Involvement of Phosphatidylinositol 3-kinase in the regulation of proline catabolism in Arabidopsis thaliana

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INTRODUCTION

As sessile organisms, plants need to cope with adverse environmental stresses. Abiotic constraints such as drought and salinity have a major impact on plant development and crop productivity (Zhu, 2002). A common feature of drought and salt stress is the lower availability of water, due to decrease of soil water potential. In addition, salt generates an ionic stress due to the presence of Na\(^+\) and Cl\(^-\). Perception of drought and salt constraints triggers complex signaling networks, which then induce the adaptive response of plants. Among these networks, various molecular components are involved, including phytohormones, protein kinases and phosphatases, and second messengers like Ca\(^{2+}\), ROS, and lipid signaling elements (Munnik and Vermeer, 2010; Huang et al., 2012; Deinelein et al., 2014; Golldack et al., 2014).

Phospholipids are important structural components of cellular membranes but can also play an essential role in the adaptation of plants to abiotic stress (Munnik and Tellerink, 2009; Xue et al., 2009; Munnik and Vermeer, 2010; McLoughlin and Tellerink, 2013). They are modified by enzymes such as phospholipase C (PLC) and D (PLD), and by lipid-kinases, such as diacylglycerol kinase (DGK), PA kinase and various phosphoinositide kinases (Meijer and Munnik, 2003). These modifications produce important second messengers that regulate various plant responses.

Phosphatidylinositol 3-kinase (PI3K) phosphorylates the D-3 position of the inositol ring of phosphoinositides. In mammals, three distinct PI3K classes (I–III) can be distinguished, differing in gene structure, enzyme regulation, and substrate preference. Class III PI3Ks are homologous to the yeast VPS34, which uses PI as a sole substrate to produce PI3P (Backer, 2008). VPS34 promotes membrane fusion and vesicle trafficking by recruiting PI3P-binding proteins to membranes. VPS34 is associated with different proteins, forming distinct protein complexes, including the regulation of mTORC1 (Target of rapamycin complex 1) that monitors the nutritional status of the cell (Backer, 2008; Ktistakis et al., 2012; Robaglia et al., 2012).

Higher plants only contain VPS34-like PI3Ks (Lee et al., 2010). In Arabidopsis, PI3K activity is encoded by a single gene (At1g60490), which is important for pollen development...
(Welters et al., 1994; Lee et al., 2010). Pollen grains harboring a 
pi3k KO allele display abnormal germination (Lee et al., 2008b; 
Gao and Zhang, 2012), preventing the acquisition of homozygous 
pi3k KO mutants. PI3P has been detected in vacuolar mem-
branes and in late endosomal and pre-vacuolar compartments, 
indicating that, like in mammals and yeast, plant PI3K is involved 
in vesicle trafficking and membrane biogenesis (Voigt et al., 2005; 
Vermeer et al., 2006; Lee et al., 2010; Simon et al., 2014). PI3P 
is considered as a second messenger that recruits PI3K-binding 
proteins to membranes (Meijer and Munnik, 2003; Van Leeuwen 
et al., 2004; Wywial and Singh, 2010). Numerous data indicate 
that PI3K and its product PI3P are involved in plant responses 
to drought and salt stress. PI3P participates in stomata closure 
response to abscisic acid (ABA) and ROS production in guard 
cells (Jung et al., 2002; Park et al., 2003; Choi et al., 2008). Also 
NADPH oxidase endocytosis leading to ROS production upon 
ionic stress is triggered by PI3K via PI3P (Leshem et al., 2007).

In plants, PI3P can be further phosphorylated by PI3P-5 kinase 
(FAB) to produce PI(3,5)P2 (Meijer et al., 1999; Munnik and 
Nielsen, 2011; Gao and Zhang, 2012).

Wortmannin (Acaro and Wymann, 1993) and LY294002 
(Vlahos et al., 1994) are pharmaceutical PI3K inhibitors that 
have frequently been used to decipher the function of PI3Ks 
and their products in various mammalian and yeast systems. 
LY294002 is derived from the flavonoid quercetin and competes 
with ATP and binds to a Lys residue in the ATP-binding pocket 
of PI3Ks (Walker et al., 2000). As such, LY294002 has been 
described to inhibit PI3K-related kinases such as TOR (Target 
of rapamycin) and DNA-PK (DNA-dependent protein kinase; 
Brunner et al., 1996), but also other protein kinases, such as Casein 
Kinase 2 (Gharbi et al., 2007). Nevertheless LY294002 is consid-
ered to be a more selective PI3K inhibitor (Walker et al., 2000; 
Jung et al., 2002; Templeton and Moorhead, 2005).

In plants, Wortmannin and LY294002 have also been used to 
show the involvement PI3K and PI3P (Jung et al., 2002; Park 
et al., 2003; Jallais et al., 2006; Vermeer et al., 2006; Leshem et 
al., 2007; Choi et al., 2008a; Takač et al., 2012, 2013). 
LY294002 was shown to inhibit stomatal closing induced by ABA, 
polar tip-growth of root hairs, and chloroplast accumulation in 
response to blue light (Jung et al., 2002; Lee et al., 2008a; Aggarwal 
et al., 2013). At the subcellular level, LY294002 blocks endocytosis 
and vacuolar trafficking and inhibits auxin-mediated ROS gen-
eration (Etxeberria et al., 2005; Joo et al., 2005). Despite LY294002’s 
frequent use, only few reports have shown an effect on PI3K 
activity in vitro (Jung et al., 2002; Joo et al., 2005), yet none in vivo.

In response to water stress, plants accumulate organic osmo-
lytes such as amino acids, sugars and polyamines, which 
play key roles in decreasing the cellular osmotic potential but 
also in preventing the aggregation and/or precipitation of macro-
molecules following the decrease of water availability (Slama 
et al., 2015). Among these organic osmolites, free proline is 
well known to rapidly increase upon water constraints (Szabados 
and Savouré, 2010; Liang et al., 2013). Proline levels represent 
a delicate balance between biosynthesis and catabolism. Proline 
is synthesized from glutamate by a two-step reaction (Szabados 
and Savouré, 2010; Liang et al., 2013). First, glutamate is reduced 
to glutamyl-5-semialdehyde (GSA) by the bifunctional enzyme 
pyrroline-5-carboxylate synthetase (P5CS). GSA is then sponta-
aneously converted to P5C, which is then reduced to proline by 
P5C reductase (P5CR). The rate-limiting enzyme of the biosyn-
thetic pathway is P5CS, which is encoded by two genes P5CS1 
and P5CS2 in A. thaliana. These two isoforms play distinct roles 
during development and stress responses (Székely et al., 2008). 
Under normal growth conditions, proline biosynthesis occurs in 
the cytosol and is mainly under the control of P5CS2. P5CS2 has 
been shown to be expressed in dividing cells, in meristematic and 
reproductive tissues (Strizhov et al., 1997; Székely et al., 2008; 
Mattioli et al., 2009). Upon salt stress and drought, proline accumu-
lation is dependent of P5CS1 expression (Savouré et al., 1995; 
Yoshib et al., 1995). P5CS1 has been shown to be localized in 
chloroplasts upon water stress by Székely et al. (2008).

Upon relief from stress, proline is rapidly oxidized in mito-
chondria by a two-step reaction. First, proline is oxidized by 
proline dehydrogenase (ProDH) to form P5C, which is then oxi-
dized to glutamate by P5C dehydrogenase (P5CDH). ProDH is 
the rate-limiting enzyme for proline catabolism and is encoded 
in Arabidopsis by two genes, ProDH1 (also named ERD5) and 
ProDH2 (Servet et al., 2012). ProDH1 is considered as the main 
isoform, ProDH2 being weakly expressed (Kiyosue et al., 1996; 
Funck et al., 2010). Under either salt or drought stress, ProDH1 
expression is repressed allowing proline accumulation (Funck 
et al., 2010). On the opposite, when stress is relieved, ProDH1 
expression is triggered leading to proline degradation in mito-
chondria (Kiyosue et al., 1996; Verbruggen et al., 1996).

Proline accumulation in response to water stress is not only 
important for osmotic adjustment, but also as scavenger for 
reactive oxygen species (ROS) and molecular chaperone to sta-
bilize proteins, antioxidant enzymes and membrane structures 
(Szabados and Savouré, 2010; Liang et al., 2013). Proline is also 
considered as a source of energy, which may be important upon 
stress recovery (Szabados and Savouré, 2010; Liang et al., 2013; 
Kavi Kishor and Sreenivasulu, 2014).

As proline accumulation and degradation result from a tight 
regulation of its metabolism, deciphering the signaling networks 
involves is of prime importance. Activation of proline biosynthe-
sis is linked to both ABA mediated-signal transduction (Strizhov 
et al., 1997; Abrahám et al., 2003) and ABA-independent sig-
naling (Savouré et al., 1997; Sharma and Verslues, 2010). P5CS1 
expression was also shown to be positively regulated by ROS, act-
ing as intermediate in ABA-mediated proline accumulation, while 
ProDH activity was repressed (Yang et al., 2009).

In Arabidopsis, lipid signaling components are involved in 
the regulation of P5CS1 expression. Under normal growth con-
dition, PLD negatively regulates P5CS1 expression, preventing 
proline accumulation (Thierry et al., 2004). Upon ionic but not 
osmotic stress, PLC triggers P5CS1 expression leading to proline 
accumulation (Parre et al., 2007). P5CS1 up-regulation by PLC 
involves Ca++ as a second messenger, which acts as a molecular 
switch to trigger downstream signaling events (Parre et al., 2007). 
Expression of both ProDH genes is regulated by bZIP transcrip-
tion factors (Weltmeier et al., 2006; Hanson et al., 2008). After 
dark treatment or in response to hypoosmolarity stress, ProDH1 
expression is induced by the heterodimer bZIP53/bZIP10 which 
recognizes the ACTCAT regulating sequence in ProDH1 promoter.
Here, the role of PI3K in the regulation of proline accumulation was investigated. Using a pharmacological and biochemical approach, we show that the decrease of PI3P by LY294002 treatment correlated with lower proline accumulation upon salt stress in Arabidopsis seedlings. The decrease of proline content was associated with both the reduction of PSCTI transcript and protein levels and the induction of ProDH1 transcript and protein levels. In a reverse genetic approach, using a hemizygous pi3k mutant, a similar pattern of ProDH1 expression was found as WT seedlings treated with LY294002. During normal growth condition, a strong expression of ProDH1 is detected in WT seedlings treated with LY294002 as well as in pi3k hemizygous mutant. These data suggest that a signaling pathway involving PI3P participates to the regulation of proline metabolism in normal growth condition and in response to salt stress through the repression of proline catabolism. A detailed metabolite profiling analysis was conducted to search for other compounds regulated by PI3P. Interestingly, raffinose exhibited a similar pattern of accumulation as proline in the presence of LY294002. In addition, hydrophobic- and aromatic amino acid contents strongly increased in presence of LY294002.

**MATERIALS AND METHODS**

**PLANT MATERIAL**

Arabidopsis (Arabidopsis thaliana) Heynh, ecotype Columbia Col-0 as wild-type (WT) and pi3k hemizygous mutant from GABI library (GK_418H02-018138) were used. In the hemizygous pi3k mutant (PI3K/pi3k), T-DNA insertion is located in the fifth exon of one allele of the gene (Lee et al., 2008b). WT seeds were sown on 0.5 x 5 Murashige and Skoog (MS) solid medium (0.8% agar) in 14-cm-diameter Petri dishes as described previously (Parre et al., 2007). pi3k mutant seeds were sown on 0.5 x MS solid medium supplemented with 19.2 μM sulfadiazine (dissolved in DMSO) in order to select hemizygous plants versus WT homozygous plants. After 16 h at 4 °C to raise dormancy, seeds were germinated and grown under continuous light with an intensity of 90 μmole photons m⁻² s⁻¹ for 12 days at 22°C.

**STRESS AND PHARMACOLOGICAL TREATMENTS**

Twelve-days-old seedlings were removed from 0.5 x MS agar plates and put onto liquid 0.5 x MS medium (control) supplemented with either 200 mM NaCl or 400 mM mannitol. After different incubation times, seedlings were collected and immediately frozen in liquid nitrogen and stored at −80°C prior analysis.

For pharmacological treatments, seedlings were pre-treated for 1 h in 0.5 x MS liquid medium with various concentrations of LY294002 dissolved in DMSO or with the same amount of DMSO as a control. Seedlings were thereafter transferred for 3 h or 24 h onto 0.5 x MS liquid medium alone (control), or supplemented with either 200 mM NaCl or 400 mM mannitol and with the same amount of DMSO or LY294002 as for the pre-treatment.

**PROLINE DETERMINATION**

Free proline contents were measured using L-proline as a standard according to Bates et al. (1973).

**PHOSPHOLIPIDS ANALYSIS**

For practical reasons this experiment was performed on 3- to 5-days-old seedlings into a 2 mL Eppendorf tube containing 200 μL of 2.5 mM MES/KOH buffer (pH 5.7) and 1 mM KCl. In order to label phospholipids, 10 μCi of 32P-inorganic phosphate were added in each tube and incubated overnight (Munnik and Zarza, 2013). Either 100 μM LY294002 or the same amount of DMSO for the control were then added for 1 h of pre-incubation. Then a volume of 2.5 mM MES/KOH buffer (pH 5.7), 1 mM KCl buffer with 400 mM NaCl was added into the tube to reach a final concentration of 200 mM NaCl. For control condition, an equivalent volume of MES/KCl buffer was added. Treatments were stopped just after the addition of NaCl (0 h), or after 30 min or 3 h, by adding perchloric acid (5% w/v, final concentration), and after 10 min shaking the total solvent was removed. To extract lipids from the seedlings, 400 μL CHCl₃/MeOH/HCl (50/100/1, v/v/v) was added and the mix was vortexed for 10 min. To induce the separation of two phases, 400 μL CHCl₃ and 200 μL 0.9% (w/v) NaCl were added, vortexed 10 s and then centrifuged for 1 min at 10,000 g. The organic lower phase was transferred to a new tube containing 400 μL CHCl₃/MeOH/1M HCl (3/48/47, v/v/v). After shaking and centrifugation, the upper phase was removed, and 20 μL isopropanol was added to the purified organic phase, which was then dried down in a vacuum centrifuge at 50°C. The residue was dissolved in 100 μL CHCl₃.

Phospholipids were separated as previously described (Munnik et al., 1994, 1995; Munnik and Zarza, 2013) by thin-layer chromatography (TLC) using heat-activated silica gel plates impregnated with a solution of 1% K-oxalate, 2 mM EDTA dissolved in MeOH/H₂O (2/3, v/v) and chromatographed with an alkaline solvent CHCl₃/MeOH/Na₂OH/H₂O (90/70/4/16, v/v/v/v). Radiolabeled phospholipids were visualized and quantified using a phosphoImager.

In order to separate PI3P and PI4P, spots corresponding to the PIP pool was scraped off from the TLC plate and deacylated with 800 μL of mono-methylamine reagent (25% mono-methylamine/MeOH/n-ButOH (42.8/45.7/11.5, v/v/v) at 53°C for 30 min as described in (Munnik et al., 1994, 1996; Munnik, 2013). Samples were centrifuged at 10,000 g for 2 min and the supernatant was collected and dried under a N₂ stream for 30 min and then by rotary evaporation. To remove the fatty-acyl groups, samples were dissolved in 500 μL H₂O and extracted twice with 600 μL n-ButOH/petroleum ether (40–60°C) ethyl for-mate (20/40/1, v/v/v). The aqueous lower phase that contains glycerophosphoinositides (GroPInsP) was dried by rotary evaporation, dissolved in 500 μL H₂O and filtered (0.22 μm).

GroPIns3P and GroPIns4P were separated by anion-exchange HPLC using a Partisil 10-SAX column and a discontinuous gradient of 1.25 M NaH₂PO₄ (pH 3.7) at a flow rate of 1 mL min⁻¹ (Munnik, 2013). Fractions were collected every 30 s and measured for radioactivity by liquid-scintillation counting.

**NORTHERN BLOT ANALYSIS**

Total RNAs were isolated from seedlings ground in liquid nitrogen using the guanidinium thiocyanate-CsCl purification method (Sambrook et al., 1989). RNAs were separated by electrophoresis in a 1.2% agarose-formaldehyde gel. After transfer to nylon membrane, RNAs were fixed by UV cross-linking. Membranes
were hybridized at 65°C with either specific 3’UTR region of
AtP5CS1 or with full length of AtProDH1 according to Church
and Gilbert (1984). The fragments were labeled with 32P-dCTP
using Ready-To-GoTM DNA labeling beads. Before hybridiza-
tion, membranes were stained with methylene blue as a con-
trast for RNA loading and transfer. The hybridization signals
were quantified using a PhosphorImager (Amersham Biosciences,
USA).

**QUANTITATIVE RT-qPCR ANALYSIS**

Total RNAs were extracted following the protocol of the RNeasy
Plant Mini Kit (Qiagen) from around 100 mg of powder obtained
after grinding a pool of seedlings. After treatment with the
RNase-free DNase (Fermentas), 1.5 μg of total RNA were reverse-
transcribed by Revertaid™ Reverse Transcriptase (Fermentas)
using 1 μM oligo(dT) following the manufacturer instructions.
The resulting first-strand cDNA was 20-fold diluted and used as
the template for real-time quantitative PCR (RT-qPCR) ampli-
fication performed on a MasterCycler® ep realplex thermocycler
(Eppendorf) with Maxima® SYBR Green/ROX qPCR Master Mix
(Thermo Scientific) following the manufacturer protocol. Each
reaction was performed with 5 μL diluted cDNA sample in a
total reaction volume of 15 μL. The relative expression of
P5CS1 (At2g39800), ProDH1 (At3g30775) and P13K (At1g60490) genes
were determined using specific primers (Supplementary Table 1).
Expression levels of the different genes were standardized to APT1
(At1g27450) used as a standard reference. The applied RT-qPCR
program was 2 min at 95°C, 40 cycles with 15 s at 95°C, 30 s at
57°C and 30 s at 72°C followed by 15 s at 95°C, 15 s at 55°C,
a gradual temperature rise of 20 min to 55°C at 95°C associ-
ated with a streaming of the plate, followed by 15 s at 95°C. The
expression level of each gene was calculated using the following
equation: 2(ΔΔCt)gene × 100.

**GEL ELECTROPHORESIS, ELECTRO-BLOTTING AND IMMUNOLOGICAL
DETECTION**

Proteins were extracted as described in Martínez-García et al.
(1999) separated by SDS-PAGE (Laemmli, 1970) and transferred
electrophoretically to a nitrocellulose membrane in a solution of
48 mM Tris, 39 mM glycine, 0.04% (w/v) SDS and 20% (v/v)
ethanol at 50 mA for 1 h. For immunodetection, the nitrocellu-
lose membrane was incubated in TBS with 0.05% (v/v) Tween 20
(TBS-T) and 5% non-fat dry milk for 1 h at 4°C and then in TBS-
T with 0.1% (v/v) rabbit antiserum for 16 h at room temperature.
Antiserums were obtained by immunization of rabbits with either
P5CS or ProDH recombinant proteins (Thiery et al., 2004). Blots
were washed with TBS-T. Detection was performed with an ECL
assay using horseradish peroxidase-conjugated secondary anti-
bodies (GE Healthcare). Equal protein loading and integrity of
protein samples were verified by Ponceau S red staining of the
blot membrane.

**METABOLITE PROFILING USING GC-MS AND METABOLIC DATA
PROCESSING**

Three independent samples of 12-days-old seedlings from each
genotype treated during 24 h in different conditions were col-
clected, and the equivalent of 50 mg of powder of each samples
were used to perform the extraction and further metabolomics
analysis. Extraction, derivatization, analysis, and data processing
were performed according to Fiehn (2006). Metabolites were ana-
alyzed by GC-MS 3 h after derivatization. One microliter of the
derivatized samples was injected in splitless mode on an Agilent
7890A gas chromatograph coupled to an Agilent 5975C mass
spectrometer. The column was an Rtx-5SilMS from Restek (30 m
with 10-m InterGuard column). The liner (Restek 20994) was
changed before each series of analyses, and 10 cm of column was
removed. The oven temperature ramp was 70°C for 7 min
then 10°C/min to 325°C for 4 min (run length 36.5 min). The
helium constant flow was 1.5231 mL/min. Temperatures were as
follows: injector, 250°C; transfer line, 290°C; source: 250°C; and
quadrupole, 150°C. Samples and blanks were randomized. Amino
acid standards were injected at the beginning and end of the analy-
sis to monitor the derivatization stability. An alkane mix (C10,
C12, C15, C19, C22, C28, C32, and C36) was injected in the mid-
dle of the queue for external calibration. Five scans per second were
acquired.

Metabolites were annotated, and their levels on a fresh
weight basis were normalized with respect to the ribitol internal
standard.

Raw Agilent data files were converted in NetCDF format and
analyzed with AMDIS (http://chemdata.nist.gov/mass-spc/ amdis/).
A home retention index/mass spectra library built from the
NIST, Golm, and Fiehn databases and standard compounds
were used for metabolite identification. Peak areas were then
determined using the Quanlynx software (Waters) after conver-
sion of the NetCDF file to masslynx format. TMEV (http://www.
tm4.org/mev.html) was used for all statistical analysis. Univariate
analysis by permutation (One-Way and Two-Way ANOVA) was
first used to select the significant metabolites. Multivariate analy-
sis (hierarchical clustering and principal component analysis) was
then performed on this subset.

**RESULTS**

LY294002 AFFECTS PROLINE ACCUMULATION ONLY IN RESPONSE TO
SALT TREATMENT

To investigate whether PI3K is involved in the regulation of
proline metabolism in response to ion and/or hyperosmotic
constraints, the effect of LY294002 on proline accumulation
was assessed in 12-days-old Arabidopsis seedlings subjected to
either 200 mM NaCl or 400 mM mannitol for 24 h. As shown in
Figure 1A, typically a 5- to 6-fold accumulation of pro-
line is observed in Arabidopsis seedlings treated with either
NaCl or mannitol in comparison to the control seedlings.
Interestingly, while LY294002 had no effect on the proline lev-
el in control seedlings or seedlings stressed with mannitol
(Figure 1), 40% less proline accumulated in the LY294002-
treated seedlings upon salt stress. When increasing concen-
trations of LY294002 were applied, proline levels decreased in a
dose-dependent manner in plants stressed with NaCl, with a
maximum effect observed for 100 μM LY294002 (Supplementary
Figure 1). In contrast, no effect on the proline levels of con-
trol or mannitol-stressed seedlings where found, whatever con-
centration of LY294002 (Figure 1 and Supplementary Figure
1). These results show that LY294002 negatively regulates
proline accumulation in response to salt stress but not to mannitol.

LY294002 REDUCES THE LEVEL OF PI3P

In order to characterize the inhibitory effect of LY294002 on PI3K activity, in vivo PI3P levels were measured. For practical reasons, 6-days-old seedlings were used. These seedlings accumulated slightly less proline than the 12-days-old seedlings after NaCl treatment but the results were consistent, indicating that the younger seedlings perceived and responded well to salt stress (data not shown).

Seedlings were $^{32}$P$_{3}$-labeled overnight and the lipids extracted and separated by TLC (Supplementary Figure 2). Phosphoinositides (PI, PIP, and PIP$_{2}$) were quantified using PhosphoImaging (Figure 2A). PI, a structural phospholipid of membranes, represented $\sim$80–90% of the total $^{32}$P-labeled phospholipids, and its levels remained fairly constant throughout our experiments. PIP and PIP$_{2}$ are minor lipid constituents and accounted for $\sim$0.05–0.15% of the total phospholipids, respectively. PIP$_{2}$ progressively increased upon salt stress, reaching a 3-fold increase at 3 h compared to control seedlings, while no significant effects of salt stress were found for PIP levels. Interestingly, LY294002 treatment caused a slight but significant decrease in PIP$_{2}$ under control conditions as well as in response to salt stress.

In plants, the PIP pool is composed of 3 isomers, PI3P, PI4P, and PI5P (Munnik and Vermeer, 2010). PI4P is the most predominant PIP species (50–90%), with PI3P and PI5P each accounting for $\sim$5–10% of the PIP pool (Meijer et al., 2001). On TLC, the PIP isomers cannot be separated but by removing their fatty acids and analysing the resulting glycerophosphoinositolphosphates (GroPlnPs) by HPLC, it is relatively easy to distinguish and quantify the GroPlnPs3P from the GroPlnPs4P and GroPlnPs5P. The latter two are rather difficult to separate (Meijer et al., 2001).

So to determine the PI3P levels under our conditions, TLC-separated $^{32}$P-labeled PIP spots from 30 min treated seedlings were scraped off, deacylated and separated by anion-exchange HPLC. At control and salt conditions, PI3P was found to account for $\sim$5% of the PIP pool. Addition of 100 µM LY294002, however, induced a 50% decrease of PI3P, whatever control or stress condition (Figure 2B). These results, and the inhibitory effect of LY294002 on proline accumulation in response to salt stress, are consistent with the involvement of PI3P as a lipid mediator on the regulation of proline metabolism.
LY294002 IMPACTS THE EXPRESSION OF GENES INVOLVED IN PROLINE METABOLISM

Proline accumulation is the consequence of a tight regulation of gene expression (Szabados and Savouré, 2010; Liang et al., 2013). We investigated transcript accumulation of two genes involved in proline metabolism, P5CS1 and ProDH1 that encode key enzymes regulating proline biosynthesis and catabolism, respectively. RT-qPCR analysis showed a 17-fold higher regulating proline biosynthesis and catabolism, respectively. RT-proline metabolism, we investigated transcript accumulation of two genes involved in proline metabolism, we aimed for Arabidopsis pi3k T o further unravel the role of PI3K in the regulation of pro-

The role of PI3K on key genes and enzymes involved in proline metabolism was investigated using LY294002. Northern and western blot analysis revealed that LY294002 affected mRNA and protein accumulation in all tested conditions. After 3 h LY294002 treatment, a modest increase of P5CS1 mRNA was observed in control condition while ProDH1 transcript level was higher (Figure 3A). A dramatic effect of LY294002 on both P5CS1 and ProDH1 expression compared to non-treated seedlings was observed at 24 h salt stress. In salt stress seedlings treated with LY294002, P5CS1 steady state transcript level was lower than in non-treated ones while ProDH1 transcript level was higher (Figure 3A). In salt stress seedlings treated with LY294002, P5CS1 transcript accumulation decreased by 60% (Figure 3C) compared to non-treated seedlings. RT-qPCR analysis confirmed the higher ProDH1 transcript accumulation in presence of LY294002 whatever the growth conditions (Figure 3C). Using western blots, LY294002 triggered P5CS accumulation in control seedlings while P5CS level diminished in salt-treated plantlets. In contrast, LY294002 strongly enhanced ProDH accumulation in both control and salt-treated seedlings (Figure 3B).

The lower proline accumulation observed in response to salt stress with LY294002 is correlated with a down-regulation of P5CS1 and up-regulation of ProDH1 at both transcript and protein levels. As LY294002 reduced P13P levels, our data suggest that PI3K is involved in the regulation of proline metabolism.

ProDH1 EXPRESSION IS INDUCED IN pi3k MUTANT

To further unravel the role of PI3K in the regulation of proline metabolism, we aimed for Arabidopsis pi3k KO mutants. Unfortunately, however, homozygous pi3k mutants are not viable (Lee et al., 2008b; Gao and Zhang, 2012). To partly resolve this issue, we selected sulfadiazine-resistant seedlings to get hemizygous pi3k mutant (Leshem et al., 2007; Lee et al., 2008a,b). Segregation analysis indicated a 1:1 ratio in sulfadiazine resistant and sensitive seedlings, respectively (data not shown), due to the male gametophytic defect (Lee et al., 2008b). Therefore PI3K/pi3k hemizygous mutants were selected and further analyzed (Figure 4A). PCR-based genotyping of GABL_418H02 pi3k mutants confirmed that no homozygous mutants were obtained from this selection (data not shown).

Expression analysis by RT-qPCR revealed a 25% decrease of steady-state PI3K transcript level in pi3k hemizygous mutants compared to WT (Figure 4B). In this mutant, ProDH1 transcript level was almost 5-fold higher than in WT seedlings in normal growth condition. On the contrary, no difference in P5CS1 transcript level was observed between WT and pi3k hemizygous mutant.

When WT and pi3k hemizygous mutant were subjected to 200 mM NaCl for 24 h, they showed a higher proline accumulation of 18-fold and 11.5-fold, respectively (Figure 5A). pi3k hemizygous mutant showed a lower proline accumulation in response to NaCl. However the proline accumulation was not significantly different from WT seedlings, probably due to the remaining PI3K wild-type allele.

P5CS1 and ProDH1 transcript levels were investigated in pi3k hemizygous seedlings in response to 3 and 24 h of salt stress (Figure 5B). P5CS1 mRNA accumulation was similar in WT and pi3k hemizygous in response to salt stress. An equivalent increase of P5CS1 transcripts was observed at 3 h of stress and a decrease at 24 h of stress. Higher ProDH1 transcript levels than WT were

FIGURE 3 | LY294002 impacts ProDH1 steady state transcript and proteins levels. Northern- and western blot analysis were performed on total RNA or proteins extracted, respectively, from 12-days-old seedlings treated with either 0.5x MS alone (control) or supplemented with 200 mM NaCl (NaCl) during 3 h or 24 h with 100 µM LY294002 (+) or DMSO (-) as described in the legend of Figure 1. (A) Total RNA (10 µg) was loaded in each lane and northern blots were probed with DNA fragments specific for AtP5CS1 and AtProDH1. Methylene blue staining of rRNAs is shown as a loading control. (B) Total proteins (20 µg) were separated on an 8% SDS-PAGE gel. Western blots were incubated with specific antibodies directed against AtP5CS1 and AtProDH proteins. Detection of the immunolabeled proteins was done by autoradiography using an ECL kit. Membranes were stained with Ponceau S Red as control for protein loading (Rubisco). (C) Relative expression levels of P5CS1 and ProDH1 genes measured by real-time quantitative PCR after 24 h treatment as described in (A). Results expressed as percentage compared to APT1 as a reference gene are means ± SD of 3 replicates. Letters indicate statistical differences in P5CS1 (blue letter) or ProDH1 (red letter) gene expression depending on treatment conditions (Two-Way ANOVA test, P < 0.05). The results presented in (A,C) were obtained from two independent experiments.
LY294002 AFFECTS SEEDLING METABOLOME

To investigate other changes induced by LY294002, we compared the metabolite profiles of 12-days old WT seedlings treated with LY294002 or DMSO upon control and salt stress conditions. Hierarchic clustering analysis indicated that DMSO did not have any significant effect on the relative metabolites contents, indicating that the difference in the metabolite patterns obtained with LY294002 is an effect of the inhibitor (Figure 6). Treatment for 24 h of salt stress significantly modified the amounts of several metabolites. Relative amounts of sucrose, ribose and maltose as well as proline, serine and raffinose increased in response to salt stress (Figures 6, 7B). On the contrary, the amounts of some other sugars, like galactose, mannose, trehalose and xylose (Figure 6) and glucose-6-phosphate (Glucose-6-P) and fructose-6-phosphate (Fructose-6-P) decreased in response to salt stress (Figure 7A). LY294002 reduced the level of proline content almost by half (Figure 7B), in accordance with our previous results (Figure 1 and Supplementary Figure 1). Interestingly, two other compatible osmolytes, serine and raffinose, exhibited an accumulation pattern similar to that of proline, i.e., higher accumulation in response to salt stress and lower when LY294002 is added (Figure 7B). Surprisingly, LY294002 addition had a strong impact on some amino-acid levels whatever the treatment. The

observed in pi3k mutant in all tested conditions whether the seedlings were subjected to salt stress or not (Figure 5B). In pi3k hemizygous mutant, ProDH protein levels were stronger than in WT at both 3 and 24 h salt stress (Figure 5C). These results indicate that pi3k hemizygous seedlings are able to respond to salt stress, triggering P5CS1 gene expression and proline accumulation, although the transcript level of ProDH1 gene is increased. Altogether, our data indicate that the higher level of ProDH1 transcripts is correlated with higher ProDH amount in pi3k mutant. These data are consistent with those obtained with LY294002 treatment (Figure 3), where LY294002 induced higher ProDH1 transcripts and proteins. These results further strengthen the participation of a PI3K-mediated pathway regulating proline catabolism.

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most dramatic effect being for tyrosine with an almost 100-fold increase in the presence of LY294002, in either control condition or NaCl stress (Figure 7C). Similarly, a 10–20-fold increase was observed for lysine, leucine, isoleucine and phenylalanine and a 4-fold increase for valine in response to LY294002.

Thus, the strong increase of those aliphatic and aromatic amino acids in presence of LY294002 suggests that PI3K, and/or its product PI3P, negatively regulates their metabolism through inhibition of their synthesis and/or promotion of their catabolism.

DISCUSSION

In this paper, we investigated the effect of the PI3K inhibitor, LY294002, on the response of Arabidopsis seedlings to salt stress. LY294002 was found to reduce PI3P levels by 50% and to dramatically decrease the accumulation of proline upon salt stress. The latter was a consequence of lower transcript- and protein levels for P5CS1 and higher transcript and protein levels for ProDH1. In the pi3k hemizygous mutant line, as also observed for WT seedlings treated with LY294002, an up-regulation of ProDH1 expression was found, suggesting that PI3K and its product PI3P are involved.
in a pathway repressing ProDH1 expression. Metabolomic profiling of Arabidopsis seedlings in response to salt stress showed that LY294002 reduced the amount of raffinose, another compatible osmolyte, and strongly increased the amount of free aliphatic and aromatic amino acids.

**NaCl stress modified phosphoinositide composition**

PI is not only a structural phospholipid of membranes, but also a precursor of several signaling phosphoinositides that are produced by distinct kinases and phosphatases, which add and remove phosphates at different positions of the inositol ring (Mueller-Roeber and Pical, 2002; Xue et al., 2009; Munnik and Vermeer, 2010; Munnik and Nielsen, 2011). Characterization of the phospholipid composition of overnight 32P-labeled Arabidopsis seedlings showed that salt stress mostly affected the PIP2 pool. This latter is predominantly composed of the PI(4,5)P2 isomer (Munnik, 2013). Some plant systems, in particular Chlamydomonas, have reported on increased PI(3,5)P2 levels (Meijer et al., 1999), but we did not observe this here for Arabidopsis seedlings. Increased PIP2 levels in response to salt and/or osmotic stress have been reported for several plant systems (Pical et al., 1999; DeWald et al., 2001; Zonia and Munnik, 2004; Darwish et al., 2009; Munnik and Zarza, 2013). Part of this PI(4,5)P2 response occurs at the plasma membrane (Van Leeuwen et al., 2007; König et al., 2008), where it is considered to be an important signaling molecule (Munnik and Nielsen, 2011), potentially through initiation of vesicle budding via its interaction with clathrin-adapter proteins. The subsequent formation of clathrin-coated vesicles during salt stress could be a mechanism for the cell to modify the plasma membrane according to water/ion movement (König et al., 2008). E.g., PI(4,5)P2 has been suggested to modulate stomatal aperture in response to water stress by regulating K+ efflux channel (Lee and Lee, 2008).

Alternatively, PI(4,5)P2 can be hydrolysed by PLC to form inositol trisphosphate (IP3) and DAG, which can both be rapidly metabolized into other signaling molecules, e.g., IP6 and phosphatidic acid (Munnik and Vermeer, 2010). PLC has been implicated in stress signaling (Drobak and Watkins, 2000; Tasma et al., 2008; Xue et al., 2009; Munnik and Vermeer, 2010). In addition, Parre et al. (2007) have demonstrated that proline biosynthesis in response to salt stress is regulated by a Ca2+-signature depending on PLC activity.

LY294002 addition had a small effect on the PIP2 pool. Theoretically, as an ATP analog, LY294002 could inhibit other enzymes (Davies et al., 2000; Walker et al., 2000; Gharbi et al., 2007), but to our knowledge there is no report regarding such an effect on PIP 5-kinases. Another explanation could be that vesicles containing the LY294002-sensitive PIP2 pool require the down-stream synthesis of PIP2, for example, to fuse with or pinch-off from membranes.

Minor changes in PIP levels were measured in Arabidopsis seedlings upon salt stress. This pool is a mixture of 3 isomers, i.e., PIP3, PIP4P, and PIP5P. The latter isomer is thought to result from PI(3,5)P2 dephosphorylation and is, like its precursor, present at very low concentrations (Meijer et al., 2001; Munnik, 2013). Since the PIP isomers cannot be separated by TLC, we removed the fatty acids by chemical deacylation and analyzed the resulting GroPInsP isomers by anion-exchange chromatography. As such, we found that only 5% of the PIP pool accounted for PIP3. The majority of the pool was composed of PIP4P pool (>90%). PIP5P levels were not determined as they were extremely difficult to separate from the 4-isomer (Meijer et al., 2001). LY294002 caused the PIP3 pool to be reduced by 50%. No variation was observed in response to salt stress. Earlier, Meijer et al. (2001) reported an increase in PIP3 after 5 min of 300 mM NaCl but this was in Chlamydomonas, which seems to exhibit a different PI3P- and PI(3,5)P2 metabolism compared to higher plants. To our knowledge, there is no other data available on the effect of salt on PIP3 levels. Nevertheless, several studies have indicated a role for PI3P in the plant’s response to salt or water stress on the basis of PI3K inhibitors. As such, PI3P has been implicated in the regulation of stomatal closure in response to ABA (Jung et al., 2002). Leshem et al. (2007) have demonstrated that PI3P triggers the endocytosis of NADPH oxidase located at the plasma membrane under salt stress and this was implicated in the formation of ROS, which are important signaling molecules for plants to cope with salt stress (Leshem et al., 2006; Ben Rejeb et al., 2014). Phosphoinositides recruit proteins through specific phosphoinositide-binding domains to particular membranes (Van Leeuwen et al., 2004; Banerjee et al., 2010; Munnik and Nielsen, 2011). As such, the interaction between PI3P and the immunophilin ROF1 could participate in the plant’s response to salt stress too (Karali et al., 2012). In addition, PI3P can participate in the regulation of vesicular trafficking and vacuole formation by recruiting proteins such as VTI11 and EPS1N that are involved in membrane fusion (Lee and Lee, 2008; Lee et al., 2010; Zheng et al., 2014a,b).

**PI3P is involved in the repression of proline catabolism**

Proline accumulation is a well-known plant response to salt, and more generally, to water stress (Szabados and Savouré, 2010; Gupta and Huang, 2014). We found that LY294002 reduced the proline accumulation in response to salt stress. This could be explained by repression of proline biosynthesis and induction of proline catabolism genes both at the mRNA and protein levels. The strong induction of ProDH1 expression and the accumulation of the corresponding protein were correlated with the inhibitory effect of LY294002 on PI3P levels in both control and stress conditions, suggesting a role of PI3P in the inhibition of proline catabolism whatever the conditions. This hypothesis was confirmed by studies on pi3k mutant. ProDH1 expression was higher in pi3k mutant than in the WT upon salt stress but also in normal growth condition. On the contrary, P5CS1 expression while being diminished by LY294002 was not affected in pi3k mutant, suggesting that LY294002 may also act on other protein kinase activity involved in signaling pathway regulating P5CS1 expression. We previously found evidence that P5CS1 expression in response to salt stress fell under the regulation of PLC activity (Parre et al., 2007). Here, seedlings treated with LY294002 had a lower PIP2 response with salt. Maybe the lower availability of PIP2 as PLC substrate contributes to the reduced P5CS1 expression. The comparison between seedlings treated with LY and pi3k mutant showed differences in P5CS expression in contrast to ProDH expression. This may be due to the fact that LY294002 could
have additional effects like inhibiting other kinases. Another possibility is that adaptation to long-term decrease in PI3K activity could occur in the hemizygous pi3k mutant whereas the effect of LY294002 is more sudden and could change some of the plant stress responses.

Lee et al. (2008b) indicated that the pi3k mutant is strongly impaired in its pollen development, leading to the inability to obtain homozygous mutant. Our segregation analysis also showed a gametophytic defect in pi3k mutant, which explained why only pi3k hemizygous mutant could be obtained. pi3k mutant showed a reduced expression of the complete gene, in agreement with Leshem et al. (2007) and to the lethal phenotype of the reported antisense transformation (Welters et al., 1994). Consequently, genetic and biochemical analyses of PI3K are very difficult to assess. The development of promoter inducible lines may be useful to further investigate the role of PI3K in plant stress adaptation.

ProDH1 expression is under the control of bZIP10 and bZIP53 transcription factors (Satoh et al., 2004; Weltmeier et al., 2006; Dietrich et al., 2011; Veerabagu et al., 2014). Their expression and activity is regulated by various abiotic constraints, and also by the nutrient status of the plant (Weltmeier et al., 2009). In mammals, nutrient deficiency induces ProDH gene expression as a consequence of mTOR complex inactivation (Liao et al., 2008). VPS34 has been shown to participate in the regulation of mTOR upon nutrient stress (Backer, 2008). TOR is a protein kinase that possesses a catalytic domain with strong homology to the kinase domain of PI3K, and has also been shown to be sensitive to LY294002 (Brunn et al., 1996). In Arabidopsis, it is therefore possible that LY294002 inhibits TOR kinase too and as a result ProDH. Nevertheless, the fact that the ProDH1 increase is also observed in the pi3k hemizygous mutant background supports a direct involvement of the PI3K pathway in repressing ProDH1. TOR could even be a downstream component of PI3K and as such participate in the regulation of ProDH expression. This has already been observed in other eukaryotes (Liao et al., 2008).

**BESIDES PROLINE ACCUMULATION, LY294002 AFFECTS PLANT METABOLISM**

Salt stress affects several metabolites, among them sugars such as hexose-phosphates, disaccharides and raffinose family oligosaccharides (RFO). Sucrose, maltrate and raffinose accumulated in response to salt stress, whereas Glu-6-P and Fru-6-P decreased. Variations of these sugar contents are well-conserved responses among various plant species upon salt stress (Kempa et al., 2008; Sanchez et al., 2008). These sugars are derived from photosynthesis activity. They have a key role in osmoprotection, in ROS scavenging and as a source of carbon storage (Keuyen et al., 2013; Gupta and Huang, 2014). Sugar could also be considered as signaling molecules regulating gene expression and triggering specific responses to abiotic stress (Keuyen et al., 2013). Raffinose is the only sugar whose accumulation was reduced by 50% in salt-stressed Arabidopsis seedlings treated with LY294002. Similar to proline, raffinose is also considered as a compatible solute for plant cells (Valluru and Van den Ende, 2011; Van den Ende, 2013). In order to determine whether the biosynthesis of raffinose upon salt stress is dependent of the same signaling pathway as proline, it will be of interest to investigate the regulation of raffinose metabolism in pi3k mutant background.

The accumulation of several amino acids other than proline was observed upon salt stress. Aliphatic (Val, Leu, Ileu) and aromatic (Phe, Tyr) amino acids increased in response to salt stress (our study; Kempa et al., 2008; Sanchez et al., 2008), but also after an extended period of darkness (Usadel et al., 2008). These authors suggested that these might be due to induction of protein catabolism and/or remobilization of nitrogen sources. Surprisingly, a dramatic increase of these amino acids was observed in seedlings treated with LY294002 grown in normal conditions. Further experiments will be required to determine the role of PI3K signaling in the regulation of amino acid metabolism.

In conclusion, our results strongly indicate that a signaling pathway involving PI3K and its product PI3P is involved in the repression of proline catabolism upon salt stress. Identifying specific PI3P targets will allow to decipher whether TOR is an intermediate signaling component in the regulation of ProDH in plants.

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**SUPPLEMENTARY MATERIAL**

The Supplementary Material for this article can be found online at: http://www.frontiersin.org/Journal/10.3389/fpls.2014.00772/abstract

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Leprince et al.

Regulation of proline catabolism by Phosphatidylinositol 3-kinase


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.

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