

File ID	uvapub:135793
Filename	Rapid_phosphatidic_acid.pdf
Version	final

---

SOURCE (OR PART OF THE FOLLOWING SOURCE):

Type	article
Title	Rapid phosphatidic acid accumulation in response to low temperature stress in Arabidopsis is generated through diacylglycerol kinase.
Author(s)	S.A. Arisz, R. van Wijk, W. Roels, J.K. Zhu, M.A. Haring, T. Munnik
Faculty	FNWI: Swammerdam Institute for Life Sciences (SILS)
Year	2013

FULL BIBLIOGRAPHIC DETAILS:

<http://hdl.handle.net/11245/1.395738>

---

*Copyright*

*It is not permitted to download or to forward/distribute the text or part of it without the consent of the author(s) and/or copyright holder(s), other than for strictly personal, individual use, unless the work is under an open content licence (like Creative Commons).*

---



# Rapid phosphatidic acid accumulation in response to low temperature stress in *Arabidopsis* is generated through diacylglycerol kinase

Steven A. Arisz<sup>1</sup>, Ringo van Wijk<sup>1</sup>, Wendy Roels<sup>1</sup>, Jian-Kang Zhu<sup>2,3</sup>, Michel A. Haring<sup>1</sup> and Teun Munnik<sup>1\*</sup>

<sup>1</sup> Department of Plant Physiology, Swammerdam Institute for Life Sciences, University of Amsterdam, Amsterdam, Netherlands

<sup>2</sup> Department of Horticulture and Landscape Architecture, Purdue University, West Lafayette, IN, USA

<sup>3</sup> Shanghai Center for Plant Stress Biology and Shanghai Institute of Plant Physiology and Ecology, Shanghai Institutes of Biological Sciences, Chinese Academy of Sciences, Shanghai, China

## Edited by:

Xuemin Wang, University of Missouri-St. Louis, USA

## Reviewed by:

Uener Kolukisaoglu, University of Tuebingen, Germany

Ying Gu, Pennsylvania State University, USA

## \*Correspondence:

Teun Munnik, Department of Plant Physiology, Swammerdam Institute for Life Sciences, University of Amsterdam, Science Park 904, NL-1098 XH Amsterdam, Netherlands.

e-mail: t.munnik@uva.nl

Phosphatidic acid (PtdOH) is emerging as an important signaling lipid in abiotic stress responses in plants. The effect of cold stress was monitored using <sup>32</sup>P-labeled seedlings and leaf discs of *Arabidopsis thaliana*. Low, non-freezing temperatures were found to trigger a very rapid <sup>32</sup>P-PtdOH increase, peaking within 2 and 5 min, respectively. In principle, PtdOH can be generated through three different pathways, i.e., (1) via *de novo* phospholipid biosynthesis (through acylation of lyso-PtdOH), (2) via phospholipase D hydrolysis of structural phospholipids, or (3) via phosphorylation of diacylglycerol (DAG) by DAG kinase (DGK). Using a differential <sup>32</sup>P-labeling protocol and a PLD-transphosphatidylation assay, evidence is provided that the rapid <sup>32</sup>P-PtdOH response was primarily generated through DGK. A simultaneous decrease in the levels of <sup>32</sup>P-PtdInsP, correlating in time, temperature dependency, and magnitude with the increase in <sup>32</sup>P-PtdOH, suggested that a PtdInsP-hydrolyzing PLC generated the DAG in this reaction. Testing T-DNA insertion lines available for the seven *DGK* genes, revealed no clear changes in <sup>32</sup>P-PtdOH responses, suggesting functional redundancy. Similarly, known cold-stress mutants were analyzed to investigate whether the PtdOH response acted downstream of the respective gene products. The *hos1*, *los1*, and *fry1* mutants were found to exhibit normal PtdOH responses. Slight changes were found for *ice1*, *snow1*, and the overexpression line *Super-ICE1*, however, this was not cold-specific and likely due to pleiotropic effects. A tentative model illustrating direct cold effects on phospholipid metabolism is presented.

**Keywords:** abiotic stress, cold stress, diacylglycerol kinase, lipid signaling, phosphatidic acid, phosphoinositide, phospholipase, plant signaling

## INTRODUCTION

The potential to survive low temperatures is one of the factors that determine the geographical distribution of plants. Moreover, freezing and cold stress restrict the arable land and yield of crops. Therefore, much effort is made to understand the mechanisms that make plants more tolerant to low temperatures. One of the most popular plant models in these studies is *Arabidopsis thaliana* (Van Buskirk and Thomashow, 2006).

Like many temperate plants, *Arabidopsis* is capable of cold acclimation, i.e., during a period of cold, non-freezing temperatures, its tolerance for freezing temperatures increases. This process, also referred to as cold hardening, involves a myriad of metabolic and developmental changes, accompanied by accumulation of proteins and compatible solutes, and alterations in membrane composition (Uemura et al., 1995; Thomashow, 1999; Cook et al., 2004; Chinnusamy et al., 2007).

Transcriptome profiling and mutant screens have resulted in the characterization of multiple genes involved in the initiation of cold acclimation and freezing tolerance. These include the

conserved CBF/DREB1 transcription factors that are responsible for activating the expression of many cold response (COR) genes via conserved C-repeat elements in their promoters. Zhu and coworkers have used *Arabidopsis* plants transfected with the *RD29A-LUC* construct to select for mutants with altered responses to cold treatment ("cold response mutants"), resulting in the identification of several genes. Enhanced cold-induced expression was found in the *fry1* (Xiong et al., 2001) and *hos1* (Ishitani et al., 1998) mutants, whereas *los1* (Guo et al., 2002) showed decreased expression. Moreover, the dominant negative *ice1* (Chinnusamy et al., 2003) mutation has been demonstrated to negatively affect cold-induced gene transcription by interfering with the function of *AtICE1*, a *myc*-type transcription factor which functions in CBF transcription in cold signaling. The *myb*-type transcription factor SNOW1/MYB15, also binds to the CBF promoter region, interacting with ICE1 (Agarwal et al., 2006). Upon exposure to cold stress (4°C), the transcript levels of CBF/DREB1 genes increase within 15–30 min, followed by the accumulation of COR gene transcripts after about 2 h (Thomashow, 1999).

Much less is known about the signal transduction pathway that precedes the gene expression changes. Nonetheless, there is mounting evidence that  $\text{Ca}^{2+}$  functions as a second messenger (Knight et al., 1996; Knight and Knight, 2000; Carpaneto et al., 2007) and that part of the pathway involves activation of a MAP kinase cascade (Jonak et al., 1996; Mizoguchi et al., 1996; Teige et al., 2004). One of the latest additions to the field of cold signaling is the formation of the lipid second messenger, phosphatidic acid (PtdOH). In suspension-cultured cells, this phospholipid was shown to accumulate within minutes of cold stress (Gawer et al., 1999; Ruelland et al., 2002; Cantrel et al., 2011). Like  $\text{Ca}^{2+}$  and MAP kinases, PtdOH is involved in the signal transduction pathways of several other plant stress responses, including drought, wounding, and pathogen infection (Li et al., 2009; Testerink and Munnik, 2011), and it is not unlikely that these pathways strongly overlap.

In stress-induced signal transduction, PtdOH responses have been mainly attributed to two pathways. It is the direct product of phospholipase D (PLD), which hydrolyses structural phospholipids like phosphatidylcholine (PtdCho) and phosphatidylethanolamine (PtdEtn), and a secondary product of the phospholipase C (PLC) pathway, which first hydrolyzes polyphosphoinositides (PPIs) to diacylglycerol (DAG), that is subsequently phosphorylated to PtdOH by diacylglycerol kinase (DGK). However, metabolism of DAG and PtdOH is more complex, since multiple sources have now been demonstrated, and PtdOH is formed *de novo* via acylation of glycerolphosphate (Gro3P) as a common intermediate in glycerolipid biosynthesis, both in the plastid and the ER. Thus, PtdOH is precursor to all phosphoglycerolipids as well as triacylglycerols and galactolipids, and its turnover is crucial in determining lipid metabolic fluxes and membrane compositions.

The *Arabidopsis* genome is predicted to encode 12 PLDs, 9 PLCs, and 7 DGKs (Gomez-Merino et al., 2004; Testerink and Munnik, 2005, 2011; Tasma et al., 2008; Arisz et al., 2009; Li et al., 2009; Munnik and Testerink, 2009). Their genetic abundance and specific gene expression patterns suggests that some of these enzymes are specific to certain locations in specific organs and/or involved in distinct processes. The PLC/DGK and PLD pathways have been implicated in the transcriptional induction of an array of cold-induced genes in *Arabidopsis* (Vergnolle et al., 2005). PLD $\delta$  has been shown to be important in the generation of freezing tolerance during acclimation (Li et al., 2004) in contrast to PLD $\alpha$ 1 which negatively influenced survival of freezing, both in cold-acclimated and in non-acclimated plants (Rajashekar et al., 2006; Chen et al., 2008; Du et al., 2010). In suspension-cultured cells, biochemical evidence was found that cold shock activated both PLC/DGK and PLD pathways (Ruelland et al., 2002). Moreover, several genes have been shown to be upregulated in response to cold stress, including *PLD $\alpha$ 1*, *PLD $\delta$* , *PLC1*, *PLC4*, *PLC5*, *DGK1*, and *DGK2* (Hirayama et al., 1995; Gomez-Merino et al., 2004; Li et al., 2004; Lee et al., 2005). Also, in maize roots and leaves several genes encoding DGK, PLC, and PLD were upregulated within 30 min of cold stress (Sui et al., 2008).

In this study we show that *Arabidopsis* seedlings and leaf disks exposed to low temperatures accumulate PtdOH within minutes.

Using a differential  $^{32}\text{P}$ -labeling strategy (Munnik et al., 1998b; Arisz et al., 2009) and PLD's ability to transphosphatidylate n-butanol to PtdBut (Munnik et al., 1995, 1998b), we provide evidence that the rapid PtdOH response does not originate from PLD but from DGK. The simultaneous decrease in the level of phosphatidylinositolphosphate (PtdInsP) suggests the involvement of a PtdInsP-hydrolyzing PLC. T-DNA insertion lines were used to address the question which DGK and PLC were involved, while the COR mutants *hos1*, *los1*, *fry1*, *ice1*, and *snow1* were analyzed to see whether PtdOH acts up- or down-stream of these genes in the COR.

## MATERIALS AND METHODS

### PLANT MATERIAL

*A. thaliana* seeds were sterilized in 70% EtOH (1 min) and 25% bleach (20 min), and sown on media in Petri dishes. For  $^{32}\text{P}$ -radiolabeling experiments, seedlings were grown on  $\frac{1}{2}$  x Murashige and Skoog (MS) basal medium at pH 5.7 (KOH), solidified with 1.0 % bacto-agar. The *ice1*, *snow1*, *los1*, *hos1*, *fry1* mutants, and their WT's were grown on 1 x MS medium supplemented with 1% sucrose. A 16 h light/8 h dark regime (150  $\mu\text{mol quanta m}^{-2}\text{s}^{-1}$ ) at 21°C was set. To promote uniform germination, plates were kept in the dark at 4°C for 2 days before transfer to a climate room.

### RT-PCR EXPRESSION ANALYSES OF DGK T-DNA INSERTION LINES

Homozygous T-DNA insertion lines of the *DGK* genes where genotyped using primer sequences found in **Table A1** (**Figure A3**). Wild type *A. thaliana Col-0* or lines containing T-DNA insertions in *DGK1*, -3, -5, -7 genes were grown on  $\frac{1}{2}$  x Murashige and Skoog (MS) basal medium supplemented with 1% w/v sucrose at pH 4.6, solidified with 1% w/v daishin agar. To promote uniform germination, plates were kept in the dark at 4°C for 2 days before transfer to a climate room. Seedlings where harvested for RNA isolation after 9 days in a climate room with light regime set at 16 h light/8 h dark at 21°C and Relative Humidity 70%. Additionally, flowers of *A. thaliana* lines containing T-DNA insertions in *DGK2*, -4, -6 genes, and wild type *Col-0* were collected from plants grown in a greenhouse under the same environmental conditions. RNA was isolated using Tri Reagents LS (Sigma) and treated with Turbo RNase free DNase (Ambion) for removal of genomic DNA. The RNA concentration and integrity was analyzed using a Nanodrop ND-1000 spectrophotometer.

cDNA was synthesized from 2  $\mu\text{g}$  total RNA using RevertAid H Minus Reverse Transcriptase (Fermentas) according to the manufacturers protocol. RT-PCR was performed using Accuprime Taq DNA polymerase (Invitrogen). **Table A2** contains the primer sequences used to amplify the different *Arabidopsis DGK* genes and the *At2g28390* (SAND family) reference gene (**Figure A4**; Hong et al., 2010). Thermal cycling was done according to the following profile; 94°C for 2 min, followed by 40 cycles of 94°C for 30 s, 50°C for 30 s, 68°C for 2 min and 1 cycle of 68°C for 6 min.

### $^{32}\text{P}$ -ORTHOPHOSPHATE RADIOLABELING *in vivo* AND ANALYSIS OF PHOSPHOLIPIDS

Five-days-old seedlings or leaf disks (5 mm  $\varnothing$ ) of 3-weeks-old plants were transferred to a 2.0 ml Eppendorf tube, containing

MES (2-[N-morpholino]ethane sulfonic acid)-based buffer of 2.56 mM MES (pH 5.7) and 1 mM KCl. To label phospholipids, 10  $\mu$ Ci carrier-free  $^{32}$ P-orthophosphate per tube was added for 16 h, unless indicated otherwise. Cold shock treatments were executed by transferring tubes to ice water. Incubations were stopped by the addition of HClO<sub>4</sub> (final concentration 5%, w/v), and 10 min of subsequent shaking.

The total solvent was removed and 375  $\mu$ l CHCl<sub>3</sub>/MeOH/HCl (50:100:1, by vol.) was added to extract the lipids. After 10 min of vigorous shaking, two phases were induced by adding 375  $\mu$ l CHCl<sub>3</sub> and 200  $\mu$ l 0.9% (w/v) NaCl. The organic lower phase was then transferred to a tube containing 375  $\mu$ l CHCl<sub>3</sub>/MeOH/1M HCl (3:48:47, by vol.). Shaking, spinning, and removing the upper phase yielded a purified organic phase, which was dried down in a vacuum centrifuge at 50°C. The residue was resuspended in 50  $\mu$ l CHCl<sub>3</sub> and sampled for lipid analysis.

Phospholipids were analyzed by thin-layer chromatography (TLC) on heat-activated silica gel 60 plates (Merck, 20 × 20 cm) using one of the following solvent systems (ratios by vol.): (A) CHCl<sub>3</sub>/MeOH/NH<sub>4</sub>OH (25%)/H<sub>2</sub>O (90:70:4:16); or (B) ethylacetate/*iso*-octane/formic acid/H<sub>2</sub>O (13:2:3:10), of which the organic phase was used for TLC. Solvent A was used for total phospholipid analysis, while B was used to quantitate PtdOH and PtdBut. Radiolabeled phospholipids were visualized and quantified by phosphoimaging (Molecular Dynamics, Sunnyvale CA, USA).

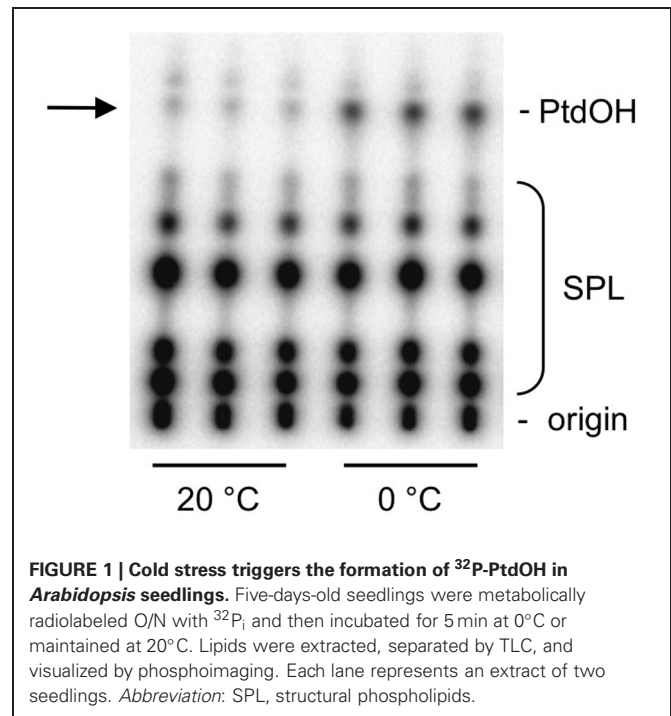
## RESULTS

### COLD STRESS RAPIDLY TRIGGERS A PtdOH RESPONSE

PtdOH levels in plants are approximately 2 mol% of total phospholipids (Walti et al., 2002) which likely represents ER- and plastid-localized PtdOH as precursor and turnover product of structural glycerolipids. To be able to see PtdOH increases during stress-signaling, plants can be metabolically radiolabeled with carrier-free  $^{32}$ P-phosphate ( $^{32}$ P<sub>i</sub>). To study phospholipid metabolism during cold shock in *Arabidopsis*, we radiolabeled 5-days-old seedlings for 16 h with  $^{32}$ P<sub>i</sub> and subsequently incubated them for 5 min at 0°C. Phospholipids were then extracted, separated by TLC and analyzed by autoradiography. A typical  $^{32}$ P-labeling pattern is shown in **Figure 1**, revealing a PtdOH increase in response to cold.

To test the temperature dependency of this response,  $^{32}$ P<sub>i</sub>-prelabeled seedlings were exposed to different temperatures for 5 min. As shown in **Figures 2A,B**, a temperature-dependent PtdOH response was found. Concomitantly, a decrease in  $^{32}$ P-PtdInsP was observed (**Figure 2A**). To investigate whether leaves of adult plants responded similarly, leaf disks of 3-weeks-old plants were subjected to the same labeling procedure and temperature treatments. Quantitation of the PtdOH levels by phosphoimaging revealed a significant response at 8°C or lower (**Figure 2C**), which is different for seedlings which already responded to a shift to 16°C (**Figure 2B**).

Next, the kinetics of the PtdOH response was investigated. As shown in **Figure 3A**, PtdOH accumulation at 0°C in seedlings reached a maximum within 2 min and then leveled off, staying up for at least 2 h. The response of leaf disks of adult plants to 0°C was found to be slightly slower but was still relatively fast, peaking



**FIGURE 1 | Cold stress triggers the formation of  $^{32}$ P-PtdOH in *Arabidopsis* seedlings.** Five-days-old seedlings were metabolically radiolabeled O/N with  $^{32}$ P<sub>i</sub> and then incubated for 5 min at 0°C or maintained at 20°C. Lipids were extracted, separated by TLC, and visualized by phosphoimaging. Each lane represents an extract of two seedlings. Abbreviation: SPL, structural phospholipids.

at 5 min after the onset of incubation after which it leveled off, approaching control levels after 2 h (**Figure 3B**).

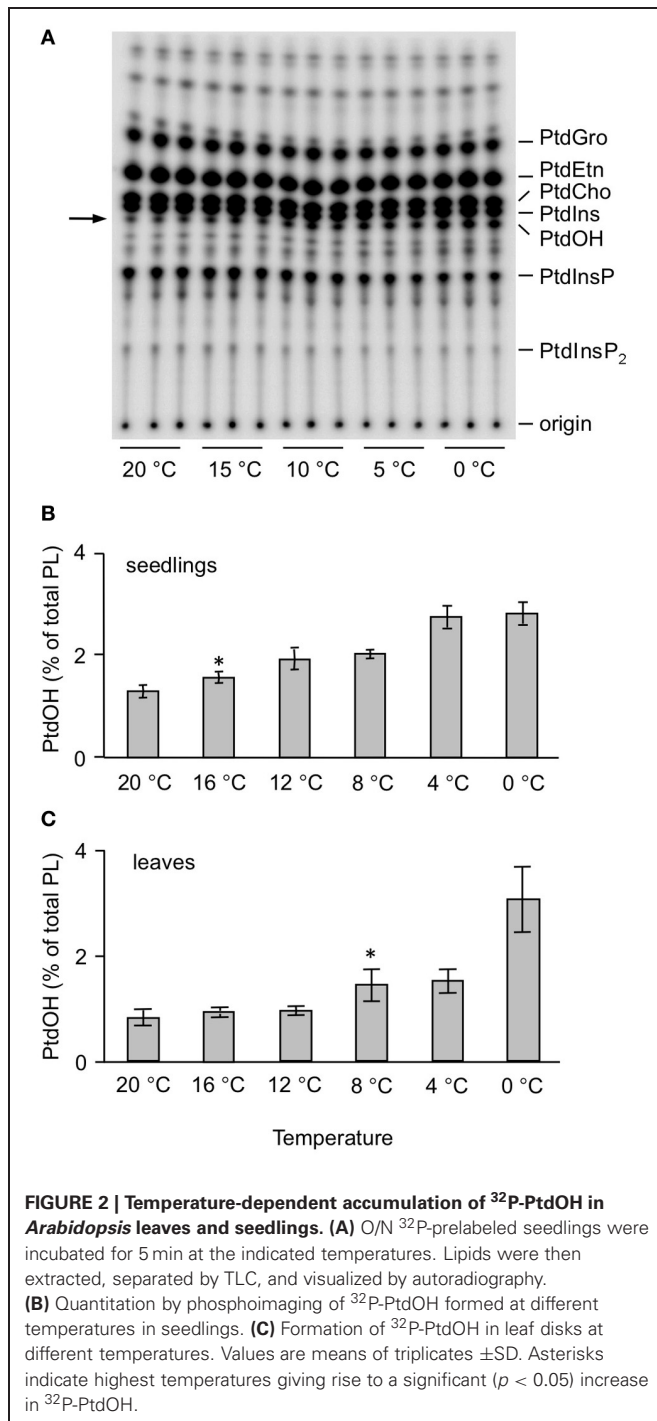
### THE RAPID COLD INDUCED-PtdOH RESPONSE IS GENERATED BY DGK, NOT BY PLD ACTIVITY

Next, we focused on the metabolic origin of the cold-induced PtdOH response. Previous studies in suspension-cultured *Arabidopsis* cells indicated that part of the cold shock-induced PtdOH response was generated by PLD activity (Ruelland et al., 2002). To investigate PLD's contribution a transphosphatidylation assay was performed, i.e., in the presence of a low concentration of a primary alcohol, such as n-ButOH, this serves as a substrate in a PLD-catalyzed reaction generating PtdBut, at the cost of PLD-catalyzed production of PtdOH (Munnik et al., 1995). The accumulation of PtdBut is a measure of PLD activity.

Thus, seedlings were prelabeled for 16 h with  $^{32}$ P<sub>i</sub>, then n-ButOH (0.5% final conc.) was added, and 30 min later the seedlings were transferred to 0°C for 5 min or kept at room temperature. As shown in **Figure 4A**, cold stress did not affect  $^{32}$ P-PtdBut levels, while  $^{32}$ P-PtdOH levels increased. These data indicate that PLD is not responsible for the initial PtdOH response.

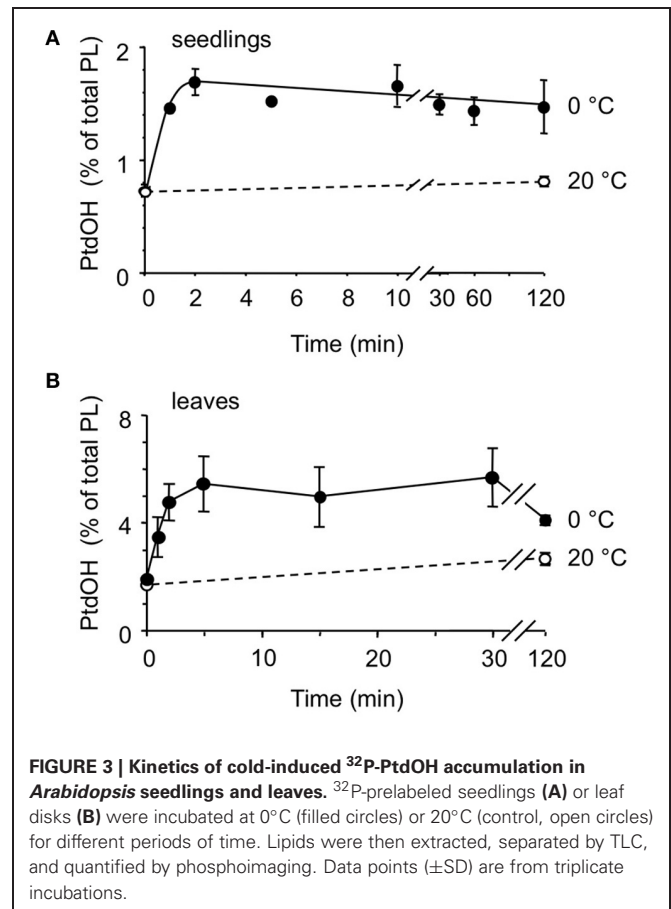
To investigate the potential involvement of DGK, a differential radiolabeling protocol was applied (Munnik, 2001; Ariz et al., 2009). In short, when cells are metabolically labeled with  $^{32}$ P<sub>i</sub>, the phospholipid classes are labeled with different kinetics, depending on the labeling of their precursors, their rates of synthesis, turnover, and pool size. Thus, DGK-derived PtdOH is labeled after relatively short labeling times because it acquires its  $^{32}$ P-phosphate directly from ATP molecules, which are rapidly labeled. This is in contrast to PtdOH arising from PLD activity, which are not labeled until the pool of its precursor, i.e., PtdEtn,





PtdCho, or PtdGro, is sufficiently labeled, which is typically O/N (Munnik et al., 1998b; Arisz and Munnik, 2011).

Thus, seedlings were  $^{32}\text{P}_i$ -prelabeled for different periods of time (20, 60, and 180 min) after which they were subjected for 5 min to  $0^\circ\text{C}$ . As shown in **Figures 4B,C**, cold stress triggered a marked increase in  $^{32}\text{P}$ -PtdOH in seedlings prelabeled for only 20 min. Under these conditions, structural phospholipids like PC and PE were hardly labeled excluding them as precursors to  $^{32}\text{P}$ -PtdOH in a PLD-catalyzed reaction. This is in

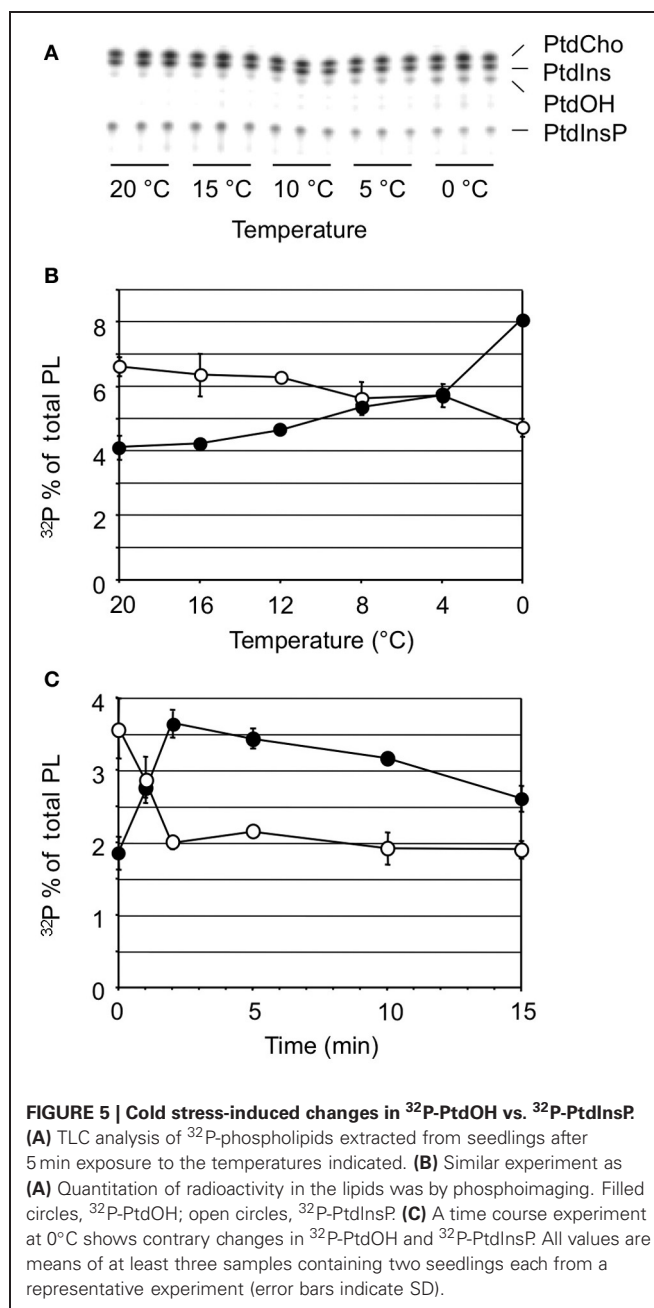
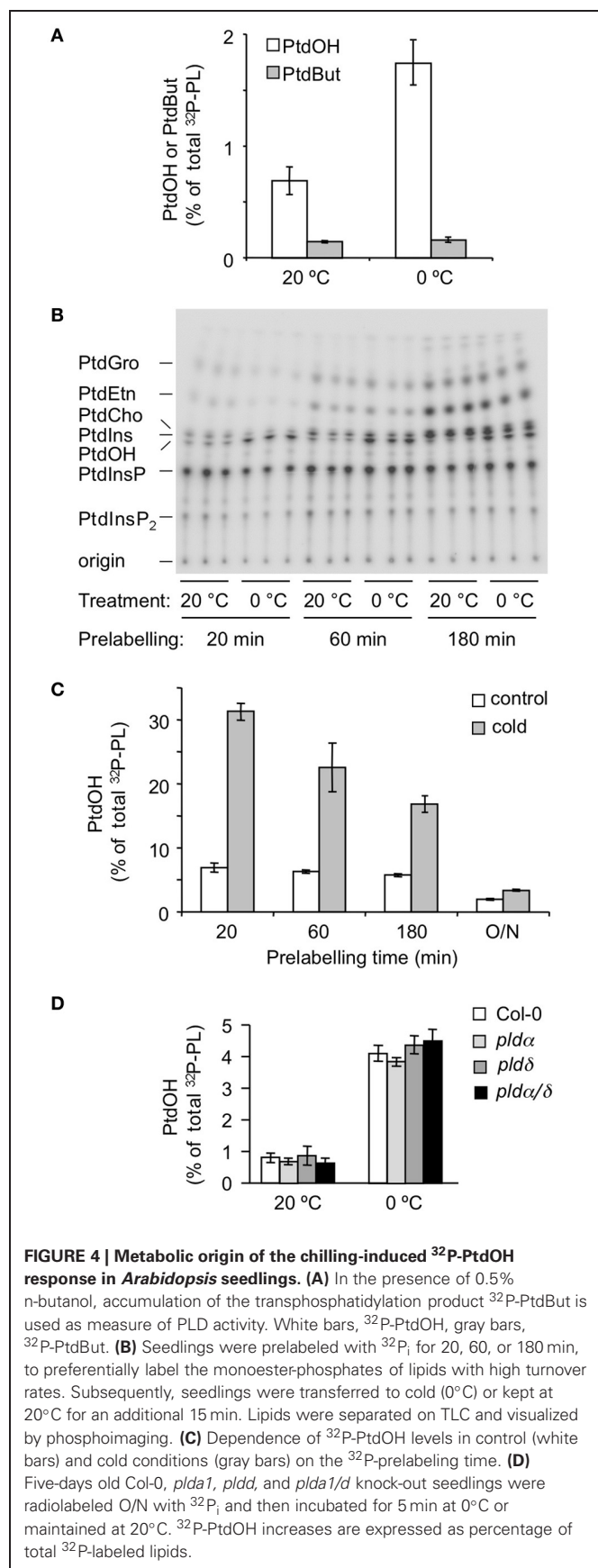


agreement with the results of the transphosphatidylase assay (**Figure 4A**). Hence, the increase in  $^{32}\text{P}$ -PtdOH is unlikely to reflect a PLD activity, and is consistent with a DGK activity. At longer prelabeling time points, the relative increases in  $^{32}\text{P}$ -PtdOH gradually diminished (**Figure 4C**), due to the decrease in the specific radioactivity of the ATP pool and the general increase in structural phospholipid labeling.

Two of the most abundant PLD isoforms in *Arabidopsis*, *PLD $\alpha$ 1* and *PLD $\delta$* , have been implicated in cold stress tolerance (Ruelland et al., 2002; Li et al., 2004; Rajashekar et al., 2006). To test their contribution to the early PtdOH response to cold stress, both single and double KO-mutants were analyzed (Bargmann et al., 2009a,b). As shown in **Figure 4D**, all mutants exhibited a normal PtdOH response upon cold stress.

Together, these results argue against the involvement of PLD in the acute cold-shock-induced PtdOH response and strongly point to a role for DGK.

The implication of DGK in the early COR raised the question of DAG's metabolic origin. One possible source of DAG is the induced PLC hydrolysis of the polyphosphoinositides, PtdInsP and/or PtdInsP<sub>2</sub>, a well-defined plant stress response, which was supported by the observation that  $^{32}\text{P}$ -PtdInsP decreased in response to cold (**Figure 2A**). Moreover, this decrease correlated closely with an equivalent increase in  $^{32}\text{P}$ -PtdOH, in a temperature- and time-dependent fashion (**Figure 5**). These results strongly argue for the scenario that cold stress activates



PLC hydrolysis of PtdInsP to form DAG, which is subsequently phosphorylated to PtdOH by DGK.

#### PtdOH RESPONSES IN ARABIDOPSIS *dgk*- AND COLD STRESS MUTANTS

*Arabidopsis* contains 7 DGK encoding genes. In an attempt to identify the isozyme involved in the cold-induced PtdOH response, a reversed genetic approach was used, screening a series of T-DNA insertion lines (Tables 1, 2). These lines carry insertions in or near the DGK encoding regions, although not all of the lines were established as knockout or knock-down mutants (Table 1; Figure A4). Seedlings of these lines were

**Table 1 | Cold-induced PtdOH formation in seedlings of T-DNA insertion lines.**

Gene	AGI ID	Line	Name	Status*	Control	Cold	Fold increase	Replicates
		wt	<i>Col-0</i>	wt	0.98 ± 0.07	3.00 ± 0.20	3.1	6
<i>AtDGK1</i>	At5g07920	SALK 053412	<i>dgk1-1</i>	KD	0.93 ± 0.07	3.11 ± 0.31	3.4	6
<i>AtDGK2</i>	At5g63770	SAIL 718_G03	<i>dgk2-1</i>	KO	1.00 ± 0.06	2.70 ± 0.48	2.7	6
		SAIL 71_B03	<i>dgk2-2</i>	KO	0.96 ± 0.10	3.19 ± 0.40	3.3	6
<i>AtDGK4</i>	At2g20900	SAIL 339_C01	<i>dgk4-1</i>	no KO	1.22 ± 0.08	3.73 ± 0.26	3.1	3
		SALK 069158	<i>dgk4-2</i>	KO	0.83 ± 0.08	3.65 ± 0.50	4.4	3
<i>AtDGK5</i>	At2g20900	SAIL 1212_E10	<i>dgk5-1</i>	KO	0.85 ± 0.07	3.73 ± 0.15	4.4	3
<i>AtDGK6</i>	At4g28130	SALK 016285	<i>dgk6-1</i>	ND	1.08 ± 0.03	4.48 ± 0.55	4.1	3
<i>AtDGK7</i>	At4g30340	SAIL 51_E04	<i>dgk7-1</i>	KD	0.90 ± 0.17	2.92 ± 0.22	3.2	8
		SALK 059060	<i>dgk7-2</i>	KD	0.91 ± 0.14	3.24 ± 0.15**	3.6	8
		SALK 007896	<i>dgk7-3</i>	no KO	0.87 ± 0.14	2.94 ± 0.19	3.4	8

5-days-old *Arabidopsis* seedlings were labeled O/N with  $^{32}\text{P}_i$  and incubated at 0°C for 5 min. Lipids were then extracted separated by TLC and quantified by phosphoimaging. PtdOH levels are expressed as a percentage of the total  $^{32}\text{P}$ -labeled lipids and values represent averages of multiple samples containing 2 seedlings each ( $\pm$  SD).

\* Transcript analysis by RT-PCR confirmed knock-down (KD) or knock-out (KO) status. Expression of *DGK6* was too low for detection hence the status of *dgk6-1* could not be determined.

\*\* Reproducible statistically significant difference of T-DNA line compared with wild type (Tukey HSD test,  $P < 0.05$ ) within the wild type control homogeneous subset.

**Table 2 | Cold-induced PtdOH formation in leaf disks of T-DNA insertion lines.**

	Control	Cold	Fold increase
<i>Col-0</i>	2.1 ± 0.7	6.0 ± 1.1	2.9
<i>dgk1-1</i>	2.2 ± 0.5	5.1 ± 0.4	2.3
<i>dgk2-1</i>	2.2 ± 0.6	6.7 ± 1.7	3.1
<i>dgk2-2</i>	1.9 ± 0.3	6.5 ± 1.8	3.3
<i>dgk4-1</i>	1.9 ± 0.4	5.9 ± 0.9	3.2
<i>dgk4-2</i>	2.0 ± 0.3	7.2 ± 1.0	3.7
<i>dgk5-1</i>	3.3 ± 0.3	8.4 ± 0.4	2.6
<i>dgk6-1</i>	2.6 ± 0.2	9.7 ± 0.6*	3.7
<i>dgk7-1</i>	2.0 ± 0.4	9.4 ± 1.4*	4.6
<i>dgk7-2</i>	2.3 ± 0.5	10.3 ± 1.0**	4.5
<i>dgk7-3</i>	1.8 ± 0.2	8.7 ± 0.5	4.9

Rosette leaf disks of 5–6-weeks-old *Arabidopsis* plants were labeled O/N with  $^{32}\text{P}_i$  and incubated at 0°C for 5 min. Lipids were extracted, separated by TLC and quantified by phosphoimaging. PtdOH levels are expressed as percentage of the total  $^{32}\text{P}$ -labeled lipids. Values are averages  $\pm$ SD ( $n \geq 3$ ).

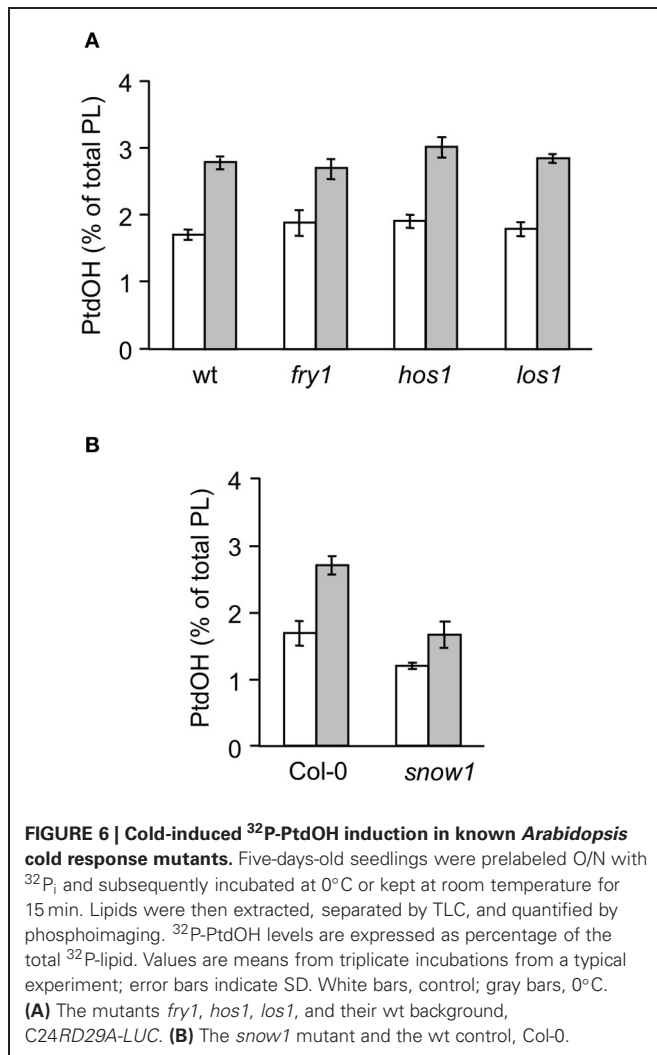
\*\*\* Reproducible statistically significant difference of T-DNA line compared with wild type (Tukey HSD test, \* $P < 0.05$ ; \*\* $P < 0.01$ ) within the wild type control homogeneous subset.

$^{32}\text{P}_i$ -prelabeled (O/N) and then exposed to 0°C for 5 min to measure their PtdOH response. Surprisingly, among the lines with normal control PtdOH levels, neither showed significant defects in the cold-induction of PtdOH, whilst slight larger PtdOH increases were found in cold-stressed *dgk7-2* seedlings (Table 1). In leaf discs, *dgk6-1*, *dgk7-1*, and *dgk7-2* revealed slightly enhanced levels of cold-induced PtdOH (Table 2). Clearly, redundancy is involved and some of the KO/KD lines may constitutively upregulate genes that even mask or enhance the response.

In *Arabidopsis*, several COR mutants have been identified, including *ice1*, *snow1*, *fry1*, *hos1*, and *los1* [(Ishitani et al., 1998; Xiong et al., 2001; Guo et al., 2002; Chinnusamy et al., 2003) mutation has been demonstrated to negatively affect cold-induced gene transcription by interfering with the function of *AtICE1*, a *myc*-type transcription factor which functions in CBF transcription in cold signaling. The *myb*-type transcription factor SNOW1/MYB15, also binds to the CBF promoter region, interacting with ICE1 (Agarwal et al., 2006)]. To gain information on the position of the PtdOH response in the cold sensing pathway, each mutant was analyzed for its cold-induced PtdOH response. As shown in Figure 6A, *fry1*, *hos1*, and *los1* all showed a normal response, but *snow1* had a lower basal and cold shock-induced level of  $^{32}\text{P}$ -PtdOH ( $p = 0.006$ ; Figure 6B); nevertheless, the relative stimulation levels were not significantly altered. Although the cold-induced PtdOH response in *ice1* generally appeared to be lower than wild-type (Figure 7), it did not reach the significance level and was not cold-specific either, because the PtdOH response induced by salt stress (300 mM NaCl, 15 min) was also decreased ( $p = 0.008$ ). Since these seedlings look stunted, pleiotropic effects are most likely to account for the observed differences. Similarly, PtdOH levels in the overexpressor of *ICE1*, *Super-ICE1*, tended to be suppressed, again indicating pleiotropic effects (Figure 7). Together these results indicate that the PtdOH response is upstream.

## DISCUSSION

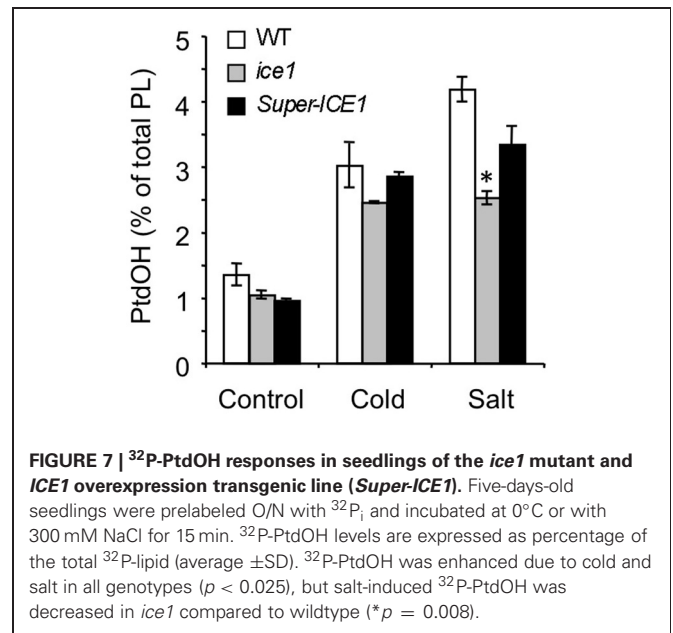
While, in *Arabidopsis* seedlings and leaves, the acclimation process in response to low temperatures is rapidly initiated, it takes 6–7 days to achieve maximal freezing tolerance (Uemura et al., 1995; Peng et al., 2007). The formation of PtdOH has been speculated to function in the regulation of this response (Ruelland et al., 2002; Xiong et al., 2002; Gomez-Merino et al., 2004; Li et al.,



2004; Vergnolle et al., 2005; Rajashekar et al., 2006). While this has previously been studied in suspension-cultured cells, we have focused on the response in whole seedlings and mature leaves. The results showed that cold shock treatment triggered a rapid and sustained (during hours) accumulation of PtdOH, both in seedlings and in leaf discs of mature *Arabidopsis* plants (Figures 1–3). The leaf response was generally more pronounced, but in seedlings the PtdOH increase was faster and already visible upon minor temperature shifts which did not lead to a response in leaves. Since the accumulation of PtdOH is emerging as a common early element in environmental stress responses, and because it is suggested to be involved in the acclimation process, it is important to have knowledge of the underlying mechanisms.

#### DGK RATHER THAN PLD ACTIVITY GENERATES EARLY, COLD-INDUCED PtdOH

Two routes have been found to generate PtdOH under conditions of environmental stress in plants, i.e., PLD hydrolysis of structural phospholipids (i.e., PtdCho/PtdEtn/PtdGro) and phosphorylation of DAG by DGK (Arisz et al., 2009; Testerink and Munnik,



2011). Using transphosphatidylation assays, the absence of a  $^{32}\text{P}$ -PtdBut increase under chilling conditions that triggered massive  $^{32}\text{P}$ -PtdOH responses indicate that PLD is not involved (Figure 4A). Using a differential  $^{32}\text{P}_i$ -labeling assay,  $^{32}\text{P}$ -PtdOH demonstrated to be rapidly labeled, in agreement with a DGK involvement, and in contrast to the labeling of structural phospholipids PtdEtn, PtdCho, and PtdGro, which required long labeling times, again suggesting a PLD-independent pathway (Figure 4B).

Although this seemed at variance with studies of suspension-cultured cells, which suggested a cold-activated PLD activity (Ruelland et al., 2002; Cantrel et al., 2011), it is well-possible that PLD plays a role at a later phase of the COR. This is for example supported by (1) the induced membrane localization of PLD $\delta$  after 1 day at  $2^\circ\text{C}$  (Kawamura and Uemura, 2003), (2) its importance in freezing tolerance (Li et al., 2004; Chen et al., 2008; Du et al., 2010) and (3) the transcriptional regulation of *Arabidopsis* PLD $\delta$  and PLD $\alpha 1$  (Welti et al., 2002; Li et al., 2004) and two PLD $\alpha$  homologs from cotton (Kargiotidou et al., 2010) during cold acclimation. Nevertheless, consistent with the present data, we have found *pld $\alpha 1$ /pld $\delta$*  seedlings to display a normal  $^{32}\text{P}$ -PtdOH response after 5 min at  $0^\circ\text{C}$  (Figure 4D).

#### THE SUBSTRATE FOR DGK MAY BE GENERATED BY A PtdINSP-HYDROLYZING PLC

Since DGK was implicated, the question was raised how the substrate DAG was formed. Several pathways could account for this. A clue was provided by the concomitant decreases of  $^{32}\text{P}$ -PtdInsP, equivalent to the increase of  $^{32}\text{P}$ -PtdOH, suggesting the former to be precursor to DAG and PtdOH via PLC and DGK, respectively (Figures 5 and 8, reactions 1 and 3). Previously, cold stress in *Arabidopsis* cells has been shown to trigger decreases in both PtdInsP $_2$  and PtdInsP (Ruelland et al., 2002). While PtdInsP $_2$  is usually considered as the substrate for



PLC, in plants PtdInsP<sub>2</sub> levels are extremely low, and, *in vitro*, PtdInsP is hydrolyzed equally well (Cho et al., 1993; Munnik et al., 1998a; Munnik and Testerink, 2009; Munnik and Vermeer, 2009).

### PtdOH RESPONSE IN T-DNA INSERTION LINES

To answer the question which of the seven DGK isozyme(s) in *Arabidopsis* was responsible for the cold shock-induced PtdOH response, we tested T-DNA insertion lines with insertions in or near the coding regions, which caused suppressed transcript levels in some but not all (Figures A3 and A4). Nonetheless, neither seedlings (Table 1), nor leaf discs (Table 2) carrying the insertion mutations displayed an abrogation of the PtdOH response. In contrast, in both systems a knock-down allele of *DGK7*, *dgk7-2*, was associated with an increased accumulation of PtdOH. Knockdown of *DGK2*, did not result in significantly reduced PtdOH levels, perhaps as a consequence of the activity of the structurally similar *DGK1* (Gomez-Merino et al., 2004; Arisz et al., 2009).

These results may indicate a functional redundancy among *Arabidopsis* DGKs, such that deficient gene functions are compensated for by related isozymes. The PtdOH “overshoot” in the *dgk7-2* KD line could reflect such a mechanism. This experimental problem will be precluded by generating multigene KO lines, e.g., *dgk1/dgk2*.

In young seedlings, but also in rosette leaves, *DGK7* is more abundantly expressed than *DGK1* and *DGK2* (Arisz et al., 2009), but its transcript levels have not been found to increase upon cold stress. Transgenically expressed *DGK7* protein has been shown to have *in vitro* DGK activity despite the lack of a C1 domain, which is thought to function in the regulation of kinase activity in cluster-I DGKs, *DGK1* and *DGK2* (Gomez-Merino et al., 2004, 2005; Arisz et al., 2009). *DGK7* belongs to cluster II, together with *DGK3* and *DGK4* whose activity may be responsible for the enhanced PtdOH production in *dgk7-2* seedlings.

### OTHER POTENTIAL SOURCES OF DAG AND PtdOH

Although biochemical evidence strongly suggested a PLC-DGK route, two alternative enzymes might be considered as contributors to the cold-induced PtdOH accumulation as well.

Inositolphosphorylceramide synthase (IPCS) transfers the inositol phosphate group from PtdIns to ceramide to generate inositolphosphorylceramide (IPC) and DAG (Figure A1). In yeast, the PtdIns substrate in this reaction is supplied through dephosphorylation of PtdIns4P by Sac1, coupling the consumption of PtdIns4P to the generation of DAG (Brice et al., 2009). An *Arabidopsis* IPCS, encoded by *ERH1*, has been implicated in pathogenic interactions (Wang et al., 2008). Interestingly, the physiological functions of PLC- and IPCS-mediated pathways may rely not only on the generation of inositol polyphosphates, IPC, and DAG/PtdOH, but also on the consumption of PtdInsP, which has novel functions in the biogenesis of secretory vesicles and the establishment/maintenance of cell polarity (reviewed by Munnik and Nielsen, 2011).

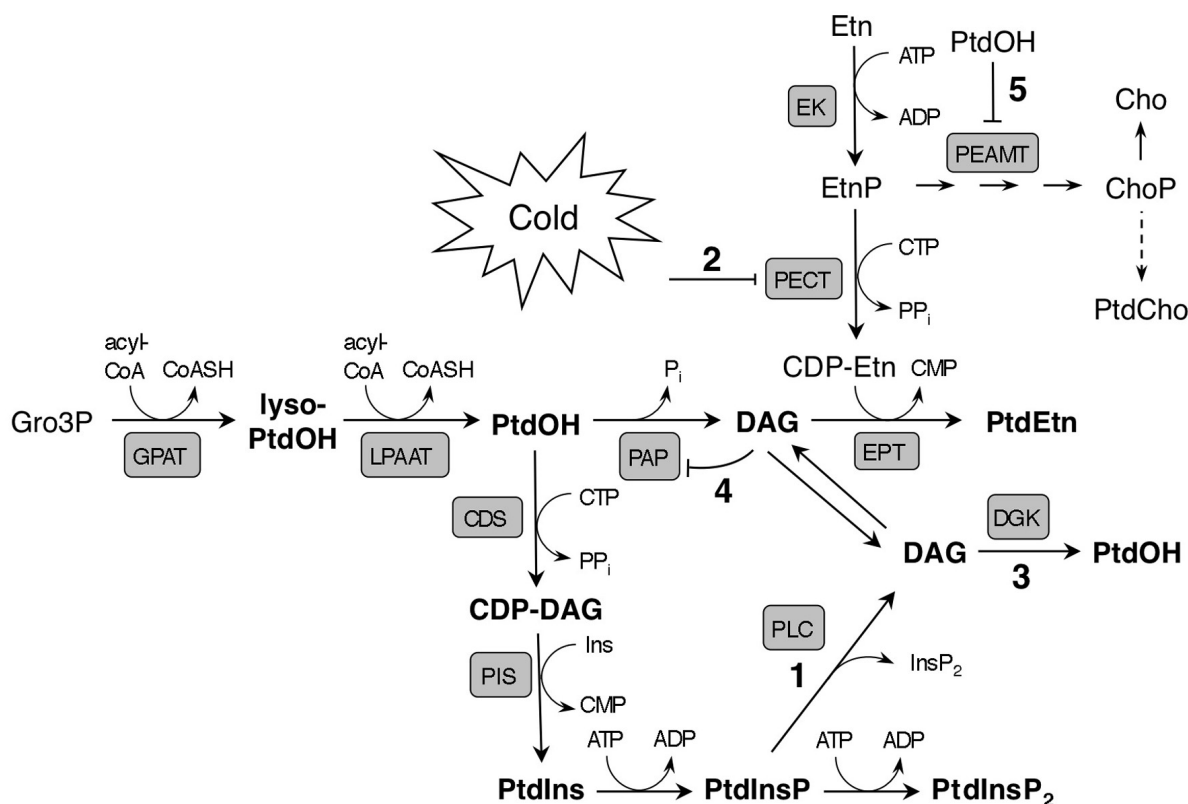
Recently, two interesting novel DAG sources have emerged in *Arabidopsis* stress responses: a PtdCho-hydrolyzing PLC, NPC4, that promotes tolerance to osmotic stresses (Peters et al., 2010), and SFR2, a galactolipid:galactolipid galactosyl transferase (GGGT), that produces DAG and oligogalactolipids to increase freezing tolerance (Thorlby et al., 2004; Moellering et al., 2010). For neither of the enzymes there is direct evidence that links their activity to stress-induced PtdOH accumulation. Rather, the enzymes could provide DAG as precursors for the synthesis of glycerolipids such as PtdCho and MGDG, or triacylglycerol, as for GGGT (Moellering et al., 2010; Moellering and Benning, 2011). It will nevertheless be interesting to subject KO mutants and overexpression lines of the corresponding genes to a differential PtdOH screen as we have applied in this study.

### PtdOH ACCUMULATION DUE TO COLD-INDUCED INHIBITION OF GLYCEROLIPID *de novo* SYNTHESIS?

Cold may have a direct impact on glycerolipid *de novo* synthesis as it lowers diffusion rates and decreases the number of substrate molecules that have sufficient energy to allow enzymatic reactions (Mahan et al., 2004). In fact, in our differential labeling experiments we noticed a cold-induced decrease in the rate of PtdEtn labeling, only visible after approximately 20–30 min of labeling when its *de novo* synthesis in seedlings can just be witnessed (Figure 4B). Also in leaf discs, <sup>32</sup>P-PtdEtn labeling was decreased upon cold incubation, suggesting a cold-induced inhibition of PE's *de novo* biosynthesis (Figure A2). This effect occurred at temperatures ≤8°C, while the decrease remained constant at lower temperatures down to 0°C. In contrast, maximum <sup>32</sup>P-PtdOH accumulation was achieved at 0°C (Figure A2).

Glycerolipid *de novo* synthesis starts with two acylations of Gro3P to generate PtdOH (Figure 8). For the synthesis of PtdIns (and PtdGro), PtdOH is converted to cytidine diphosphate-diacylglycerol (CDP-DAG), the substrate for phosphatidylinositol synthase (PIS). Alternatively, PtdOH is dephosphorylated by phosphatidic acid phosphatase (PAP) to generate DAG as substrate in a reaction by which phosphoethanolamine (EtnP) is transferred from cytidine diphosphate-ethanolamine (CDP-Etn) to the lipid moiety, yielding PtdEtn. Cold does not seem to cause a general inhibition in the uptake of <sup>32</sup>P<sub>i</sub> or its incorporation into the Kennedy pathway of glycerolipid *de novo* synthesis, since labeling of other structural phospholipids was not affected by cold stress (Figure A2). Therefore, the cause of decreased <sup>32</sup>P-PtdEtn labeling is most likely in the synthesis or supply of its headgroup.

The precursor CDP-Etn is generated through the cytidylation of EtnP by phosphoethanolamine cytidyl transferase (PECT), analogous to the PtdCho headgroup precursor CDP-Cho, being the product of phosphocholine cytidyl transferase (CCT) using ChoP as substrate. The latter is produced by repeated methylations of EtnP, catalyzed by phosphoethanolaminemethyltransferase (PEAMT). This activity, which is considered rate-limiting for PtdCho synthesis, likely accounts for the different labeling kinetics of PtdCho and PtdEtn, only the latter being radioactively detected after 30 min of <sup>32</sup>P<sub>i</sub>-labeling.



**FIGURE 8 | Model illustrating potential early effects of cold stress on phospholipid metabolism and *de novo* synthesis in *Arabidopsis*.** The main route to rapid cold-induced PtdOH formation is suggested to be based on the phosphorylation of PLC-generated DAG from PtdInsP (reactions 1/3). The activity of PECT, which produces the precursor of the polar head of PtdEtn, CDP-Etn, is proposed to be down regulated by low ambient temperature (2). This would lead to reduced PtdEtn formation, and potentially, to DAG accumulation, which might cause PtdOH to accumulate as a result from phosphorylation of DAG by a DGK (3), or due to product

inhibition of PAP by DAG (4). The major pathway of PtdCho synthesis depends on methylation of EtnP to ChoP by PEAMT, which could be inhibited by PtdOH (5). Note that the model only highlights immediate effects of cold temperature; longer exposure to cold induces a myriad of metabolic changes which impact lipid biosynthesis in different ways. **Abbreviations:** Acyl-CoA, acyl-coenzyme A; CDS, CDP-DAG synthase; EK, ethanolamine kinase; EPT, CDP-ethanolamine phosphotransferase; GPAT, glycerol 3-phosphate acyltransferase; LPAAT, lysophosphatidic acid acyltransferase; lyso-PtdOH, lysophosphatidic acid.

As previous studies have shown that low temperatures can inhibit the *in vitro* activity of recombinant CCT (Inatsugi et al., 2002), PECT activity is speculated to be similarly downregulated by cold (Figure 8, designated by “2”), resulting in a limited availability of CDP-Etn for PtdEtn synthesis. This would form a bottleneck leading to the accumulation of DAG as precursor for PtdEtn synthesis. Notably, this DAG could be an additional source for cold-induced PtdOH through DGK activity (Figure 8, reaction 3), which has been shown to be partly localized at the ER in *Arabidopsis* (Vaultier et al., 2008). Alternatively, accumulated DAG may block its own formation through feedback inhibition of PtdOH phosphatase (Figure 8, reaction 4), again promoting PtdOH accumulation. Similar regulation of PAP activity by product inhibition has been demonstrated in chloroplast envelope membranes from spinach (Malherbe et al., 1992).

In summary, we have shown a very fast (in minutes) accumulation of PtdOH in response to cold temperatures in *Arabidopsis* seedlings and leaf discs, which was not due

to PLD activity. Instead, <sup>32</sup>P-radiolabeling studies indicated a dominant role of DGK under these conditions. Using single T-DNA insertion lines, we were unable to pinpoint the *DGK* gene(s) involved but do propose that DGK acts in tandem with a PtdInsP-hydrolyzing PLC, based on the close correlation between the increase in <sup>32</sup>P-PtdOH and the decrease in <sup>32</sup>P-PtdInsP.

PtdOH accumulation was not affected by the *fry1*, *hos1*, and *los1* mutations, consistent with an independent, upstream position in cold signaling. Although the *snow1* and *ice1* mutants displayed decreased PtdOH levels they likely reflected pleiotropic effects of the mutations.

Apart from the PLC/DGK route, additional, hypothetical sources of DAG and PtdOH were discussed, viz. via IPCS (Figure A1), NPC, GGGT and lipid *de novo* synthesis (Figure 8). Although for neither of these pathways there is sufficient evidence at present, they should not be ignored when studying PtdOH responses to cold or other environmental stresses.

## REFERENCES

- Agarwal, M., Hao, Y., Kapoor, A., Dong, C. H., Fujii, H., Zheng, X., et al. (2006). A R2R3 type MYB transcription factor is involved in the cold regulation of CBF genes and in acquired freezing tolerance. *J. Biol. Chem.* 281, 37636–37645.
- Ariz, S. A., and Munnik, T. (2011). The salt stress-induced LPA response in *Chlamydomonas* is produced via PLA hydrolysis of DGK-generated phosphatidic acid. *J. Lipid Res.* 52, 2012–2020.
- Ariz, S. A., Testerink, C., and Munnik, T. (2009). Plant PA signaling via diacylglycerol kinase. *Biochim. Biophys. Acta* 1791, 869–875.
- Bargmann, B. O. R., Ariz, S. A., Laxalt, A. M., ter Riet, B., van Schooten, B., Merquiol, E., et al. (2009a). Multiple PLDs required for high salinity- and water deficit tolerance in plants. *Plant Cell Physiol.* 50, 78–89.
- Bargmann, B. O. R., Laxalt, A. M., ter Riet, B., Testerink, C., Merquiol, E., Mosblech, A., et al. (2009b). Reassessing the role of phospholipase D in the *Arabidopsis* wounding response. *Plant Cell Environ.* 32, 837–850.
- Brice, S. E., Alford, C. W., and Cowart, L. A. (2009). Modulation of sphingolipid metabolism by the phosphatidylinositol-4-phosphate phosphatase Sac1p through regulation of phosphatidylinositol in *Saccharomyces cerevisiae*. *J. Biol. Chem.* 284, 7588–7596.
- Cantrel, C., Vazquez, T., Puyaubert, J., Reze, N., Lesch, M., Kaiser, W. M., et al. (2011). Nitric oxide participates in cold-responsive phosphosphingolipid formation and gene expression in *Arabidopsis thaliana*. *New Phytol.* 189, 415–427.
- Carpaneto, A., Ivashikina, N., Levchenko, V., Krol, E., Jeworutzki, E., Zhu, J. K., et al. (2007). Cold transiently activates calcium-permeable channels in *Arabidopsis* mesophyll cells. *Plant Physiol.* 143, 487–494.
- Chen, Q. F., Xiao, S., and Chye, M. L. (2008). Overexpression of the *Arabidopsis* 10-kilodalton acyl-coenzyme A-binding protein ACBP6 enhances freezing tolerance. *Plant Physiol.* 148, 304–315.
- Chinnusamy, V., Ohta, M., Kanrar, S., Lee, B. H., Hong, X., Agarwal, M., et al. (2003). ICE1: a regulator of cold-induced transcriptome and freezing tolerance in *Arabidopsis*. *Genes Dev.* 17, 1043–1054.
- Chinnusamy, V., Zhu, J., and Zhu, J. K. (2007). Cold stress regulation of gene expression in plants. *Trends Plant Sci.* 12, 444–451.
- Cho, M. H., Shears, S. B., and Boss, W. F. (1993). Changes in phosphatidylinositol metabolism in response to hyperosmotic stress in *Daucus carota* L. cells grown in suspension culture. *Plant Physiol.* 103, 637–647.
- Cook, D., Fowler, S., Fiehn, O., and Thomashow, M. F. (2004). A prominent role for the CBF cold response pathway in configuring the low-temperature metabolome of *Arabidopsis*. *Proc. Natl. Acad. Sci. U.S.A.* 101, 15243–15248.
- Du, Z. Y., Xiao, S., Chen, Q. F., and Chye, M. L. (2010). Depletion of the membrane-associated acyl-coenzyme A-binding protein ACBP1 enhances the ability of cold acclimation in *Arabidopsis*. *Plant Physiol.* 152, 1585–1597.
- Gawer, M., Norberg, P., Chervin, D., Guern, N., Yaniv, Z., Mazliak, P., et al. (1999). Phosphoinositides and stress-induced changes in lipid metabolism of tobacco cells. *Plant Sci.* 141, 117–127.
- Gomez-Merino, F. C., Arana-Ceballos, F. A., Trejo-Tellez, L. I., Skirycz, A., Brearley, C. A., Dormann, P., et al. (2005). *Arabidopsis* AtDGK7, the smallest member of plant diacylglycerol kinases (DGKs), displays unique biochemical features and saturates at low substrate concentration: the DGK inhibitor R59022 differentially affects AtDGK2 and AtDGK7 activity *in vitro* and alters plant growth and development. *J. Biol. Chem.* 280, 34888–34899.
- Gomez-Merino, F. C., Brearley, C. A., Ornatowska, M., Abdel-Halim, M. E., Zanor, M. I., and Mueller-Roeber, B. (2004). AtDGK2, a novel diacylglycerol kinase from *Arabidopsis thaliana*, phosphorylates 1-stearoyl-2-arachidonoyl-sn-glycerol and 1,2-dioleoyl-sn-glycerol and exhibits cold-inducible gene expression. *J. Biol. Chem.* 279, 8230–8241.
- Guo, Y., Xiong, L., Ishitani, M., and Zhu, J. K. (2002). An *Arabidopsis* mutation in translation elongation factor 2 causes superinduction of CBF/DREB1 transcription factor genes but blocks the induction of their downstream targets under low temperatures. *Proc. Natl. Acad. Sci. U.S.A.* 99, 7786–7791.
- Hirayama, T., Ohta, C., Mizoguchi, T., and Shinozaki, K. (1995). A gene encoding a phosphatidylinositol-specific phospholipase C is induced by dehydration and salt stress in *Arabidopsis thaliana*. *Proc. Natl. Acad. Sci. U.S.A.* 92, 3903–3907.
- Hong, S. M., Bahn, S. C., Lyu, A., Jung, H. S., and Ahn, J. H. (2010). Identification and testing of superior reference genes for a starting pool of transcript normalization in *Arabidopsis*. *Plant Cell Physiol.* 51, 1694–1706.
- Inatsugi, R., Nakamura, M., and Nishida, I. (2002). Phosphatidylcholine biosynthesis at low temperature: differential expression of CTP: phosphorylcholine cytidylyltransferase isogenes in *Arabidopsis thaliana*. *Plant Cell Physiol.* 43, 1342–1350.
- Ishitani, M., Xiong, L., Lee, H., Stevenson, B., and Zhu, J. K. (1998). HOS1, a genetic locus involved in cold-responsive gene expression in *Arabidopsis*. *Plant Cell* 10, 1151–1161.
- Jonak, C., Kiegerl, S., Ligterink, W., Barker, P. J., Huskisson, N. S., and Hirt, H. (1996). Stress signaling in plants: a mitogen-activated protein kinase pathway is activated by cold and drought. *Proc. Natl. Acad. Sci. U.S.A.* 93, 11274–11279.
- Kargiotidou, A., Kappas, I., Tsaftaris, A., Galanopoulou, D., and Farmaki, T. (2010). Cold acclimation and low temperature resistance in cotton: *Gossypium hirsutum* phospholipase Dalpha isoforms are differentially regulated by temperature and light. *J. Exp. Bot.* 61, 2991–3002.
- Kawamura, Y., and Uemura, M. (2003). Mass spectrometric approach for identifying putative plasma membrane proteins of *Arabidopsis* leaves associated with cold acclimation. *Plant J.* 36, 141–154.
- Knight, H., and Knight, M. R. (2000). Imaging spatial and cellular characteristics of low temperature calcium signature after cold acclimation in *Arabidopsis*. *J. Exp. Bot.* 51, 1679–1686.
- Knight, H., Trewavas, A. J., and Knight, M. R. (1996). Cold calcium signaling in *Arabidopsis* involves two cellular pools and a change in calcium signature after acclimation. *Plant Cell* 8, 489–503.
- Lee, B. H., Henderson, D. A., and Zhu, J. K. (2005). The *Arabidopsis* cold-responsive transcriptome and its regulation by ICE1. *Plant Cell* 17, 3155–3175.
- Li, M., Hong, Y., and Wang, X. (2009). Phospholipase D- and phosphatidic acid-mediated signaling in plants. *Biochim. Biophys. Acta* 1791, 927–935.
- Li, W., Li, M., Zhang, W., Welti, R., and Wang, X. (2004). The plasma membrane-bound phospholipase Ddelta enhances freezing tolerance in *Arabidopsis thaliana*. *Nat. Biotechnol.* 22, 427–433.
- Mahan, J. R., Dotray, P. A., and Light, G. G. (2004). Thermal dependence of enzyme function and inhibition: implications for herbicide efficacy and tolerance. *Physiol. Plant.* 120, 187–195.
- Malherbe, A., Block, M. A., Joyard, J., and Douce, R. (1992). Feedback inhibition of phosphatidate phosphatase from spinach chloroplast envelope membranes by diacylglycerol. *J. Biol. Chem.* 267, 23546–23553.
- Mizoguchi, T., Irie, K., Hirayama, T., Hayashida, N., Yamaguchi-Shinozaki, K., Matsumoto, K., et al. (1996). A gene encoding a mitogen-activated protein kinase kinase is induced simultaneously with genes for a mitogen-activated protein kinase and an S6 ribosomal protein kinase by touch, cold, and water stress in *Arabidopsis thaliana*. *Proc. Natl. Acad. Sci. U.S.A.* 93, 765–769.
- Moellering, E. R., and Benning, C. (2011). Galactoglycerolipid metabolism under stress: a time for remodeling. *Trends Plant Sci.* 16, 98–107.
- Moellering, E. R., Muthan, B., and Benning, C. (2010). Freezing tolerance in plants requires lipid remodeling at the outer chloroplast membrane. *Science* 330, 226–228.
- Munnik, T. (2001). Phosphatidic acid: an emerging plant lipid second messenger. *Trends Plant Sci.* 6, 227–233.
- Munnik, T., Ariz, S. A., De Vrije, T., and Musgrave, A. (1995). G protein activation stimulates phospholipase D signaling in plants. *Plant Cell* 7, 2197–2210.
- Munnik, T., Irvine, R. F., and Musgrave, A. (1998a). Phospholipid signalling in plants. *Biochim. Biophys. Acta* 1389, 222–232.
- Munnik, T., van Himbergen, J. A. J., Ter Riet, B., Braun, F.-J., Irvine, R. F., van den Ende, H., et al. (1998b). Detailed analysis of the turnover of polyphosphoinositides and phosphatidic acid upon activation of phospholipases C and D in *Chlamydomonas* cells treated with non-permeabilizing concentrations of mastoparan. *Planta* 207, 133–145.
- Munnik, T., and Nielsen, E. (2011). Green light for polyphosphoinositide signals in plants. *Curr. Opin. Plant Biol.* 14, 489–497.
- Munnik, T., and Testerink, C. (2009). Plant phospholipid signaling: “in a nutshell.” *J. Lipid Res.* 50(Suppl.), S260–S265.
- Munnik, T., and Vermeer, J. E. M. (2009). Osmotic stress-induced phosphoinositide and inositol

- phosphate signalling in plants. *Plant Cell Environ.* 33, 655–669.
- Peng, Y., Lin, W., Cai, W., and Arora, R. (2007). Overexpression of a *Panax ginseng* tonoplast aquaporin alters salt tolerance, drought tolerance and cold acclimation ability in transgenic *Arabidopsis* plants. *Planta* 226, 729–740.
- Peters, C., Li, M., Narasimhan, R., Roth, M., Welti, R., and Wang, X. (2010). Nonspecific phospholipase C NPC4 promotes responses to abscisic acid and tolerance to hyperosmotic stress in *Arabidopsis*. *Plant Cell* 22, 2642–2659.
- Rajashekar, C. B., Zhou, H. E., Zhang, Y., Li, W., and Wang, X. (2006). Suppression of phospholipase D $\alpha$ 1 induces freezing tolerance in *Arabidopsis*: response of cold-responsive genes and osmolyte accumulation. *J. Plant Physiol.* 163, 916–926.
- Ruelland, E., Cantrel, C., Gawer, M., Kader, J. C., and Zachowski, A. (2002). Activation of phospholipases C and D is an early response to a cold exposure in *Arabidopsis* suspension cells. *Plant Physiol.* 130, 999–1007.
- Sui, Z., Niu, L., Yue, G., Yang, A., and Zhang, J. (2008). Cloning and expression analysis of some genes involved in the phosphoinositide and phospholipid signaling pathways from maize (*Zea mays* L.). *Gene* 426, 47–56.
- Tasma, I. M., Brendel, V., Whitham, S. A., and Bhattacharyya, M. K. (2008). Expression and evolution of the phosphoinositide-specific phospholipase C gene family in *Arabidopsis thaliana*. *Plant Physiol. Biochem.* 46, 627–637.
- Teige, M., Scheikl, E., Eulgem, T., Doczi, R., Ichimura, K., Shinozaki, K., et al. (2004). The MKK2 pathway mediates cold and salt stress signaling in *Arabidopsis*. *Mol. Cell* 15, 141–152.
- Testerink, C., and Munnik, T. (2005). Phosphatidic acid: a multifunctional stress signaling lipid in plants. *Trends Plant Sci.* 10, 368–375.
- Testerink, C., and Munnik, T. (2011). Molecular, cellular and physiological responses to phosphatidic acid formation in plants. *J. Exp. Bot.* 62, 2349–2361.
- Thomashow, M. F. (1999). PLANT COLD ACCLIMATION: freezing tolerance genes and regulatory mechanisms. *Annu. Rev. Plant Physiol. Plant Mol. Biol.* 50, 571–599.
- Thorlby, G., Fourrier, N., and Warren, G. (2004). The SENSITIVE TO FREEZING2 gene, required for freezing tolerance in *Arabidopsis thaliana*, encodes a beta-glucosidase. *Plant Cell* 16, 2192–2203.
- Uemura, M., Joseph, R. A., and Steponkus, P. L. (1995). Cold Acclimation of *Arabidopsis thaliana* (effect on plasma membrane lipid composition and freeze-induced lesions). *Plant Physiol.* 109, 15–30.
- Van Buskirk, H. A., and Thomashow, M. F. (2006). *Arabidopsis* transcription factors regulating cold acclimation. *Physiol. Plant.* 126, 72–80.
- Vaultier, M. N., Cantrel, C., Guerbet, E., Boutte, Y., Vergnolle, C., Cicek, D., et al. (2008). The hydrophobic segment of *Arabidopsis thaliana* cluster I diacylglycerol kinases is sufficient to target the proteins to cell membranes. *FEBS Lett.* 582, 1743–1748.
- Vergnolle, C., Vaultier, M. N., Taconnat, L., Renou, J. P., Kader, J. C., Zachowski, A., et al. (2005). The cold-induced early activation of phospholipase C and D pathways determines the response of two distinct clusters of genes in *Arabidopsis* cell suspensions. *Plant Physiol.* 139, 1217–1233.
- Wang, W., Yang, X., Tangchaiburana, S., Ndeh, R., Markham, J. E., Tsegaye, Y., et al. (2008). An inositolphosphorylceramide synthase is involved in regulation of plant programmed cell death associated with defense in *Arabidopsis*. *Plant Cell* 20, 3163–3179.
- Welti, R., Li, W., Li, M., Sang, Y., Biesiada, H., Zhou, H. E., et al. (2002). Profiling membrane lipids in plant stress responses. Role of phospholipase D  $\alpha$  in freezing-induced lipid changes in *Arabidopsis*. *J. Biol. Chem.* 277, 31994–32002.
- Xiong, L., Lee, B., Ishitani, M., Lee, H., Zhang, C., and Zhu, J. K. (2001). FIERY1 encoding an inositol polyphosphate 1-phosphatase is a negative regulator of abscisic acid and stress signaling in *Arabidopsis*. *Genes Dev.* 15, 1971–1984.
- Xiong, L., Schumaker, K. S., and Zhu, J. K. (2002). Cell signaling during cold, drought, and salt stress. *Plant Cell* 14(Suppl.), S165–S183.

**Conflict of Interest Statement:** The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

Received: 09 December 2011; accepted: 01 January 2013; published online: 22 January 2013.

Citation: Arisz SA, van Wijk R, Roels W, Zhu J-K, Haring MA and Munnik T (2013) Rapid phosphatidic acid accumulation in response to low temperature stress in *Arabidopsis* is generated through diacylglycerol kinase. *Front. Plant Sci.* 4:1. doi: 10.3389/fpls.2013.00001

This article was submitted to *Frontiers in Plant Physiology*, a specialty of *Frontiers in Plant Science*.

Copyright © 2013 Arisz, van Wijk, Roels, Zhu, Haring and Munnik. This is an open-access article distributed under the terms of the Creative Commons Attribution License, which permits use, distribution and reproduction in other forums, provided the original authors and source are credited and subject to any copyright notices concerning any third-party graphics etc.



APPENDIX

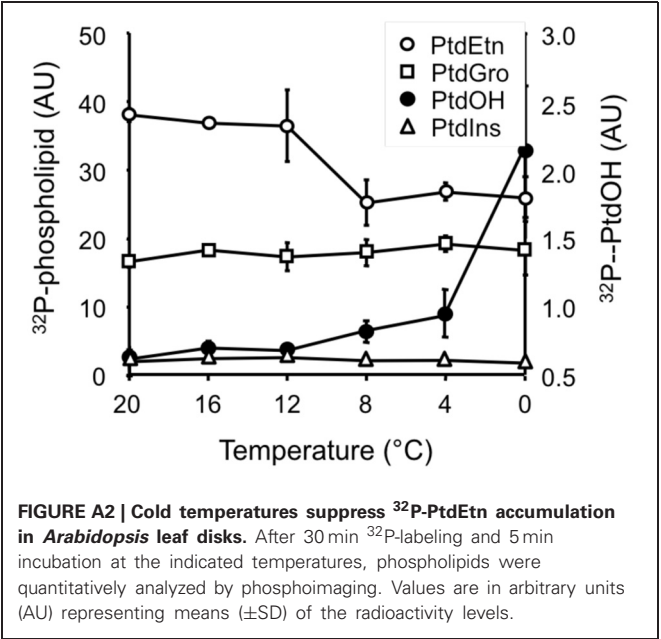
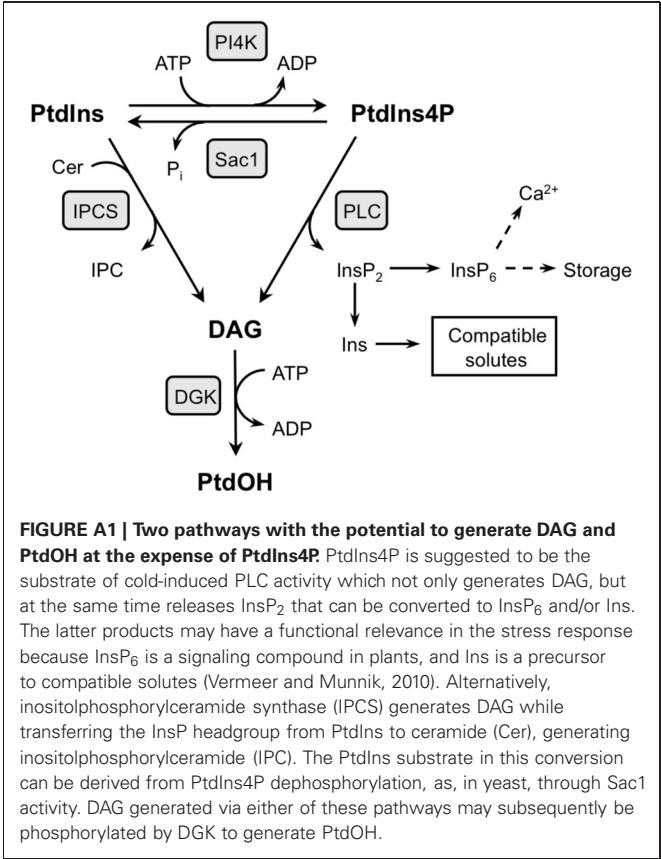
Table A1 | Primers used for genotyping the DGK T-DNA insertion lines.

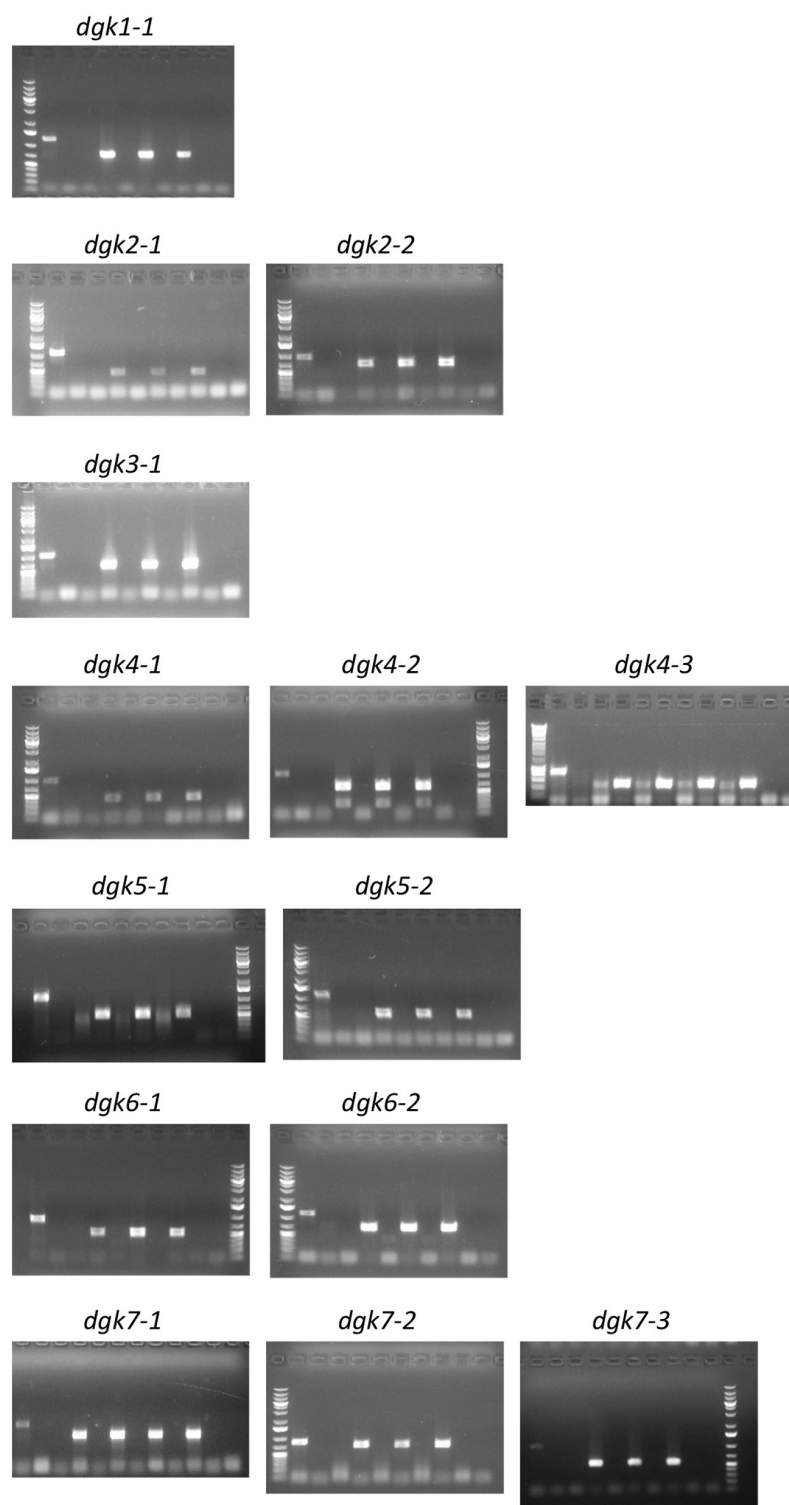
Genotype	LP	RP	Border primer	Sequence	Wild type	Insertion	LP/RP	RP/LB	Insertion
<i>dgk1-1</i>	SALK_053412 GGA TTC TCC TCC CGT AGA TTG	TCA TGC CGT ACT GGA AAA TTC	LBb1.3	ATTTTGCCGATTTCGGAAC	LP/RP	RP/LBb1.3	1205 bp	588 – 888 bp	chr5 2527025
<i>dgk2-1</i>	SAIL_718_G03 GCA AAG AAC AAA AAG GCA CAG	CAG ATG CAA GAC CGC TTT TAG	LB3	TAGCATCTGAATTTTCATAAC CAATCTCGATACAC	LP/RP	RP/LB3	1106 bp	439 – 739 bp	chr5 25520634
<i>dgk2-2</i>	SAIL_71_B03 TTG TAAC TGG ATC TGT TGG CC	CTA AAA GCG GTC TTG CAT CTG	LB3	TAGCATCTGAATTTTCATAAC CAATCTCGATACAC	LP/RP	RP/LB3	957 bp	432 – 732 bp	chr5 25520003
<i>dgk3-1</i>	SALK_082600 TGC TCT CAG TGG GAA GAG ATC	CCG GAA AAC TAT CCG GTT AAC	LBb1.3	ATTTTGCCGATTTCGGAAC	LP/RP	RP/LBb1.3	1179 bp	539 – 839 bp	chr2 8121795
<i>dgk4-1</i>	SAIL_339_C01 AAA GCG AAG CCG ATA TAA AGC	GTC TTT GGC AAA TCG TGG TAG	LB3	TAGCATCTGAATTTTCATAAC CAATCTCGATACAC	LP/RP	RP/LB3	1011 bp	447 – 747 bp	chr5 23371938
<i>dgk4-2</i>	SALK_069158 TTT GAT CCC ATC GAA ACT CAC	GCC GAT GAT GGA CTA CTT GAG	LBb1.3	ATTTTGCCGATTTCGGAAC	LP/RP	RP/LBb1.3	1096 bp	448 – 748 bp	chr5 23372207
<i>dgk4-3</i>	SALK_151239 GGG AGC AGA ATC AGC AGG AAA	GAA TCA TCC TCG CCG TCA ATG	LBb1.3	ATTTTGCCGATTTCGGAAC	LP/RP	RP/LBb1.3	929 bp	?	chr5 23374706
<i>dgk5-1</i>	SAIL_1212_E10 TTC AGA GCA CAT GTG ACC AAC	TCC AAT TCG GAC ATT TGT TTC	LB3	TAGCATCTGAATTTTCATAAC CAATCTCGATACAC	LP/RP	RP/LB3	1163 bp	505 – 805 bp	chr2 8990178
<i>dgk5-2</i>	SAIL_253_E12 GAC TTG AGC TGT TGC TGA TCC	GCG CAA CAA TTT TGG TAG AAG	LB3	TAGCATCTGAATTTTCATAAC CAATCTCGATACAC	LP/RP	RP/LB3	1164 bp	536 – 836 bp	chr2 8993229
<i>dgk6-1</i>	SALK_016285 TGG GTA AAG TGA TCA ATG CAA AAG A	TGG CAA GCG AAA TTG GAA AGA	LBb1.3	ATTTTGCCGATTTCGGAAC	LP/RP	RP/LBb1.3	919 bp	?	chr4 13973176
<i>dgk6-2</i>	SALK_054320 GGG CCA TTA GTG GAA TTA AGC	CCT CCA GAT CAA AAA CCT GAG	LBb1.3	ATTTTGCCGATTTCGGAAC	LP/RP	RP/LBb1.3	1265 bp	606 – 906 bp	chr4 13971587
<i>dgk7-1</i>	SAIL_51_E04 TTT GCA AGA ATG CAT TTT TCC	TGC TGA TGG AGA TGT ACC TCC	LB3	TAGCATCTGAATTTTCATAAC CAATCTCGATACAC	LP/RP	RP/LB3	1118 bp	434 – 734 bp	chr4 14839359
<i>dgk7-2</i>	SALK_059060 CAC GAT CTA ATA ACA CAC CAC ACC C	ACG ACC ACC ACT TTT CGG GTT	LBb1.3	ATTTTGCCGATTTCGGAAC	LP/RP	RP/LBb1.3	904 bp	?	chr4 14841224
<i>dgk7-3</i>	SALK_007896 CTC CAG GAG TTT TAG TTG GGG	CCG AAC ACG TTC TGT TAA AGC	LBb1.3	ATTTTGCCGATTTCGGAAC	LP/RP	RP/LBb1.3	1056 bp	457 – 757 bp	chr4 14839179

Table A2 | Primers used for RT-PCR expression analyses of DGK T-DNA insertion lines.

Gene	Direction	Primename	Sequence
DGK1	sense	SALK_053412_LP	GGATTCTCCTCCCGTAGATTG
	antisense	SALK_053412_RP	TCATGCCGTA CTGGAAAATTC
DGK2	sense	dgk2 fw	GACTGAGAGTTCCACTTTCTC
	antisense	dgk2 rv	GATCTACTCCACCCATATAGC
DGK3	sense	SALK_082600_LP (dgk3-1)	TGCTCTCAGTGGGAAGAGATC
	antisense	dgk3 rv	CAAAC TTCATTCTCACAACAC
DGK4	sense	dgk4 fw	GCAGTTGTTGCATTGAATCTAC
	antisense	dgk4 rv (A)	CCAAAGACTGGTGAGGGACTC
	antisense	dgk4 r3 (B)	CGCATCTTTCCAGTCTCCTC
DGK5	sense	dgk5 fw2	CCAGTGGCAGGACCTCCAC
	antisense	dgk5 rv	GGAATCTTGAAGGTATCCGCAG
DGK6	sense	dgk6 fw	CCTGGAACAGATAGGTCTTCG
	antisense	dgk6 rv	CATTGGCCATTTCATCACAAATCTG
DGK7	sense	SALK_007896_LP	CTCCAGGAGTTTTAGTTGGGG
	antisense	dgk7 rv	GTTGTTTCATGGTTCACCATCC
SAND ref	sense	At2g28390-Q-fw	CAGACAAGGCGATGGCGATA
	antisense	At2g28390-Q-rv	GCTTTCTCTCAAGGGTTTCTGGGT

Sequences of the different primer pairs used to measure the expression of the various DGK genes used in this study. SAND (At2g28390) expression was used as the reference gene (Hong et al., 2010).

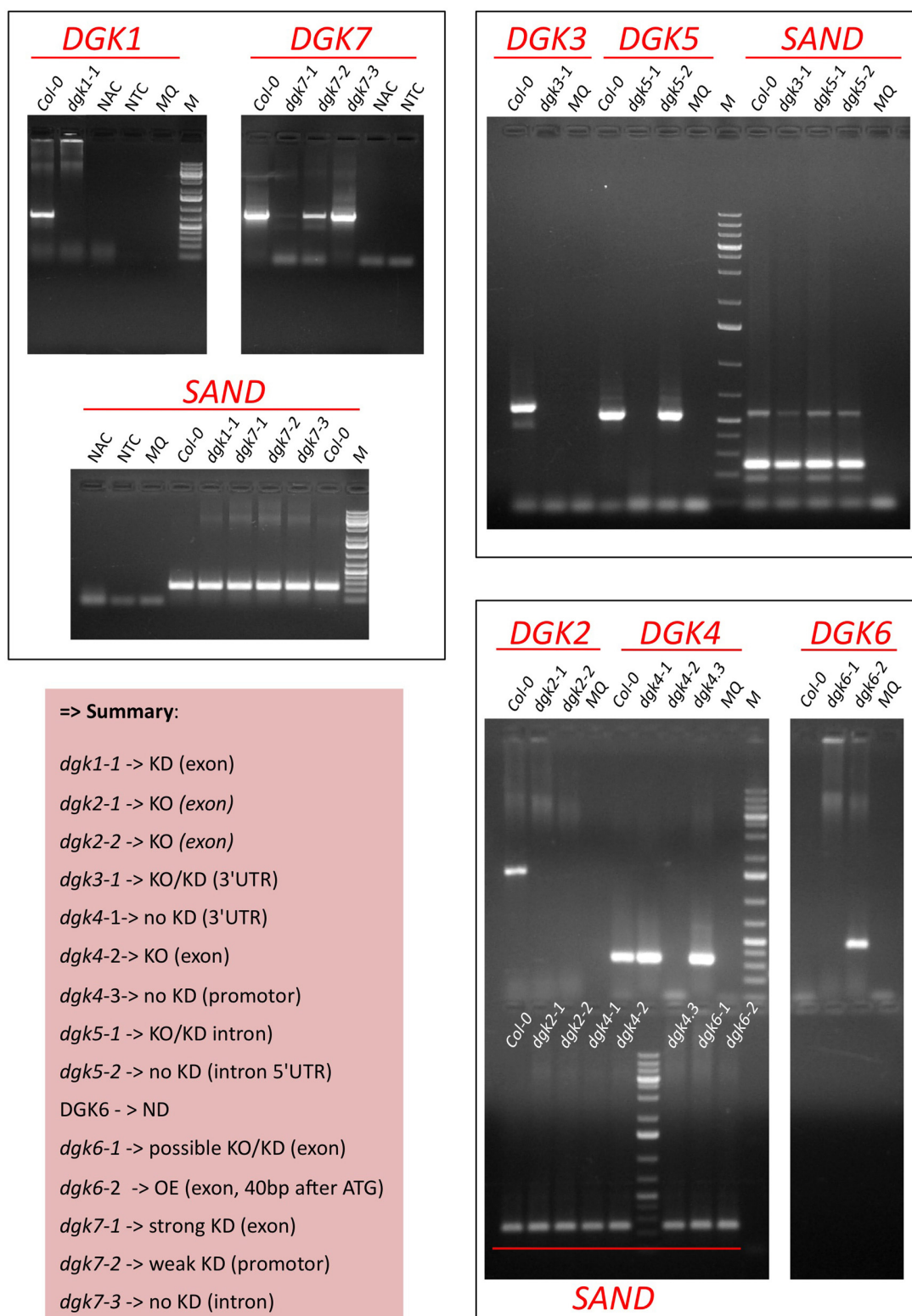




**FIGURE A3 | Genotyping of the Arabidopsis *dgk* T-DNA insertion lines.**

DNA was isolated from leaf discs from 3 to 4 independent plants and zygosity determined by PCR using specific wt- and mutant-primer sets as listed in **Table A1**. Gel sample order (from left to right): Size marker; Col-0 (wt), with first primer set, LP/RP (= wt band) and second primer set, RP/LB (= insertion band); Three (or four) independent *dgk* plants, with first primer

set, LP/RP (= wt band) and second primer set RP/LB (= insertion band); Last lane, primer control (—). Some gels end with the size marker. Results clearly show that all lines used are homozygous T-DNA insertion lines since Col-0 was the only line giving wt bands and all insertion lines gave only bands with the second primer set. For the SALK lines LBb1.3 was used as LB primer, for the SAIL lines LB3 was used as LB primer (see primer list, **Table A1**).



**FIGURE A4 | Determination of *DGK* KO/KD-expression in Arabidopsis T-DNA insertion mutants by RT-PCR.** RNA was isolated from 9-days old roots (*DGK1*, -3, -5, -7) or flowers (*DGK2*, -4, -6) since the latter genes did not reveal expression in the root (not shown). Predicted band Sizes: *DGK1*, 702 bp; *DGK3*, 592 bp; *DGK5*, 534 bp; *DGK6*, 434 bp;

*DGK7* 791 bp; *SAND* (reference gene), 244 bp. Abbreviations: ND, not detectable; NAC, non-amplification control (test for genomic DNA contamination), i.e., RT reaction without RT enzyme on Col-0 RNA; NTC, no target control (test for contamination + primer dimers), i.e., RT reaction with water sample.