

钙对蓝莓幼苗生长及生理特性的影响

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摘要:【目的】探讨钙对蓝莓幼苗生长、光合及生理特性的影响。【方法】以大果蓝金品种为材料,利用水培法,设置不同钙离子浓度梯度,测定不同梯度下蓝莓幼苗生长及生理指标。【结果】较低浓度钙($2.9 \text{ mmol} \cdot \text{L}^{-1}$ 和 $5.8 \text{ mmol} \cdot \text{L}^{-1}$)促进蓝莓叶片的气孔开放,增加胞间 CO_2 浓度,提升蒸腾速率和光合速率,有效积累叶片中钙含量,株高分别增高了5.41%和25.13%,基径分别增粗了5.13%和10.26%。而较高浓度的钙处理($11.6 \text{ mmol} \cdot \text{L}^{-1}$ 和 $14.5 \text{ mmol} \cdot \text{L}^{-1}$)则导致蓝莓基径、叶片含水率、地上生物量与总生物量显著减小,胞间 CO_2 浓度、气孔导度、蒸腾速率明显降低($p < 0.05$)。钙离子浓度为 $8.7 \text{ mmol} \cdot \text{L}^{-1}$ 和 $11.6 \text{ mmol} \cdot \text{L}^{-1}$ 时根系活力显著高于对照。超氧化物歧化酶(SOD)活性在钙离子浓度为 $8.7 \text{ mmol} \cdot \text{L}^{-1}$ 时达最高,过氧化氢酶(CAT)活性随钙离子浓度的增加而逐渐升高。【结论】适宜浓度的外源钙($5.8\sim8.7 \text{ mmol} \cdot \text{L}^{-1}$)可有效调节蓝莓幼苗叶片气孔开放,促进光合和蒸腾作用,有利于蓝莓植株的生长。此外,还对提高根系活力及SOD、CAT等抗氧化酶活性($8.7 \text{ mmol} \cdot \text{L}^{-1}$)均有积极影响。

关键词:蓝莓;钙;生长;光合作用;生理特性

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Effect of calcium application on growth and physiological property of blueberry cutting seedlings

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Abstract:【Objective】Calcium is a crucial regulator of growth and development in plants, the myriad processes dealing with the calcium ion participation are involved in nearly all aspects of plant development. The calcium supplements have increasingly been used in other fruit trees, such as apple, pear, grape, peach and so on. Blueberry, as a kind of *Vaccinium* L. in Ericaceae, vaccinioideae, is suggested to be planted only in low calcium soil because it is considered to be a calcifuge plant. However, the previous studies reported that the calcium application increased calcium-content of the leaves and improved the fruit quality in some blueberry cultivars. The mechanisms of the effects of calcium application on the growth and fruit quality in blueberry is still unclear. The study aimed to detect the effects of calcium application on the growth, photosynthesis and physiological characteristics of blueberry cutting seedlings.【Methods】The experiment was carried out during July to December, 2020 in Dalian Senmao Modern Agriculture Limited Company. The local main variety Daguo Lanjin was used as the experimental material. The cutting seedlings were uniformly cultured in Hoagland and Arnon basic nutrient solution (calcium-deficient), the pH of hydroponic nutrient solution was adjusted to 4.0-4.5 with vitriol. The experiment included six treatments of calcium application: 0 (CK), $2.9 \text{ mmol} \cdot \text{L}^{-1}$ (Ca-1), $5.8 \text{ mmol} \cdot \text{L}^{-1}$ (Ca-2), $8.7 \text{ mmol} \cdot \text{L}^{-1}$ (Ca-3), $11.6 \text{ mmol} \cdot \text{L}^{-1}$ (Ca-4), $14.5 \text{ mmol} \cdot \text{L}^{-1}$ (Ca-5). CaSO_4 was

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the test source of calcium and saturation concentration is the limiting concentration of our study. The cutting seedlings were irrigated with 500ml hydroponic nutrient solution every 7 days in the beginning of the growth, and every 4 days in the period of vigorous growth. The photosynthetic parameter [net photosynthetic rate (P_n), intercellular CO₂ concentration (C_i), stomatal conductance (G_s) and transpiration rate (T_r)] were measured on 0, 10, 20, 30 and 40 days after CaSO₄ application by the portable photosynthesis of LI-6400/XT, at 8:30-11:00 a.m. The plant height, base diameter, root volume, calcium content of the stem and calcium content of the leaf were tested on 0, 40 days after CaSO₄ application. The leave and stem samples were collected on 40 days when the experiment is accomplished, and were brought back to the laboratory for the measurements of the leave moisture content, stem moisture content, above-ground biomass, total biomass, root activity, superoxide dismutase (SOD) activity and catalase (CAT) activity. The adopted experimental design was a randomized block with five repeats of each treatments (three plants per repeat were used). The differences between means were evaluated by using Tuckey's test ($p < 0.05$), the data were all mean ± standard error, statistical analysis was performed using SPSS statistical package (SPSS Statistics for Windows, Version 26.0. SPSS Inc), prism®8 (GraphPad Software, Inc.) were used to make the figure of the test. 【Results】 Our results showed that low calcium concentrations (2.9 mmol·L⁻¹ and 5.8 mmol·L⁻¹) notably enhanced the photosynthetic properties such as transpiration rate (T_r) and stomatal conductance (G_s) ($p < 0.05$) of the cutting seedlings, the leaf net photosynthetic rate (P_n) increased by 10.47% and 11.24%, compared with the control. The root volume with calcium application of 5.8 mmol·L⁻¹ and 8.7 mmol·L⁻¹ was 3.23 and 2.84 times as much as that of the control, the plant height increased by 5.41% and 25.13%, the basal diameter increased by 5.13% and 10.26% respectively, while 5.8 mmol·L⁻¹ and 8.7 mmol·L⁻¹ calcium applications had little effect on the above-ground biomass and total biomass. However, net photosynthetic rate (P_n), transpiration rate (T_r), stomatal conductance (G_s), basal diameter, above-ground biomass and total biomass were significantly reduced with the increase of calcium concentrations ranging from 11.6 mmol·L⁻¹ to 14.5 mmol·L⁻¹. The leaf moisture content displayed the tendency of first rising and then declining with the range of calcium concentrations from 2.9 mmol·L⁻¹ to 14.5 mmol·L⁻¹, whereas calcium application had little effect on the stem moisture content and calcium content of the blueberry cutting seedlings. The root activity was significantly higher than that of the control when the plants treated with 8.7 mmol·L⁻¹ and 11.6 mmol·L⁻¹ calcium ion. However, low calcium ion (2.9 mmol·L⁻¹, 5.8 mmol·L⁻¹) and high calcium ion (14.5 mmol·L⁻¹) were unfavorable for the root activity. In addition, the calcium application had significantly enhanced the SOD, CAT activities compared with the control, the SOD activity was the strongest when the calcium ion concentration was 8.7 mmol·L⁻¹, CAT activity increased gradually with the increase of calcium concentrations. 【Conclusion】 CaSO₄ application promoted the development of blueberry seedlings with appropriate calcium ion environment (5.8–8.7 mmol·L⁻¹), effectively regulated the stomatal opening and improved the transpiration rate (T_r), net photosynthetic rate (P_n), plant height, basal diameter, root volume and leaf moisture content, it also improved the root activity and the activities of superoxide dismutase (SOD), catalase (CAT). However, the plant calcium ion concentrations (11.6–14.5 mmol·L⁻¹) significantly inhibited photosynthesis and plant growth.

Key words: Blueberry; Calcium; Growth; Photosynthesis; Physiological property

钙在植物生长发育过程中发挥重要作用,是植物生长发育的必需营养元素之一^[1],主要行使营养功能和信使功能^[2]。钙在植物中作为单一营养元素维持植物细胞壁和细胞膜结构完整性,同时通过膜内外钙离子流动维持胞内环境的稳定,增强植物的抗性^[3-4],从20世纪80年代开始,钙被作为“第二信使”的研究越来越广泛,主要参与植物激素调控过程和光调节过程^[5-6]。 Ca^{2+} 行使第二信使功能时通常与CaM结合直接或间接调控植物体内生理生化过程^[7-8]。植株缺钙使其顶芽、侧芽和根尖等分生组织发生卷曲变形,叶缘开始变黄并逐渐坏死^[9],缺钙导致多种生理病害,如辣椒、番茄脐腐病^[10-11]及苹果苦痘病^[12]。外源喷施200 mmol·L⁻¹纳米碳酸钙,增加芍药茎中的钙含量,增强茎秆强度^[13];喷施150 mg·L⁻¹ CaCl₂,可使小白菜的产量与品质显著提高^[14];向烟草叶片喷施20 mmol·L⁻¹ CaCl₂可有效提高超氧化物歧化酶(SOD)、抗坏血酸过氧化物酶(APX)、过氧化氢酶(CAT)等抗氧化酶活性^[15];培养液中添加外源钙离子可显著降低水稻中茉莉酸含量,延缓叶片衰老^[16];在黄瓜水培营养液中外源施加2 mmol·L⁻¹的CaCl₂可显著促进植株的净光合速率(P_n)、蒸腾速率(T_r)、气孔导度(G_s)、胞间CO₂浓度(C_i)、光系统II实际光化学量子效率(FPS II)的提升^[17]。但高浓度钙也会对植株生长发育有抑制作用,高钙基质(1 g·kg⁻¹ CaCl₂)栽培的玉米株高、生物量、光合速率、蒸腾速率及气孔导度显著下降^[18],高钙营养液[360 mg·L⁻¹的Ca(NO₃)₂]中甜瓜叶片叶绿素含量和类胡萝卜素含量显著降低^[19]。蓝莓(Blueberry)是杜鹃花科(Ericaceae)越橘属(*Vaccinium*)植物,被认为是嫌钙植物^[20-21],但也有研究显示土壤施加硫酸钙可增加叶片钙含量,提高果实硬度^[22-23];叶面喷施0.5%的钙制剂显著增加蓝莓果实中K、Ca、Mg含量,此外对果实中的多酚和维生素C含量具有显著影响^[24]。然而,施加不同浓度的外源钙是否通过影响蓝莓生理指标进而影响蓝莓生长状况,这一关键科学问题一直未得到较好的回答。因此,笔者利用水培技术,以3月生的大果蓝金幼苗为试验材料,设计不同钙离子供给水平,研究其对蓝莓幼苗生长发育的影响,为蓝莓钙素营养供给提供理论参考。

1 材料和方法

1.1 试验设计

试验于2020年7—11月在辽宁省大连市金普新区华家街道大连森茂现代农业有限公司温室大棚进行。以3月生的大果蓝金扦插幼苗为试验材料,采用水培方式培养,营养液以Hoagland和Arnon营养液为基础(减钙),微量元素参照其他通用配方。营养液用去离子水配置,利用硫酸调整pH为4.0~4.5。试验首先将长势一致的蓝莓幼苗转移到普通水培营养液中进行2周的适应性转水培养,在此基础上再次挑选长势一致的幼苗进行下一步正式试验。试验以CaSO₄的最高溶解度为上限,设定6个Ca²⁺浓度:0(CK)、2.9(Ca-1)、5.8(Ca-2)、8.7(Ca-3)、11.6(Ca-4)及14.5(Ca-5) mmol·L⁻¹,钙由CaSO₄提供。幼苗培养前期7 d更换1次营养液,生长旺盛期4 d更换1次。试验所用水培容器为高15 cm、直径8 cm、容量900 mL的玻璃广口瓶,外部用黑色塑料袋包裹,避免透光影响根系生长。

分别于试验处理后0、40 d测定株高、基径、根体积、茎钙含量、叶片钙含量;在试验处理后0 d(7月28日)、10 d(8月8日)、20 d(8月18日)、30 d(8月28日)、40 d(9月8日)测定光合参数;在试验处理40 d时,取样,测定叶片含水率、茎含水率、地上生物量、总生物量、根系活力、SOD活性、CAT活性。

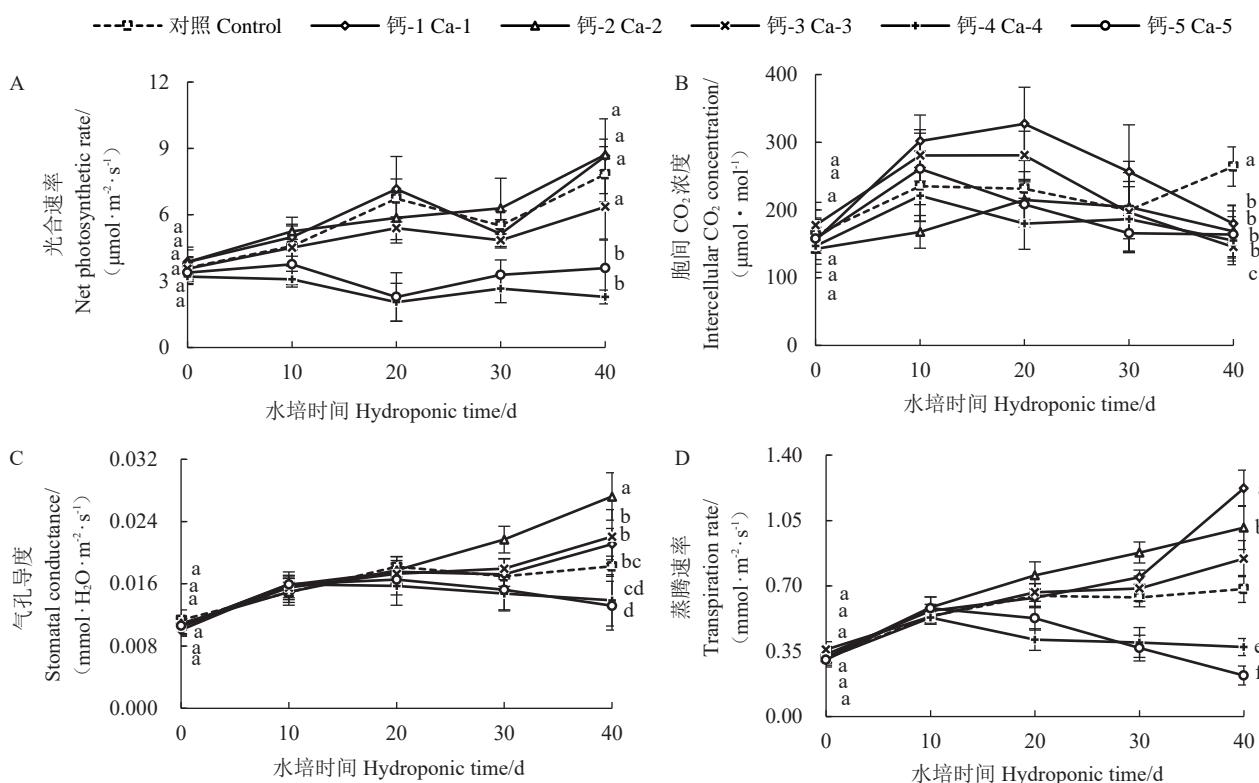
1.2 分析测定方法

分别用卷尺(精确到1 mm)、游标卡尺(精确到0.02 mm)测量株高和基径。采用LI-6400/XT光合测定系统于上午8:30—11:00测定叶片光合参数(净光合速率、蒸腾速率、胞间CO₂浓度和气孔导度),气温在18~19 °C,叶温23 °C,光强为1000 μmol·m⁻²·s⁻¹,测定部位为从顶部往下数第4片叶,每组处理随机测3株,3次重复。根体积、根系活力、SOD和CAT活性、钙含量分别采用排水法、TTC法^[25]、丙酮比色法^[26]、碘化钾法^[27]、火焰原子吸收分光光度法^[28]测定。采用Excel2016软件整理原始数据,用SPSS Statistics 26对整理好的所有数据进行单因素方差分析(ANOVA),其中 $p < 0.05$ 表示处理组之间差异显著,数据均为3次重复的平均值±标准误差,最后将分析好的数据利用Excel2016制图。

2 结果与分析

2.1 钙对蓝莓幼苗叶片气体交换参数的影响

较低浓度钙离子对蓝莓幼苗的光合速率具有一定的促进作用,随施钙时间的增加呈上升趋势,Ca-1



不同小写字母表示在 $p < 0.05$ 差异显著。下同。

Different small letters indicate significant difference at $p < 0.05$. The same below.

图 1 钙处理对蓝莓幼苗叶片净光合速率 (P_n)、胞间 CO_2 浓度 (C_i)、气孔导度 (G_s) 和蒸腾速率 (T_r) 的影响
Fig. 1 Effects of calcium application on net photosynthetic rate (P_n), the intercellular CO_2 concentration (C_i), stomatal conductance (G_s) and transpiration rate (T_r) of blueberry seedlings

处理组在 20 d 与 40 d 时的光合速率与对照相比分别增加了 6.47% 和 10.4%。而钙离子浓度高于 Ca-3 时, 40 d 时植株的光合速率与对照相比显著降低 ($p < 0.05$), 且光合速率在测定周期内变化趋势平缓(图 1-A); 胞间 CO_2 浓度随施钙时间的增加呈先上升后下降的趋势, 当钙离子浓度低于 Ca-4 时, 在施钙 20 d 时胞间 CO_2 浓度达最高之后下降, Ca-4 和 Ca-5 处理组在施钙 10 d 后胞间 CO_2 浓度就开始下降(图 1-B); 不同钙水平处理对蓝莓叶片的气孔导度和蒸腾速率影响的变化与光合速率相似, 呈上升趋势, 施钙浓度低于 Ca-4 时, 叶片的气孔导度和蒸腾速率在

40 d 时显著高于对照($p < 0.05$); 而高于 Ca-3 时, 叶片的气孔导度和蒸腾速率在 20 d 后受到抑制, 40 d 时差异显著($p < 0.05$)。因此说明, 低浓度的钙处理促进蓝莓幼苗的气孔开放和蒸腾作用, 而浓度过高时会对其产生抑制作用(图 1-C、1-D)。

2.2 钙对蓝莓幼苗生长的影响

随钙离子浓度的升高各项测定指标数值呈现先上升后下降的趋势, Ca-1、Ca-2 处理组可促进蓝莓幼苗的株高、基径、根体积等各项指标的提升, 其中根体积在 Ca-2 与 Ca-3 处理组中分别是对照的 3.23 与 2.84 倍, 差异显著($p < 0.05$)(表 1); 较高浓度的钙

表 1 钙处理对蓝莓幼苗株高、基径、根体积、地上生物量及总生物量的影响

Table 1 Effects of calcium application on plant height, basal diameter, root volume, above-ground biomass and total biomass of blueberry seedlings

因素 Factors	水平 Level/($\text{mmol} \cdot \text{L}^{-1}$)	株高 Plant height/cm	基径 Basal diameter/cm	根体积 Root volume/ cm^3	地上生物量 Above-ground biomass/g	总生物量 Total biomass/g
钙 Calcium	对照 CK(0)	9.43±1.16 ab	0.390±0.012 a	1.030±0.088 b	6.710±0.140 a	7.390±0.200 ab
	Ca-1(2.9)	9.93±0.44 ab	0.410±0.018 a	1.100±0.100 b	6.730±0.280 a	7.380±0.300 ab
	Ca-2(5.8)	11.80±0.87 a	0.430±0.021 a	3.330±0.560 a	6.680±0.120 a	7.290±0.170 ab
	Ca-3(8.7)	9.10±1.07 ab	0.310±0.019 b	2.930±0.180 a	6.850±0.240 a	7.890±0.082 a
	Ca-4(11.6)	8.40±1.32 b	0.300±0.009 b	1.400±0.310 b	6.040±0.073 b	7.130±0.064 bc
	Ca-5(14.5)	7.70±0.75 b	0.270±0.026 b	1.030±0.033 b	5.930±0.095 b	6.530±0.250 c

对蓝莓幼苗的各项指标产生一定的抑制作用,其中基径、地上生物量与总生物量在钙离子浓度高于Ca-3时明显小于对照,差异显著($p < 0.05$),Ca-4、Ca-5处理对植株各项生长指标均有不同程度的抑制作用。

2.3 钙对蓝莓幼苗茎和叶片含水率的影响

植株组织中的含水率是植物生理特征的重要指标。试验中发现,叶片含水率随钙离子浓度的增加呈现先上升后下降的趋势,各处理组中茎含水率较对照无显著变化,叶片含水率在浓度为Ca-1时显著增加,Ca-5时显著降低($p < 0.05$)。说明适宜浓度的钙离子可提高叶片含水率,较高浓度的钙离子抑制叶片含水率(图2)。

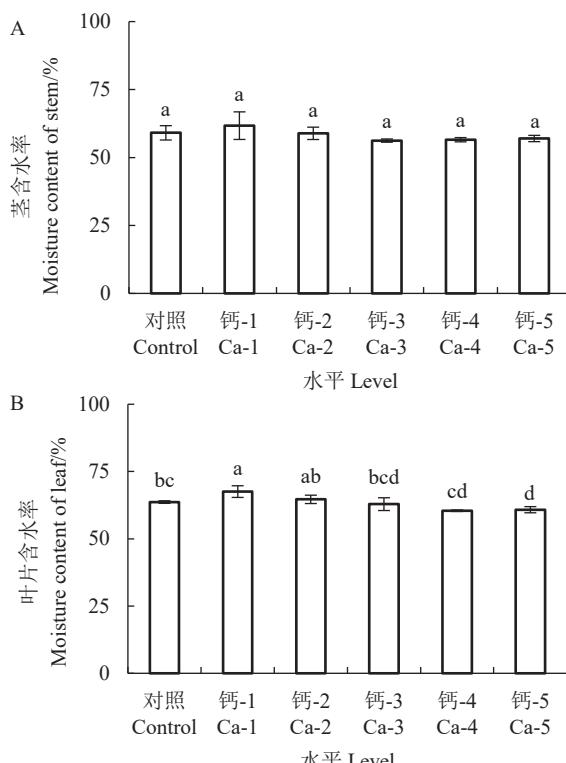


图2 钙处理对蓝莓茎、叶片含水率的影响

Fig. 2 Effects of calcium application on stem, leaf moisture content of blueberry seedlings

2.4 钙对蓝莓幼苗茎和叶片钙含量的影响

蓝莓幼苗地上各部位钙含量由高到低依次为叶片、茎。水培前各处理组中茎、叶片钙含量均无显著差异,向营养液中施加不同浓度的硫酸钙,培养40 d后各处理组中茎和叶片的钙含量虽无显著差异,但随着钙离子浓度的增加叶片钙含量呈先上升后下降的趋势,其中Ca-3钙含量达最高,Ca-5最低,即饱和浓度的硫酸钙对蓝莓叶片的钙含量积累

有抑制作用(图3)。

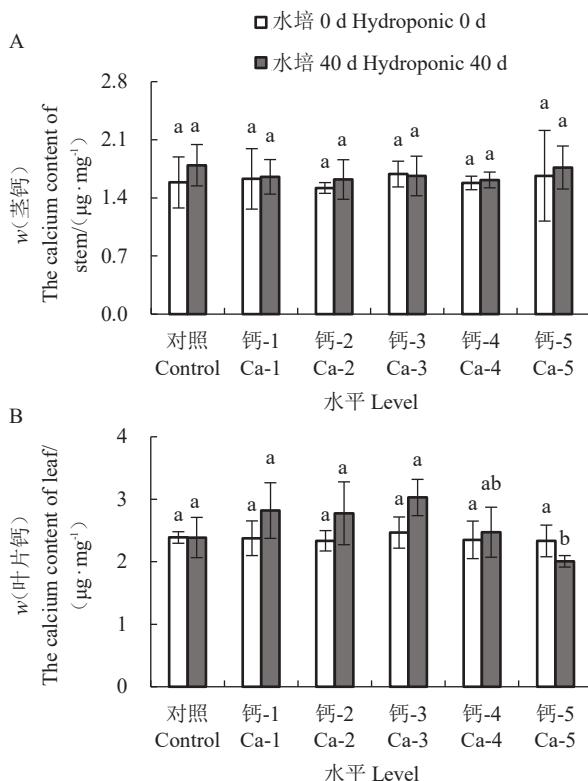


图3 钙处理对蓝莓茎、叶片钙含量的影响

Fig. 3 Effects of calcium application on stem, leaf calcium content of blueberry seedlings

2.5 钙对蓝莓根系活力的影响

四氮唑还原强度是检测植物根系活力的重要指标,钙浓度在Ca-3和Ca-4时,蓝莓幼苗的根系活力明显高于对照组;而钙离子浓度为Ca-1、Ca-2、Ca-5时,蓝莓幼苗的根系活力明显低于对照,不适宜蓝莓根系的生长(图4)。说明蓝莓根系活力的保持需要中高浓度的钙离子,而较低浓度与过高浓度钙均会降低根系活力,影响蓝莓根系的生长。

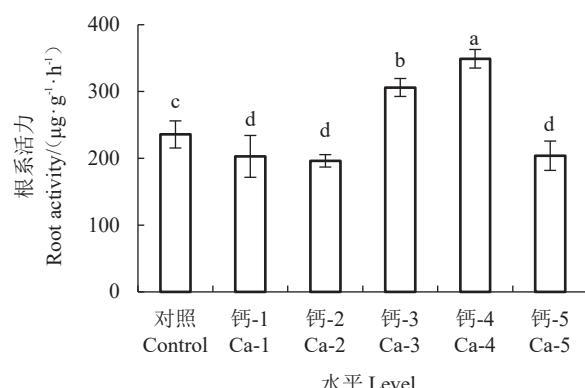


图4 钙处理对蓝莓幼苗根系活力的影响

Fig. 4 Effects of calcium application on root activity of blueberry seedlings

2.6 钙对蓝莓叶片 SOD、CAT 活性的影响

外源施加钙离子对叶片 SOD、CAT 活性均有积极影响。钙离子浓度为 Ca-2 时 SOD 活性达最大, 差异显著($p < 0.5$), 随钙离子浓度增加其活性呈现先上升后下降的趋势, 当钙离子浓度范围在 Ca-1~Ca-3 时, SOD 活性明显高于对照处理(图 5-A); 钙离子处理的蓝莓幼苗叶片 CAT 活性较对照差异显著($p < 0.05$), 随钙离子浓度的增加 CAT 活性呈上升趋势, 钙离子浓度为 Ca-5 时, 其活性最高(图 5-B)。

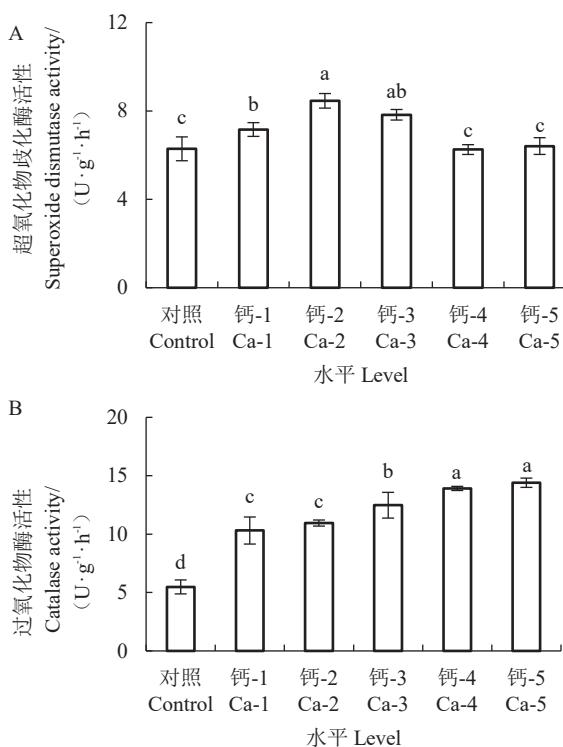


图 5 钙处理对蓝莓叶片 SOD 和 CAT 活性的影响

Fig. 5 Effects of calcium application on the activity of SOD and CAT leaf

3 讨 论

钙是植物生长所需的必要营养元素, 参与植株体内的光合作用、细胞骨架维持、离子吸收与激素调控等系列生理生化过程, 有效促进植株生长^[6,29]。越橘属植物(包括蓝莓)长期以来被认为是嫌钙植物, 叶片中钙浓度低且适宜在少钙土壤中生长^[30-31]。但在本研究中发现 Ca-1 和 Ca-2 处理使得蓝莓幼苗株高分别增高了 5.41% 与 25.13%, 基径分别增粗了 5.13% 与 10.26%, 叶片钙含量分别增加了 17.99% 与 16.32%。在水培 20 d 时, Ca-1 和 Ca-2 处理的胞间 CO₂ 浓度达到最高, 20 d 后气孔导度与蒸腾作用显

著高于对照, 且在 40 d 时的光合速率分别比对照组增加了 6.47% 和 10.4%。这与钙在玉米、拟南芥和大蒜等的研究中结果相似, 钙信号不仅参与叶片气孔的开闭^[32-34], 同时对胞间 CO₂ 浓度、蒸腾作用均有促进作用^[35-36]。但过高浓度的钙会抑制植物的生长代谢, 导致信号紊乱, 严重时引起细胞死亡^[37-38]。本试验中较高浓度钙(Ca-4, Ca-5)处理抑制了蓝莓幼苗株高和基径的生长, 叶片含水率降低了 11.76%, 植株的光合速率、胞间 CO₂ 浓度、气孔导度、蒸腾速率、生物量与总生物量显著低于对照组。说明适宜浓度钙可促进蓝莓幼苗的生长, 蓝莓并非是嫌钙植物, 可能是由于适宜蓝莓生长的土壤环境为透水性、通气性及排水性良好, 有机质含量丰富, 尤为关键的是 pH 需控制在 4.5~5.5 之间, 若 pH 过高, 造成缺铁失绿, 生长不良, 产量降低甚至植株死亡^[39-40]。但通常未改良土壤中钙大多为碳酸钙和氯化钙, pH 近于中性或呈微碱性^[41-42], 不符合蓝莓栽培条件。而本试验利用营养液水培法, 通过施加不同浓度硫酸钙, 控制 pH, 发现外源施钙促进气孔的开放, 一方面, 提高了叶片的蒸腾速率, 有利于水分和无机营养的吸收; 另一方面, 增加了胞间 CO₂ 浓度, 促进叶片的光合碳同化, 为蓝莓植株的生长提供了更多的有机营养^[43-44]。

根系是重要的营养器官, 其形态大小与活力程度影响植物对水分与矿质营养的吸收与利用, 外源钙的施加促进根的生长^[45]。Ca²⁺通过参与调控 IAA 合成过程诱导黄瓜不定根的形成^[46]; 拟南芥根毛长度与培养基中 Ca²⁺浓度相关, 适宜浓度的 Ca²⁺(0.3~3 mmol·L⁻¹) 促进根毛生长^[47]; Ca²⁺通过促进蚕豆根部 GA 和 IAA 的积累进而影响根部生长^[48]。蓝莓属于浅根性须根系植物, 钙离子对于蓝莓根系的影响结果各不相同。Austin 等^[49]研究发现蓝莓土壤中钙离子浓度较高时, 降低其根系活力, 但在 Hanson 等^[22]的研究中发现外源土壤施加硫酸钙对蓝莓植株无害。本试验中 Ca-2 和 Ca-3 处理组的根体积分别是对照组的 3.23 和 2.84 倍, Ca-3 和 Ca-4 处理组中, 蓝莓幼苗的根系活力明显高于对照组。推测是因为外源施加钙并没有改变适宜蓝莓幼苗生长的水培养养液环境, 适宜浓度钙离子(Ca-3)促进蓝莓根体积生长和根系活力的提高。钙对植物细胞膜的结构和功能具有稳定作用^[50], Zhou 等^[51], Afifyanti 等^[52]和王博伟等^[53]研究发现, 外源施钙可有效提高番茄、苜蓿、生菜叶片 SOD、POD 与 CAT 等抗氧化酶的活性,

提高其抗逆适应能力。本研究中发现,Ca-1~Ca-3范围浓度的钙离子可以促进蓝莓SOD活性的升高,CAT活性随钙离子浓度的升高而升高。说明适宜浓度的钙处理,可以促进蓝莓SOD、CAT等抗氧化酶的活性,可能提高其对逆境胁迫的适应能力。

4 结 论

适宜的外源钙可以有效调节蓝莓幼苗叶片气孔开放,促进光合和蒸腾作用,有利于蓝莓植株的叶片、根系等器官的生长,此外,还可提高SOD、CAT等抗氧化酶活性;但较高浓度的钙处理会明显抑制蓝莓的生长发育。笔者在本研究中初步探讨了钙对蓝莓生长发育的影响,以期为蓝莓栽培中外源施钙应用提供一定理论参考。

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