First Report of Glyphosate-Resistant Biotypes of Eleusine Indica (L.) Gaertn. in Europe

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Abstract: Glyphosate-resistant biotypes of Eleusine indica (L.) Gaertn. have been detected in Asia, the Americas but not in Europe. The aim of this study was to evaluate resistance levels and possible target site resistance mechanisms of an E. indica biotype (19-1) collected from a plant nursery in Southern Italy where poor glyphosate efficacy was reported. Two dose-response experiments were conducted to evaluate the sensitivity of biotype 19-1 to glyphosate in comparison with two susceptible checks. 5-enolpyruvylshikimate-3-phosphate synthase (EPSPS) sequencing was performed to identify possible mutations conferring the resistance. The susceptible biotypes were completely controlled at the glyphosate recommended field dose of 360 g ae ha⁻¹, while 50% of the plants of biotype 19-1 survived at 1440 g ae ha⁻¹. The resistance index of biotype 19-1 ranged between 5.8 and 7.3 for the response variables of fresh weight reduction and plant survival, respectively. All the plants surviving glyphosate application and sampled for DNA analyses had the point mutation P106A. The biotype 19-1 can be confirmed as glyphosate-resistant, representing the first glyphosate-resistant population of E. indica in Europe.

Keywords: goosegrass; glyphosate resistance; herbicide resistance; plant nursery

1. Introduction

Glyphosate inhibits the enzyme 5-enolpyruvylshikimate-3-phosphate synthase (EPSPS), hindering plant production of aromatic amino acids and thus leading to plant death. Glyphosate is the most used herbicide worldwide, mainly due to the rapid diffusion of genetically modified (GM) glyphosate-tolerant crops across millions of hectares in North and South America and Australia [1,2]. It has been extensively used for decades as the main weed control tool in perennial tree crops, plant nurseries, gardens, parks and other non-agricultural areas. The main reasons for the success of glyphosate are the wide spectrum of weed species controlled, including some troublesome perennials, its limited toxicity and low cost [1,2]. However, the widespread, often exclusive, use of glyphosate over a prolonged period and across huge cultivated areas is causing a continuous selection pressure on weed communities, inevitably leading to the evolution of weed biotypes resistant to this herbicide [2]. The first glyphosate-resistant (GR) biotype of Lolium rigidum Gaud. was reported in 1996 in Australia [3] and since then the number of resistant populations has been increasing in different countries. The first glyphosate-resistant (GR hereinafter) biotype was reported in 1996 in Lolium rigidum Gaud. from Australia [3] and since then the number of resistant species (or biotypes) has been increasing in different countries. GR biotypes are
currently reported for 50 weed species (24 dicots and 26 monocots), across all continents [4]. Globally, most GR biotypes evolved in GM glyphosate-tolerant crops such as maize, soybean, oilseed rape and cotton [5]. In Europe, the cultivation of GM glyphosate-tolerant crops is not allowed, therefore the situation is completely different. Most European GR weed biotypes have evolved in perennial crops, such as orchards, olive groves and vineyards, and belong to the genus Conyza [6–8] and Lolium [9,10]. More recently, GR biotypes of Sorghum halepense (L.) Pers. and Hordeum murinum subsp. leporinum (Link) Arcang were found in Spain [11,12].

Eleusine indica (L.) Gaertn., or goosegrass, is an annual diploid grass species (2n = 18). Considered native to Africa and Asia [13], it is currently distributed in warm and temperate areas worldwide and is listed among the most problematic weeds [14]. In Italy E. indica was reported for the first time in 1879 [15] but until the early 1980s it was not very widespread [16]. After that its spread increased, favored by modification of land use and climate change, reaching the current status of “invasive alien of anthropic environments” [17]. E. indica has C₄ physiology and it grows fast in full sunlight and warm conditions, while shading by plant canopy strongly limits its growth [18]. E. indica is consequently a major weed in annual row-crops with limited canopy or slow growth, but it can also colonize the edges of annual crops such as maize and rice. In Europe, E. indica is prevalent in nurseries, orchards, ridges, and roadides due to its tolerance to diverse environmental disturbances such as heat, drought and mowing.

A wide range of herbicides have been used to control E. indica in various crops; however, resistant biotypes to several herbicides with different Sites of Action (SoA) have been reported worldwide since the 1970s [4]. E. indica was the second species to evolve resistance to glyphosate, with the first GR population detected in 1997 in a fruit orchard in Malaysia [19]. Since then, GR populations have been reported in other Asian countries, such as China [20], the Philippines [21], Indonesia [22], but also Brazil [23], Mexico [24] and the USA [25]. E. indica is one of the most important weed species in terms of number of countries and crops where GR populations have been reported [5]. Moreover, E. indica populations with multiple resistance to glyphosate, glufosinate, paraquat and acetyl CoA carboxylase (ACCase) -inhibiting herbicides [26] or glyphosate and cyhalofop-butyl [27] have recently been described. Target site resistance mechanisms, i.e., single and double amino acid substitutions at positions 102 or 106 of the EPSPS are known to confer glyphosate resistance to E. indica. The single amino acid substitution at position 106 is the most common, although it confers a low-to-intermediate resistance level [28–30]. Four different amino acid substitutions at position 106 (proline to serine P106S, proline to threonine P106T, proline to leucine P106L, proline to alanine P106A) have been described [21,29,31,32]. P106S is the most frequent substitution in all areas where GR biotypes of E. indica occurred, such as Malaysia [32], the Philippines [21], Mexico [24], the USA [30], and Brazil [33].

A double mutation in EPSPS at positions 102 and 106 (T102I + P106S (TIPS)) was identified in some highly resistant E. indica populations [28,34]; however, this double EPSPS mutation has a relevant fitness cost for the species [35]. Gene amplification of EPSPS is another resistance mechanism conferring low resistance level, if present alone [29,31,36]. However, highly resistant populations, with both EPSPS amplification and P106S or P106A mutations, have recently been identified in Mexico and China [24,29].

Herbicide resistance has never been reported for E. indica in Europe or the Mediterranean area [4]. However, poor control after repeated glyphosate applications was recently reported in a plant nursery in Southern Italy. Therefore, a study was conducted to confirm the presence of a glyphosate-resistant biotype and investigate the potential involvement of a target site resistance mechanism.

2. Materials and Methods

2.1. Seed Collection and Plant Growing Conditions

Seeds of the putative GR biotype, named 19-1, of E. indica were collected at a plant nursery in Southern Italy (41°10’ N, 14°09’ E, Caserta province) in autumn 2019 from plants surviving repeated
glyphosate applications. Limited glyphosate efficacy on *E. indica* had been observed in this farm and the surrounding areas since 2018, but it only became problematic for the farm operations in 2019. *E. indica* plants were growing in an area of the farm where pots with ornamental plants were maintained in outdoor conditions and the ground covered with plastic mulch. Plants of *E. indica* emerged mostly in gaps or holes of the mulch and only seldom in the pots. The limited competition by other weeds, thanks to the presence of plastic mulches, and the absence of shading by crop canopy allowed these *E. indica* plants to grow rapidly, reach a large size and produce many seeds (Figure 1). Other plants of *E. indica* emerged along the margins of the gravel roads between the different fields of the plant nursery. Even if environmental conditions were probably less favorable here, those plants were also fully grown and developed.

![Figure 1. Plants of Eleusine indica (L.) Gaertn. that emerged from holes and gaps of plastic mulches. These plants were survivors to repeated glyphosate applications.](image-url)
The standard weed control strategy in cultivated and uncultivated areas of the plant nursery was based on weed patch spraying (3–4 applications per year) with glyphosate (2.6 g ae L\(^{-1}\) of spray solution). Herbicide application was performed using a weeding lance equipped with spray hood for localized weed control. When the farmer noticed in summer 2019 that this E. indica biotype completely survived the herbicide treatments, he decided to perform an additional application (5 applications in total) but without achieving better control. Two additional populations (18-S and 19-S), collected in autumn 2018 and 2019 at Legnaro (45°20′ N 11°57′ E, North-eastern Italy) in fields where poor control with glyphosate had never been reported, were included in the experiments as susceptible reference checks.

After collection, seeds of the three biotypes were left to dry at room temperature (20 °C) for 2 weeks and then stored at 4 °C until the start of the experiment. Seeds were put in petri dishes filled with peat substrate and placed in a germination cabinet at a temperature regime of 28 (light) / 18 (dark) °C and a 12 h photoperiod. The cabinet was equipped with neon tubes providing a photosynthetic photon flux density (PPFD) of 15–30 µmol m\(^{-2}\) s\(^{-1}\). Seedlings were transplanted at the cotyledon stage into plastic trays (volume 6 L) filled with a mixed substrate (silt loam soil 62%, sand 35%, and peat 3%). Fifteen seedlings were transplanted into each tray. Trays were then placed in a greenhouse where favorable temperature and substrate moisture were maintained throughout the experiments.

2.2. Dose-Response Experiments

Two dose-response experiments were conducted to evaluate glyphosate efficacy on the biotypes. Herbicide application (Roundup Power 2.0, Monsanto Agricoltura Italia S.p.A., Via Giovanni Spadolini, 5, 20141 Milano, Italy; glyphosate content 360 g ae L\(^{-1}\)) was done at the 1–2 tillers or BBCH 21–22 growth stage [37] using a precision bench sprayer with a boom equipped with three flat fan hydraulic nozzles (TeeJet XR11002-VK, Glendale Heights, IL, USA), with a spray volume of 300 L ha\(^{-1}\) applied at a pressure of 215 kPa and speed of 0.75 m s\(^{-1}\). E. indica plants were treated with a range of glyphosate doses (22.5, 45, 90, 180, 360, 720, 1440, 2880 and 5760 g ae ha\(^{-1}\), i.e., from 1/16 to 16 times the recommended label dose of 360 g ae ha\(^{-1}\)). Only the putative GR biotype (19-1) was treated with the two highest doses (2880 and 5760 g ae ha\(^{-1}\)). Untreated controls were also included for all biotypes. The experimental layout was a completely randomized design with three replicates, i.e., three trays of 15 plants each. Plant survival was evaluated 3 weeks after treatment (3 WAT) and expressed as percentage of the plants counted before the treatment in each tray. Plants showing no active growth were considered as dead. Fresh weight of the above-ground biomass, including both dead and alive plants, was measured for each replicate and expressed as percentage of the mean value of the untreated replicates of the same biotype.

2.3. Statistical Analyses

Means and standard errors were estimated for plant survival and fresh weight of each treatment. Non-linear regression analysis was performed to estimate ED\(_{50}\) and ED\(_{90}\), (i.e., the glyphosate dose required to kill 50% and 90% of the treated plants, respectively) and GR\(_{50}\) and GR\(_{90}\) (i.e., the glyphosate dose which causes a growth reduction of 50% and 90%, respectively in comparison to the untreated plants for each biotype) and their standard errors. This analysis was conducted using the Windows Excel\(^{®}\) macro BIOASSAY\(^{®}\) [38] and adopting a log-logistic model:

\[
Y = C + \left\{ \frac{D - C}{1 + \left( \frac{x}{I_{50} (OR I_{90})} \right)^b} \right\}
\]

where \(Y\) is the fresh weight or plant survival, \(C\) and \(D\) are the lower and upper asymptotes, \(I_{50}\) (or \(I_{90}\)) is the dose resulting in a 50% (or 90%) decrease in plant biomass or survival, and \(b\) is the slope. The upper asymptote of survival data was constrained to 100 for biological reasons and to improve the estimates.
of other parameters, whereas no constraints were imposed considering fresh weight data. Data of each biotype were analyzed as a single curve to estimate its specific parameters and the two experiments were first analyzed separately. An ad hoc lack-of-fit F-test was successively performed for each biotype, by comparing data of the two experiments, to assess whether the data could be pooled. This would allow a single set of parameters to be estimated for each biotype. In order to evaluate the level of glyphosate resistance of the putative GR biotype (19-1), a Resistance Index (RI) was estimated as the ratio between the ED$_{50}$ (or GR$_{50}$) of biotype 19-1 and the mean ED$_{50}$ (or GR$_{50}$) of the two susceptible reference biotypes (18-S and 19-S):

$$\text{RI} = \frac{\text{ED}_{50} \text{ or GR}_{50} (19 - 1)}{\text{Mean ED}_{50} \text{ or GR}_{50} (18 - S, 19 - S)}$$

(2)

2.4. EPSPS Sequencing

Genomic DNA (gDNA) was extracted from 10 plants of biotype 19-1 that were alive 3 weeks after being treated with the glyphosate dose of 360 g ae ha$^{-1}$ and four untreated plants of 19-S biotype. In brief, 2–3 fresh leaves were collected from each plant and frozen at −18 °C until DNA extraction; leaves were transferred into a 2 mL tube (with two disposable iron beads) and ground at room temperature with TissueLyser II (Qiagen, Hilden, Germany), with no buffer. After 1–2 min grinding, a cetrimonium bromide (CTAB) protocol for gDNA extraction was used [39]. The gDNA quantity and quality were determined by gel electrophoresis and analysis with a Nanodrop (Thermo Fisher Scientific, Waltham, MA, USA). Primers were designed to amplify the sites of EPSPS gene known to confer resistance to glyphosate if mutated (positions 102 and 106), using deposited EPSPS E. indica sequences (GenBank accessions JN004269.1 and JN004268.1). Codes and primer sequences (5′-3′) were: EI_EPSP_F2, TCTCAGGGAACAACTGTGGTG and EI_EPSP_R2, GGCAGTCAGTGCCAAGGAAA.

Amplifications were performed using GoTaq® G2 Hot Start Polymerase (Promega Corporation, Fitchburg, WI, USA) in a 25 µL mixture including 5 µL of 5x Green GoTaq Flexi Buffer, deoxynucleoside triphosphates (dNTPs) mix (0.2 mM each), MgCl$_2$ (3 mM), forward and reverse primers (0.4 µM), 0.125 µL GoTaq DNA Polymerase, and 25 ng DNA. The PCR program was as follows: 2 min at 95 °C, 35 cycles of 30 s at 95 °C, 30 s at 58 °C, 35 s at 72 °C, 5 min at 72 °C. PCR products were purified with NucleoSpin® Gel and PCR Clean-up kit (Macherey-Nagel GmbH & Co., Düren, Germany) following the manufacturer’s instructions. Once purified, amplicons were Sanger-sequenced by BMR Genomics (BMR Genomics S.r.l., Padova, Italy) with EI_EPSP_F2 and analyzed with FichTV (Geospiza Inc., Seattle, WA, USA). All sequences (ab1 files) were deposited in a dedicated repository [40].

3. Results

3.1. Dose-Response Experiment

Significant differences in the response to glyphosate application were observed in both experiments between the putative GR biotype (19-1) and two susceptible reference biotypes (18-S and 19-S). The susceptible biotypes were completely controlled (survival 0%) at 360 g ae ha$^{-1}$, while the putative GR biotype (19-1) showed 100% and 46 ± 5.4% of plant survival at 720 and 1440 g ae ha$^{-1}$, respectively. Plant survival significantly decreased at the dose of 2880 g ae ha$^{-1}$, but complete control of biotype 19-1 was achieved only at the dose of 5760 g ae ha$^{-1}$.

Since the lack-of-fit F-test identified no significant differences between repeated experiments, the data were pooled. The two susceptible biotypes (18-S and 19-S) had a very similar response. Biotype 19-1 had a lower response to glyphosate application in terms of both plant survival and fresh weight (Figures 2 and 3). Different values were estimated for the ED$_{50}$ and GR$_{50}$ of the putative resistant and susceptible biotypes (Table 1). ED$_{50}$ and ED$_{90}$ values of the susceptible biotypes were lower than 250 and 400 g ae ha$^{-1}$, respectively, while those of biotype 19-1 were estimated at around 1400 and 2000 g ae ha$^{-1}$, respectively. Similarly, GR$_{50}$ and GR$_{90}$ values of the susceptible
biotypes were lower than 250 and 500 g ae ha\(^{-1}\), respectively, while GR\(_{50}\) and GR\(_{90}\) of biotype 19-1 were approximately 1000 and 2000 g ae ha\(^{-1}\), respectively. The estimated Resistance Indexes (RI) of biotype 19-1, in comparison with the mean of the two susceptible biotypes, were 7.3 and 5.8 when calculated with plant survival or fresh weight reduction, respectively.

**Figure 2.** Plant survival response (%) to increasing glyphosate dose of the putative GR biotype (19-1, red line) and the two susceptible reference biotypes (18-S and 19-S, blue and green lines, respectively). Values are the mean of the two experiments, i.e., six replicates. Bars represent standard errors.

**Figure 3.** Fresh weight reduction response (expressed as % of the untreated control) to increasing glyphosate dose of the putative glyphosate-resistant (GR) biotype (19-1, red line) and the two susceptible reference biotypes (18-S and 19-S, blue and green lines, respectively). Values are the mean of the two experiments, i.e., six replicates. Bars represent standard errors.
Table 1. Values of effective dose of glyphosate for 50% and 90% estimated plant death (ED\text{50} and ED\text{90}) and fresh biomass reduction (GR\text{50} and GR\text{90}) for the putative GR biotype (19-1) and the two susceptible reference biotypes (18-S and 19-S), plus Resistance Index (RI) of the putative GR biotype.

| Biotype | ED\text{50} g ae ha\textsuperscript{-1} | ED\text{90} g ae ha\textsuperscript{-1} | RI  
<table>
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<tbody>
<tr>
<td>19-1</td>
<td>1402 ± 20.9</td>
<td>2030 ± 172.1</td>
<td>7.3</td>
</tr>
<tr>
<td>18-S</td>
<td>199 ± 30.7</td>
<td>264 ± 146.1</td>
<td></td>
</tr>
<tr>
<td>19-5</td>
<td>183 ± 7.6</td>
<td>295 ± 44.9</td>
<td></td>
</tr>
</tbody>
</table>

| Biotype | GR\text{50} g ae ha\textsuperscript{-1} | GR\text{90} g ae ha\textsuperscript{-1} | RI  
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</thead>
<tbody>
<tr>
<td>19-1</td>
<td>999 ± 49.5</td>
<td>1931 ± 206.4</td>
<td>5.8</td>
</tr>
<tr>
<td>18-S</td>
<td>192 ± 20.5</td>
<td>480 ± 124.0</td>
<td></td>
</tr>
<tr>
<td>19-5</td>
<td>50 ± 12.0</td>
<td>311 ± 56.0</td>
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Mean values plus standard errors. Resistance Index (RI) was estimated as the ratio between the ED\text{50} (or GR\text{50}) of the 19-1 and the mean ED\text{50} (or GR\text{50}) of the two susceptible reference biotypes (18-S and 19-S).

3.2. EPSPS Sequencing

All plants had the wild type codon ACT (threonine) at position 102, while all the (10) plants of biotype 19-1 sampled among the survivors from glyphosate dose of 360 g ae ha\textsuperscript{-1} had the point mutation CCA to GCA at position 106 of the EPSPS gene (Figure 4), which caused an amino acid change from proline to alanine (P106A). The presence of a single peak in the electropherograms indicated that all plants were homozygous for this mutation. No other polymorphisms were observed along the sequence obtained with that primer combination. No mutations of the EPSPS gene were detected in any plants (four) of the susceptible biotype 19-S.

Figure 4. Screenshot of chromatograms obtained from a susceptible plant of 19-S biotype (upper graph) and a resistant plant of biotype 19-1 (bottom graph). Only codons from 102 to 106 are shown.

4. Discussion

Biotype 19-1 has an intermediate level of glyphosate resistance, with an RI of 7.3 and 5.8 for plant survival and fresh weight, respectively. ED\text{50, 90}, GR\text{50, 90} and RI estimated for biotype 19-1 are in keeping with previous studies on GR populations of \textit{E. indica} with target site resistance due to single
amino acid substitution at position 106 in EPSPS gene [23,28–30]. Higher levels of resistance were instead described in E. indica populations with the double amino acid substitution at positions 102 and 106 (TIPS) [28,34] or having EPSPS amplification plus the substitution at position 106 [24,29].

All plants of biotype 19-1 were homozygous for the point mutation CCA to GCA at position 106 of the EPSPS gene, which caused an amino acid change from proline to alanine (P106A). E. indica is a self-pollinating species, thus homozygosity is expected some generations after the start of the selection process. Consistently, a previous work did not find heterozygous plants for mutations at this locus [41]. Heterozygosity was observed in the case of E. indica with the double point mutation called TIPS (T102I/P106S), but this might be due to the high fitness cost of that mutation [35]. Since all the sampled plants of biotype 19-1 had the same point mutation and consequently the same amino acid substitution (P106A), this is likely the main glyphosate resistance mechanism for this biotype.

The amino acid substitution P106A detected in the EPSPS gene of 19-1 plants has been reported only once for E. indica and in a single GR biotype collected in tea plantations in China [29], but it has already been described for other grass species such as Lolium rigidum and Poa annua [42,43]. Considering both results of the dose-response experiment and molecular analysis, the biotype 19-1 can be confirmed as glyphosate-resistant, therefore, representing the first case of GR and more generally herbicide-resistant population of E. indica in Europe.

Although glyphosate doses above 2880 g ae ha⁻¹ also controlled biotype 19-1, other control tools and tactics should be used. Control efficacy could be lower under field conditions, due to environmental factors or non-optimal application. The use of glyphosate as the only weed control tool could lead to an increased frequency of GR biotype in the local E. indica population or even to the evolution of further resistance mechanisms, such as the double 102–106 substitution or the EPSPS gene amplification, and consequently an increase in the resistance level. Integrating glyphosate with other herbicides with different SoA is recommendable for chemical control of E. indica in this area. Unfortunately, resistant biotypes to several herbicides and SoAs, such as the protoporphyrinogen oxidase (PPO) - inhibitor oxadiazon [44], ACCase-inhibitors [26,27] or the photosystem II inhibitor metribuzin [45], have already been reported in several countries. Specific strategies, such as rotation and/or a mixture of herbicides with different SoA, should be adopted to avoid the potential evolution of resistance to other herbicides. Organic acids with herbicidal effect, such as pelargonic acid, can be interesting management tools for this resistant weed. Their application provokes fast and nonselective desiccation of plant green tissue and the evolution of resistance against this effect is believed unlikely since multiple targets are affected by their action [46]. Mechanical and physical control methods could in theory play an important role for the sustainable management of this GR biotype. However, they are hardly adoptable in a plant nursery due to the presence of plant pots, irrigation pipes and plastic mulches.

Elucidating the process that led this GR biotype to establish itself is relevant for designing management strategies to limit the selection of glyphosate resistance across E. indica populations in other areas. It is suspected that biotype 19-1 evolved from a susceptible local population because of the high selection pressure exerted by the repeated use of glyphosate as the only weed control tool in the plant nursery. Otherwise, biotype 19-1 could have been introduced to that plant nursery from areas where GR populations of E. indica are common. Seeds of E. indica could be transported inside pots of ornamental plants or potting substrates that were regularly imported from abroad by the farm. However, the presence of an amino acid substitution (P106A) previously described only once in a E. indica population collected in China [29], instead of the more common P106S, might support the theory of in situ evolution instead of introduction of the GR biotype from other world areas.

To minimize the risk of GR E. indica biotype spreading, the overall management should take into consideration both processes. That is reducing the selection pressure exerted by glyphosate, by integrating alternative herbicides and control tools, to avoid in situ evolution and monitoring imported plants and substrates to promptly identify potential introduction of GR biotypes. Given that E. indica is a self-pollinating species, the spread of glyphosate resistance via pollen flow is
negligible. Efforts should be directed to avoid dissemination of GR biotype, removing all plants that survived herbicide treatments and minimizing the risk of seed transport by machinery, plant materials, or substrates. Given that *E. indica* has a longer seed longevity in comparison with other grass weed species, with high seed viability reported even after three years of burial [47], these strategies should be repeated for several years in order to significantly reduce the soil seed bank of the 19-1 GR biotype at local level.


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**References**

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23. Takano, H.K.; Oliveira, R.S.; Constantin, J.; Braz, G.B.P.; Gheno, E.A. Goosegrass resistant to glyphosate (Eleusine indica) and -susceptible biotypes of Eleusine indica (L) Gaertn) population. Pestic. Biochem. Physiol. 2017, 562–566. [CrossRef]


26. Jalaludin, A.; Yu, Q.; Powles, S.B. Multiple resistance across glufosinate, glyphosate, paraquat and ACCase-inhibiting herbicides in an Eleusine indica population. Weed Res. 2015, 55, 82–89. [CrossRef]


30. Huffman, J.L.; Riggins, C.W.; Steckel, L.E.; Tranel, P.J. The EPSPS Pro106Ser substitution solely accounts for glyphosate resistance in a goosegrass (Eleusine indica) population from Tennessee, United States. J. Integr. Agric. 2016, 15, 1304–1312. [CrossRef]


41. Molin, W.; Wright, A.; Nandula, V. Glyphosate-resistant goosegrass from Mississippi. *Agronomy* 2013, 3, 474–487. [CrossRef]


43. Cross, R.B.; McCarty, L.B.; Tharayil, N.; McElroy, J.S.; Chen, S.; McCullough, P.E.; Powell, B.A.; Bridges, W.C. A Pro 106 to Ala Substitution is associated with resistance to glyphosate in annual bluegrass (*Poa annua*). *Weed Sci.* 2015, 63, 613–622. [CrossRef]


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