CYCLIN-DEPENDENT KINASE G1 Is Associated with the Spliceosome to Regulate CALLOSE SYNTHASE5 Splicing and Pollen Wall Formation in Arabidopsis

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Arabidopsis thaliana CYCLIN-DEPENDENT KINASE G1 (CDKG1) belongs to the family of cyclin-dependent protein kinases that were originally characterized as cell cycle regulators in eukaryotes. Here, we report that CDKG1 regulates pre-mRNA splicing of CALLOSE SYNTHASE5 (CalS5) and, therefore, pollen wall formation. The knockout mutant cdkg1 exhibits reduced male fertility with impaired callose synthesis and abnormal pollen wall formation. The sixth intron in CalS5 pre-mRNA, a rare type of intron with a GC 5’ splice site, is abnormally spliced in cdkg1. RNA immunoprecipitation analysis suggests that CDKG1 is associated with this intron. CDKG1 contains N-terminal Ser/Arg (RS) motifs and interacts with splicing factor Arginine/Serine-Rich Zinc Knuckle-Containing Protein 33 (RSZ33) through its RS region to regulate proper splicing. CDKG1 and RS-containing Zinc Finger Protein 22 (SRZ22), a splicing factor interacting with RSZ33 and U1 small nuclear ribonucleoprotein particle (snRNP) component U1-70k, colocalize in nuclear speckles and reside in the same complex. We propose that CDKG1 is recruited to U1 snRNP through RSZ33 to facilitate the splicing of the sixth intron of CalS5.

INTRODUCTION

Cyclin-dependent protein kinases (CDKs) belong to an important superfamily of Ser/Thr protein kinases that require the binding of a regulatory cyclin partner to induce their kinase activity (Morgan, 1997). Although CDKs were originally identified as key regulators of cell cycle transition (CDK1, 2, 4, and 6; Rane et al., 1999; Moore et al., 2003; Kozar et al., 2004), members in this family were also found to be involved in other cellular processes, such as activation of other CDKs (CDK7; Kaldis, 1999) and regulation of gene transcription by phosphorylating RNA polymerase II (CDK7, 8, and 9; Pinheiro et al., 2004). Recent studies both in animal and plant cells suggest that a number of CDKs may function in RNA splicing regulation (Doonan and Reddy, 1999). An snRNP is composed of an snRNA and several related snRNP proteins, for example, U1 snRNA and U1-70K in U1 snRNP (Valadkhan and Jaladat, 2010). Spliceosome assembly is initiated by U1 snRNP recognition and binding of the 5’ splice site (5’sS) of an intron, which is assisted by the SR-rich proteins (Cho et al., 2011). During early spliceosome assembly, a transient base-pairing forms between U1 snRNA and the 5’ SS, which could be stabilized by the binding of splicing factors to establish the U1 snRNP-pre-mRNA complex (Black, 2003). These splicing factors also include some SR protein family members, such as 9G8 and SC35. They are essential for the recognition of suboptimal splice sites (Kraiovicova et al., 2011). Therefore, SR proteins are important early in spliceosome assembly (Shepard and Hertel, 2009). Several SR proteins interact with U1-70K, such as SR222 and SR221 (Golovkin and Reddy, 1998), SR33, and SR45 (Golovkin and Reddy, 1999).

In angiosperms, the pollen wall plays an important role in the pollen–stigma interaction and in the protection of pollen from environmental stresses (Heslop-Harrison, 1968; Meuter-Gerhards et al., 1999). The pollen wall consists of a simple inner intine layer, an intricate outer exine layer, and an outer pollen coat. Pollen wall development begins with primexine formation between the callose wall and the microspore membrane at the tetrad stage (Arizumi and Toriyama, 2011). At this stage, four microspores, the products of a meiocyte, are enclosed inside a common callose envelope to form a tetrad. Callose synthase5 (CalS5; or glucan synthase-like2) is the main enzyme for the callose layer biosynthesis (Dong et al., 2005). Subsequently,
**RESULTS**

**Plants with Loss of CDKG1 Function Show Dramatically Reduced Male Fertility**

We performed functional analysis of CDKG1 in pollen wall formation and male fertility in *Arabidopsis*. We obtained a T-DNA insertion line of CDKG1 (SALK_075762, designated as cdkg1) from the ABRC (http://abrc.osu.edu/). The annotated T-DNA was stated to be inserted in the coding sequence (CDS) of CDKG1 (Figure 1A). However, PCR detection (Figure 1B) and sequencing analysis showed that the At5g63370 (CDKG1) CDS fragment from +605 to +1406 bp, together with the annotated T-DNA, was absent in this locus in cdkg1, although the T-DNA was still in the genome of cdkg1 (Figure 1A). The CDS for this truncated form of cdkg1 contained a premature stop codon (TGA, 607 to 609 bp) that would result in a truncated protein containing only a partial CDKG1 N terminus (Figure 1A). Therefore, this mutated protein is expected to lack CDK activity.

**cdkg1** plants displayed normal vegetative growth similar to the wild type (Columbia-0 [Col-0]; Figure 1C). Wild-type siliques contained 48.8 ± 11.4 seeds (n = 75) on average, whereas cdkg1 showed reduced fertility with only 5.35% seed set (2.61 ± 5.13 per silique, n = 400). The anther dehiscence and anther length of cdkg1 were similar to those of the wild type (Figure 1D). An Alexander’s staining (Alexander, 1969) assay for pollen viability showed the cdkg1 anthers contained very few purple-stained viable pollen grains (Figure 1D), which was consistent with its low fertility.

Backcrossing with the wild type (Col-0) resulted in F1 plants with normal fertility. In the F2 population, the male-sterile mutant phenotype was inherited in a Mendelian fashion, with 25% of the progeny displaying male sterility (98 of 409, x² = 0.235; P > 0.01), indicating that the reduced male fertility phenotype was caused by a single recessive nuclear mutation that was controlled sporophytically. For genetic complementation, a 3.6-kb genomic fragment consisting of 1300-bp upstream sequence, 1839-bp CDS, and 500-bp downstream sequence of CDKG1 (Figure 1A) was introduced into homozygous cdkg1 plants. In the T1 progeny, 17 of 23 transgenic lines showed restored fertility with normal pollen development and seed set (Figures 1C and 1D). This indicates that the deletion within CDKG1 was responsible for the mutant phenotype. There is another CDKG gene, CDKG2. However, the T-DNA insertion lines of this gene had no obvious phenotype, so we did not perform further analysis of the CDKG2 mutants.

To understand the details of the developmental defects in cdkg1 pollen development, we compared semithin cross sections of anthers of cdkg1 and the wild type. In *Arabidopsis*, anther development has been divided into 14 stages based on morphological characters (Sanders et al., 1999). Before stage 7 (tetrad stage), no detectable difference was observed between cdkg1 and the wild type. However, the microspores appeared to be shrunken in cdkg1 anthers at stage 7 (Figures 1E and 1K). At stage 8, most microspores of cdkg1 were relatively round rather than the typical angular shape of wild-type microspores (Figures 1F and 1L). At stage 9, wild-type microspores had a trivalvular shape with dense cytoplasm (Figure 1G), while most cdkg1 microspores were still round and cytoplasm leakage was observed from some microspores (Figure 1M). After stage 9, wild-type microspores gradually developed into mature pollen grains (Figures 1H to 1J). However, most cdkg1 microspores were gradually degraded after stage 9 (Figures 1N to 1P).

To obtain some clues about the role of CDKG1 in microspore development, we analyzed the expression pattern of CDKG1. Real-time RT-PCR analysis with total RNA from root, stem, rosette leaf, and inflorescence indicated that CDKG1 was preferentially expressed in leaf and inflorescence and weakly expressed in the other tissues examined (see Supplemental Figure 1A online). RNA in situ hybridization with wild-type anther cross sections showed that CDKG1 exhibited relatively higher expression at the meiosis and tetrad stages (see Supplemental Figure 1B online), suggesting that CDKG1 may play a role during these two stages.

**cdkg1 Plants Are Defective in Pollen Wall Formation**

Several mutants defective in pollen wall formation also show round microspores at stage 8 (Ariizumi et al., 2004; Paxson-Sowders et al., 2001; Guan et al., 2008). Therefore, we investigated pollen wall pattern of cdkg1 via scanning electron microscopy analysis. The surviving pollen grains of cdkg1 exhibited abnormal exine pattern when compared with that of the wild type (Figures 2A and 2B). To further examine the reason for...
the pollen wall defect in cdkg1, we performed transmission electron microscopy (TEM). In the wild type, the primexine matrix was deposited between the callose wall and the microspore plasma membrane at the early tetrad stage, and the microspore membrane became regularly undulated (Figure 2C). However, in cdkg1, the primexine matrix was thinner and the regular plasma membrane undulation was not observed (Figure 2F). In the wild type, the callose wall was less electron dense at the early tetrad stage, a time when the electron-dense sporopollenin (or more probably its precursor) was deposited on the outside surface of the callose wall (Figure 2C). By contrast, in cdkg1, the callose wall was moderately electron dense, and flat sporopollenin particles were observed on the surface of the microspore plasma membrane at the early tetrad stage (Figure 2F). At the late tetrad stage of the wild type, the degenerating callose wall was electron dense, and the sporopollenin was deposited as rod-like probacula that were regularly distributed in the primexine matrix (Figure 2D). However, in cdkg1, dot-like sporopollenin particles were randomly distributed on the plasma membrane (Figure 2G). At stage 10, the exine composed of tecta and bacula was formed around wild-type microspores (Figure 2E), while this layer was absent in cdkg1, with only big sporopollenin granules around microspores (Figure 2H). These results suggest that the primexine formation is defective in cdkg1.
Callose Deposition Is Defective in cdkg1
Due to the Abnormal Splicing of CalS5 Pre-mRNA

Since the callose wall is important in primexine formation (Nishikawa et al., 2005; Dong et al., 2005), we investigated callose deposition at the tetrad stage by aniline blue staining. The callose wall fluorescence of the cdkg1 tetrad was weaker than that of the wild type, especially that of the peripheral callose wall (Figure 3A). Callose staining of tetrad stage sections showed that the callose wall of cdkg1 was much thinner than that of the wild type (Figures 1E and 1K). This was further confirmed by a callose fluorescence bleaching test. For the 30 cdkg1 tetrads randomly selected from 30 tetrad-stage buds, the average fluorescence quenching time for the callose wall was much less than that for the wild type (Figure 3A). These results suggest that callose synthesis was significantly reduced in cdkg1. In Arabidopsis, CalS5 (Figure 3B) encodes the main synthase for the peripheral callose wall around meiocytes and tetrads (Dong et al., 2005), and this callose layer is almost lost in cals5 knockout mutants (Dong et al., 2005; Nishikawa et al., 2005; Figure 3A). We therefore investigated whether the expression of CalS5 was altered in cdkg1. Quantitative RT-PCR analysis using the primer set for exon2-3 of CalS5 showed 3.9-fold upregulation in cdkg1, whereas the primer set for exon 39 showed only 17% of the wild-type level (Figure 3C). This suggests that the full-length CalS5 transcript in cdkg1 is present at most at 17% of the level in the wild type.

To further examine the abnormal transcription of CalS5 in cdkg1, additional primer pairs were designed for RT-PCR analysis. When the primer set for exons 5 to 7 was used, two PCR products were obtained in cdkg1. The smaller product was similar to the size of the PCR band from wild-type cDNA, while the larger one was in a size intermediate between the smaller one and the PCR product from the genomic DNA (Figure 3D). Sequencing showed that the larger product contained the 99-bp intron 6 of CalS5. Thus, the sixth intron was only partially spliced in cdkg1 (Figure 3E). A stop codon occurred inside intron 6, resulting in a truncated protein with only 254 amino acid residues for the abnormal CalS5 mRNA containing the intron. The other primer sets tested showed that other introns of CalS5 were spliced normally in cdkg1.

CalS5 Pre-mRNA Is Coimmunoprecipitated with CDKG1

To understand whether CDKG1 is directly associated with intron 6 of CalS5 pre-mRNA, an RNA immunoprecipitation (RIP) assay...
was performed as described (Terzi and Simpson, 2009). We made two protein fusion constructs, one tagged with Myc, PCDKG1:CDKG1-Myc (Figure 4A), and one with green fluorescent protein (GFP), PCDKG1:CDKG1-GFP (Figure 4C). Each construct complemented the phenotypic defects when introduced into cdkg1 (Figures 4B and 4D), indicating that the two fusion proteins can fulfill the function of CDKG1. In fluorescence of these transgenic plants were used for the RIP assay. In the RIP assay with CDKG1-Myc, for the primer set spanning intron 6, an expected PCR product was detected in the anti-Myc RIP (Figure 4E, lane 4). This PCR product was absent in the anti-Myc RIP with the wild type (Figure 4E, lane 3) and in both mock RIPs (with magnetic beads only) of the transgenic plants and the wild type (Figure 4E, lane 5 and 6). We used the primer sets CalS5-900 and SRZ22-CDS as negative controls. No PCR product was obtained from any of these RIP cDNAs (Figure 4F). Similar results were observed when we used PCDKG1:CDKG1-GFP transgenic plants (Figures 4G and 4H). These results indicate that CDKG1 might bind to the CalS5 pre-mRNA. To investigate whether CDKG1 can bind to other parts of CalS5 pre-mRNA, a total of nine additional primer sets for CalS5 was designed for RIP RT-PCR analysis. PCR products were obtained only for primer sets covering intron 6 and exons 5 to 7 (Figure 4I). These results indicate that CDKG1 is associated with the CalS5 pre-mRNA around intron 6. Since CDKG1 is a protein without an obvious RNA recognition motif (RRM), we hypothesize that

![Figure 3. cdkg1 Shows Reduced Callose Synthesis and Abnormal pre-mRNA Splicing of CalS5.](image-url)

![Figure 4. CDKG1 Is Associated with the Intron 6 Region of CalS5 pre-mRNA.](image-url)
CDKG1 might bind to CalS5 pre-mRNA by interacting with some other proteins, such as splicing factors that have RRM s.

**CDKG1 Interacts with Splicing Factor RSZ33**

Several CDKs have been reported to be involved in pre-mRNA splicing, including human CDK11, CDK12, and CDK13 and Arabidopsis CDKC2 (Hu et al., 2003; Chen et al., 2006, 2007; Even et al., 2006; Kitsios et al., 2008). These CDKs share two common features: (1) they colocalize or interact with splicing factors such as SR proteins; and (2) they contain at least one RS motif in their N termini, for example, CDK11 (three RS motifs), CDK12 (20 RS motifs), CDC2L5/CDK13 (15 RS motifs), and CDKC2 (one RS motif). RS motifs are thought to mediate the interaction with splicing factor SR proteins (Wu and Maniatis, 1993; Zhu and Krainer, 2000). CDKG1 has an N terminus that is longer than those of CDKA to CDKF. CDKG1 contains four RS motifs in the N terminus (Figure 5A), including the last RS motif conserved in almost in all the CDKG1 orthologs (see Supplemental Figure 2A online). We also compared the core region (as defined by Ko et al., 2001) of CDKG1 with those of Arabidopsis CDKs and the mammalian CDKs related to splicing regulation. CDKG1 is more similar to the CDKs associated with splicing than the others, including CDKA, CDKB, and CDKF (see Supplemental Figure 2B online).

In the phylogenetic tree of Arabidopsis CDKs, the CDKG clade is close to the CDKC clade (see Supplemental Figure 2C online). Therefore, as these CDKs are related to splicing regulation, CDKG1 might also interact with some SR proteins with one to two RRM s (Shepard and Hertel, 2009).

We therefore performed yeast two-hybrid assays to find potential SR protein partners of CDKG1. The Arabidopsis genome contains 19 SR genes (Reddy, 2004). We cloned the CDSs of these SR genes into the pGBKKT7 vector and the CDS of CDKG1 into the pGADT7 vector. The yeast clones bearing pGAD-CDKG1 and pGBK-RSZ33 were able to grow on the selective medium (Trp-, Leu-, adenine-, and His-), although their growth was slower than that of the positive control, containing pTAC12 and pTAC14 (Gao et al., 2011). These clones showed positive X-α-Gal reaction (Figure 5B). These results show that CDKG1 interacts with RSZ33 in yeast.

**Figure 5.** CDKG1 Interacts with Splicing Factor RSZ33.

(A) Domains of 14 CDKs in Arabidopsis. S_TKC domain, Ser/Thr, protein kinase catalytic domain; RS motif, Ser/Arg motif. (B) CDKG1 full length and an internal fragment both interact with RSZ33 in yeast cells; SR34+CDKG1 is a negative control, and pTAC12+pTAC14 is a positive control. BK, pGBK-T7; AD, pGADT7; 232-395, CDKG1 fragment amino acids 232 to 395. (C) BiFC assay between CDKG1 and RSZ33 in tobacco leaf cells. YC, YFP C terminus; YN, YFP N terminus. Bars = 10 μm. (D) mRFP-CDKG1 colocalizes with GFP-RSZ33. A root cell nucleus of a transgenic Arabidopsis plant expressing the two fusion proteins is shown. Bars = 10 μm.
To confirm and visualize the interaction between RSZ33 and CDKG1 in plant cells, a bimolecular fluorescence complementation (BiFC) assay was performed based on split yellow fluorescent protein (YFP; Hu et al., 2002). CDKG1 and RSZ33 were fused with the N and C terminus of YFP, respectively, and transiently coexpressed in tobacco (*Nicotiana tabacum*) leaves. For two combinations (YN-CDKG1+YC-RSZ33 and YC-CDKG1+YN-RSZ33), YFP signals were observed in the nucleus, with relatively stronger signals in nuclear speckles. No YFP signal was observed for the other two combinations, YN+YC-CDKG1 and YC+YN-CDKG1 (Figure 5C). The BiFC assay confirmed the interaction between CDKG1 and RSZ33. When the monomeric red fluorescent protein (mRFP)–tagged CDKG1 (mRFP-CDKG1) and GFP-RSZ33 were coexpressed in transgenic plants, we also observed their colocalization in nuclear speckles (Figure 5D), where most SR proteins localize (Fang et al., 2004). This colocalization assay therefore supports the interaction between the two proteins.

**CDKG1 Is Targeted to Splicing Machinery and Is Dependent on the Region Including the Conserved RS Motif**

RSZ33 is known to interact with SRZ22 and SRZ21 (Lopato et al., 2002), which further interact with U1-70k, a component of the U1 snRNP (Golovkin and Reddy, 1998). This suggested that CDKG1 might be targeted to the splicing machinery through physical and functional interaction with RSZ33. To further verify the involvement of CDKG1 in the splicing machinery, we performed colocalization and coimmunoprecipitation assays with CDKG1 and SRZ22. When coexpressed in tobacco leaf epidermal cells, both the cyan fluorescent protein (CFP)–tagged CDKG1 (CDKG1-CFP) and SRZ22-YFP were observed in the nucleoplasm, primarily in nuclear speckles (Figure 5A). We obtained transgenic *Arabidopsis* lines expressing both P35S:mRFP-CDKG1 and P35S:SRZ22-YFP. Nuclear extracts from the transgenic plants were immunoprecipitated using an anti-GFP antibody. Immunoblot analysis of the immunoprecipitated products using an anti-RFP antibody detected a protein band consistent with the size of the mRFP-CDKG1 fusion protein (Figure 6B). This coimmunoprecipitation assay suggests that CDKG1 and SRZ22 reside in the same complex in nuclear speckles.

To define the region in CDKG1 responsible for nuclear speckle localization, we made several constructs using GFP fused with the CDKG1 fragments (Figure 6C). CDKG1 contains a nuclear localization signal in its N terminus, according to the PredictNLS server (Cokol et al., 2000; https://rostlab.org/owiki/index.php/PredictNLS). When fused with N-terminal fragments containing various RS motifs, CDKG1 1 to 93 (RS motifs 1 and 2), 1 to 264 (RS motifs 1 to 3), and 1 to 300 (all four RS motifs), diffuse GFP signals were observed in the nuclei of the epidermal cells (Figure 6D). Similarly, for CDKG1 298-612-GFP lacking the CDKG1 N terminus, the diffuse GFP signals were observed in the cytoplasm and nucleus, similar to the localization pattern of free GFP (Figure 6D). These results suggest that both the N terminus and the S_TKC domain are required for its nuclear speckle localization. The other three fusion proteins, including 1-395-GFP,

**Figure 6.** CDKG1 Is Targeted to Splicing Machinery in a Manner Dependent on the Region Including the Conserved RS Motif.

(A) CDKG1-CFP colocalizes with SRZ22-YFP in tobacco leaf cells. Bars = 5 μm.

(B) mRFP-CDKG1 is coimmunoprecipitated with SRZ22-YFP. Left panels, inputs. The inputs detected by anti-RFP antibody are nuclear extracts and the inputs detected by anti-β-actin antibody (down) are total soluble proteins. Right panel, nuclear extracts detected by anti-RFP antibody that were immunoprecipitated by anti-GFP magnetic beads or by magnetic beads alone. RG1, mRFP-CDKG1; SY, SRZ22-YFP; WT, the wild type.

(C) Diagrammatic representation of CDKG1 and the fragments fused with GFP. Blue line, RS motifs; green line, nuclear localization signal; yellow line, PLTSLRE motif; cyan box, T-loop; white box, N terminus; black box, S_TKC domain; gray box, C terminus.

(D) The cellular localization of CDKG1-GFP fragments in tobacco leaf cells. C, cytoplasm; N, nucleus. Bars = 5 μm.
1-450-GFP, and 232-395-GFP, showed nuclear speckle localization (Figure 6D), similar to full-length CDKG1-CFP. The CDKG1 fragment includes the peptide from 232 to 395, which showed a stronger interaction with RSZ33 than did the full-length CDKG1 (Figure 5B). This 232 to 395 fragment contains a partial core region and an N-terminal RS motif (the fourth motif in the N-terminal region), which is conserved in almost all orthologs of CDKG1 in other eukaryotes (see Supplemental Figure 2A online). The core region also includes some critical residues for the function of a CDK, including the putative substrate binding pocket. These results suggest that the fragment 232 to 395 including the conserved RS motif is sufficient to define the CDKG1 nuclear speckle localization and to interact with RSZ33.

**DISCUSSION**

CDKG1 is one of the two CDKG genes in the *Arabidopsis* CDK gene family, with RS motifs in its N terminus. The *cdkg1* is a knockout mutant, presumably due to a T-DNA–induced deletion (Figure 1A). Similar T-DNA–induced deletions have been reported in dozens of T-DNA lines (Wang, 2008). Here, we present that CDKG1 regulates pollen wall formation (Figure 2) and male fertility (Figure 1C). Molecular and biochemical evidence shows that CDKG1 interacts with a splicing factor (Figures 5B to 5D) to regulate the pre-mRNA splicing of CalS5 (Figures 3D and 3E), the major enzyme for callose synthesis and pollen wall development. Double mutant analysis of *cdkg1 cals5-2* (see Supplemental Figure 3 online) suggests that CDKG1 may also regulate some other genes involved in pollen wall formation and plant fertility.

**CDKG1 Interacts with RSZ33 to Regulate the Splicing of *CalS5* Pre-mRNA**

We present several lines of evidence, including yeast two-hybrid, BIFC, and colocalization assays, to support the interaction between CDKG1 and splicing factor RSZ33 (Figures 5B to 5D). RSZ33 is known to interact with SRZ22 and SRZ21 (Lopato et al., 2002), which further interact with U1-70K, a component of U1 snRNP (Golovkin and Reddy, 1998). Additionally, CDKG1 is also colocalized with SRZ22 in tobacco leaves and coimmunoprecipitated with this SR protein in transgenic *Arabidopsis* (Figures 6A and 6B). The RIP assay showed that CDKG1 is associated with the pre-mRNA of *CalS5* (Figures 4E to 4I). Therefore, it is likely that CDKG1 binds to the *CalS5* pre-mRNA indirectly through SRZ33, a splicing factor with one RRM (Shepard and Hertel, 2009).

The sixth intron of *CalS5* is a rare type of intron with a GC 5′ SS. A GC 5′ SS is intrinsically weaker than the major GT 5′ SS because the T-C substitution introduces a mismatch in the base pairing between the splice site and U1 snRNA (Kralovicova et al., 2011). For this rare type of intron, various SR proteins bind to the U1 snRNP-pre-mRNA complex to stabilize the base pairing (Black, 2003). Human SR proteins 9G8 and SC35 can promote the splicing of introns with the GC 5′ SS (Kralovicova et al., 2011). In *cdkg1*, the low splicing efficiency of the *CalS5* intron 6 (Figure 3D) verifies the weakness of this GC 5′ SS. This suggests that U1 snRNP binding to the sixth intron of *CalS5* is not stable.

The normal splicing of this intron in the wild type and in *cdkg1* complementation lines suggests that CDKG1 is recruited to stabilize the binding of U1 snRNP to the region of *CalS5* intron 6. This facilitates spliceosome removal of this rare type of intron. Kralovicova et al. (2011) reported that the flanking sequences could enhance the splicing efficiency of an intron with the GC 5′ SS. Therefore, both CDKG1 and the flanking sequences are important for efficient splicing of the introns with a GC 5′ SS.

**CDKG1 Affects Pollen Wall Formation during Microsporogenesis**

CDKG1 is preferentially expressed in leaves and inflorescences with relatively low expression in root and stem (see Supplemental Figure 1A online). However, only a defect in male reproduction was observed in *cdkg1*. This is similar to some other genes with extensive expression patterns that are involved in microsporogenesis, such as CDKA (Iwakawa et al., 2006) and NPU (Chang et al., 2012). It is probable that a lack of gene redundancy in anthers allows observation of the *cdkg1* phenotype during anther...
development. CDKG1 may be involved in other biological processes that cannot be observed morphologically in other tissues. CalS5 shows highest expression in meiocytes, tetrads, microspores, and mature pollen (Dong et al., 2005; Nishikawa et al., 2005). The relatively higher expression of CDKG1 at the meiosis and the tetrad stages (see Supplemental Figure 1B online) is consistent with the regulation of CalS5 pre-mRNA splicing during these stages.

Pollen development includes several cell division processes: male meiosis, tapetal cell endoreproduction, and pollen mitosis. We analyzed these cell division processes with 4',6-diamidino-2-phenylindole staining (Ross et al., 1996). No obvious defects were found in cdkg1 in these cell cycle events. During meiosis I and II, the nuclear division and chromosome distribution in cdkg1 were indistinguishable from those in the wild type. In cdkg1, the tapetal cells contained two nuclei at the meiosis stage, and the surviving pollen grains contained three nuclei (see Supplemental Figure 4 online). These data indicate that CDKG1 is not or not solely involved in cell cycle regulation of anther cells during anther development.

Pollen wall development initiates at the tetrad stage with primexine formation beneath the callose wall. At this stage, the callose wall may also act as a molecular filter to protect the haploid microspores in tetrads from the influence of the diploid tissues in anther (Heslop-Harrison and Mackenzie, 1967). CalS5 is responsible for callose layer synthesis (Dong et al., 2005). Both knockout (Dong et al., 2005) and reduced expression of CalS5 (Nishikawa et al., 2005) cause an exine developmental defect. cdkg1 showed downregulated CalS5 expression (Figure 3C) and improper callose deposition around tetrads (Figure 3A), which are quite similar to those in CalS5 mutants. Based on the molecular filter function of callose wall and the TEM results (Figures 2C and 2D), we speculated that at the early tetrad stage in the wild type, the sporopollenin or its precursors can hardly pass through the dense callose wall until the callose wall begins to be degenerated at the late tetrad stage, as indicated by the electron density of callose wall and the probacula formation (Figures 2C and 2D). By contrast, in cdkg1 tetrads, the sporopollenin seems to be deposited around microspores in advance (Figure 2F). It is possible that the lower callose content (Figure 3A) results in a bigger mesh size of the molecular filter of the cdkg1 callose wall. Therefore, it cannot protect the sporopollenin or its precursors from invading at the early tetrad stage, when the primexine, the mold for exine formation, is not fit to accept them. Simultaneously, the primexine is also thinner in cdkg1 tetrads than in the wild type, even at the late tetrad stage (Figures 2F and 2D). CDKG1 may regulate the synthesis and/or the deposition of primexine. It was proposed that callose could conceivably trap primexine subunits, increasing their local concentration and preventing them from diffusing into the anther locule (Ariizumi and Toriyama, 2011). Probably the bigger mesh size of the molecular filter of the callose wall in cdkg1 leads to leakage of the primexine materials. In cdkg1, the probacula do not have enough space to develop because of the thinner primexine. As a result, sporopollenin is deposited randomly into flat particles (Figure 2F) and then into incomplete dot-like structure around the microspores of cdkg1 (Figures 2G and 2H). Finally, most developing microspores of cdkg1 undergo degradation after stage 9 (Figures 1M to 1P) as they lose protection from the complete and regular exine layer (Figure 2H).

In cdkg1, the CalS5 5' region was upregulated to 3.9-fold and its 3' region was downregulated to ~17% of the wild-type level (Figure 3C). One possibility for the abnormal expression is a pause in gene transcription owing to the disturbed splicing as reported (Bentley, 1999; Kitsios et al., 2008). Alternatively, the abnormal CalS5 transcript might undergo noncoding mRNA decay due to the premature termination codon in the nonspliced intron 6. Previous studies revealed that abnormal mRNA containing a premature termination codon could be degraded by the noncoding mRNA decay mechanism (Culbertson and Leeds, 2003; Isken and Maquat, 2007). Both of the above possibilities could result in reduced expression of mature CalS5 and the accumulation of the 5' region of CalS5 mRNA in cdkg1 (Figure 3C).

The phenotype rescue by CalS5 CDS is important in revealing the relationship between the cdkg1 phenotype and the abnormal CalS5 splicing. When we obtained the full-length CDS of CalS5, all of the fragments were mutated. This is in agreement with a previous report that T-DNA constructs containing a full-length genomic copy of CalS5 are lethal in Agrobacterium tumefaciens (Nishikawa et al., 2005). Therefore, we constructed double mutants to analyze the relationship between CDKG1 and CalS5 in this work. Both cdkg1 and CalS5 knockout mutants were able to yield a limited number of seeds. If CalS5 is the only target of CDKG1 in microsporogenesis, the phenotype of cdkg1 calss-2 should be the same as that of calss-2. However, the double mutant had a more severe phenotype (i.e., almost complete male sterility and pollen loss; see Supplemental Figure 3 online). This suggests that CDKG1 may regulate some other genes important for primexine development beyond CalS5. In Arabidopsis, DEX1, NEF1, RPG1, and NPU are involved in primexine formation (Paxson-Sowders et al., 2001; Ariizumi et al., 2004; Guan et al., 2008; Chang et al., 2012; Sun et al., 2013), but the splicing patterns of these genes were not altered in cdkg1, and their transcription levels were similar to those in the wild type (see Supplemental Figure 5 online). CDKG1 may therefore regulate some novel gene(s) important for pollen wall formation and plant fertility.

Putative Working Model of CDKG1 Function in Pre-mRNA Splicing

We propose a working model for CDKG1 function in pre-mRNA splicing of CalS5 (Figure 7). Normally, introns with the GT 5' SS are spliced without the help of CDKG1. For rare introns with the GC 5' SS, such as intron 6 of CalS5, CDKG1 is recruited to U1 snRNP through RSZ33 and SRZ22 to stabilize the base paring between the GC 5' SS and U1 snRNA. This recruitment facilitates the efficient splicing of intron 6. The mature CalS5 mRNA is then transferred to the cytoplasm for translation of CalS5.

METHODS

Plant Growth Conditions

The Arabidopsis thaliana plants used in this study are in a Col-0 background. Seeds were sown on vermiculite and allowed to imbibe for 3 d at 4°C. Plants were grown under long-day conditions (16 h light/8 h dark) in an ~22°C growth room. Transgenic plants were generated via Agrobacterium tumefaciens–mediated transformation (Clough and Bent, 1998) and selected
on PNS (Plant Nutritional Solution) media containing 20 mg/L hygromycin B and/or 50 mg/L kanamycin. Transient expression for subcellular localization and BiFC assays in tobacco (Nicotiana tabacum) leaves were performed as previously described (Hu et al., 2002).

Phenotype Characterization and Microscopy

Plant photos were taken with a Cybershot T-20 digital camera (Sony). Flower pictures were taken using an Olympus dissection microscope with an Olympus digital camera. Alexander’s staining and 4’’,6-diamidino-2-phenylindole staining were performed as described (Alexander, 1969; Ross et al., 1996). Callose staining and semithin sections were performed as described in a previous report (Zhang et al., 2007). For the callose fluorescence bleaching assay, the inflorescences were fixed in Carnoy’s fixative for 3 h. After rinsing with phosphate buffer, pH 7.2, three times, the inflorescences were stained with 0.1% aniline blue at 4°C for 3 d. The tetrads were isolated from the anthers of the inflorescences and kept under UV. The quenching time was recorded when the callose fluorescence of each tetrad disappeared. Pictures were taken with an Olympus BX-51 microscope (Olympus).

For scanning electron microscopy examination, the opened flowers of the wild type and cdkg1 were dissected and the dehiscent anthers were coated with 8-nm gold. Observation and photography were performed on a JSM-840 microscope (Jeol). TEM samples were prepared as previously described (Zhang et al., 2007). Sections were observed on an H-600 transmission electron microscope (Hitachi).

For cdkg1 complementation, the 3.6-kb CDKG1 genomic fragment was amplified using LA-Taq polymerase (Takara) with primer set CMP-F/ CMP-R. The oligonucleotides used for complementation and other assays in this study are listed in Supplemental Table 1 online. Primers were synthesized by Life Technologies. After verification by sequencing (Genomics), the fragment was cloned into the pCAMBIA1300 binary vector (Cambia) and subsequently introduced into homozygous cdkg1 plants. For cdkg1 background verification, SC-F/SC-R primers were used to validate DNA deletion of CDKG1 for the transformants. LP/RP primers were used to detect either CDKG1 genomic sequence or transgenic complementation fragment, and the genomic-specific primers CHC-F/ CHC-R were used to validate the homozygous cdkg1 background. As CHC-R primer was designed 180 bp downstream of CMP-R primer, PCR with the CHC-F/CHC-R primer set was not able to amplify a 1.7-kb fragment in homozygous plants even if the complementation fragment was integrated into the genome.

Expression Analysis

RNA extraction was performed using Trizol (Life Technologies) following the protocol in the user’s manual. A total of 30 cycles of PCR was performed for RT-PCR if not mentioned. Col-0 and cdkg1 inflorescence cDNAs were used for the expression analysis of CDKG1, CalS5, and the pre-mRNA splicing of CalS5. Real-time quantitative PCR were performed using gene-specific primers and IQ SYBR Green Master Mix (Bio-Rad) on the ABI 7300 platform (Applied Biosystems). The experiment was repeated thrice and data were averaged. β-Tubulin gene was used as an internal normalization control. Fold change in gene expression were calculated using the ΔΔCT (cycle threshold) values.

Subcellular Localization, Colocalization Analysis, and BiFC Assay

GFP, CFP, YFP, and mRFP fluorescence was detected under an Olympus IX70 inverted microscope system. For GFP fusion, CDKG1 and its fragments were amplified from Arabidopsis wild-type genomic DNA with KOD polymerase (Toyobo). The fragments were cloned into the binary vector pMON530-GFP or pCAMBIA1300-35S-GFP, driven by the 35S promoter. These constructs were introduced into wild-type plants or transiently expressed in tobacco leaf via Agrobacterium strain ASE or GV3101 as described (Clough and Bent, 1998; Hu et al., 2002). For CFP and mRFP fusions, the CDKG1 CDS was amplified with KOD polymerase (Takara) using the primer set G1-Bam-F/G1-Sal-R. The CDKG1 CDS was cloned into the binary vectors pC131-35S-CFP and pCAMBIA1300-35S-mRFP. SRZ22 CDS was amplified from Arabidopsis wild-type inflorescence cDNA with the primer set SRZ22-Kpn-Eco-F/SRZ22-Bam-Sal-R and then cloned into pMON530-YFP. The two constructs were introduced into wild-type plants or transiently expressed in tobacco leaves as described above.

For BiFC assays, the CDSs of CDKG1 and RSZ33 were cloned into pCAMBIA1300-35S-YN and pCAMBIA1300-35S-YC vectors and transiently expressed in tobacco leaf cells as described by Hu et al. (2002).

In Vivo Communoprecipitation

In vivo communoprecipitation assay was performed as previously described (Fill et al., 2008). The leaves of 4-week-old transgenic plants of the wild type and 35S:SRZ22-YFP::3Ss:mRFP-CDKG1 were used as protein sources. After fixation, grinding, and sonication treatment, nuclear proteins were extracted from 4 g of leaves. Protein G-coupled magnetic beads (Life Technologies) and an anti-GFP antibody (Cwbiotech) were used for immunoprecipitation. Eluted proteins were run on SDS-PAGE, immunoblotted using the Pierce Fast Western Blot Kit (Thermo Scientific), and probed with an anti-RFP (MBL).

RIP Assay

The 3.1-kb CDKG1 genomic sequence before the terminal code was amplified with the primer set PG1-Eco-F/G1-Sal-R and cloned into pCAMBIA1300-Myc and pCAMBIA1300-GFP, following a 3’-Nos sequence. The RIP assay was performed as described (Terzì and Simpson 2009). Briefly, 3 to 5 g of inflorescence from transgenic and wild-type (as control) plants was fixed, and the total soluble nuclear extracts were isolated after sonication. A portion of each nuclear extract was immunoprecipitated with the corresponding antibody-coupled magnetic protein G beads (Life Technologies) or protein G beads as control. Sixty microliters of each nuclear extract was stored at −70°C for input preparation. RNAs were isolated from the immunoprecipitated products and each input. The resulting cDNAs were used for RT-PCR analysis. The anti-Myc antibody was obtained from Genomics.

Yeast Two-Hybrid Analysis

Yeast two-hybrid assay was performed using the Clontech two-hybrid system according to the manufacturer’s instructions. The CDS of CDKG1 and its fragment 232 to 395 were amplified and cloned into the pGADT7 vector. The CDSs of 19 SR proteins were cloned into pGBK7 vector individually. The pGADT7-CDKG1 was cotransformed into the yeast strain AH109 with the pGBK7-SR individually. The transformants were screened on supplemented synthetic dextrose medium lacking Leu and Trp or on supplemented synthetic dextrose medium lacking Leu, Trp, His, and adenine hemisulfate salt with X-a–Gal. The positive clones were verified both by CDKG1-specific primers and by SR protein-specific primers.

Phylogenetic Analysis

The multiple sequence alignment of full-length protein sequences was performed using the ClustalW tool online (http://www.genome.jp/tools/clustalw/) and displayed using Boxshade (http://www.ch.embnet.org/software/BOX_form.html). The alignment is available as Supplemental Data Set 1 online. Phylogenetic trees were constructed and tested by MEGA4.0 based on the neighbor-joining method (bootstrap method with 1000 trials was used to calculate the statistical support for the nodes).
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AUTHOR CONTRIBUTIONS

X.-Y.H. and Z.-N.Y. designed the research. X.-Y.H., J.N., M.-X.S., J.Z., and Z.-N.Y. analyzed the data. X.-Y.H. and Z.-N.Y. prepared the article. X.-Y.H. and Z.-N.Y. designed the research. X.-Y.H., J.N., M.-X.S. J.Z., and Z.-N.Y. analyzed the data. X.-Y.H. and Z.-N.Y. prepared the article.

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CYCLIN-DEPENDENT KINASE G1 Is Associated with the Spliceosome to Regulate CALLOSE SYNTHASE5 Splicing and Pollen Wall Formation in Arabidopsis
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