The plastoglobule-localized protein AtABC1K6 is a Mn$^{2+}$-dependent kinase necessary for timely transition to reproductive growth

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Roberto Espinoza-Corral$^{1,2,*}$ and Peter K. Lundquist$^{1,2,**}$

From the $^1$Department of Biochemistry and Molecular Biology, and $^2$Plant Resilience Institute, Michigan State University, East Lansing, Michigan, USA

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The Absence of bc$_3$ Complex (ABC1) is an ancient, atypical protein kinase family that emerged prior to the archaeal-eubacterial divergence. Loss-of-function mutants in ABC1 genes are linked to respiratory defects in microbes and humans and to compromised photosynthetic performance and stress tolerance in plants. However, demonstration of protein kinase activity remains elusive, hampering their study. Here, we investigate a homolog from Arabidopsis thaliana, AtABC1K6, and demonstrate in vitro autophosphorylation activity, which we replicate with a human ABC1 ortholog. We also show that AtABC1K6 protein kinase activity requires an atypical buffer composition, including Mn$^{2+}$ as a divalent cation cofactor and a low salt concentration. AtABC1K6 associates with plastoglobule lipid droplets of A. thaliana chloroplasts, along with five paralogs. We show that the protein kinase activity associated with isolated A. thaliana plastoglobules was inhibited at higher salt concentrations, but could accommodate Mg$^{2+}$ as well as Mn$^{2+}$, indicating salt sensitivity, but not the requirement for Mn$^{2+}$, may be a general characteristic of ABC1 proteins. Finally, loss of functional AtABC1K6 impairs the developmental transition from vegetative to reproductive growth. This phenotype was complemented by the wild-type sequence of AtABC1K6, but not by a kinase-dead point mutant in the unique Ala-triad of the ATP-binding pocket, demonstrating the physiological relevance of the protein's kinase activity. We suggest that ABC1s are bona fide protein kinases with a unique regulatory mechanism. Our results open the door to detailed functional and mechanistic studies of ABC1 proteins and plastoglobules.

Plastoglobule lipid droplets are a unique compartment of plant chloroplasts that are physically associated with the photosynthetically active thylakoid membrane (1). They hold essential but poorly understood roles related to photosynthesis, development, and stress tolerance (2-4). Despite the persistent physical association, the plastoglobule maintains a proteome and lipidome that is distinct from the thylakoid, comprising a rich store of various prenyl-lipid compounds related to photosynthesis (i.e., tocochromanols, quinones, and carotenoids) and several dozen proteins largely related to prenyl-lipid metabolism or redox reactions (5, 6). We have previously proposed that the plastoglobule serves as a dynamic hub that orchestrates rapid thylakoid membrane remodeling, sequesters and neutralizes toxic (oxidized) lipid and protein, and scavenges reactive oxygen species (5, 6). These functions are thought to be engaged routinely to regulate photosynthesis under constantly changing environmental conditions and to promote developmental transitions. This model presumes rapid functional modulation of the plastoglobule proteome; however, it remains unclear how such rapid modulation may be effected.

Prime candidates to achieve this modulation are six members of an ancient, atypical protein kinase (aPK) family known as the activity of bc$_3$ complex (ABC1) family (alternately known as COQ8, UbiB, or aarF domain-containing kinase [ADCK]) (7). ABC1s are conserved among the archaea, eubacteria, and eukaryotes where mutant phenotypes have indicated a critical role in ubiquinone biosynthesis (7-9), reflected in the various names ascribed to the family. The ABC1 atypical protein kinase family is unrelated to the ATP-binding cassette (ABC) transporter family. Mutants lacking functional ABC1 in bacteria, yeast, and human fail to accumulate appreciable amounts of ubiquinone, leading to deficiencies in cellular respiration. The yeast and human ubiquinone biosynthetic pathways employ an enzyme complex at the mitochondrial inner membrane termed Complex Q, whose stability is dependent on functional ABC1 (10). Significantly, both Arabidopsis thaliana and human orthologs were able to complement the respiratory deficiency in yeast, indicating a broad evolutionary conservation of ABC1 proteins (8, 11). While 1 to 2 ABC1 homologs are encoded in prokaryote genomes, and up to five in nonphotosynthetic eukaryotes, they have proliferated particularly in plants. Seventeen homologs are encoded in the model plant, A. thaliana, six of which localize to chloroplast plastoglobules where ubiquinone is neither biosynthesized nor accumulated. This suggests that these proteins have roles related to metabolism of other
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quinone and prenyl lipid compounds, likely including multiple compounds employed in photosynthesis (7).

In spite of the significance of the ABC1 family to organismal survival across a wide evolutionary space, it is still disputed whether they are bona fide protein kinases. With limited sequence conservation to the “classical” eukaryotic protein kinases (ePKs), the ABC1s nevertheless have several conserved key kinase motifs, such as the DFG motif responsible for divergent cation coordination and nucleotide binding, and the invariant Lys that binds and transfers the γ-phosphate. However, a number of other key motifs are missing, and multiple conserved motifs unique to ABC1s, with unclear catalytic roles, are evident (7, 12). Additionally, the Gly-rich loop, central in ATP binding, is found instead as a conserved Ala triad (AAAS sequence) in what is known as the A-rich loop.

It has been difficult to demonstrate conclusive protein kinase activity among ABC1 proteins. Protein kinase activity may be facilitated in ABC1s through unique catalytic or regulatory mechanisms that demand unique conditions. Alternatively, protein kinase activity may be precluded by the unique features of the ABC1 domain, existing instead as a so-called unorthodox kinase, i.e., pseudokinase or lipid/metabolite kinase. On the one hand, the crystal structure of a truncated sequence of the human ABC1 ortholog, HsADCK3Δ254, revealed a classical biloled protein kinase architecture but with a KxGQ domain, unique to the ABC1 family, laying over the active site and preventing access of substrate (13) (Fig. S1). No protein kinase activity was found with HsADCK3Δ254 by in vitro assay, but curiously, an A-to-G point mutant in the Ala triad (A339G) did exhibit clear protein kinase activity (13). In addition, phosphoproteomics experiments with mice either did not identify evidence of phosphorylation on ubiquinone biosynthesis proteins or did not find a statistically significant decrease of phosphorylation in the absence of the ABC1 homolog using MS/MS-based phosphoproteomics (10). Collectively, the authors concluded that endogenous HsADCK3 represents a pseudokinase capable of ATPase consumption but not protein kinase activity (10, 13, 14). On the other hand, in vitro assays with the plastoglobule-localized AtABC1K1 and AtABC1K3 provided evidence for weak trans-phosphorylation of the plastoglobule-localized substrate, tocopherol cyclase (15, 16). Our own recent work has also demonstrated indirect evidence for in vitro ABC1 kinase activity associated with isolated A. thaliana plastoglobules that is substantially reduced in an abc1k1/abc1k3 double mutant (5). Furthermore, isoelectric point shifts on several ubiquinone biosynthesis proteins in yeast, ascribed to phosphorylation, appeared in a COQ8/ABC1-dependent manner (8). These shifts could also be rescued by heterologous expression of HsADCK3. These results support the idea that plastoglobule-localized and other ABC1 proteins hold protein kinase activity. Here, we report our investigation of the plastoglobule-localized AtABC1K6. We demonstrate that with proper buffer conditions, clear protein kinase activity is exhibited as autophosphorylation and that this activity is associated with a functional role in plant development. These results provide strong support for a central regulatory role of the ABC1s on plastoglobules and present the experimental means to investigate the targets, functions, and regulation of ABC1s.

Results

Loss of AtABC1K6 impairs the transition from vegetative to reproductive growth

The A. thaliana plastoglobule harbors six members of the ABC1 protein family (6, 17–19). A coexpression network of plastoglobule genes previously demonstrated that AtABC1K6 was positioned as a network hub, suggesting a central role in the regulation of the plastoglobule (6). To investigate the function of AtABC1K6, we isolated a SALK T-DNA insertion mutant line in the sixth exon of the AtABC1K6 gene (At3g24190, hereafter referred to as abc1k6−1), which fails to accumulate detectable transcript (Fig. S2). Analysis of photosynthetic activity did not reveal a difference between wild-type (Columbia-0) and abc1k6−1 (Fig. S3). However, under standard growth chamber conditions, we observed that abc1k6−1 plants bolted and flowered several days later than the wild-type plants (Fig. 1A). The developmental rates of abc1k6−1 and wild-type were quantified according to the A. thaliana developmental stage descriptors established by Boyes et al. (20), which demonstrated that abc1k6−1 reached stage 5.10 and stage 6.0 (first flower buds and first flower opening, respectively) around 3 days later than wild-type, whether grown under long day (LD; 16/8 h photoperiod) or constant light (CL) conditions (Fig. 1B). Furthermore, under CL, mutant plants were also delayed in reaching stage 6.9 (end of senescence) compared with the wild-type. In contrast, the rate of leaf emergence and the rosette size throughout vegetative growth were indistinguishable. Furthermore, the time from bolting (stage 5.10) to first flower open (stage 6.0) was not altered in abc1k6−1. Together, this indicates that abc1k6−1 is impacted specifically in the developmental transition from vegetative to reproductive growth, rather than the rate of vegetative or reproductive development per se.

To verify that this developmental phenotype is associated with disruption of AtABC1K6, we transformed abc1k6−1 with AtABC1K6 cDNA fused C-terminally to GFP under the cauliflower mosaic virus 35S promoter (Fig. 1C). Introduction of AtABC1K6-GFP fully complemented the developmental phenotype as seen in the complementation line 3 (Comp3), which overexpressed the ABC1K6-GFP product (Fig. 1), confirming that the delayed developmental transition arises due to loss of AtABC1K6. In investigating the subchloroplast localization of the AtABC1K6-GFP fusion in complementation line 3 (Comp3), we discovered that the fusion protein was predominantly associated with thylakoid and stromal fractions when assaying with anti-GFP antibody (Fig. S4). Similarly, confocal microscopy results showed a diffuse localization of GFP fluorescence throughout chloroplasts. However, our proteomic analyses of isolated plastoglobules from wild-type, abc1k6−1, and Comp3 (see below) revealed that AtABC1K6-GFP is also prevalent in plastoglobule samples in Comp3, and that its magnitude of overaccumulation in leaf tissue matches its overaccumulation in plastoglobule samples
We speculate that the high expression level of AtABC1K6-GFP causes spillover of the heterologous protein from its natural location at the plastoglobule leading to ectopic accumulation in additional subcompartments of the chloroplast.

Plastoglobule-localized ABC1 proteins are induced during development

To test how changes to the leaf proteome may influence the developmental phenotype, we performed quantitative proteomic analyses of leaf samples from wild-type, abc1k6-1, and Comp3, at 4 weeks and 5 weeks (28 and 35 days after germination [DAG], respectively) in LD conditions. These time points represent several days before and several days after wild-type has initiated bolting (stage 5.10), providing an easily discernible developmental marker of the transition from vegetative to reproductive growth. Importantly, at the week 4 time point both genotypes are still at the same developmental stage, i.e., late vegetative growth, shortly before the wild-type will transition to reproductive growth. Hence, our comparison at week 4 can provide insight into what proteomic differences may account for the delay in transition of abc1k6-1, independent of developmental stage effects, while our inclusion of the week 5 time point can reveal how the developmental transition in wild-type effects the leaf proteome and how these changes may or may not be reproduced in abc1k6-1.

The quantitative proteomic analysis identified and quantified 4076 proteins or protein groups (hereafter referred to collectively as proteins) across all samples (Table S1). Quantification of proteins employed label-free quantification (LFQ) of MS1 peptide ion intensity as implemented in the MaxQuant proteomics software pipeline (21, 22). To determine those proteins whose abundance was effected by the mutation, we identified proteins that fit both of the following criteria: (i) a
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We explored functional categorization of the DAPs using the MapMan bin classification system (24). DAPs at week 4 were particularly enriched in chlorophyll and protein biosynthesis (Table S3). Nine proteins of chlorophyll biosynthesis, encompassing five distinct enzymatic steps, were all elevated approximately 1.75-fold in the abc1k6-1, relative to wild-type. Similarly, protein biosynthesis was elevated to a similar magnitude across a broad set of plastid and cytosolic ribosomal proteins (Fig. S5). We suggest that these patterns reflect an initial downregulation of chlorophyll and protein biosynthesis in wild-type rosette leaf tissue in anticipation of the transition to reproductive growth. At week 5, tetrapyrrole metabolism was prominently represented among the DAP proteins. However, only one of the chlorophyll biosynthesis proteins represented at week 4 was also found to be differentially accumulated at week 5. Instead, the tetrapyrrole proteins represented at week 4 was also found to be differentially accumulated at week 5. A heatmap illustrates the changes of each plastoglobule-localized ABC1 homolog in the plastoglobule proteome at week 4. Values are normalized to the highest value among the dataset (i.e., FBA-1 in wild-type). D, levels of AtABC1K6 and AtABC1K7 in the plastoglobule proteome closely track with each other across each of the genotypes. Each protein is normalized to its wild-type value. E, a heatmap illustrates the changes of each plastoglobule-localized ABC1 homolog in the plastoglobule proteome at week 4. Values are normalized to the highest value among the dataset (i.e., AtABC1K9 in wild-type). F, protein quantification suggests lipoxygenase 2 of the JA biosynthetic pathway is remobilized to the plastoglobule in abc1k6-1. Two plastoglobule-localized enzymes of jasmonic acid biosynthesis were reduced in leaf tissue of abc1k6-1 at week 4, while the lipoxygenase specifically was elevated about tenfold at the plastoglobule in the presence of over-expressed AtABC1K6 in the Comp3 line. Values are normalized to each protein’s wild-type week 4 level (total leaf samples) or to the wild-type level (plastoglobule samples). In all panels, n = 3 biological replicates; error bars indicate ±1 standard error of the mean; where error bars are not shown they are smaller than the data point. In panels, B and F, significant differences between samples are indicated with a red line and/or asterisks; one asterisk denotes p value < 0.05, two asterisks denote a p value < 0.01, three asterisks denote a p value < 0.001.

As expected, AtABC1K6 could be readily identified and quantified in wild-type and Comp3 genotypes, but not in abc1k6-1 (Table S1). From the proteomics results, we could estimate that AtABC1K6 accumulation in leaf tissue was about 16-fold higher than wild-type levels at 4 weeks and about threefold higher at 5 weeks (Fig. 2A). The greater differential at 4 weeks was due to a tripling of endogenous AtABC1K6 levels in wild-type from week 4 to week 5. Interestingly, we found that the other plastoglobule-localized ABC1 homologs also were significantly increased from week 4 to week 5, in particular AtABC1K9 whose accumulation increased over 25-fold (Fig. 2B). We also looked for the accumulation patterns of proteins with well-known roles in flowering (e.g., Flowering Locus T, Flowering Locus Y, Suppressor of Overexpression of Constans 1, and Leafy). However, these proteins were not identified in our dataset, presumably because of low abundance within the leaf proteomes that prevented their detection with the LC-MS/MS.

significantly different normalized LFQ (nLFQ) in abc1k6-1 relative to wild-type, [p value < 0.05, controlling for multiple hypotheses using the Benjamini–Hochberg method (23)]; and (ii) an nLFQ restored at least half-way to wild-type levels in Comp3. With this in mind, a total of 313 differentially accumulated proteins (DAPs) were affected at 4 weeks and 379 proteins at 5 weeks. Surprisingly, only 56 of these DAPs were affected at both time points, indicating that effects on the proteome are highly dependent on developmental stage.

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Significantly, three of the enzymatic steps of jasmonic acid (JA) biosynthesis were significantly reduced in abc1k6-1 at week 4, but not week 5 (Table 1). JA signaling can promote
chlorophyll degradation and senescence, and the plastid-localized steps have been found to associate with plastoglobules under specific conditions (4, 25–31). We considered whether a possible deficiency in JA synthesis, and hence signaling, at week 4 may account for the developmental phenotype in abc1k6-1. To test this possibility, we treated wild-type and abc1k6-1 plants with 25 μM of Me-JA as a foliar spray occurring each day for 7 days from 21 to 28 DAG. While leaves of the treated samples accumulated anthocyanins, indicating perception of JA, the time at which each genotype reached bolting was unchanged relative to mock-treated plants, indicating that the JA treatment failed to complement the developmental phenotype (Fig. S6).

Quantitative proteomics reveals specific changes to the plastoglobule proteome

Comparison of protein localizations among the total leaf proteomes revealed that the sum of plastoglobule protein levels was reduced by about 20% relative to wild-type, which cannot be accounted for solely by the loss of AtABC1K6 (Table S2). To identify specific changes to the plastoglobule core proteome associated with loss of AtABC1K6, we compared the proteomes of isolated plastoglobules from wild-type, abc1k6-1 and Comp3 at week 4 (6). Ten proteins of the plastoglobule core proteome were affected by loss of AtABC1K6, according to the same criteria used with leaf samples (Fig. 2C and Table S4). Only a single tryptic peptide of AtABC1K6 was identified in abc1k6-1 plastoglobules, which came from near the N-terminus (residues 175–183), upstream of the T-DNA insertion site. This suggests that truncated AtABC1K6 accumulates at low levels and associates with the plastoglobule in abc1k6-1. About 14-fold more AtABC1K6 protein was associated with plastoglobules in Comp3 than wild-type, consistent with the enrichment of AtABC1K6 seen at the leaf tissue level (Fig. 2A). Strikingly, nLFQ levels of AtABC1K7, but not the other plastoglobule-localized ABC1 homologs, closely tracked with that of AtABC1K6 over the genotypes; no AtABC1K7 was found in isolated plastoglobules of abc1k6-1, while AtABC1K7 associated with plastoglobules to nearly 50-fold higher levels in Comp3 than in wild-type (Fig. 2, D and E). This suggests that AtABC1K6 may stabilize AtABC1K7 or be necessary for its association with plastoglobules. Lipoxygenase 2 (AtLOX2), involved in JA biosynthesis, and Plastoglobular Metalloprotease M48 (AtPGM48), a positive regulator of leaf senescence, also were more abundant in plastoglobule samples of Comp3 (Fig. 2C). The tenfold enrichment of AtLOX2 at the Comp3 plastoglobule was not reflected at the leaf level where total levels of AtLOX2 were the same at the week 4 time point when plastoglobules were isolated (Fig. 2F). This is indicative of a recruitment of AtLOX2 to the plastoglobule upon overaccumulation of AtABC1K6-GFP, reminiscent of recruitment of this and other JA biosynthesis enzymes to the plastoglobule in other conditions (4, 5). In contrast, evidence did not support recruitment of AtAOS and AtPGM48 to the plastoglobule, since AtAOS levels on the plastoglobule were comparable in all genotypes, while AtPGM48 levels were below the detection limit in leaf samples (Fig. 2F). Curiously, three proteins were conspicuously absent from plastoglobules of Comp3, despite their prominent accumulation in plastoglobule samples of wild-type and abc1k6-1: (i) a SOUL heme-binding protein (AtHBP3), (ii) the Plastoglobule protein of 18 kDa (AtPG18), and (iii) an uncharacterized protein called Unknown 2 (Fig. 2C). The elevated levels of AtPGM48 metalloprotease, in tandem with the specific loss of the above three proteins, present these three proteins as candidate substrates for proteolytic cleavage by AtPGM48. None of the three proteins have known functions, although it seems unlikely that their absence from plastoglobules is related to the phenotypic complementation, since they are prominent components of the wild-type plastoglobules.

AtABC1K6 is a protein kinase that requires low salt and Mn$^{2+}$ as cofactor

Multiple conserved residues and kinase motifs shared with canonical ePKs suggest that the ABC1 family encodes protein or lipid kinase activity; however, clear demonstration of any kinase activity remains elusive (13, 14). To test for protein kinase activity of AtABC1K6, we heterologously expressed the mature AtABC1K6 protein (i.e., lacking only the chloroplast transit peptide; residues 1–19) in Escherichia coli, fused at the N-terminus to the maltose-binding protein (MBP), containing a Factor Xa protease cleavage site, and at the C-terminus to a 6xHis tag (MBP-AtABC1K6-His; Fig. 3C). Inclusion of the MBP tag was necessary to support solubility of the expressed protein. Purified MBP-AtABC1K6-His was incubated in vitro kinase buffer with γ-32P-labelled ATP for 30 min and separated with SDS-PAGE. No in vitro kinase activity could be detected in the presence of 150 mM NaCl (data not shown). However, in buffer lacking NaCl, in vitro kinase activity was readily apparent from a spontaneous cleavage product of the recombinant protein (Fig. 3A). This cleavage product appeared during the purification despite our exclusion of Factor Xa or any other protease and also was seen when purified using a wheat germ expression system (Fig. S7). Furthermore, heterologous expression of two other plastoglobule-localized ABC1 proteins from E. coli also displayed spontaneous cleavage,

### Table 1

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$^a$ The three proteins depleted in abc1k6-1 are all complemented (at least half restored to wild-type levels) in the Comp3 line, indicating that their change in abundance is due to loss of AtABC1K6.

## References

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Figure 3. AttABC1K6 displays in vitro kinase activity dependent on Mn2+. A, AtABC1K6 protein fused to MBP was expressed in E. coli and purified to perform in vitro autophosphorylation assays using γ-32P-ATP. Purified protein samples were incubated with γ-32P-ATP for 30 min and gel was exposed to autoradiography film for 7 days. On the left panel is shown the autophosphorylation activity of the heterologously expressed protein, while MBP alone was used as a negative control. A spontaneously occurring, truncated form of MBP-AtABC1K6-His migrates at ca. 75 to 80 kD, is indicated as ‘AtABC1K6Δ71-His’, and displays in vitro kinase activity. Two microgram of protein were loaded in each lane. The Coomassie stained band (CBB) is indicated as a loading control.

B The purified MBP-AtABC1K6-His was assayed for in-vitro kinase activity with increasing concentrations of Mn2+ in buffer containing 20 mM Tris (pH 7.5) and no NaCl. As a negative control for kinase activity, 20 mM EDTA was added to the assay with 20 mM Mn2+. Sample was incubated with γ-32P-ATP for 45 min and the gel was exposed to autoradiography film for 3 days. The two spliced panels are from the same image of the same film/gel. The values in red indicate the relative intensity of radiography at the AtABC1K6Δ71-His band, quantified using Image J, and normalized to the 20 mM Mn2+ lane. Five microgram of protein were loaded in each lane. C, the identity of the spontaneous cleavage product, and location of the spontaneous cleavage, was made by exciting bands from an SDS-PAGE gel and analyzing by LC-MS/MS, as described in the main text and illustrated in Fig. S6. A schematic illustration of the MBP-AtABC1K6-His recombinant protein, and the locations of the encoded Factor Xa cleavage site and the spontaneous cleavage site occurring during expression or purification is given.

Kinase activity is associated with isolated plastoglobules

We have previously demonstrated in vitro protein kinase activity with isolated plastoglobules that target multiple substrate proteins of varying molecular weights (5). In order to understand the cation dependency in the context of purified plastoglobules, we tested in vitro kinase activity with wild-type plastoglobules under different cation concentrations. Interestingly, although 20 mM Mn2+ showed detectable kinase activity, a stronger signal was obtained under 20 mM of Mg2+, while no kinase activity was observed under 20 mM of Ca2+ (Fig. 4A). We then performed in vitro kinase assays using isolated wild-type plastoglobules in kinase buffer with gradients of NaCl and KCl. Consistent with results from purified AtABC1K6, kinase activity associated with plastoglobules was sensitive to both salts, preferring low salt concentrations of less than 50 mM (Fig. 4B). Importantly, the proteomic data of our isolated plastoglobules, described above, reveal that all six expected ABC1 proteins are present, but no other known or putative protein kinases are found (Table S4). This strongly whereas expression of two unrelated plastoglobule-localized proteins did not display evidence of spontaneous cleavage (Fig. S7). We conclude that the spontaneous cleavage during heterologous expression is a characteristic of the plastoglobule-localized ABC1 proteins, but not of plastoglobule proteins in general.

Mass-spectrometry-based proteomic analysis of the cleaved protein fragment from MBP-AtABC1K6-His, separated by SDS-PAGE, identified the cleavage site after K443 (counting from the first residue of the MBP-AtABC1K6-His recombinant protein) (Fig. 3C). This indicates the cleavage product receiving γ-32P contains AtABC1K6 protein minus the 71 N-terminal residues, which eliminates the predicted intrinsically disordered region (Fig. S8). This conclusion is consistent with the immunoblotting data that indicates that the cleavage product retains the 6xHis tag but not the MBP tag, and its migration at ca. 75 kD is consistent with the theoretical molecular weight of the putative protein fragment, calculated at 80.7 kD. Hence, we refer to this kinase-active cleavage product as AtABC1K6Δ71-His for the remainder of this article. Importantly, full-length MBP-AtABC1K6-His migrating at ca. 130 kD did not display kinase activity, suggesting that either the presence of the MBP tag or the disordered region inhibits the kinase activity in the in vitro system.

To closely mimic the in vivo stromal environment of AtABC1K6, we initially used 5 mM Mg2+ and 0.5 mM Mn2+ in our in vitro kinase buffer, concentrations matching those estimated in chloroplast stroma during photosynthesis (32–34). We then tested whether the kinase activity depended on the Mg2+ or Mn2+ by providing only one of the two cations. Assays revealed dose-dependent activity in the presence of Mn2+; however, no kinase activity could be detected even at the highest tested concentration of Mg2+ (Figs. 3B and S9). We conclude that a truncated form of AtABC1K6, lacking a predicted disordered region at its N-terminus, encodes protein kinase activity that employs Mn2+ as cofactor and is regulated by ionic strength of the buffer.
indicates that all kinase activity identified with the isolated plastoglobules derives from one or more of the associated ABC1 proteins.

To determine if any of the detectable protein kinase activity from isolated plastoglobules might derive from AtABC1K6, we compared protein kinase activity of plastoglobules and chloroplasts isolated from wild-type and abc1k6-1. However, band patterns were essentially unchanged between the two genotypes, indicating that AtABC1K6 does not account for the detectable protein kinase activity in our in vitro kinase assays (Fig. 4C). Thus, the salt sensitivity appears to be a general phenomenon among plastoglobule-localized ABC1 paralogs, while the requirement for Mn2+ as divalent cation cofactor is not absolute across all plastoglobule-localized ABC1 homologs.

In vitro kinase activity and in vivo functionality of AtABC1K6 are dependent on the conserved Ala triad of the ATP-binding pocket

A unique feature of the ABC1 protein kinase domain is the substitution of a highly conserved Ala triad in the ATP-binding pocket, in place of the canonical GxGxxG motif (the Gly loop, where x indicates any residue) seen in ePKs (Fig. 5B) (7, 12, 35). We evaluated the role of the Ala triad in AtABC1K6 with in vitro kinase assays using point mutants in the triad. As with the wild-type recombinant protein, point mutants were spontaneously cleaved during purification and migrated at the same apparent molecular weight. Strikingly, the A268G mutation (mutated in the first Ala of the triad, equivalent to the A339G mutation of HsADCK3Δ254) showed severely diminished kinase activity (Fig. 5A). Furthermore, point mutants of the next two Ala residues (A269G and A270G) and the triple A-G point mutant also showed severely diminished kinase activity.

To test if the in vitro kinase activity of AtABC1K6 is relevant to in vivo function, we attempted complementation of abc1k6-1 with the A268G point mutant of AtABC1K6. To mimic the conditions of the Comp3 line that fully complemented the developmental phenotype, we retained the C-terminal GFP tag and 35S promoter sequences. Despite strong overexpression of the A268G point mutant in the abc1k6-1 background, no complementation of the developmental phenotype was apparent (Fig. 5, C–E), contrasting with the full complementation seen when using the wild-type sequence in Comp3 (Fig. 1). This indicates that the in vitro kinase activity detected with AtABC1K6Δ71-His recombinant protein is necessary for its in vivo function related to the developmental transition.

The human ABC1 ortholog, HsADCK3Δ254, also exhibits protein kinase activity but with differing requirements

In light of the distinct requirements for protein kinase activity that we found with AtABC1K6, we considered whether previous efforts may not have detected kinase activity with HsADCK3Δ254 due to the chosen buffer conditions. Thus, we tested HsADCK3Δ254 for divalent cation dependency and salt
sensitivity. As for AtABC1K6, HsADCK3ΔΔ254 exhibited protein kinase activity when Mn+2 was provided as divalent cation (Fig. 6A). However, activity was also clearly seen when providing solely Mg+2, albeit not as strong as with Mn2+. This indicates that HsADCK3ΔΔ254 is more active with Mn+2 but can also accommodate Mg2+ as divalent cation cofactor, contrasting with the apparent absolute requirement for Mn2+ seen with AtABC1K6.

When testing salt sensitivity in the presence of both 5 mM Mg2+ and 0.5 mM Mn2+ the recombinant HsADCK3ΔΔ254 surprisingly displayed clear in vitro kinase activity at both 0 and 150 mM NaCl, whether the MBP tag is

Figure 5. The highly conserved Ala triad of the ABC1K kinase domain is essential for in vitro kinase activity and in vivo function of AtABC1K6. A, in vitro kinase assays with the wild-type sequence, and various point mutants of the Ala triad, of MBP-AtABC1K6-His are presented. The values in red indicate the relative intensity of radiography at the AtABC1K6NΔ71-His band, quantified using Image J, and normalized to the wild-type lane. Three microgram of protein was loaded in each lane. Sample was incubated for 30 min with γ-P32-ATP and the gel was exposed to autoradiography film for 1 week. B, a sequence alignment of several ABC1 homologs, along with two canonical protein kinases, AtSTN7 and MmPKA from A. thaliana and Mus musculus, respectively. The alignment is intended to illustrate the sequence context of the point mutants tested in (A) and illustrate the conservation of the unique ATP-binding pocket of the ABC1 family. AtABC1K6, AtABC1K1, and AtABC1K3 are plastoglobule-localized ABC1 homologs from A. thaliana, HsADCK3 is an ABC1 homolog from mitochondria of Homo sapiens, AtSTN7 is State Transition 7 kinase of A. thaliana thylakoid membranes, and MmPKA is Protein Kinase Alpha of M. musculus. C, an anti-GFP immunoblot demonstrates expression of the AtABC1K6-GFP in various lines stably transformed with either the wild-type (Comp3), or A268G point mutant (Comp6 & Comp7) variant of AtABC1K6. Anti-PsaF, detecting a membrane protein of the photosystem II complex, was used as a loading control. Values in red represent the relative ratio of GFP intensity normalized by PsaF. D, transformation with the point mutants fails to complement the developmental phenotype, in contrast to transformation with the wild-type sequence. Schematic illustrates the developmental stage transitions of each genotype following the nomenclature described by Boyes et al., 2001 (20), given as the average time from ten individual plants. Arrows indicate the time in which wild-type plants reached the respective developmental stage; 1.04: four rosette leaves; 1.10: ten rosette leaves; 5.10: first flower buds visible; 6.00: first flower open; 9.70: senescence complete. Asterisks indicate a statistically significant difference in reaching a given developmental stage compared to wild-type. n = 10; Student's t test with p value less than 0.05. E, representative photos of genotypes demonstrating the lack of complementation seen in the lines transformed with the A268G point mutant.
retained or cleaved off (Fig. 6B). This contrasts with previous results from Stefely et al. that did not identify in vitro kinase activity with HsADCK3\(\Delta\)254 at 150 mM NaCl and 20 mM Mg\(^{2+}\) (13, 14).

Both the wild-type and A339G point mutant of HsADCK3\(\Delta\)254 displayed strong kinase activity, indicating that the HsADCK3\(\Delta\)254 ortholog does not share the delicate sensitivity of the Ala triad seen in the AtABC1K6 under our in vitro kinase conditions (Fig. 6A). These results indicate possible structural differences in the ATP-binding pocket between the A. thaliana and human orthologs that may also account for the differential sensitivity to salt concentration shown above. To investigate this possibility, we compared the crystal structure of HsADCK3\(\Delta\)254 (PDB: 5i35) with a modeled structure of AtABC1K6 generated using Alphafold (ID: AF-Q9LRN0-F1), which generates models de novo, i.e.,
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independent of any homology template (36). Threedimensional alignment of the kinase domain structures from AtABC1K6 and HsADCK3 revealed considerable architectural conservation with a root-mean-square deviation (rmsd) of 2.868 Å for the atomic positions of 1305 atoms from the aligned residue pairs (37). Both structures revealed that the Ala-triad forms the last strand of a β-sheet, which cups the ATP analog (Fig. 6C). The similar structural features of the ATP-binding pocket preclude an explanation for the differing effects of the A-to-G mutations. However, the adjacent conserved Ser is positioned to form a hydrogen bond with the β-phosphate of the ATP (Fig. 6D). Hence, the β-sheet, comprised of the Ala-triad, is important for the positioning of the Ser residue and stability of the ATP. We suggest that the provision of Gly within this triad introduces an enhanced flexibility, which destabilizes the strand and removes a part of the ATP-binding pocket explaining the loss of kinase activity in the Ala-triad point mutants of AtABC1K6.

Discussion

The study of the ABC1 family has been hampered by the previous inability to conclusively demonstrate catalytic function. Here, we report that an A. thaliana ortholog, AtABC1K6, possesses in vitro protein kinase activity requiring two modifications from standard in vitro assay buffers: (i) provision of Mn2+ rather than Mg2+, and (ii) low salt concentration. The requirement for Mn2+ suggests that it, rather than Mg2+, is used as divalent cation cofactor. There is precedent for Mn2+-dependency among protein kinases, including a two-component sensor kinase of the chloroplast stroma (38). The effect of removing NaCl from the assay buffer, on the other hand, is more surprising. In vitro kinase data from the literature corroborate an apparent salt sensitivity of ABC1 protein kinase activity since the report of weak kinase activity from AtABC1K1 and AtABC1K3 used 50 mM NaCl (15, 16). This intermediate level of activity is consistent with the gradient of activity we report with our isolated plastoglobules (Fig. 4A).

Complete removal of buffer salt cannot be physiologically relevant to ABC1s, since the chloroplast stroma is thought to contain >100 mM KCl (39). However, the stroma can modulate ionic strength, primarily through transport of K+, which may impact ABC1 activity (40, 41). Further investigations will be necessary to determine whether physiological changes in stromal ionic strength can impact ABC1 activity in vivo. Regardless, we suggest that in manipulating ionic strength of the buffer, we have identified a way to artificially induce regulatory rearrangements of the kinase structure that are a part of the natural regulatory cycle of AtABC1K6, leading to activation of kinase activity. All protein kinases are characterized by a spatially conserved arrangement of hydrophobic residues that create a pair of “spines” providing a strong but flexible scaffold (42–44). Rearrangements of these hydrophobic interactions are central to the structural dynamics that regulate activity, but are complex and specific to each kinase family. Reduced ionic strength in our buffer may influence the hydrophobic interactions within the AtABC1K6 spines, leading to activation of protein kinase activity.

Some protein kinases have been shown to hold dual kinase/phosphatase activity within the same catalytic domain (45–49). In the case of AceK from E. coli, the bifunctional kinase-phosphatase activity can be fine-tuned by swapping Mn2+ and Mg2+ as divalent metal (48, 49). It is worth considering whether the ABC1 proteins may also hold dual kinase/phosphatase activity. In our hands, protein kinase activity with the heterologously expressed protein is weaker than many canonical eukaryotic protein kinases. This could be accounted for if competing kinase and phosphatase activity exists within the expressed protein. Our modification of divalent metal and/or salt concentration may tip the balance of such competing activities to favor protein phosphorylation leading to our discovery of protein kinase activity with AtABC1K6. The possibility of dual activity within ABC1 proteins would also nicely explain the conspicuous absence of phosphatases at the plastoglobule. At this point, it is not clear whether AtABC1K6, or any other ABC1 protein, may hold phosphatase activity, and further investigation is necessary to support or refute this prospect.

Our results with the isolated plastoglobules also indicate that a proper membrane environment might support protein kinase activity. Importantly, our proteomic analysis of the same isolated plastoglobule samples that were used for the in vitro kinase assays demonstrated that all six ABC1 proteins— but no other known or putative protein kinases— could be detected. This strongly indicates that detected activity derives from one or more of the ABC1 proteins. While our test of abc1k6-1 plastoglobules did not indicate that detectable activity arises from AtABC1K6, we have previously identified a substantial reduction in activity using isolated plastoglobule from an abc1k1/abc1k3 double mutant (5). This indicates that at least one of these two plastoglobule-localized ABC1s generates a significant proportion of the in vitro kinase activity we could detect. Furthermore, the substrate proteins receiving γ-P32 migrate at molecular weights considerably lower than the ca. 75 kD size of the ABC1 proteins, indicating the detectable in vitro protein kinase activity represents “trans” phosphorylation. The use of isolated plastoglobules may promote an active kinase conformation, for example, through specific lipid binding. However, this possibility awaits further investigation.

An insert motif within the N-lobe called the KxGQ domain, unique to the ABC1 family, tightly packs over the active site cleft via hydrophobic interactions, occluding access of substrate (13). The placement of the KxGQ domain is comparable to the inactive conformations of ePKs, in which an Activation Loop is also folded over the active site cleft, preventing access of substrate (50, 51). In fact, two of the key distinguishing features of ABC1 proteins are their absence of an Activation Loop and the presence of the KxGQ domain. We thus propose that the KxGQ domain of ABC1 proteins is functionally analogous to the mobile Activation Loop of canonical ePKs, serving as an autoregulatory feature that can expose or obscure the active site cleft to activate/inactivate the kinase. Moreover, the presence of the conserved Ala-triad adjacent to the ATP-binding site, and the impact of mutating to Gly, highlights
the importance of this adjoining region. In this work, we show that such mutations greatly diminish kinase activity, which we suggest is due to a destabilization and unfolding of the β-sheet forming the ATP-binding pocket. However, further experimental and computational support is required to fully explain this effect on kinase activity of AtABC1K6.

Our results with HsADCK3Na254 present a puzzle: why are the requirements for enzymatic activity that we demonstrate with AtABC1K6—the salt sensitivity, the absolute dependency for Mn2+, or the significance of the intact Ala triad—not seen with HsADCK3Na254? The answer may lie in the extension sequence(s) flanking the conserved ABC1 kinase domain, which is removed on the N-terminal end from the assays with HsADCK3Na254 but left mostly intact with that of AtABC1K6Na71. In contrast to the highly conserved ABC1 domain, the extension sequences are highly variable in length and sequence and often encode predicted lipid or protein-binding motifs. Thus, the extensions may define the distinct functional identity of each ABC1 and regulate autoinhibition that can be relieved only by a proper trigger. In this scenario, the absence of the N-terminal extension of HsADCK3Na254 may have weakened its autoinhibitory capacity. Indeed, the crystal structure of the protein (PDB: 5I35) reveals multiple hallmarks of an active conformation, including DFGin, forming an Asp-His salt bridge, and aC-helixin, forming a Lys-Glu salt bridge (50, 51), indicating that the HsADCK3Na254 structure is primed for activation. Alternatively, unique catalytic or regulatory mechanisms may have emerged in each of the two proteins, reflecting their distinct functional roles and subcellular localizations.

Plastoglobule morphology is very dynamic during developmental transitions, which has suggested that they may play a role in mediating development (17, 52–55). Our demonstration that the plastoglobule-localized AtABC1K6 is necessary for the timely transition from vegetative to reproductive growth supports this notion and identifies a putative regulatory player. Although our assay with abc1k6-1 plastoglobules and chloroplasts did not detect appreciable activity coming from the AtABC1K6, the protein may not be active at the developmental stage or condition from which our plastoglobules were isolated, or the AtABC1K6 activity is dwarfed by that of other plastoglobule ABC1 kinases. It is curious that, with 17 homologs in A. thaliana and six at the plastoglobule alone, single gene knockouts result in clear developmental or stress-related phenotypes, as seen in abc1k6, or abc1k1/abc1k3 mutants, respectively (3, 4, 15, 16, 56). This indicates a limited degree of functional redundancy that presumably reflects unique target substrates, possibly dictated by the unique flanking sequences mentioned above. It remains unclear what substrates AtABC1K6 may target in affecting initiation of reproductive growth, but the biosynthetic enzymes of JA biosynthesis that partially localize to the plastoglobule (4, 5, 18, 19), particularly AtLOX2, which our results suggest is remobilized to the plastoglobule in abc1k6-1, present an intriguing possibility. The phytohormone JA is a positive regulator of developmental senescence and tightly intertwined with the early steps in the switch to reproductive growth (28, 29, 57–63).

In sum, by modifying buffer conditions, we have been able to demonstrate in vitro protein kinase activity with the plastoglobule-localized AtABC1K6. Our discovery of proper buffer conditions to facilitate protein kinase activity among AtABC1K6 and HsADCK3 presents the experimental means to explore the substrates, activity, and regulation of this protein family, opening up a long-awaited opportunity to understand the functions of this enigmatic protein family.

**Experimental procedures**

**Plant material and growth conditions**

The T-DNA insertion line SALK_129956C (abc1k6-1, At3g24190) was obtained from the Salk collection, and homozygous mutants were screened using the following oligonucleotides: K6F: 5’-ATCGGCTATTGCAGGGGAT-3’, K6R: 5’-CGAGTTTAACAAAACTTTTACTAG-3’ and LBb1 5’-GCGTGAGACCCTTGCTGCAACT-3’. Additionally, oligonucleotides for OE16 (Atg2g8900) were used as positive control for PCR reactions: OE16 F 5’-TGTAGACGCCC CGAAG-3’ and OE16 R 5’-CTTACAAACCGCTGAG-3’. Homozygous mutant plants were isolated using antibiotic selection on Murashige and Skoog agar media supplemented with kanamycin (50 μg/ml). For complementation, the coding sequence of AtABC1K6 was cloned into the vector pB7FWG2 under the control of the 35S promoter (64). The construct was introduced into Agrobacterium tumefaciens strain GV3101 and abc1k6-1 mutants were transformed by floral dip (65). The T1 progeny were selected on soil by spraying BASTA herbicide 1 and 2 weeks after sowing. Homozygous plants for the transgene were selected based on their resistance to BASTA of the T3 progeny, which came from a T2 progeny with 75% of resistance against the herbicide to ensure a single insertion event. If not indicated otherwise, A. thaliana plants were grown on soil under normal light conditions (LD, 16/8 h light/dark, 21 °C, and 120 μE m⁻² s⁻¹). Continuous light was applied using the same conditions but with a 24-h light period.

**Measurement of photosynthetic parameters**

*A. thaliana* wild-type and mutant plants grown under long day conditions for 3 weeks were dark adapted for 15 min and analyzed according to their chlorophyll fluorescence as described in Espinoza-Corral et al. (2) using a pulse-modulated fluorometer (Imaging PAM and DUAL-PAM100; Walz).

**SDS-PAGE and immunoblot analyses**

Total leaf or chloroplast subcompartment fractions were separated on 12% SDS–polyacrylamide gels (66) and transferred to a nitrocellulose membrane (Amersham, Protran). The membrane was then incubated with monospecific polyclonal antisera (anti-FBN1a; AS06 116, anti-PsBA; AS05 084, anti-PsaF; AS06 104, anti-SBPase; AS15 2973, anti-LHC1a; AS01 005, anti-PsiA; AS09 461, anti-Cyt f; AS08 306, anti-PoA; AS11 1783 [Agrisera], anti-GFP; PA1 980A [Invitrogen], anti-His; TA150087 [OriGene]), followed by incubation with secondary polyclonal antisera HRP (Fisher scientific,
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were pelleted with 5-min centrifugation at 1500

glycerophosphate, 1 mM Na-OrthoVanadate, and 10 mM Na-

inhibitors cocktail containing 50 mM NaF, 25 mM

substrate and phosphatases inhibitors cocktail (P9599, Sigma)

and incubated for 30 min on

chloroplasts with subsequent separation of their subcompart-

ments as previously described in (5). Plant rosettes were

ground with isolation buffer (13 mM Tris-HCl, 20 mM MOPS

phosphate, 1 mM EDTA, 0.6 M sucrose, and a cocktail of phospha-

tases and proteases inhibitors containing 74

and 20% wattage from a Fisher Scienti-

c). After bacterial transformation, the

mature form of AtABC1K6 was

terminus of the protein an additional His-tag was added by

these constructs were used to transform BL21 pLys

cells, which were induced for overexpression with 1 mM IPTG

when they reached an OD_{600} of 0.6 to 0.8. The overexpression

was performed overnight at 12 °C. After cell lysis by sonication

buffer containing 20 mM Tris-HCl pH 7.5 and 15 mM imidazole,

the solution was centrifuged at 20,000g and 4 °C for

30 min. The supernatant was transferred to equilibrated Ni^{2+}–

NTA beads (Thermo scientific) in 20 mM Tris-HCl pH 7.5 and

15 mM imidazole and incubated under rotation for 2 h at 4 °C.

Beads were pelleted by centrifugation at 100g and 4 °C for

2 min and then washed with buffer 20 mM Tris-HCl pH 7.5

and 15 mM imidazole for four times. The recombining proteins

were eluted by washing the beads with 20 mM Tris-HCl

pH 7.5 and 500 mM imidazole. Eluted proteins were then
dialyzed over night against kinase buffer (20 mM Tris-HCl pH

7.5, 5 mM MgCl2, and 0.5 mM MnCl2) and quantified by using

Pierce BCA Protein Assay Kit (23227, Thermo scientific).

The in case of protease digestions, Factor Xa (P8010S, New

England BioLabs) or TEV protease (P8112S, New England Bio-

Labs) was used for the cleavage of constructs in the backbone

of pMAL-c5x and for His-MBP-[TEV]-ADCK3. The diges-
tions were performed for 1 h under room temperature for

Factor Xa with the addition of 2 mM CaCl2 or 1 h under 37 °C

for TEV protease with the addition of 0.1 mM DTT.

Generation of point mutants

The constructs containing the AtABC1K6 gene for over-

expression in E. coli using the vector pMAL5-x, as well as in

A. thaliana using the vector pB7FWG2, were used as templates

for the generation of point mutants. Point mutant variants were

generated by inverse-PCR using the primers indicated in

Table S5. The inverse-PCR reaction was cleaned up and the

template plasmid digested with DpnI for 1 h at 37 °C. After

inactivating DpnI at 80 °C for 10 min, the inverse-PCR product

was ligated by adding T4 DNA ligase and T4 polynucleotide

kinase (Thermo scientific). After bacterial transformation,

the point mutants were confirmed by sequencing.

In vitro kinase assays

Heterologously expressed and purified recombinant protein

or isolated plastoglobules were incubated with 3 μCi of [γ-^{32}P]

ATP in 40 μl kinase reaction buffer (20 mM Tris-HCl pH 7.5,

5 mM MgCl2, and 0.5 mM MnCl2). The reaction was incubated

for 30 min to 1 h (as described in the figure legend) at room

temperature and stopped by adding 5 μl of SDS loading buffer

without boiling the samples. The proteins were separated on

12% SDS–polyacrylamide gels followed by exposure to autora-
diography films from overnight to up to 1 week depending on the

signal intensity, as indicated in the figure legend.

LC-MS/MS proteomics and data analysis of heterologous

MBP-AtABC1K6-His protein

To delineate the boundaries of the MBP-AtABC1K6-His

cleavage products, expressed and purified protein was
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resuspended in SDS-PAGE sample buffer and heated at 60 °C for 10 min. The sample was cooled and loaded onto a 12.5% pre-cast BioRad Criterion 1D gel and electrophoresed until the dye front began to run off the bottom of the gel. Electrophoresis was stopped and the gel fixed in 40% Methanol/20% Acetic Acid for 4 h and stained with colloidal Coomassie Blue overnight. The gel was then destained using 10% Acetic Acid until the background was clear. Gel bands were digested in-gel according to Shevchenko et al. (68) with modifications. Briefly, gel bands were dehydrated using 100% acetonitrile and incubated with 10 mM dithiothreitol in 100 mM ammonium bicarbonate, pH ~8, at 56 °C for 45 min, dehydrated again, and incubated in the dark with 50 mM iodoacetamide in 100 mM ammonium bicarbonate for 20 min. Gel bands were then washed with ammonium bicarbonate and dehydrated again. Sequencing grade modified trypsin was prepared to 0.01 μg/μl in 50 mM ammonium bicarbonate and ~100 μl was added to each gel band so that the gel was completely submerged. Bands were then incubated at 37 °C overnight. Peptides were extracted from the gel by water bath sonication in a solution of 60% Acetonitrile/1% Trifluoroacetic acid and vacuum dried to ~2 μl. An injection of 5 μl was automatically made using a Thermo (www.thermo.com) EASYnLC 1000 onto a Thermo Acclaim PepMap RSLC 0.1 mm × 20 mm C18 trapping column and washed for ~5 min with Buffer A. Bound peptides were then eluted over 35 min onto a Thermo Acclaim PepMap RSLC 0.075 mm × 250 mm resolving column with a gradient of 5% B to 40% B in 24 min, ramping to 90% B at 25 min and held at 90% B for the duration of the run (Buffer A = 99.9% Water/0.1% Formic Acid, Buffer B = 80% Acetonitrile/0.1% Formic Acid/19.9% Water) at a constant flow rate of 300 nl/min. Column temperature was maintained at a constant temperature of 50 °C using an integrated column oven (PRSOV1, Sonation GmbH). Eluted peptides were then sprayed into a ThermoScientific Q-Exactive mass spectrometer using a FlexSpray spray ion source. Survey scans were taken in the Orbitrap (70,000 resolution, determined at m/z 200), and the top 15 ions in each survey scan were then subjected to automatic higher-energy collision-induced dissociation (HCD) with fragment spectra acquired at 17,500 resolution. The resulting MS/MS spectra were processed with the MaxQuant software program, version 1.6.11.0 (69). Peak lists were searched with the embedded Andromeda search engine (70) against the E. coli UNIPROT database concatenated with the MBP-AtABC1K6-His heterologous protein sequence plus the common contaminant list appended by MaxQuant. Oxidation of methionine, deamidation of asparagine and glutamine, and N-terminal acetylation were set as variable modifications, carbamidomethylation was set as a fixed modification. Digestion mode was Trypsin/P with a maximum of two missed cleavages. MS/MS tolerance of the first search was 20 ppm, and main search was 4.5 ppm, with individualized peptide mass tolerance selected. False discovery rate (FDR) at peptide spectrum match and protein levels was set as 0.01. Filtering of resulting protein groups was performed manually at a fixed FDR of 0% by accepting protein IDs with the highest MaxQuant Scores until the first decoy protein ID was reached. The mass spectrometry proteomics data have been deposited to the ProteomeXchange Consortium via the PRIDE partner repository (https://www.ebi.ac.uk/pride/) in MIAPE-compliant format with the dataset identifier PXD030384.

LC-MS/MS proteomics and data analysis of plastoglobule samples

Lyophilized samples in triplicate (three biological replicates) were resuspended in 50 μl of 125 mM Tris, pH 6.8, 4% SDS, 20% glycerol, heated to 60 °C for 5 min and electrophoresed into a 12% Tris-HCl Criterion gel from BioRad at 50 V for ~20 min or until the dye front moved just below the bottom of the sample wells. Electrophoresis was stopped and the gel stained with Coomassie Blue protein stain until bands appeared. Protein bands were excised from the gel and digested according to Shevchenko et al. (71) with modifications. Briefly, gel bands were dehydrated using 100% acetonitrile and incubated with 10 mM dithiothreitol in 100 mM ammonium bicarbonate, pH ~8, at 56 °C for 45 min, dehydrated again, and incubated in the dark with 50 mM iodoacetamide in 100 mM ammonium bicarbonate for 20 min. Gel bands were then washed with ammonium bicarbonate and dehydrated again. Sequencing grade modified trypsin was prepared to 0.01 μg/μl in 50 mM ammonium bicarbonate and ~100 μl of this was added to each gel band so that the gel was completely submerged. Bands were then incubated at 37 °C overnight. Peptides were extracted from the gel by water bath sonication in a solution of 60% Acetonitrile/1% Trifluoroacetic acid and vacuum dried to ~2 μl. An injection of 5 μl was automatically made using a Thermo (www.thermo.com) EASYnLC 1200 onto a Thermo Acclaim PepMap RSLC 0.1 mm × 20 mm C18 trapping column and washed for ~5 min with Buffer A. Bound peptides were then eluted over 65 min onto a Thermo Acclaim PepMap RSLC 0.075 mm × 250 mm resolving column with a gradient of 5% B to 40% B in 24 min, ramping to 90% B at 25 min and held at 90% B for the duration of the run (Buffer A = 99.9% Water/0.1% Formic Acid, Buffer B = 80% Acetonitrile/0.1% Formic Acid/19.9% Water) at a constant flow rate of 300 nl/min. Column temperature was maintained at a constant temperature of 50 °C using an integrated column oven (PRSOV1, Sonation GmbH). Eluted peptides were then sprayed into a ThermoScientific Q-Exactive HF-X mass spectrometer (www.thermo.com) using a FlexSpray spray ion source. Survey scans were taken in the Orbitrap (60,000 resolution, determined at m/z 200), and the top 15 ions in each survey scan were then subjected to automatic higher-energy collision-induced dissociation (HCD) with fragment spectra acquired at 7500 resolution. The resulting MS/MS spectra were processed with the MaxQuant software program, version 1.6.11.0 (69). Peak lists were searched with the embedded Andromeda search engine (70) against the E. coli UNIPROT database concatenated with the MBP-AtABC1K6-His heterologous protein sequence plus the common contaminant list appended by MaxQuant. Oxidation of methionine, deamidation of asparagine and glutamine, and N-terminal acetylation were set as variable modifications, carbamidomethylation was set as a fixed modification. Digestion mode was Trypsin/P with a maximum of two missed cleavages. MS/MS tolerance of the first search was 20 ppm, and main search was 4.5 ppm, with individualized peptide mass tolerance selected. False discovery rate (FDR) at peptide spectrum match and protein levels was set as 0.01. Filtering of resulting protein groups was performed manually at a fixed FDR of 0% by accepting protein IDs with the highest MaxQuant Scores until the first decoy protein ID was reached. The mass spectrometry proteomics data have been deposited to the ProteomeXchange Consortium via the PRIDE partner repository (https://www.ebi.ac.uk/pride/) in MIAPE-compliant format with the dataset identifier PXD030698.
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LC-MS/MS proteomics and data analysis of leaf tissue samples

The triplicate samples (three biological replicates of ~100 μg each) were precipitated using chloroform:methanol (1:4) to remove buffer salts and SDS. Protein pellets were then resuspended in 100 mM Tris–HCl, pH 8.0, supplemented with 4% (w/v) sodium deoxycholate and digested according to Humphrey et al. (72). Following digestion, sodium deoxycholate was removed by phase extraction, peptides desalted using C18 StageTips and eluates dried by vacuum centrifugation. Each sample was then resuspended in 20 μl of 2% acetonitrile/0.1% trifluoroacetic acid.

Injections of ~2 μg were automatically made using a Thermo (www.thermo.com) EASYnLC 1200 onto a Thermo Acclaim PepMap RSLC 0.1 mm × 20 mm C18 trapping column and washed for ~5 min with Buffer A. Bound peptides were then eluted over 125 min onto a Thermo Acclaim PepMap RSLC 0.075 mm × 500 mm resolving column with a gradient of 8% B to 40% B in 114 min, ramping to 90% B at 115 min and held at 90% B for the duration of the run (Buffer A = 99.9% Water/0.1% Formic Acid, Buffer B = 80% Acetonitrile/0.1% Formic Acid/19.9% Water) at a constant flow rate of 300 nl/min. Column temperature was maintained at a constant temperature of 50 °C using integrated column oven (PRSO-V2, Sonation GmbH).

Eluted peptides were sprayed into a ThermoScientific Q-Exactive HF-X mass spectrometer (www.thermo.com) using a FlexSpray spray ion source. Survey scans were taken in the Orbitrap (35,000 resolution, determined at m/z 200), and the top 15 ions in each survey scan are then subjected to automatic HCD with fragment spectra acquired at 7500 resolution. The resulting MS/MS spectra were converted to peak lists using MaxQuant, v1.6.11.0, and searched against a database containing all of A. thaliana protein sequences available from ARAPORT11 (downloaded from www.arabidopsis.org), with common laboratory contaminants using the Andromeda search algorithm, a part of the MaxQuant environment. The mass spectrometry proteomics data have been deposited to the ProteomeXchange Consortium via the PRIDE partner repository (https://www.ebi.ac.uk/pride/) in MIAPER-compliant format with the dataset identifier PXD030750.

Data availability

All mass spectrometry proteomics data have been deposited at the ProteomeXchange Consortium via the PRIDE partner repository (https://www.ebi.ac.uk/pride/) in MIAPER-compliant format with the following unique identifiers: PXD030384, PXD036098, and PXD030750.

Supporting information—This article contains supporting information.

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Abbreviations—The abbreviations used are: ABC1, activity of bc1 complex; ACN, acetonitrile; ADCK, aarF domain-containing kinase; CL, constant light; ePK, eukaryotic protein kinase; FDR, false discovery rate; HCD, higher-energy collision-induced dissociation; JA, jasmonic acid; TFA, trifluoroacetic acid.

References


Protein kinase activity of plastoglobule-localized AtABC1K6


