

1 **Characterization of the single *FERONIA* homolog in *Marchantia***
2 ***polymorpha* reveals an ancestral role of CrRLK1L receptor kinases**
3 **in regulating cell expansion and morphological integrity**

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25 **Short title: Deciphering the ancestral role of CrRLK1L receptors in land plants**

26 **Summary**

27 Plant cells are surrounded by a cell wall, a rigid structure rich in polysaccharides and
28 glycoproteins. The cell wall is not only important for cell and organ shape, but crucial for
29 intercellular communication, plant-microbe interactions, and as a barrier to the
30 environment. In the flowering plant *Arabidopsis thaliana*, the 17 members of the
31 *Catharanthus roseus* RLK1-like (*CrRLK1L*) receptor kinase subfamily are involved in a
32 multitude of physiological and developmental processes involving the cell wall, including
33 reproduction, hormone signaling, cell expansion, innate immunity, and various stress
34 responses. Due to genetic redundancy and the fact that individual *CrRLK1L*s can have
35 distinct and sometimes opposing functions, it is difficult to assess the primary or ancestral
36 function of *CrRLK1L*s. To reduce genetic complexity, we characterized the single
37 *CrRLK1L* gene of *Marchantia polymorpha*, *MpFERONIA* (*MpFER*). Plants with reduced
38 *MpFER* levels show defects in vegetative development, i.e., rhizoid formation and cell
39 expansion, but also affect male fertility. In contrast, *Mpfer* null mutants and
40 overexpression lines severely affect cell integrity and morphogenesis of the gametophyte.
41 Thus, the *CrRLK1L* gene family originated from a single gene with an ancestral function
42 in cell expansion and the maintenance of cellular integrity. During land plant evolution,
43 this ancestral gene diversified and was recruited to fulfil a multitude of specialized
44 physiological and developmental and roles in the formation of both gametophytic and
45 sporophytic structures essential to the life cycle of flowering plants.

46

47 **Key Words**

48 cell wall integrity, cell elongation, *CrRLK1L* receptor kinases, *Marchantia polymorpha*,
49 signaling pathway evolution

50 Introduction

51 The functioning of plant cells is contingent upon the cell wall acting as a barrier between
52 the cell and its environment. This complex matrix of polysaccharides, glycoproteins, and
53 other organic compounds defines the growth and shape of a cell and provides protection
54 against biotic and abiotic stresses [1,2]. Moreover, its stiffness has to resist turgor
55 pressure while also allowing cell expansion during growth. Thus, sensing and controlling
56 cell wall integrity (CWI) is crucial for plant cells and receptor kinases (RKs) play an
57 important role in sensing extracellular cues that activate intracellular pathways. Different
58 RK subfamilies are defined by their extracellular domain (ECD), which is more variable
59 than their transmembrane and kinase domains [3]. Over the last decade, the *CrRLK1L*
60 subfamily emerged as important CWI sensors [4,5].

61 Functions have been assigned to 15 of the 17 *CrRLK1L* members encoded in the *A.*
62 *thaliana* genome: they are required for cell elongation and cell shape, sensing
63 environmental cues, and cell-cell communication in diverse contexts [6–22]. For instance,
64 *FERONIA* (*AtFER*), *THESEUS1* (*AtTHE1*), *HERCULES1* (*AtHERK1*), and *AtHERK2* are
65 required for cell elongation as *Atfer* single, *Atthe1Atherk1* double, and
66 *Atthe1Atherk1Atherk2* triple mutants exhibit stunted growth [7,9–11]. Root hairs burst in
67 mutants affecting *AtFER* and *[Ca²⁺]_{cyt}-ASSOCIATED PROTEIN KINASE1*
68 (*AtCAP1*)/*ERULUS* (*AtERU*) [12,16]. Moreover, *AtFER* plays a role in powdery mildew
69 resistance [11], innate immunity [15,22], calcium signaling [23], phytohormone signaling
70 [10,24–26], and mechano- and heavy metal sensing [20,27]. In most contexts, the
71 *CrRLK1L*s tend to promote cellular growth and cell elongation. In contrast, *AtTHE1*
72 actively inhibits cell elongation in hypocotyls when cell wall perturbations occur, *e.g.*, due
73 to mutations in the cell wall biosynthesis machinery or upon treatment with isoxaben, a
74 cellulose synthesis inhibitor [7,28].

75 In addition to regulating cellular growth during the vegetative phase of the life cycle,
76 several *CrRLK1*Ls play a role during fertilization by controlling the growth and reception
77 of the pollen tube. In the female gametophyte, *AtFER* is highly expressed in the two
78 synergid cells that flank the egg cell and are important for double fertilization [6,29].
79 *AtFER* is crucial for pollen tube reception and the release of the two sperm cells
80 [6,10,23,30,31], a process also involving two other, redundantly acting synergid-
81 expressed *CrRLK1*Ls, *AtHERK1* and *ANJEA* (*AtANJ*) [17]. In contrast to these synergid-
82 expressed *CrRLK1*Ls, four pollen-expressed *CrRLK1*Ls, named *ANXUR1* (*AtANX1*),
83 *AtANX2*, *BUDDHA'S PAPER SEAL1* (*AtBUPS1*) and *AtBUPS2* [8,14,18,32], are
84 redundantly required for pollen tube integrity and tip growth.

85 To transduce extracellular cues, *CrRLK1*Ls bind secreted peptides of the RAPID
86 ALKALINIZATION FACTOR (RALF) family [22,32–35], which induce complex formation
87 with members of LORELEI (LRE) family of GPI-anchored proteins [32,35–37]. Hence, it
88 was suggested that at least some *CrRLK1*Ls act as CWI sensors that coordinate cell
89 elongation in response to changes in the cell wall [38–43].

90 The 17 *A. thaliana* *CrRLK1*Ls act partially redundantly and can have opposite effects on
91 cellular growth (reviewed in [19,41,43,44]). To investigate the original function of
92 *CrRLK1*Ls, we characterized them in a system with reduced genetic complexity: the
93 liverwort *Marchantia polymorpha* encodes a single *CrRLK1*L [45,46]. Liverworts represent
94 an early diverging land plant lineage and have been hypothesized to retain, at least in
95 part, characteristics of the earliest land plants [45,47–52]. The presence of similar types
96 of gene families in all plant lineages suggests that the differences in development evolved
97 by co-opting and modifying existing developmental programs and genetic networks,
98 rather than through the evolution of novel genes [47,48,53]. *M. polymorpha* is thus an

99 ideal system to study the ancestral role of genes and how it was modulated and diversified
100 during land plant evolution [45,54,55].

101 We refer to the a single *CrRLK1L* family member encoded in the *M. polymorpha* genome,
102 Mp4g15890 (Mapoly0869s0001), as Mp*FERONIA* (Mp*FER*) (aka Mp*THESEUS*, Mp*THE*)
103 [44–46,56]. An Mp*fer/the* mutant was identified in a large T-DNA screen for defective
104 rhizoid formation, developing short and irregularly shaped rhizoids with brown tips,
105 indicating rhizoid rupture [56]. In the same screen, a mutant affecting the ortholog of the
106 *A. thaliana* receptor-like cytoplasmic kinase *MARIS* (*AtMRI*), known to act downstream of
107 *AtFER*, was identified with a similar phenotype (named Mpmri aka Mppti, Mp7g17560
108 [Mapoly0051s0094]) [56,57]. As in *A. thaliana*, a dominant active version of Mp*MRI* can
109 partially rescue the Mp*fer* rhizoid phenotype [57]. This suggests that at least some of the
110 machinery associated with cell elongation and CWI sensing is conserved between *A.*
111 *thaliana* and *M. polymorpha*.

112 Here, we report that *CrRLK1L* function is required in a variety of aspects of *M. polymorpha*
113 gametophyte development. Phylogenetic analyses suggest that *CrRLK1L* genes first
114 appeared in the common ancestor of land plants. The characterization of lines with
115 reduced Mp*FER* levels indicates that, in addition to its function in rhizoid formation,
116 Mp*FER* controls cell size and organ growth and is involved in male gametogenesis and
117 fertility. Analysis of the Mp*FER* expression pattern points to an involvement in female
118 sexual organogenesis and sporophyte development. Our data suggests that the broad
119 involvement of *CrRLK1L* function in the development and physiology of land plants is an
120 ancestral and conserved characteristic, although the functions of *CrRLK1L*s were
121 extended and adapted to control additional developmental processes in the sporophyte
122 in the course of land plant evolution.

123

124 **Results**

125 **CrRLK1L is conserved in land plants and arose in this lineage**

126 The *M. polymorpha* genome encodes a single CrRLK1L homolog (Mp4g15890
127 [Mapoly0869s0001]) [44–46]. Like other CrRLK1L members, this gene encodes an
128 intracellular serine/threonine kinase domain, a transmembrane domain, and an
129 extracellular malectin-like domain with two malectin domains (Figure 1A). Given that the
130 ECD of CrRLK1Ls in *A. thaliana* is crucial for their function and, in contrast to the kinase
131 domain, not interchangeable [58], we performed a phylogenetic analysis with the amino
132 acid sequences of the predicted malectin-like domain of Mp4g15890, the five members
133 of the moss *Physcomitrella patens*, the two of the lycophyte *Selaginella moellendorffii*,
134 and the 17 CrRLK1Ls of *A. thaliana* (Figure 1B). The *M. polymorpha* CrRLK1L member
135 grouped together with the CrRLK1Ls of basal land plants and was in the same clade as
136 AtFER, while AtTHE1 was in a different one (Figure 1B). Based on this phylogenetic
137 position and the fact that the malectin-like domain of the *M. polymorpha* CrRLK1L shared
138 the highest amino acid identity with AtFER (Figure S1A), we named it MpFER [45,46,57],
139 but it is also known as MpTHE [56].

140 Interestingly, no significant similarity to the MpFER malectin-like domain was found when
141 compared to available genomic data of six Chlorophyte algae (*Chlamydomonas*
142 *reinhardtii*, *Dunaliella salina*, *Volvox carteri*, *Coccomyxa subellipsoidea*, *Micromonas*
143 *pusilla*, *Ostreococcus lucimarinus*) and expression data of six Charophycean algae
144 (*Klebsormidium nitens*, *Nitella mirabilis*, *Coleochaete orbicularis*, *Spirogyra pratensis*,
145 *Mesostigma viride*, *Closterium peracerosum–strigosum–littorale* complex). In *Closterium*
146 complex, an RK named CpRLK1 expressed during sexual reproduction was suggested
147 to be a CrRLK1L homolog [59]. However, comparing the full sequence or the ECD of
148 CpRLK1 with the predicted *M. polymorpha* proteome, higher identity to another RK was

149 found (Figures S1B and S1C). Moreover, no RALF orthologs have been reported in algae,
150 suggesting the origin of the *CrRLK1L* pathway in land plants [45].

151 To further support MpFER as a member of the *CrRLK1L* subfamily, the three-dimensional
152 structure of its ECD was modelled [60] using the structure of the AtFER ECD (PDB
153 6a5b.1A, [35]) as a template (Figure 1C). Overall, the predicted structure of the MpFER
154 ECD is highly similar to that of AtFER, keeping an almost identical number of secondary
155 structures (7 alpha-helices and 33 beta-sheets in the MpFER ECD [35] and 8 alpha-helix
156 and 34 beta-sheets in the AtFER ECD, respectively), and a similar spatial disposition for
157 32 of the beta-sheet structures and 6 of the alpha-helices (Figure 1C). When comparing
158 the structural superposition of the AtFER and MpFER ECDs (Figures 1D and S1D), we
159 observed that most conserved parts of the malectin-like domain are in the core of the
160 protein, while the more variable parts are located peripherally. Moreover, the 3D structure
161 of the CpRLK1 ECD could not be properly modelled using the AtFER ECD (Figures S1D
162 and S1E). Taken together, these results suggest that the malectin-like domain of MpFER
163 could interact with similar proteins as AtFER does in *A. thaliana*, and may thus be involved
164 in similar pathways. Sequence comparison of the MpFER kinase domain showed a
165 conservation of the activation loop and a Lys important for catalytic activity (Figure 1E).
166 Furthermore, several determined phosphorylation sites found in AtFER, AtERU, and
167 *CrRLK1* were conserved in MpFER (Figure 1E) [13,33,61,62]. Thus, we propose MpFER
168 to be basal and orthologous to all other land plant *CrRLK1L*s, and that the *CrRLK1L* family
169 arose as plants conquered land.

170

171 **MpFER is broadly expressed throughout the *M. polymorpha* life cycle**

172 In *A. thaliana*, the expression patterns of the 17 *CrRLK1L* members span vegetative and
173 reproductive stages and both generations of the life cycle (reviewed in [43]), pointing to

174 the importance of *CrRLK1Ls* for basic cellular functions. To assess whether this pattern
175 is reflected in the land plant with the most ancestral characteristics, a promoter fragment
176 of 3.2 kb (*proMpFER*) was used to drive expression of a triple yellow fluorescent protein
177 (*trpVNS*) with a nuclear localization signal (*proMpFER:trpVNS*) and transformed into *M.*
178 *polymorpha* sporelings. All transformants expressing the *trpVNS* fluorescent protein (45
179 of 48 independent lines) exhibited an indistinguishable expression pattern (Figure 2).
180 During vegetative stages of gametophyte development, expression was observed in most
181 cells of the thallus (Figure 2A). In concordance with the importance of *MpFER* for rhizoids
182 [56], high *trpVNS* expression was observed in these tip-growing cells (Figure 2B). While
183 *trpVNS* was expressed in gemma cups, no expression was observed in mature gemmae,
184 possibly due to their dormancy (Figure 2C).
185 In mature female sexual organs, strong expression was observed in the entire
186 archegoniophore (Figure 2D). The archegonia expressed *trpVNS* in most cells, except for
187 the egg cell (Figure 2E). However, after fertilization, *proMpFER:trpVNS* became active in
188 the zygote (Figure 2F). Inside the sporogonium, the sporogenous cells and subsequently
189 the developing spores, but not the elater cells, strongly expressed *trpVNS* (Figure 2I). In
190 male sexual organs, *trpVNS* was broadly detected in the antheridial splash platform
191 (Figure 2G) and, specifically, in the spermatogenous tissue of the antheridia and the non-
192 reproductive jacket cells surrounding them at different developmental stages (Figure 2H).
193 Taken together, *MpFER* is expressed in most tissues during vegetative stages of
194 gametophyte development as well as in the antheridia and archegonia during the sexual
195 reproduction. Interestingly, the *proMpFER* promoter is not active in mature gemmae and
196 the unfertilized, quiescent egg cell, although expression is initiated after dormancy and in
197 the zygote. In concordance with the collectively ubiquitous expression of *CrRLK1Ls* in *A.*

198 *thaliana*, the MpFER expression pattern reinforces the importance of this family for basic
199 cellular functions in land plants.

200

201 **MpFER controls cell expansion during vegetative gametophyte development**

202 To analyze the function of MpFER, three independent artificial microRNA constructs
203 targeting MpFER were designed (amiR-MpFER), based on the endogenous microRNA
204 MpmiR160 (Figure S2, [63]) and driven by the ubiquitous *pro*MpEF1 promoter [64]. While
205 amiR-MpFER1 and amiR-MpFER2 target sites in the ECD coding sequence, amiR-
206 MpFER3 targets the sequence encoding the kinase domain (Figures 3A and 3B). For
207 each amiR-MpFER construct, several independent transgenic lines were obtained that
208 showed a reduction in thallus size (Figures 3C, 3D and S3A). MpFER levels were
209 quantified by qRT-PCR in two independent amiR-MpFER2 and amiR-MpFER3 lines
210 each, which showed a similar phenotype (Figure 3E). MpFER expression ranged from
211 10% to 20% of the wild-type level in all amiR-MpFER lines analyzed (Figure 3E).

212 Previously, the T-DNA insertion mutant *Mpfer-1* was identified in a screen for mutants
213 with defective rhizoids ([56], referred to as *Mpthe*). As the T-DNA inserted into the 3' UTR
214 of the MpFER gene (Figure 3A; [56]), MpFER expression was analyzed by qRT-PCR but
215 it showed no reduction compared to the wild type (Figure 3F), suggesting that the T-DNA
216 insertion does not affect MpFER transcription. However, thallus size of *Mpfer-1* mutants
217 is similar to that of amiR-MpFER lines (Figures 3C and 3D), suggesting that MpFER
218 function is partially affected by the T-DNA insertion.

219 Rhizoids are tip-growing cells analogous to the root hairs in flowering plants
220 (angiosperms). Reduced MpFER activity strongly impaired rhizoid formation and the
221 rhizoids collapsed (Figure 3C). This had also been reported for the *Mpfer-1* mutant [56]
222 and is in concordance with the function of AtFER and AtANX1/2 in tip-growing root hairs

223 [12,16,33,65] and pollen tubes [8,14], respectively. As *CrRLK1L* members are important
224 for cell expansion in angiosperms [7,9,11,19,34,66,67], cell size of epidermal thallus cells
225 in fully differentiated regions with minimal growth was measured in *Mpfer-1* and amiR-
226 *MpFER* lines, showing a reduction in cell area as compared to the wild type (Figures 3G,
227 S3B and S3C). Although amiR-*MpFER* lines produced more intact rhizoids than *Mpfer-1*
228 mutants (Figure 3C), the average epidermal cell area was similarly reduced, although the
229 distribution of cell areas differed between amiR-*MpFER* lines and *Mpfer-1* (Figures 3G,
230 S3B and S3C).

231 In conclusion, *MpFER* has a fundamental role in rhizoid formation and growth during
232 vegetative development. These results suggest an ancestral and conserved function of
233 the *CrRLK1Ls* in polar cell growth and cell expansion.

234

235 ***MpFER* is important for the morphological integrity of the gametophyte**

236 Given that *Mpfer-1* and amiR-*MpFER* lines retain some *MpFER* activity, we generated
237 *Mpfer* knock-out mutants with the goal to unveil potentially hidden *CrRLK1L* functions by
238 generating a plant without any *CrRLK1L* activity. Using CRISPR/Cas9, two sites in the
239 ECD-coding sequence of *MpFER* were targeted (Figure 4A). At least 10 independent
240 lines were selected and analyzed for each target site. All plants with severely affected
241 thallus development contained a frame shift mutation at the respective target site (Figures
242 4A-4C) while *MpFER* was not affected in transgenic plants with normal development.

243 The thallus area of the newly generated, amorphic knock-out mutants was more strongly
244 reduced compared to *Mpfer-1* [56], confirming that, like the amiR-*MpFER* lines, *Mpfer-1*
245 is a hypomorphic mutant (Figures 4C and 4D). In contrast, rhizoid integrity was similarly
246 affected in hypo- and amorphic *Mpfer* alleles, suggesting that rhizoid formation is more
247 sensitive to reduced *MpFER* activity than thallus growth (Figures 4E-4F). Despite of the

248 strong impact on thallus development, both hypo- and amorphic *Mpfer* mutants form
249 gemma cups.

250 The strong disruption of thallus growth in amorphic *Mpfer* mutants prompted us to
251 investigate whether cells of the thallus died in the absence of *MpFER* activity. Indeed,
252 using trypan blue staining, we detected dead cells in these mutants, indicating a loss of
253 morphological integrity (Figure 4G).

254

255 **Some reproductive but not all CWI sensing functions of *CrRLK1Ls* are conserved**
256 **in land plants**

257 In *A. thaliana*, at least seven *CrRLK1L* family members play a role in reproduction, with
258 *AtANX1/2* and *AtBUPS1/2* being important for pollen tube growth [8,14,18,32] and
259 *AtFER*, *AtHERK1*, and *AtANJ* for pollen tube reception by the synergid cells [6,17]. To
260 characterize the function of *MpFER* in reproductive development, amiR-*MpFER3-2* lines
261 were transferred to sexual organ-inducing conditions [68]. The number of
262 antheridiophores produced was significantly less in amiR-*MpFER* lines compared to the
263 wild type (Figures 5A and S3D) and the antheridiophore splash platforms were also
264 smaller (Figure 5B). The reduction in *MpFER* expression in antheridiophores of amiR-
265 *MpFER* lines was confirmed by qRT-PCR and results were comparable to those using
266 thallus tissue (Figure 5C). A pronounced reduction in spermatogenous tissue was
267 observed in many antheridia isolated from amiR-*MpFER3-2* lines (Figures 5D and 5E).
268 However, although the plants were grown under optimal conditions in axenic culture
269 boxes, the effects on reproductive structures could be partly indirect due to the reduction
270 in plant size and/or rhizoid function.

271 To determine male fertility of these amiR-*MpFER* lines with reduced spermatogenous
272 tissue, we performed crosses with wild-type female plants. The spermatocyte

273 concentration harvested from wild-type antheridia was almost 3-fold higher than from
274 amiR-MpFER lines and was adjusted using a hemocytometer. In just one of the crosses
275 (n=20), a single sporophyte was formed, resulting in 0.05 ± 0.05 (mean \pm SE) sporophytes
276 per archegoniophore, whereas the same females crossed with wild-type males yielded
277 9.95 ± 2.33 (mean \pm SE) sporophytes per archegoniophore. In summary, MpFER plays
278 a role in antheridiophore development and spermatocytes of plants with reduced MpFER
279 activity are poorly fertile. Thus, CrRLK1Ls have a conserved role in reproduction in
280 addition to their role in cell expansion and integrity during vegetative development.

281 Some *A. thaliana* CrRLK1Ls are involved in surveying CWI and in inhibiting growth when
282 CWI is impaired. Cell wall defects caused either by mutations affecting cellulose
283 biosynthesis or by treatment with isoxaben, a cellulose synthesis inhibitor, can be
284 suppressed by a mutation in *AtTHE1* [7,28]. To investigate whether MpFER also acts as
285 CWI sensor, we treated wild-type, *Mpfer*, and amiR-MpFER3-2 gemmae with isoxaben
286 (Figure 5F). As in the wild type, isoxaben affected the development of gemmae with
287 reduced MpFER activity, all growing progressively less with increasing isoxaben
288 concentration (Figure 5F). These results suggest that the function of *AtTHE1* in
289 repressing cellular growth when CWI is impaired appeared later in the course of land
290 plant evolution or was lost in the *Marchantia* lineage.

291

292 **The CrRLK1L signaling pathway is conserved in land plants**

293 As downregulation of MpFER or AtFER produce similar phenotypes concerning polarized
294 growth and cell expansion in *M. polymorpha* and *A. thaliana*, respectively, interspecific
295 complementation was attempted. First, the coding sequence of MpFER fused to the
296 Green Fluorescent Protein (GFP) gene under the control of the viral 35S promoter
297 (*pro35S:MpFER-GFP*) was transformed into heterozygous *Atfer-2* mutants. Then, GFP

298 expression at the cell periphery and complementation of the bursting root hair and
299 reduced rosette size phenotypes were assessed in transgenic plants (Figure 6A). *Atfer-2*
300 homozygotes expressing the MpFER-GFP protein look similar to *Atfer-2* mutants,
301 indicating that MpFER does not complement these *Atfer-2* loss-of-function phenotypes
302 (Figure 6A). In a complementary experiment, the coding sequence of AtFER fused to the
303 Citrine gene under the control of the MpEF1 promoter (*proMpEF1:AtFER-Cit*) was
304 transformed into amiR-MpFER3-2 plants. Ten independent lines with Citrine expression
305 at the cell periphery were phenotypically characterized (Figures 6B and 6C). All had
306 bursting rhizoids and a reduced thallus size, similar to the parental amiR-MpFER3-2 line,
307 suggesting that AtFER does not rescue the vegetative phenotypes produced by down-
308 regulation of MpFER.

309 In *A. thaliana*, it was shown that AtFER forms a complex with AtRALF peptides and the
310 LRE/LLG co-receptors [35]. As no interspecific complementation was observed, the
311 conservation of binding surfaces between CrRLK1L, RALF, and LRE homologs were
312 analyzed. The *M. polymorpha* genome encodes three RALF peptides (MpRALF:
313 Mp1g27120 [Mapoly0002s0166]; MpRALF2: Mp2g21670 [Mapoly0040s0047];
314 MpRALF3: Mp7g07270 [Mapoly0076s0067]) [44,45]. We performed a phylogenetic
315 analysis using the amino acid sequences of the predicted mature peptides of *A. thaliana*
316 and *M. polymorpha* RALFs (Figure S4A). All MpRALF peptides clustered together with
317 AtRALFs known to interact with CrRLK1L receptors, including AtRALF1 and AtRALF23,
318 which serve as AtFER ligands [22,33] (Figure S4A). The MpRALFs shared the four
319 conserved Cys residues as well as the YXXY and YY motifs with the AtRALF1 subgroup
320 (Figure S4B). However, only MpRALF1 and MpRALF3 had an RRXL motif important for
321 S1P cleavage (Figures S4B and S4C)[69], consistent with the presence of one S1P
322 ortholog in basal plants (Mp8g07990 [Mapoly0155s0018], Figure S4D). There are two

323 MpLRE/LLG proteins encoded in the *M. polymorpha*, genome (Figure S5A). Both the
324 MpLRE1 (Mp5g09600 [Mapoly0048s0110]) and MpLRE2 (Mp4g22100
325 [Mapoly0090s0020]) proteins showed conservation of the eight Cys and the ND motif
326 distinctive of the family (Figure S5B); however, MpLRE1 did not contain a GPI anchoring
327 site (Figure S5B). Structure prediction using AtLLG2 as a template [35] showed a
328 conserved general structure (Figures S5C and S5D). Some of the MpRALF and MpLRE
329 family members show a similar expression pattern as MpFER (Figure S5F), suggesting
330 that the corresponding proteins could form a complex similar to that described in *A.*
331 *thaliana*.

332 To gain insights into the formation of a putative CrRLK1L signaling complex, the
333 MpFER/MpRALF/MpLRE complex was modelled. The general structure appeared
334 conserved (Figure S5E); however, analysis of conserved sites between *M. polymorpha*
335 and *A. thaliana* showed a low amino acid conservation in the interacting surfaces of the
336 complex subunits (Figure 6D). This suggests that the lack of interspecific
337 complementation may be due to differences in amino acids that are important for complex
338 formation, and that the proteins forming the complex have diverged, following distinct
339 routes of co-evolution in the two lineages.

340 Because interspecific complementation of the amiR-MpFER phenotypes was
341 unsuccessful, we asked whether suppression using a downstream component of
342 CrRLK1L signaling identified in *A. thaliana* was possible. AtMRI, a receptor-like
343 cytoplasmic kinase, acts downstream of AtFER or AtANX1/2 in the regulation of polar tip
344 growth [70,71]. The dominant AtMRI^{R240C} mutation can suppress pollen tube rupture in
345 the *Atanx1/Atanx2* double mutant and the root hair defects of *Atfer-4* [70,71]. A mutation
346 in MpMRI (aka MpPTI) with a similar bursting rhizoid phenotype as observed in *Mpfer-1*,
347 was identified in the same genetic screen for *M. polymorpha* mutants with defective

348 rhizoids [56]. To test whether MpMRI is a conserved downstream component of the
349 MpFER signal transduction pathway, a dominant-active form of the protein equivalent to
350 AtMRI^{R240C} was transformed into the amiR-MpFER3-2 line, driven by 2 kb of the
351 endogenous promoter (*proMpMRI:MpMRI^{R240C}*). Several independent lines showed a
352 partial restoration of rhizoid growth; however, we also observed defects in thallus
353 development of lines with higher levels of MpMRI expression, mainly abnormalities in the
354 epidermis (Figures 6E and S6A-S6E). This suggest that MpMRI acts downstream of
355 MpFER in the signal transduction pathway controlling polarized growth since the origin of
356 land plants, in agreement with a recent report on the functional characterization of MpMRI
357 [57].

358

359 **Overexpression of MpFER affects morphological integrity**

360 Because overexpressing AtFER in *M. polymorpha* did neither lead to any obvious
361 phenotypes nor complementation of amiR-MpFER lines (Figures 6B and 6C), we also
362 overexpressed AtFER and MpFER in the wild type. As in the amiR-MpFER background,
363 overexpression of AtFER (*proMpEF1:AtFER-Cit*) in a wild-type background had no effect,
364 consistent with the idea that AtFER is unable to form a complex with the corresponding
365 *M. polymorpha* proteins. However, when expressing an MpFER-Citrine fusion protein
366 (*proMpEF1:MpFER-Cit*) in wild-type plants, thallus development was strongly affected
367 (Figure 7A). As expected, MpFER-Cit localized to the membrane (Figure 7B), and higher
368 protein levels correlated with more severe phenotype (Figures 7A and 7C). Scanning
369 electron microscopy of epidermis of plant expressing high levels of MpFER showed
370 defects in the formation of air chambers and air pores (Figure 7D), similar to what we
371 observed when expressing the dominant-active MpMRI^{R240C} (Figure S6A-S6C).
372 Moreover, overexpression of MpFER affects the number of lobes in the antheridial

373 receptacle, producing only 4 instead of 8 (Figure S6F), supporting the importance of
374 MpFER during sexual development. Considering the importance of MpFER for normal
375 rhizoid formation, we also analyzed rhizoids in the *proMpEF1:MpFER-Cit* lines: while
376 rhizoid morphology looked normal, a decrease in rhizoid number was observed (Figures
377 7E and 7F).

378 Taken together, the results obtained by overexpressing MpFER support the role of
379 MpFER in cell expansion and maintaining tissue integrity during plant morphogenesis.

380

381 **Discussion**

382 During land plant evolution, many developmental aspects have changed in order to adapt
383 to new environments, producing the enormous diversity observed in the plant kingdom.
384 However, the control and maintenance of CWI remained a key aspect for the biology of
385 a plant cell. CWI sensing is not only important for vegetative growth by cell expansion,
386 but it is also essential for sexual reproduction and defense responses.

387 The *CrRLK1L* family was first characterized for its importance during angiosperm
388 fertilization, a process comprising aspects of polar cell elongation, CWI control, and cell-
389 cell communication [6,30,31]. Lately, different members of this family were found to carry
390 out diametrically opposite roles in diverse aspects of plant development, which has made
391 it difficult to define the core or ancestral function of this gene family comprising 17-
392 members in *A. thaliana* [6–12,14–20,32].

393 Here, we report the characterization of the single *CrRLK1L* gene encoded in the genome
394 of *M. polymorpha*. Structurally, the MpFER ECD and kinase domain share similarities
395 with previous characterized *CrRLK1L* members, including the malectin-like domain and
396 conserved phosphorylation sites. Conservation of RALF and LRE members in *M.*
397 *polymorpha* suggests that MpFER also forms a complex at the plasma membrane [35].

398 While modeling predicted that the general structure of the complex was conserved, the
399 interaction surfaces seem to have diverged and co-evolved independently in the
400 respective lineages.

401 We observed a broad involvement of Mp*FER* at both vegetative and reproductive stages
402 of development: During vegetative growth of the gametophyte, Mp*FER* is required for
403 rhizoid formation and cell expansion but it also plays a role in male gametogenesis and
404 is expressed in female reproductive organs. These findings suggest that the *CrRLK1Ls*
405 have, also in ancestral land plants, held roles in various aspects of both vegetative and
406 reproductive development. Thus, the role of *CrRLK1Ls* in angiosperm reproduction does
407 not represent a derived feature from a purely vegetative function in cell elongation in
408 bryophytes, but constitutes a conserved feature of this gene family.

409 Moreover, the almost ubiquitous expression of Mp*FER* in *M. polymorpha* is consistent
410 with the collective expression of different *CrRLK1L* members in essentially all tissues and
411 organs of *A. thaliana* [43]. Thus, gene duplication allowed the expansion of the *CrRLK1L*
412 family in angiosperms, diversifying their expression patterns and functions but also
413 leading to genetic redundancy [8,17,32,43].

414 *CrRLK1L* members are important sensors of CWI during polarized growth, both in pollen
415 tubes [8,32] and root hairs [16]. In early divergent land plants, rhizoids are tip-growing cell
416 with a function analogous to that of root hairs, important for taking up nutrients and water
417 [56,72]. Based on the conservation of many genes controlling the development and
418 growth of tip-growing cells forming rooting systems, it was suggested that these pathways
419 were active in the earliest land plants that existed about 470 million years ago [56]. That
420 Mp*FER* also controls rhizoid integrity points to the importance of the *CrRLK1L* pathway
421 for polarized growth since the origin of land plants. Moreover, the fact that amorphic Mp*fer*
422 mutants contain many dead cells demonstrates the importance of the *CrRLK1L* pathway

423 for cellular integrity during vegetative growth. This aspect has not yet been reported for
424 *A. thaliana* CrRLK1L mutants, possibly due to genetic redundancy among family
425 members.

426 The signaling mechanisms downstream of MpFER in *M. polymorpha* development are
427 still unclear. MpMRI, the *M. polymorpha* ortholog of AtMRI, which acts downstream of
428 AtFER and AtANX1/2 in *A. thaliana* during polarized growth [70], does have a similar
429 rhizoid phenotype as MpFER [56,57]. The suppression of the rhizoid phenotype in amiR-
430 MpFER transformants with the constitutively active form MpMRI^{R240C} corroborates a
431 conserved CrRLK1L signaling cascade during polarized growth in *M. polymorpha* but
432 whether this pathway also relies on Ca²⁺ and reactive oxygen species (ROS) as second
433 messengers as it does in *A. thaliana* remains to be determined.

434 In *A. thaliana*, loss-of-function mutations in the motor protein AtKINESIN-13a, a
435 microtubule-based motor, cause cell elongation in petals, leaves, and hypocotyls
436 concomitant with changes in cell wall composition in leaves [73]. This effect depends on
437 functional AtTHE1 and does not occur in *Atkinesin-13-a/Atthe1* double mutants.
438 Interestingly, in this case, AtTHE1 stimulates cell elongation in response to defects in cell
439 wall deposition [73]. Based on our results, we have no indication that MpFER acts as a
440 CWI sensor and regulates growth similar to AtTHE1 in response to cell wall disturbances.
441 We have not observed a context in *M. polymorpha* development in which MpFER inhibits
442 cell expansion. Therefore, the ability to restrict growth in response to disturbances in CWI
443 could represent an evolutionary derived feature, as gene duplication allowed for the
444 diversification and specialization of CrRLK1Ls to regulate cellular growth in a more
445 complex and context-dependent way. The capacity to flexibly regulate growth appears to
446 be fundamentally context-dependent, as a similar discrepancy in promoting or inhibiting
447 growth is also known for some well-studied growth promoting agents, like the

448 phytohormones auxin and brassinosteroid, which can have growth-inhibitory effects
449 dependent on tissue and concentration. The reverse was noted for the phytohormones
450 ethylene and abscisic acid, which are usually considered growth inhibitors but can also
451 promote growth, dependent on the context [74–76].

452 Taken together, our results suggest an ancestral and conserved function of the *CrRLK1Ls*
453 in polar cell growth and cell expansion, but also during sexual reproduction. In addition,
454 we probe the relevance of *CrRLK1L* members for cell integrity and morphogenesis. In
455 angiosperms, *CrRLK1Ls* occupy an important position at the interface of developmental
456 and environmental inputs, which are integrated by a downstream signaling machinery
457 that controls cell shape and polar growth to ensure normal development, *e.g.* by
458 preventing cell rupture upon CWI defects. Whether the same downstream machinery is
459 utilized in a similar fashion in *M. polymorpha* remains to be shown, even though the
460 similarity in function points to a related mechanism. Even less is known about upstream
461 aspects of *CrRLK1L* signaling. Future studies on the transcriptional regulation and the
462 upstream components, such as the putative RALF ligand(s), in the highly simplified *M.*
463 *polymorpha* system could provide fundamental insights into the molecular mechanism of
464 the *CrRLK1L* pathway, which is so central to land plant physiology, growth, and
465 development.

466

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470 cell area, Célia Baroux and Ethel Mendocillo-Sato for instructions on confocal microscopy
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474

475 **Author contributions**

476 MAM: Conceptualization, Formal analysis, Investigation, Methodology, Validation,
477 Visualization, Writing – original draft, Writing – review & editing

478 MR: Conceptualization, Formal analysis, Investigation, Methodology, Validation,
479 Visualization, Funding acquisition, Writing – original draft, Writing – review & editing

480 AF-G: Data curation, Formal analysis, Software, Validation, Visualization, Writing –
481 original draft, Writing – review & editing

482 DM: Formal analysis, Investigation, Validation, Writing – review & editing

483 PG: Formal analysis, Investigation, Validation, Writing – review & editing

484 JLB: Funding acquisition, Resources, Supervision, Validation, Writing – review & editing

485 UG: Conceptualization, Formal analysis, Funding acquisition, Project administration,
486 Supervision, Validation, Writing – original draft, Writing – review & editing

487

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496

497 **Competing interests**

498 The authors have declared that no competing interests exist.

499

500 **Main figure titles and legends**

501 **Figure 1. CrRLK1Ls Are Conserved among Land Plants**

502 (A) Representation of AtFER and MpFER proteins. Malectin-like domains are
503 represented in blue and kinase domains in green. SP, signal peptide; TM, transmembrane
504 domain (yellow). Amino acids important for regulation and activity are marked in red,
505 violet, and light blue as described in Figure 1E.

506 (B) A rooted neighbor-joining tree of the amino acid sequence of the predicted malectin-
507 like domain. CrRLK1L members from *Marchantia polymorpha* (Mp), *Physcomitrella*
508 *patens* (Pp), *Selaginella moellendorffii* (Smo), and *Arabidopsis thaliana* (At) were used.
509 The numbers indicate the bootstrap values (%) from 1000 replications. The given scale
510 represents a substitution frequency of 0.1 amino acids per site.

511 (C) Cartoon representation of the predicted three-dimensional structure of the MpFER
512 malectin-like domain, showing predicted alpha-helix and beta-sheet structures.

513 (D) Structural superposition of the malectin-like domains of AtFER (blue) and MpFER
514 (green).

515 (E) Alignment of the cytoplasmic domains of MpFER, AtERU, AtFER, and CrRLK. Kinase
516 domains are in green, putative phosphorylation sites in light blue (Ser) and violet (Thr),
517 and the conserved catalytic Lys in red.

518 See also Figure S1

519

520 **Figure 2. MpFER Is Broadly Expressed in most Tissues throughout the *M.***
521 ***polymorpha* Life Cycle**

522 Expression of *proMpFER:trpVNS* in different organs of male and female *M. polymorpha*
523 plants. Fluorescence is visualized as either green (epifluorescence microscope) or yellow
524 (confocal microscope) color, depending on the panel.

525 (A to D) Bright field (left) and epifluorescence (right) images of a thallus, (A) rhizoid (C),
526 gemmae cups (C), and an archegoniophore (D). Red arrows indicate meristematic zones,
527 the yellow arrow gemmae. Scale bars, 0.5 mm.

528 (E and F) Confocal images of archegonia before (E) and 2 days after fertilization (F).
529 Scale bar, 50 μ m.

530 (G) Bright field (left) and epifluorescence (right) images of an antheridiophore. Scale bar,
531 1 mm.

532 (H) Confocal image of antheridia. Scale bar, 100 μ m.

533 (I) Confocal image of sporophyte and spores. Scale bar, 25 μ m.

534

535 **Figure 3. MpFER Controls Cell Size during Gametophyte Development**

536 (A) Representation of the mature MpFER mRNA. Coding sequence are in green,
537 locations of the amiRNA target sites in blue.

538 (B) Base complementary between mature amiR-MpFER1, amiR-MpFER2, and amiR-
539 MpFER3 with the MpFER transcript.

540 (C) Relative expression level of MpFER in thallus tissue from wild-type (WT; Tak1) and
541 two independent insertion lines of amiR-MpFER2 and amiR-MpFER3, as measured by
542 qRT-PCR. MpEF1 was used as internal control. Shown are means \pm standard errors of
543 the means (SEM) of three biological replicates. *P < 0.01, one-way analysis of variance
544 (ANOVA).

545 (D) Relative expression level of Mp*FER* in WT and Mp*fer-1* thallus tissue, as measured
546 by qRT-PCR. Mp*EF1* was used as internal control. Shown are means \pm SEM of three
547 biological replicates. *P < 0.01, one-way ANOVA.

548 (E) Representative pictures of 10-day old gemmalings of WT, Mp*fer-1*, and amiR-
549 Mp*FER3-2* lines. Scale bar, 1 mm.

550 (F) Thallus area of WT, Mp*fer-1*, and amiR-Mp*FER3-2* lines. n = 30, outliers are indicated
551 as black dots.

552 (G) Violin plot of cell size in WT, Mp*fer-1*, and amiR-Mp*FER3-2* lines. Significant
553 differences are indicated according to the non-parametric Kruskal-Wallis test and linear
554 regression models (**p<0.001).

555 See also Figures S2 and S3

556

557 **Figure 4. The Integrity of Thalli from Mp*fer* Knock-out Lines Is Severely Affected**

558 (A) Representation of the mature Mp*FER* mRNA with the coding sequence in green, and
559 the location of the gRNA target sites in red.

560 (B) Sequences of gRNA target sites in the confirmed Mp*fer* knock-out mutants. Deletions
561 and insertions are highlighted in red.

562 (C) Representative pictures of 10-day old gemmalings of WT, Mp*fer-1*, and two
563 independent Mp*fer* knock-out mutants. Scale bar, 0.5 cm.

564 (D) Thallus area of 10-day old gemmalings of WT, Mp*fer-1*, and two independent Mp*fer*
565 knock-out mutants. n = 30.

566 (E) Representative pictures of 3-day old gemmalings growing in upside-down plates of
567 WT, Mp*fer-1*, and two independent Mp*fer* knock-out mutants. Scale bar, 500 μ m.

568 (F) Rhizoid number in 3-day old gemmalings of WT, Mp*fer-1*, and two independent Mp*fer*
569 knock-out mutants. n = 30.

570 (G) Trypan blue staining of 7-days old gemmalings of WT, *Mpfer-1*, and two independent
571 *Mpfer* knock-out mutants. Scale bar, 1 mm.

572

573 **Figure 5. Conservation of CrRLK1L Function in Land Plants**

574 (A) Number of antheridiophores produced per plant for wild-type (WT, n=9) and two
575 independent amiR-MpFER3 lines (amiR-MpFER3-2, n = 9; amiR-MpFER3-3, n = 12).
576 Shown are means \pm SEM of two biological replicates. Means with same letter do not differ
577 significant different with $P > 0.05$, one-way ANOVA, Duncan test.

578 (B) Antheridiophore splash platform size distribution of WT (n = 32), amiR-MpFER2-3 (n
579 = 34), and amiR-MpFER3-3 (n = 11) lines. For all antheridia with a stalk >8 mm, length
580 and width of the platform were recorded.

581 (C) qRT-PCR of MpFER levels in antheridiophores in a WT and amiR-MpFER3-3 line.
582 MpEF1 was used as internal control. Shown are means \pm SEM of three biological
583 replicates. * $P < 0.01$, ANOVA.

584 (D and E) Mature antheridia of a WT (D) and amiR-MpFER3-3 line (E). Bright field images
585 (left) and in cross-section (right). The spermatogenous areas are indicated in red. Scale
586 bar, $100\mu\text{m}$.

587 (F) Thallus area of 15-day old gemmalings of WT, *Mpfer-1* mutant, and an amiR-
588 MpFER3-2 line growing on media containing different isoxaben concentrations (nM). n =
589 30, outliers are indicated as black dots. Shown are means \pm SEM of two biological
590 replicates. Means designated by the same letter do not significantly differ at $P > 0.05$,
591 one-way ANOVA, Duncan test.

592 See also Figure S3

593

594 **Figure 6. Interspecific Complementation of *A. thaliana* and *M. polymorpha* Plants**
595 **with Reduced Levels of *FER* Activity**

596 (A) Roots of independent *A. thaliana* lines expressing 35S:Mp*FER-GFP* in *Ler* wild-type
597 (WT) and *Atfer-2* plants. Scale bar, 200 μ m.

598 (B) 10-day old gemmalings of WT and amiR-Mp*FER3-2* *M. polymorpha* plants with and
599 without expression of *AtFER-Cit*. Scale bar, 0.5 mm.

600 (C) Citrine expression in 7-day old gemmalings of WT and amiR-Mp*FER3-2* plants with
601 and without expression of *AtFER-Cit*. Scale bar, 100 μ m.

602 (D) Cartoon representation of the structural conservation of the *CrRLK1L*^{ECD}/LRE/RALF
603 complex by comparing *AtFER*/*MpFER* and *AtLLG2*/*MpLRE1*. Blue and green represent
604 spatially highly conserved regions of *CrRLK1L* and LRE homologs, respectively. Red
605 indicates structural regions with low spatial conservation, lilac the C-terminal part of
606 *AtRALF23*, and black arrows the interaction regions of the three proteins.

607 (E) Complementation of amiR-Mp*FER* lines with a dominant-active version of *MpMRI*
608 under its own promoter (*proMpMRI:MRI*^{R240C}). Scale bar, 1 mm.

609 See also Figures S4-S7

610

611 **Figure 7: Overexpression of *MpFER* Affects Morphological Integrity**

612 (A) Representative pictures of 10-day old gemmalings of wild-type (WT, Tak1) and
613 different lines overexpressing *MpFER* (*proMpEF1:MpFER-Cit*). Scale bar, 1 mm.

614 (B) Citrine expression at the plasma membrane in WT and *proMpEF1:MpFER-Cit* lines #5
615 and #9 as observed under confocal microscopy. Scale bar, 20 μ m.

616 (C) Western blot analysis of *proMpEF1:MpFER-Cit* lines using an anti-GFP antibody.
617 *proMpEF1:Cit* and WT lines were used as positive and negative controls, respectively. The

618 Ponceau membrane staining of the most intense band at 55 kDa (presumably Rubisco)
619 was used as a loading control.

620 (D) Scanning electron microscopical images of thalli from 20-day old plants of WT and
621 *proMpEF1:MpFER-Cit* lines #5 and #9. Scale bar, 500 μ m.

622 (E) Representative pictures of 3-day old gemmalings growing in upside-down plates of
623 WT and *proMpEF1:MpFER-Cit* lines #5 and #9. Scale bar, 1 mm.

624 (F) Number of rhizoids in 3-day old gemmalings of WT and *proMpEF1:MpFER-Cit* lines #5
625 and #9. n = 30.

626 See also Figure S6

627

628 **Supplemental figure titles and legends**

629

630 **Figure S1. The CrRLK1L Family Appeared together with Land Plants. Related to**
631 **Figure 1.**

632 (A) Percentage of identity of the malectin-like domain, the complete ECD, and the full
633 protein of MpFER with AtFER, AtANX1, AtANX2, and AtTHE1.

634 (B) Percentage of identity of the ECD and full protein of CpRLK1 with MpFER, AtFER,
635 and Mapoly001s0111.

636 (C) A rooted neighbor-joining tree of the amino acid sequence of the predicted malectin-
637 like domain was generated using ClustalW. CrRLK1L members from *Marchantia*
638 *polymorpha* (Mp), *Physcomitrella patens* (Pp), *Selaginella moellendorffii* (Smoe), and
639 *Arabidopsis thaliana* (At) were used. Algal CpRLK1 and Mapoly001s0111 were also
640 included. The numbers indicate the bootstrap values (%) from 1000 replications. The
641 given scale represents a substitution frequency of 0.1 amino acids per site.

642 (D) Qmean value for prediction of MpFER or CpRLK1 3D structures of the ECD based
643 on the AtFER ECD. Green numbers indicate a good modelling fit, red numbers indicates
644 a bad modelling fit.

645 (E) Structural superposition of the ECD of AtFER (blue) and CpRLK1 (purple).

646

647 **Figure S2. Design of amiR-MpFER Precursors. Related to Figure 3.**

648 (A) *MpmiR160* (Mapoly0002s0211, Mp1g26670) and amiR-MpFER1, amiR-MpFER2,
649 and amiR-MpFER sequences. miRNA sequences are in red and miRNA* in blue. Cloning
650 sequences from amiR-MpFER constructs are in bold.

651 (B) Drawing of the minimum free energy structure of *MpmiR160* and amiR-MpFER3
652 constructs predicted by the RNAfold web server ([http://rna.tbi.univie.ac.at/cgi-](http://rna.tbi.univie.ac.at/cgi-bin/RNAWebSuite/RNAfold.cgi)
653 [bin/RNAWebSuite/RNAfold.cgi](http://rna.tbi.univie.ac.at/cgi-bin/RNAWebSuite/RNAfold.cgi)). Red arrows indicate location and orientation of the
654 mature miRNA in the precursor. The structures are colored by base-pairing probabilities;
655 for unpaired regions the color denotes the probability of being unpaired.

656

657 **Figure S3. Reduction in Thallus and Cell Area in Plants with Reduced MpFER**
658 **Levels. Related to Figures 3 and 5.**

659 (A) Thallus area of wild-type (WT), *Mpfer-1*, and amiR-MpFER3-2 plants at different days
660 after putting gemmae on plates. n = 30, outliers are indicated as black dots. Areas were
661 estimated using ImageJ software.

662 (B) Representative images from cell surface areas measured in WT, amiR-MpFER2-1,
663 and amiR-MpFER3-2 plants. Scale bar, 50 μ m.

664 (C) Difference in cell size between WT, amiR-MpFER2-1, and amiR-MpFER3-2 plants is
665 significant ($p < 0.001$) based on the nonparametric Kruskal-Wallis test and a linear
666 regression model with a highly significant interaction ($p < 0.001$).

667 (D) Induction of antheridiophores in WT and two independent amiR-Mp*FER3* lines. Three
668 plants were grown in each sterile plastic box under far-red light induction.

669

670 **Figure S4. MpRALF Peptides Belong to the AtRALF1-clade of the RALF Family.**
671 **Related to Figure 6.**

672 (A) A rooted neighbor-joining tree of the amino acid sequence of the predicted mature
673 RALF peptides was generated using ClustalW. RALF members from *M. polymorpha* and
674 *A. thaliana* were used. Red arrows indicate RALFs that are known ligands of CrRLK1Ls.
675 The numbers indicate the bootstrap values (%) from 1000 replications. The given scale
676 represents a substitution frequency of 0.1 amino acids per site.

677 (B) Amino acid sequence comparison of AtRALF1 and MpRALF1-3. Predicted signal
678 peptides are in black, predicted mature peptides in light blue, conserved Cys in red, and
679 predicted S1P recognition sites in violet.

680 (C) Processing pathway AtRALF1 by the S1P protease.

681 (D) A rooted neighbor-joining tree of the amino acid sequence of the S1P orthologs was
682 generated using ClustalW. S1P members from *M. polymorpha*, *P. patens*, and *A. thaliana*
683 were used. The numbers indicate the bootstrap values (%) from 1000 replications. The
684 given scale represents a substitution frequency of 0.1 amino acids per site.

685

686 **Figure S5. Two LORELEI-like Proteins Are Encoded in the *M. polymorpha* Genome.**
687 **Related to Figure 6.**

688 (A) A rooted neighbor-joining tree of the amino acid sequence of LRE orthologs was
689 generated using ClustalW. LRE members from *M. polymorpha*, *P. patens*, and *A. thaliana*
690 were used. The numbers indicate the bootstrap values (%) from 1000 replications. The
691 given scale represents a substitution frequency of 0.1 amino acids per site.

692 (B) Amino acid sequence of AtLRE, MpLRE1, and MpLRE2. Conserved Cys are in light
693 blue, the ND motif in red, and the GPI-anchoring site in green.

694 (C) Cartoon representation of the predicted 3-dimensional structure of MpLRE1, showing
695 predicted alpha-helices.

696 (D) Structural superposition of AtLGG2 (blue) and MpLRE1 (green).

697 (E) Cartoon representation of the predicted 3-dimensional structure of the
698 MpFER/MpLRE1/MpRALF1 complex, showing predicted alpha-helices and beta-sheets.

699 (F) Heatmap depicting relative gene expression based on RNAseq data (row-Z-score of
700 vs normalized counts) of MpFER and the *M. polymorpha* orthologs of AtMRI, AtRALF1,
701 and AtLRE across different tissues. Vegetative and reproductive tissues are grouped by
702 green and orange, respectively. Averaged expression values are represented with colors
703 of increasing red and blue intensity indicating upregulation and downregulation of gene
704 expression, respectively.

705

706 **Figure S6. Expression of MpMRI^{R240C} Suppresses the Bursted Rhizoid Phenotype**
707 **of amiR-MpFER3 Lines but Leads to Aberrant Epidermis Development. Related to**
708 **Figures 6 and 7.**

709 (A to C) Epidemical pictures of thalli from the wild-type (WT, Tak-1) (A), and amiR-
710 MpFER3 + _{pro}MpMRI:MpMRI^{R240C} lines #6 (B) and #8 (C), which both partially suppressed
711 the bursting rhizoid phenotype (Fig. 6).

712 (D and E) Relative expression of MpFER (D) and MpMRI (E) against the geometric mean
713 of the reference genes MpACT1, MpACT7, and MpAPT3 in WT, amiR-MpFER3-2, and 5
714 lines (#5 to #9) co-transformed with the amiR-MpFER3-2 and _{pro}MpMRI:MpMRI^{R240C}
715 constructs. Expression levels of three biological replicates were assessed by droplet
716 digital PCR (ddPCR). The y-axis corresponds to the log₂-ratio between the test and the

717 geometric mean of the reference genes. Shown are means \pm SEMs of three biological
718 replicates. *P < 0.01, one-way ANOVA.

719 (F) Representative images of the antheridial receptacle of WT and *pro*MpEF1:MpFER-Cit
720 plants. Scale bar, 2 mm.

721

722 **Figure S7. Structure Assessment and Quality Estimations of the MpFER and**
723 **MpLRE 3-dimensional Models. Related to Figure 6.**

724 QMEAN score barplot indicating quality estimates of the predicted models, across their
725 aminoacidic sequence. Below the barplots, the corresponding secondary structures are
726 displayed for both target and template sequences, with alpha-helices represented by light
727 purple boxes and beta-sheets by green arrows. All plots were adapted from the model
728 report produced by the SWISS-MODEL workspace.

729 (A) MpFER^{ECD}, for which AtFER was used as a template (PDB 6a5b.1A).

730 (B) MpLRE1, for which AtLLG2 was used as a template (PDB 6a5d.1A).

731

732 **RESOURCE AVAILABILITY**

733

734 **Lead Contact**

735 Further information and requests for resources and reagents should be directed to and
736 will be fulfilled by the Lead Contact, Ueli Grossniklaus (grossnik@botinst.uzh.ch).

737

738 **Materials Availability,**

739 Requests for other strains and plasmids should be directed to the Lead Contact.

740

741 **Data and Code Availability.**

742 This study did not generate datasets or code.

743

744 **EXPERIMENTAL MODEL AND SUBJECT DETAILS**

745 *M. polymorpha* subsp. *ruderalis* plants were grown on sterile culture on half-strength
746 Gamborg's B5 basal medium (PhytoTechnology Laboratories), supplemented with 1%
747 phytoagar in Petri dishes sealed with air-permeable tape. The plants were kept under
748 fluorescent light and long-day conditions (16 h light at 22°C, 8 h dark at 20 °C) in Percival
749 growth cabinets (models AR-41L3 and AR-41/L2). Alternatively, to induce sexual
750 reproduction, plants were cultivated in jiffy pots filled with a 1:1 mixture of soil
751 ("Einheitserde D73 + Bims", Universallerde) and sand or in culture boxes in sterile culture
752 on half-strength Gamborg's B5 basal medium (Duchefa Eco2 Box, #E1654), under
753 fluorescent light supplemented with far-red light (GreenPower LED module HF far-red,
754 #929000464503 and #929000632103 Philips).

755 Wild-type (WT) plants in all experiments were *Arabidopsis thaliana* L. (Heyn) accession
756 Ler-0. The *Atfer-2* mutant was obtained from the Signal Collection at the Salk Institute.
757 Plants were grown in soil (ED73; Universallerde), covered with a thin quartz sand layer,
758 under long photoperiods (16 hs light/8 hs dark) at 23°C and 60% relative humidity.

759

760 **METHOD DETAILS**

761 **Phylogenetic analysis**

762 Protein sequences were identified in plant genomes via BLASTp searches in
763 <https://phytozome.jgi.doe.gov/pz/portal.html>. To elucidate the evolutionary relationship
764 across land plant evolutionary history, we focused on *A. thaliana*, *Selaginella*

765 *moellendorffii*, *Physcomitrella patens* and *M. polymorpha*, as representative species for
766 angiosperms, lycophytes, mosses and liverworts, respectively. Complete or partial coding
767 protein sequences were aligned using the ClustalW parameters and were conducted in
768 MEGA7. Phylogenetic trees were constructed with Neighbor-joining method, with a
769 bootstrap test of 1000 replicates. The evolutionary distances were computed using the
770 Poisson correction method.

771

772 **Vector construction**

773 Expression constructs: a BJ36 plasmid [77] containing a tripleVENUS-NLS (trpVNS)
774 fragment was modified by adding a ligation-independent cloning (LiC) adapter site [78].
775 A promoter fragment of Mp*FER* of 3.2 kb was amplified using primers specified below
776 and cloned into the BJ36 vector via LiC cloning. *pro*Mp*FER* promoter fragment fused to
777 trpVNS was then shuttled via *NotI* restriction sites into the HART01 [77] expression
778 vector. The expression vector was also directly modified to contain the LiC sites, so
779 promoter fragments can now be directly cloned in front of the trpVNS in the HART01
780 vector via LiC-cloning. This vector was called VHL (trpVNS-containing HART01 vector
781 with LiC sites).

782 amiRNA constructs: For functional analyses three independent artificial microRNA
783 (amiRNA) constructs were made (Figure S2). The endogenous microRNA precursor
784 Mp*miR160* was used as both backbone and template to design the constructs which
785 generate single species small RNA molecules of 21 nt length, complementary to the
786 target gene transcript [63,79]. Three miR160 backbones containing independent amiR-
787 Mp*FER*/amiR-Mp*FER** duplexes were synthesized by GenScript and fused to the
788 *pro*Mp*EF1* promoter in the BJ36 shuttle vector and shuttled to the HART01 expression
789 vector as described previously [79]. The folding structure and design procedures were

790 explained in detail in [63,79]. The sequences for the amiRNAs were designed to retain
791 the exact physical properties of the endogenous *MpmiR160* template. The folding
792 patterns were analyzed in the Mfold software [80] and the sequences synthesized (Figure
793 S2).

794 CRISPR/Cas gRNA: Selection of target sites was done using CasFinder [81].
795 Complementary oligos were designed and annealed for ligation into gRNA, in the
796 pMpGE_En03 vector, previously digested with Bsal restriction enzyme. Resulting gRNAs
797 were incorporated into binary vector pMpGE011 through gateway recombination.

798 Over-expression of *MpFER* and *AtFER*: The full-length *MpFER* sequence was amplified
799 from genomic *M. polymorpha* DNA with attB sites and Gateway-cloned via pDONR207.
800 For *A. thaliana* expression, *MpFER* was introduced into the expression vectors
801 pMDC201, which contains a *2X35S* promoter fragment and an in-frame C-terminal
802 mGFP6 [82]. For *M. polymorpha* expression, *MpFER* was introduced into the binary
803 vector pMpGWB308, which contains a *MpEF1* promoter and an in-frame C-terminal
804 citrine [83]. Similarly, for expression of *AtFER* in *M. polymorpha*, coding sequence of
805 *AtFER* was introduced into pMpGWB308 binary vector, for the generation of
806 *proMpEF1:AtFER-Cit* construct.

807

808 **Quantitative real-time PCR (qRT-PCR) and droplet digital PCR (ddPCR)**

809 RNA extraction from *M. polymorpha*: RNA extraction from thallus tissue (ca. 100mg
810 including apical notch) was done using the Rneasy plant mini kit (#74904, Qiagen). To
811 remove contaminating DNA, the TURBO DNAfree™ Kit (AM1907 Ambion) was used
812 according to manufacturer's recommendations. To extract RNA from the gametophores
813 (2 antheridiophores > 5mm diameter were pooled, respectively, for each replicate), the
814 Direct(-zol) RNA MiniPrep (#R2050, ZymoResearch) was used with TRIzol Reagent

815 (#15596026, Ambion) according to the manufacturer's protocol, including on-column
816 DNase digestion and subsequent TURBO DNA-free™ Kit (AM1907 Ambion) treatment.
817 RNA samples were quantified using the Qubit® RNA HS Assay Kit (#Q32852, Life
818 Technologies), a Nanodrop ND-1000 Spectrometer or an Agilent 2100 BioAnalyzer.
819 cDNA synthesis: cDNA was synthesized from 0.5 µg or 1.0 µg of total RNA. Reverse
820 transcription was performed in 25 µl with 20 µg/ml Oligo(dT) 12-18 primers and 200 units
821 of SuperScript® II Reverse Transcriptase (#18064-014, Invitrogen) according to the
822 manufacturer's protocol. The resulting cDNA was diluted 1:9 by adding nuclease-free
823 water. To control for genomic DNA contamination, control replicates of all samples were
824 incubated without SuperScript® II reverse transcriptase.
825 Primer tests and quantitative RT-PCR: Primer efficiency and concentration tests were
826 carried out for *MpFER* and suitable reference genes as described previously (Rövekamp
827 et al., 2016, Althoff et al., 2014). Amplification experiments were carried out using a 7500
828 Applied Biosystem Fast Real-Time PCR System. Reactions were performed in 20 µl
829 volumes containing 10 µl 2X SYBR-green mastermix (SsoAdvanced™ Universal
830 SYBRGreen Supermix), 200 nM (*MpEF1* and *MpFER*) or 250 nM (*MpACTIN*) forward and
831 reverse primers, and 1 µl diluted cDNA. Where possible, three technical and biological
832 replicates were performed for each reaction. The primers used for the qRT-PCR are
833 summarized in Table S1.
834 For ddPCR analysis, individual PCR reactions were performed in a total volume of 25 µl,
835 using 1X ddPCR EvaGreen Supermix, with droplets generated according to
836 manufacturer's recommendations. Reading of the PCR-amplified droplets was carried out
837 by the QX200 Droplet Reader (Bio-Rad) and analysed by the QuantaSoft™ Software
838 (v1.4, Bio-Rad). The counts from ddPCR were normalised through a log₂ transformation.
839 Afterwards, relative expression was calculated against the geometric mean of the counts

840 for all three reference genes (*MpACT1*, *MpACT7*, and *MpAPT3*) [84]: ($\log_2(\text{gene tested}$
841 $+ 1) - \log_2(\text{geom. mean of all references} + 1)$). The primers used for the ddPCR are
842 summarized in Table S1.

843

844 **Transformation into *M. polymorpha* and *A. thaliana***

845 *Agrobacterium*-mediated transformation using regenerating thalli of *M. polymorpha* was
846 done by co-cultivation 15-days old Marchantia fragments with transformed *Agrobacterium*
847 *tumefaciens* (GV3101) cells [85,86]. After three days of incubation, positive transformants
848 were selected on Gamborg's B5 plates supplemented with 10 $\mu\text{g/ml}$ Hygromycin B
849 (Invitrogen), 0.5 μM Chlorosulfuron or Kanamycin, and 100 $\mu\text{g/ml}$ of Cefotaxime sodium.
850 Isogenic lines were obtained by using plants derived from gemmae of the T1 generation
851 (G1 generation). G1 or subsequent gemmae generations were used for all experiments,
852 as they are derived from single cells [87,88]. Transformation of *A. thaliana* via *A.*
853 *tumefaciens* (GV3101) was performed by floral dipping according to [89]. Primers used
854 for amplification of promoter fragments of *MpFER* are listed in Table S2.

855

856 **Microscopy**

857 Plants were observed in a Lumar.V12 (Zeiss) or a Leica MZFLIII dissection microscope
858 and photographed with an AxioCam HRc (Zeiss) or Leica DFC 420C camera. Clearings
859 were observed using a Leica DMR microscope and photographed with an AxioCam 105
860 color camera. Fluorescence reporter expression were analyzed using a Leica SP5
861 confocal microscope. Fiji [90] and GIMP (version 2.8.10) software were used for
862 adjustments of brightness, contrast, channel selection, and image size.

863 Tissue clearing for bright field microscopy: *M. polymorpha* tissues were fixed in Carnoy's
864 solution and incubated at 4°C overnight, followed by rehydration in an ethanol series of

865 85%, 70%, 50%, and 30%. Samples were incubated for 1 hour at 4°C for each step and
866 then incubated in chloral hydrate solution at 4°C overnight. Samples were mounted in
867 chloral hydrate.

868 Calcofluor-white staining for cell size measurements: Pieces of fresh thallus tissue
869 located in fully differentiated zones [91] of plants grown on plates were dissected out and
870 put into water. The pieces were transferred to PBS pH 6.1 containing 100 µg/ml
871 Calcofluor-white and vacuum-infiltrated for 1 hour before being mounted on slides. The
872 area of the epidermal cells was measured using Imaris 8.3.1. software. Cells of air pores
873 and the two adjacent cell layers were excluded from the analysis, as well as the spiralling
874 cells surrounding newly developing air pores.

875 For cell-death staining, seven-days old gemmalings were stained with lactophenol-
876 trypanblue (10 mL of lactic acid, 10 mL of glycerol, 10 g of phenol, 10 mg of trypan blue,
877 dissolved in 10 mL of distilled water) and boiled for approximately 1 min in the stain
878 solution and then decolorized in chloral hydrate (2.5 g of chloral hydrate dissolved in 1
879 mL of distilled water) for at least 30 min. They were mounted in chloral hydrate and
880 imaged in dissecting microscope.

881

882 **Western blot analysis**

883 *M. polymorpha* explants were ground in liquid nitrogen and resuspended in 300 µL of
884 Laemmli buffer [92]. After centrifugation at 6,000 rpm for 5 min at 4°C, the supernatant
885 was recovered. Protein accumulation was confirmed by Western blotting, using the anti-
886 GFP antibody (1:3000, Torrey Pines Biolabs). As secondary antibodies, the Agrisera
887 S09602 goat anti-rabbit antibody was used with the ECL-chemistry of FUSION FX -
888 Western Blot & Chemi Imaging (Vilber Lourmat).

889

890 **Growth rate experiments**

891 Gemmae of wild-type, *Mpfer-1*, and independent transformation lines of amiR-*MpFER*
892 lines were grown on plates and the thallus area was measured over time. 10 or 30
893 gemmae of four independent transformation lines were grown on half-strength Gamborg's
894 B5 and scanned on an Epson 2450 Photo scanner at 600dpi. Plant area was measured
895 using color threshold settings and particle analysis in the Fiji software [90].

896

897 **Expression map**

898 Gene expression of *CrRLK1L* signaling pathway components was calculated from
899 publicly available RNA-seq data of 39 samples of *M. polymorpha* Tak-1 and Tak.2.
900 Samples were grouped by tissue and analyzed under the same pipeline: Nextera paired-
901 end adapters were trimmed from sequencing reads using the bbdduk tool embedded in
902 the BBMap software package [93]. Read ends with quality below 20 were also trimmed
903 and the minimum read length was set to 25. The rest of parameters of bbdduk were set as
904 default. Reads were mapped to *M. polymorpha* reference genome (v3.1) [45] using the
905 Tophat software (v2.1.1) [94], designed for RNA-sequencing read mapping. Tophat
906 contains Bowtie (v2.3.2.0) [95] as the aligner software. All parameters were set to default
907 values except for the RAM memory and number of used Cores. Mapped reads were
908 sorted using samtools (v1.3.1) [96] and counted with the software "HTSeq-count" (v.0.9.1)
909 [97]. Gene expression values were calculated using the package DESeq2 (Release 3.6)
910 [98] for R software (v3.4.4) [99]. Another R package, ggplot2 (release 2.2.1) [100] was
911 used for producing the figures. A summary of the samples that were used in this study is
912 provided in Table S3.

913

914

915 **Three-dimensional (3D) protein structure modeling**

916 Comparative modeling of protein 3D structures was performed using the SWISS-MODEL
917 online-tool [60]. The software was fed with the three protein sequences of MpFER,
918 MpRALF1, and MpLRE1 as input to model, and one PDB file as a 3D template, which
919 contains the published crystal structure of the protein complex formed by AtFER,
920 AtRALF23, and AtLLG2 [35]. Quality control of the model was assessed using the same
921 tool and the results are summarized in Figure S5E. Spatial comparisons of the *M.*
922 *polymorpha* predicted structures with the crystal structures of *A. thaliana* were performed
923 in PyMol [101].

924

925 **QUANTIFICATION AND STATISTICAL ANALYSIS**

926 Analyses were performed in InfoStat (<http://www.infostat.com.ar>). Tests are indicated in
927 the corresponding figure legend.

928

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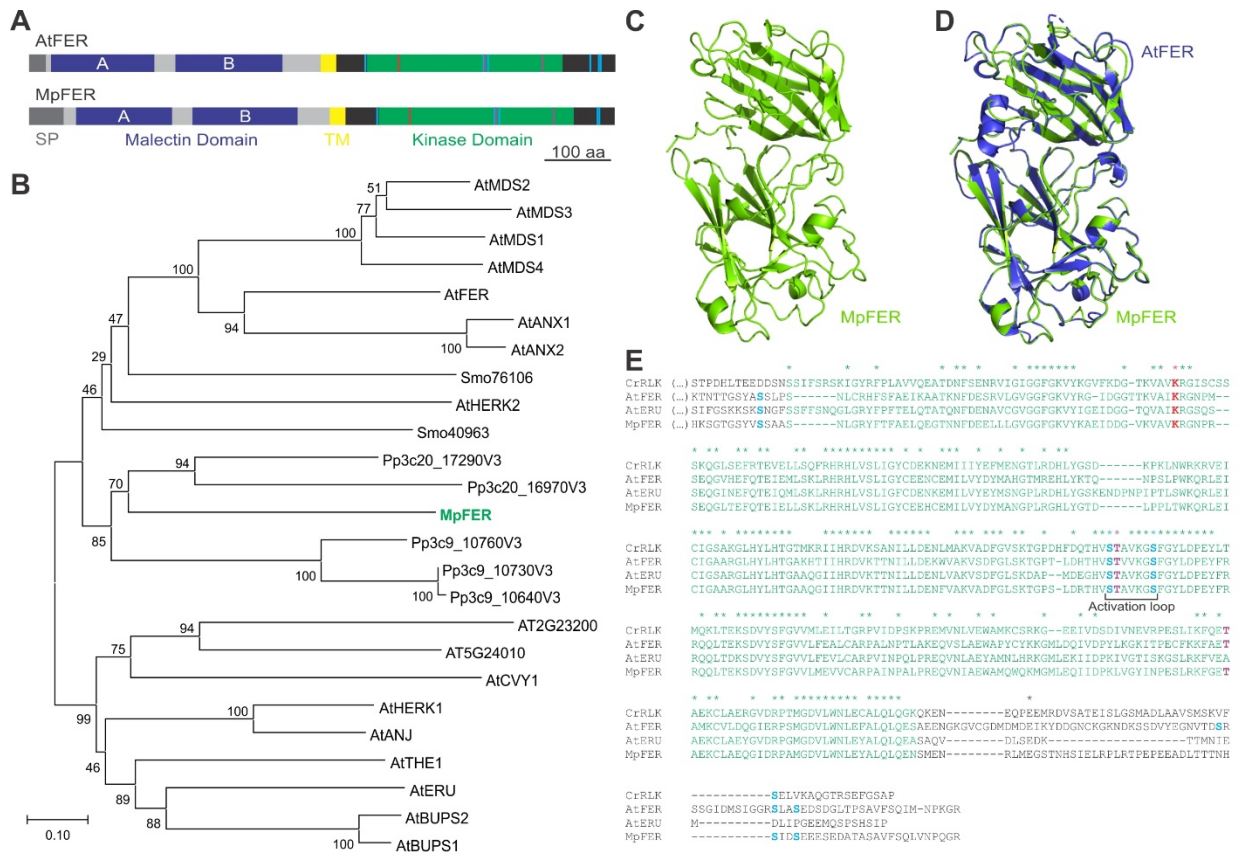
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1180 **Figure 1. CrRLK1Ls Are Conserved among Land Plants**

1181 (A) Representation of AtFER and MpFER proteins. Malectin-like domains are
 1182 represented in blue and kinase domains in green. SP, signal peptide; TM, transmembrane
 1183 domain (yellow). Amino acids important for regulation and activity are marked in red,
 1184 violet, and light blue as described in Figure 1E.

1185 (B) A rooted neighbor-joining tree of the amino acid sequence of the predicted malectin-
 1186 like domain. CrRLK1L members from *Marchantia polymorpha* (Mp), *Physcomitrella*
 1187 *patens* (Pp), *Selaginella moellendorffii* (Smo), and *Arabidopsis thaliana* (At) were used.
 1188 The numbers indicate the bootstrap values (%) from 1000 replications. The given scale
 1189 represents a substitution frequency of 0.1 amino acids per site.

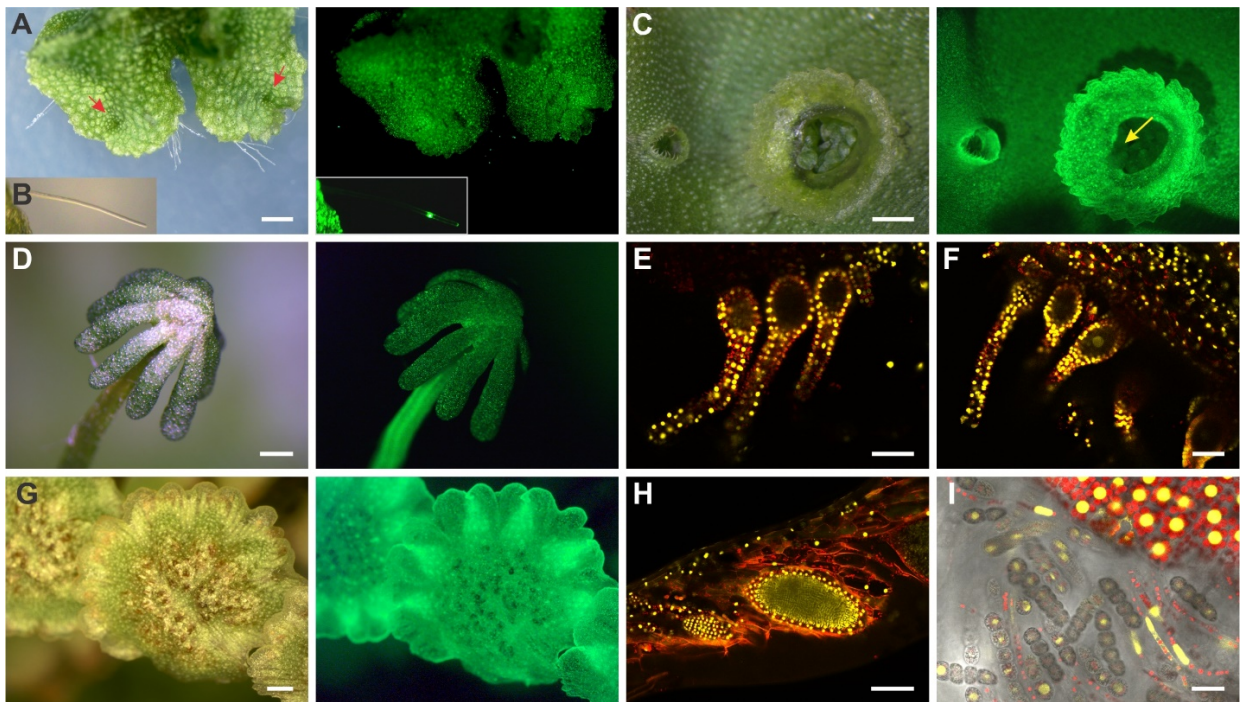
1190 (C) Cartoon representation of the predicted three-dimensional structure of the MpFER
1191 malectin-like domain, showing predicted alpha-helix and beta-sheet structures.

1192 (D) Structural superposition of the malectin-like domains of AtFER (blue) and MpFER
1193 (green).

1194 (E) Alignment of the cytoplasmic domains of MpFER, AtERU, AtFER, and CrRLK. Kinase
1195 domains are in green, putative phosphorylation sites in light blue (Ser) and violet (Thr),
1196 and the conserved catalytic Lys in red.

1197 See also Figure S1

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1201 **Figure 2. MpFER Is Broadly Expressed in most Tissues throughout the *M.***
1202 ***polymorpha* Life Cycle**

1203 Expression of *proMpFER:trpVNS* in different organs of male and female *M. polymorpha*
1204 plants. Fluorescence is visualized as either green (epifluorescence microscope) or yellow
1205 (confocal microscope) color, depending on the panel.

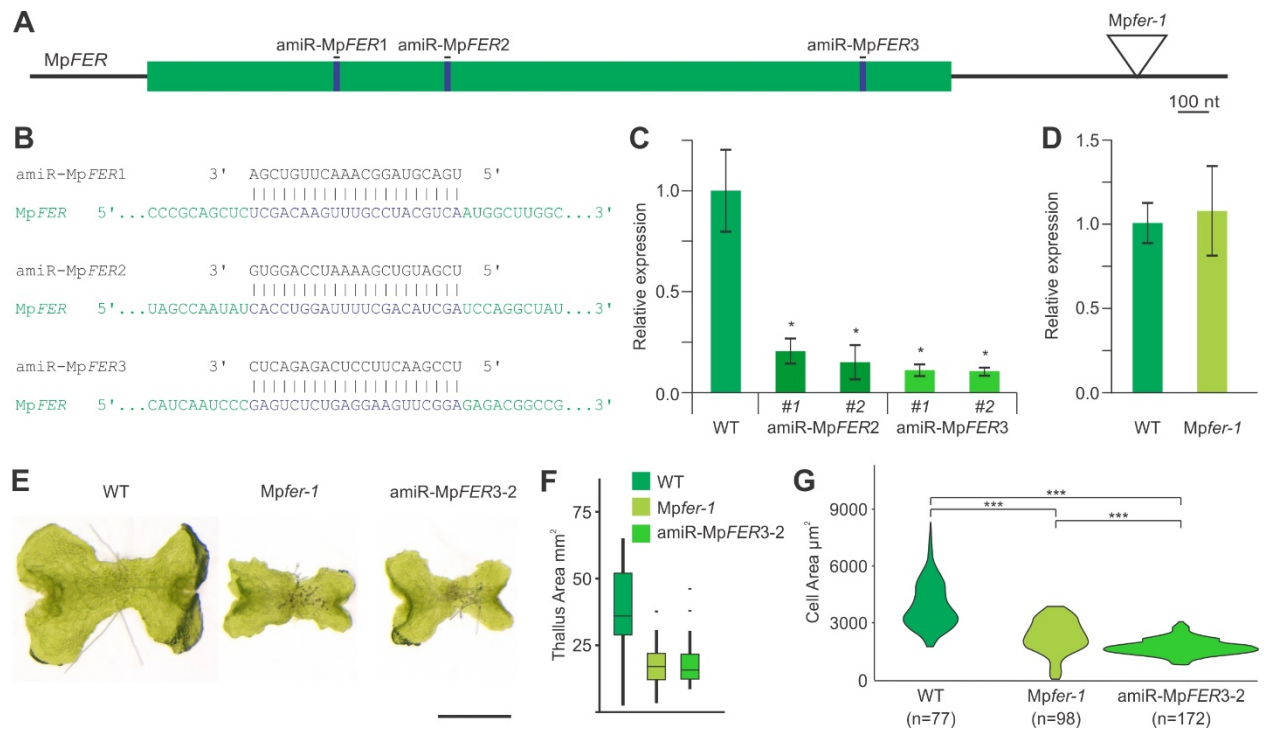
1206 (A to D) Bright field (left) and epifluorescence (right) images of a thallus, (A) rhizoid (C),
1207 gemmae cups (C), and an archegoniophore (D). Red arrows indicate meristematic zones,
1208 the yellow arrow gemmae. Scale bars, 0.5 mm.

1209 (E and F) Confocal images of archegonia before (E) and 2 days after fertilization (F).
1210 Scale bar, 50 μ m.

1211 (G) Bright field (left) and epifluorescence (right) images of an antheridiophore. Scale bar,
1212 1 mm.

1213 (H) Confocal image of antheridia. Scale bar, 100 μ m.

1214 (I) Confocal image of sporophyte and spores. Scale bar, 25 μ m.



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1217 **Figure 3. MpFER Controls Cell Size during Gametophyte Development**

1218 (A) Representation of the mature *MpFER* mRNA. Coding sequence are in green,
 1219 locations of the amiRNA target sites in blue.

1220 (B) Base complementarity between mature amiR-*MpFER1*, amiR-*MpFER2*, and amiR-
 1221 *MpFER3* with the *MpFER* transcript.

1222 (C) Relative expression level of *MpFER* in thallus tissue from wild-type (WT; Tak1) and
 1223 two independent insertion lines of amiR-*MpFER2* and amiR-*MpFER3*, as measured by
 1224 qRT-PCR. *MpEF1* was used as internal control. Shown are means \pm standard errors of
 1225 the means (SEM) of three biological replicates. * $P < 0.01$, one-way analysis of variance
 1226 (ANOVA).

1227 (D) Relative expression level of *MpFER* in WT and *Mpfer-1* thallus tissue, as measured
 1228 by qRT-PCR. *MpEF1* was used as internal control. Shown are means \pm SEM of three
 1229 biological replicates. * $P < 0.01$, one-way ANOVA.

1230 (E) Representative pictures of 10-day old gemmalings of WT, *Mpfer-1*, and amiR-
1231 *MpFER3-2* lines. Scale bar, 1 mm.

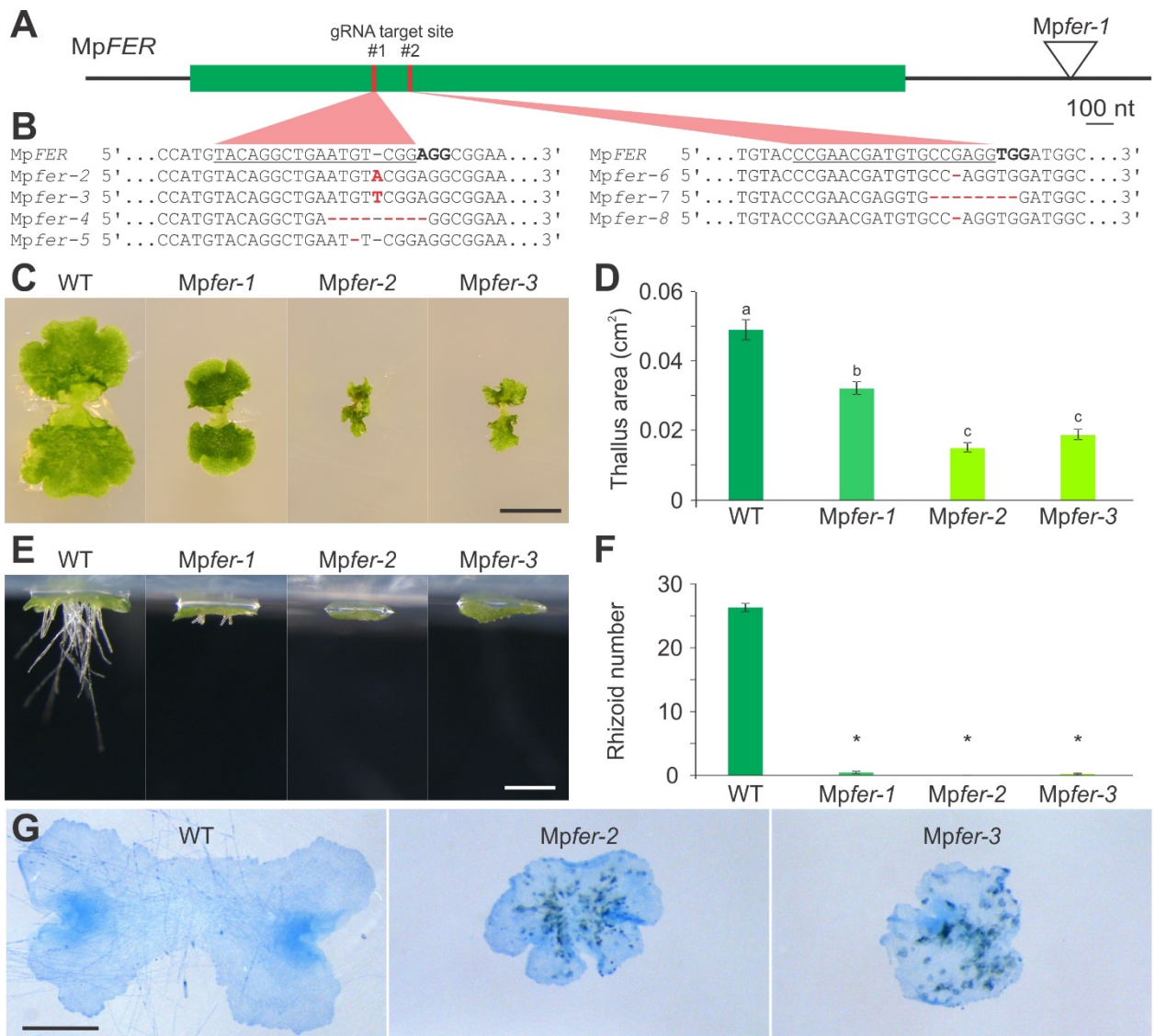
1232 (F) Thallus area of WT, *Mpfer-1*, and amiR-*MpFER3-2* lines. n = 30, outliers are indicated
1233 as black dots.

1234 (G) Violin plot of cell size in WT, *Mpfer-1*, and amiR-*MpFER3-2* lines. Significant
1235 differences are indicated according to the non-parametric Kruskal-Wallis test and linear
1236 regression models (**p<0.001).

1237 See also Figures S2 and S3

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1242 **Figure 4. The Integrity of Thalli from *Mpfer* Knock-out Lines Is Severely Affected**

1243 (A) Representation of the mature *MpFER* mRNA with the coding sequence in green, and
 1244 the location of the gRNA target sites in red.

1245 (B) Sequences of gRNA target sites in the confirmed *Mpfer* knock-out mutants. Deletions
 1246 and insertions are highlighted in red.

1247 (C) Representative pictures of 10-day old gemmalings of WT, *Mpfer-1*, and two
 1248 independent *Mpfer* knock-out mutants. Scale bar, 0.5 cm.

1249 (D) Thallus area of 10-day old gemmalings of WT, *Mpfer-1*, and two independent *Mpfer*
1250 knock-out mutants. n = 30.

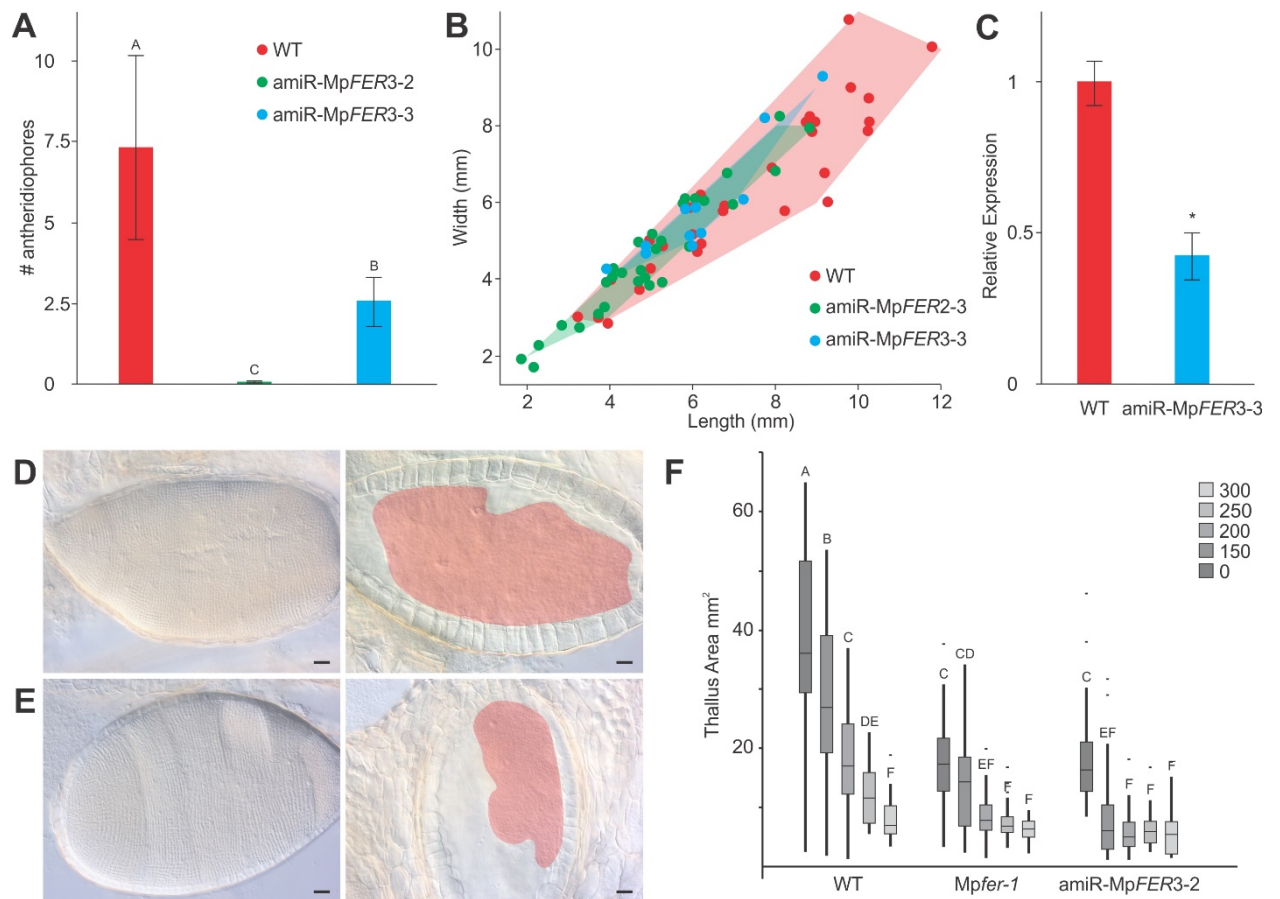
1251 (E) Representative pictures of 3-day old gemmalings growing in upside-down plates of
1252 WT, *Mpfer-1*, and two independent *Mpfer* knock-out mutants. Scale bar, 500 μ m.

1253 (F) Rhizoid number in 3-day old gemmalings of WT, *Mpfer-1*, and two independent *Mpfer*
1254 knock-out mutants. n = 30.

1255 (G) Trypan blue staining of 7-days old gemmalings of WT, *Mpfer-1*, and two independent
1256 *Mpfer* knock-out mutants. Scale bar, 1 mm.

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1261 **Figure 5. Conservation of CrRLK1L Function in Land Plants**

1262 (A) Number of antheridiophores produced per plant for wild-type (WT, n=9) and two
 1263 independent amiR-MpFER3 lines (amiR-MpFER3-2, n = 9; amiR-MpFER3-3, n = 12).
 1264 Shown are means \pm SEM of two biological replicates. Means with same letter do not differ
 1265 significant different with $P > 0.05$, one-way ANOVA, Duncan test.

1266 (B) Antheridiophore splash platform size distribution of WT (n = 32), amiR-MpFER2-3 (n
 1267 = 34), and amiR-MpFER3-3 (n = 11) lines. For all antheridia with a stalk >8 mm, length
 1268 and width of the platform were recorded.

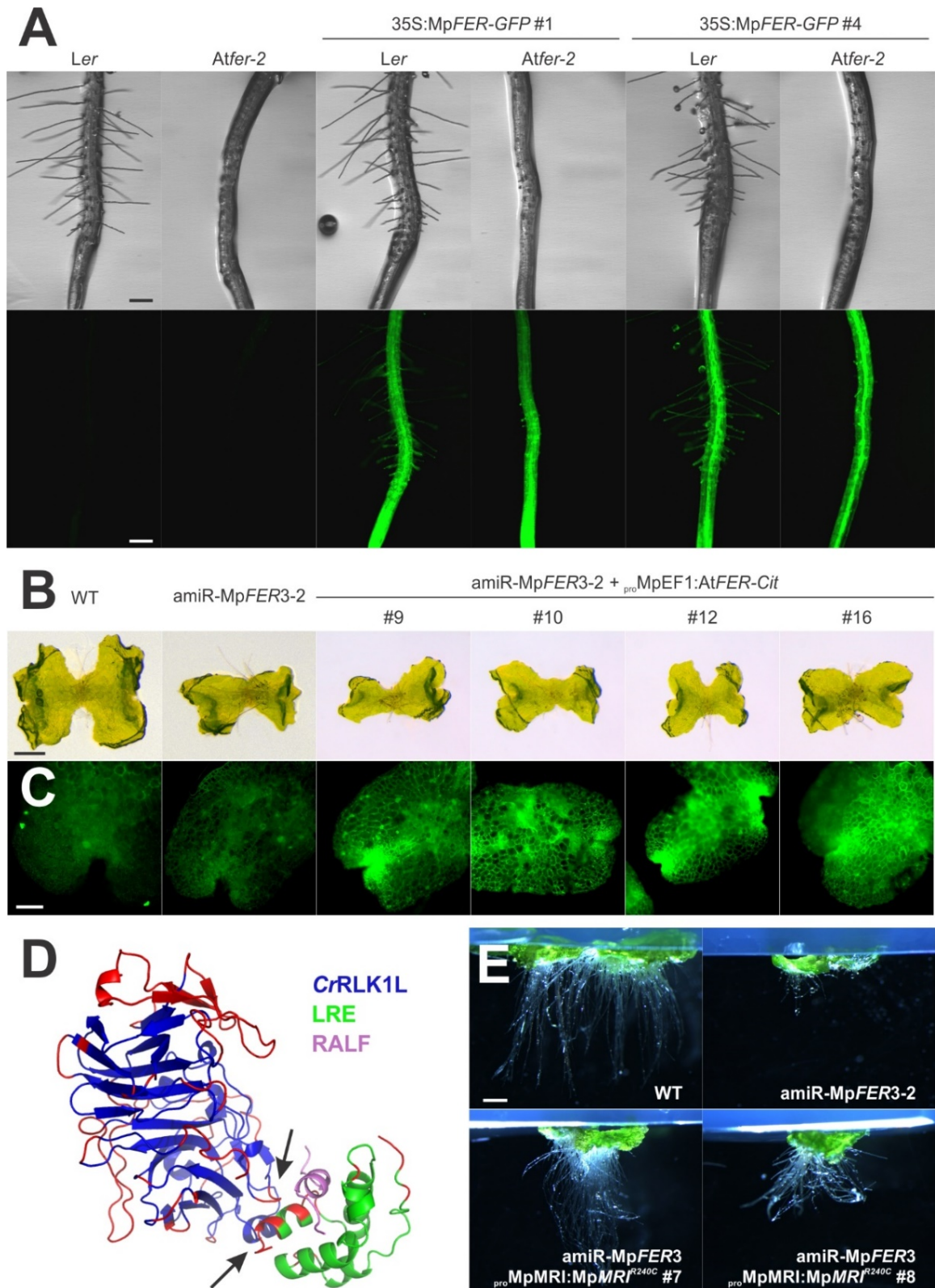
1269 (C) qRT-PCR of MpFER levels in antheridiophores in a WT and amiR-MpFER3-3 line.
 1270 MpEF1 was used as internal control. Shown are means \pm SEM of three biological
 1271 replicates. * $P < 0.01$, ANOVA.

1272 (D and E) Mature antheridia of a WT (D) and amiR-Mp*FER3-3* line (E). Bright field images
1273 (left) and in cross-section (right). The spermatogenous areas are indicated in red. Scale
1274 bar, 100 μ m.

1275 (F) Thallus area of 15-day old gemmalings of WT, *Mpfer-1* mutant, and an amiR-
1276 Mp*FER3-2* line growing on media containing different isoxaben concentrations (nM). n =
1277 30, outliers are indicated as black dots. Shown are means \pm SEM of two biological
1278 replicates. Means designated by the same letter do not significantly differ at $P > 0.05$,
1279 one-way ANOVA, Duncan test.

1280 See also Figure S3

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1284 **Figure 6. Interspecific Complementation of *A. thaliana* and *M. polymorpha* Plants**

1285 **with Reduced Levels of *FER* Activity**

1286 (A) Roots of independent *A. thaliana* lines expressing 35S:MpFER-GFP in Ler wild-type
1287 (WT) and *Atfer-2* plants. Scale bar, 200 μ m.

1288 (B) 10-day old gemmalings of WT and amiR-MpFER3-2 *M. polymorpha* plants with and
1289 without expression of *AtFER-Cit*. Scale bar, 0.5 mm.

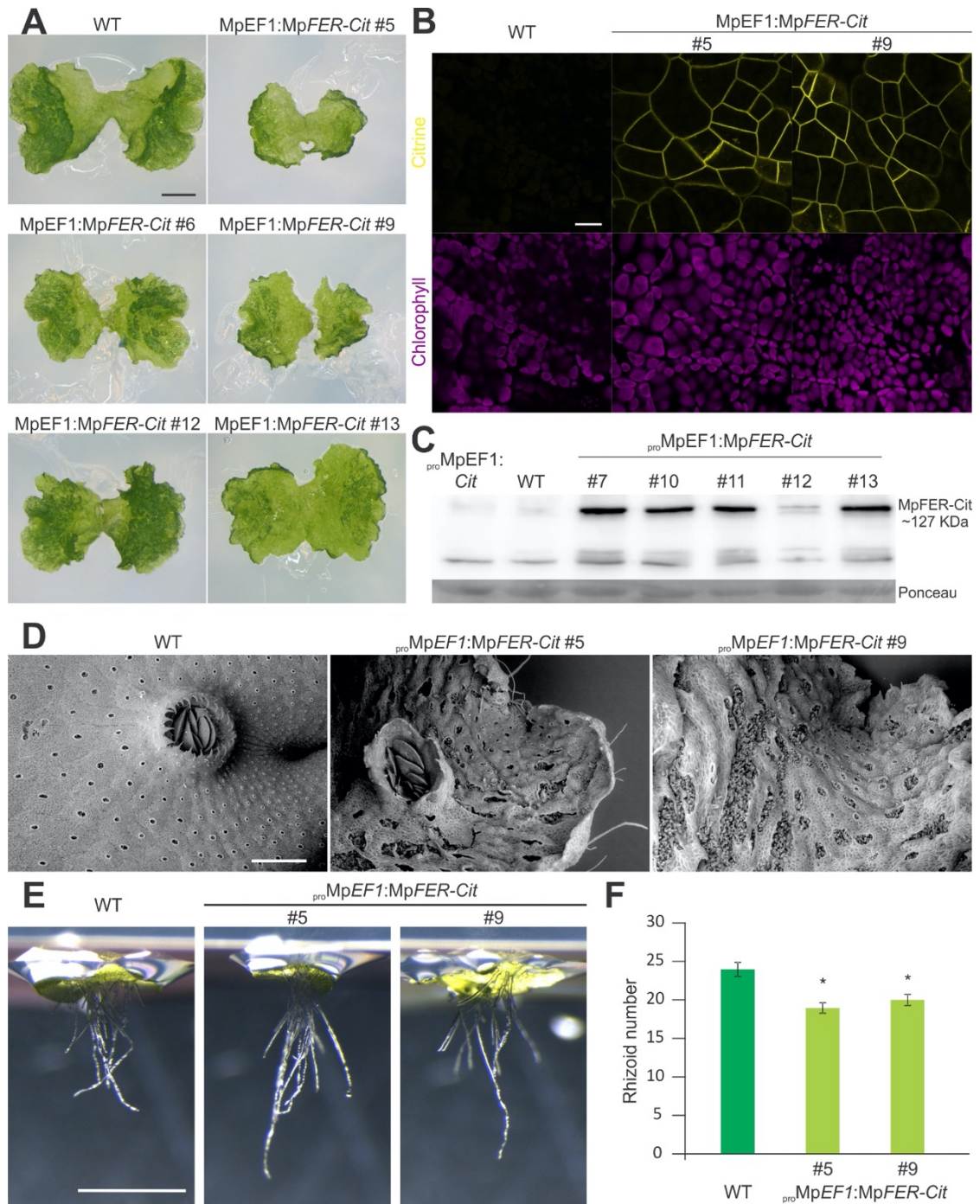
1290 (C) Citrine expression in 7-day old gemmalings of WT and amiR-MpFER3-2 plants with
1291 and without expression of *AtFER-Cit*. Scale bar, 100 μ m.

1292 (D) Cartoon representation of the structural conservation of the CrRLK1L^{ECD}/LRE/RALF
1293 complex by comparing AtFER/MpFER and AtLLG2/MpLRE1. Blue and green represent
1294 spatially highly conserved regions of CrRLK1L and LRE homologs, respectively. Red
1295 indicates structural regions with low spatial conservation, lilac the C-terminal part of
1296 AtRALF23, and black arrows the interaction regions of the three proteins.

1297 (E) Complementation of amiR-MpFER lines with a dominant-active version of MpMRI
1298 under its own promoter (*proMpMRI:MRI^{R240C}*). Scale bar, 1 mm.

1299 See also Figures S4-S7

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1303 **Figure 7: Overexpression of MpFER Affects Morphological Integrity**

1304 (A) Representative pictures of 10-day old gemmalings of wild-type (WT, Tak1) and
1305 different lines overexpressing MpFER (*proMpEF1:MpFER-Cit*). Scale bar, 1 mm.

1306 (B) Citrine expression at the plasma membrane in WT and *proMpEF1:MpFER-Cit* lines #5
1307 and #9 as observed under confocal microscopy. Scale bar, 20 μ m.

1308 (C) Western blot analysis of *proMpEF1:MpFER-Cit* lines using an anti-GFP antibody.
1309 *proMpEF1:Cit* and WT lines were used as positive and negative controls, respectively. The
1310 Ponceau membrane staining of the most intense band at 55 kDa (presumably Rubisco)
1311 was used as a loading control.

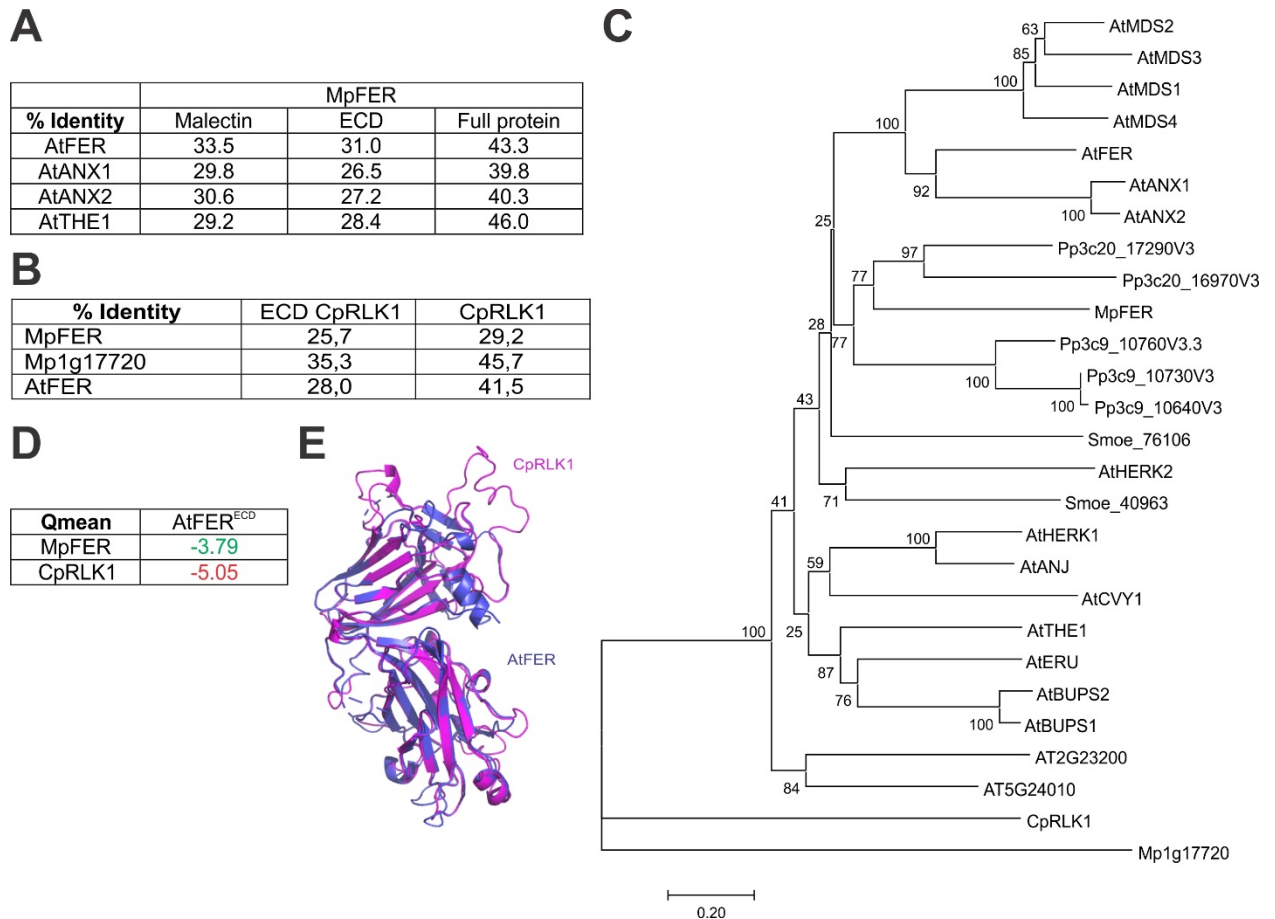
1312 (D) Scanning electron microscopical images of thalli from 20-day old plants of WT and
1313 *proMpEF1:MpFER-Cit* lines #5 and #9. Scale bar, 500 μ m.

1314 (E) Representative pictures of 3-day old gemmalings growing in upside-down plates of
1315 WT and *proMpEF1:MpFER-Cit* lines #5 and #9. Scale bar, 1 mm.

1316 (F) Number of rhizoids in 3-day old gemmalings of WT and *proMpEF1:MpFER-Cit* lines #5
1317 and #9. n = 30.

1318 See also Figure S6

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1321 **Figure S1. The CrRLK1L Family Appeared together with Land Plants. Related to**
 1322 **Figure 1.**

1323 (A) Percentage of identity of the malectin-like domain, the complete ECD, and the full
 1324 protein of MpFER with AtFER, AtANX1, AtANX2, and AtTHE1.

1325 (B) Percentage of identity of the ECD and full protein of CpRLK1 with MpFER, AtFER,
 1326 and Mapoly001s0111.

1327 (C) A rooted neighbor-joining tree of the amino acid sequence of the predicted malectin-
 1328 like domain was generated using ClustalW. CrRLK1L members from *Marchantia*
 1329 *polymorpha* (Mp), *Physcomitrella patens* (Pp), *Selaginella moellendorffii* (Smoe), and
 1330 *Arabidopsis thaliana* (At) were used. Algal CpRLK1 and Mapoly001s0111 were also
 1331 included. The numbers indicate the bootstrap values (%) from 1000 replications. The
 1332 given scale represents a substitution frequency of 0.1 amino acids per site.

1333 (D) Qmean value for prediction of MpFER or CpRLK1 3D structures of the ECD based
1334 on the AtFER ECD. Green numbers indicate a good modelling fit, red numbers indicates
1335 a bad modelling fit.

1336 (E) Structural superposition of the ECD of AtFER (blue) and CpRLK1 (purple).

1337

A *MpmiR160* Mp1g26670
 GCACCTCCTCTCTCCGACTGCAGCCCGTTTTTCGAGATCCGAGGACTTGCTCGACGCGACTAATTGGGGAGGCCAGACTG
 CACTT**GGCTGGCTCCCTGTATGCCA**ACTGAGGAGCTCCTCAGAGACCTTGACAGGCTCCGTAGCTGGCATT**CAGGGGG**
CCATGCAGGAGGAAGTCGCTACCTCCCGCAAGGTGCGACTAGCTTTCTGTCTTGGGTGCACACCTCACTGATGTTTGA
 TAGATTTACTTA

amiR-Mp*FER1*
TGCAAGCTTGCACCTCCTCTCTCCGACTGCAGCCCGTTTTTCGAGATCCGAGGACTTGCTCGACGCGACTAATTGGGGAG
 GCCAGACTGCACT**TGACGTAGGCAA**ACT**TGT**CGAACTGAGGAGCTCCTCAGAGACCTTGACAGGCTCCGTAGCT**CGAC**
ATGTTTGTCTACCTCAGGAGGAAGTCGCTACCTCCCGCAAGGTGCGACTAGCTTTCTGTCTTGGGTGCACACCTCACT
 GATGTTT**GATAGATTTACTTAGGATCCATA**

amiR-Mp*FER2*
TGCAAGCTTGCACCTCCTCTCTCCGACTGCAGCCCGTTTTTCGAGATCCGAGGACTTGCTCGACGCGACTAATTGGGGAG
 GCCAGACTGCACT**TCGATGTCGAAAATCCAGGTG**ACTGAGGAGCTCCTCAGAGACCTTGACAGGCTCCGTAGCT**CCACCT**
GCATTTTTGACAACGAGGAGGAAGTCGCTACCTCCCGCAAGGTGCGACTAGCTTTCTGTCTTGGGTGCACACCTCACT
 GATGTTT**GATAGATTTACTTAGGATCCATA**

amiR-Mp*FER3*
TGCAAGCTTGCACCTCCTCTCTCCGACTGCAGCCCGTTTTTCGAGATCCGAGGACTTGCTCGACGCGACTAATTGGGGAG
 GCCAGACTGCACT**TCGGA**ACT**TCCTCAGAGACTC**ACTGAGGAGCTCCTCAGAGACCTTGACAGGCTCCGTAGCT**CGAGTC**
TGTGAGGGAGTTGGGAGGAGGAAGTCGCTACCTCCCGCAAGGTGCGACTAGCTTTCTGTCTTGGGTGCACACCTCACT
 GATGTTT**GATAGATTTACTTAGGATCCATA**

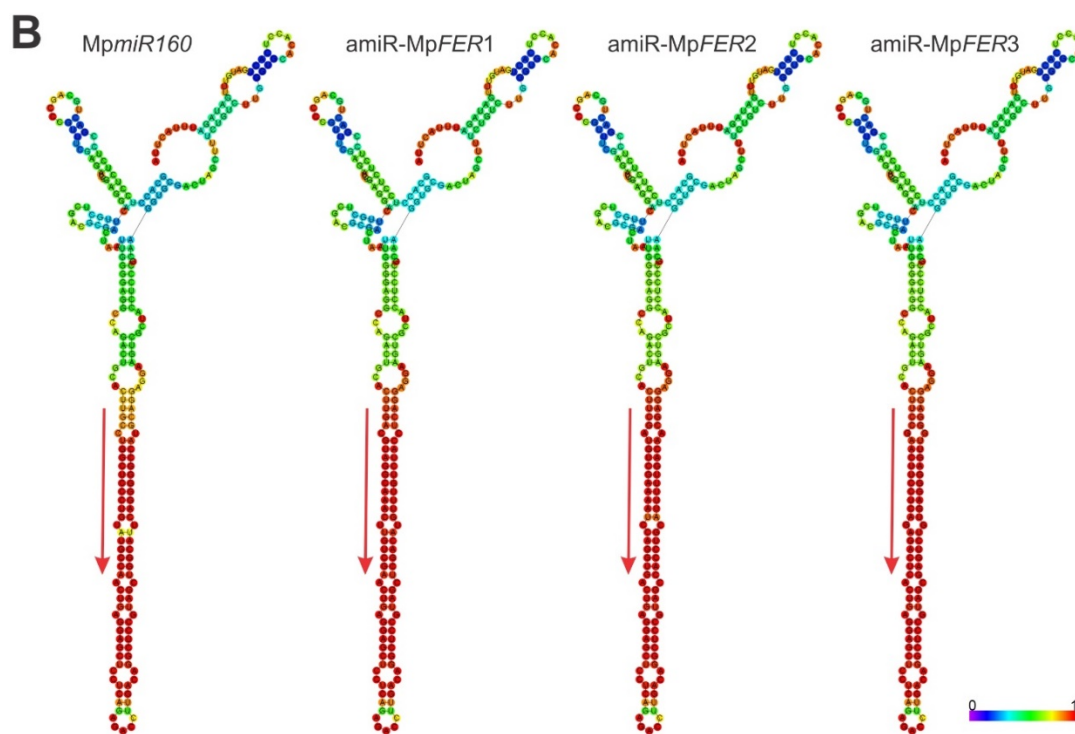
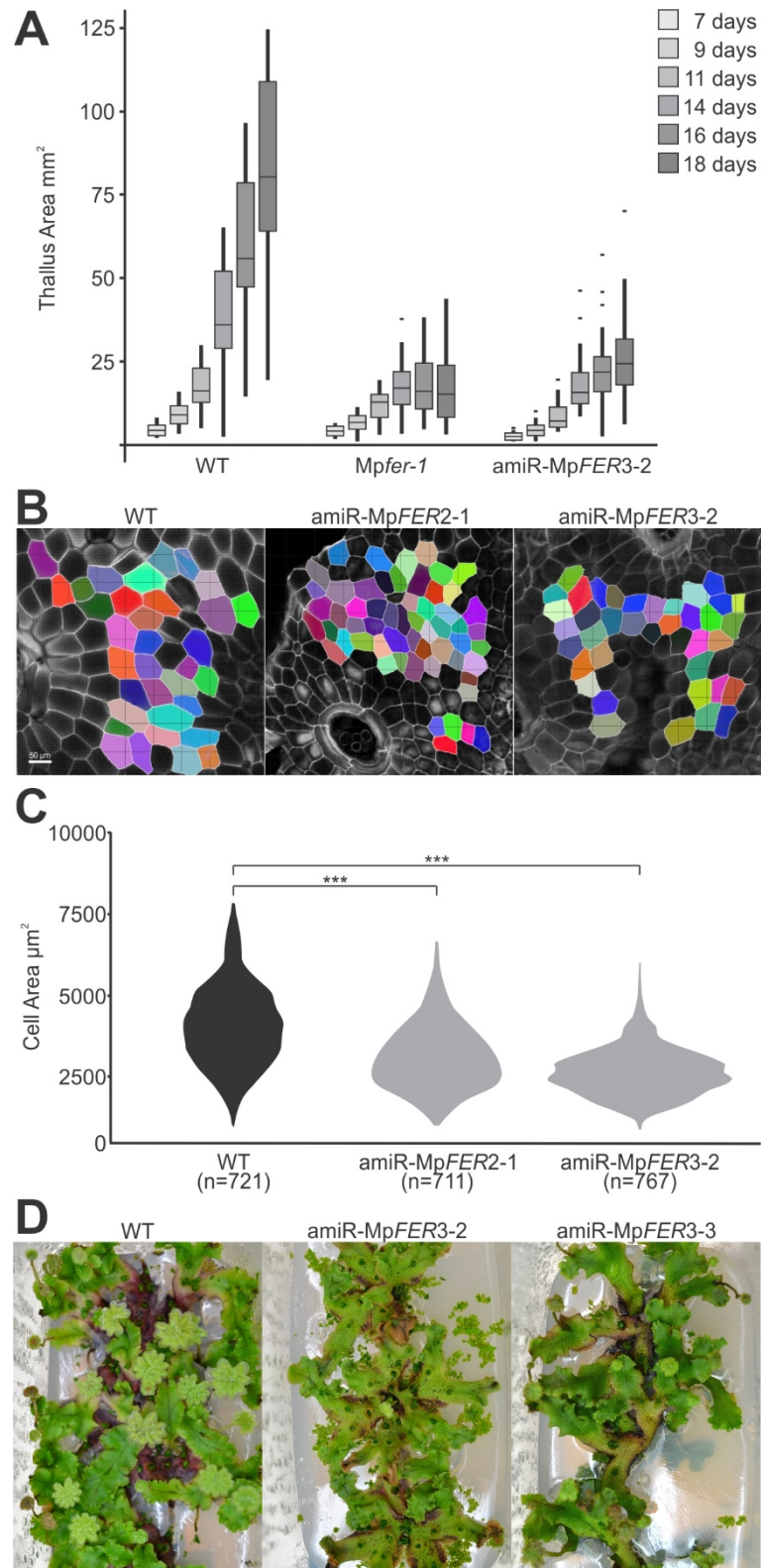


Figure S2. Design of amiR-Mp*FER* Precursors. Related to Figure 3.

(A) *MpmiR160* (Mapoly0002s0211, Mp1g26670) and amiR-Mp*FER1*, amiR-Mp*FER2*, and amiR-Mp*FER3* sequences. miRNA sequences are in red and miRNA* in blue. Cloning sequences from amiR-Mp*FER* constructs are in bold.

1344 (B) Drawing of the minimum free energy structure of *MpmiR160* and *amiR-MpFER3*
1345 constructs predicted by the RNAfold web server ([http://rna.tbi.univie.ac.at/cgi-](http://rna.tbi.univie.ac.at/cgi-bin/RNAWebSuite/RNAfold.cgi)
1346 [bin/RNAWebSuite/RNAfold.cgi](http://rna.tbi.univie.ac.at/cgi-bin/RNAWebSuite/RNAfold.cgi)). Red arrows indicate location and orientation of the
1347 mature miRNA in the precursor. The structures are colored by base-pairing probabilities;
1348 for unpaired regions the color denotes the probability of being unpaired.
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1352 **Figure S3. Reduction in Thallus and Cell Area in Plants with Reduced MpFER**

1353 **Levels. Related to Figures 3 and 5.**

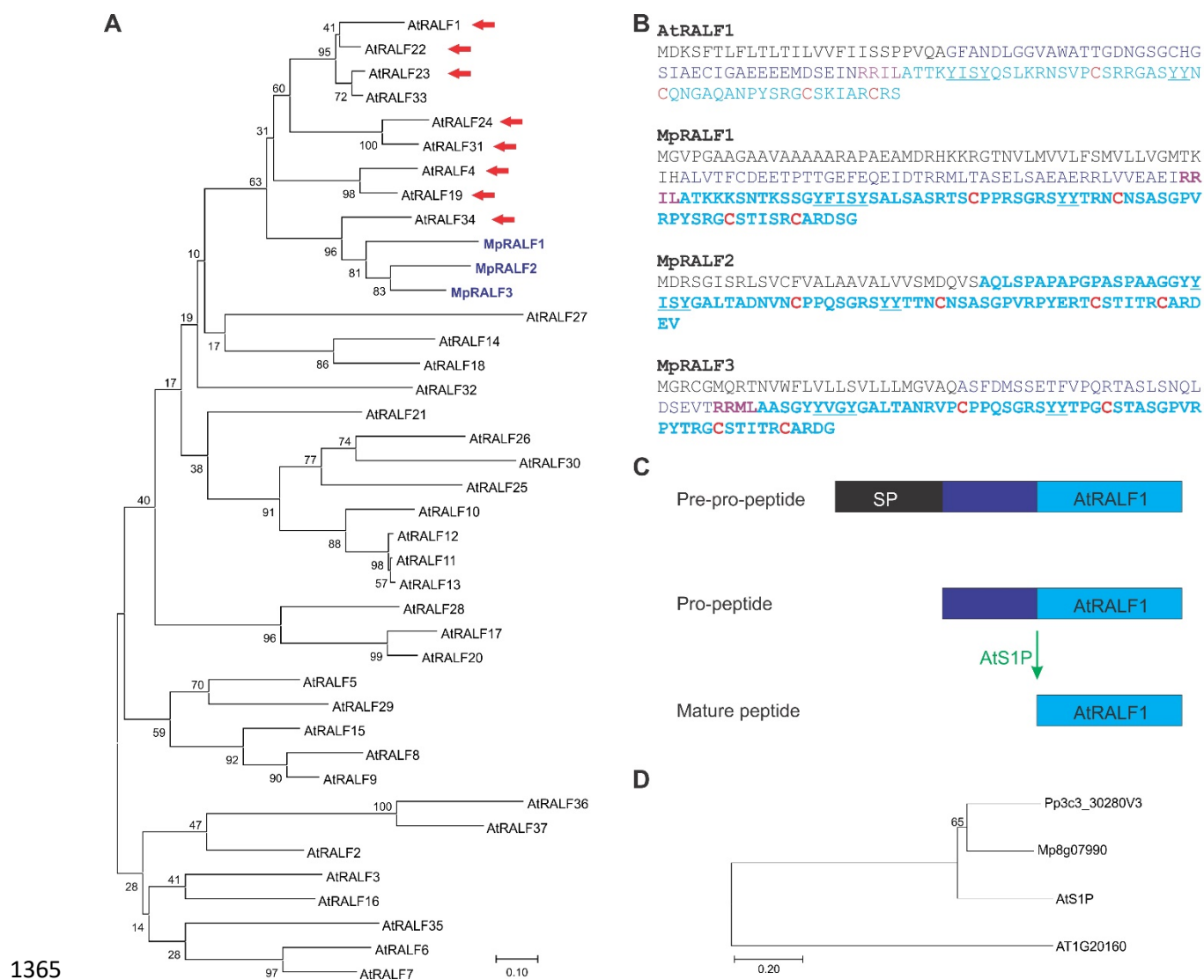
1354 (A) Thallus area of wild-type (WT), *Mpfer-1*, and *amiR-MpFER3-2* plants at different days
1355 after putting gemmae on plates. $n = 30$, outliers are indicated as black dots. Areas were
1356 estimated using ImageJ software.

1357 (B) Representative images from cell surface areas measured in WT, *amiR-MpFER2-1*,
1358 and *amiR-MpFER3-2* plants. Scale bar, $50 \mu\text{m}$.

1359 (C) Difference in cell size between WT, *amiR-MpFER2-1*, and *amiR-MpFER3-2* plants is
1360 significant ($p < 0.001$) based on the nonparametric Kruskal-Wallis test and a linear
1361 regression model with a highly significant interaction ($p < 0.001$).

1362 (D) Induction of antheridiophores in WT and two independent *amiR-MpFER3* lines. Three
1363 plants were grown in each sterile plastic box under far-red light induction.

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1367 **Figure S4. MpRALF Peptides Belong to the AtRALF1-clade of the RALF Family.**

1368 **Related to Figure 6.**

1369 (A) A rooted neighbor-joining tree of the amino acid sequence of the predicted mature

1370 RALF peptides was generated using ClustalW. RALF members from *M. polymorpha* and

1371 *A. thaliana* were used. Red arrows indicate RALFs that are known ligands of CrRLK1Ls.

1372 The numbers indicate the bootstrap values (%) from 1000 replications. The given scale

1373 represents a substitution frequency of 0.1 amino acids per site.

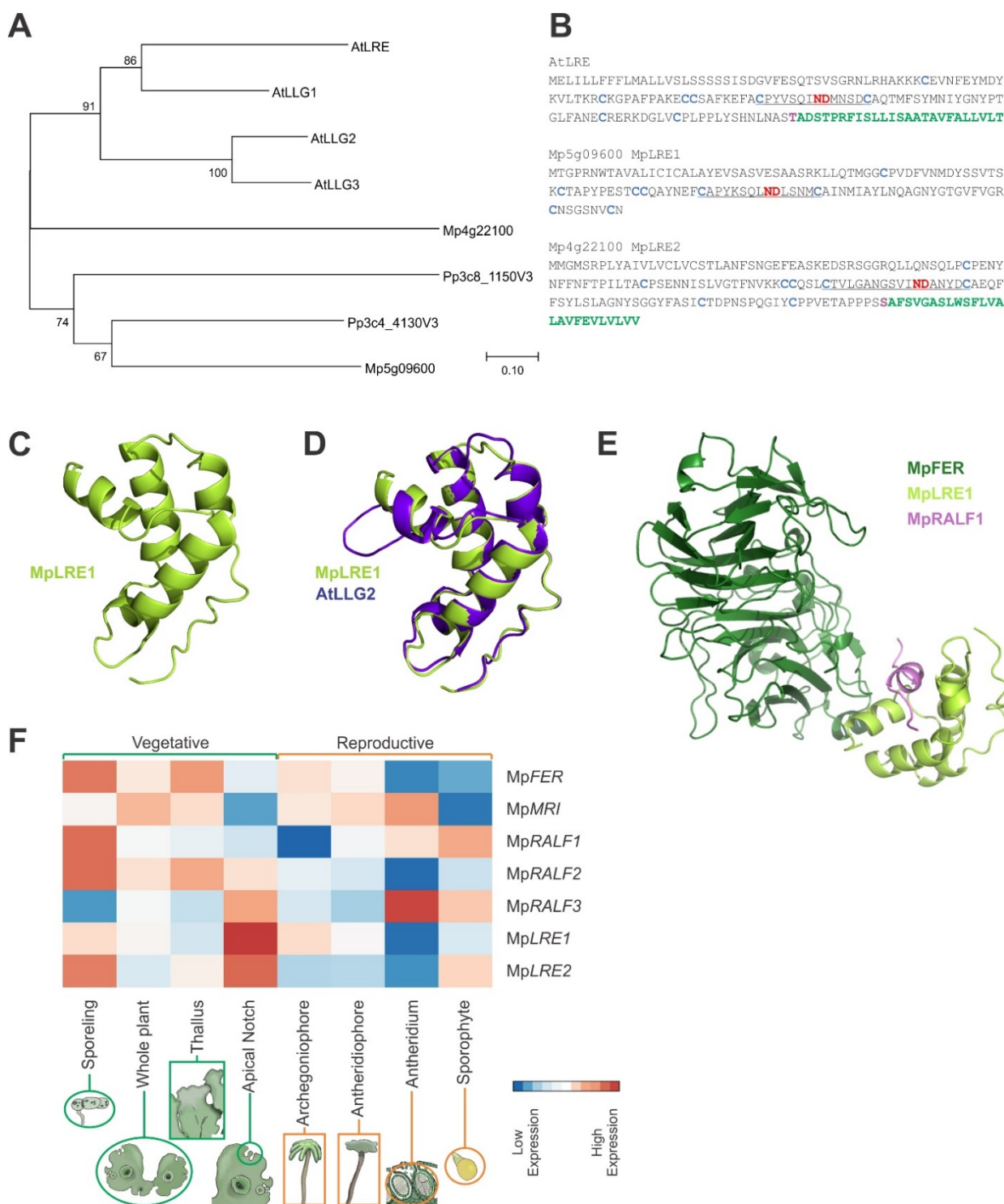
1374 (B) Amino acid sequence comparison of AtRALF1 and MpRALF1-3. Predicted signal
1375 peptides are in black, predicted mature peptides in light blue, conserved Cys in red, and
1376 predicted S1P recognition sites in violet.

1377 (C) Processing pathway AtRALF1 by the S1P protease.

1378 (D) A rooted neighbor-joining tree of the amino acid sequence of the S1P orthologs was
1379 generated using ClustalW. S1P members from *M. polymorpha*, *P. patens*, and *A. thaliana*
1380 were used. The numbers indicate the bootstrap values (%) from 1000 replications. The
1381 given scale represents a substitution frequency of 0.1 amino acids per site.

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1386 **Figure S5. Two LORELEI-like Proteins Are Encoded in the *M. polymorpha* Genome.**

1387 **Related to Figure 6.**

1388 (A) A rooted neighbor-joining tree of the amino acid sequence of LRE orthologs was
1389 generated using ClustalW. LRE members from *M. polymorpha*, *P. patens*, and *A. thaliana*
1390 were used. The numbers indicate the bootstrap values (%) from 1000 replications. The
1391 given scale represents a substitution frequency of 0.1 amino acids per site.

1392 (B) Amino acid sequence of AtLRE, MpLRE1, and MpLRE2. Conserved Cys are in light
1393 blue, the ND motif in red, and the GPI-anchoring site in green.

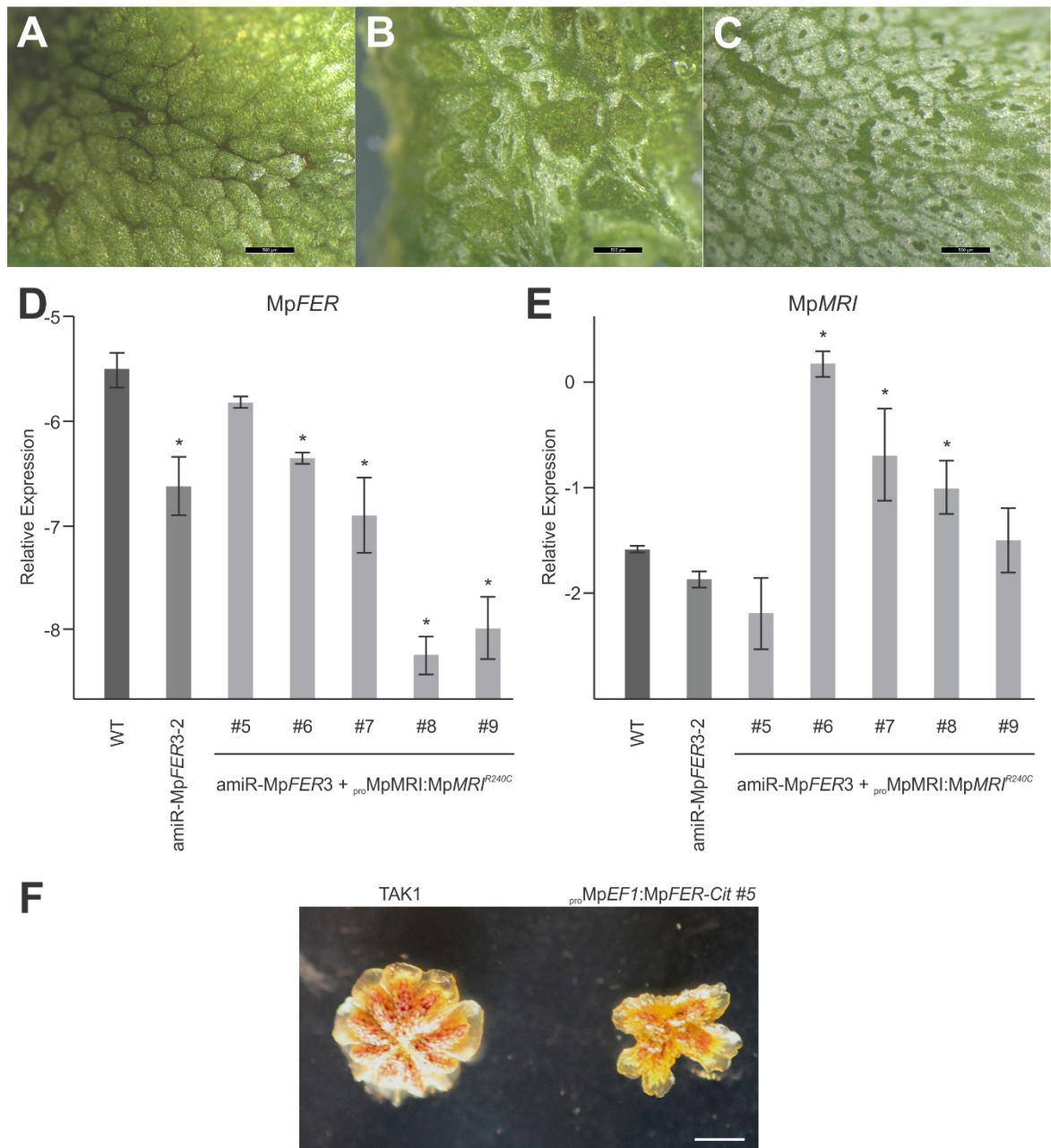
1394 (C) Cartoon representation of the predicted 3-dimensional structure of MpLRE1, showing
1395 predicted alpha-helices.

1396 (D) Structural superposition of AtLGG2 (blue) and MpLRE1 (green).

1397 (E) Cartoon representation of the predicted 3-dimensional structure of the
1398 MpFER/MpLRE1/MpRALF1 complex, showing predicted alpha-helices and beta-sheets.

1399 (F) Heatmap depicting relative gene expression based on RNAseq data (row-Z-score of
1400 vs normalized counts) of MpFER and the *M. polymorpha* orthologs of AtMRI, AtRALF1,
1401 and AtLRE across different tissues. Vegetative and reproductive tissues are grouped by
1402 green and orange, respectively. Averaged expression values are represented with colors
1403 of increasing red and blue intensity indicating upregulation and downregulation of gene
1404 expression, respectively.

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1408 **Figure S6. Expression of $MpMRI^{R240C}$ Suppresses the Bursted Rhizoid Phenotype**

1409 **of $amiR-MpFER3$ Lines but Leads to Aberrant Epidermis Development. Related to**

1410 **Figures 6 and 7.**

1411 (A to C) Epidemical pictures of thalli from the wild-type (WT, Tak-1) (A), and amiR-
1412 Mp*FER3* + *pro*MpMRI:MpMRI^{R240C} lines #6 (B) and #8 (C), which both partially suppressed
1413 the bursting rhizoid phenotype (Fig. 6).

1414 (D and E) Relative expression of Mp*FER* (D) and MpMRI (E) against the geometric mean
1415 of the reference genes Mp*ACT1*, Mp*ACT7*, and Mp*APT3* in WT, amiR-Mp*FER3-2*, and 5
1416 lines (#5 to #9) co-transformed with the amiR-Mp*FER3-2* and *pro*MpMRI:MpMRI^{R240C}
1417 constructs. Expression levels of three biological replicates were assessed by droplet
1418 digital PCR (ddPCR). The y-axis corresponds to the log₂-ratio between the test and the
1419 geometric mean of the reference genes. Shown are means ± SEMs of three biological
1420 replicates. *P < 0.01, one-way ANOVA.

1421 (F) Representative images of the antheridial receptacle of WT and *pro*Mp*EF1*:Mp*FER-Cit*
1422 plants. Scale bar, 2 mm.

1423



Figure S7. Structure Assessment and Quality Estimations of the MpFER and MpLRE 3-dimensional Models. Related to Figure 6.

QMEAN score barplot indicating quality estimates of the predicted models, across their aminoacidic sequence. Below the barplots, the corresponding secondary structures are displayed for both target and template sequences, with alpha-helices represented by light purple boxes and beta-sheets by green arrows. All plots were adapted from the model report produced by the SWISS-MODEL workspace.

(A) MpFER^{ECD}, for which AtFER was used as a template (PDB 6a5b.1A).

(B) MpLRE1, for which AtLLG2 was used as a template (PDB 6a5d.1A).