The R2R3 MYB transcription factors FOUR LIPS and MYB88 regulate female reproductive development

Srilakshmi Makkena1,2, Eunkyoung Lee3, Fred D. Sack3 and Rebecca S. Lamb2,*

1 Plant Cellular and Molecular Biology Graduate Program, The Ohio State University, Columbus, OH 43210, USA
2 Department of Molecular Genetics, The Ohio State University, Columbus, OH 43210, USA
3 Department of Botany, The University of British Columbia, Vancouver, BC, Canada V6T 1Z4

*To whom correspondence should be addressed. E-mail: lamb.129@osu.edu

Received 9 March 2012; revised 21 June 2012; accepted 25 June 2012

Abstract

Gamete formation is an important step in the life cycle of sexually reproducing organisms. In flowering plants, haploid spores are formed after the meiotic division of spore mother cells. These spores develop into male and female gametophytes containing gametes after undergoing mitotic divisions. In the female, the megaspore mother cell undergoes meiosis forming four megaspores, of which one is functional and three degenerate. The megaspore then undergoes three mitotic cycles thus generating an embryo sac with eight nuclei. The embryo sac undergoes cellularization to form the mature seven-celled female gametophyte. Entry into and progression through meiosis is essential for megasporogenesis and subsequent megagametogenesis, but control of this process is not yet well understood. FOUR LIPS (FLP) and its parologue MYB88, encoding R2R3 MYB transcription factors, have been extensively studied for their role in limiting the terminal division in stomatal development by direct regulation of the expression of cell cycle genes. Here it is demonstrated that FLP and MYB88 also regulate female reproduction. Both FLP and MYB88 are expressed during ovule development and their loss significantly increases the number of ovules produced by the placenta. Despite the presence of excess ovules, single and double mutants exhibit reduced seed set due to reduced female fertility. The sterility results at least in part from defective meiotic entry and progression. Therefore, FLP and MYB88 are important regulators of entry into megasporogenesis, and probably act via the regulation of cell cycle genes.

Key words: Arabidopsis thaliana, female sterility, FLP, gametogenesis, MYB88, plant reproduction, transcription

Introduction

Plants are characterized by the alternation of haploid (gametophyte) and diploid (sporophyte) generations. In angiosperms, such as Arabidopsis thaliana, the gametophyte is short lived and develops within the sporophytic tissue of the flower. The female gametophyte, or embryo sac, develops within ovules contained in the ovary of the pistil. Ovule primordia develop from placental tissue in the ovary. Along the proximal–distal axis an ovule primordium consists of three distinct regions: the funiculus, the chalaza, and the nucellus. Within the nucellus the megaspore mother cell (MMC) is formed, which divides first meiotically and then mitotically to form the embryo sac (reviewed in Berger and Twell, 2011). The MMC divides meiotically to make four haploid megaspores, of which three degenerate and one, the proximal chalazal megaspore, continues to develop. This functional megaspore increases in size and undergoes a series of nuclear divisions to form an eight-nuclei embryo sac. Then cellularization takes place, dividing the embryo sac into seven cells with four cell types: three antipodal cells at the chalazal end, a diploid central cell, and two synergids and the egg cell at the micropylar end. Antipodal cells, which have no currently known function,
degenerate before fertilization in *Arabidopsis*. The central cell, the egg cell, and the synergid cells form the female germ unit.

Proper specification of the MMC, entry into and completion of meiosis, control of mitotic divisions, and cell fate determination and differentiation are essential for development of the embryo sac. In *Arabidopsis*, a number of genes are known to regulate these processes. For example, mutations in a gene encoding a type I MADS-box transcription factor, *AGL23*, block the first nuclear division of the functional megasporo (Colombo et al., 2008). Mutations in the *PROLIFERA* (*PRL*) gene, encoding the DNA replication licensing factor subunit *MC7*, arrest the embryo sac with one nucleus (Springer et al., 2000). Temporal and spatial regulation of cellularization of the syncytial embryo sac is critical for gametophytic development and cell fate. In the female sterile mutant *hadad* (*hdd*), the gametophyte becomes prematurely cellularized after the first or second gametophytic division (Moore et al., 1997). Simultaneously with cellularization, cell fate within the embryo sac is acquired and differentiation begins. *MYB98* encodes an R2R3-MYB transcription factor specifically expressed in the synergid cells. *myb98* mutants fail to form the filiform apparatus and are unable to guide the pollen tube to the micropyle (Punwani et al., 2007). However, although many genes are known that are involved in meiotic recombination and other aspects of the meiotic cycle (reviewed in Ma, 2006), how MMC entry into meiosis is controlled is still not known.

The atypical R2R3-MYB transcription factor *FOUR LIPS* (*FLP*) and its parologue *MYB88* have been well characterized in *Arabidopsis*. In loss-of-function *flp* single and *flp; myb88* double mutants, the placenta produces extra ovule primordia, suggesting these genes normally regulate the proliferation of this tissue. How *FLP* and *MYB88* function as positive transcriptional regulators of entry into meiosis is still unknown. The *FLP/MYB88* transcriptome in *flp-1; myb88* double mutants, the placenta produces extra ovule primordia, suggesting these genes normally regulate the proliferation of this tissue. However, seed set is reduced. The reduction is the result of defects in female megasporogenesis. The expressivity of the defect is influenced by genetic background. In a subset of mutant ovules the MMC either fails to divide meiotically or divides abnormally, leading to a lack of female gametophyte development. In contrast to the embryo sac defects, mutant pollen developed and functioned normally. Therefore, *FLP* and *MYB88* function as positive transcriptional regulators of entry into megasporogenesis.

**Materials and methods**

Plant material and growth conditions

The following allelic series of *FLP* were used in this study: *flp-1, flp-7, flp-8,* and *SALK 033970*. In addition, two double mutants were also analysed: *flp-1; myb88* and *flp-7; myb88*. *flp-7 and flp-8* are in the Landsberg *erecta* (*L. er*) background whereas *flp-1, flp-1; myb88, SALK 033970*, and *flp-7; myb88* are all in the Columbia-0 (*Col-0*) background as previously reported (Lai et al., 2005). Genotyping was performed as described, using primer combinations listed in Supplementary Table S1 available at JXB online (primer sequences are given in Supplementary Table S2). Various female gametophyte-specific marker lines (Supplementary Table S3) were introgressed into the *flp-7* background by crossing homozygous mutant plants to the marker lines and allowing the F1 and F2 plants to self-fertilize. PCR genotyping of the segregating F2 population was performed and the presence of the marker gene was identified by selecting seedlings on Murashige and Skoog (MS) plates containing kanamycin at 50 µg ml−1.

*Arabidopsis* seeds were cold treated for 3 d at 4 °C and germinated and grown on Fafard 2 mix soil (Fafard) under long-day (16 h, 80 µmol m−2 s−1) irradiance either in controlled growth chambers (Enconair Ecological Chambers Inc., Manitoba, Canada) or in growth rooms with subirrigation at 22 °C with 60% relative humidity. Seeds grown on plates were sterilized with 70% ethanol followed by 10% (v/v)
hypocthrile and 0.1% SDS, and placed on Petri dishes containing MS media (RP1 Corp.) with 1% plant agar with or without antibiotic. The plates were incubated in the dark at 4 °C for 5 d to achieve uniform germination and then moved to a CU-36L growth chamber (Percival Scientific Inc., Perry, IA, USA) and grown under long-day conditions (22 °C; 16 h photoperiod) unless noted. Seedlings were transplanted to soil at ~2 weeks old.

PCR-based genotyping

Genomic DNA was extracted as previously described (Teotia and Lamb, 2009). Primer combinations and primer sequences are shown in Supplementary Tables S1 and S2 at JXB online, respectively. PCR was done using Biolase RED DNA Polymerase (Bioline) on a conventional PCR machine (Bio-Rad-iCycler Thermal Cycler). flp-1, flp-7, and er alleles were identified by derived cleaved amplified polymorphic sequences (dCAPS; see Supplementary Table S1) (Neff et al., 1998, 2002). The flp-1 allele was genotyped as described (Xie et al., 2016a).

Seed set and fertility analysis

In order to analyse the seed set of the mutants and wild type, plants were grown under long-day conditions. Seeds of various genotypes were sown in 4 inch round pots. Four plants per genotype were used for comparing the percentage seed set. The total number of ovules and the number of seeds present in the first 15 siliques on the primary inflorescence only were counted, as previously described (Alvarez and Smyth, 1999). The percentage seed set was calculated by taking into account the total number of ovules and the number of seeds present in a silique. Statistical significance of the values was calculated as above using a Student’s t-test.

Reciprocal crosses were done to test male and female fertility by emasculating and hand pollinating 15 flowers of each genotype, as previously described (Unte et al., 2003). The number of seeds made in each silique was counted after the siliques were fully matured. Statistical significance of the values was calculated as above.

Aniline blue staining of pollen tubes and mucilage staining of seeds

Aniline blue staining of pistils was done according to Jiang et al. (2005). Flowers were emasculated just prior to pollination (late stage 12) and were grown for another 18–24 h to allow pollinating tract and ovule development to finish. Pistils were hand pollinated and grown for a further 24 h to allow pollen tube growth. Pollinated pistils were fixed in a solution of ethanol/acetic acid (3:1) for 2 h at room temperature, washed three times with ddH2O, softened in 8 M NaOH overnight, and washed in ddH2O several times before staining. Pistils were stained in aniline blue solution (0.1% aniline blue in 0.1 M K2HPO4–KOH buffer, pH 11) for 3 h in the dark. The stained pistils were observed and photographed with a Nikon Eclipse 80i compound microscope.

Mucilage production by flp-7 and flp-1; myb88 seeds was analysed according to Debeaujon et al. (2000). In brief, seeds were soaked in 0.03% ruthenium red (w/v) for 15 min and washed in water before being mounted on a slide and observed using a Nikon Digital Sight DS-5M camera on a Nikon SMZ800 dissecting microscope.

Ovule clearing and differential interference contrast (DIC) optics

Definitions for floral, ovule, and gametophyte development stages were as described in Smyth et al. (1990), Christensen et al. (1997), and Schneitz et al. (1995). Embryo sacs were collected from the entirety of the primary inflorescence and also secondary inflorescences. To examine mature ovules, flowers were emasculated just prior to pollination (late stage 12) and were grown for another 24 h. Ovule clearing and microscopy was done according to Rodrigo-Peiris et al. (2011), with some modifications. Pistils were opened along the carpel margins and fixed in a solution of 9:1 absolute ethanol:glacial acetic acid overnight at 4 °C followed by washing in 90% ethanol for 1 h, and stored in 70% ethanol until examination. Ovules were cleared in chloral hydrate (8 g of chloral hydrate and 2 ml of ddH2O to each ml of glycerol) on microscope slides for 2 h before microscopic analysis with DIC optics on a Nikon Eclipse 90i Microscope. Pictures were taken using the attached Nikon camera and analysed with NIS elements Advanced Research software version 3.0.

β-Glucuronidase (GUS) staining

Histochemical staining for GUS activity was performed as described (Jefferson et al., 1987). In brief, ovules were collected from pistils at different stages of flower development. The ovules were incubated in GUS staining buffer (50 mM sodium phosphate buffer pH 7.0, 10 mM EDTA, 2 mM potassium ferricyanide, 2 mM potassium ferrocyanide, 0.1% Triton X-100, and 2 mM 5-bromo-4-chloro-3-indolyl-β-d-glucuronic acid) at 37 °C for 72 h. Ovules were then washed in 90% ethanol for 1 h and stored in 70% ethanol until examination. Photographs were taken using a Nikon Digital Sight DS-5M camera attached to a Nikon Eclipse 80i compound microscope.

Confocal microscopy of ovules

Ovules were collected from pistils at different stages of flower development and were incubated in 0.5% propidium iodide in 50 mM phosphate buffer at room temperature or in 10 µg ml–1 of propidium iodide. Ovules were observed using either a Nikon D-Eclipse C1si or Nikon Eclipse 80i confocal microscope at excitation wavelengths of 488 nm and 543 nm. Emission was collected at 620–720 nm and 488–562 nm to visualize propidium iodide and green fluorescence protein (GFP) fluorescence, respectively.

Results

FLP and MYB88 are expressed in the flower

To characterize FLP and MYB88 expression during reproductive development, transgenic plants containing either a pFLP::GUS-GFP construct (Lai et al., 2005) or a pMYB88::GUS-GFP construct (Vanneste et al., 2011; a kind gift of Steffen Vanneste) were used. FLP expression was detected in unopened flower buds, at the bases of sepal, petals, and stamens and in the receptacle of carpels (Fig. 1A, 1B, and data not shown). pFLP::GUS-GFP is strongly expressed in the placenta within the ovary (Fig. 1C). Additionally, expression of FLP can be seen in both the style (Fig. 1D) and stigma (Fig. 1E) of the pistil. During ovule development, FLP has a dynamic expression pattern. During early stages of ovule development (before integuments are morphologically distinct), FLP expression was barely detectable (Fig. 1F). However, once integument outgrowth has begun, strong FLP expression is seen in the funiculus (Fig. 1G), which persists into later stages (Fig. 1H, 1I). Notably, the FLP promoter drives expression in the nucellus in younger ovules, where it is specifically expressed in the MMC and in epidermal cells (Fig. 1J). FLP is also expressed in the integuments, starting at a low level when they initiate (Fig. 1H). Later in ovule development it is expressed in the endothelial layer (the adaxial layer of the inner integument) and the outer layer of the outer integument, which will form the mucilage-containing seed coat cells (Fig. 1J). In contrast, little to no expression was seen in older anthers (Fig. 1B), consistent with Genevestigator data (Zimmermann et al., 2004, 2005). MYB88 is expressed at a much lower level than FLP (Lai et al., 2005). However, MYB88...
expression is detected in ovules (Fig. 1K) and in the embryo sac of stage 13 flowers (Fig. 1L).

**Loss of FLP and MYB88 reduces female fertility**

To investigate the roles of *FLP* and *MYB88* in reproductive development, fruit size and seed set were examined in different *flp* alleles and double mutants with *myb88*. The siliques of both *flp-7* and *flp-1; myb88* appeared shorter than their respective wild types (Fig. 2A, 2B). Mutant siliques contain small, white ovules that appear to be either aborted or not fertilized (Fig. 2A, 2B, 2D; Table 1), consistent with the presence of shorter siliques. The seed set of *flp* and *myb88* mutants was then compared with that of their respective wild type (*L. er* or Col-0). All examined *flp* and *flp; myb88* plants except *flp-1* have significantly reduced seed set (Table 1). Interestingly, *flp-7*, *flp-8*, *SALK_033970* (which is a loss-of-function allele in the *FLP* locus), and *flp-7; myb88* have significantly more ovules than the wild type, but still have reduced seed set (Table 1). This suggests that the ovary and placenta are larger in pistils of *flp* mutants, but that the reduced fertility leads to smaller fruits. These results show that *FLP* and *MYB88* regulate reproductive development. None of the available *flp* alleles is RNA null, although the *myb88* allele is a knockdown (Lai et al., 2005). This makes it difficult to assess
in flp-7 are more severe than those of flp-1 (Lai et al., 2005). As flp-7 (L. er) and flp-1; myb88 (Col-0) display significantly reduced seed set and are in two different genetic backgrounds, they were selected for further studies.

The strong expression of FLP in integuments (Fig. 11, 1J) suggests that this gene may be involved in development of the seed coat. Mature seeds of flp-7 and flp-1; myb88 were compared with their respective wild types. Although some minor variability in seed shape was observed in the mutant seeds (Supplementary Fig. S1 at JXB online), particularly flp-7, no noticeable germination defects were observed (data not shown). In addition, mucilage was produced by both flp-7 and flp-1; myb88 seeds (Supplementary Fig. S1), suggesting that differentiation of the seed coat is not significantly altered.

To determine whether the reduced seed set is due to male and/or female infertility, reciprocal crosses were made using flp-7 or flp-1; myb88 homozygous plants and the wild type. Both flp-7 and flp-1; myb88 siliques harboured fewer seeds than their respective wild types when pollinated with either wild-type or mutant pollen (Table 2), indicating that female fertility is compromised. flp-7 pollen, when used to fertilize wild-type pistils, resulted in slightly reduced seed set (Table 2). However, the fertility of flp-1; myb88 pollen was comparable with that of the wild type. Taken together, these results suggest that FLP and MYB88 function in female reproduction.

Embryo sac development is altered by loss of FLP and MYB88

The aborted ovules seen in siliques of flp mutants could be caused by defects in ovule and/or embryo sac development or by lack of fertilization. Stigmatic and transmitting tract tissues in the carpel are required for proper pollen tube growth and fertilization. Mutants with reduced growth of these tissues have reduced fertility due to poor pollen tube growth (Heisler et al., 2001; Gremski et al., 2007). FLP is strongly expressed in the stigma and style of the pistil (Fig. 1D, 1E), suggesting that it could function in these tissues. In order to examine whether pollen tube growth is affected in flp mutants, carpels were stained with aniline blue 24 h after pollination. In both L. er and flp-7 pistils, pollen tubes grew throughout the transmitting tract and into ovules (Supplementary Fig. S2 at JXB online), indicating that the reduced seed set in flp alleles is not due to defects in either the stigma or transmitting tract.

Since pollen tubes were able to travel to the ovules, the reduced fertility seen in flp-7 and flp-1; myb88 plants is probably due to ovule and/or female gametophyte defects. Therefore, ovules were examined 2 d after emasculating flowers at stage 13 (according to Smyth et al., 1990) to determine whether ovule development proceeded normally. The morphology of mutant ovules at female gametophyte developmental stage 7 (FG7; Christensen et al., 1997) appeared normal, with fully developed outer and inner integuments and well-differentiated proximal-distal polarity (Fig. 3B, 3D), suggesting that ovule development is intact in these mutants. The FG7 embryo sacs of flp-7 and flp-1; myb88 were then compared with their respective wild types, L. er and Col-0. Ninety percent of L. er (n = 281) and 97% of Col-0 (n = 607) ovules display four visible

---

**Fig. 2.** Fertility is lowered in flp-7 and flp-1; myb-88 plants. Dissecting microscope images of siliques. (A) flp-7 fruit are shorter than those of the wild type and contain aborted ovules (arrowheads). (B) flp-1; myb88 fruit are shorter than those of the wild type and contain aborted ovules. (C) L. er siliques containing many seeds. (D) flp-7 siliques containing a few seeds and several aborted ovules (arrowheads).
and 78% of synergids (Fig. 3E). Although Polygonum-type embryo sacs, nuclei, corresponding to the egg cell, central cell, and two synerids (Fig. 3E). Although Polygonum-type embryo sacs, such as those found in Arabidopsis, contain seven cells of four types (Yadegari and Drews, 2004), in Arabidopsis the three antipodal cells have degenerated by FG7 (Christensen et al., 1997). In contrast to the wild type, only 56% of flp-7 (n=337) and 78% of flp-1; myb88 (n=589) embryo sacs had four discernible cells. Strikingly, 44% and 22% of the mutant ovules appeared to contain cellular structures harbouring one or more prominent large cells and did not resemble mature female gametophytes (Fig. 3B, 3D). Another growth difference between the abnormal flp-7 and flp-1; myb88 ovules and wild-type ovules was also observed. In the wild type, the growth and expansion of the embryo sac is accompanied by the degeneration of cells in the proximal nucellar region. However, in the abnormal flp ovules, these proximal cells persist (Fig. 3B, 3D).

In order to identify and compare the cell types present in embryo sacs at FG7 in wild-type and mutant ovules, specific markers were crossed into the flp-7 mutant background. ET884 expression, while in flp-7 only 14% of the observed ovules (n=356) expressed the synergid marker. No expression of ET884 was observed in those flp-7 ovules in which no embryo sac could be distinguished (Fig. 4B). The egg cell-specific marker ET1119 (Gross-Hardt et al., 2007) was also examined. Relatively few flp-7 ovules sampled expressed this marker (3%, n=430) (Fig. 4F). In contrast, 48% of wild-type ovules sampled (n=250) displayed ET1119 expression while the remainder did not. ET1119 expression was never seen in ovules with abnormal embryo sac development (Fig. 4E). Finally, expression of two markers of central cells, pMEA::GUS (Gross-Hardt et al., 2007) and AGL61::GFP (Steffen et al., 2008), was analysed in wild-type and flp-7 ovules. Consistent with the above results, the expression of the central cell markers was less frequent in flp-7 than in wild-type ovules (Fig. 4I; Supplementary Fig. S3 at JXB online). pMEA::GUS expression in the wild type was found in 49% of ovules (n=279) but only in 12% (n=243) of flp-7 ovules. No pMEA::GUS expression was seen in ovules containing abnormal embryo sacs (Fig. 4H). Results with AGL61::GFP were similar (Supplementary Fig. S3). Taken together, these data suggest that the aborted flp-7 ovules lack differentiated female gametophytes and thus are incapable of being fertilized.

**Comparison of structural development of the female gametophyte in the flp-7 mutant and wild type**

To investigate the basis of the flp-7 phenotype and the origin of the large cells seen in FG7 of ovules of the mutant, a stage wise comparison of development was carried out by examining cleared ovules using DIC microscopy. Stages of ovule and female gametophyte development are summarized in Supplementary Table S4 at JXB online, based on Smyth et al. (1990) and Schneitz et al. (1995). Early stages were normal in flp-7, such as the appropriate initiation of ovule primordia (data not shown). In addition, both wild-type and flp-7 pre-meiotic ovules (stage 2-III of ovule development; see Supplementary Table S4) displayed clearly differentiated MMCs (Fig. 5A, 5D). In L. er wild type stage 2-V, after meiosis, a clear tetrad of megaspores can be seen (Fig. 5B, Table 3). However, many of the

---

### Table 1. flp and flp; myb88 mutants have reduced fertility that is influenced by genetic background

<table>
<thead>
<tr>
<th>Genotype</th>
<th>Ovules per silique</th>
<th>Seeds per silique</th>
<th>Aborted ovules per silique</th>
<th>Seed set</th>
</tr>
</thead>
<tbody>
<tr>
<td>L. er</td>
<td>45 ± 0.6</td>
<td>42 ± 1.0</td>
<td>3 ± 0.6</td>
<td>94%</td>
</tr>
<tr>
<td>L. er/pER::ER</td>
<td>58 ± 0.8</td>
<td>53 ± 0.7</td>
<td>5 ± 0.3</td>
<td>92%</td>
</tr>
<tr>
<td>flp-7</td>
<td>61 ± 0.9**</td>
<td>4 ± 0.7**</td>
<td>57 ± 1.3**</td>
<td>6%</td>
</tr>
<tr>
<td>flp-7/pER::ER</td>
<td>54 ± 1.4***</td>
<td>17 ± 0.5****</td>
<td>40 ± 1.3****</td>
<td>31%</td>
</tr>
<tr>
<td>flp-8</td>
<td>66 ± 0.8**</td>
<td>4 ± 2.1</td>
<td>20 ± 0.6**</td>
<td>70%</td>
</tr>
<tr>
<td>Col-0</td>
<td>48 ± 0.5</td>
<td>39 ± 1.5</td>
<td>9 ± 1.5</td>
<td>80%</td>
</tr>
<tr>
<td>flp-1</td>
<td>47 ± 0.7</td>
<td>34 ± 1.7</td>
<td>13 ± 1.6</td>
<td>63%</td>
</tr>
<tr>
<td>SALK_033970/flp</td>
<td>54 ± 0.8**</td>
<td>37 ± 3.4</td>
<td>17 ± 2.7*</td>
<td>63%</td>
</tr>
<tr>
<td>flp-1; myb88</td>
<td>49 ± 0.7</td>
<td>14 ± 2.4**</td>
<td>35 ± 2.1**</td>
<td>27%</td>
</tr>
<tr>
<td>flp-7; myb88</td>
<td>68 ± 0.7**</td>
<td>47 ± 3.1*</td>
<td>21 ± 3.2**</td>
<td>69%</td>
</tr>
</tbody>
</table>

**a** Values are means ±SE (n = 60) or means ±SE of n = 60 for three independent transgenic lines (180 siliques in total) for L. er/pER::ER and flp-7/pER::ER.

**b** Defined as small white ovules with no evidence of embryo or seed development.

*Values that are significantly different from wild type at P < 0.05. **Values that are significantly different from the wild type at P < 0.01. ***Values that are significantly different from L. er/pER::ER at P < 0.01.

---

### Table 2. Loss of FLP and MYB88 compromises female fertility

<table>
<thead>
<tr>
<th>Female parent</th>
<th>Male parent</th>
<th>Seeds per crossed silique</th>
</tr>
</thead>
<tbody>
<tr>
<td>L. er</td>
<td>L. er</td>
<td>43.7 ± 6.1**</td>
</tr>
<tr>
<td>flp-7</td>
<td>L. er</td>
<td>37.8 ± 25.0*</td>
</tr>
<tr>
<td>flp-7</td>
<td>flp-7</td>
<td>14.6 ± 6.2**</td>
</tr>
<tr>
<td>Col-0</td>
<td>Col-0</td>
<td>40.3 ± 4.7</td>
</tr>
<tr>
<td>flp-1; myb88</td>
<td>flp-1; myb88</td>
<td>7.3 ± 9.4**</td>
</tr>
</tbody>
</table>

**a** Values are the mean ±SE (n = 15).

*Values that are significantly different from wild type at P < 0.05.

**Values that are significantly different from the wild type at P < 0.01.
Fig. 3. Loss of FLP and/or MYB88 leads to abnormal nucellar structures. (A–D) DIC micrographs of FG7 ovules containing mature female gametophytes. (A) Col-0. (B) flp-1; myb88. (C) L. er. (D) flp-7. (E) Quantification of embryo sac defects. CC, central cell; CE, chalazal end of the embryo sac; EC, egg cell; ME, micropylar end of the embryo sac. Arrows indicate synergid cells, and arrowheads indicate large cells found in abnormal flp ovules in the region where an embryo sac would normally form.
flp-7 ovules at comparable stages contain an abnormal number of cells in the region where the megaspores should be, including one large cell (Fig. 5E, Table 3). Normally during stage 3-I, the three non-functional megaspores begin to degenerate, leaving a mononuclear embryo sac (Fig. 5C). However, in similarly staged flp-7 ovules, no cellular degeneration was found (Fig. 5F, Table 3). At stage 3-II, the embryo sac normally undergoes its first mitotic division producing a cell with two nuclei (Fig. 5G). Abnormal flp-7 ovules display no such nuclear division and contain a single larger MMC- or megaspore-like cell as well as no (Fig. 5M), two (Fig. 5J), or three (Fig. 5N) other cells nearby. By stage 3-IV, the wild-type female gametophyte contains four nuclei (Fig. 5H), but these were not present in flp-7 ovules (Fig. 5K). At stage 3-VI the embryo sac begins differentiation and cellularization in L. er (Fig. 5I). However, in abnormal flp-7 ovules, instead of an embryo sac forming, a single large cell can be seen and is associated with a variable number of other cells (Fig. 5L, 5O). These results suggest that the MMC meiosis either did not occur or was abnormal.

The lowered expressivity of embryo sac defects in flp-1; myb88 mutants made a detailed developmental study of embryo sac defects difficult. However, examination of mature embryo sacs of this genotype reveals that many (22%) contained abnormal structures that do resemble embryo sacs but rather contain a large cell resembling the MMC in position and size (Fig. 3D). This suggests that, similar to flp-7, flp-1; myb88 MMCs undergo abnormal meiosis.

The expressivity of female gametophytic defects caused by loss of FLP and/or MYB88 is influenced by genetic background

Although a number of flp alleles and allelic combinations with myb88 show significant reductions in fertility from their wild
types, there is variability in the expressivity of the phenotype, with seed set varying from 6% to 72% (Table 1), although the penetrance is complete (data not shown). Examination of the pattern of severity reveals that flp alleles in the L. er ecotype background exhibit more phenotypic severity than those in Col-0 (Table 1). In addition, crossing flp-7 into the Col-0 background reduces the severity of the fertility defect (Table 1), supporting the hypothesis that either the L. er background harbours enhancer(s) of the flp phenotype or Col-0 has suppressors of the flp phenotype, or both.

Many genetic differences exist between L. er and Col-0, including both single nucleotide polymorphisms and larger scale indels (Schmid et al., 2003; Ziolkowski et al., 2009). A prominent difference is the presence of a mutated ERECTA (ER) locus in L. er. ER and its family members, ERECTA LIKE1 (ERL1) and ERECTA LIKE2 (ERL2), encode leucine-rich repeat (LRR) domain-containing receptor-like kinases. They function in the stomatal patterning along with the TOO MANY MOUTHS (TMM) LRR receptor-like protein (Nadeau and Sack, 2002; Shpak et al., 2005). ER and its family members act upstream of FLP and MYB88. ER is the most important gene in its family and masks the functions of its paralogues. The ER family members also function in female reproductive development (Pillitteri et al., 2007). In er-105; erl1-2; erl2-1/+ ovules, cell proliferation in integuments is reduced, gametophytes abort, and cyclin A-encoding genes are misregulated. It

Fig. 5. Stages of female gametophyte development in the wild type and flp-7 mutant. Ovule stages were determined from the development of the sporophyte using the nomenclature of Schneitz et al. (1995); see Supplementary Table S4 at JXB online. (A–O) DIC micrographs of ovules at different stages. (A, B, C, G, H, I) L. er. (D, E, F, J, K, L, M, N, O) flp-7. (A, D) Stage 2-III (pre-meiotic) with large MMCs near the tip of the nucellus (arrows). (B) Stage 2-V with tetrad of four megaspores (primary megaspore indicated by a dashed circle) in L. er. (C) At stage 3-I, the three non-functional megaspores begin to degenerate in L. er. (E) Varying numbers of cells are seen in flp-7 ovules. Here, three cells can be seen (two toward the apex, arrowheads; and a larger cell more basally, asterisk). (F) The abnormal cells formed in flp-7 ovules do not appear to be degenerating.
was hypothesized that the mutant er locus in L. er influences the expressivity of flp alleles and may sensitize the ovule to loss of FLP function. To test these hypotheses, the er allele was complemented in the flp-7 mutant background by transforming L. er and flp-7 plants with an ER genomic rescue construct (a kind gift of Dr Keiko Torii). Complementing the defect in er in the flp-7 background increased the seed set from 6% to 31% (Table 1), suggesting that this mutation accounts for some of the increased expressivity of female gametophyte defects in flp-7 compared with flp-1; myb88. Restoration of wild-type ER activity also reduced the number of ovules per silique (Table 1) and reduced the severity of the stomatal defect (data not

Fig. 5. Continued

(G) At stage 3-II, the first mitotic division of the megaspore to give rise to a two-nuclei (circled by dashed lines) female gametophyte in L. er. (J, M, N) No corresponding nuclear division can be seen in abnormal stage 3-II flp-7 ovules; instead, a large cell remains with either zero (M), two (J), or three (N) other cells attached. (H) At stage 3-IV, a female gametophyte with four nuclei (dashed circles) is seen in L. er. In addition, the other cells of the nucellus have begun degenerating. (K) No such structures were seen in flp-7 ovules; instead the abnormal cells remain. (I) By stage 3-VI, the embryo sac has begun differentiating and cellularizing in L. er. (L, O) In flp-7 a single large cell is associated with a variable number of other cells (three in L, two in O). Asterisks indicate MMC-like/functional megaspore-like cells seen in flp-7 ovules. Arrowheads indicate megaspores in L. er and cells associated with the MMC-like/functional megaspore-like cell in flp-7.
shown). However, ER complementation did not restore fertility to wild-type levels, suggesting that other genetic differences account for the remaining difference in expressivity between L. er and Col-0.

Discussion

In all sexually reproducing organisms, haploid gametes are formed after meiosis, either directly as in animals or indirectly after mitotic divisions as in plants. Therefore, meiosis is the key step leading to gametogenesis (van Werven and Amon, 2011). In angiosperms, although there is sexual dimorphism between the MMC and the male microspore mother cell, most mutations affecting meiosis affect both sexes. However, not all genes implicated in meiosis affect both male and female reproductive development (Ma, 2006). Broadly speaking, genes that have roles in megasporogenesis and megagametogenesis can be divided into sporophytic-acting and gametophytic-acting groups. Sporophytically acting genes with specific roles in regulating gametogenesis exhibit defects in spore and/or gamete formation but not in sporophytic parts of the ovule (Schneitz et al., 1997).

In this work, it was shown that a variety of loss-of-function, although not RNA null, flp alleles alone and in combination with myb88 exhibit reduced seed set due to reduced female fertility (Fig. 2, Tables 1, 2). A defect in female but not male fertility is consistent with the low expression of FLP and MYB88 in stamens and pollen (Zimmermann et al., 2004, 2005; Lai et al., 2005; this study). The female infertility of flp and flp; myb88 mutants is not due to major defects in overall ovule morphology (Fig. 3B, 3D). In addition, mucilage is produced normally by mutant seed coats (Supplementary Fig. S1 at JXB online), suggesting that seed coat differentiation is relatively normal. However, examination of mature ovules revealed that abnormal flp-7 and flp-1; myb88 ovules did not contain embryo sacs; rather, they contained cellular structures with a larger single cell located close to the micropylar end of the ovule (Fig. 3B, 3D). These structures do not contain differentiated synergids, egg cells, or central cells (Fig. 4; Supplementary Fig. S3). Examination of earlier stages in ovule development revealed that these abnormal ovules arise due to a partially expressed defect in entry into meiosis by the MMC and/or abnormal meiosis and lack of megaspore differentiation (Fig. 5). Several lines of evidence support this interpretation. First, mature mutant ovules harboured a single large cell that resembles an MMC or a megaspore in position and size which was accompanied by variable numbers of cells that appeared to arise from the same division (Figs 3, 5). Secondly, the retention of proximal nucellar cells in abnormal flp-7 and flp-1; myb88 suggests that embryo sacs are absent, consistent with phenotypes seen before in other embryo sac absent mutants, including those with defects in meiosis of the MMC (Siddiqi et al., 2000). Thirdly, the smaller of the presumed division products of the MMC seen in flp mutants are positioned nearer the antipodal end of the ovule, where the non-functional megaspores are positioned in the wild type, while the large cell is nearer to the chalazal end, where the functional megaspore operates (Figs 3, 5). However, these smaller cells do not degenerate as normally the smaller megaspores would. Similar observations have been reported in Arabidopsis mutants where meiosis is incomplete or otherwise abnormal (Chen et al., 2011). Taken together, these results suggest that the paralogous FLP and MYB88 transcription factors function in the ovule to control entry into and progression through megasporogenesis, although they do not appear to function in male reproductive development, consistent with expression analysis.

FLP and MYB88 are required to limit cell divisions in the staminal lineage; flp and flp; myb88 double mutant GMCs undergo extra divisions, resulting in staminal clusters (Yang and Sack, 1995; Lai et al., 2005; Xie et al., 2010a). They perform this function by regulating expression of cell cycle genes (Xie et al., 2010a; Vanneste et al., 2011). FLP directly represses expression of the CDKB1;1 gene (encoding a cyclin-dependent kinase that promotes entry into mitosis) after the symmetric division of the GMC, thereby preventing further division and forming a functional two-celled stoma (Xie et al., 2010a). The A2-type cyclin CYCA2;3 can form a functional complex with CDKB1;1 (Boudolf et al., 2009) and its expression is also directly and coordinately repressed by FLP/MYB88 in young guard cells (Vanneste et al., 2011). It has been demonstrated here that a larger number of ovules are formed in flp and flp; myb88 mutants (Table 1). The molecular events that regulate development of the placenta, from which ovules are formed, is not well understood; however, an increase in the number of ovules suggests either

<table>
<thead>
<tr>
<th>Ovule stage</th>
<th>Genotype</th>
<th>One cell</th>
<th>Two cells</th>
<th>Four cells</th>
<th>&gt;4 cells</th>
<th>Total ovules observed</th>
</tr>
</thead>
<tbody>
<tr>
<td>3-I L. er</td>
<td>57% (69)</td>
<td>31% (54)</td>
<td>10% (17)</td>
<td>2% (3)</td>
<td>173</td>
<td></td>
</tr>
<tr>
<td>3-II flp-7</td>
<td>68% (186)</td>
<td>15% (40)</td>
<td>17% (47)</td>
<td>NO</td>
<td>273</td>
<td></td>
</tr>
<tr>
<td>3-IV L. er</td>
<td>6% (3)</td>
<td>NO</td>
<td>55% (27)</td>
<td>39% (19)</td>
<td>49</td>
<td></td>
</tr>
<tr>
<td>3-VI L. er</td>
<td>1% (2)</td>
<td>67% (137)</td>
<td>26% (64)</td>
<td>205</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

- a Stages according to Schneitz et al. (1995).
- b The number in parentheses indicates the number of ovules observed.
- c NO, none observed.
increased or extended proliferation to create a larger placenta or more entry into organogenesis by cells of this tissue. It is possible that the placental phenotype seen in flp; myb88 mutants might also be due to lack of repression of similar cell cycle genes they repress during stomatal development. Alternatively, it has been shown that down-regulation of cytokinin signalling and metabolism takes place within the stomatal lineage and that this might be important for the transition from proliferation to differentiation (Pillitteri et al., 2011). Mutations in the CKX3 and CKX5 genes, which encode cytokinin oxidase/dehydrogenase enzymes functioning in degradation of this hormone, lead to increased proliferation in the placenta, supernumerary ovules, and increased seed set (Bartrina et al., 2011), suggesting that levels of cytokinin are important to regulate the size of this tissue. Therefore, FLP and MYB88 might also influence ovule number by regulating cytokinin signalling or homeostasis. Orthologues of FLP and MYB88 in seed crop plants could be involved in control of ovule number (and therefore yield) and would be interesting targets for modification in such crops.

The role of cell cycle genes in controlling Arabidopsis meiosis is not well understood. In mammals, type A1 cyclins have been shown to function in the meiotic cell cycle (Wolgemuth, 2011). This appears to be true for Arabidopsis. CYCA1;2/TARDY ASYNCHRONOUS MEIOSIS (TAM) is required for entry into both the first and second meiotic divisions (d’Erfurth et al., 2010). Such genes may be potential targets of FLP/MYB88. Suppression of mitotic genes such as CDKB1;1 and/or CYCA2;3 by FLP/MYB88 could also be necessary to allow entry into meiosis. This would be consistent with the necessity to suppress these genes for entry into the endocycle (Boudolf et al., 2009) and is also consistent with the finding that meiotic arrest can be caused by a failure to regulate CDK activity appropriately (Bulankova, 2010). It is reasonable to hypothesize that FLP and MYB88 function to regulate expression of cell cycle genes controlling meiotic entry and progression. During both stomatal development (Lai et al., 2005; Xie et al., 2010a) and placenta development (this study), FLP and MYB88 appear to inhibit expression of genes that positively regulate cell division. Based on the defects seen in meiosis in flp mutants (Fig. 5), they would appear to promote meiotic division, but might also do this by repression of mitotic promoting factors.

Loss of FLP or FLP/MYB88 function does not lead to complete loss of female fertility. This incomplete expressivity could be due to a number of factors. The FLP alleles available are not RNA nulls (Lai et al., 2005). Therefore, it is possible that none of them is functionally null and there is residual FLP function present that is able to support meiosis in many ovules. Alternatively, other as yet unidentified genes could be partially functionally redundant with FLP/MYB88, allowing some female fertility in the flp; myb88 mutants. However, it is clear that genetic background impacts the expressivity of the loss of FLP and/or MYB88 function. Although all the flp alleles examined have reduced seed set, the flp-7 allele in the L. er ecotype is the most severe, even more than the similar flp-1 allele in Col-0 or flp-7; myb88 in which flp-7 has been introgressed into the Col-0 background (Table 1). Interestingly, the flp-7 stomatal phenotype is also stronger than that of flp-1 (Lai et al., 2005). Differences between genetic backgrounds in mutant penetrance, expressivity, and/or phenotype have been documented in Arabidopsis previously (e.g. Sedbrook et al., 2004; Sugliani et al., 2009). Many differences are present between L. er and Col-0 (Schmid et al., 2003; Ziolkowski et al., 2009), including in the ER gene. ER and its family members ERL1 and ERL2 have diverse roles coordinating cell proliferation with differentiation (Tori et al., 1996; Shpak et al., 2003, 2004, 2005; Pillitteri et al., 2007; Hord et al., 2008). ER family members act in complexes with TMM in GMCs (Lee et al., 2012) and act synergistically in enforcing stomatal patterning by interacting with the secreted peptides EPIDERMAL PATTERNING FACTORS (EPFs) (Hara et al., 2007, 2009; Hunt and Gray, 2009; Lee et al., 2012). These peptides function upstream of FLP/MYB88 in the stomatal lineage (Shpak et al., 2005). ER family members also ensure the proper growth of integuments and the progression of the mitotic cell cycle in the female gametophyte in a dosage-dependent manner (Pillitteri et al., 2007). The mutant er allele found in the L. er background might increase sensitivity to loss of FLP function. Indeed, when this mutation was complemented in the flp-7 background, ovule number was reduced, seed set was improved (Table 1), and the stomatal defects were ameliorated (data not shown). The smaller number of ovules which form when ER function is restored is expected since ER also can control ovule number per fruit (Alonso-Blanco et al., 1999). The present results imply that ER also regulates entry of the MMC into meiosis. Complementation results support the hypothesis that loss of ER function provides a sensitized background for the loss of other genes, such as FLP, that function in common pathways. However, additional genetic changes must also contribute to the differential phenotypic expressivity, since ER complemented flp-7 plants still display more severe phenotypes than similar mutations in the Col-0 background.

**Supplementary data**

Supplementary data are available at *JXB* online.

Supplementary Figure S1. Mucilage production is intact in flp-7 and flp-1; myb88 seeds.

Supplementary Figure S2. Pollen tube growth is not inhibited in flp-7 pistils.

Supplementary Figure S3. Abnormal flp-7 ovules do not express AGL61:GFP.

Supplementary Table S1. Primer combinations used for PCR genotyping.

Supplementary Table S2. Primers used in this study.

Supplementary Table S3. Female gametophyte markers used in this study.

Supplementary Table S4. Stages of floral, ovule, and gametophyte development.

**Acknowledgements**

The authors thank Dr Zidian Xie for advice and sharing of data before publication, Qin Lei for technical assistance, and two anonymous reviewers, members of the Lamb lab, and Dr Iris Meier (Ohio State University) for discussions. Dr Steffen Vanneste (Ghent University), Dr Ueli Grossniklaus (University of Zurich), and Dr Gary Drews (University of Utah) provided seeds, and
Dr Keiko Torii provided the ERp::ER plasmid used for complementation experiments. This work was supported in part by a grant from the National Science Foundation to RSL and funds from The Ohio State University.

References


