Vegetative and Fecundity Fitness Benefit Found in a Glyphosate-Resistant *Eleusine indica* Population Caused by 5-Enolpyruvylshikimate-3-Phosphate Synthase Overexpression

Zhiling Li¹, Xiangju Li¹, Hailan Cui¹, Guodong Zhao², Dan Zhai³ and Jingchao Chen¹*

¹ Institute of Plant Protection, Chinese Academy of Agricultural Sciences, Beijing, China, ² School of Life Sciences, Inner Mongolia University, Hohhot, China, ³ College of Plant Protection, Henan Agricultural University, Zhengzhou, China

Fitness is an important trait in weed species that have developed herbicide resistance, including resistance to the popular herbicide glyphosate. Fitness cost is commonly found in weeds with glyphosate resistance, which is caused by target-site mutations. In this study, the vegetative and fecundity fitness traits in a glyphosate-resistant (GR) *Eleusine indica* population caused by 5-enolpyruvylshikimate-3-phosphate synthase (*EPSPS*) overexpression were investigated under glyphosate-free conditions. The results showed that the resistance index of the population resistant (R) to glyphosate compared with that of the population susceptible (WT) to it was approximately 4.0. Furthermore, *EPSPS* expression level in the R plants was 20.1–82.7 times higher than that in the WT plants. The dry weight of the R population was significantly higher than that of the WT population at the later growth stage after planting; a similar trend was observed for leaf area. In addition, seed production in the R population was 1.4 times higher than that in the WT population. The R and WT populations showed similar maximum germination rates and T50 values. UPLC-MS/MS was performed for the metabolic extracts prepared from the leaves of R and WT populations to address changes in the metabolome. A total of 121 differential metabolites were identified between R and WT individuals. The levels of 6-hydroxy-1H-indole-3-acetamide and indole acetaldehyde, which are associated with auxin synthesis, were significantly higher in plants of the R population than in those of the WT population. However, some secondary metabolite levels were slightly lower in the R population than in the WT population. To conclude, in this study, vegetative and fecundity fitness benefits were found in the GR *E. indica* population. The results of metabolome analysis indicate that the increase in 6-hydroxy-1H-indole-3-acetamide and indole acetaldehyde levels may be the result of fitness benefit. Further studies should be conducted to confirm the functions of these metabolites.

**Keywords:** fitness, benefit, glyphosate, *EPSPS*, metabonomics, auxin
**INTRODUCTION**

Glyphosate is a non-selective herbicide that has been extensively used worldwide for weed control in crop and non-crop situations (Duke and Powles, 2008). This herbicide clearly inhibits the target 5-enolpyruvylshikimate-3-phosphate synthase (EPSPS; E.C. 2.5.1.19), which plays an important role in the formation of aromatic amino acids (Steinrücken and Amrhein, 1980). The annual Gramineae weed goosegrass [Eleusine indica (L.) Gaertn.,] originated from Asia, is now widely distributed in temperate and tropical regions (Holm et al., 1977). This weed can lead to a 20–50% reduction in the yield of infested crops (Ma et al., 2015; Song et al., 2019). Glyphosate is the most commonly used herbicide to control weed species, including goosegrass, in infested acreage, especially orchards (Duke and Powles, 2008). However, after many years of selection pressure, weed species have evolved to develop resistance to glyphosate; 54 glyphosate-resistant (GR) weed species have been reported (Heap, 2021). In China, most of the GR goosegrass species have been reported in orchards; in recent years, GR goosegrass has been reported to occur in cotton fields and tea gardens, as well as in direct sowing rice fields (Chen et al., 2015a, 2020a,b; Deng et al., 2020).

One or more amino acid substitution in the conserved regions of EPSPS or overexpression of this target gene is considered the cause of target-site resistance (TSR) to glyphosate in weeds (Sammons and Gaines, 2014). The mutation commonly occurs at positions Thr102 or Pro106, and EPSPS overexpression is commonly caused by the amplification of this gene in the chromosome (Gaines et al., 2010; Yu et al., 2015; Li et al., 2018; Perotti et al., 2019). Non-target-site resistance (NTSR) is another resistance mechanism that is associated with increased metabolism, reduced translocation or uptake, and enhanced vacuolar sequestration of glyphosate (Délye, 2013). For goosegrass, both target site mutations at amino acid residues and EPSPS overexpression have been reported (Baerson et al., 2002; Chen et al., 2015b; Yu et al., 2015). In addition, our previous study has found that EPSPS overexpression and Pro-106-Ala mutation have evolved in the same GR individuals (Chen et al., 2020b).

Fitness is defined as the reproductive success of a plant in a given environment; therefore, seed production, germination rate, and vegetative growth are important parameters that determine fitness (Vila-Aiub et al., 2009). Fitness cost is a characteristic of herbicide-resistant weeds, where plants show a competitive disadvantage in the absence of herbicide selection pressure (Vila-Aiub et al., 2009). The GR weed species, caused by either NTSR or TSR mechanisms, are associated with fitness costs (Yanniccare et al., 2015; Fernández-Moreno et al., 2017; Vila-Aiub et al., 2019). A GR Lolium perenne population, due to high EPSPS activity, showed a 33% reduction in leaf blade area and a 55% reduction in shoot biomass, compared with susceptible plants (Yanniccare et al., 2015). A Lolium multiflorum population exhibited altered uptake and translocation, which reduced seed output by at least 38% compared with that of the susceptible population (Fernández-Moreno et al., 2017). Except for the fitness cost, high fecundity rate (30% increase in silique and seed number per plant) was found in GR Oryza sativa and Arabidopsis thaliana, in which transgenic technology was used to overexpress the endogenous gene EPSPS (Yang et al., 2017; Fang et al., 2018). A mutated EPSPS gene which was overexpression in O. sativa produced 17%–19% more grains compared to the wild type in the absence of glyphosate application (Achary et al., 2020). For goosegrass, a significant fitness cost occurred in the GR population with a double mutation in EPSPS (Han et al., 2017). However, there have been few reports on fitness traits in GR goosegrass with EPSPS overexpression. Here, the characteristics of fitness of a GR goosegrass population caused by EPSPS overexpression, as confirmed in our previous study, was assessed. The objectives of this research were to: (1) confirm the resistance level and mechanism for the selected population, (2) to investigate vegetative growth, seed production, and germination rate of resistant versus susceptible plants sharing a common genetic background, and (3) to elucidate the differences in metabolites between resistant and susceptible plants using metabolomics.

**MATERIALS AND METHODS**

**Plant Material and Genetic Background**

**Control**

We selected a GR goosegrass population caused by EPSPS overexpression, as confirmed in our previous study (Chen et al., 2015b). To minimize variability in genetic backgrounds, we selected resistant and susceptible individuals from within a population (Vila-Aiub et al., 2015; Keshkhar et al., 2019). In detail, seeds from more than 100 individuals in the resistant population were collected and then grown to the tillering stage under (27 ± 4)/(20 ± 4)°C day/night temperature with a 14 h photoperiod (Chen et al., 2020b). At least 40 seedlings were cloned into two groups and labeled. One group was sprayed with glyphosate at a dosage of 900 g ai ha⁻¹ to detect resistant and susceptible plants. Twenty-one days after treatment, based on the response of the corresponding clones, the individuals of the other group not treated with glyphosate were classified. The individuals for which sensitivity was confirmed were grown for seed production in two isolated places to prevent cross-pollination. The seeds were collected for fitness studies and named R (resistant) and WT (susceptible).

**Excluding the Non-target Site Resistance Mechanism**

Seeds of the R and WT populations were cultured in plastic pots (8 cm × 8 cm) and kept in a greenhouse (Beijing, China) under the same condition described above. All individuals were thinned to eight plants per pot around the two-leaf stage and watered as needed (Chen et al., 2020b). To confirm the EPSPS overexpression resistance mechanism and no other NTSR mechanisms exist in R population, different treatments were conducted for these two populations. One group treated a set of doses (0, 56, 112, 225, 450, and 900 g ai ha⁻¹ for WT population; 0, 450, 900, 1800, 3600, and 7200 g ai ha⁻¹ for R population) of glyphosate (Roundup, isopropylamine salt of glyphosate, 410 g ae L⁻¹, Bayer Crop Science, St. Louis, MO, United States). All the individuals were applied glyphosate at four- to six-leaf stage using a moving TeeJet® XR8002 flat fan nozzle cabinet.
sprayer (Beijing Research Center for Information Technology in Agriculture, Beijing, China) (Wang et al., 2018). For the other group, the P450 inhibitor piperonylbutoxide (10 mM) and the GST inhibitor 4-chloro-7-nitro-1,2,3-benzooxadiazole (3 mM) were sprayed for all individuals before glyphosate treatment for 3 and 72 h, respectively. Twenty-one days after treatment, fresh weight of the shoots in each pot under different treatments was measured (Jugulam and Shyam, 2019; Chen et al., 2020b).

Expression of 5-Enolpyruvylshikimate-3-Phosphate Synthase

The expression level of EPSPS was measured in these two populations using a standard method. In brief, leaf tissue of 24 individuals from each population was collected when the plants were at the five- to six-leaf stage. Total RNA was extracted from each sample, and the quality was detected using agarose gel electrophoresis. First-strand complementary DNA was synthesized using EasyScript® All-in-One First-Strand cDNA Synthesis SuperMix (TransGen Biotech, Beijing, China). We used confirmed primer pairs to amplify EPSPS, with chloroplast acetolactate synthase (ALS) as the reference gene (Chen et al., 2017a). Quantitative real-time PCR (qPCR) was conducted in a 20 µL reaction with the 2× SYBR Green PCR master mix and run on an ABI 7500 PCR machine (Applied Biosystems Inc., Foster City, CA, United States). The expression of the target gene relative to that of the reference gene was analyzed using the 2−ΔΔCt method and the formula −2ΔCt = −[(Ct, EPSPS − Ct, ALS)trat − (Ct, EPSPS − Ct, ALS)control] (Livak and Schmittgen, 2001). This experiment was repeated twice.

Assessment of Vegetative and Fecundity Traits Without Competition

The purified seeds of R and WT were cultured using the method described above, and only one individual was maintained in each pot. At 20, 45, and 70 days after the individuals emerged, leaf area for all leaves was measured using a leaf area scanner (YMJ-pot. At 20, 45, and 70 days after the individuals emerged, leaf area was measured (Jugulam and Shyam, 2019; Chen et al., 2020b).

Characterization of Seedling Emergence Rate

Thirty seeds each of the R and WT populations, were placed on a 9 cm × 9 cm petri dish lined with filter paper and moistened with 5 mL distilled water. They were kept in a growth chamber with a 16 h photoperiod; 21 µmol m−2 s−1 light intensity; and day/night temperatures of 30 and 20°C, respectively. Seedling emergence (appearance of the coleoptile) was recorded daily for 10 days and this experiment was repeated twice (Matzrafi et al., 2017).

Metabolites Analysis Using LC-MS/MS

Five individuals each of the R and WT populations were selected, and the leaf tissue was prepared using a standard method. In brief, 50 mg of sample was ground using liquid nitrogen and homogenized using an extraction solution with an isotopically labeled internal standard mixture. The samples were then incubated for 1 h at −40°C and centrifuged at 12,000 rpm at 4°C for 15 min. The resulting supernatant was transferred to a fresh glass vial for further analysis.

LC-MS/MS analyses were performed using a UHPLC system (Thermo Fisher Scientific, Waltham, MA, United States) with a UPLC HSS T3 column (2.1 mm × 100 mm, 1.8 µm) coupled to a Q Exactive HFX mass spectrometer (Orbitrap MS, Thermo Fisher Scientific, CA, United States). The mobile phase consisted of 5 mmol L−1 ammonium acetate and 5 mmol L−1 acetic acid in water (A) and acetonitrile (B). The auto-sampler temperature was 4°C, and the injection volume was 3 µL. A standard method for QE HFX mass spectrometry was performed, and all parameters were the same as those used in the previous study (Li et al., 2021).

Statistical Analyses

The data from the whole-plant assay and seedling emergence analysis were analyzed using the log-logistic model: $Y = Y_0 + a/[1 + (X/X_0)^b]$, using SigmaPlot 12.0 (Systat Software, San Jose, CA, United States) (Seefeldt and Fuerst, 1995; Ritz et al., 2013). In this model, $b$ is the slope of the curve, $Y_0$ is the lower limit, $a$ is the difference between the upper and lower limits, and $X_0$ is the herbicide dose required for 50% plant growth reduction (GR50) or the time required for 50% of the seeds to be germinated (T50). The resistance index (RI) reflects the resistance level for the R population and is the ratio of the GR50 of the R population to that of the WT population.

5-Enolpyruvylshikimate-3-phosphate synthase expression and fitness trait data were analyzed using Student’s t-test for R and WT populations. All analyses were performed using SPSS (version 13.0; SPSS, Chicago, IL, United States). The raw data from metabolite analysis were converted to the mzXML format using ProteoWizard software and processed with an in-house program, which was developed using R package and
based on XCMS, for peak detection, extraction, alignment, and integration. Then, an in-house MS2 database (BiotreeDB) was used for metabolite annotation. The cutoff for annotation was set at 0.3. Principal component analysis (PCA) and orthogonal projections to latent structures discriminant analysis (OPLS-DA) were performed using the SIMCA software (V16.0.2, Sartorius Stedim Data Analytics AB, Umea, Sweden). Metabolites with variable importance in the projection (VIP) values >1 and a false discovery rate <0.05 were considered differentiated (Zhang et al., 2020).

RESULTS

Response of R and WT Populations to Glyphosate

The P450 inhibitor piperonylbutoxide and the GST inhibitor 4-chloro-7-nitro-1,2,3-benzoxadiazole were used to identify possible NSTR in the R population. The response to glyphosate was similar between resistant individuals that were treated with the inhibitors and those that were not. A slight growth inhibition was observed after administration of the recommended dose (Figure 1). The GR50 value for glyphosate calculated from the fresh weights of the R population was 1539.4 g ae ha⁻¹, and a similar result (1517.0 g ae ha⁻¹) was calculated after treatment with the inhibitors. Compared with the WT population, the RI of R to glyphosate was 4.0 and 4.8, respectively.

Expression Level of 5-Enolpyruvylshikimate-3-Phosphate Synthase

Under glyphosate-free conditions, expression of EPSPS relative to that of the reference gene ALS was determined for the R and WT populations. The expression level of EPSPS (relative to that of ALS) ranged from 0.7 to 3.6 in the WT individuals (Figure 2). EPSPS expression level in R individuals showed a wide range, with the expression level in some of them reaching up to 150.0 (relative to that of ALS). EPSPS expression level in the R plants was 20.1–82.7 times higher than that in the WT plants.

Vegetative Growth and Fecundity

The dry weight of above ground tissues and the total leaf area were recorded at 20, 45, and 70 days after planting. No significant differences were found in the dry weight and leaf area between the R and WT populations at 20 and 45 days after planting (Table 1). However, the dry weight of the R population was significantly
higher than that of the WT population at 70 days after planting, and a similar trend was found for the leaf area. Furthermore, seed production in the R population was much higher than that in the WT population, at 658 and 466 per plant, respectively (Figure 3).

Seedling Emergence Rate
The GR population R and susceptible population WT showed similar maximum germination rate and $T_{50}$ under the selected conditions (Figure 4). From 24 to 72 h, the seedlings from both the R and WT populations emerged quickly and at a high percentage. At approximately 96 h, maximum germination was reached in both R and WT populations.

Metabolites Analysis
Metabolite differences between the R and WT populations were detected using UPLC-MS/MS, and five individuals were selected from each population. The results of PCA and OPLS-DA indicate that the metabolome between R and WT shows an identical pattern (Supplementary Figure 1). The values of $R^2_Y$ and $Q^2$, calculated by a permutation test, were 0.97 and 0.01, respectively (Supplementary Figure 1). All the above findings indicate that the established model has no overfitting phenomenon and can be used in subsequent analyses.

A total of 13,056 metabolites were detected after preprocessing the raw data, and 3,624 of them were identified using different databases. Under the criteria of statistical significance (VIP > 1 and $p < 0.05$), 121 differential metabolites were identified between R and WT individuals (Supplementary Table 1 and Table 2). Among these metabolites, 95 were downregulated and 26 were upregulated in R relative to those in WT (Supplementary Table 1 and Table 2). The levels of metabolites associated with auxin synthesis were significantly higher in the R population than in the WT population (Table 2). However, the levels of some secondary metabolites were lower in the R individuals than in the WT individuals. In addition, metabolites in the tryptophan (Trp) and tyrosine (Tyr) metabolism pathways were significantly different between the R and WT populations (Supplementary Figure 2).

DISCUSSION
In this study, the fitness characteristics of a GR goosegrass population (caused by EPSPS overexpression) were investigated in the absence of glyphosate. To estimate the fitness cost, individuals or genotypes must share the same genome, except for

| ID   | MS2 name                                           | MS2 score | VIP value | P-value | Log2 fold change |
|------|----------------------------------------------------|-----------|-----------|---------|-----------------|
| 221  | Arbutin                                            | 0.93      | 2.18      | 0.005   | 3.45            |
| 382  | 6-Hydroxy-1H-indole-3-acetamide                    | 0.79      | 1.82      | 0.001   | 2.96            |
| 224  | Serotonin                                          | 0.92      | 2.30      | 0.001   | 2.04            |
| 305  | (7alpha,10beta)-1(10 -> 19)-abeo-7-acetoxyisobacun-3,10-olide | 0.86      | 1.83      | 0.023   | 1.97            |
| 160  | Indole acetaldhyde                                 | 0.91      | 2.25      | 0.001   | 1.87            |
| 86   | Mustakone                                          | 0.99      | 1.71      | 0.027   | 1.58            |
| 761  | Coumestrin                                         | 0.46      | 2.24      | 0.000   | −2.25           |
| 352  | Chrysoeriol 7-O-(6''-malonyl-glucoside)            | 0.82      | 2.27      | 0.010   | −2.13           |
| 584  | Tricin arabinoside                                 | 0.63      | 2.25      | 0.002   | −1.97           |
| 581  | Vicenin 2                                          | 0.64      | 2.32      | 0.010   | −1.85           |
| 145  | Aflatoxin G                                        | 0.97      | 2.06      | 0.010   | −1.79           |
| 213  | Cyanidin 3-rutinoside                              | 0.94      | 2.23      | 0.020   | −1.72           |
| 639  | Cyanidin 3-(3''-malonyl-glucoside)                 | 0.59      | 2.14      | 0.009   | −1.67           |
| 679  | Erysodine                                          | 0.56      | 2.02      | 0.030   | −1.64           |
| 55   | Luteolin                                           | 1.00      | 2.00      | 0.006   | −1.61           |
| 196  | Trifolian                                          | 0.94      | 2.00      | 0.008   | −1.59           |

*The MS2 score ranged from 0 to 1 and the metabolite is matched more accurate when the MS2 score near to 1.*
the gene or genes endowing resistance (Vila-Aiub et al., 2015). Creating near-isogenic lines, selecting resistant and susceptible individuals from within a population, segregating F2 populations, and conducting experiments by comparing many resistant and susceptible populations are common methods to avoid the effects of genetic background (Vila-Aiub et al., 2015; Keshhtkar et al., 2019). For goosegrass, the second method is the most preferred way to eliminate the influence of genetic background and was also used in this study (Han et al., 2017; Chen et al., 2021). In addition, the mechanism of EPSPS overexpression was confirmed, and its expression level was 20.1–82.7 times higher in the R population than in the WT population. Similar expression levels were also found in other GR weeds that developed resistance via the same mechanism (Gaines et al., 2010; Malone et al., 2015; Ngo et al., 2018). Many studies use P450 and GST inhibitors to successfully explore the potential NTSR in weeds (Yuan et al., 2007; Han et al., 2021). These findings in this study suggest that no NTSR mechanisms involving P450s or GSTs exist in the R population.

At the later growth stage, a fitness benefit was found according to dry weight, leaf area, and seed production in this study. These results were similar to those of transgenic A. thaliana overexpressing EPSPS via the CaMV35S promoter, which resulted in a 30% increase in silique and seed number (Fang et al., 2018). In contrast to our results, plant growth and reproductive fitness traits in Amaranthus palmeri and Kochia scoparia were not affected by EPSPS overexpression, which was caused by the amplification of the gene in the genome (Vila-Aiub et al., 2014; Giacomini, 2015; Kumar and Jha, 2015). Furthermore, a fitness cost in seed weight production was found in a K. scoparia population with EPSPS amplification (Martin et al., 2017). The GR individuals of L. perenne, exhibiting 15-fold more EPSPS transcripts and 3-fold more EPSPS activity than the susceptible individuals, displayed a 40% reduction in the total number of seeds produced under field conditions (Yanniccarri et al., 2015). Overexpression of EPSPS can substantially increase EPSPS enzyme production, as confirmed in our previous studies (Chen et al., 2017b). In theory, the extra copies of EPSPS gene, transcript, and protein may lead to material and energy expenses (Akashi and Gojobori, 2002; Lynch and Marinov, 2015). However, different fitness traits in different weed species with EPSPS overexpression make it more complex. Fang et al. (2018) found that a fecundity advantage occurred in a transgenic A. thaliana lineage over-expressing EPSPS. This fitness trait was associated with increased levels of the auxin indole-3-acetic acid (IAA), which plays an important role in the regulation of plant growth and development (Fang et al., 2018). Studies have indicated that Trp is the main precursor for IAA in plants, and four proposed pathways for the biosynthesis of IAA from Trp have been identified in plants (Mashiguchi et al., 2011). In this research, metabolites in the Trp metabolism pathway showed significant differences between the R and WT populations. The levels of two important precursors of IAA synthesis – namely, 6-hydroxy-1H-indole-3-acetamide and indole acetaldehyde – were significantly higher in the R population than in the WT population (Table 2). This indicates that auxin also plays an important role in imparting beneficial fitness traits to the selected R goosegrass population in this study. The high concentration of free amino acids can be regulated, and catabolism of Tyr has been shown to return the highest energy in ATP currency in plants (Hildebrandt et al., 2015). It was hypothesized that the energy cost invested in the massive EPSPS amplification of glyphosate-resistant weeds would be compensated by catabolism of the excess amino acids, particularly Tyr, produced by the amplified EPSPS activity. To support this hypothesis, in this study, Tyr in the R population was slightly lower than that in the WT population, indicating that most of the excess Tyr has been catabolized.

In this study, vegetative and fecundity fitness benefits were found in the GR E. indica population. The results of metabolome analysis suggest that the increase in levels of 6-hydroxy-1H-indole-3-acetamide and indole acetaldehyde, which are related to auxin synthesis, is due to glyphosate resistance. Further studies should be conducted to confirm the functions of these metabolites.

DATA AVAILABILITY STATEMENT

The original contributions presented in the study are included in the article/Supplementary Material, further inquiries can be directed to the corresponding author.

AUTHOR CONTRIBUTIONS

JC designed the research. ZL, GZ, and DZ performed the experimental work and the data analysis. XL and HC provided helpful suggestions for the data analysis and manuscript revision. JC and ZL wrote the manuscript. All authors read and approved the final manuscript.

FUNDING

This research was funded by the National Transgenic Major Program of China (No. 2019ZX08013007-003).

ACKNOWLEDGMENTS

The authors are thankful to the National Transgenic Major Program of China for financial support.

SUPPLEMENTARY MATERIAL

The Supplementary Material for this article can be found online at: https://www.frontiersin.org/articles/10.3389/fpls.2021.776990/full#supplementary-material
REFERENCES

Achary, V. M. M., Sheri, V., Mann, M., Panditi, V., Borphukan, B., Ram, B., et al. (2020). Overexpression of improved EPSPS gene results in field level glyphosate tolerance and higher grain yield in rice. Plant Biotechnol J. 18, 2504–2519. doi: 10.1111/pbi.13428

Akashi, H., and Gojobori, T. (2002). Metabolic efficiency and amino acid composition in the proteomes of Escherichia coli and Bacillus subtilis. Proc. Natl. Acad. Sci. U.S.A. 99, 3695–3700. doi: 10.1073/pnas.062526999

Baerson, S. R., Rodriguez, D. J., Tran, M., Feng, Y., Biest, N. A., and Dill, G. M. (2002). Glyphosate-resistant goosegrass Identification of a mutation in the target enzyme 5-enolpyruvylshikimate-3-phosphate synthase. Plant Physiol. 129, 1265–1275. doi: 10.1097/wpp.0100000000001560

Chen, J., Cui, H., Ma, X., Ma, Y., and Li, X. (2020a). Distribution differences in the EPSPS gene in chromosomes between glyphosate-resistant and glyphosate-susceptible goosegrass (Eleusine indica). Weed Sci. 66, 33–40. doi: 10.1017/wsc.2019.72

Chen, J., Huang, W., Wei, S., Cui, H., Li, X., and Zhang, C. (2020b). Glyphosate resistance in Eleusine indica: EPSPS overexpression and P106A mutation evolved in the same individuals. Pestic. Biochem. Phys. 164, 203–208. doi: 10.1016/j.psbio.2020.01.014

Chen, J., Huang, W., Wei, S., Zhang, C., and Huang, Z. (2015a). Characterization of glyphosate-resistant goosegrass (Eleusine indica) populations in China. J. Integr. Agric. 14, 919–925. doi: 10.1016/S2095-3119(14)60910-2

Chen, J., Huang, H., Zhang, C., Wei, S., Huang, Z., Chen, J., et al. (2015b). Mutations and amplification of EPSPS gene confer resistance to glyphosate in goosegrass (Eleusine indica). Planta 242, 859–868. doi: 10.1007/s00218-015-2324-2

Chen, J., Huang, Z., Huang, W., Wei, S., Yan, L., Jiang, C., et al. (2017a). Selection of relatively exact reference genes for gene expression studies in goosegrass (Eleusine indica) under herbicide stress. Sci. Rep. 7:46494. doi: 10.1038/srep46494

Chen, J., Jiang, C., Huang, W., Wei, S., Huang, Z., Wang, H., et al. (2017b). Characterization of Eleusine indica gene with mutation or amplification in EPSPS to glyphosate. Pestic. Biochem. Phys. 143, 201–206. doi: 10.1016/j.pestbp.2017.09.012

Chen, J., Wei, S., Huang, H., Cui, H., Zhang, C., and Li, X. (2021). Characterization of glyphosate and quinazolof-p-ethyl multiple resistance in Eleusine indica. Pestic. Biochem. Phys. 176, 104862–104862. doi: 10.1016/j.psbio.2021.104862

Delye, C. (2013). Unravelling the genetic bases of non-target-site-based resistance (NTSR) to herbicides: a major challenge for weed science in the forthcoming decade: unravelling the genetic bases of non-target-site-based resistance to herbicides. Pest Manag. Sci. 69, 176–187. doi: 10.1002/ps.3318

Deng, W., Yang, Q., Chen, Y., Yang, M., Xia, Z., Zhu, J., et al. (2020). Cyhalofop-butyl and glyphosate multiple-herbicide resistance evolved in an Eleusine indica population collected in Chinese direct-seeding rice. J. Agric. Food Chem. 68:2623. doi: 10.1021/acs.jafc.9b07342

Duke, S. O., and Powles, S. B. (2008). Glyphosate: a once-in-a-century herbicide. Pest Manag. Sci. 64, 319–325. doi: 10.1002/ps.1518

Fang, J., Nan, P., Gu, Z., Ge, X., Feng, Y.-Q., and Lu, B. R. (2018). Overexpression exogenous 5-enolpyruvylshikimate-3-phosphate synthase (EPSPS) genes increases fecundity and auxin content of transgenic Arabidopsis plants. Front. Plant Sci. 9:233. doi: 10.3389/fpls.2018.00233

Fernández-Moreno, P. T., Alcántara-de la Cruz, R., Smeda, R. I., and De Prado, R. (2017). Differential resistance mechanisms to glyphosate result in fitness cost for Lolium perenne and L. multiflorum. Front. Plant Sci. 8:1796. doi: 10.3389/fpls.2017.01796

Gaines, T. A., Zhang, W., Wang, D., Bukan, B., Chisholm, S. T., Shaner, D. L., et al. (2010). Gene amplification confers glyphosate resistance in Amaranthus palmeri. Proc. Natl. Acad. Sci. U.S.A. 107, 1029–1034. doi: 10.1073/pnas.0906691107

Ghanizadeh, H., and Harrington, K. C. (2019). Fitness costs associated with multiple resistance to dicamba and atrazine in Chenopodium album. Planta 249, 787–797. doi: 10.1007/s00425-018-3040-3

Giacomini, D. A. (2015). EPSPS Gene Duplication in Palmer Amaranth: Relative Fitness, Inheritance, and Duplication Mechanism of the Glyphosate Resistance Trait. Fort Collins, CO: Colorado State University.

Han, H., Vila-Aiub, M. M., Jalaludin, A., Yu, Q., and Powles, S. B. (2017). A double EPSPS gene mutation endowing glyphosate resistance shows a remarkably high resistance cost. Plant Cell Environ. 40, 3031–3042. doi: 10.1111/pce.13067

Han, H., Yu, Q., Belfa, R., González, S., Maiwald, F., Wang, J., et al. (2021). Cytochrome P450 CYP81A10v7 in Lolium rigidum confers metabolic resistance to herbicides across at least five modes of action. Plant J. 105, 79–92. doi: 10.1111/tpj.15040

Heap, I. (2021). The International Survey of Herbicide Resistant Weeds. [Online]. Available online at: www.weedsciences.org (Accessed August 31, 2021)
Steinrücken, H., and Amrhein, N. (1980). The herbicide glyphosate is a potent inhibitor of 5-enolpyruvylshikimic acid-3-phosphate synthase. *Biochem. Biophys. Res. Commun.* 94, 1207–1212. doi: 10.1016/0006-291X(80)90547-1

Vila-Aiub, M. M., Goh, S. S., Gaines, T. A., Han, H., Busi, R., Yu, Q., et al. (2014). No fitness cost of glyphosate resistance endowed by massive EPSPS gene amplification in *Amaranthus palmeri*. *Planta* 239, 793–801. doi: 10.1007/s00425-013-2022-x

Vila-Aiub, M. M., Gundel, P. E., and Preston, C. (2015). Experimental methods for estimation of plant fitness costs associated with herbicide-resistance genes. *Weed Sci.* 63, 203–216. doi: 10.1614/WS-D-14-00062.1

Vila-Aiub, M. M., Neve, P., and Powles, S. B. (2009). Fitness costs associated with evolved herbicide resistance alleles in plants. *New Phytol.* 184, 751–767. doi: 10.1111/j.1469-8137.2009.03055.x

Vila-Aiub, M. M., Yu, Q., and Powles, S. B. (2019). Do plants pay a fitness cost to be resistant to glyphosate? *New Phytol.* 223, 532–547. doi: 10.1111/nph.15733

Wang, J., Li, X., Li, D., Han, Y., Zheng, L., Yu, H., et al. (2018). Non-target-site and target-site resistance to AHAS inhibitors in American sloughgrass (*Beckmannia syzigachne*). *J. Integr. Agric.* 17, 2714–2723. doi: 10.1016/S2095-3119(18)62021-0

Yang, X., Li, L., Jiang, X., Wang, W., Cai, X., Su, J., et al. (2017). Genetically engineered rice endogenous 5-enolpyruvoylshikimate-3-phosphate synthase (epspS) transgene alters phenology and fitness of crop-wild hybrid offspring. *Sci. Rep.* 7:6834. doi: 10.1038/s41598-017-07089-9

Yannicciari, M., Vilaaiub, M., Istilart, C., Acciaresi, H., and Castro, A. M. (2015). Glyphosate resistance in perennial ryegrass (*Lolium perenne*) is associated with a fitness penalty. *Weed Sci.* 64, 71–79. doi: 10.1614/WS-D-15-00065.1

Yu, Q., Jalaludin, A., Han, H., Chen, M., Sammons, R. D., and Powles, S. B. (2015). Evolution of a double amino acid substitution in the EPSP Synthase in *Eleusine indica* conferring high level glyphosate resistance. *Plant Physiol.* 167, 1440–1447. doi: 10.1104/pp.15.00146

Yuan, J. S., Tranel, P. J., and Stewart, C. N. (2007). Non-target-site herbicide resistance: a family business. *Trends Plant Sci.* 12, 6–13. doi: 10.1016/j.tplants.2006.11.001

Zhang, X., Zhang, R., Li, L., Yang, Y., Ding, Y., Guan, H., et al. (2020). Negligible transcriptome and metabolome alterations in RNAi insecticidal maize against *Monolepta hieroglyphica*. *Plant Cell Rep.* 39, 1539–1547. doi: 10.1007/s00299-020-02582-4

**Conflict of Interest:** 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.

**Publisher’s Note:** All claims expressed in this article are solely those of the authors and do not necessarily represent those of their affiliated organizations, or those of the publisher, the editors and the reviewers. Any product that may be evaluated in this article, or claim that may be made by its manufacturer, is not guaranteed or endorsed by the publisher.

Copyright © 2021 Li, Li, Cui, Zhao, Zhai and Chen. This is an open-access article distributed under the terms of the Creative Commons Attribution License (CC BY). The use, distribution or reproduction in other forums is permitted, provided the original author(s) and the copyright owner(s) are credited and that the original publication in this journal is cited, in accordance with accepted academic practice. No use, distribution or reproduction is permitted which does not comply with these terms.