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# Sulfur Application Amends Detoxification Processes in Eggplant in Response to Excessive Doses of Thiacloprid <sup>+</sup>

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Abstract: Sulfur is considered an essential macronutrient during plant growth and is found to play 11 critical roles in xenobiotics detoxifying processes in plants. In the present study, the effects of exog-12 enous sulfur treatment as additional fertilization on detoxifying enzymes activities and plant health 13 indicators were investigated in eggplant (Solanum melongena) seedlings exposed to excessive doses 14 of thiacloprid. Eggplant seedlings (cultivar Hansel F1) were irrigated with ammonium sulfate (140 15 mg L<sup>-1</sup>) at 14 days after sowing in combination with the spraying of a 4-fold recommended dose of 16 thiacloprid. In another treatment, seedlings received ammonium sulfate (70 mg L<sup>-1</sup>) as a minimum 17 sulfur need in their growth in combination with a mentioned dose of thiacloprid. After 14 days of 18 treatment, leaves were collected to determine their physiological parameters. Based on results, plant 19 health indicators including malondialdehyde, hydrogen peroxide, and electrolyte leakage index 20 were significantly lower in treatments that were received additional amounts of sulfur than other 21 ones. While, the activities of glutathione S-transferase, glutathione reductase, glutathione peroxi-22 dase, thioredoxin reductase, and cytochrome P450 monooxygenase were higher in them. Our find-23 ings suggest that the sulfur can decrease membrane permeability and increase cell viability as well 24 as magnify their detoxification capacity which consequently leads to the reduction of oxidative 25 damage in plants. It can be concluded that the sulfur supply in eggplant farms that thiacloprid is 26 intensively used against sap feeder insects should be considered because it can lead to reducing 27 potential risk to the environment by decreasing pesticide damage to host plants as non-target or-28 ganisms. 29

Keywords: detoxifying enzymes; eggplant; health indicator; sulfur; thiacloprid

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# 1. Introduction

Sulfur is an essential macronutrient that is necessary for plants' growth and devel-33 opment. Changes in plants' sulfur content disrupt their tolerance against biotic and abiotic 34 stresses [1]. Also, sulfur is the main element of xenobiotic detoxification processes in 35 plants especially in thiolic compounds-based pathways [2]. Glutathione, as a non-protei-36 nous thiol, plays a key role in xenobiotic detoxification processes [3]. Of note, glutathione 37 S-transferase, glutathione reductase, and glutathione peroxidase enzymes are involved in 38 glutathione production, consumption, and detoxification processes [4]. Also, thioredoxin 39 reductase acts as a reducer for oxidized thioredoxin [5] and helps glutathione oxidation-40 reduction cycle to incorporate with glutathione reductase and glutathione peroxidase [6]. 41 Cytochrome monooxygenase P450 is found to be another important detoxifying enzyme 42 that catalysis oxidation reaction and changes chemical compounds of pesticides to sec-43 ondary metabolites [7]. 44

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In the present study, we hypothesized that the ammonium sulfur can mitigate the 1 phytotoxicity effects of thiacloprid high doses in eggplant seedlings. To investigate this, 2 detoxifying enzymes activity was examined in response to the 4-fold recommended dose 3 of thiacloprid in combination with ammonium sulfate. Furthermore, plant health indices 4 including malondialdehyde, hydrogen peroxide, and electrolyte leakage index were eval-5 uated to scrutinize plant health status and gain more understanding about oxidative stress 6 situation. 7

#### 2. Materials and Methods

Eggplant seeds (cultivar Hansel F1) were planted in 15 cm-diameter plastic pots of 9 sterilized soil composed of 1:1:2 cocopeat: peat moss: perlite. Plants were grown in a 10 greenhouse under controlled conditions of L16:D8 photoperiod, the temperature of  $26 \pm 2$ 11 °C, and 30–40% relative humidity. 12

The eggplant seedlings were irrigated with ammonium sulfate (140 mg L<sup>-1</sup>) 14 days 13 after growth. At the same time, they were sprayed with a 4-fold recommended dose (0.4 14 gr a.i./L) of thiacloprid (Actara<sup>®</sup> 25 WG, Syngenta, Switzerland). Simultaneously, a group 15 of control plants was irrigated with 70 mg L<sup>-1</sup> ammonium sulfate (the crucial sulfur con-16 tent for eggplants growth), in combination with the same dose of thiacloprid spray. Egg-17 plant leaves were collected 14 days after treatment to determine their physiological pa-18 rameters. 19

Glutathione S-transferase activity was evaluated using CDNB as a substrate [8]. The 20 absorbance of the reaction mixture that consisted of GSH (5 mM), CDNB (1 mM), and 21 phosphate buffer (50 mM, pH 7) was measured at 340 nm and its activity was calculated 22 using the extinction coefficient equal to 9.6 mM<sup>-1</sup> cm<sup>-1</sup>. Glutathione reductase activity was 23 measured according to Homayoonzadeh et al. [8]. An increase in absorbance at 412 nm 24 was observed due to the formation of TNB resulting from the reaction of DTNB (1 mM) 25 with GSH (10 mM) ( $\varepsilon$  = 14.15 mM<sup>-1</sup> cm<sup>-1</sup>), as a measure of enzyme activity. The glutathione 26 peroxidase activity was analyzed based on Herbette et al. method [9], which was meas-27 ured by monitoring NADPH oxidation at 340 nm. The reaction mixture contained Tris-28 HCl (100 mM, pH 7.5), EDTA (5 mM), NADPH (0.2 mM), and GSH (3 mM) ( $\epsilon$  = 6220 mM<sup>-1</sup> 29 cm<sup>-1</sup>). Thioredoxin reductase activity was measured following the method of Holmgren 30 and Bjornstedt [10]. This enzyme reduces DTNB (8 mM) to TNB by NADPH (0.25 mM) 31 and has an absorbance maximum at 412 nm ( $\varepsilon$  = 13,600 M<sup>-1</sup> cm<sup>-1</sup>). The activity of cyto-32 chrome P450 monooxygenase was investigated according to the method described by 33 Guengerich et al. [11]. The reaction mixture consisted of phosphate buffer (100 mM, pH 34 7.1), EDTA (1 mM), glycerol (20%, v/v), and sodium cholate (0.5%, w/v). Optical density of 35 mixture was recorded at 450 nm ( $\epsilon$  = 91 mM<sup>-1</sup> cm<sup>-1</sup>). 36

Malondialdehyde content was quantified using the TBA test as described by 37 Homayoonzadeh et al. [12]. The spectrophotometric measurement was performed at 532 38 using an extinction coefficient equal to 155 mM<sup>-1</sup> cm<sup>-1</sup>. Hydrogen peroxide content was 39 estimated according to the method of Homayoonzadeh et al. [13], which is based on KI 40oxidation by  $H_2O_2$  in an acidic medium. The absorbance of the reaction mixture including 41 phosphate buffer (10 mM, pH 7), KI (1 M), and TCA (0.1%, w/v) was measured at 390 nm 42 while using a standard curve of hydrogen peroxide. Also, the electrolyte leakage index 43 was estimated according to the method of Homayoonzadeh et al. [14]. The electrolyte 44 leakage index was estimated as the percentage of initial to final conductivity after placing 45 leaf discs at 25 °C for 3 h and then boiling at 105 °C for 4 min. 46

The experiments were designed and carried out in a completely randomized design 47 using five independent biological replicates. An unpaired t-test was used to compare the 48results between treatments. The trait means were compared with the Tukey test at the 0.05 probability level. All analyses were performed in GraphPad Prism version 8.2.0. 50

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## 3. Results

Results demonstrated that detoxifying enzymes activities were higher in plants 2 treated with the 4-fold recommended dose of thiacloprid combined with ammonium sul-3 fate (140 mg L<sup>-1</sup>) than in control. Specific activities of glutathione S-transferase, glutathi-4 one reductase, and glutathione peroxidase were significantly higher in treated eggplant 5 seedlings compared to control in 1.35-, 1.39-, and 1.40-fold, respectively. In addition, the 6 specific activities of thioredoxin reductase and cytochrome P450 monooxygenase experi-7 enced a similar trend with 1.60- and 1.62-fold significant increase, respectively in treated 8 eggplant seedlings (Table 1). 9

The analyses of plant health indicators illustrated significantly more accumulation of 10 malondialdehyde (1.66-fold), hydrogen peroxide (1.11 fold), and electrolyte leakage percent (1.23-fold) in control plants than treated ones (Table 2).

**Table 1.** Mean (±SE) specific activities of glutathione S-transferase (GST), glutathione reductase13(GR), glutathione peroxidase (GPX), thioredoxin reductase (TrxR), and cytochrome P450 monooxy-14genase (CYT P450) (µmol min<sup>-1</sup> mg<sup>-1</sup> protein) in tomato seedlings after exposure to ammonium sul-15fate (70 mg<sup>-1</sup> L) + 4-fold recommended dose of thiacloprid (control) and ammonium sulfate (14016mg<sup>-1</sup> L) + 4-fold recommended dose of thiacloprid (treated). Asterisks were used to show statistically17significant differences between treated and non-treated plants.18

Enzyme	Control	Treated	<i>t</i> -Value	<i>p</i> -Value
GST	$1.320 \pm 0.086$	1.789 ± 0.127 *	3.546	0.025
GR	$1.111 \pm 0.072$	1.546 ± 0.203 *	2.643	0.048
GPX	$1.006 \pm 0.058$	1.414 ± 0.313 *	3.619	0.034
TrxR	$1.239 \pm 0.095$	1.986 ± 0.441 *	4.217	0.043
CYT P450	$0.452 \pm 0.032$	$0.735 \pm 0.043$ *	4.186	0.024

**Table 2.** Mean (±SE) contents of malondialdehyde (MDA), hydrogen peroxide (H2O2) ( $\mu$ g g<sup>-1</sup> fresh19weight), and electrolyte leakage index (ELI) (%) in tomato seedlings after exposure to ammonium20sulfate (70 mg<sup>-1</sup> L) + 4-fold recommended dose of thiacloprid (control) and ammonium sulfate (14021mg<sup>-1</sup> L) + 4-fold recommended dose of thiacloprid (treated). Asterisks were used to show statistically22significant differences between treated and non-treated plants.23

Parameter	Control	Treated	t-Value	<i>p</i> -Value
MDA	$0.578 \pm 0.037$	$0.347 \pm 0.046$ *	2.766	0.014
H <sub>2</sub> O <sub>2</sub>	$8.565 \pm 0.407$	7.653 ± 0.291 *	4.619	0.025
ELI	$19.42\pm0.924$	$15.72 \pm 0.804$ *	3.802	0.037

### 4. Discussion

This study revealed that the sulfur application in eggplants could induce detoxifying 25 enzymes activity and then mitigate phytotoxicity in response to high doses of thiacloprid 26 as a common insecticide in sap feeder insects' control. Sulfur, as the most abundant ele-27 ment in thiol groups, is essential in redox reactions and also acts as the modulator of de-28 toxifying' enzymes structure [15]. When sulfur is uptake by plants, it is inverted to amino 29 acids synthesis cycle especially methionine and cysteine that act as the intersection of pri-30 mary metabolism to form S-containing defense compounds. Also, excessive sulfur is 31 transported to leaves and stored in vacuoles to make a sulfur reserve for plant metabolism 32 such as detoxification processes [16]. 33

Thiacloprid, as a neonicotinoid insecticide, is metabolized in plants by a sulfoxidation reaction that results in producing —SO metabolite [17]. Thus, it seems that sulfur application enhances eggplant seedlings to overcome the detrimental impacts of thiacloprid high doses. 37

In this study, the specific activity of cytochrome P450 monooxygenase was increased 38 in eggplant seedlings that received an additional amount of sulfur in combination with a 39 high dose of thiacloprid. It has clear that cytochrome P450 monooxygenase is found to be 40

one of the key detoxifying enzymes in phase one and catalyzes the oxidation reaction to 1 make products of phase one reactions that are more hydrophilic than the parent xenobi-2 otic [18]. Thus, it seems that the additional sulfur application in eggplants results in an 3 activated phase one reaction. 4

Then, phase one products are detoxified through conjugation with plant metabolites 5 such as glutathione [18]. Glutathione reductase, glutathione peroxidase, and thioredoxin 6 reductase are enzymes that make a glutathione redox system to provide GSH for gluta-7 thione S-transferase activity. Glutathione S-transferase catalyzes the conjugation of the 8 GSH with phase one electrophilic compounds [19]. Demonstrating increased content of 9 glutathione reductase, glutathione peroxidase, thioredoxin reductase, and glutathione S-10 transferase activity, the present study indirectly reflects phase two reactions induction in 11 eggplants. 12

Based on observed results, plant health indices including malondialdehyde, hydro-13 gen peroxide, and electrolyte leakage index were improved in response to additional sul-14 fur access. Thus, it can be concluded that sulfur application is able to decrease mem-brane 15 permeability as well as increase cell viability. 16

To put it in a nutshell, sulfur application in eggplant farms where the thiacloprid is 17 used intensively should receive more attention. The sulfur solution can be used to mitigate 18 the deleterious effects of high doses of neonicotinoid insecticides on host plants. This may 19 benefit moderating pesticide potential risk to the environment, especially to the non-tar-20 get plants. 21

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