Plant Glyoxylate/Succinic Semialdehyde Reductases: Comparative Biochemical Properties, Function during Chilling Stress, and Subcellular Localization

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Plant NADPH-dependent glyoxylate/succinic semialdehyde reductases 1 and 2 (cytosolic GLYR1 and plastidial/mitochondrial GLYR2) are considered to be of particular importance under abiotic stress conditions. Here, the apple (Malus × domestica Borkh.) and rice (Oryza sativa L.) GLYR1s and GLYR2s were characterized and their kinetic properties were compared to those of previously characterized GLYRs from Arabidopsis thaliana [L.] Heynh. The purified recombinant GLYRs had an affinity for glyoxylate and succinic semialdehyde, respectively, in the low micromolar and millimolar ranges, and were inhibited by NADP⁺. Comparison of the GLYR activity in cell-free extracts from wild-type Arabidopsis and a glyr1 knockout mutant revealed that approximately 85% and 15% of the cellular GLYR activity is cytosolic and plastidial/mitochondrial, respectively. Recovery of GLYR activity in purified mitochondria from the Arabidopsis glyr1 mutant, free from cytosolic GLYR1 or plastidial GLYR2 contamination, provided additional support for the targeting of GLYR2 to mitochondria, as well as plastids. The growth of plantlets or roots of various Arabidopsis lines with altered GLYR activity responded differentially to succinic semialdehyde or glyoxylate under chilling conditions. Taken together, these findings highlight the potential regulation of highly conserved plant GLYRs by NADPH/NADP⁺ ratios in planta, and their roles in the reduction of toxic aldehydes in plants subjected to chilling stress.

Keywords: abiotic stress, aldehyde detoxification, mutants, overexpression, redox homeostasis, substrate promiscuity

INTRODUCTION

Under stress conditions such as chilling, drought and salinity, toxic aldehydes can accumulate in plants and interact with DNA, lipids and proteins, or influence the transcription of stress-related genes, thereby causing cellular and developmental problems (Weber et al., 2004; Kotchoni et al., 2006; Yamauchi et al., 2011; Mano, 2012; Biswas and Mano, 2015; Srivastava et al., 2017; see references therein). Altering activities of enzymes that modify the aldehyde chemical grouping has been shown to affect plant tolerance to various stresses (Oberschall et al., 2000; Bouché et al., 2003;
Sunkar et al., 2003; Zarei et al., 2016), indicating that detoxification of aldehydes is important for maintaining plant health.

Succinic semialdehyde (SSA) is a mitochondrially generated intermediate in the metabolism of 4-aminobutyrate (GABA), which accumulates in response to abiotic stresses (Bown and Shelp, 1997; Shelp et al., 1999; Kinnersley and Turano, 2000). SSA is typically oxidized via SSA dehydrogenase (Tuin and Shelp, 1994; Bouché et al., 2003), but evidence is available for the generation of 4-hydroxybutyrate (GHB) from SSA in response to hypoxia, high light, salinity, drought and chilling (Allan et al., 2003, 2008, 2012; Breitkreuz et al., 2003; Fait et al., 2005). Glyoxylate can be generated in various subcellular compartments and biochemical processes: the peroxisome during the photosynthetic or non-photosynthetic synthesis of serine (Bauwe et al., 2010; Hoover et al., 2013; Ros et al., 2014); the glyoxysome during fatty acid catabolism (Kunze et al., 2006); the cytosol during the interconversion of organic acids (Eprintsev et al., 2015); and the endoplasmic reticulum during purine catabolism (Werner and Witte, 2011). Glyoxylate, the precursor of glyoxylate in photosynthesis, is known to accumulate under hypoxic conditions, and both glyoxylate and glyoxylate accumulate with the suppression of glycolate oxidase (Narsai et al., 2009; Lu et al., 2014).

Several mechanisms can be involved in the detoxification of glyoxylate: transamination to glycine (e.g., GABA transaminase); decarboxylation to formate; oxidation to oxalate; and, reduction to glycolate (Givan and Kleczkowski, 1992; Bauwe et al., 2010; Peterhansel et al., 2010; Shelp et al., 2012a; Igamberdiev and Eprintsev, 2016; see references therein). Our laboratory has biochemically characterized two NADPH-dependent glyoxylate reductases (EC 1.1.1.79) from Arabidopsis thaliana (L.) Heynh. (GLYR1, GenBank accession no. AAK94781; GLYR2, (GenBank accession no. AAP42747) that efficiently convert glyoxylate into glycolate (Hoover et al., 2007a,b; Simpson et al., 2008). Notably, they also convert SSA into GHB, albeit with lower catalytic efficiency than for the glyoxylate to glycolate conversion (2–12 vs. 660–2870 s⁻¹ mM⁻¹, see Table 1 for additional information), and constitutive expression of AtGLYR1 in an SSA dehydrogenase-deficient yeast enables growth on 20 μM GABA and significantly enhances cellular GHB levels (Breitkreuz et al., 2003). Furthermore, glyr1 and glyr2 single mutants of Arabidopsis accumulate less SSA with submergence stress than the wild-type (WT) (Allan et al., 2012). A gaba transaminase/ssa dehydrogenase double mutant of Arabidopsis accumulates both SSA and GHB, and is more sensitive to exogenous SSA or GHB than WT plants (Ludewig et al., 2008). Both WT and glyr1/2 double mutants of Arabidopsis are more sensitive to exogenous SSA than GHB under ambient conditions, suggesting that GHB accumulation in ssa dehydrogenase mutants and during abiotic stress is a response for avoiding SSA toxicity (Mekonnen and Ludewig, 2016). Further research is required to establish whether a differential phenotype is associated with the glyr1/glyr2 double mutants under abiotic stress conditions.

Arabidopsis GLYR1 and GLYR2 activities have strong affinity for NADPH ($K_m = 1.2$–$2.6$ μM; see Table 1 for additional information), regardless of whether glyoxylate or SSA is the substrate, and AtGLYR1 is known to be competitively inhibited by NADP⁺ (Hoover et al., 2007a,b; Simpson et al., 2008). The ratio of NADPH/NADP⁺ in mature leaves of Arabidopsis and tobacco increases from approximately 2 to 3–5 with submergence and heat stresses, and to 8 with chilling (Allan et al., 2008). In the absence of applied stress, the NADPH/NADP⁺ ratio in an NAD KINASE1 overexpression line of Arabidopsis is increased by threefold, and there is a corresponding increase in GHB accumulation (Allan et al., 2012).

Arabidopsis GLYR1 and GLYR2 are moderately expressed throughout the plant, including roots and imbibed seed, but GLYR2 expression is more highly associated than GLYR1 with rosette leaves, which are known to highly express photosynthetic genes (Foyer et al., 2009; Shelp et al., 2012b). With the exception of two studies (Allan et al., 2008, 2012), there is little evidence for the stress-induced expression of GLYRs (Shelp et al., 2012b; Zarei et al., 2017; see references therein). Furthermore, transient GHB accumulation in cold-acclimated Arabidopsis plants (Kaplan et al., 2007) and rice seed germinated under hypoxia (Narsai et al., 2009) seems to be independent of GLYR expression (see Shelp et al., 2012b). Overall, these in vitro and in planta studies support the operation and redox regulation of glyoxylate- and SSA-dependent GLYR activities under abiotic stress conditions, as suggested previously (Allan et al., 2008, 2012).

Early research indicated that NADPH-dependent GLYR activity is located in both cytosol and plastids (Givan et al., 1988; also see review by Givan and Kleczkowski, 1992). Recently, Simpson et al. (2008), Ching et al. (2012), Brikis et al. (2017) established, using various transient and stable expression systems, in combination with appropriate organelle markers, that GLYR1s from apple (Malus × domestica Borkh.) and rice (Oryza sativa L.), as well as Arabidopsis, are exclusively cytosolic, whereas the GLYR2s are localized to both mitochondria and plastids. Apple and rice GLYR2s have not yet been biochemically characterized.

In the present study, we demonstrated that: (i) GLYRs from apple and rice, like those from Arabidopsis, display higher affinity and catalytic efficiency for glyoxylate than for SSA; (ii) the activity of apple GLYRs is inhibited by NADP⁺; (iii) the growth of plantlets or roots of various Arabidopsis lines with altered GLYR activity respond differentially to exogenous SSA, GHB or glyoxylate under chilling conditions; and (iv) approximately 85 and 15% of the total GLYR activity in Arabidopsis is present in the cytosol and plastids/mitochondria, respectively. Taken together, these findings highlight the potential roles of GLYRs in the reduction of toxic aldehydes within distinct compartments of the plant cell during exposure to chilling conditions.

**MATERIALS AND METHODS**

**Plant Material, and Gene Expression and Southern Analysis**

*Arabidopsis thaliana* (L.) Heynh. ecotype Columbia (Col-0) was the genetic background for the WT and all transgenic/mutant lines. Seeds were sterilized in a closed container using a...
seeds for 3 days at 4◦C, the plates were transferred to a controlled environment chamber (Enconair, Winnipeg, Canada) and supplied with a 16-h light (Sylvania Model 3500K) period at 60 μmol s⁻¹ m⁻² at 23°C and an 8-h dark period at 21°C.

Total RNA was extracted from 100 to 1000 mg of liquid N₂-frozen Arabidopsis rosette leaves (Zarei et al., 2011), rice (O. sativa L.) leaves (Brauer et al., 2011) and mature apple (Malus × domestica Borkh.) fruit (Gasic et al., 2004). RNA was treated with DNase I using the TURBO DNA-free kit (Applied Biosystems, Austin, TX, United States) according to the manufacturer's manual. One microgram total RNA was used for first-strand cDNA synthesis with Oligo(dT)20 and Superscript III (Invitrogen, Carlsbad, CA, United States) at 50°C according to the manufacturer's protocol. Primers used for quantitative real-time PCR (qPCR) analyses were designed using Primer Express 3 software (Applied Biosystem, Austin TX, United States). Primers with 90–105% efficiency were selected for further analysis.

Quantitative PCR was performed according to standard methods with an iQ5 Multicolor Real-Time PCR Detection system (Bio-Rad Laboratories). The primer sequences used to determine the abundance of GLYR1, GLYR2 and housekeeping transcripts are listed in Supplementary Table S1 (RTatGLYR1-F/R, RTatGLYR2-F/R, RTaEF-1-F/R, and 18S rRNA- F/R). The housekeeping gene ELONGATION

### TABLE 1 | Comparison of the kinetic parameters for purified recombinant GLYRs from apple, rice, and Arabidopsis.

| Varied substrate | Fixed substrate | $K_m$ μM | $V_{max}$ μmol min⁻¹ mg⁻¹ protein | $k_{cat}$ s⁻¹ | $k_{cat}/K_m$ s⁻¹ mM⁻¹ |
|------------------|----------------|----------|----------------------------------|--------------|------------------------|
| MvGLYR1          | Glyoxylate     | 14.1 ± 2.1 | 54.7 ± 6.7                      | 30.9 ± 3.8   | 2230 ± 153             |
|                  | NADPH         | 3.3 ± 0.5 | 39.4 ± 6.1                      | 22.3 ± 3.4   | 6603 ± 278             |
|                  | SSA           | 1133 ± 273 | 8.1 ± 2.9                       | 4.6 ± 1.6    | 5.1 ± 2.3              |
|                  | NADPH         | 7.4 ± 1.9 | 9.8 ± 1.0                       | 5.5 ± 0.6    | 832 ± 166              |
| OsGLYR1          | Glyoxylate     | 23.9 ± 1.9 | 34.3 ± 3.7                      | 19.1 ± 2.1   | 796 ± 33               |
|                  | NADPH         | 1.8 ± 0.5 | 28.2 ± 2.7                      | 15.7 ± 1.5   | 10070 ± 2275           |
|                  | SSA           | 6500 ± 327 | 18.6 ± 3.4                      | 10.3 ± 1.9   | 1.6 ± 0.2              |
|                  | NADPH         | 11.7 ± 3.5 | 15.5 ± 1.5                      | 8.6 ± 0.9    | 833 ± 183              |
| OsGLYR2          | Glyoxylate     | 53.2 ± 11.1 | 66.8 ± 16.5                     | 33.7 ± 7.5   | 636 ± 99               |
|                  | NADPH         | 8.8 ± 2.5 | 28.5 ± 5.9                      | 14.2 ± 2.9   | 1729 ± 201             |
|                  | SSA           | 4003 ± 341 | 43.9 ± 8.5                      | 21.9 ± 4.2   | 5.4 ± 0.9              |
|                  | NADPH         | 36.2 ± 2.1 | 21.4 ± 5.0                      | 10.7 ± 2.5   | 290 ± 51               |
| OsGLYR2          | Glyoxylate     | 19.1 ± 9  | 18.6 ± 7.9                      | 10.8 ± 4.6   | 349 ± 16               |
|                  | NADPH         | 8.4 ± 1.2 | 15.3 ± 2.9                      | 8.9 ± 1.7    | 1040 ± 73              |
|                  | SSA           | 1457 ± 124 | 12.4 ± 3.5                      | 7.2 ± 2.0    | 5.1 ± 1.8              |
|                  | NADPH         | 12.5 ± 0.6 | 11.3 ± 0.3                      | 6.5 ± 0.2    | 525 ± 40               |
| AgGLYR1          | Glyoxylate     | 23.2 ± 3.2 | 56.7 ± 5.6                      | 28.4 ± 2.8   | 1259 ± 188             |
|                  | NADPH         | 4.5 ± 0.9 | 15.2 ± 2.3                      | 13.1 ± 1.9   | 2870 ± 453             |
|                  | NADPH¹         | 2.2 ± 0.3 | 21.6 ± 2.4                      | 9.3 ± 0.1    | 4370 ± 446             |
|                  | SSA¹           | 870 ± 40  | 23.5 ± 1.5                      | 10.1 ± 0.4   | 11.6 ± 2.2             |
|                  | NADPH¹         | 2.6 ± 0.1 | 18.8 ± 3.3                      | 8.1 ± 0.0    | 3500 ± 208             |
| AgGLYR2          | Glyoxylate     | 19.3 ± 4.0 | 36.8 ± 19.9                     | 18.4 ± 10.0  | 906 ± 353              |
|                  | NADPH         | 34.2 ± 3.3 | 40.6 ± 8.7                      | 22.5 ± 4.8   | 660 ± 130              |
|                  | SSA²           | 8960 ± 710 | 30.7 ± 4.0                      | 17.0 ± 2.2   | 1.9 ± 0.3              |
|                  | NADPH²         | 1.2 ± 0.2 | 19.3 ± 1.5                      | 10.7 ± 0.8   | 9180 ± 1620            |

Data from this study were collected using a Spectramax Plus384 Absorbance Microplate Reader and represent the mean ± SE of two (Arabidopsis) or three (apple and rice) enzyme preparations. The $V_{max}$ values obtained with glyoxylate and NADPH or SSA and NADPH were not influenced by whether the substrate or co-factor was varied or fixed, as indicated by a t-test with $P$ = 0.05 as the significance threshold. Previously published data were collected using a Cary 300 double-beam absorption spectrophotometer (¹Hoover et al., 2007; ²Simpson et al., 2008) and represent the mean ± SE of three enzyme preparations.
FACTOR-1α (EF-1α, At5g60390) was chosen based on the research by Czechowski et al. (2005). Relative expression and data analysis were determined using the 2^−ΔΔCt method (Livak and Schmittgen, 2001). Two technical replicates were conducted for each biological replicate, and the mean ± SE of three biological replicates was determined for each genotype.

The presence and copy number of transgene inserts in GLYR1 overexpression (Ox) lines were determined by Southern hybridization as described by McLean et al. (2003). Genomic DNA was isolated as described by Montiel et al. (2011). About 5 μg of genomic DNA was digested with NcoI, separated on a 1% (w/v) agarose gel, and transferred to a positively charged nylon membrane (Roche Applied Science, Laval, QC, Canada). The membrane was probed, processed, and labeled bands were detected according to digoxigenin (DIG)-labeling kit protocols (Roche Applied Science). The probe was generated by PCR from the cDNA using Probe-GLYR1-F and Probe-GLYR1-R primers (Supplementary Table S1) in the presence of DIG-deoxy-uridine triphosphate. The forward primer flanked an intron/exon boundary in the GLYR1 gene to prevent GLYR1 genomic DNA amplification during probe synthesis. Hybridization was performed with a DIG-labeled probe according to the most stringent hybridization protocols of the manufacturer (Roche Applied Science). The membrane was exposed under a Fluorchem 8800 Imaging System (Alpha Innotech Inc., Miami, FL, United States) for 16 min to visualize labeled bands.

Cloning of cDNAs Encoding Apple and Rice GLYRs and Arabidopsis GLYR2

The predicted amino acid sequences of Arabidopsis GLYR1 (At3g25530, GenBank Acc. No. NM_113449) and GLYR2 (At1g17650, GenBank Acc. No. NM_101628) were obtained from The Arabidopsis Information Resource. A truncated AtGLYR2 lacking its N-terminal 58 amino acid-long predicted plastid targeting sequence (AtGLYR2Δ58) was amplified from the Arabidopsis cDNA using primers VB-F3 and VB-R2 and cloned into pET15b using NdeI and BamHI restriction enzymes to yield pET15b-atGLYR2Δ58 construct. All primer sequences used to clone the various genes are listed in Supplementary Table S1. When expressed, the truncated AtGLYR2 possesses an N-terminal 6×His tag for purification. The AtGLYR1 expression construct was made elsewhere (Hoover et al., 2007b).

Two apple GLYRs were identified in the apple genome database. Two products (958- and 1166-bp), designated as MdGLYR1 (GenBank Acc No. KT202799) and MdGLYR2 (GenBank Acc No. KT202800), respectively, were amplified from apple fruit cDNA cv. Empire and appropriate primers (CT-F12 and CT-R12 for the MdGLYR1 gene, and CT-F13 and CT-R13 for the MdGLYR2 gene). The resulting open reading fragments (ORF) were cloned into pCR2.1-TOPO (Invitrogen). The predicted translation products are 58% identical to each other. MdGLYR1 is 80% identical to AtGLYR1, and MdGLYR2 is 78% identical to AtGLYR2. The primers CT-F17 and CT-R17 were used to amplify MdGLYR1 with 5’ NdeI and 3’ BamHI restriction sites. TargetP4 (v1.1; Emanuelsson et al., 2000) predicted that MdGLYR2 has an N-terminal 52 amino acid plastid targeting sequence. The primers CT-F20 and CT-R20 were used to amplify the MdGLYR2 ORF minus the sequence encoding the putative N-terminal plastid transit peptide with 5’ NdeI and 3’ BamHI restriction sites. Both sets of PCR products were digested with NdeI and BamHI and ligated into pET15b expression vector (Novagen, Cambridge, MA, United States) to produce pET15b-MdGLYR1 and pET15b-MdGLYR2Δ54. When expressed, the truncated MdGLYR2 possesses an N-terminal 6×His tag.

The AtGLYR1 and AtGLYR2 cDNAs were blasted against the rice genome annotation project5 to reveal two full-length cDNA sequences, LOC_Os02g35500 and LOC_Os01g39270, which are 81 and 65% identical to AtGLYR1 and AtGLYR2, respectively. These sequences were designated as putative OsGLYR1 and OsGLYR2, respectively. Full-length OsGLYR1 and OsGLYR2 were amplified from the rice cv. Kaybonnet cDNA using forward and reverse primers, VB-F1 and VB-R1 for OsGLYR1, and VB-F2 and VB-R2 for OsGLYR2. TargetP5 (v1.12; Emanuelsson et al., 2000) predicted that OsGLYR2 contains an N-terminal 35 amino acid-long plastid targeting sequence. The primers VB-F3 and VB-R2 were used to amplify OsGLYR2, minus the sequence encoding the protein’s putative plastid transit peptide. The last two sets of PCR products were cloned into pET15b with NdeI and BamHI enzymes to yield pET15b-OsGLYR1 and pET15b-OsGLYR2Δ35. When expressed, the truncated OsGLYRs possess an N-terminal 6×His tag for purification. All GLYR genes PCR-amplified from cDNAs were sequenced after cloning.

Production, Purification and Kinetic Analysis of Recombinant Apple, Rice and Arabidopsis GLYRs

The constructs pET15b-MdGLYR1 and pET15b-MdGLYR2Δ54 were individually expressed in Escherichia coli BL-21(DE3) Rosetta (pLysS) cells (Novagen) co-expressing the GroES/GroEL chaperone complex, and the resulting soluble recombinant proteins were subjected to affinity chromatography on a Ni2+ column essentially as described previously (Clark et al., 2009). Also, the pET15b-atGLYR1, pET15b-atGLYR2Δ58, pET15b-OsGLYR1, and pET15b-OsGLYR2Δ35 constructs were expressed in E. coli BL-21(DE3) cells (Novagen) and the recombinant proteins purified as described in Hoover et al. (2007a). Each eluted protein was precipitated with 80% (w/v) ammonium sulfate, stored at −80°C, and the pellet was resuspended, as necessary, in 100 μL of reaction buffer [50 mM 4-(2-hydroxyethyl)-1-piperazinethanesulfonic acid (HEPES), 10% sorbitol (w/v), pH 7.8 for AtGLYR1 and AtGLYR2, pH 7.5 and 7.3 for MdGLYR2 and MdGLYR2, 2009).

http://www.cbs.dtu.dk/services/TargetP/
http://rice.plantbiology.msu.edu/
and pH 6.5 and 7.1 for OsGLYR1 and OsGLYR2Δ35, respectively.

Enzymatic activity was monitored as the oxidation of NAD(P)H at room temperature using a Spectramax Plus384 Absorbance Microplate Reader (Molecular Devices, Sunnyvale, CA, United States). The 250-µL assay was initiated with a pre-made master mix of buffer, co-factor and substrate. Reaction rates were proportional to the enzyme concentrations used in the assays (2–50 nM). The optimal pH for enzymatic activity at 50 µM NADPH and 50 µM glyoxylate was determined using buffers (50 mM each) with overlapping pH range [2-morpholinoethanesulfonic acid (MES) for pH 5.5–6.8; HEPES for pH 6.8–8.2; and N-tris(hydroxymethyl)methyl-4-aminobutanesulfonic acid (TABS) for pH 8.2–9.6]. Initially, the three buffers were tested separately for the apple isoforms, then they were combined for rice. The influence of redox ratio on the activities of MdGLYR1 and MdGLYR2 was investigated using 50 µM glyoxylate, 50 µM NADPH and 0–0.5 mM NADP⁺. The kinetic parameters (V_{max}, K_{m}, k_{cat}, k_{cat}/K_{m}) for the purified enzymes were determined using a mixture of the MES, HEPES, and TABS (50 mM each) at the pH optimum for the enzyme source under consideration, with saturating co-factor (50 µM NADPH) and varying substrate (up to 200 µM glyoxylate or 10 mM SSA) or varying co-factor (up to 100 µM NADPH) and saturating substrate (50 µM glyoxylate or 3.75–10 mM SSA) as described previously (Hoover et al., 2007a). For each enzyme source the concentration of substrate or co-factor was varied to give five to six data points both above and below the K_{m}. The dependence of enzymatic activity on pH or redox ratio was determined as the mean ± SD of three to four technical measurements of a typical enzyme preparation (i.e., one biological replicate), whereas the kinetic parameters were expressed as the mean ± SE of three biological replicates, each measured using four technical replicates.

**Generation of Arabidopsis Lines with Altered GLYR Expression**

Arabidopsis glyr1 and glyr2 single mutants (Salk_057410 and G3316040, respectively) were twice backcrossed to the WT. Plants homozygous for glyr1 T-DNA were identified using the glyr1-RP and glyr1-LP primers for the WT allele, and LBb1.3 and glyr1-RP for T-DNA insertion (as recommended by the Salk Institute). The primers were designed using the Salk iSECT tool online software. Similarly, homozygous glyr2 was identified using glyr2-RP and glyr2-LP primers for the WT allele, and GABI-Kat T-DNA border primer and glyr2-RP for the T-DNA. The double glyr1/glyr2 mutant was generated by crossing healthy plants of the homozygous glyr1 and glyr2 single mutants. With the exception of three to five closed flowers with white petals on the main shoot of the mother plant, all flowers and siliques were removed using scissors and forceps. The intact flowers on the female parent were emasculated with the aid of a dissecting microscope. Open flowers with yellow anthers were removed from the male parent and brushed against the stigmatic surface of carpels on the female parent. Female plants were labeled and covered with porous plastic bags until fruit maturity (approximately 4 weeks later). Screening of the T₂ population was based on the gene-specific and T-DNA primers given above.

Arabidopsis glyr1RNAi and glyr2 RNAi lines were generated in glyr2 T-DNA and glyr1 T-DNA mutant backgrounds, respectively. To prepare hairpin RNA constructs, a 233-bp fragment (from +722 to +955 relative to the transcriptional initiation site) for AtGLYR1 and a 382-bp fragment (from +630 to +1012) for AtGLYR2 were each amplified by PCR from cDNA with the following primer pairs containing a LB clone site: Clonase-GLYR1-F/Clonase-GLYR1-R and Clonase-GLYR2-F/Clonase-GLYR2-R, respectively. These amplified fragments were separately cloned into the intermediate clonase vector pDONR221, followed by assembling the fragments into a gateway binary RNAi vector pB7GWIWG2 (II) (by the LR clonase site reaction⁴ (Karimi et al., 2002). The binary vectors GLYR1-pB7GWIWG2(II) and GLYR2-pB7GWIWG2(II) were introduced into Agrobacterium tumefaciens strain EHA105 according to Hood et al. (1993). The GLYR1 and GLYR2 RNAi constructs were introduced, respectively, into the Arabidopsis glyr2 and glyr1 T-DNA mutant backgrounds using the floral dip method (Clough and Bent, 1998). Seeds were selected on solid half-strength MS medium containing 20 mg L⁻¹ Basta (as a T-DNA selection marker) and 100 mg L⁻¹ timentin (to prevent Agrobacterium growth), and then 14-days-old plants were transferred to Sunshine LC1 potting mix (Sun Gro Horticulture, Canada) and placed in a controlled environment growth chamber (Enconair Model GC8-2H set at 60 °C for 16 h and no light at 21 °C for 8 h), where they were allowed to self-pollinate and mature. T₂ progenies were again selected on half-strength MS medium containing Basta and then transferred to LC1 potting mix. Thirty seedlings were separately screened at the rosette stage for the GLYR1 and GLYR2 genes using qPCR (Supplementary Figure S1); the primer sets are given in Supplementary Table S1. Seeds of selected lines were collected for further experiments.

For generation of AtGLYR1 overexpression (Ox) lines, the full-length cDNA sequence for AtGLYR1 was amplified using SpeI-GLYR1-F and SpeI-GLYR1-R primers and cloned into a prepared pMDM7 binary vector (Ni et al., 1995) and transformed into A. tumefaciens strain LB4404. Plant transformation, transformant screening, tissue culture and growth conditions were similar to above, except that seed were selected on medium containing 50 mg L⁻¹ kanamycin. The three transgenic lines that were chosen for further analysis had two to three times the GLYR1 transcript level of the WT (Supplementary Figure S2). Assays of GLYR activity in desalted cell-free leaf extracts with 50 µM glyoxylate and NADPH (Simpson et al., 2008) revealed that two of the three lines had 70–90% higher activity than the WT. Southern blot analysis revealed weak signals, if at all, in the WT lane, perhaps as a results of only partial binding between the unprocessed endogenous gene and the probe. However, it was clear that the two lines with highest GLYR activity carried two to three copies, of the transgene. The line with the highest activity

⁴http://www.psb.ugent.be/gateway/
GLYR1

In preliminary chilling trials, five sterilized seeds from the Arabidopsis WT and each of the GLYR1 Ox, glyr1/2 and glyr1/2-RNAi lines were sown along the center of 14-cm Petri plates containing half-strength MS medium with 0.6% (w/v) agar, 30 g L⁻¹ sucrose, 1 mM MES (pH 5.8) and SSA or GHB (0–3 mM) and then subjected to a stratification regime (3 days, 4°C, dark). Thereafter, the plates were stored vertically in a tissue culture chamber under 12-h period at 22°C for 12 days or 10°C for 27 days (photosynthetic flux density of 60 μmol m⁻² s⁻¹). The experimental design was completely randomized and comprised of three plate replicates for each treatment.

In follow up trials, sterilized seeds from the WT and glyr1/2 and GLYR1 Ox lines were sown as a lawn on 150-mm Petri plates containing half-strength MS medium with 0.6% (w/v) agar, 30 g L⁻¹ sucrose, and 1 mM MES (pH 5.8), subjected to a stratification regime (3 days, 4°C, dark), and then placed in a tissue culture chamber (22°C, 14 h light/18°C, 10 h dark, photosynthetic flux density of 120 μmol m⁻² s⁻¹) for 8 days. Seedlings from all three genotypes were placed along the center of individual 14-cm plates containing the same medium supplemented with glyoxylate (0–2 mM) or SSA (0–3 mM) and the primary root apex was noted on the exterior surface of the plate. Then the plates were stored vertically at 10°C for another 8 days and photographed. The gain in root growth was estimated using ImageJ software. The experimental design was completely randomized and comprised of 10 plate replicates for each treatment.

Distribution of GLYR Activity in Cytosolic, Plastidial, and Mitochondrial Fractions from Arabidopsis

Crude homogenates of rosette leaf tissue from 4-week-old Arabidopsis WT or glyr1 mutant (grown as described above), were prepared using the extraction medium [0.3 M sucrose, 25 mM Na₄P₂O₇, 2 mM sodium ethylenediaminetetraacetate, 10 mM KH₂PO₄, 1% (w/v) polyvinylpyrrolidone-40, 1% bovine serum albumin] described by Taylor et al. (2014). Then the homogenates were sonicated for 10 s using a sonic dismembrator (Fisher Scientific, Model 120), and pelleted by centrifugation. The supernatants were desalted using Econo-Pac 10DG columns (Bio-Rad) and assayed for GLYR activity with 50 μM glyoxylate and NADPH. Intact mitochondria were also isolated from 50 g of homogenized rosette tissue of the glyr1 mutant grown under a 23°C 11-h light (120 μmol m⁻² s⁻¹)/19°C 13-h dark regime according to Taylor et al. (2014). Briefly, the crude homogenate was differentially centrifuged to give an organelle suspension, which was then applied to a linear polyvinylpyrrolidone-40 gradient (0–4.4% v/v) in 28% (w/v) Percoll and centrifuged at 40,000 × g for 40 min. This enabled recovery of a white/pale brown mitochondrial band toward the bottom of the gradient (separated from plastid thylakoid and peroxisomal fractions), which was then concentrated and washed repeatedly using differential centrifugation. Aliquots of the organelle suspension and purified mitochondria fractions were sonicated, pelleted by centrifugation, and desalted as described above. GLYR activity was assayed with 50 μM glyoxylate and NADPH, and mitochondrial fumarase and plastid phosphoribulokinase were assayed as described previously (Taylor et al., 2014), with the exception that the reaction volume was 250 μL and the wavelength used for the fumarase assay was 240 nm (Puchegger et al., 1990), rather than 340 nm as given.

RESULTS

Characterization of Recombinant Apple, Rice, and Arabidopsis GLYRs

The predicted N-terminal targeting presequences in GLYR2s from apple, rice, and Arabidopsis (Brikis et al., 2017) were removed and the resulting truncated sequences (i.e., MdGLYR2Δ54, OsGLYR2Δ35, and AtGLYR2Δ58), as well as the corresponding full-length GLYR1s from the same three plant species (i.e., MdGLYR1, OsGLYR1, and AtGLYR1), were individually expressed in E. coli. This allowed for the recovery of a significant portion of the recombinant GLYR2 proteins, similar to the GLYR1 proteins, in the corresponding soluble fraction (Supplementary Figure S3). In all fractions eluted from Ni²⁺-affinity columns, the recombinant protein was purified to near homogeneity. Predicted protein masses for MdGLYR1, MdGLYR2Δ54, OsGLYR1, OsGLYR2Δ35, AtGLYR1, and AtGLYR2Δ58 were 31.8, 33.1, 30.5, 31.4, 30.7, and 33.2 kDa, respectively. The mass of recombinant AtGLYR2Δ58 appeared slightly larger than predicted. The scientific literature contains many examples of cytosolic proteins that migrate during SDS polyacrylamide electrophoresis at rates that are inconsistent with their molecular mass (see Shi et al., 2012). While such “gel shifting” could be caused by single amino acid substitutions, we repeatedly cloned and sequenced the gene and prepared the recombinant AtGLYR2Δ58 protein with similar results. It is possible that interaction between SDS and the mutant protein is modified by the presence of the N-terminal 6×His motif. The activities of MdGLYR1 and OsGLYR1 were maximal at pH 6.8 and 6, respectively, although both enzymes retained at least half of maximal activity over the pH range of 5.5–7. The maximal activities of MdGLYR2 and OsGLYR2 were found to be at approximately pH 6.8 and at least 50% of maximal activity was retained in the pH range of 6–8 (Figure 1).

In the present study, enzymatic activity for the various recombinant proteins was determined using a microplate reader. A preliminary study of glyoxylate-dependent activity in rice GLYR1 revealed that 50 and 100 μM NADPH gave approximately 4.5 times the activity with corresponding levels of NADH (data not shown); therefore, detailed NADH kinetics for both rice and apple GLYRs were not determined here. The kinetic data for the apple and rice proteins with NADPH as the co-factor

http://rsbweb.nih.gov/ij/
represent the mean (±SE) of three replicate preparations, and the $V_{max}$ values obtained with glyoxylate or SSA were not significantly influenced by whether the substrate or co-factor was varied or fixed ($P > 0.05$, t-test) (Table 1). Glyoxylate-dependent kinetics with NADPH were also determined here for the AtGLYR1 and the AtGLYR2 using the microplate reader, so that direct comparisons could be made with published values obtained with the use of a Cary 300 spectrophotometer. Overall, the catalytic efficiencies ($k_{cat}/K_m$) of the apple and rice GLYRs, like those for the Arabidopsis GLYRs (Hoover et al., 2007a; Simpson et al., 2008), were within the same order of magnitude for each substrate, and the $K_m$ values for glyoxylate and NADPH were in the micromolar range, whereas the $K_m$ values for SSA were in the millimolar range (see Table 1 and Supplementary Figures S4–S6). For all GLYRs, the highest catalytic efficiencies were observed for NADPH, mostly at fixed glyoxylate concentrations, followed by glyoxylate at fixed NADPH concentrations, whereas catalytic efficiencies for SSA were 0.2 to 1.4% of those for glyoxylate (Table 1). Overall, the catalytic efficiencies of GLYR1 and GLYR2 for glyoxylate or SSA in the NADPH-dependent reaction ranged approximately three- to four-fold only across the species compared here, and there were no identifiable catalytic differences between the GLYR1 and GLYR2 isoforms, indicating that all GLYRs prefer glyoxylate over SSA, and have a high affinity for their co-substrate NADPH. The activities of MdGLYR1 and MdGLYR2 displayed inhibitory responses to an increasing NADP$^+$/NADPH ratio (Figure 2), and their reaction rates were reduced by half with an NADP$^+$/NADPH ratio of 5.

**FIGURE 1** | Dependence of recombinant apple and rice GLYR activities on pH. The left and right panels, represent apple (A,B) and rice (C,D) GLYRs, respectively, whereas the upper and lower panels represent GLYR1a (A,C) and GLYR2a (B,D), respectively. The data, expressed as specific activity ($\mu$mol min$^{-1}$ mg$^{-1}$ protein), were determined using 50 $\mu$M glyoxylate and NADPH in three buffers (50 mM each): 2-morpholino-ethanesulfonic acid, 4-(2-hydroxyethyl)-1-piperazineethanesulfonic acid and N-tris(hydroxymethyl)methyl-4-aminobutanesulfonic acid either separately (A,B) or together (C,D). Each datum represents the mean (±SD) of three technical measurements of a typical enzyme preparation.

**Phenotypic Response to Chilling of Various GLYR Lines Supplied with Exogenous Glyoxylate, Succinic Semialdehyde, or γ-Hydroxybutyrate**

To investigate whether GLYR activity is linked to a phenotypic response to chilling, seeds of Arabidopsis WT, GLYR1 Ox, glyr1/2 and glyr1/2-RNAi were sown directly on agar medium and grown at 22°C or 10°C in separate tissue culture chambers. At 22°C, the early growth of the different genotypes was similar in the absence or presence of added SSA or GHB up to 12 days (data not shown). The early growth of the genotypes at 10°C was also similar in the absence of glyoxylate or GHB after 27 days (Figures 3A,E). However, when SSA or GHB (up to 3 mM each) was added to the medium, a differential response was observed among the genotypes at 10°C. That is, increasing concentrations of SSA progressively inhibited growth and development (Figures 3B–D). At 1 mM SSA, the inhibition was greater with glyr1/2-RNAi and glyr1/2 than the WT and GLYR1 Ox, and purple tingeing of the leaves was repeatedly observed in the WT (Figure 3B). At 2 mM SSA, radicle development in both glyr1/2-RNAi and glyr1/2 was halted, and shoot production was entirely absent (Figures 3C,D). Higher magnification observations from a replicate experiment indicated that 2 mM SSA stunted radicle growth in all four lines, and confirmed that shoot and leaf development was hindered in glyr1/2-RNAi and glyr1/2 (Figures 3C,D). In general, the plantlets were more tolerant to GHB than SSA. Root length was reduced with 2 and 3 mM GHB, and shoot reduction and yellowing were particularly evident at 3 mM GHB; notably, the WT and GLYR1 Ox line were more sensitive to GHB than glyr1/2 and glyr1/2-RNAi (Figures 3E–G).

To further investigate whether GLYR activity could be linked to a phenotypic response, Arabidopsis WT, and transgenic GLYR1 Ox and glyr1/2 plantlets were transferred to a chilling temperature of 10°C for an 8-days period in the presence of glyoxylate or SSA. In the absence of glyoxylate, all genotypes displayed a similar gain in root length; however, the addition of glyoxylate generally decreased this gain (Figure 4A). Notably, the GLYR1 Ox line gained more root length than WT and glyr1/2 mutant plants with 1 mM glyoxylate, whereas GLYR1 Ox gained more root length than glyr1/2 but not the WT in the presence of 2 mM glyoxylate (Figure 4B). We were unable to demonstrate the same differential response to SSA using this experimental design (data not shown).

**Subcellular Localization of GLYR Activity in Arabidopsis**

As expected, AtGLYR1 expression in leaves of the Arabidopsis gylr1 mutant was not detectable, and the total GLYR activity in crude tissue homogenates was markedly less than that in the WT (Figures 5A,B), indicating that cytosolic AtGLYR1 and plastidial/mitochondrial AtGLYR2 could be responsible for approximately 85 and 15% of the total GLYR activity, respectively, in leaves. To further assess whether AtGLYR2 is located in mitochondria, as well as plastids, mitochondria were purified from a leaf organelle suspension from the gylr1
FIGURE 2 | Dependence of recombinant apple GLYR activities on NADP+/NADPH ratio. The activities of MdGLYR1 (A) and MdGLYR2Δ54 (B) were determined using 50 µM glyoxylate and NADPH, and increasing concentrations of NADP⁺. Data represent the mean ± SD of four technical measurements of a typical enzyme preparation.

FIGURE 3 | Impact of Arabidopsis genotype on the early phenotypic response to succinic semialdehyde or γ-hydroxybutyrate under chilling stress. Seeds of wild-type (WT) plants and glyr1/2-RNAi, glyr1/2, and GLYR1 Ox were sown on plates containing half-strength MS medium with up to 3 mM SSA or GHB and grown at 10°C for 27 days. Levels of SSA or GHB are 0 (A,E), 1 (B), 2 (C,D,F), and 3 (G) mM.

mutant. The organelle suspension and purified mitochondria were assayed for marker enzymes and GLYR activity. As shown in Figure 5C, the activities of plastidial phosphoribulokinase, mitochondrial fumarase and GLYR were present in the organelle suspension. Phosphoribulokinase activity was not detectable in the purified mitochondria, confirming that this fraction was essentially free of plastid contamination and, hence, plastidial AtGLYR2. The specific activity of
fumarase was enriched approximately 53-fold in the purified mitochondria, compared to the organelle suspension, whereas the specific activity of GLYR was enriched approximately 2.1-fold. These findings lend further support for the notion that AtGLYR2 is indeed localized in mitochondria, as well as plastids.

DISCUSSION

The recombinant apple and rice GLYR1s and GLYR2s investigated here were highly active with glyoxylate and NADPH over a relatively broad pH range from slightly acidic to slightly basic, regardless of whether the different pH buffers were combined or tested separately in the assay mixtures (Figure 1). These results are in agreement with previous studies of the recombinant Arabidopsis GLYRs (Hoover et al., 2007b; Simpson et al., 2008), and the purified spinach enzyme (Kleckzowski et al., 1986). Our kinetic analyses showed that recombinant GLYRs from the monocotyledonous rice, as well as the dicotyledonous apple and Arabidopsis, behaved similarly with respect to their catalytic efficiencies (Table 1), probably because of their conserved active site residues and NADPH-binding sequence (Hoover et al., 2013; Brikis et al., 2017). The $K_m$ values for glyoxylate and SSA were 5–53 µM and 0.8–9 mM, respectively, with NADPH as the co-factor (Table 1), whereas the $K_m$ for glyoxylate was 60 µM for the purified spinach enzyme (Kleckzowski, 1995). Thus, it can be concluded that all plant GLYRs have a preference for glyoxylate over SSA. There are few reliable data on the pool sizes of glyoxylate and SSA in plants, perhaps in part because of their propensity to form adducts (Nikiforova et al., 2014; also see references therein). The level of glyoxylate in rice seedlings is 0.27 µmol g$^{-1}$ fresh mass (FM) (Yu et al., 2010) or 0.27 mM (assuming 1 g FM equals 1 mL H$_2$O), which is within the micromolar range known to inhibit photosynthesis in intact chloroplasts (Campbell and Ogren, 1990). The product of glyoxylate reduction, glycolate, is known to accumulate in rice seedlings germinated under hypoxic conditions (Narsai et al., 2009). SSA levels are low (10–16 µM) in various parts of Arabidopsis plants (Palanivelu et al., 2003; Toyokura et al., 2011). However, accumulation of GHB, the product of SSA reduction, increases in Arabidopsis and tobacco plants from approximately 0.04 µmol g$^{-1}$ FM to 0.12–2.4 µmol g$^{-1}$ FM with the imposition of salinity, submergence, heat and chilling stresses (Allan et al., 2008), and submergence-induced GHB accumulation is markedly reduced in Arabidopsis glyr1 and glyr2 mutants (Allan et al., 2012).

The $K_m$s for NADPH were 2–9 µM with the recombinant apple, rice, and Arabidopsis GLYRs (Table 1), and 3–6 µM for the purified spinach enzyme (Kleckzowski et al., 1986; Kleczkowski, 1995). NADPH was preferred over NADH as the co-factor for both OsGLYR1 and AtGLYR1 (see Results; Hoover et al., 2007a), results in general agreement with those for the purified spinach enzyme (note: $K_m$s for glyoxylate and NADH are 0.3 and 1.1 mM, respectively, in combination with a lower estimated $V_{max}$, Kleczkowski, 1995). Our previous results indicated that the $K_i$ for NADP$^+$, a competitive inhibitor of the glyoxylate- and SSA-dependent reactions, is 3.1–9.5 µM for AtGLYR1 (Hoover et al., 2007b). This is much lower than the corresponding value ($K_i = 60$ µM) for the purified spinach enzyme (Kleckzowski, 1995), and seemingly at odds with the slight inhibition of apple GLYRs by NADP$^+$ (i.e., approximately 50% inhibition with a fivefold increase in NADP$^+$, Figure 2). The estimated ratios of NADPH:NADP$^+$ in intact leaves of Arabidopsis (Allan et al., 2008, 2012), cytosol of spinach, barley, and pea leaves (Heineke et al., 1991; Wigge et al., 1993; Kleczkowski, 1995; Igamberdiev and Gardeström, 2003), and spinach chloroplasts (Heineke et al., 1991) are approximately 2, 1–4, and 0.5,
respective. Thus, it can be concluded that all plant GLYRs have a high affinity for NADPH, and NADP⁺ is not likely to be an important regulator of GLYR activity under ambient conditions, as suggested earlier (Kleczkowski, 1995). Notably, the estimated NADPH/NADP⁺ ratios in Arabidopsis and tobacco leaves have been shown to increase by 0.5- to 4-fold with submergence, heat and chilling stresses (Allan et al., 2008). In the absence of applied stress, the NADPH/NADP⁺ ratio in an NAD KINASE1 Ox line of Arabidopsis is increased by threefold, and there is a corresponding increase in GHB accumulation (Allan et al., 2012).

Taken together, these findings are in agreement with the suggestion that GLYR activities are stimulated by stress, likely in response to the elevated supply of one or both substrates, glyoxylate and SSA, as well as the co-factor NADPH (Hoover et al., 2007a; Allan et al., 2008, 2012), despite the low affinity of GLYRs for SSA and the low tissue levels of SSA under ambient conditions. Mekonnen and Ludewig (2016) have shown the growth of both glyr1/2 double mutants and WT plants is more sensitive to exogenous SSA than GHB at ambient temperature. Since chilling is known to cause the accumulation of both GHB and its precursor GABA in Arabidopsis within hours of exposure, and greatly enhance the leaf NADPH:NADP⁺ ratio (Allan et al., 2008), we predicted that chilling would generate a differential phenotype among the GLYR genotypes. Indeed, plantlets of the various genotypes did display a qualitative difference in the early growth response when grown on increasing levels of exogenous SSA or GHB at 10°C, but not 22°C (Figure 3). It was also possible to quantitatively demonstrate that the root growth of established plantlets of the various genotypes responded differentially to exogenous glyoxylate (but not SSA) with chilling (Figure 4). Since roots are unlikely to generate photorespiratory glyoxylate, these findings suggest that GLYRs function in the reduction of toxic aldehydes, including glyoxylate from multiple metabolic pathways (e.g., glyoxylate from non-photorespiratory serine synthesis, fatty acid catabolism, interconversion of organic acids, purine catabolism (see Introduction for citations), in plants subjected to chilling stress.

The reduction of glyoxylate can also be catalyzed by various hydrogenpyruvate reductases (Kleczkowski et al., 1991; Timm et al., 2011). For example, in Arabidopsis, there are three isoforms: a peroxisomal HPR1 with a preference for NADH and hydrogenpyruvate over NADPH and glyoxylate, respectively; a cytosolic HPR2 with a preference for NADPH over NADH, but similar preference for hydrogenpyruvate and glyoxylate; and, a plastidial HPR3 with a preference for NADPH and glyoxylate over NADH and hydrogenpyruvate, respectively (Timm et al., 2011). Our research has used transient and stable expression systems to unambiguously demonstrate that apple, rice, and Arabidopsis GLYR1s are localized to the cytosol, whereas the corresponding GLYR2s are localized in both plastids and mitochondria (Ching et al., 2012; Brikis et al., 2017). Here, approximately 85% of the total glyoxylate-reducing activity in Arabidopsis leaf extracts was lacking in the glyr1 mutant (Figure 5), indicating that GLYR1, rather than HPR2, is responsible for the majority of the cytosolic activity. Furthermore, the recovery of glyoxylate-reducing activity in purified mitochondria from the glyr1 mutant, free from cytosolic GLYR1, plastidial GLYR2 (Figure 5) or plastidial HPR3 (Timm et al., 2011) contamination, provided in planta support for the dual targeting of GLYR2. The percentage distribution of GLYR2 between mitochondria and plastids remains uncertain, although oxalate may be a useful tool to distinguish between plastidial GLYR2 and HPR3 activities (Kleczkowski et al., 1991). Early studies, using the photosynthesizing protist Euglena gracilis, reported the presence of NADPH-dependent GLYR activity in the intermembrane space of mitochondria, which constitutes approximately 2% of total mitochondrial protein (Yokota and Kitaoka, 1979; Yokota et al., 1985). The $K_m$ values for NADPH and glyoxylate for this enzyme are in the low micromolar range and the pH optimum is 6.45 (Yokota et al., 1985), which are similar to our findings with GLYRs from Arabidopsis, apple and rice (Table 1).

In summary, the ability of GLYRs to promiscuously catalyze the reduction of SSA, as well as glyoxylate, appears to have physiological implications for the plant response to chilling conditions and could be relevant to other abiotic stresses. The generation of SSA is generally restricted to mitochondria with the catabolism of GABA (Shelp and Zarei, 2017; see references therein), whereas glyoxylate can be derived from multiple biochemical processes and subcellular compartments. With abiotic stresses, the detoxification of SSA could be diverted from the generation of succinate to GHB as SSA dehydrogenase activity becomes restricted with the accumulation of NADH (António et al., 2016). With the exception of hypoxia, abiotic stresses generally cause stomatal closure and result in corresponding increases in Rubisco oxygenase activity (Allan et al., 2009). This could increase the production and levels of glyoxylate,
so that its detoxification could require capacity in addition to the photorespiration-associated transaminase activities (i.e., serine/asparagine-glyoxylate and glutamate-glyoxylate transaminases) (Peterhansel et al., 2010; Modde et al., 2017); reduction to glycolate (GLYR and HPR); decarboxylation to formate, and, oxidation to oxalate (Igamberdiev and Eprintsev, 2016; see references therein). On the other hand, with hypoxia glycolate oxidase and glycine decarboxylase activities could be inhibited (Allan et al., 2009), and detoxification of non-photorespiratory sources of glycolate could occur via a combination of reduction (GLYR and HPR), GABA transamination, and decarboxylation and oxidation activities.

**AUTHOR CONTRIBUTIONS**

BS conceived and supervised the project. AZ, CB, VB, and GC conducted the experiments and data analysis. JS prepared and identified the Arabidopsis GLYR1 overexpression lines. JD and GB discussed the project. AZ, CB, GB, and BS wrote and/or edited the manuscript. All authors read and approved the final manuscript.

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**SUPPLEMENTARY MATERIAL**

The Supplemental Material for this article can be found online at: http://journal.frontiersin.org/article/10.3389/fpls.2017.01399/full?supplementary-material
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