MAGNESIUM IN THE DYNAMICS OF CARBOHYDRATES AND ANTIOXIDANT METABOLISM OF COFFEE SEEDLINGS IN TWO IRRADIANCE LEVELS

Kaio Gonçalves de Lima Dias¹, Paulo Tácito Gontijo Guimarães², Antônio Eduardo Furtini Neto³, Valdemar Faquin⁴, Eduane José de Pádua⁵, Helbert Rezende Oliveira de Silveira⁶

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ABSTRACT: The aim of this study was to verify the physiological impacts and the carbohydrate dynamics of Coffea arabica, seedlings subjected to increasing concentrations of magnesium (Mg) and two irradiance levels. Methods: The experiment was carried out in growth chambers with nutrient solution. The treatments were five concentrations of Mg (0, 48, 96, 192, and 384 mg L⁻¹) and two irradiance levels (80 and 320 µmol of photons m⁻² s⁻¹). The coffee seedlings were under the treatments for 90 days. Results: The leaves with deficiency or excess of Mg exposed to the irradiance of 320 µmol of photons m⁻² s⁻¹ accumulated more carbohydrates than those exposed to 80 µmol of photons m⁻² s⁻¹. The accumulation of carbohydrates in the leaves increased the activity of antioxidant enzymes due to the increased production of reactive oxygen species (ROS). Leaves exposed to 320 µmol of photons m⁻² s⁻¹ exhibited symptoms of scald by the sun caused by photo-oxidation. The scald was more intense in plants with abnormal concentrations of Mg. The antioxidant system of the coffee tree is closely related to the Mg supply and irradiance levels. Concentrations of Mg between 48 and 96 mg L⁻¹ functioned as a mitigating agent of oxidative stress under stressful conditions caused by high irradiance level.

Index terms: Oxidative stress, enzyme activity, scald, coffee nutrition.

1 INTRODUCTION

Magnesium (Mg) is an important nutrient for plants because compose the chlorophyll molecules, which contribute to the energy metabolism of the plants (photosynthesis). Consequently, Mg deficiency affects many biochemical and physiological processes that decreasing the growth and yield of crops (CAKMAK; YAZICI, 2010).

Magnesium has a primary role in the transport of carbohydrates, especially sucrose. The accumulation of carbohydrates in leaves seems to be one of the first symptoms of Mg deficiency (SILVA et al., 2014). This elevated concentration of carbohydrates in leaves of plants deficient in Mg, accompanied by an increase in the ratio of the mass of the aerial part by the root is indicative of severe inhibition in the export of photoassimilates (sugars) in the phloem (CAKMAK; KIRKBY, 2008; CAKMAK; YAZICI, 2010). The reduction of transportation of carbohydrates and, consequently, the growth of the roots, decrease the water and nutrients absorption and thus decreasing the productivity of the crops.

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Additionally, the accumulation of carbohydrates in the leaves could stimulate the production of reactive oxygen species (ROS) which are toxic to plant species (MARSCHNER, 2012).

The irradiance levels also influence the biochemical and physiological processes in the plants, influencing severely the transport of carbohydrates and the growth of roots. Typical visual symptoms of Mg deficiency are clearly noted at high irradiance level (CAKMAK; KIRKBY, 2008).

The antioxidant skill of plants is considered an important factor in their protection against different environmental stresses (GILL; TUTEJA, 2010). The fundamental difference between the sensitive plants and those resilient to oxidative stress is the capability to reduce the damage caused by free radicals produced during the environmental stress (SÁNCHEZ-RODRIGUEZ et al., 2010).

Different stresses suffered by plants increase the production of ROS, for example, superoxide, hydrogen peroxide, and hydroxyl radicals that become harmful to the plant’s organism when their production is higher than the antioxidant agent’s production, thus resulting in oxidative stress (HUSSAIN et al., 2011).

An efficient degradation of ROS requires the joint action of enzymes of the antioxidant system — mainly superoxide dismutase (SOD), catalase (CAT), and ascorbate peroxidase (APX) (NEILL et al., 2008). Despite the presence of an effective antioxidant system, oxidative damage still occurs in the plant cells due to either the uncontrolled production or inefficient removal of ROS.

The climate change, especially in the last decade, may modify the areas of expansion of agriculture, and it could influence the plant’s growth, mainly in areas like Brazilian Cerrado, where plants would have to grow under a much higher irradiance than the light saturation point of the photosynthetic apparatus. Thus, a detailed study of Mg nutrition in coffee as a function of irradiance is required.

Therefore, the objective in this work was to determine the physiological impacts and carbohydrate dynamics in *Coffea arabica* L. seedlings with increasing concentrations of Mg and different irradiance levels.

**2 MATERIAL AND METHODS**

The experiment was carried out in growth chambers, under controlled conditions and with nutrient solution by Hoagland and Arnon (1950). The treatments were five concentrations of Mg (0, 48, 96, 192, and 384 mg L⁻¹) and two irradiance levels (80 and 320 µmol of photons m² s⁻¹); the standard concentration of Mg is 48 mg L⁻¹ (HOAGLAND; ARNON, 1950), and the irradiance level of 80 µmol of photons m² s⁻¹ simulating the conditions of low incident radiation, such as in shaded coffee trees within the plant canopy or in coffee trees in high-density plantations, whereas 320 µmol of photons m² s⁻¹ simulated normal conditions of photosynthetic activity.

Six repetitions of three-liter pots were used as growth chambers, with one coffee seedling per growth chamber, organized in randomized blocks of a 5 X 2 factorial arrangement (n=60).

Coffee seedlings of Mundo Novo IAC 379/19 cultivar with four pairs of true leaves cultivated in substrate not limed were transferred to trays containing deionized water for 10 days until the growth of new roots. The plants were transferred to three-liter pots containing 50% ionic strength of the Hoagland and Arnon (1950) solution, but excluding Mg, where they remained under constant aeration for 15 days.

After this period, the nutrient solution was replaced by 100% ionic strength of the Hoagland and Arnon (1950) solution and Mg was added at the concentrations of 0, 48, 96, 192, and 384 mg L⁻¹. The volume of the pots were fill with deionized water daily, pH was adjusted to 5.0–5.5 by addition of HCl 0.1 mol L⁻¹ or NaOH 0.1 mol L⁻¹, and the pots were kept aerated continually.

The lighting was provided by special daylight tubular fluorescent bulbs (Osram 20W). The levels of irradiance were controlled using shelves to regulate the distance between the plants and the light source. The irradiance levels at the different heights were measured using a quantum sensor (Licol LI-190SA, Li-Cor Biosciences Inc., Lincoln, USA). The plants were subjected to a photoperiod of 12 h of light and 12 h of darkness simulating a natural condition of tropical regions.

Fully expanded leaves were collected ninety days after the application of the treatments to performing the physiological analyses. After the material was collected, it was immediately preserved in liquid nitrogen and then stored at -80°C in a deep freezer. Subsequently, the plants were exposed to full sun for three days, in an irradiance of 1500 µmol of photons m² s⁻¹ and one more fully expanded leaf was sampled for the physiological analyses. The plants were then harvested and divided into leaves, stems and roots that were wash in deionized water, packed in paper bags, and dried at 60 °C until constant weights.
The dried leaves, stems, and roots were milled and a portion of the dry-milled material was sample for nutritional analysis following EMBRAPA (2009). Samples of dry-milled leaves were take for carbohydrate analysis.

2.1 Carbohydrates

The carbohydrates were extract from 25 mg of the dry mass of leaves, homogenized in 100 mM potassium phosphate buffer, pH 7.0, followed by a water bath at 40 ºC for 30 min (ZANANDREA et al., 2010). Subsequently, 1.0 mL of water was extracts through ultrasonication at 60 ºC for 15 min. An aliquot of 500 µL was transfers to 1.5 mL tube and centrifuged at 5,500 rpm for 5 min. The supernatant was diluted and filtered through a PES membrane with 0.22 µm pores (adapted from MELLINGER, 2006).

The following standards were use: for glucose and galactose analysis — Sigma-Aldrich (St. Louis, MO, USA); for sucrose and fructose analysis— Fluka (St. Louis, MO, USA). The mobile phase was prepared from 50–52% NaOH solution sourced from Sigma-Aldrich (St. Louis, MO, USA). All solutions were prepared using water obtained from a Milli-Q Biocel system from Millipore (Billerica, MA, USA).

The samples were analysed using an HPLC system (Shimadzu, Kyoto, Japan) coupled to an Antec DECADE II detector (Zoeterwoude, Netherlands) — the electrochemical cell was equipped with a gold electrode. The HPLC system, controlled by CBM-20A and with PEEK piping, was set to the isocratic flow of the 20 mM NaOH solution — which had been previously degassed in DGU-A3 — supplied at 0.2 mL.min -1 by an LC-10Ai pump. A volume of 20 µL of the samples was injects (model 7725i Rheodyne injector) — the separation was performed by a 250 x 4 mm internal diameter DIONEX CarboPac PA1 column (Sunnyvale, CA) equipped with pre-column and maintained at 44 ºC. The saccharides were analysed by pulsed amperometric detection, with the following detection conditions: E1 = +0.05 V, t1 = 500 ms, and t2 = 60 ms; E2 = + 0.75V and t2 = 130 ms; and E3 = -0.80 V and t3 = 120 ms (MARTINS et al., 2005).

The total soluble sugar (TSS) content was calculates by summing the sucrose, fructose, glucose, and galactose contents.

2.2 Antioxidant enzymes

The enzyme extract was obtained through the maceration in liquid nitrogen of 0.1 g of leaves, to which was added 1.5 ml of the extraction buffer containing 1.47 ml of 0.1 M potassium phosphate buffer (pH 7.0), 15 µL of 0.1 M EDTA (pH 7.0), 6 µL of 0.5 M DTT, 12 µL of 0.1 M PMSF, 0.001 M ascorbic acid, and 22 mg of PVPP. The extract was centrifuged at 12,000 g for 10 min at 4 ºC, and the supernatant was collected, stored at -20 ºC and used in the following enzymatic analyses: catalase (CAT), superoxide dismutase (SOD) and ascorbate peroxidase (APX) (BIEMELT et al., 1998).

a) Catalase

CAT activity was evaluated using a 5 µL aliquot of the enzyme extract, which was added to 950 µL of the incubation medium containing 200 mM potassium phosphate (pH 7.0) and 12.5 mM H2O2 and incubated at 28 ºC. The activity was determines by the decrease in the absorbance at 240 nm every 15 seconds for 3 min, monitored by the consumption of the hydrogen peroxide (HAVIR; MCHALE, 1987).

b) Superoxide dismutase

SOD activity was evaluated by the ability of the enzyme to inhibit the photoreduction of nitroblue tetrazolium (NBT) (MAROUANE et al., 2011) in an incubation medium consisting of 50 mM potassium phosphate at pH 7.8, 14 mM methionine, 0.1 mM EDTA, 75 mM NBT, and 2 mM riboflavin. The tubes with the reaction medium and the sample were illuminate for 7 min using a 20 W fluorescent lamp. The same reaction medium without the sample was illuminate for the control and the blank was kept in the dark. The readings were taken at 560 nm, and the concentration of SOD was calculated using the following equation:

\[
\text{SOD U} = \frac{A_{\text{sample}} - A_{\text{blank}}}{A_{\text{control}} - A_{\text{blank}}} \times \text{Sample Concentration} \times \text{SOD Unit}\]

c) Ascorbate peroxidase

APX activity was determined by monitoring the oxidation rate of the ascorbate at 290 nm every 15 s for 3 min. An aliquot of 5 µL of the enzyme extract was added to 950 µL of incubation buffer consisting of 500 µL of 200 mM potassium phosphate (pH 7.0), 50 µL of 10 mM ascorbic acid, and 50 µL of 2 mM hydrogen peroxide (NAKANO; ASADA, 1981). The molar extinction coefficient used was 2.8 mM·cm⁻¹.
2.3 Statistical analyses

Regression equations were estimate for the Mg concentrations. Mean tests (Scott-Knott) were complete to examine the differences in the irradiance levels at each Mg concentration. All analyses were performed using the Sisvar software (Ferreira, 2011), and graphs were built using the SigmaPlot 11.0 software. The maximum and minimum points of the quadratic functions were get through making the first order derivative equal to zero.

3 RESULTS AND DISCUSSION

3.1 Nutrient content

The content of Mg in leaves increased while K in leaves decreased with the concentration of Mg applied (Figure 1), fitted by quadratic functions for both irradiance levels. This reduction in the K leaf contents is due to the antagonistic effect of these nutrients (Figure 1b). In general, increasing the amount absorbed from one cation can result in the reduction of the absorption of another cation (MARSHNER, 2012).

3.2 Carbohydrate content

The levels of sucrose, reducing sugars fructose, glucose, and galactose and total soluble sugars (TSS) were significantly influence by the concentrations of magnesium (Mg) and by the applied irradiance levels, as well as by the interactions between these factors (Figure 2). The interaction between the Mg concentrations and the irradiance levels was significant for sucrose (Figure 2A), fructose (Figure 2B), glucose (Figure 2C), galactose (Figure 2D), and TSS (Figure 2E).

The leaves of coffee seedlings with 0 mg of Mg L⁻¹ (control) showed sucrose accumulation (Figure 2A). One of the first symptoms of Mg deficiency is the accumulation of sucrose (HERMANS et al., 2010; SILVA et al., 2014). The accumulation of sucrose in the leaves of plants deficient in Mg is related to the reduction in its transport to the roots via phloem, which causes a drastic reduction in root growth (CAKMAK et al., 1994; CAKMAK; YAZICI, 2010; HERMANS et al., 2010; SILVA et al., 2014) and decreasing absorption of water and nutrients, thus hampering the growth and productivity of the coffee plants.

Sucrose contents decreased with increasing concentrations of Mg up to 96 and 48 mg L⁻¹ for irradiance levels of 80 and 320 µmol of photons m⁻² s⁻¹, respectively. The influence of the reduction in photosynthesis on the slower transport of sucrose via the phloem is questionable because the decrease in the photosynthesis rate occurs in stages following Mg deficiency (CAKMAK; KIRKBY, 2008). Accumulation of carbohydrates in the leaves in low Mg content seems to result in adverse effects on photosynthetic gene activity, such as the gene responsible for the coding of chlorophylls a and b (HERMANS et al., 2010) which is partially responsible for the decline in both the chlorophyll contents and the photochemical performance in more advanced stages of Mg deficiency (CAKMAK et al., 1994; HERMANS et al., 2010).

![Figure 1](image_url)

**FIGURE 1** - Contents of Mg (a) and K (b) in leaves of coffee seedlings as a function of the application of different concentrations of Mg at two irradiance levels, 80 (closed circle) and 320 (empty circle) µmol of photons m⁻² s⁻¹. Significance according to t-test is indicated at 5% (*) and 1% (**).
FIGURE 2 - Levels of (a) sucrose, (b) fructose, (c) glucose, (d) galactose, and (e) TSS in leaves of coffee seedlings as a function of the application of different concentrations of Mg at two irradiance levels (80 and 320 µmol of photons m⁻² s⁻¹). Significance according to t-test is indicated at 5% (*) and 1% (**).
Thus, Mg has a direct effect on the transport of carbohydrates, especially sucrose, via the phloem (CAKMAK et al., 1994; CAKMAK; KIRKBY, 2008).

The function that Mg performs in the transport of carbohydrates could be related to the decrease in the metabolic activity of the source organs (WARAICH, et al., 2011). However, it is most likely relates to the decrease in the Mg-ATP concentration in the transport locations of the phloem (CAKMAK; KIRKBY, 2008; HERMANS et al., 2010). Depending on the species that transport may occur actively and selectively (symplastic pathway), thus requiring energy in the form of ATP for co-transport via the plasma membrane (H+) (TAIZ; ZEIGER, 2009).

The increases in the sucrose levels from the previously mentioned concentrations are probably related to the reduced absorption of K due to the excess Mg in the solution (figure 1). The higher sucrose accumulations were observed at the 320 µmol of photons m⁻² s⁻¹ of irradiance and the highest rate of photosynthesis was also related to the lower K levels observed.

K deficiency causes a reduction in the transport and use of photoassimilates (MARSCHNER, 2012). Cakmak et al. (1994) observed higher concentrations of sucrose and reducing sugars in the leaves of bean plants deficient in Mg and K. In addition to the accumulation of carbohydrates in the leaves, the role of K in the transport of carbohydrates is supported by the lower concentrations of sucrose in the roots of plants deficient in K compared with the roots of plants with adequate supply of this nutrient (CAKMAK et al., 1994). Therefore, K is required for the efficient transport of carbohydrates (sucrose particularly) via the phloem (PILOT et al., 2003).

In plants deficient in K, soluble carbohydrates and soluble nitrogen compounds accumulate, and the starch content decreases (MARTINEZ et al., 2014). These changes are related to the high requirement of K for the functioning of certain regulatory enzymes, in particular, pyruvate kinase and phosphofructokinase (MARSCHNER, 2012). According to Hermans et al. (2010) these reductions in photosynthesis rates observed in plants deficient in K may occur as a consequence of sucrose accumulation in the leaves due to their role in the synthesis and transport of carbohydrates. Moreover, K is extremely important in the activation of the carboxylase function of ribulose-1,5-bisphosphate carboxylase/oxygenase (Rubisco) (PRADO, 2008). Increases in photosynthetic rates due to the adequate supply of K have been attributed to these functions (CATUCHI et al., 2011; JIA et al., 2008).

At the higher Mg concentrations, there was greater accumulation of sucrose (Figure 2a), fructose (Figure 2b), glucose (Figure 2c), galactose (Figure 2d), and TSS (Figure 2e), especially in the plants subjected to the higher irradiance level (320 µmol of photons m⁻² s⁻¹). This likely indicates that plants grown under high irradiance levels require a greater supply of K to maintain transport of carbohydrates via the phloem at normal rates.

Fructose and glucose (sucrose precursors) were present at higher levels than galactose. This behavior was linear for the fructose, glucose, and galactose contents at the two studied irradiance levels (Figures 2b, 2c, and 2d).

The levels of TSS resulting from the Mg concentrations behaved quadratically at the lower irradiance level and exponentially at 320 µmol of photons m⁻² s⁻¹ of irradiance.

Seedlings in 0 mg L⁻¹ of Mg, showed higher contents of fructose (Figure 2b) and glucose (Figure 2c) at the highest irradiance level, and TSS (Figure 2e) increased at both irradiance levels. That indicates that Mg also plays a role in the transport of these sugars and/or in the sucrose transformation reactions. Silva et al. (2014) observed increases in the levels of sucrose, reducing sugars, and TSS as a function of Mg deficiency in coffee seedlings; however, the effect of excess Mg on the levels of reducing sugars (fructose, glucose, and galactose) — see Figures 2b, 2c, and 2d — was more significant. Perhaps K has a more important role in the transport and/or transformation of these sugars. The synthesis of carbohydrates is dependent on sucrose-cleaving enzymes, sucrose synthase (SuSy), and invertases, which also influence the velocity of transport in the phloem in the source/drain direction — some of these enzymes are activated by K (TAIZ; ZEIGER, 2009).

In general, there was a greater accumulation of carbohydrates at the extremes of Mg supply (0 and 384 mg L⁻¹), at the highest irradiance level, due to the higher photosynthesis rate probably (Dias 2015).

3.3 Relative distribution of carbohydrates

The Mg concentrations also altered the relative distribution of carbohydrates in the coffee leaves; that is, the ratio of the content of each carbohydrate relative to the total soluble sugar (TSS) — see Figure 3 — fit a decreasing quadratic function for the sucrose and increasing quadratic for the reducing sugars.

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The highest percentage of sucrose in relation to the other sugars was observed in the control, indicating that Mg deficiency has more influence on the accumulation of sucrose proportionally.

Regardless of the Mg concentration, sucrose was the sugar with the highest concentration in the leaves, followed by fructose, glucose, and then galactose, which had values below the others. According to Ding and Xu (2011) sucrose and starch are the main photosynthetic products in almost all of the higher plants.

Irradiance levels had no effect on the relative distribution of carbohydrates. On the other hand, there were changes in the carbohydrate content ratios with the different Mg concentrations. The proportions of sucrose: fructose: glucose: galactose were 74: 15: 10: 1% with 0 g L\(^{-1}\) of Mg to the plants. A supply of 48 g L\(^{-1}\) of Mg resulted in 59: 24: 15: 3%, while 96 g L\(^{-1}\) of Mg resulted in 57: 22: 18: 3%, respectively. The concentration of 192 g L\(^{-1}\) of Mg resulted in 56: 23: 19: 2% of sucrose, fructose, glucose and galactose of 384 g L\(^{-1}\) of Mg in 64: 20: 14: 2% sucrose, fructose, glucose and galactose.

Sucrose has a fundamental role in plant metabolism, acting as a source of carbon and energy, especially for non-photosynthetic tissues (BASSON et al., 2010). Reduction in the sucrose synthesis rate or increase in its degradation affects plant physiology, root development, and fruit quality (KÜHN; GROF, 2010).

The main enzymes that catalyze the sucrose dynamics reactions are sucrose synthase (SuSy), sucrose phosphate synthase (SPS), and the invertases (alkaline invertase and acid invertase). SPS controls the biosynthesis and accumulation of sucrose and it plays an important role in the translocation and distribution of photoassimilates in higher plants (TAIZ; ZEIGER, 2009; WANG et al., 2013). SuSy and alkaline invertase are primarily located in the cytosol, where as acid invertase is mainly associated with cell walls and vacuoles, where pH is approximately 5 (TAIZ; ZEIGER, 2009).

The catalyzed equations are:

- **Sucrose synthase:** Sucrose + UDP \(\rightarrow\) Fructose + UDP-Glucose
- **Sucrose phosphate synthase:** Sucrose + UDP \(\rightarrow\) Fructose + UDPG
- **Invertase:** Sucrose + H\(_2\)O \(\rightarrow\) Fructose + Glucose

In addition to the reactions shown, SPS could also irreversibly convert sucrose-6-phosphate into sucrose (PARK et al., 2009).

The importance of these enzymes depends on where sucrose is being metabolizes. SuSy is likely the main enzyme that degrades sucrose in organs that store starch (e.g., developing seeds or tubers) and in tissues undergoing rapid growth, which require translocated sucrose in the respiration process for the production of energy and
carbon skeletons. However, when the unloading of the phloem occurs passively (via apoplast), the acid invertase — because it is present in the cell wall — can convert the sucrose into hexoses (fructose and glucose) before entering the cell. In mature cells, cytosolic invertase may be important in the degradation of sucrose, supplying glucose and fructose for respiration (HOPKINS, 2000).

SPS is a key enzyme in the carbohydrate dynamics of plants — it is primarily responsible for the participation of sucrose in the physiological processes of different metabolisms. Previous studies of different crops have found a correlation between sucrose accumulation and increases in SPS activity or decreases in invertase activity (HIROTSU et al., 2007; ISHIMARU et al., 2008; ZHANG et al., 2010).

The relationships between sucrose and fructose and between sucrose and glucose were inversely proportional, suggesting that Mg and/or K participate in the activities of these enzymes. K has an established connection to SuSy activity (TAIZ; ZEIGER, 2009). Mg is possibly relates to invertase activity because the activity of these enzymes is generally inversely proportional to sucrose accumulation. However, more research is required regarding the role of these nutrients in the activity of these enzymes.

### 3.4 Antioxidant metabolism

The increase in Mg concentrations in the solution decreased the enzyme activity of the plant antioxidant system (SOD, CAT, and APX) — see Figures 4 and 5. For the SOD activity, the interaction between the Mg concentrations and the irradiance was not significant — the activity of this enzyme as a function of the concentrations was represent by a decreasing quadratic function (Figure 4). Increasing Mg concentrations up to 245 mg L⁻¹ caused a reduction in SOD activity, and from this point onward, the activity of this enzyme increased. SOD is responsible for the dismutation of O₂⁻ to form H₂O₂ and O₂. It is considered the first line of defense against ROS (KARUPPANAPANDIAN et al., 2011). CAT and APX are enzymes that catalyze the conversion of H₂O₂ to water and O₂ (ABEDI et al., 2010).

Increases in the activity of the antioxidant system resulting from Mg deficiency have been observed in the following crops: rice (CHOU et al., 2011), bean (CAKMAK; YAZICI, 2010), and coffee (SILVA et al., 2014).

Silva et al. (2014) observed the greater activity of the SOD, CAT, and APX enzymes in coffee seedlings deficient in Mg compared with those with an adequate supply of Mg.

Before the appearance of visual Mg deficiency symptoms, the activity of the enzymes from the antioxidant complex of the plant reduces the photooxidative damage caused by ROS and the inactivation of photosynthetic enzymes, which results in decreased photosynthetic activity only at more advanced stages of deficiency (KAISER, 1976).

The interaction between the Mg concentrations and the irradiance levels was significant for both APX and CAT activities. The activity of these enzymes as result of the Mg concentrations fit a descending quadratic function within each irradiance level, for both the growth chamber samples and the samples exposed to full sunlight (Figure 5). The significant interaction indicates that the activity of these enzymes as a function of the Mg concentrations depends on the irradiance.

The CAT and APX activities were both greater at lower Mg concentrations. According to Cakmak and Yazici (2010), high levels of components from the antioxidant metabolism are a physiological response of the plants to the effects of Mg deficiency.

The activation of antioxidant metabolism, under Mg deficiency, likely occurs in chloroplasts, the location of a reduction in O₂⁻ and H₂O₂ as a result of the restricted consumption of reducing potential in the fixation of CO₂ (Silva et al., 2014).

Excessive Mg also caused an increase in the enzyme activity of the antioxidant system (Figures 3 and 4) likely due to the physiological disorders caused by the excess of this nutrient itself, the reduction in the absorption of other nutrients, especially K, and the increase in the saline concentration of the solution.

Several studies have demonstrated the role of enzymatic antioxidant mechanisms in protecting against oxidative stress induced by salinity (RUBIO et al., 2009). The increase in the activity of enzymes such as SOD, APX, and CAT is associated with maintaining levels of lipid peroxidation under salt stress (ASHRAF, 2009).

K deficiency can increase the production of ROS in conditions of environmental stress particularly, for example, drought, high light intensity, heat, and nutrient limitation, and the improvement in potassium nutrition could greatly reduce the production of ROS through the reduction of NADPH oxidase activity and the maintenance of electron transport (Cakmak et al., 1994). K deficiency causes a reduction in the photosynthetic fixation of CO₂ and losses in the transport and use of photoassimilates. Such disorders can result in an excess of electrons, thereby stimulating the production of ROS (MARSCHNER, 2012).
In addition to the Mg content, light intensity also influences biochemical and physiological processes in which Mg is involved, causing visual symptoms typical of deficiency, the reduction of the transport of carbohydrates, root growth (CAKMAK; KIRKBY, 2008) and reductions in crop productivity consequently.

The saturation irradiances of photosynthesis in coffee trees are relatively low (300 to 700 µmol of photons m⁻² s⁻¹) (DAMATTA et al., 2004). Irradiances higher than those needed to saturate the photosynthetic complex can cause photoinhibition of photosynthesis. Additionally, they often lead to decreases in the rate of electron transport through photosystem II (PSII) and to increases in the spin rate of D1, the main polypeptide of the PSII reaction centers (NISHIYAMA, et al., 2011).
In general, during sampling within the chamber where the irradiance was controlled, the highest irradiance level (320 µmol of photons m\(^{-2}\) s\(^{-1}\)) triggered greater CAT and APX activities at 0 mg L\(^{-1}\) of Mg. With increasing Mg concentrations, the activity of these enzymes at the higher (320 µmol of photons m\(^{-2}\) s\(^{-1}\)) and lower (80 µmol of photons m\(^{-2}\) s\(^{-1}\)) irradiance levels tended to be equal (Figure 4). The effect of irradiance antioxidants enzyme activity became even more evident in the plants in the lower irradiance (80 µmol of photons m\(^{-2}\) s\(^{-1}\)) that were exposed to full sun, where the irradiance reached 1500 µmol of photons m\(^{-2}\) s\(^{-1}\). This change in irradiance expose caused stress, which increased the CAT and APX activities of the control; however, less activity occurred due to the increase in the Mg concentrations.

Cakmak and Yazici (2010) observed a rapid increase in the antioxidant mechanisms in bean plants with Mg deficiency, especially those exposed to high light intensities.

Increases in the formation of ROS occur when the absorption of light energy captured by the plant exceeds the utilization capacity during photosynthesis (Murchie and Niyogi, 2011). These excess ROS are toxic to the cells and destroy chlorophyll, the membranes, DNA, and other organelles. Under normal growth conditions, the accumulation of ROS in cells is low. However, adverse environmental factors that disturb cellular homeostasis induce the production of ROS, thus leading to oxidative stress (Miller, et al., 2010).

Additionally, even for the control, the coffee seedlings did not display the typical visual symptoms of deficiency in the growth chamber. After being exposed to full sun for three days, the typical visual symptoms of deficiency appeared, together with symptoms of scald, which appeared more intensely in the control plant and at the highest concentration of Mg due to the restriction in the absorption of K.

The oldest part of the leaf, which was shade by the younger leaf, remained green, whereas the portion that was exposed to full irradiance exhibited symptoms of scald (Figure 6). Note also that the newest leaf that was fully expose to irradiance, even with less physical protection, remained green (Figures 6 and 7). This occurred because the scald is the effect of photooxidative damage and is not a purely physical process. Mg and K, which are mobile in the plant, most likely have an important role in protecting newer leaves against scald.

At high light intensities, the development of chlorosis increases, together with some reddish spots on the leaf blade, and the parts of the leaves that did not receive the full luminosity were asymptomatic (Cakmak and Kirkby, 2008). Plants growing in high-intensity light conditions appear to have higher Mg demands than plants grown under low light intensity.

The increased oxidative stress observed as a function of the Mg deficiency, which was more evident in the plants exposed to higher irradiance levels, can be link to several factors:

–With the onset of the stress caused by Mg deficiency, carbohydrates accumulated in the leaves (Figure 1), as has been previously observed by several authors (Cakmak et al., 1994; Silva et al., 2014). This accumulation of carbohydrates changes the photosynthetic metabolism and reduces the use of the light energy absorbed in photosynthesis, leading to a saturation of the electron transport chain with the accumulation of NADPH (Hermans et al., 2010). High levels of reducing equivalents and components of the saturated electron chain offer favorable conditions for the formation of ROS (Biemelt et al., 1998; Mittler, 2002). 

–Mg also affects the activity of Rubisco, and the binding of this enzyme to Mg increases the affinity for CO\(_2\) and doubles the maximum reaction velocity (Sugiyama et al., 1968). Therefore, under Mg deficiency, the photosynthetic rate will decrease, thereby generating the accumulation of oxygen and reducing equivalents, which leads to the generation of ROS and, consequently, triggers oxidative stress. This process is more serious in high irradiance conditions because the generation of reducing equivalents and water absorption are both higher.

The results demonstrated that a relationship exists between the antioxidant complex of the coffee tree and the Mg supply as a function of the irradiance to which the plants are subject. Mg appears to work as a mitigating agent of oxidative stress under stress conditions caused by increased irradiance.

The results indicate the need for regional field studies, given that coffee is cultivated in various regions of Brazil and other countries with different irradiance levels, and thus, the requirements for Mg should not be the same for these different conditions.
4 CONCLUSIONS

Available Mg levels influence the carbohydrate dynamics in the leaves of coffee trees. Both the deficiency and excess of Mg cause increases in their carbohydrate content, especially sucrose in the coffee leaves. Coffee seedlings leaves are highest in sucrose content, followed by fructose and glucose, which are in turn greater than the amount of galactose present in the leaves.

The accumulation of carbohydrates in coffee tree leaves is dependent on the irradiance to which the plants are subject. Under conditions of deficiency or excess of Mg, leaves exposed to higher irradiances accumulate more carbohydrates.

The accumulation of carbohydrates in coffee tree leaves caused increases in antioxidant enzyme activity due to the greater production of ROS.

Deficiency or excess of Mg under high radiation conditions leads to intense photooxidation and symptoms of sun scalding in the coffee leaves.

There is a close relationship between the antioxidant complex of coffee trees and the Mg supply due to the irradiance to which the plants are subject where Mg acts as a mitigating agent of oxidative stress caused by increased irradiance.
Magnesium in the dynamics of carbohydrates ...

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