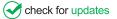


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## ARTICLE

# The Physiological and Molecular Responses of Exogenous Selenium to Selenium Content and Fruit Quality in Walnut

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#### **ABSTRACT**

To study the effect of exogenous selenium on fruit quality in walnut (Juglans regia L.), 8-year-old walnut (Qingxiang) was taken as the research object. In the fruit expansion stage, 300 mg/L of sodium selenate, yeast selenium and sodium selenite solutions were applied on the leaf of walnut, and the selenium levels in leaves, pericarp and kernel were determined at the ripening stage. The fruit quality, mineral nutrient content, antioxidant enzyme activity, and related genes' expression were analyzed. The results showed that the three exogenous selenium increased the selenium levels in leaves, pericarp and kernel of walnut. They also significantly increased fruit and kernel weights, and kernel linoleic acid, but markedly decreased kernel crude fat and saturated fatty acid. Selenium spraying promoted the absorption of mineral nutrients such as potassium and zinc, but inhibited the absorption of calcium, and had no significant effect on iron and magnesium in the kernel. Three exogenous selenium increased the activities of superoxide dismutase (SOD), peroxidase (POD), ascorbate peroxidase (APX), and catalase (CAT) significantly in the kernel. Except for sodium selenate treatment significantly reduced malondialdehyde (MDA) content in the kernel, the other two selenium sources treatments had no significant effect on MDA and hydrogen peroxide (H<sub>2</sub>O<sub>2</sub>) levels. They also increased the expression of JrCu/Zn-SOD, Jr2-Cys POD, JrCyt-APX and JrCAT in kernel, to different extents. These implies that, in the walnut fruit enlargement period, the foliar spraying selenium could increase the selenium content of walnut, affect the mineral nutrient absorption, improve the antioxidant capacity and related genes' expression, and reduce the degree of peroxide, and then improve the quality of fruit. Furthermore, yeast selenium showed the comprehensive effect of the best.

## **KEYWORDS**

Antioxidant system; fruit quality; mineral nutrient; selenium; walnut

#### 1 Introduction

As a trace element in the human diet, Selenium affects all stages of human growth and development [1]. Selenium intake could enhance the ability of oxidation resistance in the humans and prevent some diseases, including hepatopathy, tumor, cardiovascular and Kaschin-Beck disease [2,3]. Selenium as an antidote to heavy metals, it detoxifies arsenic, lead, cadmium and mercury in the human body, and also can combine



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with these heavy metals to form compounds that be excreted from the body [4]. In addition, selenium can prevent and treat diseases, regulate human immunity, and inhibit the occurrence of tumors and cardiovascular diseases [2]. It has various biological effects on the human body, such as delaying aging [3]. As a protective agent of life, selenium cannot be made by people themselves and must be obtained from food [5]. Therefore, how to supplement selenium safely and effectively becomes an urgent problem to be solved. There are mainly 2 forms of inorganic selenium and organic selenium in nature. Inorganic selenium is more toxic, and organic selenium can be safely absorbed by the human body [6]. In consequence, the conversion of inorganic selenium into organic selenium for human absorption and utilization has become an essential way of human safety selenium supplement. In the aspect of selenium transformation, plants have significant advantages because of their high selenium conversion rate and rich organic selenium content. Especially as human food sources, crops can be used as a simple and effective carrier of human safe selenium supplementation [7].

In recent years, with the improvement of national economic strength and people's living standard, the selenium-rich industry has received extensive attention from all walks of life. Australia, Denmark, the United States, Sweden, Germany and other countries have developed a variety of selenium-containing health products, selenium-rich agricultural products [8]. In China, Selenium-rich oil-tea, selenium-rich rice, selenium-rich potato, selenium-rich wheat, selenium-rich tea, selenium-rich edible fungus and other products have also been marketed and enjoyed by a vast number of consumers [9]. However, selenium-rich products and related research in the fruit field are still lacking. Therefore, it is particularly urgent to study the cultivation technology and regulation of selenium-rich fruit, explain the related mechanism, process selenium-rich fruit and research and develop high-value selenium-rich functional products.

The walnut, an important dried and oil fruit tree, is the most economically important member of the Juglandaceae family [10–13]. With the improvement of people's consumption level, healthy walnut fruits are paid more and more attention by the market, such as selenium-rich pecan kernel. Selenium-enriched walnut cultivation can not only improve the selenium level in fruits, but also improve the fruit quality [14]. Therefore, this study took 8-year-old walnut (Qingxiang) as the research object, and studied the effects of three different selenium sources on the selenium content in leaves, pericarp and kernel by spraying during the fruit expansion period. The results of three selenium sources on fruit quality, mineral nutrient content and antioxidant enzyme activity were analyzed. Hence, this study aimed to investigate the physiological responses of exogenous selenium to selenium content and fruit quality in walnut. Thus, it may set a new perspective and provide a firm theoretical basis for future about selenium-fertilizer application in walnut production.

## 2 Materials and Methods

## 2.1 Materials and Treatment

The 8-year-old walnut (Qingxiang) were provided by walnut demonstration orchard of Hubei Academy of Forestry, China. The soil fertility data were in Table 1.

рН Organic matter Nitrogen Phosphorus Potassium Calcium Magnesium Inorganic  $(mg \cdot kg^{-1})$  $(g \cdot kg^{-1})$  $(mg \cdot kg^{-1})$  $(mg \cdot kg^{-1})$  $(mg \cdot kg^{-1})$  $(mg \cdot kg^{-1})$ selenium  $(mg \cdot kg^{-1})$ 0.35 5.16 42.8 319.2 524.8 276.6 2.32 0.51

Table 1: The soil fertility of walnut orchard

Purified water, sodium selenate (Na<sub>2</sub>SeO<sub>4</sub>, 300 mg/L), yeast selenium (300 mg/L) and sodium Selenite (Na<sub>2</sub>SeO<sub>3</sub>, 300 mg/L) were sprayed on the leaf surface of walnut at its fruit enlargement period (May 20th,

2021). Na<sub>2</sub>SeO<sub>4</sub>, yeast selenium and Na<sub>2</sub>SeO<sub>3</sub> were purchased from Sigma Co., Ltd., Vienna, Austria. Each treatment was arranged in a complete randomized block design and replicated 8 times, produced a total of 32 walnut trees. Spray once a week with 4 times from 05/20/2021, each tree spray 3 L solution.

#### 2.2 Variables Measurement

Samples (leaves and fruits) were collected on September 20, 2021 (fruit ripening stage), while fruits were divided into pericarp and kernel. The leaf samples were selected from the top to bottom fourth leaf of the fruiting branch in the middle of peripheral branches of the tree crown, and the fruit samples were selected from the fruits in the middle of the tree crown with normal development and no diseases and insect pests. The collected samples were immediately placed in dry ice when washed with deionized water, brought back to the laboratory within 6 h, washed with deionized water, and stored in ultra cold storage freezer (-80°C) for later use.

Selenium contents in leaves, pericarp and kernel were determined by hydride-generation atomic fluorescence spectrometry (HG-AFS) (AFS8510; Beijing Haiguang Instrument Co., Beijing, China) according to Gui et al. [1]. Fruit and kernel weights were weighed by electronic balance. The rate of crude, linoleic, unsaturated fatty acid and saturated fatty acid in the kernel were determined as described by Velazquez et al. [15]. The concentrations of zinc (Zn), iron (Zn), potassium (K), calcium (Ca) and magnesium (Mg) in the kernel were measured by the inductively complied plasma-atomic emission spectrometry based on the protocol of Sun et al. [16].

Kernel samples (0.2 g) from 4 treatments were homogenized in 5 mL of 0.1 M phosphate buffer (pH 7.8) and centrifuged at 4,000  $\times$  g for 10 min at 4°C. And the supernatant was used to assay the concentrations of MDA and H<sub>2</sub>O<sub>2</sub>, and the activities of SOD, POD, APX and CAT, according to Liu et al. [17].

The kernel RNA was extracted in 0.1 g fresh sample using the RNA prep pure plant kit (DP441, TIANGEN Biotech Co., Ltd., China), and reverse transcription was done with the PrimescriptTM RT reagent kit with the gDNA Eraser (RR047A, TaKaRa Bio Inc., Japan). All the steps followed the manufacturer's instructions. The specific primers (Table 2) of relevant genes for qRT-PCR analysis were designed using Primer Premier 5.0 software (Palo Alto, USA), according to the Walnut genome database (http://aegilops.wheat.ucdavis.edu/Walnut/data.php). A housekeeping gene, *JrGAPDH*, was magnifified as an endogenous control along with the target genes. The relative expression of genes was calculated by the  $2^{-\Delta\Delta Ct}$  method.

Table 2: The specific primers of relevant genes designed for real time quantitative PCR amplification					
Gene name	Accession	Sequence (5'-3')-forward	Sequence (5'-3')-reverse		
JrCu/Zn-SOD	Jr1SG00045600	TGTACCCCTCCGAACCCA	CGGCTATTTCGTGATGG		
Jr2-Cys POD	Jr3SG00056000	TCGCTTGGAGACCACGCC	AATACTGACTGATCAAT		
JrCyt-APX	Jr4SG00075300	TTAGCTCGATCACGGATCT	AACAATTCAATGATTCC		

GGTTCTATTGCACAAGGA

ATTGTCCATGGTCAGGCC

GATGTTTATCATACCCTA

ATTTTGATGCAATCCCA

## 2.3 Data Statistical Analysis

Jr7SG00080400

Jr8SG00046000

**JrCAT** 

*JrGAPDH* 

SAS software (9.1.3) was used to analyze the difference significance of data by Duncan's new complex range method ( $P \le 0.05$ ). The data were analyzed using one-way Anova. When F tests were significant, means were separated by Duncan's new complex range method ( $P \le 0.05$ ).

#### 3 Results and Analysis

## 3.1 The Selenium Content in Walnut Leaves, Pericarp and Kernel

As shown in Table 3, three selenium sources improved the selenium content in varying degrees in leaves, pericarp and kernel of walnut. Compared with the water control group, yeast selenium treatment significantly increased selenium content by 191, 37 and 67 times in leaves, pericarp and kernel (Table 3). Na<sub>2</sub>SeO<sub>4</sub> and Na<sub>2</sub>SeO<sub>3</sub> treatments also dramatically increased its levels by 100 and 95 times in leaves, 24 and 23 times in pericarp, 64 and 60 times in kernel, respectively (Table 3).

Table 3: Selenium content in walnut leaves, pericarp and kernel under different selenium source treatments

Treatments	Leaves (µg·kg <sup>-1</sup> )	Pericarp (µg·kg <sup>-1</sup> )	Kernel (μg·kg <sup>-1</sup> )
Control	$104.3 \pm 10.1^{d}$	$80.7 \pm 5.2^{c}$	$15.6 \pm 1.3^{c}$
$Na_2SeO_4$	$10522.6 \pm 500.1^{b}$	$2010.1 \pm 104.8^{b}$	$1010.2 \pm 30.1^{ab}$
Yeast selenium	$20055.6 \pm 904.2^a$	$3004.2 \pm 200.8^a$	$1055.6 \pm 43.2^{a}$
Na <sub>2</sub> SeO <sub>3</sub>	$10011.3 \pm 508.1^{\circ}$	$2000.3 \pm 104.3^{b}$	$955.2 \pm 24.2^{b}$

Note: Control-purified Water,  $Na_2SeO_4$ -300 mg/L  $Na_2SeO_4$ , Yeast selenium-300 mg/L yeast selenium,  $Na_2SeO_3$ -300 mg/L  $Na_2SeO_3$ . Data (means  $\pm$  SD, n = 8) followed by different letters in same column are significantly different at P < 0.05. The same as below.

## 3.2 Fruit Quality Parameters

Fruit and kernel weights, the rate of crude, linoleic, unsaturated fatty acid and saturated fatty acid in the kernel were shown in Table 4. Na<sub>2</sub>SeO<sub>4</sub>, yeast selenium and Na<sub>2</sub>SeO<sub>3</sub> notably increased the weight of fruit and kernel by 14.1%, 25.3%, 9.1%, and 18.2%, 36.4%, 20.5%, respectively, compared to the water control (Table 4). With regards to the quality of kernel, Na<sub>2</sub>SeO<sub>4</sub>, yeast selenium and Na<sub>2</sub>SeO<sub>3</sub> dramatically decreased the kernel crude fat and kernel saturated fatty acid by 9.1%, 18.6%, 8.0%, and 7.5%, 13.6%, 6.3%, respectively, compared to the water control (Table 4). However, three exogenous seleniums markedly increased the rate of kernel linoleic acid at the same level. It is worth noting that there were no significant differences in the rate of kernel unsaturated fatty acid among all treatments.

Table 4: Walnut fruit quality under different selenium sources

Treatments	Fruit weight (g)	Kernel weight (g)	Kernel crude fat (%)	Kernel linoleic acid (%)	Kernel unsaturated fatty acid (%)	Kernel saturated fatty acid (%)
Control	$24.1 \pm 1.4^{c}$	$4.4\pm0.2^{c}$	$75.32 \pm 5.05^{a}$	$18.1 \pm 1.1^{b}$	$91.4 \pm 5.7^{a}$	$8.84 \pm 0.28^{a}$
$Na_2SeO_4$	$27.5\pm1.1^b$	$5.2\pm0.2^b$	$68.46 \pm 4.03^b$	$21.8\pm1.2^a$	$91.5 \pm 4.8^{a}$	$8.18 \pm 0.31^{b}$
Yeast selenium	$30.2 \pm 2.1^{a}$	$6.0\pm0.3^a$	$61.32 \pm 4.22^{c}$	$22.4 \pm 1.8^{a}$	$91.1 \pm 5.0^{a}$	$7.64 \pm 0.29^{c}$
Na <sub>2</sub> SeO <sub>3</sub>	$26.3 \pm 1.2^{b}$	$5.3 \pm 0.2^{b}$	$69.31 \pm 4.56^{b}$	$21.9 \pm 1.7^{a}$	$90.2 \pm 5.7^{a}$	$8.28 \pm 0.22^{b}$

# 3.3 The Mineral Nutrient Content in the Kernel

Table 5 shows the results of mineral nutrient contents in the kernel treated by three selenium sources. As can be seen from Table 3, the three selenium source treatments increased the contents of Zn and K in the kernel to varying degrees, but significantly reduced the level of Ca, while having no significant effect on Fe and Mg. Compared with water control, Na<sub>2</sub>SeO<sub>4</sub>, yeast selenium and Na<sub>2</sub>SeO<sub>3</sub> treatments increased Zn and K content by 13.7%, 22.4% and 12.5%, 30.5%, 50.2% and 30.5%, respectively (Table 5). In the

case of Zn, the yeast selenium was the only treatment significantly higher than the control. However, the Ca content in the kernel was reduced by 17.2%, 17.6% and 17.6%, respectively (Table 5).

Treatments	$Zn (mg \cdot kg^{-1})$	Fe (mg·kg <sup>-1</sup> )	$K(g \cdot kg^{-1})$	$Ca (g \cdot kg^{-1})$	$Mg (g \cdot kg^{-1})$
Control	$81.19 \pm 7.10^{b}$	$38.13 \pm 3.11^{a}$	$10.40 \pm 1.11^{c}$	$2.27\pm0.12^{\rm a}$	$2.13 \pm 0.16^{a}$
$Na_2SeO_4$	$92.33 \pm 8.09^{ab}$	$37.16 \pm 3.23^{a}$	$13.58 \pm 1.12^{b}$	$1.88\pm0.14^{b}$	$2.22\pm0.17^a$
East selenium	$99.41 \pm 9.07^{a}$	$38.20 \pm 2.92^a$	$15.62 \pm 1.14^{a}$	$1.87\pm0.12^{b}$	$2.24\pm0.21^a$
Na <sub>2</sub> SeO <sub>3</sub>	$91.32 \pm 8.11^{ab}$	$37.18 \pm 3.01^{a}$	$13.57 \pm 1.08^{b}$	$1.87 \pm 0.12^{b}$	$2.23 \pm 0.19^{a}$

Table 5: Mineral nutrient contents of kernel treated with different selenium sources

## 3.4 The Concatenation of MDA and the Activities of SOD, POD and APX in the Kernel

Table 6 shows the results of the activities of POD, SOD APX and CAT, and the contents of MDA and  $H_2O_2$  in the kernel of walnut treated with three selenium sources. Compared to the appropriate control,  $Na_2SeO_4$ , yeast selenium and  $Na_2SeO_3$  notably increased the activity of SOD, POD, APX and CAT by 28.9%, 76.3%, 28.1%, and 60.8%, 129.4%, 86.3%, and 9.7%, 24.5%, 17.8%, and 14.3%, 24.5%, and 12.9%, respectively (Table 6). However, yeast selenium and  $Na_2SeO_3$  had no significant effects on MDA and  $H_2O_2$  contents, while  $Na_2SeO_4$  decreased the level of MDA by 26.9%, compared to the water control (Table 6). It is worth noting that there were no significant differences among the three treatments and the control for  $H_2O_2$  (Table 6).

**Table 6:** SOD, POD and APX activities and MDA content in walnut kernel treated with different selenium sources

Treatments		POD (U·min <sup>-1</sup> g <sup>-1</sup> )	MDA (nmol·g <sup>-1</sup> )	APX $(U \cdot g^{-1})$	$\begin{array}{c} \text{CAT} \\ (\text{U} \cdot \text{min}^{-1} \text{g}^{-1}) \end{array}$	$\begin{array}{c} H_2O_2\\ (mg\cdot g^{-1}) \end{array}$
Control	$1.14 \pm 0.11^{c}$	$0.051 \pm 0.002^{c}$	$25.29 \pm 2.14^{a}$	$601.56 \pm 54.09^{c}$	$80.21 \pm 4.09^{c}$	$8.12 \pm 0.44^{a}$
Na <sub>2</sub> SeO <sub>4</sub>	$1.47\pm0.12^b$	$0.082 \pm 0.005^b$	$18.49 \pm 1.11^{b}$	$659.78 \pm 55.71^{b}$	$91.64 \pm 5.19^{b}$	$8.08 \pm 0.64^{a}$
Yeast selenium	$2.01 \pm 0.17^{a}$	$0.117 \pm 0.011^{a}$	$24.89 \pm 2.06^{a}$	$748.34 \pm 61.04^{a}$	$99.89 \pm 6.07^{a}$	$8.05 \pm 0.71^{a}$
$Na_2SeO_3$	$1.46\pm0.13^b$	$0.095 \pm 0.007^b$	$23.41 \pm 2.01^{a}$	$708.56 \pm 62.05^{ab}$	$90.55 \pm 5.05^{b}$	$8.01 \pm 0.69^{a}$

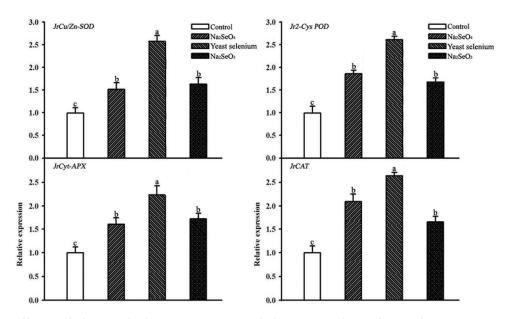
## 3.5 Relative Expression of Antioxidant System Responsive Genes

Fig. 1 shows the relative expression of antioxidant system responsive genes. Compared to the control, Na<sub>2</sub>SeO<sub>4</sub>, yeast selenium and Na<sub>2</sub>SeO<sub>3</sub> notably increased the expression of *JrCu/Zn-SOD*, *Jr2-Cys POD*, *JrCyt-APX* and *JrCAT* by 50.2%, 156.3% and 55.8%, 80.2%, 159.3% and 62.7%, 54.5%, 127.8% and 61.3%, 114.5%, 168.2%, and 68.9% respectively. Furthermore, there has no significant change of the expression of *JrCu/Zn-SOD*, *Jr2-Cys POD*, *JrCyt-APX* and *JrCAT* in between Na<sub>2</sub>SeO<sub>4</sub> and Na<sub>2</sub>SeO<sub>3</sub> treatments (Fig. 1). Interestingly, yeast selenium showed the positive effect of the best of this four genes' expression in this study (Fig. 1).

#### 4 Discussion

Compared with soil application of selenium fertilizer, foliar spraying of selenium can reduce selenium loss and effectively increase organic selenium content in plants, which is also a common method to increase organic selenium content in fruits or leaves in horticultural crop cultivation [18]. The suitable period and

species of selenium spraying on leaves of different horticultural crops are different. For example, it is the best time to apply selenium fertilizer to grape and strawberry at the beginning of fruit expansion [19,20]. Previous study showed that spraying selenate or selenite during the blueberry fruit expansion could effectively increase the selenium content and the anthocyanin concentration in fruit [21]. Therefore, in this study, three selenium sources were sprayed on the leaves during the walnut fruit expansion stage to study the effects of different selenium sources on walnut selenium content and fruit quality.



**Figure 1:** Effects of three selenium sources on relative expression of *JrCu/Zn-SOD*, *Jr2-Cys POD*, *JrCyt-APX* and *JrCAT* in walnut kernel

Spraying sodium selenite solution not only increased the selenium content of lettuce leaves, but also increased the transpiration rate, the net photosynthetic rate, and the stomatal conductance of leaves, improved photosynthesis and antioxidant capacity, and promoted its growth [22]. Similar study [23] on passion fruit has demonstrated that selenium can improve the photosynthesis, thus promoting the accumulation of sugar, and has a significant effect on the increase of soluble solid content, which can further improve the fruit weight, which is consistent with this study. The study of selenium application on cotton showed that selenium mineral fertilizer could advance cotton boll opening period, improve the cotton nutrient level, and promote boll growth and development [24]. The application of bio-selenium in the tomato cultivation process can obtain higher organic selenium content in the fruit, and improve its fruit quality significantly [25]. Similar results were observed in *Broccoli Florets* [26,27], where there was induced selenium level upon yeast selenium addition. This study showed that spraying sodium selenate, yeast selenium and sodium selenite at the expansion stage of walnut fruit could increase the selenium content in its leaves, pericarp and kernel, and yeast selenium-nich walnut.

Exogenous selenium fertilizer not only increased the selenium content in plants, but also affected the development and growth in plants, and then affected the quality of fruit. Studies showed that spraying sodium selenite on leaves could significantly increase grape fruit quality (the single grain weight, grain length, Vc content, selenium content, etc.) and delay fruit and leaf senescence [28]. Selenium rich liquid fertilizer significantly increased the content of soluble solids, fruit weight, less skin lesions, better luster, and 7 days earlier the ripening stage of peach [29]. Zahedi et al. [30] demonstrated that foliar spraying of

bio-nano selenium on a pomegranate could increase the soluble solid content of fruits, and its freshness and shelf life could be extended for 15 days. The results of this study were consistent with previous studies, three selenium sources all improved walnut fruit quality to varying degrees. In this study, fruit and kernel weights and the rate of kernel linoleic acid were increased, while the rates of kernel crude fat and kernel saturated fatty acid were decreased. Linoleic acid could reduce the risk of cardiovascular disease, as reflected by current dietary recommendations, while saturated fatty acid affects cholesterol metabolism [31,32]. So, exogenous selenium could improve walnut fruit's health value. However, the kernel crude fat was decreased by three selenium treated, which is necessary to maintain normal metabolism. Relevant mechanisms of it need to be further explored.

With regards to the effects of selenium on the uptake of mineral nutrients in plants, the results of this research showed that the foliar spraying of selenium promoted the uptake of Zn and K in kernel fruits, inhibited the uptake of Ca, but had no significant effect on the content of Fe and Mg. The results of study on selenium application in watermelon showed that spraying selenium at a lower concentration could promote the absorption of Ca and Mg, while spraying selenium at a higher concentration could inhibit the absorption of Ca and Mg [33]. Related mechanisms and the relationship between selenium and mineral nutrient absorption remain to be further studied and discussed.

SOD, POD and APX are scavengers of reactive oxygen species (ROS) in plants, which can remove excessive ROS in plant cells in the process of stress or aging process [34]. H<sub>2</sub>O<sub>2</sub> is one of the main components of ROS [34]. They can maintain the balance of cell metabolism, and improve the stress resistance of plants, which is related to the antioxidant capacity of plants [34,35]. MDA is one of the important products of membrane lipid peroxidation, and the higher content along with the stronger peroxidation degree [36]. In this study, the activities of SOD, POD and APX in walnut kernel treated with selenium spraying were significantly higher than those treated with clean water. At the same time, the contents of MDA and H<sub>2</sub>O<sub>2</sub> were lower than that treated with clean water to varying degrees. JrCu/Zn-SOD, Jr2-Cys POD, JrCyt-APX and JrCAT are regulate the plant antioxidant system [34]. At present study, exogenous selenium induced these genes' expression resulting in strengthen the activities of SOD, POD, APX and CAT in walnut kernel. It indicates that exogenous selenium can improve the antioxidant capacity of plants and protect the integrity of cell membrane by removing MDA and H<sub>2</sub>O<sub>2</sub>. This is consistent with the study in strawberry that selenium can not only protect the integrity of cell membrane, but also reduce the content of heavy metal ions, and effectively inhibit the absorption of heavy metal cadmium and lead in leaves and fruits [37]. In addition, among the three selenium source treatments in this study, yeast selenium had the strongest effect on improving the activities of SOD, POD, APX and CAT. So, yeast selenium can be applied to improve the antioxidant capacity of walnut.

#### 5 Conclusions

In a word, in the walnut fruit enlargement period, spraying three kinds of selenium fertilizer (sodium selenate, yeast selenium and sodium selenite) on the leaves can increase the selenium content of leaf and fruit, thus affecting the absorption of mineral nutrient, improve the antioxidant capacity and related genes' expression of the fruit, reduce the degree of peroxide, then improve the quality of walnut fruit. Among them, foliar spraying with yeast selenium had the best comprehensive effect. So it could be widely used in the cultivation of selenium-rich walnut. Future studies should pay more attention to the effect and mechanism of selenium on walnut quality.

**Authorship:** The authors confirm contribution to the paper as follows: study conception and design: Yuan L. Y., Zhang D. J.; data collection: Sun M. F., Hui X. R.; analysis and interpretation of results: Tong C. L.; draft manuscript preparation: Sun M. F. All authors reviewed the results and approved the final version of the manuscript.

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**Conflicts of Interest:** The authors declare that they have no conflicts of interest to report regarding the present study.

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