



E-ISSN: 2663-1067
P-ISSN: 2663-1075
IJHFS 2019; 1(2): 27-34
Received: 16-05-2019
Accepted: 18-06-2019

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Influence of macronutrient accumulation of “Magnum” watermelon by phosphate doses and application manners in calcareous soil

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Abstract

Information on the nutritional need of the watermelon crop is required particularly for P, due to the dynamics of this element in the soil and its low availability in many soils in the semi-arid region. The objective of this work was to evaluate the biomass production and the macronutrient extraction of the cv. Magnum irrigated under the influence of phosphate fertilization. The experiment was carried out in a medium textured Cambisol in Upanema, state of Rio Grande do Norte, in a randomized block experiment design, using the split-plot factorial scheme with four replications. The treatments consisted of 76, 168, 275 and 397 kg ha⁻¹ of P₂O₅, which were applied in two manners: F0 – 100% in pre-planting and F1 – in pre-planting + fertirrigation (50 kg ha⁻¹ as MAP). Two additional treatments were also applied: Adc1 – no phosphate; and Adc2 – with phosphate only in fertirrigation. Fertilization F1 provided greater accumulations of P and S, but, in general, the effects of the doses were dependent on the fertilization system, especially the treatments F0₂₇₅ and F1₁₆₈ regarding the accumulation of other nutrients.

Keywords: *Citrullus lanatus*, Fertirrigation, phosphorus

1. Introduction

Plant growth and nutrient uptake patterns are essential in crop nutrition and fertilization, especially for short-cycle crops such as watermelon, therefore meeting the plant demand over periods of the greatest demand (ALMEIDA *et al.*, 2014) ^[1]. However, the absorption and proportion between nutrients depend on both the cultivar and external factors that influence the process.

According to Fernandes and Soratto (2012) ^[7] several studies have shown that the nutritional level of phosphorus (P) may affect the absorption of other nutrients and consequently, the nutrition and production of the crops. Despite that, many studies consider only the ontogenetic variations in the absorption and utilization of the nutrients by the crop, neglecting the effects resulting from its nutritional management.

In the watermelon crop, source-drain relations undergo major changes after pollination, since the fruits are the favored drains in relation to the vegetative organs (LINS *et al.*, 2013) ^[12]. Under these conditions, mineral nutrients gain importance as productivity limiting factors, particularly when associated with a decline in the uptake by the roots (MARSCHNER, 2012) ^[15]. In general, P has been considered one of the most limiting nutrients, even in calcareous soils (MOHAMMAD *et al.*, 2004) ^[17] due to the reduced solubility of P compounds at higher pH and Ca and Mg precipitations (HOPKINS *et al.*, 2014) ^[10].

Silva *et al.* (2012) ^[22] explain that the absorption of P by the watermelon crop is small because of its low mobility in the soil and the great proportion that is adsorbed to the colloids. To reduce this effect, the general recommendation is to apply the total dose of phosphate in a localized way, in order to reduce contact with the soil.

For Mohammad *et al.* (2004) ^[17] the conventional application of P in pre-planting has the advantage of providing high concentrations in the soil solution at the initial stages of the crops, but a continuous supply of P through irrigation water (fertirrigation) may increase its absorption at the end of the production cycle, especially for crops with narrower root volume. Barreto and Fernandes (2002) ^[5] hypothesized that the mixture of phosphate fertilizer with higher soil fractions could in some cases increase the efficiency of use of phosphate fertilizer, especially in soils with low clay and iron and aluminum oxide contents, in contrast to typical tropical acid soils.

Thus, the objective of this work was to evaluate the macronutrient extraction of the cv. Magnum watermelon in calcareous soils under the influence of phosphate fertilization management.

2. Material and Methods

This study was conducted from September to November 2013 in a Cambisol of calcareous origin of the Jandaíra formation, in the municipality of Upanema, state of Rio Grande do Norte, (5°35'04" S, 37°12'08" W and 122 m above sea level). The predominant climate in the region is hot and dry – BSw^h type, according to the climatic classification of Köppen.

Over the experimental period, the average daily air temperature ranged between 26.9 and 28.2 °C and the relative humidity of the air was 65% (data from the automatic station of Mossoró A318 - INMET). Before the installation of the experiment, the 0-20 cm layer was sampled for physicochemical characterization (Table 1).

An initial sampling of the 0-20 cm layer was carried out for chemical characterization according to Silva (2009) [21] showing the following values: pH (H₂O) = 7.4; organic matter (in g kg⁻¹) = 23.86; P Mehlich (in mg dm⁻³) = 4; K⁺, Na⁺, Ca²⁺, Mg²⁺, Al³⁺ and H + Al³⁺ (in mmol_c dm⁻³) and V (%) = 5.60; 0.70; 48.4; 21.1; 0.0; 14.9 and 84. The contents of Cu, Fe, Mn, and Zn were extracted using DTPA solution pH 7.3, which was 0.8, 0.7, 17.9 and 7 mg dm⁻³, respectively and 24 mg L⁻¹ for the remaining P dosage obtained in 0.01 M CaCl₂ solution using 60 mg L⁻¹ of P of initial concentration in the 1:10 soil: solution ratio. The granulometric analysis indicated 228.87 and 685 g kg⁻¹ of clay, silt, and sand, respectively, in which the clay fraction was obtained by the pipette method; silt by the difference of mass between the total sample and the sand by sieving.

The water used in the irrigations had the following chemical characteristics: pH = 7.8; EC = 0.47 dS m⁻¹; Ca²⁺ = 2.25; Mg²⁺ = 0.89; K⁺ = 0.44; Na⁺ = 2.16; Cl⁻ = 1.31; HCO₃⁻ = 4.00; CO₃²⁻ = 0.16 (mmol_c L⁻¹).

The experiment was conducted in a complete randomized block design, in a factorial scheme (2 x 4 + 2) x 4, with two forms of fertilization management and four doses of P and two additional treatments using subdivided plots, corresponding to four periods.

Phosphate fertilization forms consisted of pre-planting application (F0) and pre-planting + topdressing (F1) of four P₂O₅ (76, 168, 275 and 397 kg ha⁻¹) doses. The pre-planting doses were hand applied using triple superphosphate (TSP) (41% P₂O₅), while the topdressing fertilization was carried out via irrigation water (fertirrigation) using mono-ammonium phosphate (MAP) (60% P₂O₅). Two additional treatments were also applied. Those treatments were the lack of P application (Adc₁) and application of P only in topdressing 50 kg ha⁻¹ P₂O₅ (Adc₂).

The soil was previously tilled by means of plowing and harrowing, rising mounds of 0.30 m in height and 0.60 m in width. The area of the experimental plots was 27.36 m² (14.4 mx 1.9 m), with a useful plot of 13.68 m² (7.2 mx 1.9 m), containing a total of eight plants spaced by 0.90 x 1.9 m. The 'Magnum' watermelon seedlings were produced in expanded polystyrene trays with 200 cells filled with Plantimax commercial farming substrate and coconut fiber. When the seedlings displayed two definitive leaves, they were transplanted to the area, seven days after emergence (DAE) or twelve days after sowing (DAS), on September

28, 2013.

Irrigation was performed daily through a drip system, with emitters spaced at 0.30 m. The irrigation system was evaluated three times over cultivation, according to Merriam and Keller (1978) [16] with the average flow and emission uniformity coefficients of 1.20 L h⁻¹ and 93.2%, respectively. Irrigation management was carried out by monitoring the soil moisture with the aid of tensiometers, in order to maintain soil with matrix potential higher than -30 kPa.

The fraction of the phosphate fertilization in pre-planting was hand applied in pits at each 0.30 m and next to each emitter. It was applied in all treatments 100 kg ha⁻¹ Barimicro fertilizer (FTE BR12) containing 1.8% B; 0.8% Cu; 2.0% Mn; 9.0% Zn; and 4.0% of S.

Nitrogen and potassium were applied via fertirrigation for all treatments, with totals estimated by the models developed by Paula *et al.* (2011) [18]. It was applied 140 and 100 kg ha⁻¹ N and K₂O, respectively, which were distributed from the week following transplanting. A further 7.6 kg ha⁻¹ CaO was applied between the 3rd and 4th week after transplanting. The sources used were urea, ammonium sulfate, potassium chloride, potassium nitrate, and calcium nitrate.

One plant was collected per plot at 28, 38, 54 and 69 DAE. The collected plant was divided into stem, leaves, and fruits, which were weighed, and the respective subsamples removed and taken to drying in a forced air circulation oven at 65 °C. After drying, they were weighed and the data used to deduction of the estimate of the total dry biomass.

For the evaluation of nutrient concentrations, the dry matter was first milled in Willey-type mill for chemical analysis according to Malavolta *et al.* (1997) [13]. Nitrogen concentrations were determined by the Kjeldahl semi-micro method, K by flame emission photometry, Ca and Mg by atomic absorption spectrophotometry, S by turbidimetry and P by metavanadate colorimetry. The accumulations of nutrients were obtained from the product between the dry biomass and their respective concentration in the material.

Data were submitted to analysis of variance, by the F test, at 5% of significance using the System for Analysis of Variance – SISVAR software, version 5.3 (FERREIRA, 2010) [8] and calculation worksheet. Afterward, regression analysis was performed and the models whose coefficients were significant by the test at 5% probability were selected, favoring those with a higher F and R² and that provided a simpler and more coherent explanation for biological behavior, using * and ** to indicate significance at 1% and 5% probability, respectively.

3. Results and Discussion

An increase was observed in the N accumulation rates until 44 DAE for most treatments, which can be explained by the stimulation of N accumulation in the vegetative organs, functioning as a nutrient reserve for the reproductive phase that is about to start, so that half of the total N is accumulated until 38-44 DAE (Figure 1A-1B).

For F0₂₇₅ treatment, an increase from 0.42 to 0.79 g day⁻¹ was found in the N accumulation rate per plant, between 28 and 42 DAE, and then, it reduced from there up to 0.20 g day⁻¹ at the end of the cycle (Figure 5A). On the other hand, in the F1₁₆₈ treatment, higher N accumulation rates were observed at 28 DAE (1.22 g day⁻¹), which was reduced to 0.13 g day⁻¹ at the end of the cycle (Figure 5B).

A higher maximum point for N accumulation was observed in F1₃₉₇, (23.08 g plant⁻¹ at 55 DAE), corresponding to approximately 134.95 kg ha⁻¹ N (Figure 5B), but it showed a significant reduction from that point, which can be explained by the loss of biomass (data not shown).

The F0₃₉₇ treatment presented a pattern similar to F1₃₉₇, but with less pronounced curvature at the point of maximum (53 DAE). The same occurs for F0₁₆₈ and F1₇₆, both with maximums at 57 DAE (Figure 5A-5B). This is more likely to have been caused by fruit abortion rather than leaf senescence since N can be easily remobilized from leaves.

In a study with the 'Tide' hybrid, Grangeiro and Cecilio Filho (2004) [9] found positive increases in N accumulation until the end of the cycle but observed a reduction in the vegetative part that was explained by a strong translocation for fruit development. The authors also indicated that N was the second most absorbed nutrient, accumulating 138.8 kg ha⁻¹ or 71 g plant⁻¹ (1,960 plants ha⁻¹).

In this work, N accumulations at the end of the cycle (69 DAE) followed the following decreasing order: 20.36 (F0₂₇₅), 16.94 (F1₁₆₈), 16.66 (F1₂₇₅) and 13.20 g plant⁻¹ (F1₃₉₇) or 119.04; 99.04; 97.41 and 77.18 kg ha⁻¹ N (5.847 plants ha⁻¹), respectively (Figure 5A-5B).

Working with cv. Olimpia, Silva *et al.* (2012) [22] reported a maximum N accumulation at 69 days after sowing (DAS) of 77.5 kg ha⁻¹ or 10.04 g plant⁻¹ (7,716 plants ha⁻¹), while Vidigal *et al.* (2009) [24] estimated 21.16 g plant⁻¹ or 105.8 kg ha⁻¹ N for cv. Crimson Sweet (at 89 DAS) with a density of 5,000 plants ha⁻¹.

The highest maximum N accumulation in Adc₂ (12.12 g plant⁻¹) (Figure 5B) in relation to F0₇₆ (11.70 g plant⁻¹) (Figure 5A) can be attributed to the synergistic effect between P and N, by the use of MAP in fertirrigation.

Araújo and Machado (2006) [4] discuss the role of P on the general regulatory effects of N absorption and assimilation. Fernandes and Soratto (2012) [7] found increased N concentrations in potato plants as P supply was increased, even after the stabilization of production from a certain level of P in the nutrient solution (>8 mg L⁻¹).

When studying the effects of the partial P supply to the roots of corn grown in nutrient solution, Alves *et al.* (1999) [3] observed a reduction in the accumulation of N in the aerial part when it is supplied separately from the P, attributing this result to the low concentration of P in the root.

Figure 2A shows a greater accumulation of P with F1 fertilization than F0, with a difference of 0.81 g plant⁻¹ at 69 DAE, therefore accumulating 50% of the total at 41 and 38 DAE, respectively. However, the application of 50 kg ha⁻¹ P₂O₅ only in the fertirrigation (Adc₂) resulted in slower initial P absorption since 50% of the respective maximum accumulation occurs at 44 DAE.

The application of all P in fertirrigation may result in nutrient deficiencies in the initial phase of the crop, as observed by Marouelli *et al.* (2015) [14] in a work with tomato. According to Mohammad *et al.* (2004) [17] different studies indicate that the P applied in pre-planting is available earlier to the plants, but its concentration in the soil solution declines over time due to the adsorption and precipitation reactions.

In this respect, despite the known benefits of localized application of phosphate fertilizers, the entire root zone needs P, so that the combination of fertilizer application techniques can be favorable, particularly in soils with low to medium levels of P (HOPKINS *et al.*, 2014) [10].

Around 50% of P totals was accumulated between 36 and 42 DAE and between 40 and 45 DAE at doses of 76 to 397 kg ha⁻¹ P₂O₅ in F0 and F1 fertilizations, respectively (Figure 2A-2B). This is the result of the effects of localized phosphate fertilization on the initial availability of P, as already discussed since the increases in the doses for both fertilizations are due to pre-planting phosphate increments.

Overall, the rapid observed accumulation of nutrients should be related to the experimental conditions, coinciding with the period in which 50% of the maximum accumulations occur for total biomass: between 37 and 46 DAE and between 41 and 47 DAE for the F0 and F1 treatments, respectively (data not shown)

The total variation in the P maximum accumulation in the factorial treatments is from 1.32 to 2.84 g plant⁻¹ (7.72 to 16.61 kg ha⁻¹), which is attributable to the differences between F1 and F0 fertilizations (0.81 g plant⁻¹) (Figure 2A) and between the doses of 397 and 76 kg ha⁻¹ P₂O₅ (0.71 g plant⁻¹) (Figure 2B). The accumulation of P in Adc₂ (1.48 g plant⁻¹) is equivalent to the 89 kg ha⁻¹ dose of P₂O₅ in F0 fertilization.

In work with cv. Magnum in sandy soil (8,333 plants ha⁻¹), Pereira *et al.* (2018) [19] also observed effects of phosphate fertilization on P accumulation, indicating an increase of 0.29 g plant⁻¹ when using the pre-planting + topdressing combination (0.96 g plant⁻¹), in relation to fertilization in pre-planting (0.67 g plant⁻¹), for a dose of 137 kg ha⁻¹ P₂O₅. The authors explained that the combination of application forms was most likely to have contributed to the increase in P availability in the soil.

It is worth noting that the effects on nutrient accumulation observed in the present work are derived from the combination of fertilization techniques rather than from the isolated application of P. Moreover, although some authors have concluded in favor of the topdressing application of P for better distribution in the root system (BARRETO and FERNANDES, 2002) [5] and to enlarge the availability of P in the soil, particularly in the application via fertirrigation (MAROUELLI *et al.*, 2015) [14] an increment in P absorption by the plant does not necessarily imply in proportional productivity gains (PEREIRA *et al.*, 2018) [19].

The application of part of the phosphate in topdressing may have favored a better distribution of the root system and thus, greater absorption of nutrients. Barreto and Fernandes (2002) [5] found greater root biomass and root distribution of maize under the effect of topdressing fertilization (broadcast) in comparison to pre-planting fertilization.

The accumulations of P found in this work are generally higher than those observed by Silva *et al.* (2012) [22] for cv. Olimpia, 8.12 kg ha⁻¹ or 1.58 g plant⁻¹ P with a density of 5,144 plants ha⁻¹, but lower than the maximum accumulation presented by Vidigal *et al.* (2009) [24] from 3.62 g plant⁻¹ (18.1 kg ha⁻¹ for 5,000 plants ha⁻¹) in cv. Crimson Sweet.

However, the work of Silva *et al.* (2012) [22] was carried out in Eutrophic Cambisol of limestone origin using 11.6 mg dm⁻³ P (Mehlich-1), where a total of 220 kg ha⁻¹ of P₂O₅ was applied, while that of Vidigal *et al.* (2003) was carried out in a sandy soil with 70.65 mg dm⁻³ of P (ion exchange resin), and the plants received the equivalent of 300 kg ha⁻¹ of P₂O₅ in addition to organic fertilization.

Differences in the accumulation of P in the literature may be the response of the genotype-environment interaction, therefore interfering in the speed of plant development and,

consequently, in the potential of accumulation of nutrients. The high harvest index (between 83% and 92% - data not shown), associated to a short cycle (69 DAE, with flowering and fruiting at 28 and 38 DAE, respectively) found in this study may have impaired the balance between source and sink, in order to adjust the input of assimilates to their demand (MARSCHNER, 2012; DUARTE and PEIL, 2008) ^[15, 6]. Lins *et al.* (2013) ^[12] explain that the watermelon crop undergoes great changes in source-drain relations after pollination, where the fruits are the favored drains.

In work using cv. Magnum in sandy soil, in Mossoró, state of Rio Grande do Norte, Pereira *et al.* (2018) ^[19] explained that the climatic conditions favored the shortening of the cycle (55 DAE, beginning of fruiting at 27 DAE) and rapid accumulation of nutrients (50% up to 2/3 of the cycle).

The highest K accumulations are found in treatments F0₂₇₅ (52.89 g plant⁻¹), F1₁₆₈ (47.17 g plant⁻¹), F1₃₉₇ (45.55 g plant⁻¹) and F1₂₇₅ (41.84 g plant⁻¹), corresponding to values between 245 and 309 kg ha⁻¹ K (Figure 3A-3B). Nevertheless, it should be noted that the maximum point for F1₃₉₇ occurred around 58 DAE, reducing from there to approximately 31.92 g plant⁻¹ K at 69 DAE, which is lower than that estimated for the F1₇₆ at the same period (34.56 g plant⁻¹). Because K is a high mobility nutrient, this reduction can be attributed mainly to the abortion of already formed fruits.

The K accumulations in this study are higher than those presented in a study developed in Cambisol Eutrophic with cv. Olímpia (5,114 plants ha⁻¹), 10.30 g plant⁻¹ (SILVA *et al.*, 2012) ^[22] and in Quartzarenic Neosol with cv. Crimson Sweet (5,000 plants ha⁻¹), 24.19 g plant⁻¹ (VIDIGAL *et al.*, 2009) ^[24].

On the other hand, Grangeiro and Cecílio Filho (2004) ^[9] found a K accumulation of 79.3 g plant⁻¹ by the hybrid Tide in a medium-textured dystrophic Red-Yellow Argisol, but with an expectation per area of 155.5 kg ha⁻¹ of K, due to the density of planting used in the experiment (1960 plants ha⁻¹).

This interference of plant density on the accumulation of nutrients was also observed by Paula *et al.* (2011) ^[18] who chose to use the relationships between productivity and nutrient exports per plant instead of area unit, in a study about the nutritional requirement of N, P and K in melon and watermelon crops.

At the intermediate doses of F1: F1₁₆₈ and F1₂₇₅ phosphate fertilizer, K accumulation rates were increasing, ranging from 0.88 to 1.38 g day⁻¹ and from 0.43 to 1.65 g day⁻¹, respectively. In F0₁₆₈, the K accumulation rate decreased from 1.05 to 0.54 g day⁻¹, between 28 and 69 DAE and in F0₂₇₅, the absorption rate increased to 0.93 g day⁻¹ at 40 DAE, reduced thereafter to 0.41 g day⁻¹, at 69 DAE (Figure 5E-5F).

The acceleration or stabilization of K uptake rates over time should be related to better distribution of the root system, which may increase nutrient uptake efficiency to a certain extent via diffusion (MARSCHNER, 2012) ^[15] although K uptake via mass flow can be more representative in conditions of its high concentration in the soil solution (RUIZ *et al.*, 1999) ^[20].

On the other hand, the reduction in K accumulation rates may be a consequence of the decline of nutrient uptake efficiency by the roots as the cycle advances, particularly at the beginning of the reproductive phase, where nutrient partitioning is prioritized and assimilated to the fruits, to the

detriment of root growth (MARSCHNER, 2012) ^[15].

Regarding Ca, the following order of total accumulation was found at the 69 DAE: F0₂₇₅>F1₁₆₈>F1₂₇₅>F1₇₆ (Figure 4A-4B). (Figure 4A-4B). Accumulation peaks of Ca ranged between 50 and 54 DAE, and an intense reduction occurred, ranging from 43% to 82% of the respective maximum (Figure 4A-4B), which should be associated with different proportions of leaf biomass caused by senescence.

This is because the movement of Ca in the plant occurring mainly in the xylem, via the transpiratory current, and because of its low redistribution by the phloem (GRANGEIRO and CECÍLIO FILHO, 2004; MALAVOLTA *et al.*, 1997; VIDIGAL *et al.*, 2009) therefore contributing for its concentration in the leaves rather than in the fruits ^[9, 13, 24].

The maximum Ca content is higher at the greatest doses of phosphate, corresponding to 20.32 and 14.36 g plant⁻¹, or 118.81 and 83.96 kg ha⁻¹ Ca, to F1₃₉₇ and F0₃₉₇, respectively (Figure 4A-4B), and can be explained mainly by the composition of the triple superphosphate (12% CaO).

Silva *et al.* (2012) ^[22] reported a Ca maximum equivalent to 3.46 g plant⁻¹ (17.8 kg ha⁻¹) for cv. Olympia and 5.58 g plant⁻¹ (28.7 kg ha⁻¹) for cv. Leopard grown in Eutrophic Cambisol originating from the Jandaíra limestone, in Baraúna, state of Rio Grande do Norte. On the other hand, Vidigal *et al.* (2009) ^[24] in a work with the Crimson Sweet hybrid in sandy soil in the northern region of Minas Gerais, indicated a maximum Ca accumulation of 19.23 g plant⁻¹ (96.15 kg ha⁻¹ of Ca).

It was observed in this work that the maximum accumulations of Ca between the treatments that received phosphate were between 9.71 and 22.32 g plant⁻¹ (Figure 4A-4B). In the absence of phosphate (Adc₁), the maximum Ca accumulation (6.62 g plant⁻¹) (Figure 4A) corresponds between 32% and 64% of the maximum values observed for the other treatments (Figure 4A-4B) due to a smaller growth of the plant. Under these conditions, nutrient absorption is seriously impaired, particularly those that are more dependent on transpiration flow (mass flow), such as Ca.

At 69 DAE, the decreasing order of the highest Mg accumulations observed was as it follows: F1₁₆₈>F0₂₇₅>F1₂₇₅>F1₃₉₇ (Figure 5A-5B). However, the highest Mg accumulations were obtained with F1₃₉₇ (2.58 g plant⁻¹ at 52 DAE), followed by F0₂₇₅ (2.02 g plant⁻¹ at 56 DAE), F0₃₉₇ (1.67 g plant⁻¹ at 51 DAE) and F1₁₆₈ (1.64 g plant⁻¹ at 69 DAE) (Figure 5A-5B).

Maximum accumulation of 1.51 g plant⁻¹ of Mg (at 57 DAE) was observed in Adc₂, which is equivalent to that obtained with F0₁₆₈ (56 DAE) and higher than F0₇₆ (1.28 g plant⁻¹ at 56 DAE) and F1₇₆ (1.43 g plant⁻¹ at 55 DAE). This can be explained by the lower K accumulation in Adc₂ in relation to these treatments, therefore reducing the effect of competitive inhibition with Mg (Malavolta *et al.*, 1997) ^[13]. The maximum Mg accumulation in Adc₁ corresponds to only 0.75 g plant⁻¹ (Figure 6C), or between 29% and 58% of that estimated for the other treatments.

In general, the effects of treatments on the Mg uptake can be attributed mainly to plant growth, thus, reflecting the potential for nutrient accumulation. In a study with grapevines, Skinner and Matthews (1990) ^[23] suggested that Mg uptake would be less sensitive than its translocation in response to the P supply.

The S accumulations increase until the end of the cycle, and for F₀, due to the asymptotic behavior of the fertilization

response, larger differences occur between the 76 and 168 kg ha⁻¹ doses of P₂O₅ (Figure 6A).

From the equations shown in Figures 6A and 6B, it can be deduced that the F1 fertilization provided greater S accumulations for the same dose in relation to F0. For doses between 76 and 397 kg ha⁻¹ P₂O₅, total accumulations were estimated from 0.88 to 1.12 g plant⁻¹ for F1 fertilization and from 0.71 to 0.90 g plant⁻¹ for F0 fertilization.

The S accumulation in Adc₂ (0.77 g plant⁻¹ at 69 DAE) (Figure 6C) is equivalent to the 96 kg ha⁻¹ P₂O₅ dose at F0 fertilization (Figure 6A), whereas, for Adc₁, a model was not adjusted because of the non-significance of the F test for the unfolding as a function of time.

The highest S accumulations in F1 fertilization in relation to F0 must be related to the greater P availability in the former. Kumar and Singh (1980)^[11] identified positive P interaction

over S uptake in a work using soybean crop. Alvarez *et al.* (2007)^[2] reported the existence of several studies that point to a strong positive interaction between P and S, whose effects may be related to several metabolic processes in the plant and also as a result of the displacement of the adsorbed S in the soil due to P, which presents higher bonding energy. The general order of the maximum nutrient accumulation in F0 and F1 fertilizations was: K> N> Ca> P> Mg> S, except for F0₂₇₅ treatment, where Mg accumulation is slightly higher than that of P. While in the additional treatments a higher Ca proportion was found by modifying the order of nutrient accumulation in Adc₂ for K> Ca> N> Mg ≈ P> S. In work with cv. Magnum in sandy soil, Pereira *et al.* (2018)^[19] found a lower P proportion in relation to Mg when applying phosphate only on pre-planting (137 kg ha⁻¹): K> N> Ca> Mg> P> S.

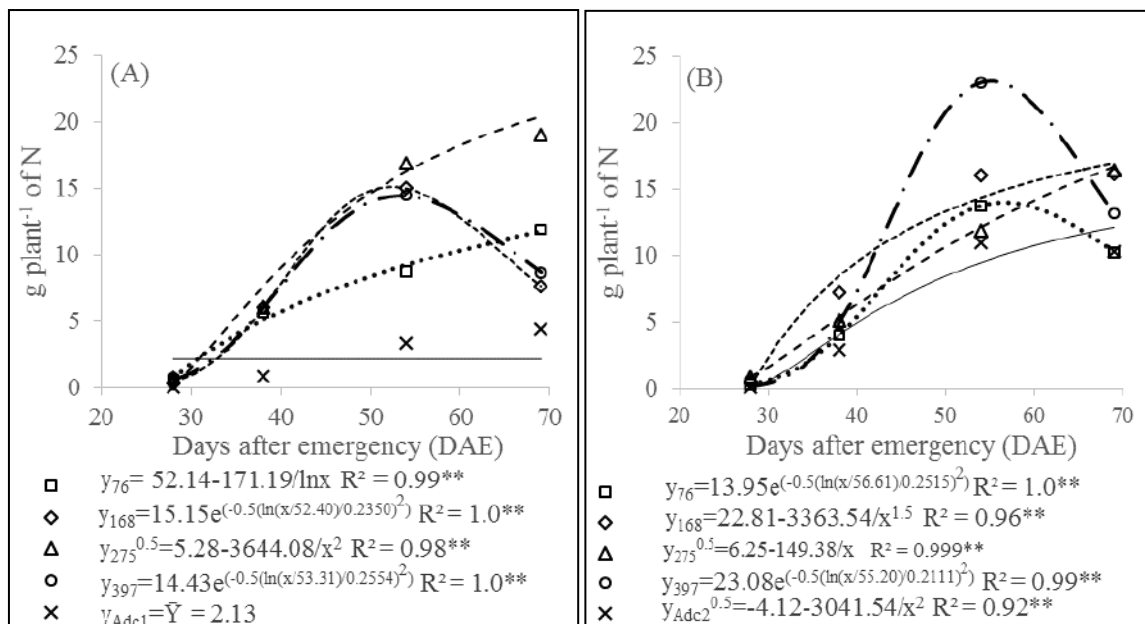


Fig 1: Total N accumulation in the aerial part of cv. Magnum watermelon plant in calcareous soil: without (A) and with P in the fertirrigation (B).

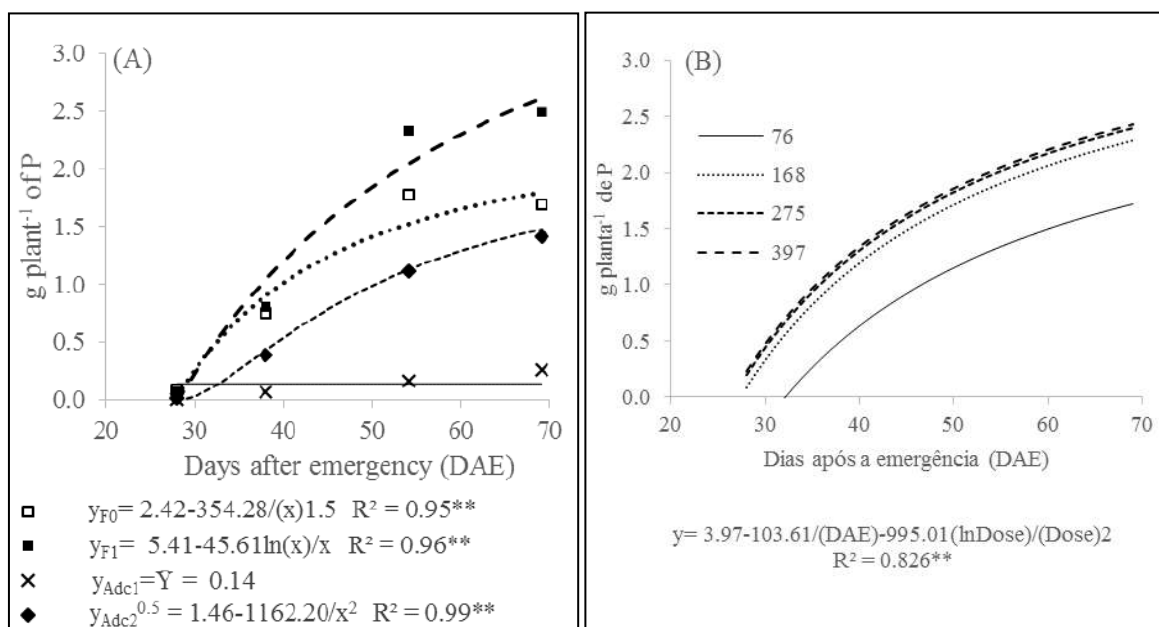


Fig 2: Total P accumulation in the aerial part of cv. Magnum watermelon in calcareous soil: without P in fertirrigation (A) and with P in the fertirrigation (B).

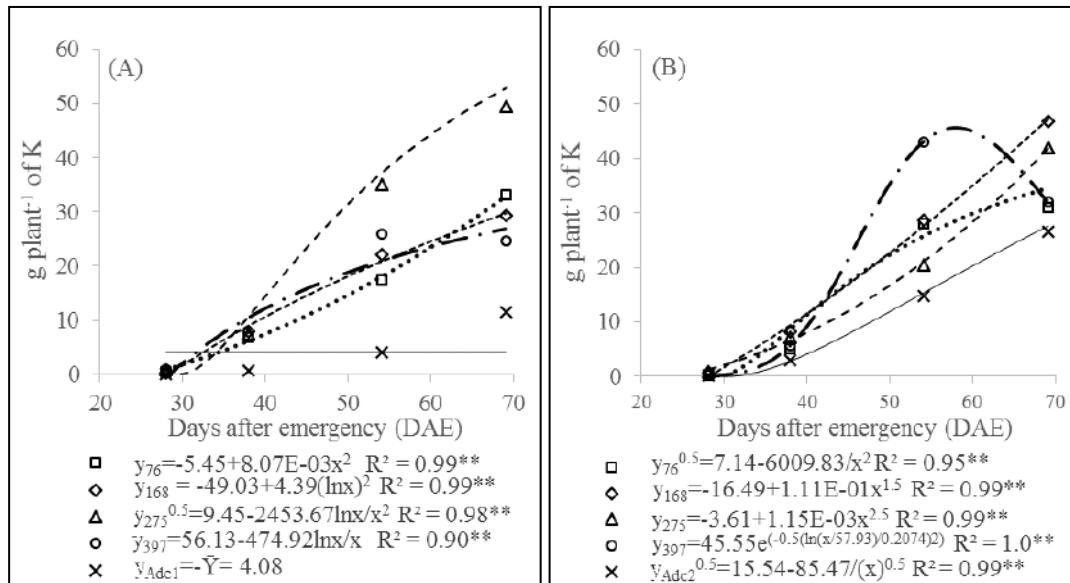


Fig 3: Total K accumulation in the aerial part of cv. Magnum watermelon in calcareous soil: without P in the fertirrigation (A) and with P in the fertirrigation (B).

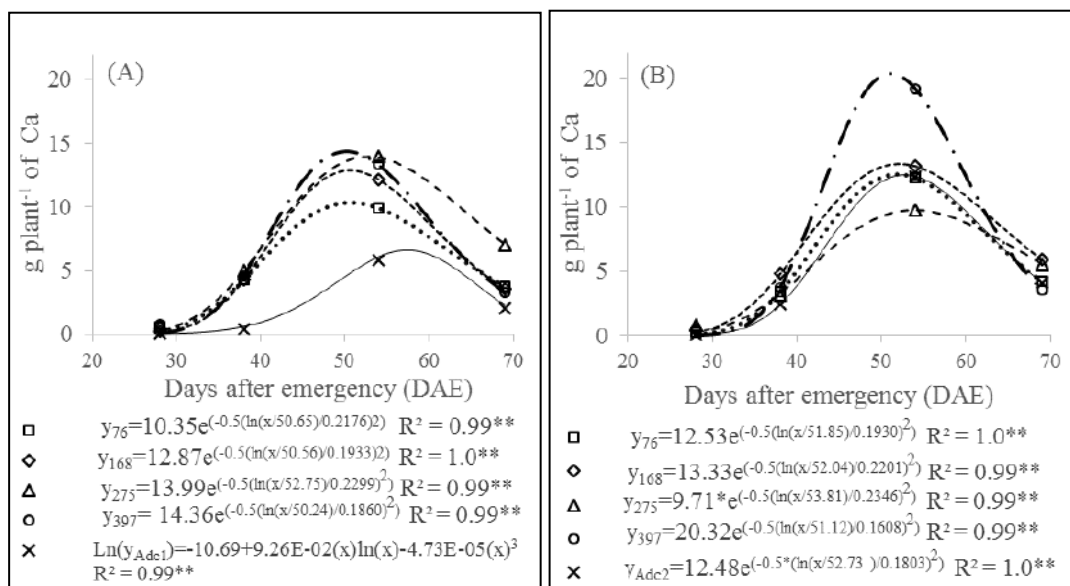


Fig 4: Total Ca accumulations in the aerial part of cv. Magnum watermelon in calcareous soil: without P in the fertirrigation (A) and with P in the fertirrigation (B).

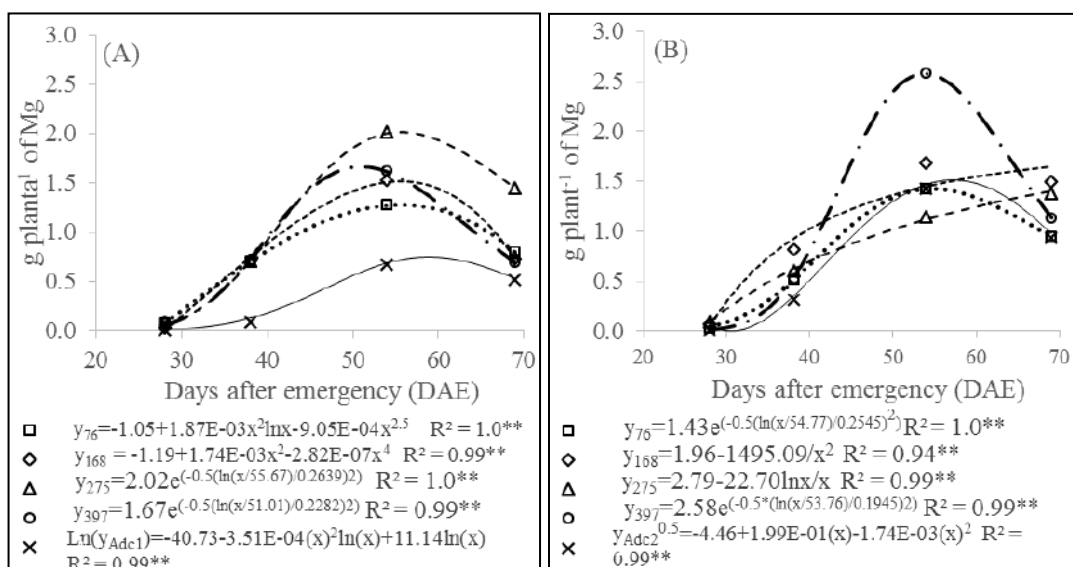


Fig 5: Total Mg accumulation in the aerial part of cv. Magnum watermelon in calcareous soil: without P in the fertirrigation (A) and with P in the fertirrigation (B).

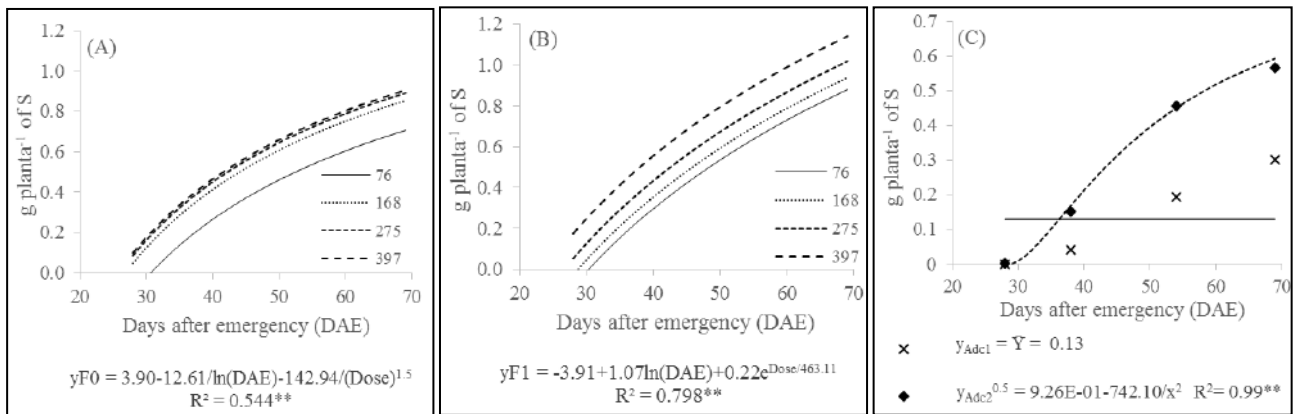


Fig 6: Total S accumulations in the aerial part of cv. Magnum watermelon in calcareous soil.: A, B, C correspond to the estimation of accumulation by the profiles of the response surfaces for total S in F0 and F1 fertilizations and in the models adjusted for the additional treatments, respectively. Upanema-RN, UFERSA, 2013.

4. Conclusions

The greatest total accumulations of P and S occur with F1 fertilization (pre-planting + fertirrigation), but in general, the characteristics evaluated in the study were dependent on the interaction between rates and manners of fertilization, where F0₂₇₅ and F1₁₆₈ stood out for the accumulation of other nutrients.

5. References

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