



# JASSY, a chloroplast outer membrane protein required for jasmonate biosynthesis

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Edited by David C. Baulcombe, University of Cambridge, Cambridge, United Kingdom, and approved April 5, 2019 (received for review January 11, 2019)

**Jasmonates are vital plant hormones that not only act in the stress response to biotic and abiotic influences, such as wounding, pathogen attack, and cold acclimation, but also drive developmental processes in cooperation with other plant hormones. The biogenesis of jasmonates starts in the chloroplast, where several enzymatic steps produce the jasmonate precursor 12-oxophytodienoic acid (OPDA) from  $\alpha$ -linolenic acid. OPDA in turn is exported into the cytosol for further conversion into active jasmonates, which subsequently induces the expression of multiple genes in the nucleus. Despite its obvious importance, the export of OPDA across the chloroplast membranes has remained elusive. In this study, we characterized a protein residing in the chloroplast outer membrane, JASSY, which has proven indispensable for the export of OPDA from the chloroplast. We provide evidence that JASSY has channel-like properties and propose that it thereby facilitates OPDA transport. Consequently, a lack of JASSY in *Arabidopsis* leads to a deficiency in accumulation of jasmonic acids, which results in impaired expression of jasmonate target genes on exposure to various stresses. This results in plants that are more susceptible to pathogen attack and also exhibit defects in cold acclimation.**

jasmonate | plant hormones | chloroplast | membrane pore | cold acclimation

Jasmonates (JAs) play an important role in various cellular responses, including the reaction to biotic and abiotic stresses as well as the formation of reproductive organs. JA biosynthesis has been investigated in some detail, and thus the enzymes involved are well understood even with respect to mechanisms and regulation (1, 2). JAs are derived from  $\alpha$ -linolenic acid ( $\alpha$ -LeA), which is released by phospholipase 1 from galactolipids in the chloroplast thylakoid membrane (3). As an initial step,  $\alpha$ -LeA is oxygenated by a lipoxygenase (LOX). Among six LOXs in *Arabidopsis*, LOX2 is thought to drive the bulk of JA formation during the first 2 h after initiation, for example, by wounding (4, 5). Subsequently, 13-allene oxide synthase (AOS) and 13-allene oxide cyclase introduce dioxygen and cyclize the fatty acid molecules to 12-oxophytodienoic acid (OPDA) (6, 7). Of note, OPDA is also found esterified to galactolipids (8, 9). However, it has not been entirely clarified whether esterified oxylipins are produced from free fatty acid intermediates or whether synthesis can occur on fatty acids while they are esterified into lipids (10).

In any case, OPDA is exported from the chloroplast, by as-yet unknown components. After import of OPDA into peroxisomes, possibly by an ATP-binding cassette (ABC) transporter, it is reduced by the peroxisomal OPDA reductase (OPR3) (11–13). These cyclic intermediates are then processed by the peroxisomal fatty acid  $\beta$ -oxidation machinery, producing JA. JA is in turn exported to the cytosol, where JAR1 forms the bioactive compound (+)-7-*iso*-jasmonoyl-L-isoleucine (JA-Ile) (14).

With identification of the JASMONATE-ZIM-DOMAIN (JAZ) proteins, the mechanistic role of JA-Ile in gene activation has been elucidated. While endogenous levels of JA-Ile are low, the JAZ proteins bind to various transcription factors, thereby

repressing their activity (15). MYC2, a basic helix-loop-helix transcription factor, was the first transcription factor identified in this process and was found to regulate the expression of multiple genes that aid the reaction to various stresses, such as insects/herbivores and wounding (16). On a rise in endogenous JA-Ile levels due to an abiotic or a biotic stimulus, JAZ is bound by an F-box protein, CORONATINE INSENSITIVE 1 (COI1), wherein JA-Ile acts as a molecular glue. COI1 in turn is part of the ubiquitin proteasome degradation machinery and forms a Skp1/Cullin/F-box (SCF<sup>COI1</sup>) complex, which has E3 ubiquitin ligase activity. This complex formation thus results in the ubiquitination and degradation of JAZ, thereby allowing the transcription factors to bind to their targets and activate gene expression (17).

Not only do JAs function in biotic stress response, but several studies also have identified JAZ as a repressor of the transcription factors INDUCER OF CBF EXPRESSION 1 and 2 (*ICE1* and *ICE2*), which activate the C-repeat binding factor (CBF) pathway. Consequently, several genes responsible for cold and freezing tolerance are activated (18–20).

Recently, a protein of as-yet unknown function was identified in a proteomics study of chloroplast outer envelopes (OEs). Intriguingly, this protein was found to be coexpressed with a number of genes involved in the JA response (21). This protein, which we termed JASSY, contains a steroidogenic acute regulatory protein-related lipid transfer (START) domain, suggesting a function in the binding and/or transport of hydrophobic molecules. Loss of functional JASSY results in JA deficiency and thus influences cold

## Significance

**Jasmonates are indispensable for plant growth, as well as for adaptation to biotic and abiotic stresses. The synthesis of jasmonates is initiated in the chloroplast, where the precursor, 12-oxophytodienoic acid (OPDA) is derived from plastid lipids. Despite the fact that much information has been gained on the enzymes involved in the biosynthesis and signaling pathways of jasmonates, the export of OPDA from plastids remains enigmatic. In the present study, we showed that JASSY, a protein localized to the outer chloroplast envelope, facilitates export of OPDA from the chloroplast. Loss-of-function mutants lead to abolished jasmonate accumulation, which in turn results in increased susceptibility to cold treatment, as well as pathogen attack.**

Author contributions: J.S., M.M., and S.S. designed research; L.G., N.D., A.E., M.L., I.S., and A.W. performed research; M.L. contributed new reagents/analytic tools; L.G., N.D., A.E., M.L., I.S., A.W., J.S., M.M., and S.S. analyzed data; and M.M. and S.S. wrote the paper with contributions from all authors.

The authors declare no conflict of interest.

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This article contains supporting information online at [www.pnas.org/lookup/suppl/doi:10.1073/pnas.1900482116/-DCSupplemental](http://www.pnas.org/lookup/suppl/doi:10.1073/pnas.1900482116/-DCSupplemental).

Published online May 8, 2019.

acclimation as well as pathogen susceptibility in *Arabidopsis* mutants lacking JASSY. In this study, we investigated the role of JASSY in the export of OPDA from chloroplasts.

## Results

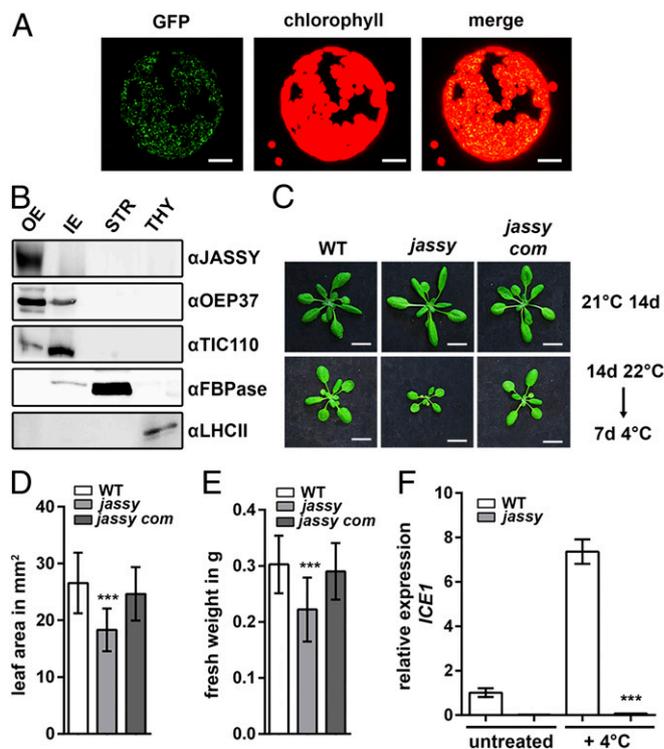
**JASSY Is Localized to the Chloroplast OE.** Chloroplasts contribute to several cellular metabolic pathways by performing crucial enzymatic reactions. Since they are surrounded by two envelope membranes, multiple channels and transporters are required within these membranes to facilitate the exchange of metabolites. Due to recent advances in analysis of subfractionated membranes using proteomics approaches, a number of as-yet uncharacterized proteins potentially residing in the chloroplast envelope membranes have been identified. Among these is a novel potential OE protein, here termed JASSY (At1g70480) (21). We started by investigating the subcellular localization of JASSY via the expression of a GFP fusion protein, as well as chloroplast fractionation and immunologic detection analyses. As a first step, we generated a C-terminal GFP fusion construct, which was used for the transfection of tobacco leaves via *Agrobacterium*. The fusion protein showed a clear chloroplast localization, indicating that JASSY is targeted to the chloroplast (Fig. 1A).

Interestingly, the GFP expression resulted in a ring-shaped signal, again indicating that JASSY might be associated with the chloroplast envelope. To investigate the sublocalization, we fractionated pea chloroplasts into OEs, inner membrane (IEs), thylakoids, and stroma. By doing so, we were able to locate JASSY exclusively in the OE fraction (Fig. 1B). Antisera against the OE protein 37 (OEP37, OE), the translocon of the chloroplast IE 110 (TIC110, IE), the fructose-1,6-bisphosphatase (FBPase, stroma) and the light-harvesting complex II (LHCII, thylakoids) served as markers to show the purity of the isolated chloroplast fractions (Fig. 1B).

To further analyze the localization, we performed an in vitro import assay using radiolabeled JASSY and isolated chloroplasts. We did not observe a shift in size of the translation product after the import reaction, indicating that the protein is not processed in the chloroplast stroma (SI Appendix, Fig. S1A). To determine whether JASSY is merely attached to the chloroplast surface or is indeed efficiently imported, chloroplasts were incubated with thermolysin after the import reaction. The radiolabeled JASSY protein was resistant to this protease treatment (SI Appendix, Fig. S1A). As a control, the ferredoxin-NADP<sup>+</sup> reductase (FNR), which contains a transit peptide and is targeted to the stroma, was treated in the same manner. In this case, the unprocessed translation product was fully digested, whereas the mature protein was protected from the peptidase (SI Appendix, Fig. S1A). Both experiments were performed in parallel, showing that thermolysin treatment was efficient.

Since JASSY does not contain any predicted hydrophobic  $\alpha$ -helical transmembrane domains, we treated pea OEs with several chaotropic reagents to analyze the mode of membrane interaction (SI Appendix, Fig. S1B). After treatment with 1 M NaCl, 6 M urea, 0.1 M Na<sub>2</sub>CO<sub>3</sub>, and 2 M NaBr, JASSY remained entirely in the insoluble fraction after centrifugation in all cases. Only treatment with 0.1 M NaOH released a portion of the JASSY protein from the membrane. As a control, membranes were incubated with 1% SDS, which resulted in total solubilization of JASSY. The integral,  $\beta$ -barrel protein Toc75 served as control and, as expected, was released to the supernatant only after treatment with 1% SDS (SI Appendix, Fig. S1B). Therefore, we conclude that JASSY is stably inserted into the OE.

**Loss of JASSY Decreases Cold Tolerance and Increases Susceptibility to Pathogen Attack.** A homozygous T-DNA insertion mutant of JASSY was isolated, in which the expression of JASSY was entirely abolished as shown on both protein and mRNA levels (SI Appendix, Fig. S1 C and D). Isolated chloroplasts from *Arabi-*



**Fig. 1.** JASSY is localized to the OE of chloroplasts and is important for cold tolerance. (A) Transient expression of JASSY GFP fusion proteins in tobacco. Protoplasts were isolated after the transfection, and fluorescence was observed at 488 nm. (Scale bar: 5  $\mu$ m.) (B) Pea chloroplasts were fractionated into OEs, IEs, stroma (STR), and thylakoids (THY). Fractions were separated by SDS/PAGE and subjected to immunoblotting with antisera against JASSY, OEP37, TIC110, FBPase, and LHCII. (C) WT, *jassy*, and *jassy com* were grown under standard conditions (Upper) or shifted to 4 °C for 7 d after growth for 14 d under standard conditions (Lower). (Scale bar: 1 cm.) (D) The leaf areas of WT, *jassy*, and *jassy com* after 7 d of cold treatment (C, Lower) was determined in square millimeters. The t test was used to indicate significance between WT and *jassy*.  $n > 20$ . (E) Fresh weights of WT, *jassy*, and *jassy com* after 7 d of cold treatment (C, Lower) was measured in grams. The t test was used to indicate significance between WT and *jassy*.  $n > 20$ . (F) Relative mRNA expression levels of *ICE1* in WT and *jassy* monitored by qPCR in untreated plants and after 24 h at 4 °C; the t test indicated significance. Similar results were obtained in three biological replicates.

*dopsis* wild type (WT) and mutant were fractionated into envelopes (mixed fractions containing IEs and OEs), stroma, and thylakoids. The fractions were subjected to SDS/PAGE, and immunoblotting was performed with an antiserum raised against the recombinant *Arabidopsis* JASSY protein. A band at the expected size of 36.3 kDa was again detected in the envelope fraction, which was absent in the mutant. The antibody recognized two additional bands in the stromal fraction; however, these were also present in the *jassy* knockout mutant and thus are assumed to represent cross-reactions of the antiserum (SI Appendix, Fig. S1E). Treatment with antisera against the translocon of chloroplast OE 75 (Toc75, OE), FBPase, and LHCII served as markers to show the purity of the isolated chloroplast fractions (SI Appendix, Fig. S1E).

Under normal growth conditions, *jassy* does not show an altered phenotype compared with the WT (Fig. 1C). However, when 14-d-old plants grown under standard long-day conditions were transferred to 4 °C for 7 d, a significant reduction in growth was observed (Fig. 1C). This was monitored by measurements of the leaf area as well as of the total fresh weight, both of which were significantly reduced in the mutant after cold treatment (Fig. 1 D and E). To ensure that the phenotype was caused by

disruption of *JASSY*, we complemented the mutant by expressing the *JASSY* cDNA. Several independent complemented lines were obtained in which the cold phenotype was fully rescued. The phenotype of a representative line, *jassy com*, is shown in Fig. 1C, along with full recovery of the leaf area and weight in Fig. 1D and E. Gene expression was likewise restored in *jassy com*, as shown by RT-PCR (SI Appendix, Fig. S1D).

Since coexpression data obtained from ATTED-II (atted.jp) suggested that *JASSY* is coexpressed with genes involved in JA metabolism, and JA-deficient mutants are known to be susceptible to cold stress (19, 22), we analyzed expression of the transcription factor *ICE1* by quantitative PCR (qPCR). *ICE1* is activated by JA and plays an important role in cold acclimation by inducing the expression of CBF3. CFBs in turn activate downstream targets mediating the cold acclimation response. Intriguingly, expression levels of *ICE1* were below the limit of detection in the *jassy* mutant, whereas in the WT, *ICE1* mRNA expression was up-regulated almost eightfold after 24 h of cold treatment (Fig. 1F).

Apart from its role in response to abiotic stresses, JA is an important player in the reaction to biotic stresses, such as defense against pathogen attack. Therefore, we treated WT and mutant plants with the *Arabidopsis* pathogen *Botrytis cinerea* for 2 d. Whereas the WT showed only mild infection, the leaves of *jassy* exhibited large lesions (Fig. 2A). The lesion size was quantified, thus verifying the observed phenotype (Fig. 2B). Plant defensin 1.2 (*PDF1.2*) is a well-characterized marker gene specifically induced by JA on pathogen treatment or wounding (23). Therefore, we tested the expression of *PDF1.2* before and after pathogen treatment to determine whether the observed pathogen susceptibility is due to a defect in the JA signaling pathway. Indeed, whereas *PDF1.2* expression was enhanced in the WT on pathogen treatment, no changes in expression level were observed in *jassy* (Fig. 2C).

**The Expression of JA-Responsive Genes Is Not Activated in the *jassy* Mutant on Wounding.** To further investigate whether the loss of *JASSY* induces a defect in JA signaling, we analyzed the expression of several JA-regulated genes in response to wounding (Fig. 3). Similar to the reaction following pathogen attack, no increase in *PDF1.2* expression was observed in *jassy* at 90 min after wounding, in contrast to WT, which showed a 14-fold higher expression (Fig. 3A). A similar result was obtained for *MYC2*, a basic helix-loop-helix transcription factor that is a well-described master regulator of the JA signaling pathway (24) (Fig.

3B). Moreover, JAZ repressor proteins are known to be up-regulated on wounding. A 30-fold up-regulation was observed in the WT, whereas no reaction was observed on the expression level in the *jassy* mutant (Fig. 3C). Not only transcription factors and defense genes are activated by JAZ degradation, but also expression of the JA biosynthesis enzymes is induced in a positive feedback loop by JA after wounding. In line with our previous results, we found that *LOX2* as well as *AOS* were lacking such induction in the *jassy* mutant (Fig. 3D and E). The reduced expression was completely restored in the complemented mutant line; the expression of *PDF1.2* is shown as a representative example (SI Appendix, Fig. S1F). Therefore, *JASSY* seems to be crucial for the initiation of JA-induced signaling pathways.

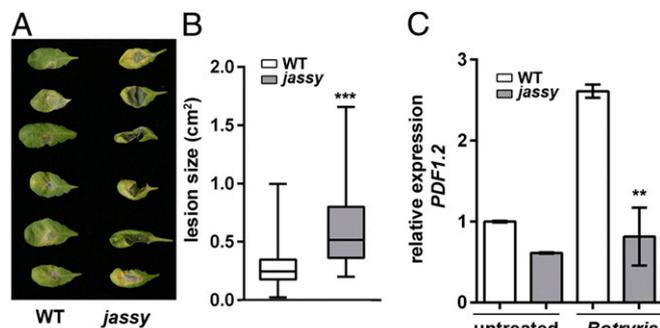
**Lack of *JASSY* Prevents JA Accumulation.** Since we observed no activation of the JA-responsive pathway on a transcriptional level, we aimed to determine whether a lack of *JASSY* leads to a general defect in the accumulation of JA. Since JA is barely detectable under standard conditions, JA levels were measured before and 90 min after wounding. Strikingly, in *jassy*, no increase in basal levels upon wounding was observed, whereas in the WT, a clear accumulation of JA was detected (Fig. 4A).

To strengthen the hypothesis that indeed the lack of JA was responsible for the observed defects in gene expression, we treated WT and *jassy* mutants externally by spraying plants with JA-Ile. Strikingly, the expression of *PDF1.2* could be fully recovered in *jassy* after 30 min (Fig. 4B). While expression in the WT was induced even further at 120 min after the JA-Ile treatment, expression levels were slightly decreased in the mutant. The observed induction clearly demonstrates that *JASSY* functions not in JA-induced gene expression per se, but rather in a step related to the biosynthesis of JA. The lowered transcript abundance of *PDF1.2* at 120 min after the JA treatment likewise indicates that the positive feedback loop in response to JA accumulation, which results in an increased de novo synthesis of JA, is hampered in *jassy*.

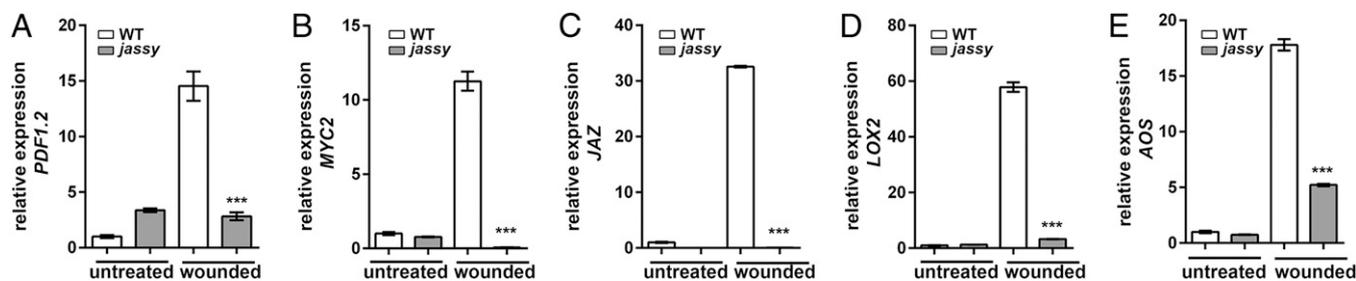
Considering that *JASSY* localizes to the chloroplast OE, it is reasonable to assume that it is likely involved in the export of OPDA. In this case, external treatment with OPDA is likewise expected to restore the transcription of JA-responsive genes. Thus, OPDA was subjected to WT and *jassy* plants, and expression levels of *PDF1.2* were determined at 90, 120, and 180 min after the treatment. Indeed, it was again observed that the expression of *PDF1.2* was induced in *jassy*, almost to WT levels (Fig. 4C). In contrast, treatment solely with  $\alpha$ -LeA resulted in *PDF1.2* expression in the WT, but not in the mutant (Fig. 4D). Taken together, these data strongly suggest that *JASSY* mediates OPDA export from the chloroplast.

***JASSY* Binds to OPDA and Functions as a Membrane Channel.** Since *JASSY* belongs to the Bet v1-like superfamily, which comprises domains known to function in binding hydrophobic components, we analyzed a potential direct interaction between OPDA and recombinant soluble *JASSY* protein purified from *Escherichia coli* using microscale thermophoresis (MST). Binding occurred, even though relatively high concentrations of OPDA had to be applied, resulting in a  $K_D$  of approximately 1 mM (SI Appendix, Fig. S2A). Moreover, no binding curve could be fitted for the tested interaction between *JASSY* and JA (SI Appendix, Fig. S2B). To analyze whether the affinity increased in a membrane environment, the recombinant *JASSY* protein was incorporated into liposomes. Indeed, MST measurements with *JASSY* proteoliposomes and OPDA revealed a much stronger binding event with a  $K_D$  of  $12.8 \pm 3.6 \mu\text{M}$  (Fig. 5A). Again, no binding occurred between *JASSY* liposomes and JA (Fig. 5A).

To elucidate whether *JASSY* has the capacity to form a membrane pore, we performed a liposome leakage assay. To this end, carboxyfluorescein-containing liposomes were generated, into which



**Fig. 2.** Loss of *JASSY* results in advanced susceptibility to *B. cinerea* in *Arabidopsis*. (A) Disease severity in WT and *jassy* plants at 2 d after *B. cinerea* treatment. (B) *Botrytis*-induced lesion size on leaves as shown in A was determined in square centimeters. The *t* test was used to indicate significance.  $n = 100$ . (C) Relative mRNA expression levels of *PDF1.2* in WT and *jassy* plants was monitored by qPCR in untreated and *B. cinerea*-treated plants. The *t* test was used to indicate significance.  $n = 3$ . Similar results were obtained in three biological replicates.



**Fig. 3.** mRNA expression levels of JA-related genes. Expression levels of the indicated transcripts were determined in untreated plants and at 90 min after wounding by qPCR. (A) *PDF1.2*. (B) *MYC2*. (C) *JAZ*. (D) *LOX2*. (E) *AOS*. Similar results were obtained in three biological replicates in all cases. The *t* test was used to indicate significance for all datasets. *n* = 3.

the recombinant JASSY protein was incorporated. During the reconstitution, fluorescence emission of the carboxyfluorescein dye was recorded. Indeed, fluorescence increased continuously over time, indicating release of the carboxyfluorescein dye from the liposomes. The IE protein TIC110 behaved in a similar manner and served as a positive control. In contrast, adding buffer to the liposomes did not result in leakage of carboxyfluorescein (Fig. 5B).

Since the liposome leakage assay indeed suggested that JASSY is able to form membrane pores, we used the planar lipid bilayer technique for an electrophysiological characterization to investigate this observation further. This technique has been used to study pore formation, activity dynamics, and substrate sensitivity of various other membrane channels, including protein translocases, metabolite transporters, pore-forming toxins, and ion channels (25–29) at single-molecule resolution. The technique uses ion conductivity as an activity readout for pore formation. To this end, we inserted JASSY into preformed liposomes via detergent-mediated reconstitution.

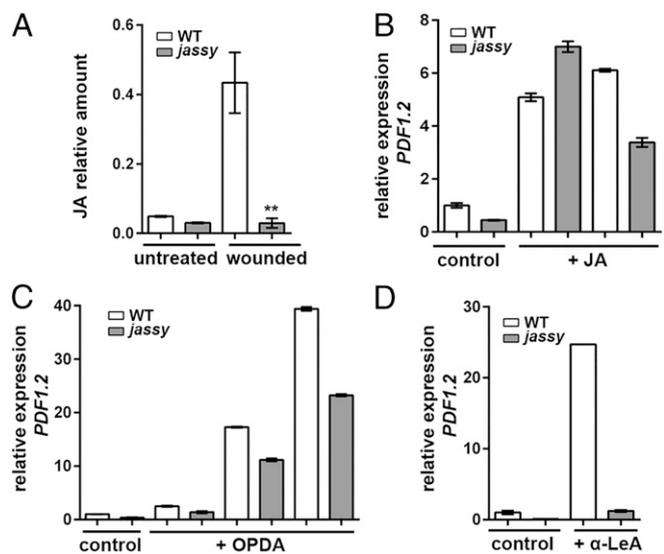
To assess incorporation success, we analyzed comigration of the protein with liposomes in a density flotation assay and found the JASSY-containing floating proteoliposomes. We confirmed that incorporated JASSY was resistant to carbonate extraction and as such behaved as an integral membrane protein (Fig. 6A). In the next step, JASSY-containing vesicles and identically treated liposomes with a purified sample of an empty plasmid mock expression were separately fused with planar lipid bilayers and subjected to high-resolution electrophysiological characterizations. While JASSY-containing proteoliposomes readily fused with bilayers and showed stable channel activity (Fig. 6A and B), the mock sample did not lead to channel insertion even after prolonged incubation of several hours (Fig. 6C). JASSY channels exhibited frequent voltage-dependent gating with dynamic conductance states between 100 and 600 pS at 250 mM KCl (Fig. 6D). A reversion potential of  $31.4 \pm 2.0$  mV at a 12.5-fold KCl gradient was measured; thus, the channels displayed a mild preference for potassium over chloride (SI Appendix, Fig. S3). It will be interesting to see in future studies how exactly OPDA is passing the channel and if different routes for different charged molecules exist, as has been suggested for other relatively wide pores (30, 31).

As we have shown that JASSY binds to OPDA, we probed possible substrate specific alterations of OPDA on the channel properties. Ethanol-solubilized OPDA was added to preinserted channels in situ, and channel characteristics before and after OPDA addition were evaluated. OPDA addition led to significantly increased gating frequencies of incorporated JASSY channels (Fig. 6E). Gating frequency analysis suggested that all conductance states were enhanced similarly, with an additional increased abundance of a high conductance state above 600 pS (Fig. 6F). While OPDA addition led to a sixfold increase in gating frequency on average, no significant change in gating frequency or

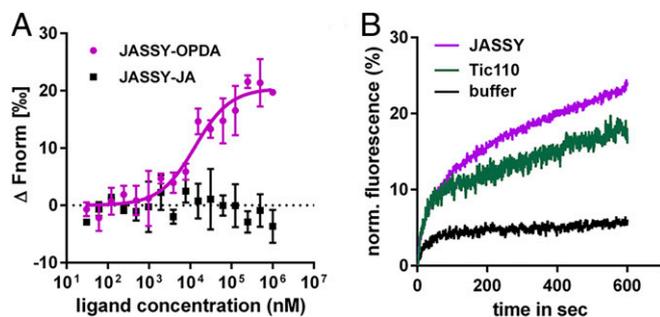
any other determined channel characteristics was observed after addition of ethanol or JA (Fig. 6G).

## Discussion

Whereas the enzymatic steps involved in JA biosynthesis have been investigated and studied in detail for decades now, only slow progress has been made concerning the transport of JAs and its precursors. Recently, JAT1 was identified as a transporter for JA-Ile into the nucleus (14). As to OPDA, the ABC transporter COMATOSE (CTS) has been suggested to be at least partially involved in OPDA import into peroxisomes. CTS is thought to transport a wide range of peroxisomal fatty acids or acyl-CoA-coupled fatty acids. Levels of JAs are reduced in the mutants but are not absent; thus, it is expected that other pathways for OPDA entry into peroxisomes exist in parallel (12, 32, 33). However, so far nothing is known about the export of OPDA from the chloroplast. In this study, we provide evidence that JASSY is a novel OE protein that fulfils this function.



**Fig. 4.** Measurement of JA levels in WT and *jassy* and treatment with JA-Ile, OPDA, and  $\alpha$ -LeA. (A) The relative amount of JA was determined in untreated WT and *jassy* plants as well as at 90 min after wounding. The *t* test was used to indicate significance. *n* = 3. (B) WT and *jassy* mutants were sprayed with 0.4% ethanol (control) or with JA-Ile. The mRNA expression of *PDF1.2* was determined at 30 and 120 min after the treatment. Similar results were obtained in three biological replicates. (C) WT and *jassy* mutants were sprayed with 0.3% ethanol (control) or with OPDA, and the mRNA expression of *PDF1.2* was determined at 90, 120, and 180 min after the treatment. (D) WT and *jassy* mutants were sprayed with 0.3% ethanol (control) or with  $\alpha$ -LeA, and the mRNA expression of *PDF1.2* was determined at 30 min after the treatment. Similar results were obtained in three biological replicates.



**Fig. 5.** JASSY binds to OPDA and forms a pore in proteoliposomes. (A) Recombinant JASSY protein was reconstituted into liposomes, and MST measurements were performed with OPDA and JA. A binding curve could be fitted only with OPDA and a  $K_D$  value of  $12.7 \pm 3.6 \mu\text{M}$  was calculated.  $n = 3$ . (B) Liposome leakage assay. JASSY or Tic110 protein was added to liposomes, and fluorescence was recorded every millisecond (excitation at 494 nm and emission at 515 nm). Initial fluorescence was set to 0, and total liposome dequenching resulting from 0.5% Triton X-100 was set to 100%.  $1 \times$  PBS buffer, pH 7.4 was added to the liposomes as a negative control.

Due to the defect in OPDA export, JA accumulation in *Arabidopsis* leaves is impaired entirely in the mutant, leading to disturbed responses to wounding, pathogen attack, and cold acclimation. Not only did we observe these effects on a phenotypic level, but we could also show that gene expression in the respective JA-responsive signaling cascades was not induced in mutants lacking JASSY. Experiments showing that gene expression could be reactivated with JA-Ile as well as OPDA spraying clearly demonstrate that JASSY functions upstream of JA perception in the nucleus as well as upstream of OPDA to JA conversion in the cytosol and peroxisome.

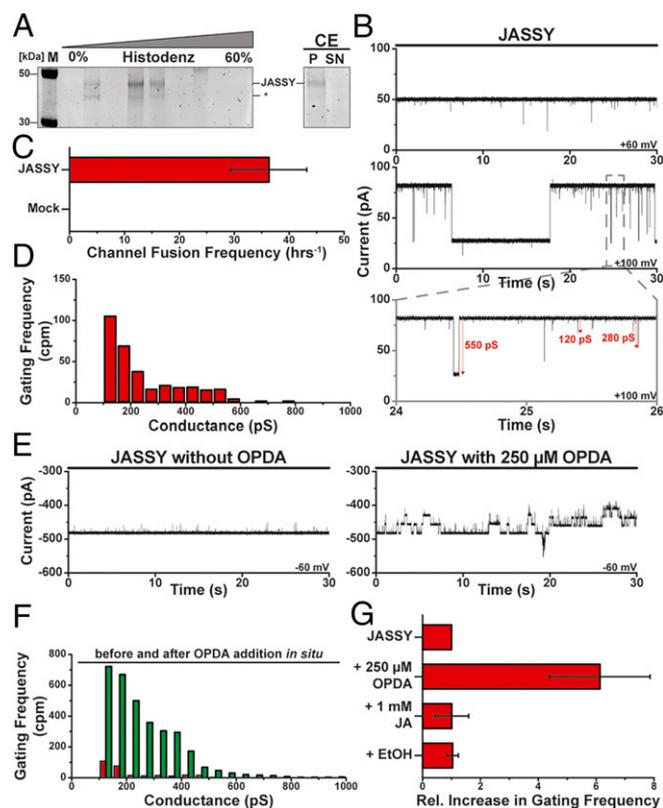
Over the last few years, a number of JA-related mutants have been analyzed to help elucidate the JA biosynthesis pathway as well as the role of JA during stress response and development. For example, the *fad3/fad7/fad8* mutant is deficient in JA synthesis due to lack of  $\alpha$ -LeA, the JA precursor in the chloroplast (34, 35). Moreover, this mutant, just like many other mutants involved in JA biosynthesis or perception, such as *dad1* (36), *aos* (37), *opr3* (38), and *coi* (39), are male sterile. Interestingly, we did not observe problems with either male or female fertility in *jassy*; however, McConn and Browse showed that a leaky allele of the *fad3/fad7/fad8* triple mutant, which accumulates traces of  $\alpha$ -LeA, is fertile (40). Since we also observed a small amount of JA in *jassy*, this might be sufficient for flower development and seed production. Alternatively, the fertility could be explained by the fact that the *Arabidopsis* genome harbors a second gene—At1g23560; *JASSY-2*—with 46% identity on the protein sequence level to JASSY. However, *JASSY-2* is solely transcribed in flower buds, in contrast to JASSY, which is ubiquitously expressed in all developmental stages ([www.bar.utoronto.ca](http://www.bar.utoronto.ca)). Therefore, *JASSY-2* might take over the function of JASSY in flowers and play a role in flower maturation.

JASSY belongs to the Bet v1-like superfamily, a large protein family containing proteins that despite their low sequence similarity share a similar 3D structure (41). The common structure is characterized by a  $\beta$ - $\alpha_2$ - $\beta_6$ - $\alpha$  fold, forming a U-shaped incomplete  $\beta$ -barrel wrapped around a long  $\alpha$ -helix, thus providing a large hydrophobic binding cavity. The Bet v1 family is divided into 11 subfamilies, including the pathogenesis-related protein 10 (PR10), START domain proteins, and oligoetide/cyclase/dehydrases, in the latter of which JASSY shows the highest similarity to that described previously (41). Moreover, a domain of unknown function (DUF220) is predicted in JASSY, partially overlapping with the START domain. The hydrophobic cavity of most of the Bet v1 proteins functions in binding ligands, such as

lipids, sterols, or secondary metabolites (41). In line with this, we observed that JASSY is able to bind OPDA, an interaction likely mediated by this conserved domain and likely important for the transport process.

Moreover, we found that JASSY is not only able to permeabilize membranes, but also forms a voltage-gated channel in planar lipid bilayers. These channels are substrate sensitive and show an OPDA-dependent activation, implicating a mode of transport of JA precursors across the OEs through JASSY. Although the relatively high OPDA concentrations applied in these experiments might not reflect the overall OPDA concentration in chloroplasts, local concentrations of OPDA in the vicinity of JASSY may be significantly elevated in vivo. Moreover, regulation of the channel activity remains to be elucidated. Acylation has been observed to play roles in fatty acid export from the chloroplast as well as OPDA import into peroxisomes and might be used to ensure unidirectional transport of OPDA (42, 43).

Where the function of JAs has been well established in all higher land plants, the presence of JAs in lower land plants and algae is still controversial (44). JAs have been detected in a broad range of bryophytes, but not in the moss *Physcomitrella*



**Fig. 6.** JASSY forms an OPDA-sensitive membrane channel. (A) JASSY proteoliposomes were subjected to density gradient flotation and subsequent sodium carbonate extraction (CE). \*The JASSY degradation band. (B) Current flux through channels formed by JASSY was recorded at various indicated membrane potentials. Conductance changes of selected gating events are indicated. (C) Frequency of channel appearance due to proteoliposome fusion with the lipid bilayer was determined for both JASSY- and mock-expressed incorporations ( $n = 3$ , 2 h per experiment); error bars represent SEM. (D) Gating event frequency was calculated from current recording sets as shown in B.  $n = 6$ . (E) Current trace of multiple JASSY channels was recorded before (Left) and after (Right) in situ incubation with OPDA. (F) Frequency of gating events before (red) and after (green) incubation of JASSY with OPDA calculated from three independent experiments, as shown in D. (G) Average increase in gating frequency of events  $>100$  pS was quantified for OPDA, JA, and ethanol.  $n = 3$ . Error bars represent SEM.

*patens* (45, 46). Nevertheless, OPDA can be detected in the latter and seems to play a role in fertility of the moss (47). Interestingly, JASSY is conserved among land plants, bryophytes, and green algae, supporting the concept that OPDA functions as a signaling molecule in *Physcomitrella* as well as the idea that JA may also trigger responses in other mosses, as well as algae (*SI Appendix, Fig. S4*).

## Materials and Methods

**Plant Material and Growth Conditions.** *Arabidopsis thaliana* WT Columbia ecotype and the mutant were grown on soil under long-day conditions (16-h light/8-h dark, 22 °C, 120  $\mu\text{E m}^{-2} \text{s}^{-1}$ ), unless indicated otherwise. Homozygous *jassy* mutants were isolated and confirmed by genotyping PCR. The WT allele was amplified with At1g70480 LP (Ex5 rev) and At1g70480 RP (Ex3 for), the mutant allele with LB2 and At1g70480 RP (Ex3 for). RT-PCR was used to confirm the absence of the transcript in the mutant. Oligonucleotide sequences are listed in *SI Appendix, Table S1*. The *jassy* mutant (SAIL\_860179) was obtained from the European *Arabidopsis* Stock Centre. For complementation, the coding sequence was cloned into the vector pAUL11 (48). The construct was introduced into *Agrobacterium tumefaciens* strain GV3101, and *jassy* was transformed by floral dip (49).

For cold treatment, plants were first grown under long-day conditions for 14 d and then transferred to 4 °C, 16-h light/8-h dark, 22 °C, 120  $\mu\text{E m}^{-2} \text{s}^{-1}$  for the indicated time periods. Wounding of *Arabidopsis* leaves was performed as follows. Leaves of 4-wk-old plants were wounded by cutting with sharp razor blades. Each leaf was wounded three times on each occasion. No leaf was wounded more than once. Wounded plants were covered and harvested leaves were immediately stored in liquid nitrogen. JA, OPDA, and  $\alpha$ -LeA for feeding experiments were obtained from Cayman Chemical. Plants were sprayed with either 200  $\mu\text{M}$  JA-Ile, 50  $\mu\text{M}$  OPDA or 300  $\mu\text{M}$  of  $\alpha$ -LA. Untreated plants were sprayed with 0.3% ethanol as a control. Plants were harvested for RNA isolation at the indicated time points.

Pea plants (*Pisum sativum* L., cv. Arvica) were grown under a 12-h light (120  $\mu\text{mol m}^{-2} \text{s}^{-1}$ ) and 12-h dark regimen at 21 °C. *N. benthamiana* was grown in soil under greenhouse conditions.

**Pathogen Assay.** Four-week-old plants grown under short day conditions (8 h light/16 h dark, 22 °C, 120  $\mu\text{E m}^{-2} \text{s}^{-1}$ ) were inoculated by applying one drop of 20  $\mu\text{L}$  *B. cinerea* spore suspension with  $2 \times 10^5$  spores per milliliter to each leaf (50). The extent of disease was analyzed by quantifying lesion size using ImageJ software.

**Agrobacterium-Mediated Transient Expression of Fluorescent Proteins in Tobacco.** Four- to 6-wk-old *Nicotiana benthamiana* leaves were infiltrated with *Agrobacterium* for transient expression of gene constructs. The *A. tumefaciens* strain Agl1 was transformed with the respective gene constructs, and infiltration was performed as described previously (51). Protoplasts were prepared essentially as described previously (52), except cell walls were digested for 90 min at 40 rpm in 1% cellulase R10 and 0.3% macerase R10 after vacuum infiltration. Fluorescence was observed with a Leica TCS SP5 confocal laser scanning microscope at 20 °C.

**Quantitative Real-Time RT-PCR Analysis.** RNA was extracted from untreated and treated plants as indicated using the Qiagen RNeasy Plant Kit. After quantification of the RNA and digestion of DNA with TURBO DNA-free kit (Life Technologies), first-strand cDNA was synthesized using M-MLV reverse transcriptase (Promega) from 1  $\mu\text{g}$  of RNA. qPCR was performed in a Bio-Rad CFX96 real-time PCR detection system with SYBR Green Real-Time PCR Master Mix (Roche). Expression levels were normalized to the expression of ACTIN2 (AT3g18780). Gene-specific oligonucleotides are listed in *SI Appendix, Table S1*.

**Chloroplast Isolation, Subfractionation, and in Vitro Import.** Pea chloroplast isolation for in vitro protein import and subfractionation was performed as described previously (53, 54). Isolation and subfractionation of chloroplasts from 4-wk-old *Arabidopsis* plants was also performed as described previously (55). Templates for in vitro transcription were cloned into pF3A (*FNR*) and pSP65 (*JASSY*) (Promega), and in vitro transcription was performed with SP6 polymerase (Fermentas). In vitro translation in reticulocyte lysate (Promega) was performed according to the manufacturer's instructions.

For in vitro import chloroplasts equivalent to 10  $\mu\text{g}$  chlorophyll and in vitro translation product were suspended in a total volume of 100  $\mu\text{L}$  of import mix buffer [250 mM methionine, 250 mM cysteine, 5% BSA, 100 mM ATP, 1 M calcium gluconate, 1 M  $\text{NaHCO}_3$ , and 1 $\times$  HMS (50 mM Hepes, 3 mL

$\text{MgSO}_4$ , and 0.3 M sorbitol)]. The reaction was incubated at 25 °C for 15 min. Chloroplasts were reisolated with a 300- $\mu\text{L}$  Percoll cushion (330 mM sorbitol, 1 M Hepes/KOH, 40% Percoll) and centrifugation at 7,000  $\times g$  for 5 min at 4 °C. The pellet was washed twice in washing buffer I (330 mM sorbitol, 1 M Hepes, and 1 M  $\text{MgCl}_2$ ). The pellet was resuspended in 100  $\mu\text{L}$  of wash buffer II (330 mM sorbitol, 50 mM Hepes, and 0.5 mM  $\text{CaCl}_2$ ) that contained 1 mg/mL thermolysin where indicated, followed by incubation on ice for 20 min. Then 0.5 M EDTA was added to stop the reaction. Chloroplasts were washed once in washing buffer III (330 mM sorbitol, 50 mM Hepes, and 5 mM EDTA). The pellet was resuspended in 20  $\mu\text{L}$  of protein. Chloroplasts were separated by SDS/PAGE and analyzed by autoradiography.

**Antisera.** For production of antisera in rabbits, full-length JASSY was expressed in LB medium in Rosetta 2(DE3) cells (Novagen), and JASSY was purified from inclusion bodies with Ni-NTA affinity chromatography (GE Healthcare). Antisera were generated by Pineda. Other antisera used included Tic110 (56), FBpase (57), OEP37 (58), Toc75 (59), and LHClI (60). Proteins were separated by SDS/PAGE, and immunodetection was performed and with enhanced chemiluminescence as described previously (61).

**JA Measurements.** JA was extracted from frozen leaf material before and after wounding as described by Glauser and Wolfender (48). In brief, 200 mg fresh weight frozen material was ground in a 2-mL reaction tube using a ball mill (TissueLyser II; Qiagen) at 30 Hz for 1 min. For extraction, 1.5 mL of pre-cooled isopropanol was added, along with 0.5% (vol/vol) formic acid and 5 ng  $\text{mL}^{-1}$  chloramphenicol as an internal standard. After extensive mixing to fully resuspend the tissue, the samples were mixed using the ball mill at 30 Hz for 4 min, then centrifuged at 16,400 rpm (Centrifuge 5417R; Eppendorf). The supernatant was transferred to a new 2-mL reaction tube and dried using a vacuum concentrator (Concentrator 5301; Eppendorf). Dry pellets were resuspended in 85% methanol using sonication (Sonifier B<sub>12</sub>; Branson Sonic Power). The solution was centrifuged at 16,400 rpm (Centrifuge 5417R; Eppendorf) for 90 s, after which 1 mL of the supernatant was loaded onto a C18 solid-phase extraction (SPE) column (Sep-Pak Vac 1 cc 50 mg; Waters). SPE columns were equilibrated previously using 1 mL of 100% and 1 mL of 85% (vol/vol) methanol. After SPE separation, the total flow through was collected, followed by a 1-mL 85% (vol/vol) methanol washing step. The flow-through and the methanol wash were combined and dried overnight using the vacuum concentration. Dry samples were filled with argon and stored at -80 °C until further measurement.

Liquid chromatography-mass spectrometry (LC-MS) analysis was done using a Dionex UltiMate 3000 ultra-high-performance liquid chromatograph (Thermo Fisher Scientific) along with an Impact II ultra-high-resolution Qq time-of-flight mass spectrometer (Bruker). The dry samples were resolved in 100  $\mu\text{L}$  of 85% (vol/vol) methanol. For separation, 20  $\mu\text{L}$  was injected onto a C18 reversed phase column (Ultra Aqueous C18, 3  $\mu\text{m}$ , 100  $\times$  2.1 mm; Restek) with a 400- $\mu\text{L min}^{-1}$  flow at 30 °C. The solvents used were water (A) and acetonitrile (B), both including 0.1% (vol/vol) formic acid. The 30-min gradient started at 5% B for 2 min, followed by a ramp to 95% B within 20 min. After a 3-min washing step at 95% B, the gradient turned back to 5% B within 1 min and remained there for a 4-min equilibration.

For MS detection, an electrospray ionization source was used in positive mode at 4 kV capillary and 0.5 kV endplate voltage. Nitrogen was the dry gas at 8 L  $\text{min}^{-1}$ , 8 bar, and 200 °C. The Impact II mass spectra were recorded in MS mode from 50 to 1,300  $m/z$  with 40,000 resolution, 1 Hz scan speed, and 0.3 ppm mass accuracy. Compounds were annotated in a targeted approach using specific mass ( $m/z$ ) at retention time and the isotopic pattern. All data were acquired with otofControl 4.0 and HyStar 3.2 (Bruker). Data evaluation was performed with DataAnalysis 5.1, ProfileAnalysis 2.3, and MetaboScape 1.0 (Bruker). Additional data evaluation was done with Microsoft Excel. Bio-Solve or Sigma-Aldrich supplied all solvents and standards in LC-MS grade.

**Protein Expression and Purification of Recombinant JASSY.** The coding sequence of JASSY was cloned into pET51b<sup>+</sup>. Soluble JASSY protein used for MST, electrophysiology, and carboxyfluorescein assays was expressed overnight at 18 °C in *E. coli* in Rosetta 2(DE3) cells, and the protein was purified with Ni-NTA and Strep-Tactin (IBA Lifesciences).

**Generation of Liposomes for MST and Carboxyfluorescein Assay.** Here 20 mg of phosphatidylcholine lipids (Larodan Fine Chemicals) was washed with chloroform/methanol (1:1) and dried under  $\text{N}_2$ . For MST analysis, dried liposomes were resuspended in an MST-suitable buffer (50 mM Tris-HCl pH 7.4, 150 mM NaCl, 10 mM  $\text{MgCl}_2$ , and 0.05% Tween-20) to a concentration of 20 mg/mL. Lipids were subjected five freeze/thaw cycles, and the mixed

lamellar liposomes were incubated with JASSY (1 mg/mL) at a 1:1 ratio for 1.5 h at 4 °C under agitation.

For the carboxyfluorescein assay, the lipids were resuspended in 1× PBS, pH 7.4 to a final concentration of 20 mg/mL, and 20 mM of carboxyfluorescein was added to 20 mg/mL PC lipids. Subsequently, lipids were subjected to five freeze/thaw cycles, and the mixed lamellar liposomes were extruded through a membrane with 200-nm pore size to generate unilamellar vesicles. To remove nonencapsulated dye, the liposomes were dialyzed against 1× PBS buffer, pH 7.4 overnight at 4 °C.

**Microscale Thermophoresis.** Soluble JASSY protein was diluted in MST buffer (50 mM Tris-HCl pH 7.4, 150 mM NaCl, 10 mM MgCl<sub>2</sub>, and 0.05% Tween-20) to 200 nM. JASSY proteoliposomes were diluted to 2.4 μM in the same buffer. Tris-NTA dyes were diluted in PBST buffer (137 mM NaCl, 2.5 mM KCl, 10 mM Na<sub>2</sub>HPO<sub>4</sub>, 2 mM KH<sub>2</sub>PO<sub>4</sub> pH 7.4, and 0.05% Tween-20) to a final concentration of 100 nM. Then 100 μL of protein was mixed with 100 μL of each dye separately, and the reaction mixtures were then incubated for 30 min at room temperature in the dark. Final concentrations of pure JASSY protein and the JASSY proteoliposomes were 50 nM and 600 nM, respectively. The concentrations of OPDA and JA varied between 30 nM and 1 mM. Reactions were incubated for 10 min at room temperature and then loaded into standard Monolith NT.115 glass capillaries (NanoTemper).

**Carboxyfluorescein Assay.** For fluorescence measurements, 5 μL of liposomes were mixed with 995 μL of 1× PBS pH 7.4 to generate a suitable fluorescent signal. After the addition of purified proteins, fluorescence was recorded every millisecond for 300 or 600 s with a PerkinElmer LS55 fluorescence spectrometer with an excitation wavelength of 494 nm and an emission wavelength of 515 nm.

**Electrophysiology and Liposome Preparation.** To prepare the samples for electrophysiological characterization, L-α-phosphatidylcholine and L-α-phosphatidylethanolamine (both from Avanti Polar Lipids) were mixed at an 80:20 molar ratio, dried under nitrogen, then desiccated under vacuum. The dried lipids were fully resuspended in liposomes buffer (150 mM NaCl and 50 mM Tris/HCl, pH 7.5) to 5 mg/mL and subjected to seven freeze/thaw cycles before being extruded through a 200-nm filter. The liposomes and purified recombinant JASSY, or the identical treated mock expression using an empty plasmid, were solubilized with 80 mM of the dialyzable detergent MEGA-9 (Glycon) separately for 15 min at room temperature. Both parts

were pooled and incubated for another 30 min at room temperature with final lipid and protein concentrations of 1.5 mg/mL and 0.2 mg/mL, respectively. The mixture was dialyzed in a 3.5-kDa cutoff dialysis tube against 5 L of liposome buffer to remove detergent, first for 2 h at room temperature and then overnight at 4 °C. The success of incorporation was monitored by density gradient flotation assay and sodium carbonate extraction as described previously (62).

The electrophysiological experiments were performed using the planar lipid bilayer technique as described previously (63). In brief, liposomes with incorporated JASSY were added below the lipid bilayer to allow osmotically driven fusion in the cis chamber (250 mM KCl and 10 mM Mops/Tris, pH 7), while the trans chamber contained a low salt buffer (20 mM KCl and 10 mM Mops/Tris, pH 7). After insertion of a channel into the bilayer, buffers in both chambers (cis and trans) were perfused to symmetrical standard buffer conditions using 20× chamber volumes. For determination of the reversal potential, asymmetric buffer conditions as used for liposome fusion were applied. Electrical recordings were performed using Ag/AgCl electrodes in a 2 mM KCl agar bridge, connected to a GeneClamp 500B amplifier via a CV-5-1GU headstage (Molecular Devices), with the cis electrode connected to the ground. The signal was digitized by a Digidata 1440a A/D converter and recorded with AxoScope 10.3 and Clampex 10.3 software (Molecular Devices). Automated data analysis was performed as described previously (30). To determine substrate effects, OPDA in ethanol, with corresponding amounts of ethanol or JA in ethanol were added to both sides of bilayer-incorporated JASSY channels. The buffer in both chambers was circulated by stirring for 2 min after addition, then left alone for another 2 min before recording.

**Computational Analyses.** Sequences were obtained from Phytozome (<https://phytozome.jgi.doe.gov/pz/portal.html>). Trees were generated by using CLC Main Workbench software (Qiagen). Alignments were generated using the algorithm provided by CLC Main Workbench (developed by Qiagen). Graphs and statistical analyses were generated using GraphPad Prism version 6.0 ([www.graphpad.com](http://www.graphpad.com)).

**ACKNOWLEDGMENTS.** We thank Tamara Hechtl for excellent technical assistance and Annabel Mechela for help with *Arabidopsis* envelope preparations. This work was supported by the Chinese Scholarship Program (L.G.) and the German Research Foundation (SFB-TR 175 Projects B05 and B06 to J.S. and S.S.; SFB 1190 Project P12 to M.M.; and FOR1905 Project P01 to N.D.).

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