regulatory elements; CrRLK: Catharanthus receptor-like kinase; Dbl: diffuse B-cell lymphoma; GTP: guanosine triphosphate; GP: germinated pollen; MP: mature pollen; ROP: Rho-type GTPases of plants; RopGAP: GTPase-activator protein for ROP; RopGEF: guanine nucleotide exchange factors for ROP; ROS: reactive oxygen species; PM: plasma membrane; PRK: pollen receptor-like kinase; PRONE: plant-specific ROP nucleotide exchanger;

Declarations

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Availability of data and materials

The datasets supporting the conclusions of this article are included with in the article and its Supplementary files.

Authors' contributions

KHJ and YJK designed the research and acquire funding; KHJ, YJK, EJK, and SWP co-wrote the manuscript and figure arrangement. YJK, EJK, SWP, WJH, and JS performed the experiments and analyzed the data.

Ethics approval and consent to participate

Not applicable.

Consent for publication

Not applicable.

Competing interests

The authors declare that they have no competing interests.

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Figures

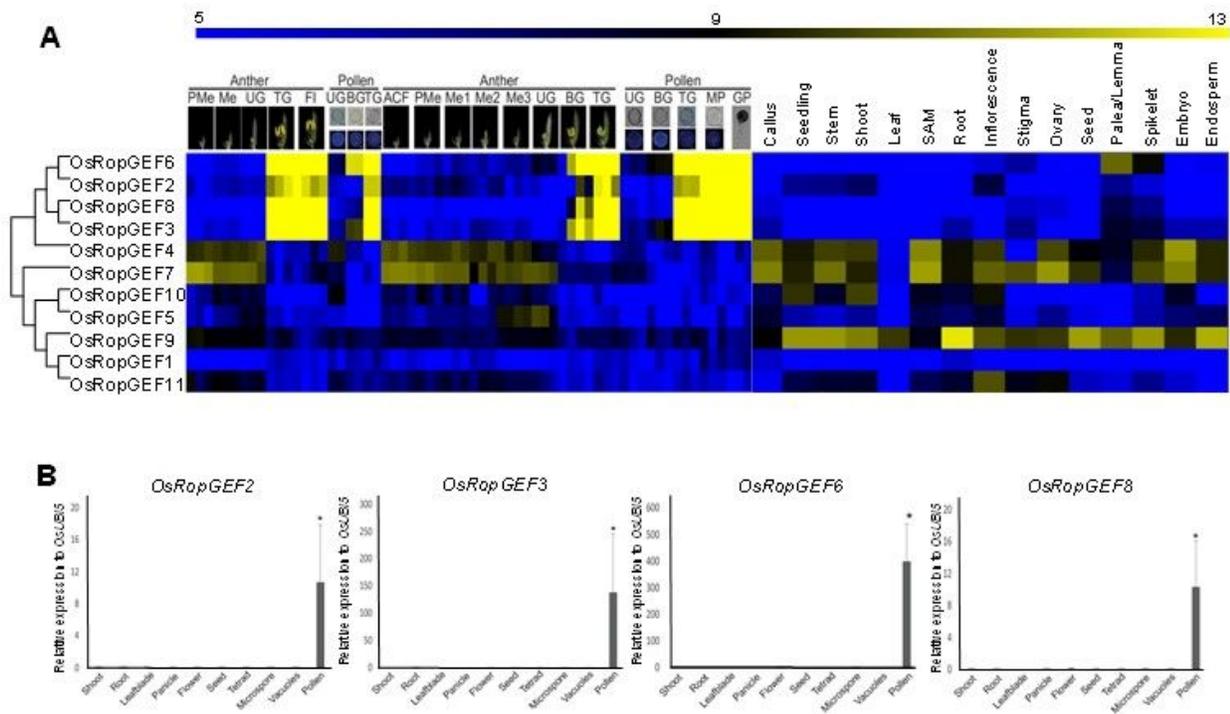


Fig. 1

Figure 1

Meta-expression analysis and genome-wide identification of OsRopGEF and validation of meta-expression patterns of pollen-specific OsRopGEFs using qPCR. (A) Heatmap expression and phylogenetic analysis of 11 OsRopGEF genes revealed four genes are specifically expressed in MP. 22 microarray data for indica rice including five stages for anthers and three for pollen and 42 microarray data for japonica rice including eight stages for anthers and five for pollen were used. ACF, archesporial cell-forming stage; BG, bi-cellular gametophyte stage; FI, flowering stage; GP, germinate ng pollen; Me, meiotic stage; Me1, meiotic leptotene stage; Me2, meiotic zygotene-pachytene stage; Me3, meiotic diplotene-tetrad stage; MP, mature pollen stage; PMe, pre-meiosis; TG, tricellular pollen stage; UG, uni-cellular gametophyte stage. Yellow color in heatmap indicates high level of expression; dark blue, low expression. Numeric values indicate an average of the normalized log₂ intensity of microarray data. (B) Expression of pollen-preferred OsRopGEF genes was analyzed by qPCR in various tissues of rice. Rice ubiquitin 5 (OsUbi5, LOC_Os01g22490) was used as an internal control. Y-axis, expression level relative to OsUbi5; X-axis, samples used for analyses. Error bars represent the standard errors of three biological replicates.

Significant differences are indicated by asterisks, *, p-value < 0.01. Data were analyzed by employing one-way ANOVA with repeated measures, using Tukey's pairwise comparison test.

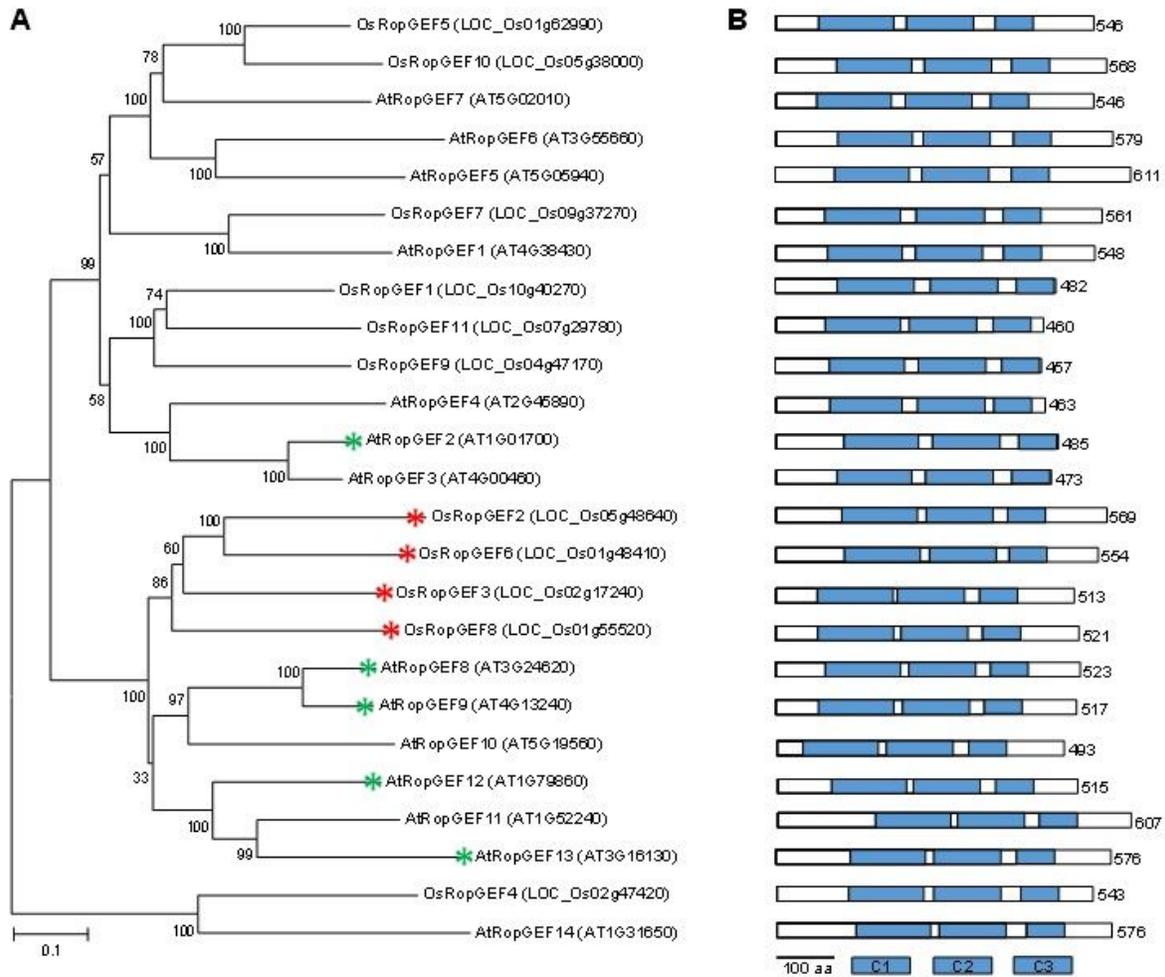


Fig. 2

Figure 2

Phylogenetic analysis of RopGEF family from *Oryza sativa* and *Arabidopsis thaliana*. (A) The phylogenetic tree was constructed by annotating the protein sequences for each gene by using MEGA7. Protein sequence alignment was performed by ClustalW and phylogeny by Neighbor-joining methods, with the reliability being estimated by the 1000 times bootstrap test. The number between the trees represents the bootstrap value. Os, *Oryza sativa*; At, *Arabidopsis thaliana*. Red and green asterisks indicate RopGEF genes with high expression in pollen in *O. sativa* (in this study) and *Arabidopsis thaliana* [8], respectively. (B) A scheme of protein structure. We arranged the sequence of the protein domain scheme to be the same as the gene sequence of the phylogenetic tree of A. Length of each RopGEF protein is shown on the right and its unit is aa (amino acid). C1, C2, and C3 (blue boxes) indicate position and length of the conserved PRONE subdomain of the RopGEF family.

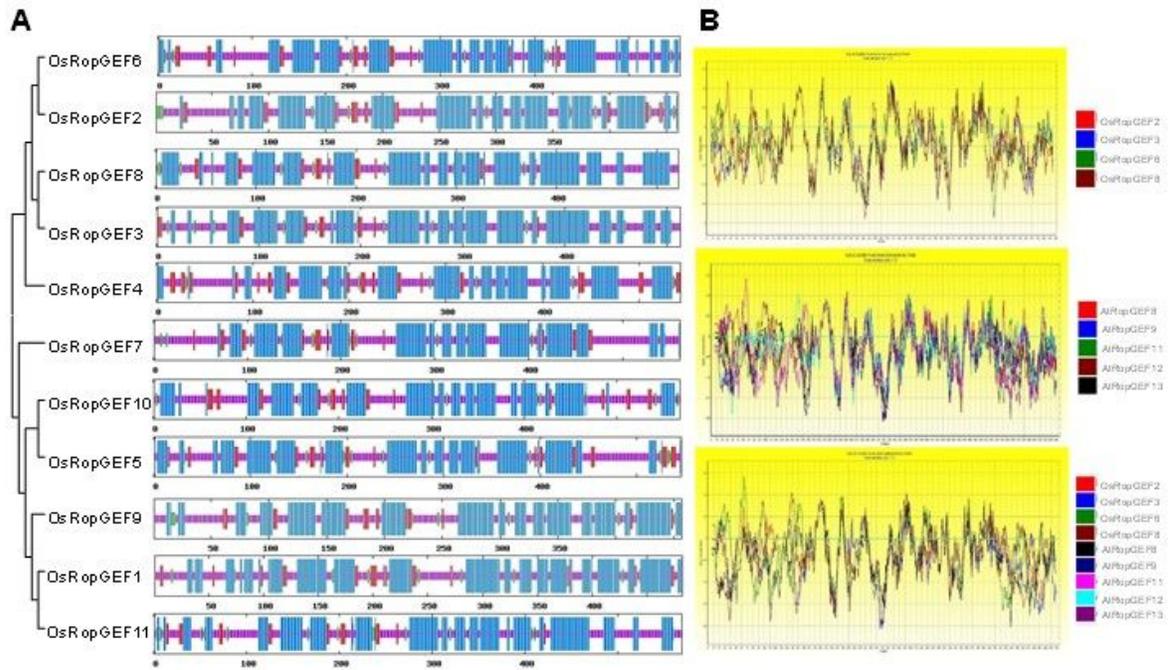


Fig. 3

Figure 3

Protein sequence analysis. (A) Secondary structure of RopGEF using the deduced amino acid sequences from OsRopGEFs. Blue lines indicate alpha helices, purple lines indicate random coils, red lines denote extended strands, and green lines represent beta turns. (B) Hydropathy plot analysis. The upper part of the figure is the hydropathy plot of four OsRopGEFs which showed pollen-specific expression, and the central part is a hydropathy plot of five AtRopGEF genes that are pollen-specific expressed in the Genevestigator. The last figure is the sum of four OsRopGEF and five AtRopGEF hydropathy plots. Profiles of rice and Arabidopsis RopGEFs showed similarities in the PRONE domain, but not for the N and C-termini.

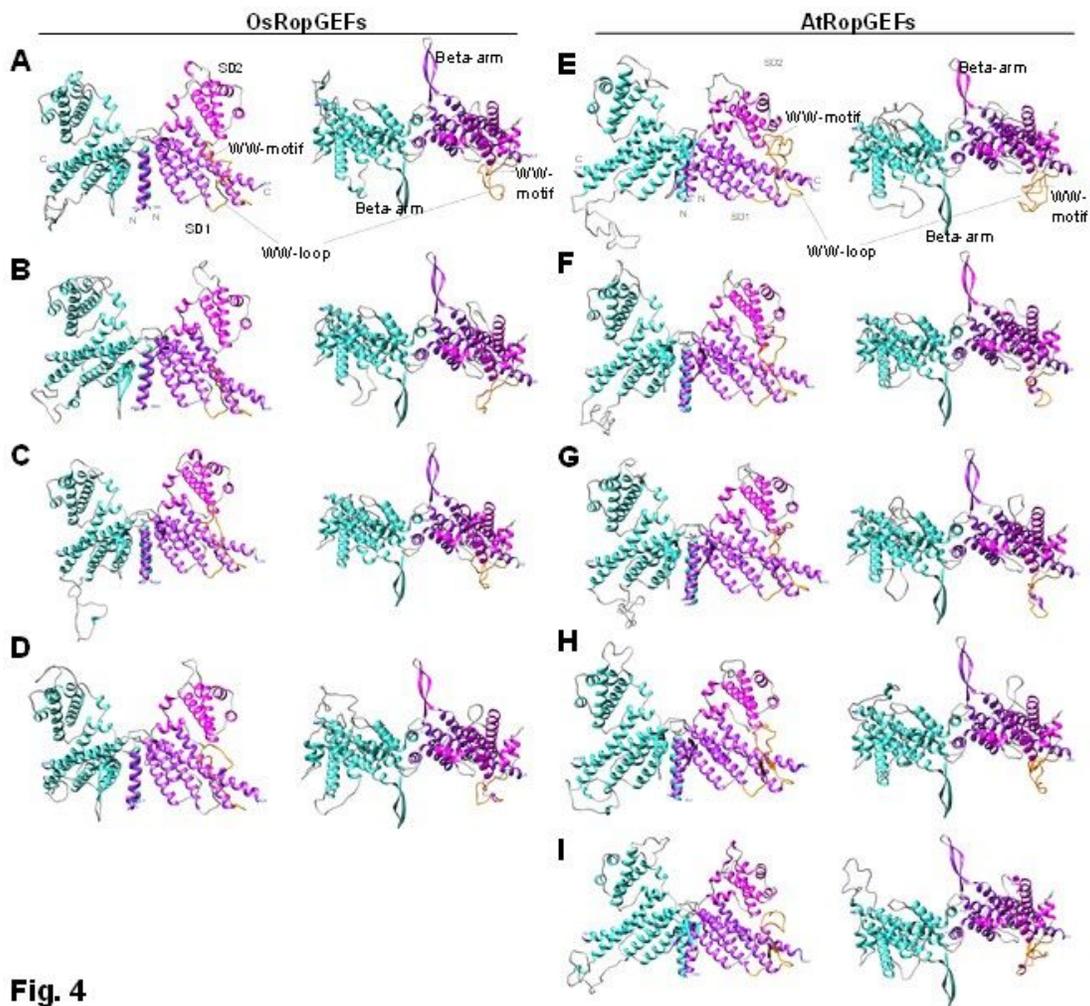


Fig. 4

Figure 4

Predicted 3-D structures of RopGEF genes in rice and Arabidopsis. (A) OsRopGEF2, (B) OsRopGEF3, (C) OsRopGEF6, (D) OsRopGEF8, (E) AtRopGEF2, (F) AtRopGEF8, (G) AtRopGEF9, (H) AtRopGEF12, and (I) AtRopGEF13. Comparative representation was performed using the UCSF Chimera package. Coil, left-promoter, subdomain 1, subdomain 2, WW-loop, and WW-motif structures are depicted in gray, sky-blue, purple, magenta, orange, and red, respectively. N- and C-termini are depicted in blue.

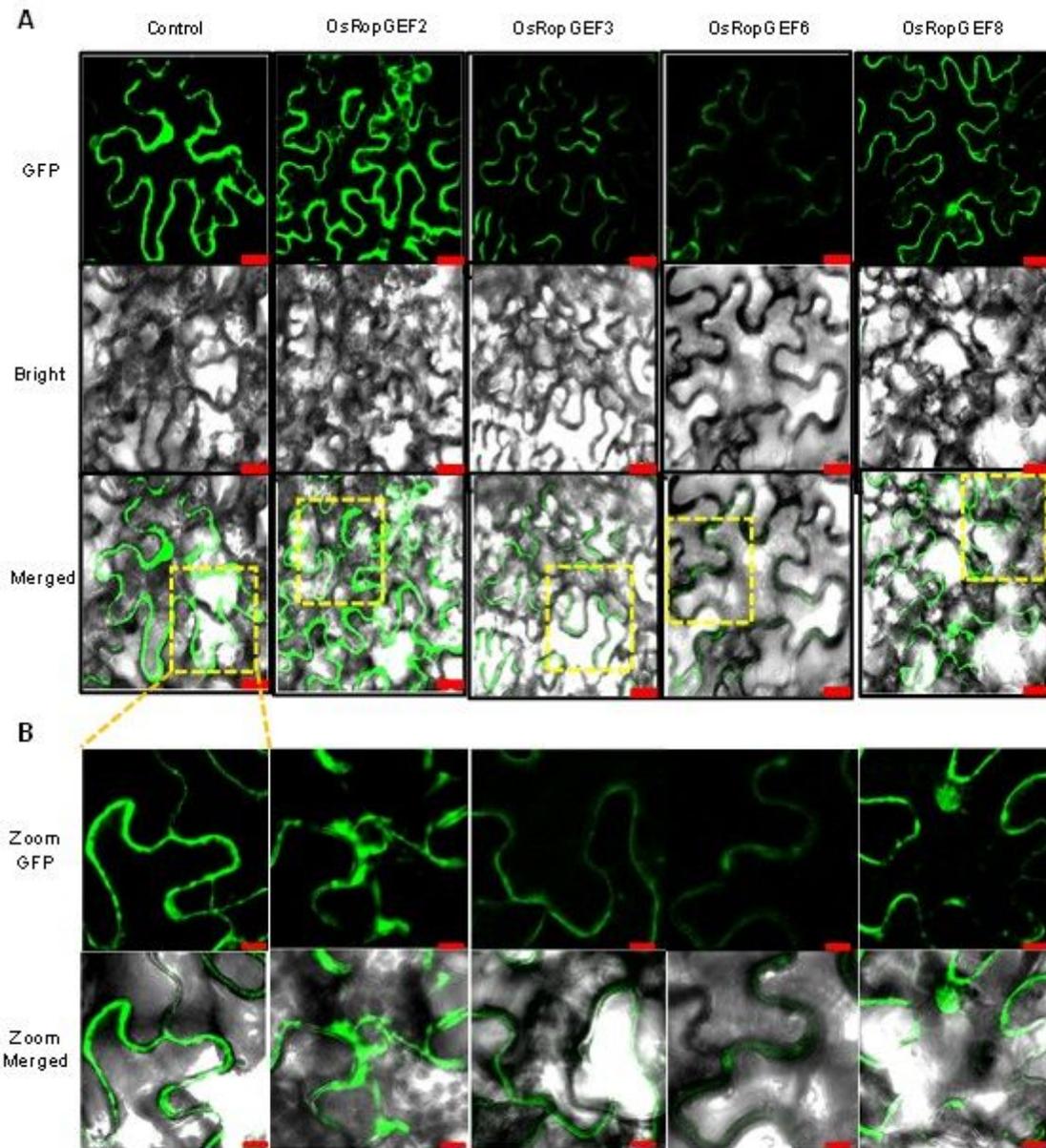


Fig. 5

Figure 5

Subcellular localization observation of OsRopGEF proteins in tobacco plants using laser scanning confocal fluorescence microscopy. (A) Control-GFP, and four pollen-specific expressed OsRopGEF GFP signals were observed in tobacco epidermal cells in the upper panel, the center panel showed the bright field image, and the lower panel showed the merged image. Bar = 20 μ m. (B) The focused zoom-image to

the plasma membrane (PM). It shows an enlarged portion of the yellow box displayed in the merged picture of A, and the order of picture is the same as in Figure A. Bar = 20 μ m.

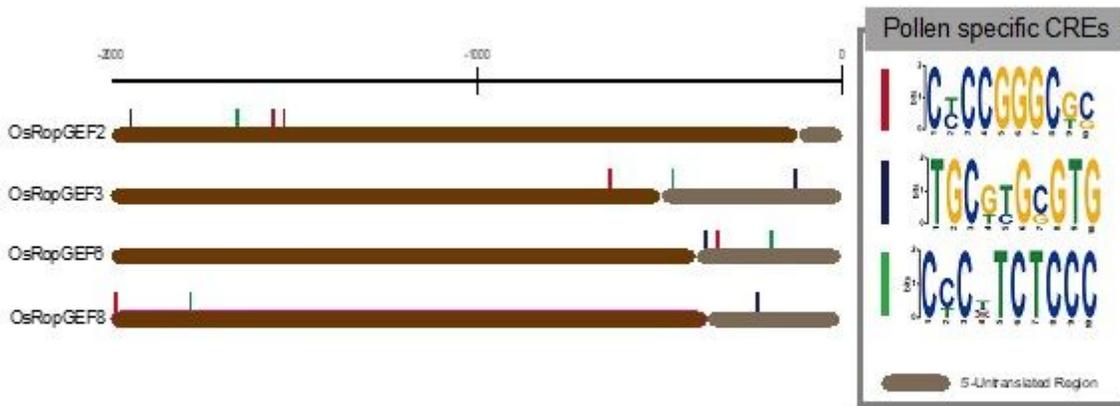


Fig. 6

Figure 6

Promoter analysis of four pollen-specific-expressed OsRopGEF genes. Identification of putative cis-acting regulatory elements (CREs) specifically found in pollen-specific-expressed OsRopGEF genes, using the MEME-suite. The number of scale bars above the figure shows the upstream position of the promoter base pair when taken at +1 of ATG. We analyzed up to the upstream 2,000-base pair. Gray bar indicated the 5'-untranslated regions and brown bars indicated the promoter regions.

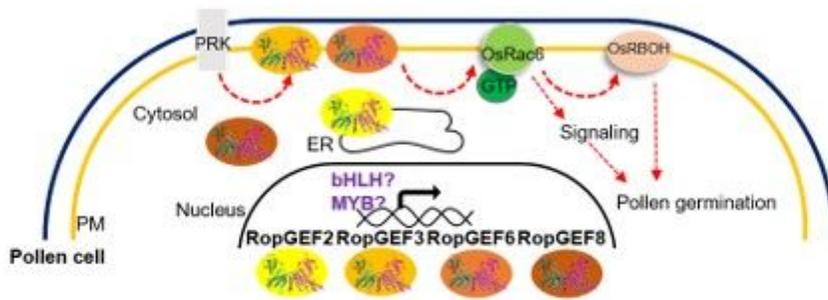


Fig. 7

Figure 7

A model summarizing four pollen-specific-expressed OsRopGEFs in pollen cells. PM, Plasma membrane; ER, Endoplasmic reticulum.

Supplementary Files

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