Short chain chito-oligosaccharides promote arbuscular mycorrhizal colonization in Medicago truncatula

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Abstract: During the establishment of arbuscular mycorrhizal (AM) symbiosis, the fungus and the host plant exchange chemical signals that are crucial to reciprocal recognition. Short-chain chitin oligomers (CO) released by AM fungi are known to trigger symbiotic signaling in all host plant species tested. Here we applied exogenous CO, derived from crustacean exoskeleton, to pot-grown Medicago truncatula inoculated with the AM fungus Funneliformis mosseae and investigated root colonization, plant gene regulation and biomass production. CO treatment strongly promoted AM colonization with significant increases in arbuscule development, biomass production and photosynthetic surface compared to untreated mycorrhizal plants. Gene expression analyses indicated that CO treatment anticipated the expression of MtBCP and MtPT4 plant symbiotic markers, during the first two weeks post inoculation. Altogether, our results provide evidence that plant treatment with symbiotic fungal elicitors, anticipated and enhanced AM development, encouraging the use of CO to promote AM establishment in sustainable agricultural practices.

Cover Letter

Torino, 30.7.2019

Dear Editor,

We are submitting the manuscript entitled «*Short chain chito-oligosaccharides promote arbuscular mycorrhizal colonization in Medicago truncatula*» to be considered for publication in Carbohydrate Polymers**.**

Over 80% of plants, including most crops, live in a mutualistic symbiotic association, called arbuscular mycorrhiza (AM), with an ancient group of soil fungi that provide their hosts with soil water and mineral nutrients in change for sugars and lipids. For this key role in plant nutrition, the introduction of AM as biofertilizers in **sustainable agricultural practices** has been proposed for decades as a promising natural solution to the increasing worldwide demand of food safety. Nevertheless, field applications tests have delivered contrasting results, and experimental data indicate a partial recalcitrance of cultivated species toward AM interaction, likely due to decades of excessive use of fertilizers and to farming practices that have been unaware of the plant underground interactions.

We now know that plant-fungus recognition is an essential step in AM establishment and is based on an exchange of chemical signals: root-exudates alerts the AM fungus of host proximity, while AM fungi release **short-chain chito-oligosaccharides** (CO) eliciting the plant symbiotic responses, such as gene regulation, starch reallocation and cell rearrangement for fungal accommodation.

In the light of this role for CO as AM fungal signals, we decided to **test whether their exogenous application could have a positive impact on symbiosis establishment**. This would on the one hand demonstrate unambiguously the biological role of CO as symbiosis elicitors and, on the other hand, open the way to their use as promoters of AM colonization in crops for sustainable, food safety-oriented agricultural applications.
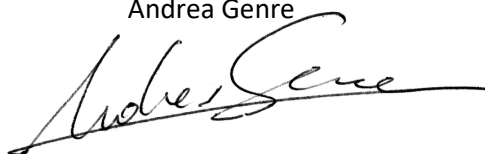
To this aim, we treated the model legume *Medicago truncatula* with a mixture of short-chain CO (ranging between 2 and 5 N-acetyl-glucosamine residues) and observed how their application influenced plant mycorrhizal status and development.

Indeed, our analyses of root colonization intensity, plant gene expression and biomass development consistently showed that **CO treatments strongly enhance and anticipate AM colonization**. This represents a **breakthrough** on the way to the large-scale introduction of AM in agricultural practices, by positively acting on the mycorrhizal aptitude of the host plants.

For these reasons we consider that the findings in this article are of primary interest in the field of carbohydrate polymer applications to plant-microbe interactions, and appropriate for timely publication in your Journal.

Yours sincerely

Andrea Genre

- Arbuscular mycorrhizal (AM) symbiosis supports life of most crop plants.
- The fungus is recognized as a symbiont via water-soluble short-chain chitin oligomers (CO)
- We tested whether exogenous CO application to *Medicago truncatula* could impact on the symbiosys.
- CO treatment strongly promoted AM colonization with significant increases in arbuscule development, biomass production and photosynthetic surface compared to untreated mycorrhizal plants.
- Plant treatment with symbiotic fungal elicitors anticipated and enhanced AM development, encouraging the use of CO to promote AM establishment in sustainable agricultural practices

1   **Short chain chito-oligosaccharides promote arbuscular mycorrhizal colonization in *Medicago***

2   ***truncatula***

3

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11

12   **Abstract**

13   During the establishment of arbuscular mycorrhizal (AM) symbiosis, the fungus and the host plant

14   exchange chemical signals that are crucial to reciprocal recognition. Short-chain chitin oligomers

15   (CO) released by AM fungi are known to trigger symbiotic signaling in all host plant species tested.

16   Here we applied exogenous CO, derived from crustacean exoskeleton, to pot-grown *Medicago*

17   *truncatula* inoculated with the AM fungus *Funneliformis mosseae* and investigated root colonization,

18   plant gene regulation and biomass production. CO treatment strongly promoted AM colonization with

19   significant increases in arbuscule development, biomass production and photosynthetic surface

20   compared to untreated mycorrhizal plants. Gene expression analyses indicated that CO treatment

21   anticipated the expression of *MtBCP* and *MtPT4* plant symbiotic markers, during the first two weeks

22   post inoculation. Altogether, our results provide evidence that plant treatment with symbiotic fungal

23   elicitors, anticipated and enhanced AM development, encouraging the use of CO to promote AM

24   establishment in sustainable agricultural practices.

25

1. Introduction

The life of most plants, including the majority of crops, is supported by a mutualistic root symbiosis with Glomeromycotina, an ancient group of soil fungi (Spatafora et al., 2016) that provide their hosts with a preferential access to soil water and mineral nutrients, while taking advantage of plant-photosynthesized sugars and lipids (Smith & Read, 2008; Wewer et al. 2014; Keymer et al., 2017).

Plant-fungus recognition is essential for AM establishment and is based on an exchange of chemical signals (Bonfante & Requena, 2011; Zipfel & Oldroyd, 2017). Root-exuded strigolactone alerts the AM fungus of host proximity (Akiyama et al., 2005); its perception activates spore germination, hyphal metabolism and branching (Besserer et al., 2006). In turn, AM fungi release diffusible molecules eliciting the plant symbiotic response (Maillet et al, 2011; Genre et al., 2013). These include local and systemic changes in gene expression (Kosuta et al., 2003), root-directed starch reallocation (Gutjahr et al., 2009), and the activation of a signal transduction pathway, partially shared with other symbiotic interactions (Oldroyd, 2013; Delaux et al., 2015; Barker et al., 2017; Genre and Russo, 2016). This pathway includes the triggering of repeated oscillations (spiking) in nuclear calcium concentration, making the detection of nuclear Ca^{2+} spiking an acknowledged benchmark for the induction of symbiotic responses in AM hosts.

Indeed, Ca^{2+} spiking is observed in root epidermal nuclei upon adhesion of AM fungal hyphopodia (Chabaud et al., 2011), application of AM fungal exudate (Navazio et al, 2007; Chabaud et al., 2011; Genre et al., 2013) or purified molecules isolated from fungal exudate, such as short-chain chito-oligosaccharides (CO) and lipo-chito-oligosaccharides (LCO) (Maillet et al., 2011; Genre et al., 2013). In particular, fungal release of tetrameric and pentameric CO is boosted upon strigolactone treatment (Genre et al., 2013). Furthermore, their activity has been demonstrated in legumes (Genre et al., 2013), carrot (Genre et al., 2008), rice (Sun et al., 2015; Carotenuto et al., 2017), *Discaria trinervis* and *Casuarina glauca* (Chabaud et al., in press).

In the light of this universal role for CO in eliciting early plant responses to AM fungi (Genre &

51 Russo, 2016), we decided to test whether their exogenous application could have a positive impact

52 on symbiosis establishment. This would on the one hand demonstrate the biological role of CO as

53 symbiosis elicitors and, on the other hand, open the way to their use as promoters of AM colonization

54 in crops for sustainable, food safety-oriented agricultural applications (Berruti et al., 2015).

55   To this aim, we treated the model legume *Medicago truncatula* with a mixture of short CO (ranging

56 below 6 N-acetyl-glucosamine residues), obtained from crustacean manufacturing industry, and

57 observed how their application influenced plant development and mycorrhizal status.

58

59 **2. Materials and Methods**

60 *2.1. Plant growth and mycorrhization.*

61   Seeds of *M. truncatula* (cv. Jemalong, genotype A17) were scarified for 3 min in H_2SO_4, rinsed

62 several times in distilled water, sterilized for 2 min in diluted commercial bleach (1:5), rinsed 4 times

63 in sterile distilled water and pre-germinated for 48 hours on agar plates (0.6% w/v) at 23° C in the

64 dark. Seedlings were grown for ten days in a photoperiod of 16h day (23°C) and 8h night (21°C)

65 before transferring them to 10x10x12cm pots containing quartz sand.

66   The AM fungus *Funneliformis mosseae* (strain BEG 12) was inoculated in mycorrhizal pots by

67 replacing 10% of the pot substrate with a commercial inoculum (MycAgroLab, Bretenière, France)

68 and mixing. All plants were watered once a week with half-strength Long-Ashton nutrient solution

69 containing 3,2 μM KH_2PO_4 as P source (Hewitt, 1966).

70

71 *2.2. Chito-oligosaccharides*

72   We used a commercial mixture of short chain CO purified from crustacean manufacturing side-

73 products (Zhengzhou Sigma Chemical Co., Ltd., Henan, China). *HPLC-MS/MS* analysis of the CO

74 mixture (File S1) confirmed the presence of deacetylated, mono-acetylated and di-acetylated CO

75 molecules composed of 2 to 5 N-acetyl-glucosamine residues (Fig. S1).

76

2.3. Chito-oligosaccharide solutions

1g/L and 1mg/L CO solutions in sterile distilled water were tested for the triggering of Ca^{2+} spiking in the root epidermis. Only the 1g/L solution was chosen for pot treatments, to counteract molecule dilution/absorption and compensate for the presence of biologically inactive CO2 and CO3 in the mix, alongside the AM signals CO4 and CO5 (Fig. S1). CO application to pot-grown plants was done either by irrigating the substrate with the CO solution, or by spraying the solution on plants and soil surface.

2.4. Analysis of CO-induced Ca^{2+} signals

Confocal microscopy was performed on 2cm-long lateral roots of *Agrobacterium rhizogenes*-generated root organ cultures (ROC) of *M. truncatula* expressing NUP-YC2.1, a nuclear localized Ca^{2+}-sensing yellow cameleon construct (Chabaud et al., 2011). Detection and plotting of relative changes in Ca^{2+} concentration in epidermal atrichoblasts was done according to Genre et al. (2013), as described in File S1. At least 100 atrichoblasts from seven independent root samples were analyzed for each condition and the t-Student test with a probability level of $P < 0.05$ was used for statistical validation.

2.5. Plant treatment with CO

In order to evaluate CO effect on plant development and mycorrhization, four experimental conditions were set up: control (Ctr), lacking both CO treatment and AM inoculation; CO-treated control (Ctr+CO); mycorrhizal (Myc), where AM inoculum was added in the absence of CO treatments; and CO-treated mycorrhizal (Myc+CO), where plants were both inoculated and exposed to the CO solution.

For CO application by irrigation, residual water was carefully removed from the pot saucers before

101 watering each Ctr+CO and Myc+CO pot with 30 mL of CO solution. Distilled water was used for Ctr

102 and Myc pots. Treatments were applied weekly, and plants were sampled 48 days post inoculation

103 (dpi), after six CO treatments (Fig. S2), for morphometric analyses and mycorrhizal intensity

104 determination.

105 For spray treatments, 5 mL of CO solution containing 0.005% Tween20 as a surfactant, was sprayed

106 weekly over Ctr+CO and Myc+CO pots from a 10 cm distance, using plastic shields to confine spray

107 diffusion. Ctr and Myc pots were sprayed with 5 mL of 0.005% Tween20 in sterile distilled water.

108 Plants treated with CO spray were divided in two groups (Fig. S2): one was grown for 28 dpi (with

109 two CO treatments) and the other for 48 dpi (with six CO treatments) before measuring their

110 morphological and physiological parameters and quantifying AM colonization.

111

112 2.6. *Analyses of plant development after CO treatment*

113 Irrigation-treated plants were harvested at 48 dpi; root and shoot fresh weight was immediately

114 measured. To determine the leaf area, green tissues were carefully isolated and placed on an Epson

115 Perfection V800 Photo scanner, to obtain black and white scanned images of the leaf surface at 600

116 dpi resolution. The resulting PNG images were then analyzed using Fiji/ImageJ

117 (http://imagej.nih.gov/;http://fiji.sc/Fiji) to calculate total leaf surface in mm^2. Following image

118 acquisition, shoot dry weight was measured after exsiccation at 60°C for 24h.

119 Spray-treated plant growth was monitored (Fig. S3) by photography at 7 dpi (prior to CO treatment),

120 28 dpi (two treatments) and 48 dpi (six treatments). Five Ctr, five Ctr+CO, ten Myc and ten Myc+CO

121 plants were harvested at 28 dpi and 48 dpi, to measure their fresh and dry weight and photosynthetic

122 surface, as described above.

123

124 *2.7. Nitrogen balance index*

125 The physiological status of spray-treated plants was estimated before sampling using a Dualex 4

126 (FORCE-A, Orsay, France) to determine flavonoid and chlorophyll content by measuring leaf UV

127 absorbance and fluorescence, respectively, in 10 randomly chosen leaves from 5 Ctr and 5 Ctr+CO

128 plants, and 20 leaves from 10 Myc and 10 Myc+CO plants. Measures were done on both adaxial and

129 abaxial leaf surface and the two values were averaged (Cartelat et al., 2005). The

130 chlorophyll/flavonoid ratio, or Nitrogen Balance Index (NBI), is considered an indicator of the plant

131 nitrogen nutritional status (Cerovic et al., 2012).

132

133 *2.8. Quantification of mycorrhizal colonization*

134 Following fresh weight measure, each root system was stained with 0.1% cotton blue in lactic acid

135 and quantitative parameters of mycorrhizal intensity were calculated as described by Trouvelot et al.

136 (1986), using the online tool MYCOCALC (http://www2.dijon.inra.fr/mychintec/Mycocalc-

137 prg/download.html).

138

139 *2.9. Gene expression analysis following CO treatment*

140 To investigate the ability of CO treatment to stimulate host receptiveness for AM fungi, a time-

141 course experiment was set up where plants received two rounds of spray treatment before AM

142 inoculation, 21 and 23 days post transplant (dpt). The inoculum was added at 25 dpt, and plants were

143 sampled at 7, 10, 14, 21 and 28 dpi. Five leaf replicates per treatment were sampled to analyse

144 phosphorus (P) content and five root replicates from Myc and Myc+CO were sampled and frozen in

145 liquid nitrogen for gene expression analyses.

146

147 *2.10. RNA isolation and quantitative RT-PCR analysis*

148 Total RNA was isolated from roots, using the RNeasy™ Plant Mini kit (Qiagen, Hilden, Germany).

149 Samples were disrupted with a TissueLyser (Qiagen Retsch GmbH, Hannover, Germany) and treated

150 following manufacturer instructions. RNA quality and quantity were evaluated using a Nanodrop and

151 agarose gel electrophoresis.

152   To remove genomic DNA contamination, RNA samples were digested with TURBO™ DNase

153 (Ambion) according to manufacturer instructions and checked for DNA traces by conventional PCR

154 using *MtTEF* primers, before cDNA synthesis and qRT-PCR analyses (File S1). The sequences of all

155 genes studied are listed in Table S1.

156

157 *2.11. Assessment of arbuscule morphology*

158   Root colonized segments from Myc and Myc+CO plants were embedded in agarose (8%) and cut

159 into 200 μm-thick sections using a Vibratome (Oxford Vibratome®). Sections were collected on

160 microscope slides and stained with fluorescein isothiocyanate-conjugated wheat germ agglutinin

161 (WGA-FITC; Sigma-Aldrich, Milan, Italy), as described in Volpe et al. (2016). Twenty root segments

162 from five biological replicates were analyzed for each treatment.

163

164 *2.12. Phosphorus quantification*

165   Phosphorus quantification was done in 5 Ctr, 5 Ctr+CO, 10 Myc and 10 Myc+CO plants, by

166 collecting the third leaf from the tip of the main stem. Collected leaves were lyophilized, weighed

167 and digested in 1 mL of 6M HNO_3 for one hour at 95°C. The digestion product was diluted in 6 mL

168 of sterile distilled water. A control solution, without any sample, was treated with the same procedure.

169 All solutions were analysed by Inductively Coupled Plasma Atomic Emission Spectrometry (ICP-

170 AES) to determine P content.

171

172 *2.13. GenBank accession numbers*

173   AY116211 (*MtPT4*); XM_003592405.2 (*MtBCP*); XM_013595882.1 (*MtTEF*); XM_013606824

174 (*MtPR10*); XM_003604044.2 (*MtChit1*); XM_003618769.2 (*MtChitIII.1*); XM_003597718.2

175 (*MtChitIII.2*); AY238969.2 (*MtChitIII.3*); AY238970.1 (*MtChitIII.4*)

7

176

3. Results

3.1. Crustacean-derived CO triggered Ca^{2+} spiking in M. truncatula atrichoblasts

179   In order to assess the commercial CO effectiveness as symbiotic elicitors, we recorded nuclear Ca^{2+}

180   signals in atrichoblasts of NUP-YC2.1 *M. truncatula* ROCs treated with 1g/L and 1mg/L CO solutions

181   (Fig. 1). Both solutions triggered comparable spiking patterns, with frequent, well-defined peaks, in

182   analogy with those induced by analogous concentrations of purified CO4, CO5 or raw AM spore

183   exudate (Chabaud et al., 2011; Genre et al., 2013). As expected, no spiking was recorded upon control

184   treatment with sterile distilled water. The percentage of responding atrichoblasts and average peak

185   numbers confirmed the lack of statistically significant differences in the cell response to either CO

186   concentration (Fig. 1B).

187   In conclusion, our bioassay confirmed that both 1g/L and 1mg/L solutions of the commercial CO

188   mix was able to mimic AM fungal signals in triggering Ca^{2+}-mediated symbiotic signaling.
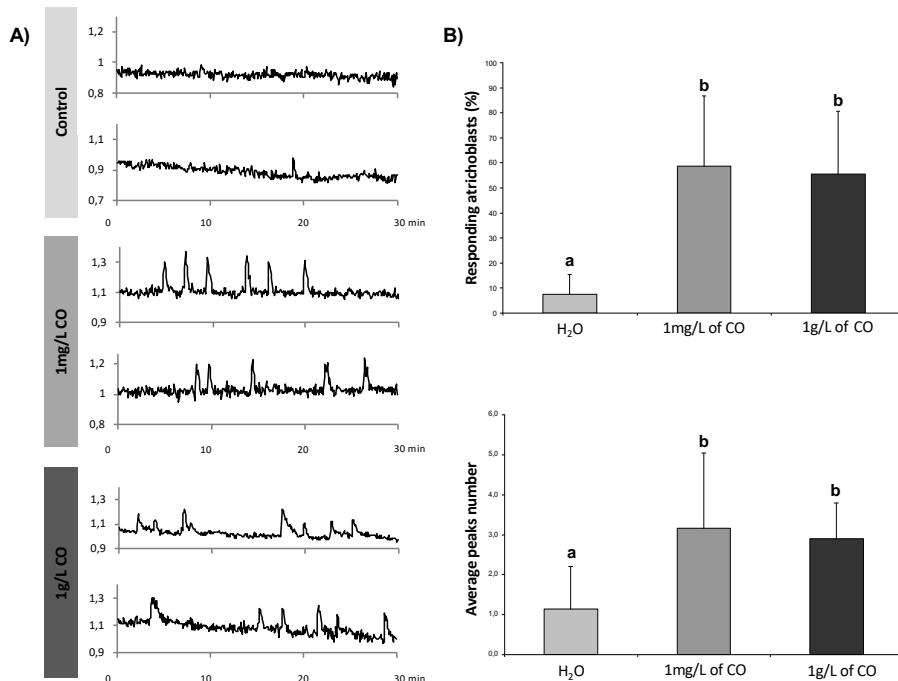
Fig. 1. Nuclear Ca^{2+} spiking in response to crustacean-derived CO. A) Representative plots of nuclear Ca^{2+} variations over 30 minutes in atrichoblast nuclei of *M. truncatula* root organ culters (ROCs) treated with water (control), 1mg/L and 1g/L CO. Both CO concentrations induced a series of Ca^{2+} peaks (spiking), characteristic of AM fungal recognition. B) Statistical analyses of the spiking response (t-Student test with $P < 0.05$) did not highlight significant differences between CO treatments, in terms of either the percentage of responding atrichoblasts (top panel) or average peak number in responding cells (bottom). Bars represent standard errors. At least 100 atrichoblasts from seven independent root samples were analyzed for each experimental condition.

189

190   *3.2. CO application by irrigation boosted AM colonization*

191   In a first attempt to investigate the effect of exogenous CO supply, we grew inoculated and non-

192   inoculated *M. truncatula* in pots, with or without a weekly supplement of CO in the watering solution,

193   and analyzed their biomass, photosynthetic surface and mycorrhizal colonization at 48 dpi. While

194   root development was comparable in Ctr, Myc and Myc+CO plants, statistically significant

195   differences were observed in the shoot fresh and dry biomass between Ctr and both mycorrhizal

196   conditions, Myc and Myc+CO (Fig. S4). The same trend was confirmed for photosynthetic surface,

197   with similar top values for Myc and Myc+CO plants and a significantly smaller area for Ctr and

198   Ctr+CO plants. Remarkably, Ctr+CO plants showed a significant reduction in biomass and

199   photosynthetic surface, when compared to all other treatments, suggesting that CO perception in the

200   absence of fungal inoculation could induce a stress condition inhibiting plant growth.

201   AM colonization was then assessed in Myc and Myc+CO plants through optical microscopy.

202   Isolated groups of arbuscules were distributed along the root axis of Myc plants, spaced by

203   uncolonized cortical cells (Fig. S5). By contrast, Myc+CO plants displayed a more extensive and

204   homogeneous distribution of arbuscules, with a continuous mass of arbusculated cells, which also

205   involved multiple layers of the root cortex (Fig. S5). This was confirmed by our quantitative analysis,

206   with significantly higher frequency (F), intensity of mycorrhization (M) and arbuscule abundance (a,

207   A) in Myc+CO compared to Myc plants (Fig. S5).

208   In conclusion, beside a partial growth inhibition in uninoculated plants, exogenous CO application

209   by irrigation strongly improved the mycorrhizal status of AM inoculated plants at 48 dpi.

210

211   *3.3. CO spray treatment enhanced AM colonization*

212   These encouraging results, coupled with the possible induction of a stress response in Ctr+CO,

213   prompted us to test an alternative treatment. We chose spray application as an amenable method of

214   CO supply in view of possible field treatments. For these experiments, we sampled our plants at 28

9

215 and 48 dpi to obtain a clearer picture of symbiosis development.

216   Our first focus was to check whether the spray treatment could be as efficient in promoting AM

217 development as the irrigation method. Indeed, F, M and A values were all significantly higher in

218 Myc+CO compared to Myc plants at 28 dpi (Fig. 2). Also in this case, CO treatment caused arbuscule

219 proliferation across multiple cortical layers, compared to the fewer arbuscules, with spotted

220 distribution, found in Myc plants. At 48 dpi, all mycorrhization parameters surged in both conditions,

221 reducing the difference between Myc and Myc+CO plants, which anyway remained significant (Fig.

222 2).

223   In short, the quantification of AM colonization based on microscopic observations confirmed a

224 positive effect on mycorrhizal intensity also for CO spray treatment.
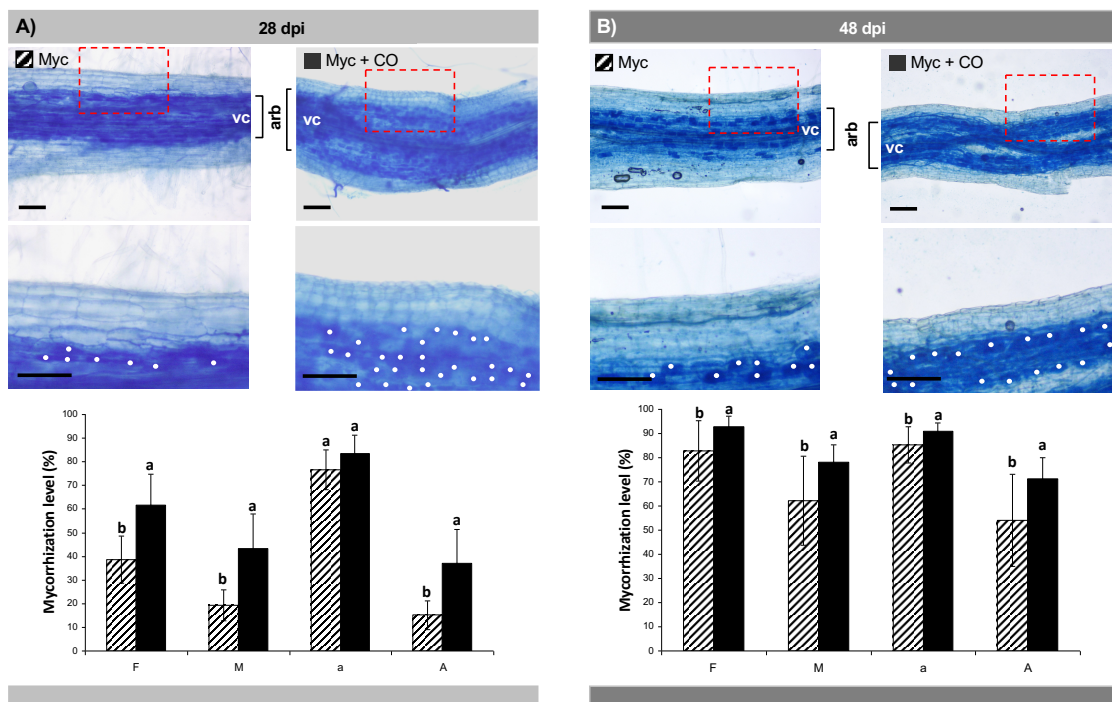
Fig. 2. Effect of spray CO treatment on mycorrhizal plants at 28 dpi (A) and 48 dpi (B). Images of root sections of mycorrhizal plants untreated (Myc) and treated with CO (Myc+CO). A strong proliferation of arbuscules (white dots) was evident in Myc+CO plants, with the colonization of multiple cortical layers at both time points, as evident in the magnified areas (red square) in the lower panels. This was confirmed by the quantitative analysis of root colonization, shown in the bottom graphs: mycorrhization frequency (F), mycorrhization intensity (M), arbuscule abundance within colonized root fragments (a) and arbuscule abundance in the whole root system (A) all reported significantly higher values in Myc+CO compared to Myc plants. The effect of CO on root colonization was anyway more evident at 28 dpi. Mean values and SDs of ten biological replicates for each treatment are shown. Bars = 100 µm.

225

226

227

228 *3.4. CO spray treatment and AM inoculation synergistically increase plant growth*

229   Photographic monitoring of shoot growth showed that prior to any CO treatment - at 7 dpi - all

230 plants had a comparable development (Fig. S3). By contrast, significant differences became evident

231 at 28 dpi (two CO treatments) and even more pronounced at 48 dpi (six CO treatments): at both time

232 points, Myc and Myc+CO plants were larger than Ctr plants. Interestingly, CO application by spray

233 to uninoculated plants (Ctr+CO) did not affect their growth at any time point (Fig. S3), in contrast

234 with the results of CO treatment by irrigation.

235   Such macroscopical observations were partially confirmed by biometric measurements. At 28 dpi

236 (Fig. 3A), the photosynthetic surface was significantly more extensive in mycorrhizal treatments

237 compared to both Ctr and Ctr+CO, with the highest value for Myc+CO plants, albeit not significantly

238 larger than in Myc plants. An analogous trend was recorded for shoot dry biomass, with the highest

239 values for the Myc and Myc + CO plants, even if differences were not statistically significant.
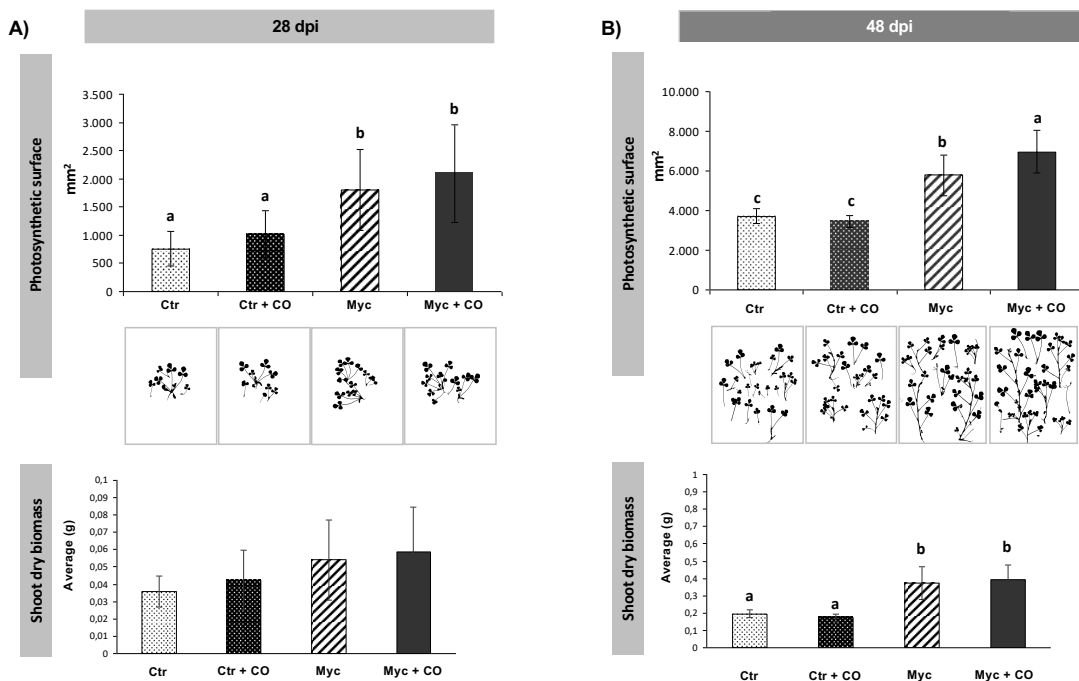

Fig. 3. Aboveground development in response to spray CO treatment. A) Average photosynthetic surface (including representative scans of the plant epigeral parts) and shoot dry biomass at 28 dpi: the photosynthetic surface was significantly more developed in Myc and Myc+CO plants compared to Ctr and Ctr+CO; a similar – albeit statistically non significant - trend was observed for shoot dry weight. B) The photosynthetic surface was significantly more development in Myc+CO plants at 48 dpi, when both mycorrhizal plants showed a significant increase of shoot dry biomass. Mean values and SDs are presented for five and ten biological replicates from control and mycorrhizal plants, respectively.

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11

241   At 48 dpi (Fig. 3B), both mycorrhizal conditions (Myc and Myc+CO) showed a significantly larger

242   photosynthetic surface than both Ctr and Ctr+CO, but in addition, Myc+CO significantly outclassed

243   Myc plants at this later time point. An analogous significant increase was observed in shoot dry

244   biomass between mycorrhizal and uninoculated plants, albeit Myc and Myc+CO had very similar

245   values.

246   In conclusion, two CO spray treatments were sufficient to induce a significant increase in AM

247   colonization at 28 dpi, while a CO-dependent increase in plant development was evident at 48 dpi,

248   after six CO treatments.
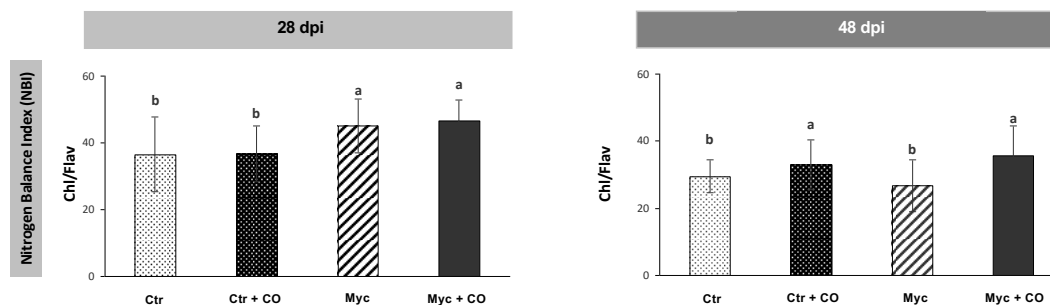
Fig. 4. Leaf nitrogen (N) status at 28 and 48 dpi. The chlorophyll/flavonoid ratio (Chl/Flav), or Nitrogen Balance Index (NBI), is presented here as an indicator of leaf N content. NBI value was significantly higher in both mycorrhizal lines (Myc and Myc+CO) than controls (Ctr and Ctr+CO) at 28 dpi. By contrast, NBI was significantly higher in all CO treated plants at 48 dpi. Mean values and SDs are presented for five and ten biological replicates from control and mycorrhizal plants, respectively.

249

250

251   *3.5. Effect of CO spray on plant nitrogen status*

252   We measured the nitrogen balance index (NBI) to indirectly monitor plant nitrogen status and

253   overall stress. At 28 dpi, the NBI was significantly higher for both mycorrhizal conditions, indicating

254   a better metabolic performance compared to Ctr and Ctr+CO plants (Fig. 4). No significant difference

255   was observed between the latter, confirming that CO spray alone did not cause significant stress in

256   uninoculated plants. By contrast, at 48 dpi, the NBI was significantly higher in CO-treated plants

257   (Ctr+CO and Myc+CO) compared to Ctr and Myc.

258   Altogether, NBI analysis indicated a comparable increase in leaf nitrogen content for both

12

259 mycorrhizal conditions at 28 dpi, and a further increase in Myc+CO at 48 dpi, in line with CO effects

260 on plant development.

261

262 *3.6. Plant AM marker genes are regulated in response to CO treatment*

263   Since CO supply had effectively enhanced root colonization at 28 and 48 dpi, we decided to

264 investigate in more detail the effect of CO with a time-course experiment that included earlier time

265 points. In addition, CO ability to stimulate host plant receptiveness for AM fungi was assessed by

266 spraying plants twice with the CO, before fungal inoculation (Fig. S2). We consequently analyzed

267 gene expression for two established AM markers: *MtBCP* (*Blue Copper-binding Protein*), expressed

268 during arbuscule development, and *MtPT4* (*Phosphate Transporter 4*), encoding an AM-specific P

269 transporter localized on the periarbuscular membrane of active, mature arbuscules (Pimprikar &

270 Gutjahr, 2018).

271   *MtBCP* expression was more strongly induced in Myc+CO plants at 7, 10 and 14 dpi, compared to

272 a later activation in Myc plants (Fig. 5), with a significant difference at 7 dpi. A drastic

273 downregulation was recorded at 21 and 28 dpi, in line with the expected reduction of *MtBCP*

274 expression in terminal stages of root colonization. Similarly, *MtPT4* was also activated earlier in

275 Myc+CO plants, but its expression surprisingly decreased at 21 dpi, apparently anticipating the

276 analogous decrease observed in Myc plants at 28 dpi.
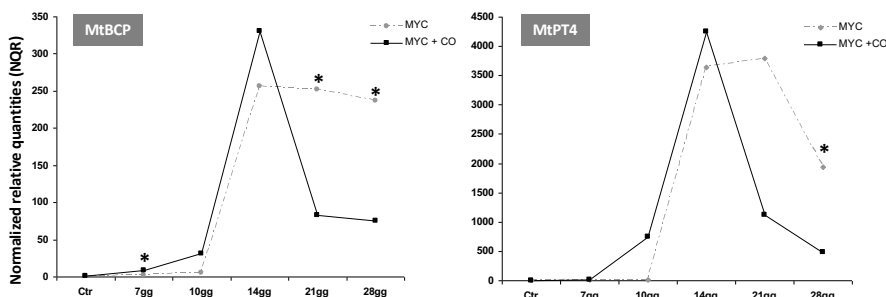
Fig. 5. Time-course analysis of *MtBCP* and *MtPT4* expression. Asterisks indicate the significant early induction of *MtBCP* at 7 dpi and the significant downregulation of both genes at 21 and 28 dpi in CO-treated plants (Myc+CO). Average values from at least 5 biological replicates are presented.

277

278   To better investigate this unexpected difference in AM marker gene expression, we anayzed the

279 regulation of three AM-induced chitinases (Salzer et al, 2000; Kremer et al., 2013; Malolepszy et al,

280 2018) in fully developed mycorrhizal roots, at 28 dpi: *MtChitIII.2*, *MtChitIII.3* and *MtChitIII.4*. In

281 our hands, only *MtChitIII.2* and *MtChitIII.4* were upregulated in inoculated plants compared to

282 controls (Fig. 6), but both chithinases were less expressed in Myc+CO compared to Myc plants. In

283 conclusion, two additional markers of arbuscule maturity displayed the same downregulation as

284 *MtPT4* in Myc+CO plants.
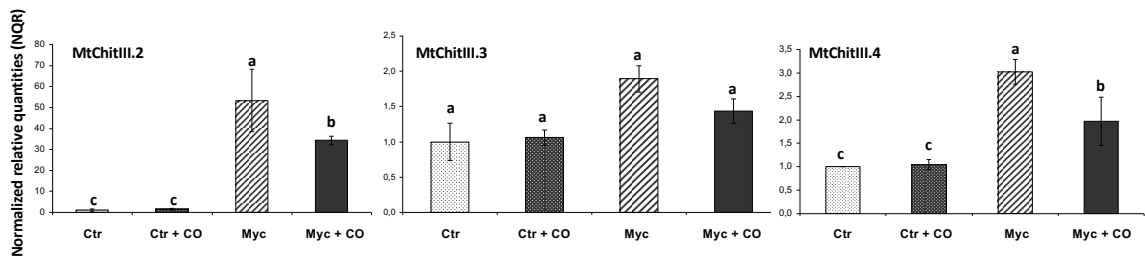
Fig. 6. Molecular analysis on roots at 28 dpi after two CO treatments. Two of the three AM-induced chitinases are strong induced in the mycorrhizal plants, but the transcripts levels for *MtChitIII.2* and *MtChitIII.4* are significant reduced in the Myc+CO compared to the Myc untreated plants, in line with the expression profile of *MtPT4*. Mean values and SDs in three biological replicates for each treatment are shown.

285

286   On this basis, we wondered whether this pattern of gene regulation could be indicative of anticipated

287 arbuscule degeneration in CO-treated mycorrhizal plants - an aspect that could not be highlighted by

288 our quantitative analyses of mycorrhizal colonization. We therefore compared arbuscule morphology

289 in Myc and Myc+CO plants (Fig. S6). No obvious difference was observed in arbuscule shape and

290 general organisation: the expected arbuscule morphology, with a large trunk and a mass of fine

291 branches occupying most of the host cell volume, was consistently recorded for both conditions,

292 suggesting that the vast majority of arbuscules was fully developed, independent of CO application.

293   Altogether, CO treatment on the one hand accelerated root colonization by the AM fungus,

294 providing a convincing demonstration of CO role as elicitors of the host plant symbiotic response.

295 Exogenous CO application led to an anticipated expression of early AM markers such as *MtBCP* and

296 the development of a higher number of arbuscules that appeared functional, based on their

297 morphological features. On the other hand, CO treatment reduced the late expression of AM marker

298 genes that are normally expressed in active arbusculated cells (Harrison et al., 2002; Elfstrand et al.,

14

299   2005; Pumplin & Harrison, 2009). This unexpected result prompted us to compare the nutritional

300   status of Myc and Myc+CO plants.

301

302   *3.7. Leaf phosphorus content was not affected by CO treatment*

303   We compared P content in leaf samples from both mycorrhizal conditions. Surprisingly, no

304   significant difference was detected for any time point: plants reached their highest P content at 14 dpi

305   under both conditions (Fig. 7), with a progressive decrease in the following time points.

306   In conclusion, the increase in arbuscule abundance in Myc+CO plants was not reflected by an

307   improvement in plant P uptake under our experimental conditions.
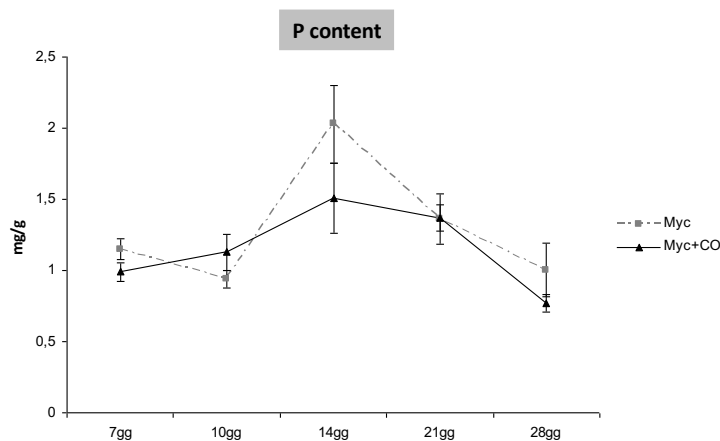
Fig. 7. Phosphorous (P) content level in untreated (Myc) and treated (Myc+ CO) mycorrhizal plants. Both lines reached the highest leaf P content at 14 dpi but no significant difference was detected in any time point. Mean values for five biological replicates are shown. Bars represent standard errors.

308

309   *3.8. CO spray does not upregulate plant defense markers*

310   To have a more comprehensive view of CO effects on the plant, we also checked whether CO

311   perception could activate plant defense responses, by analyzing the expression of the pathogenesis

312   related gene *MtPR10* and two pathogenesis-related chitinases, *MtChitI* and *MtChitIII.1* (Salzer et al.,

313   2000; Lipka et al., 2005). None of these defense markers was induced by CO treatment (Fig. 8), in

314   line with literature data about the scarce activity of short chain CO as defense elicitors (Shibuya &

315   Minami, 2001). Moreover, the significant reduction of *MtChitI* expression in Ctr+CO, Myc and

316   Myc+CO plants compared to Ctr, suggests a possible partial inhibition of defense responses in the

317   presence of either fungal or externally supplied (or both) signals.
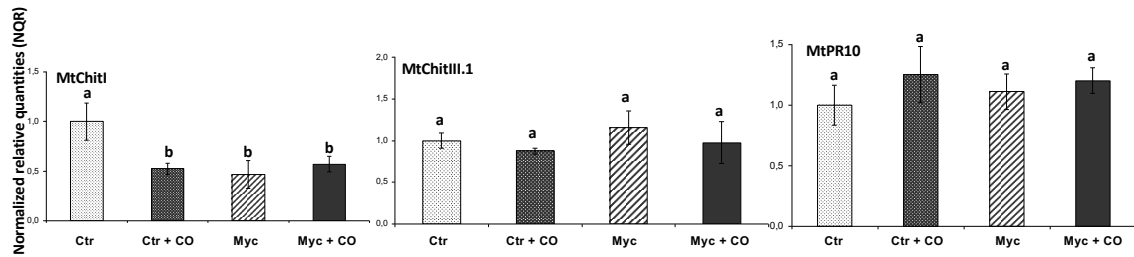
Fig. 8. Expression of pathogenesis-related genes in roots at 28 dpi after two CO treatments. The three analysed pathogenesis-induced genes, *MtChitI*, *MtChitIII.1* and *MtPR10*, were not upregulated upon treatment with crustacean-derived CO, suggesting that they were not perceived as a pathogenesis signal. Mean values and SDs in three biological replicates for each treatment are shown.

318

4. Discussion

320   In spite of several consistent reports of CO role as elicitors of early AM signaling and a few

321   investigations of CO-regulated gene expression (Weidmann et al, 2004; Giovannetti et al, 2015; Sun

322   et al., 2015), surprisingly few studies have investigated the effect of CO on symbiosis development

323   (Ramírez et al., 2010). In the light of our results, CO now appear particularly promising for large

324   scale agricultural applications for several reasons: among currently characterized AM fungal signals,

325   CO can be obtained at a fraction of the cost of the more complex LCO (lipo-chito-oligosaccharides,

326   similar to rhizobial Nod factors), from waste products of crustacean fishing industry. As natural and

327   easily bio-degradable products, CO are considered environmentally friendly and the use of chitin-

328   derived molecules has a relatively long history in agriculture, for their stimulation of plant growth

329   and defense (Benhamou et al., 1998; Sharp, 2003; Zhang et al., 2016; Winkler et al., 2017).

330

331   *4.1. Exogenous CO application promotes AM development*

332   The most striking result of this study was the observation that CO treatment induced a remarkable

333   increase in arbuscule abundance. This evidence represents a crucial advance in support of the role of

334   CO as positive elicitors of symbiotic responses in the host plant, demonstrating their effect - beyond

335   early signaling and gene expression – on the development of arbuscules, the functional core of AM

336   symbiosis.

16

337   In addition, our results highlighted that spraying the plants with CO prior to AM fungal inoculation

338 anticipated the upregulation of *MtBCP* and *MtPT4* at 7 and 10 dpi, suggesting the ability of CO to

339 accelerate AM establishment: a very promising aspect in view of their application under agricultural

340 conditions. Interestingly, CO did not anticipate symbiosis senescence, as suggested by the arbuscule

341 regular morphology in both treated and untreated samples. The fact that the longer time-span of AM

342 colonization in CO-treated plants did not correlate with a higher P content could appear surprising.

343 Nevertheless, unchanged P contents are also reported in studies on hypermycorrhizal mutants of *M.*

344 *truncatula* (Truong et al., 2014) and *Pisum sativum* (Jones et al., 2015), where arbuscule proliferation

345 appears analogous to what we have observed upon CO treatment. A few hypotheses can be made to

346 explain this, each of which deserves further investigation. Firstly, P uptake might not be the main

347 function of the morphologically active arbuscules that we have observed at 28 dpi. Under our standard

348 growth conditions, fewer arbuscules - as in untreated mycorrhizal plants - might be sufficient to grant

349 *M. truncatula* an optimal P nutrition. The plant might instead take advantage of such an abundant

350 number of arbuscules for other functions, such as water uptake. Challenging CO-treated and untreated

351 plants with different levels of drought will shed light on this possibility, but our observation of a

352 larger biomass in CO-treated mycorrhizal plants at 48 dpi - when the root system had extended to the

353 whole pot volume – hints at a better performance of this experimental line under incipient stress

354 conditions.

355   Secondly, prolonged CO elicitation could stimulate sustained fungal accommodation, while

356 arbuscule functioning and senescence could be regulated by different - and yet unexplored - signaling

357 processes. Lastly, one cannot exclude that such an abundant root colonization in a mature stage of the

358 symbiosis is favoring the AM fungus absorption of plant-derived organic compounds, with CO

359 elicitation possibly weakening the plant control mechanisms on over-colonization. Under this respect,

360 it will be extremely important to investigate C flux toward the fungus, as well as extraradical

361 mycelium development or spore production.

362

4.2. Plant responses to CO treatments

The use of different chitin-derived molecules in agricultural practices or laboratory conditions is not new. Benefits to the plant have been described in terms of better growth and resistance to biotic or abiotic stresses. Nevertheless, a clear picture of their multiple effects on plants is far from being understood and most agricultural applications rely on the heuristic observation of a correlation between chitin-based treatments and beneficial effects. In this frame, our results provide a possible explanation to such positive effects on crops: if such mixes of compounds include short-chain CO, then part of the benefits to the plant may derive from its improved symbiotic status, which in turn is well known to support plant development, defense and stress resistance.

In the field of plant-microbe interactions, the length of CO molecules is acknowledged as a major determinant of plant responses, with the following - likely simplistic - scenario: chito-biose and -triose are reported to produce little or no stimulation in plant cells; -tetraose and -pentaose have been characterized as symbiotic signals in all AM host plants (Genre et al., 2013; Giovannetti et al, 2015; Sun et al., 2015); whereas longer CO progressively shift toward defense elicitation (Liu et al., 2012; Hayafune et al., 2014), with chito-octamers being the best characterized chitin-based pathogen-associated molecular pattern (Zhang et al., 2002; Cao et al., 2017). In this frame, CO have been proposed as primers of plant defenses in agriculture, even if limited attention has been put on the oligomer composition of the mixture. The picture is even less clear about the effects of CO on plant development. A stimulating effect of chito-pentaose was demonstrated on Arabidopsis growth (Khan et al., 2011). Similarly, Winkler and colleagues (2017) demonstrated that a mixture of short-chain CO, increased Arabidopsis and poplar shoot explant development, and chito-tetraose treatment of Arabidopsis seedlings upregulated development, cell organization, biogenesis and transport-related genes. By contrast, no stimulating effect on plant growth was observed for CO by Chatelain et al. (2014), who rather reported a decrease in *Phaseolus vulgaris* biomass. Zhang et al (2016) indicated

18

387 CO length as a critical determinant of their effect also on plant development, with chito-hexaose and

388 -heptaose inducing the strongest elicitation of biomass development in wheat.

389   This complex scenario is difficult to compare with our experimental system, based on CO treatment

390 of mycorrhizal plants. Nevertheless, a few parallels may be drawn. The reduction of uninoculated

391 plant development observed in CO irrigated plants, appears in line with the inhibition of plant growth

392 described by Zhang et al. (2016), since we were using a mix of CO that did not include the growth-

393 promoting hexa- and heptamers. By contrast, a synergistic positive effect of CO and inoculation on

394 plant development was observed near the end of our experimental time frame. In this case, fungal

395 presence is introducing an additional level of complexity to the experimental system, complicating

396 the interpretation of direct and combined effects of CO, symbiotic status and stress conditions

397 possibly induced by the space limitation of the pot culture.

398

399 **5. Conclusion**

400   The introduction of AM as biofertilizers in sustainable agricultural practices has been proposed for

401 decades as a promising natural solution to the increasing worldwide demand of food safety. Results

402 have anyway been contrasting, with two major obstacles emerging. First, even if AM fungi have low

403 host specificity, studies quickly highlighted that not all fungal inocula are suitable for all agricultural

404 applications: local climate, soil characteristics, as well as the species (if not the variety) of the selected

405 crop generate a matrix of variables that only customized inocula could address, with the time- and

406 resource-consuming selection of performing fungal strains for each condition. Second, a partial

407 recalcitrance has been observed in cultivated species toward AM interaction, likely due to decades of

408 excessive use of fertilizers (Duhamel & Vandenkoornhuyse, 2013) and to farming practices that have

409 always been attentive to the aerial part of the plant but unaware of its underground interplay.

410   Our study showed that CO treatments strongly enhance AM colonization, anticipating the

411 upregulation of AM marker genes, but not arbuscule senescence. Even if several aspects of CO effects

412 on plants remain to be explored, our demonstration of their positive impact on AM establishment

413 represents a breakthrough on the way to the large scale introduction of AM in agricultural practices,

414 by positively acting on the mycorrhizal aptitude of the host plants.

415

420

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569

570

571 **Figure legends.**

572 **Fig. 1.** Nuclear Ca^{2+} spiking in response to crustacean-derived CO. A) Representative plots of nuclear

573 Ca^{2+} variations over 30 minutes in atrichoblast nuclei of *M. truncatula* root organ culters (ROCs)

574 treated with water (control), 1mg/L and 1g/L CO. Both CO concentrations induced a series of Ca^{2+}

575 peaks (spiking), characteristic of AM fungal recognition. B) Statistical analyses of the spiking

576 response (t-Student test with $P < 0.05$) did not highlight significant differences between CO

577 treatments, in terms of either the percentage of responding atrichoblasts (top panel) or average peak

578 number in responding cells (bottom). Bars represent standard errors. At least 100 atrichoblasts from

579 seven independent root samples were analyzed for each experimental condition.

580

581 **Fig. 2.** Effect of spray CO treatment on mycorrhizal plants at 28 dpi (A) and 48 dpi (B). Images of

582 root sections of mycorrhizal plants untreated (Myc) and treated with CO (Myc+CO). A strong

583 proliferation of arbuscules (white dots) was evident in Myc+CO plants, with the colonization of

584 multiple cortical layers at both time points, as evident in the magnified areas (red square) in the lower

585 panels. This was confirmed by the quantitative analysis of root colonization, shown in the bottom

586 graphs: mycorrhization frequency (F), mycorrhization intensity (M), arbuscule abundance within

587 colonized root fragments (a) and arbuscule abundance in the whole root system (A) all reported

588 significantly higher values in Myc+CO compared to Myc plants. The effect of CO on root

589 colonization was anyway more evident at 28 dpi. Mean values and SDs of ten biological replicates

590 for each treatment are shown. Bars = 100 μm.

591

592 **Fig. 3.** Aboveground development in response to spray CO treatment. A) Average photosynthetic

593 surface (including representative scans of the plant epigeral parts) and shoot dry biomass at 28 dpi:

594 the photosynthetic surface was significantly more developed in Myc and Myc+CO plants compared

595 to Ctr and Ctr+CO; a similar – albeit statistically non significant - trend was observed for shoot dry

596 weight. B) The photosynthetic surface was significantly more development in Myc+CO plants at 48

597 dpi, when both mycorrhizal plants showed a significant increase of shoot dry biomass. Mean values

598 and SDs are presented for five and ten biological replicates from control and mycorrhizal plants,

599 respectively.

600

601 **Fig. 4.** Leaf nitrogen (N) status at 28 and 48 dpi. The chlorophyll/flavonoid ratio (Chl/Flav), or

602 Nitrogen Balance Index (NBI), is presented here as an indicator of leaf N content. NBI value was

603 significantly higher in both mycorrhizal lines (Myc and Myc+CO) than controls (Ctr and Ctr+CO) at

604 28 dpi. By contrast, NBI was significantly higher in all CO treated plants at 48 dpi. Mean values and

605 SDs are presented for five and ten biological replicates from control and mycorrhizal plants,

606 respectively.

607

608 **Fig. 5.** Time-course analysis of *MtBCP* and *MtPT4* expression. Asterisks indicate the significant early

609 induction of *MtBCP* at 7 dpi and the significant downregulation of both genes at 21 and 28 dpi in

610 CO-treated plants (Myc+CO). Average values from at least 5 biological replicates are presented.

611

612 **Fig. 6.** Molecular analysis on roots at 28 dpi after two CO treatments. Two of the three AM-induced

613 chitinases are strong induced in the mycorrhizal plants, but the transcripts levels for *MtChitIII.2* and

614 *MtChitIII.4* are significant reduced in the Myc+CO compared to the Myc untreated plants, in line

615 with the expression profile of *MtPT4*. Mean values and SDs in three biological replicates for each

616 treatment are shown.

617

618 **Fig. 7.** Phosphorous (P) content level in untreated (Myc) and treated (Myc+CO) mycorrhizal plants.

619 Both lines reached the highest leaf P content at 14 dpi but no significant difference was detected in

620 any time point. Mean values for five biological replicates are shown. Bars represent standard errors.

621

622 **Fig. 8.** Expression of pathogenesis-related genes in roots at 28 dpi after two CO treatments. The three

623 analysed pathogenesis-induced genes, *MtChitI*, *MtChitIII.1* and *MtPR10*, were not upregulated upon

624 treatment with crustacean-derived CO, suggesting that they were not perceived as a pathogenesis

625 signal. Mean values and SDs in three biological replicates for each treatment are shown.

626

627 **Supplementary data**

628 **Fig. S1.** Composition of crustacean-derived CO identified by HPLC-MS/MS. CO molecules were

629 detected with a chain length ranging between 2 and 5 residues and different degrees of deacetylation.

630 A) Since CO detection threshold raises with increasing chain length, only the presence (+) or absence

631 (-) of each type of molecule is presented here. NS = not searched. B) Structure of mono-deacetylated

632 (beta anomer) CO4 and theorical precursor/product ions forming in LC-MS/MS the positive mode.

633 C) LC-MS/MS chromatogram in the MRM mode (M+H+). The two peaks correspond to alpha and

634 beta anomers of the molecule.

635

636 **Fig. S2.** Schemes representing the experimental set-up for CO treatment and sampling used for this

637   work. The camera symbol indicates photographic recording of plant development; dpi, days post

638   inoculation; dpt, days post transplant.

639

640   **Fig. S3.** Monitoring of plant growth during CO spray treatment. The growth of control (Ctr), CO-

641   treated control (Ctr+CO), mycorrhizal (Myc) and CO-treated mycorrhizal plants (Myc+CO) was

642   monitored at 7, 28 and 48 dpi. AM inoculation alone increased plant growth at both 28 and 48 dpi,

643   but aerial part development at both time points was further enhanced when AM inoculation was

644   combined with CO application (Myc+CO) at 14, 21, 28, 34, 40 and 46 dpi as summarized in Fig. S2.

645

646   **Fig. S4.** Effect of CO treatment by irrigation on plant development at 48 dpi. Shoot and root biomass

647   production was monitored in control (Ctr), CO-treated control (Ctr+CO), mycorrhizal (Myc) and CO-

648   treated mycorrhizal plants (Myc+CO). A reduced development of both shoots and roots was observed

649   in Ctr+CO compared to Ctr plants. While root biomass was not affected by AM inoculation, Myc and

650   Myc+CO plants displayed an increased shoot biomass production compared to Ctr, with consistent

651   results for fresh weight, dry weight and photosynthetic surface. Mean values and SDs of ten biological

652   replicates for each treatment are shown.

653

654   **Fig. S5.** Effect of CO treatment by irrigation on mycorrhizal development at 48 dpi. Images of root

655   sections from untreated (Myc) and CO-treated mycorrhizal plants (Myc+CO). Arbuscules

656   proliferation was strongly stimulated in Myc+CO plants, with the involvement of multiple cortical

657   layers, compared to a more limited colonization, surrounding the vascular cylinder (vc), in Myc plants.

658   This observation was confirmed by the quantitative analysis of root colonization, reported in the

659   bottom graph: mycorrhization frequency (F), mycorrhization intensity (M), arbuscule abundance

660   within colonized root fragments (a) and arbuscule abundance in the whole root system (A) all reported

661   significantly higher values in Myc+CO compared to Myc plants. Mean values and SDs of ten

662   biological replicates for each treatment are shown. Bars = 200 µm

663

664   **Fig. S6.** Arbuscule phenotype in Myc and Myc + CO plants. In both cases, fully developed and highly

665   branched arbuscules were observed at 28 dpi. Twenty root colonized sections from five biological

666   replicates were used for each treatment. Bars = 30 µm.

667

668   **Table S1.** Primers used for qRT-PCR analysis
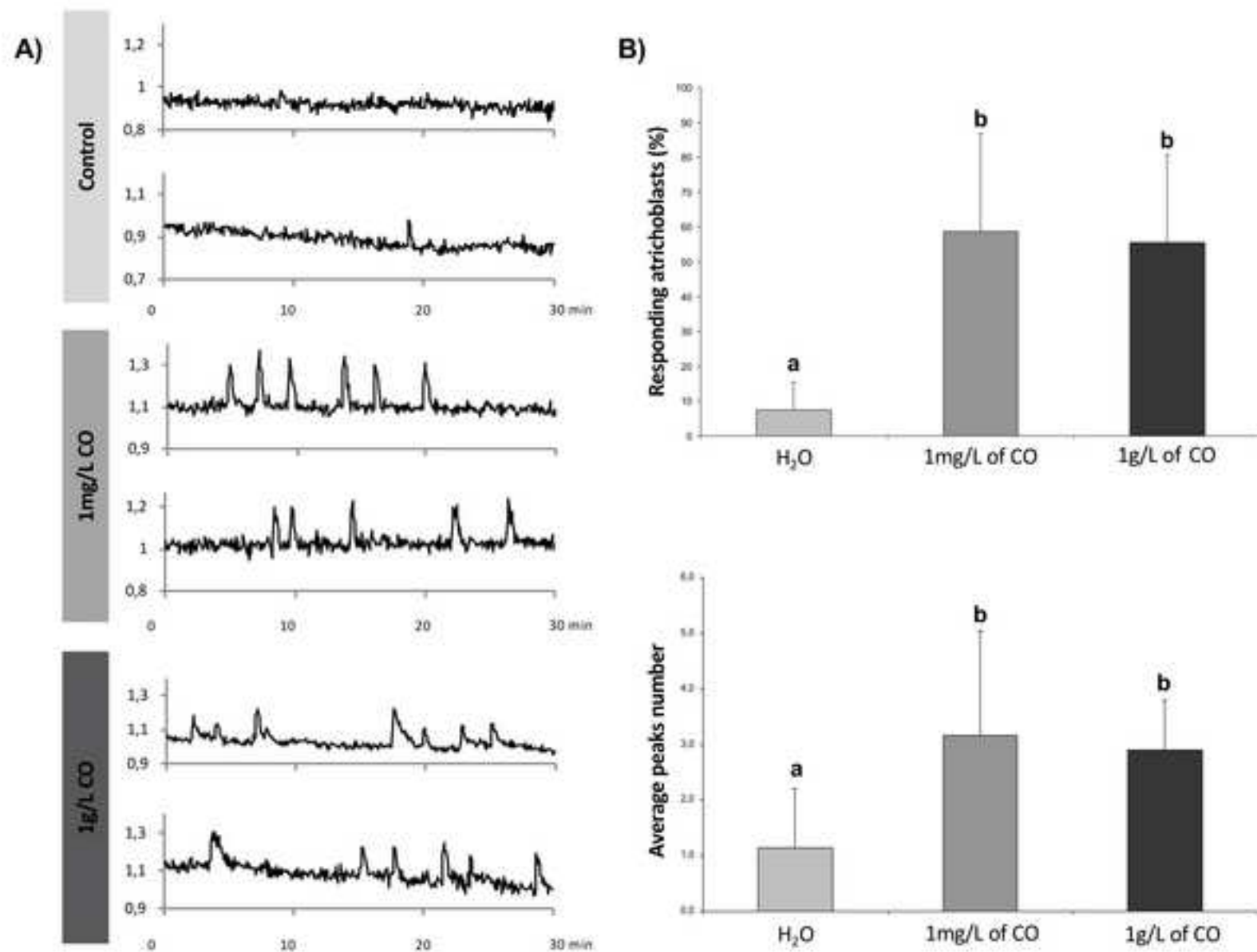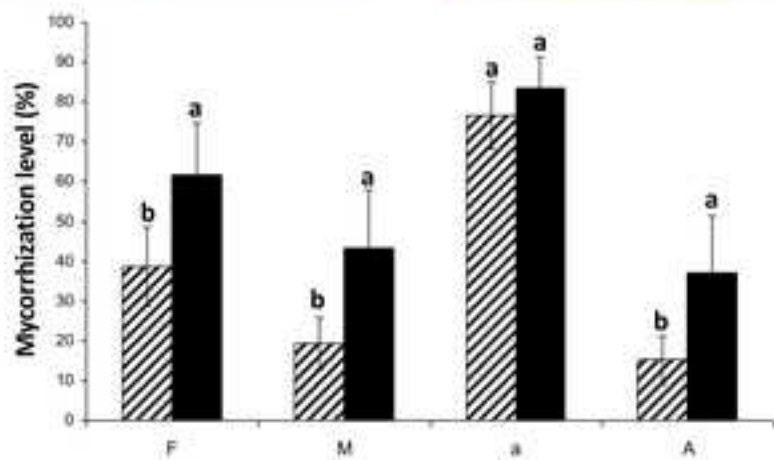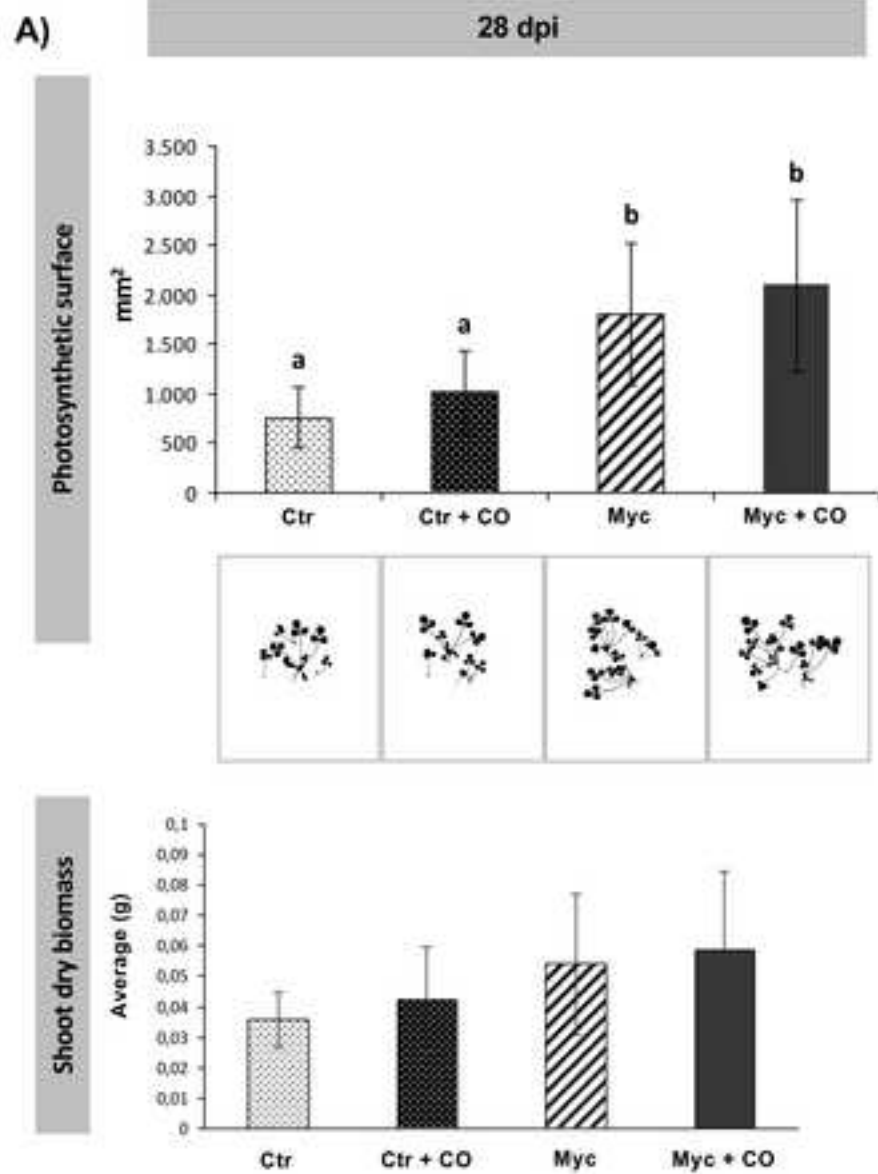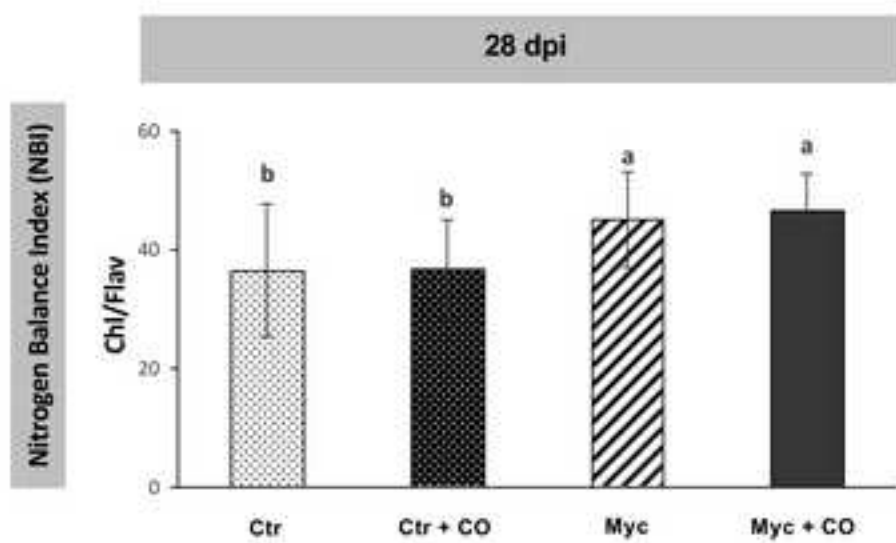
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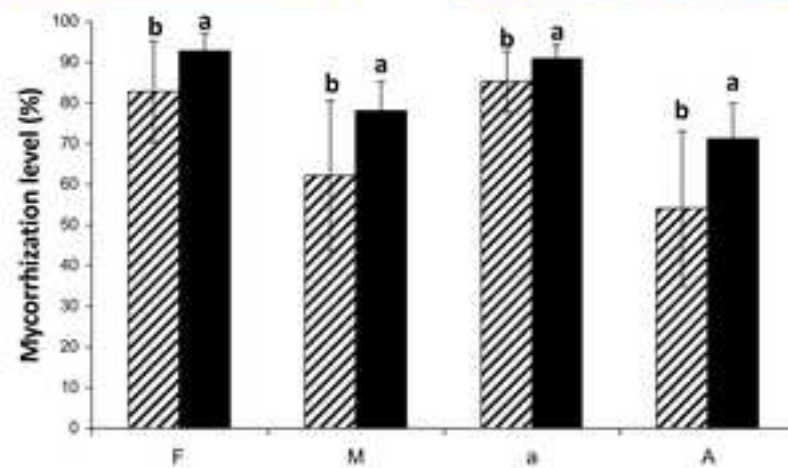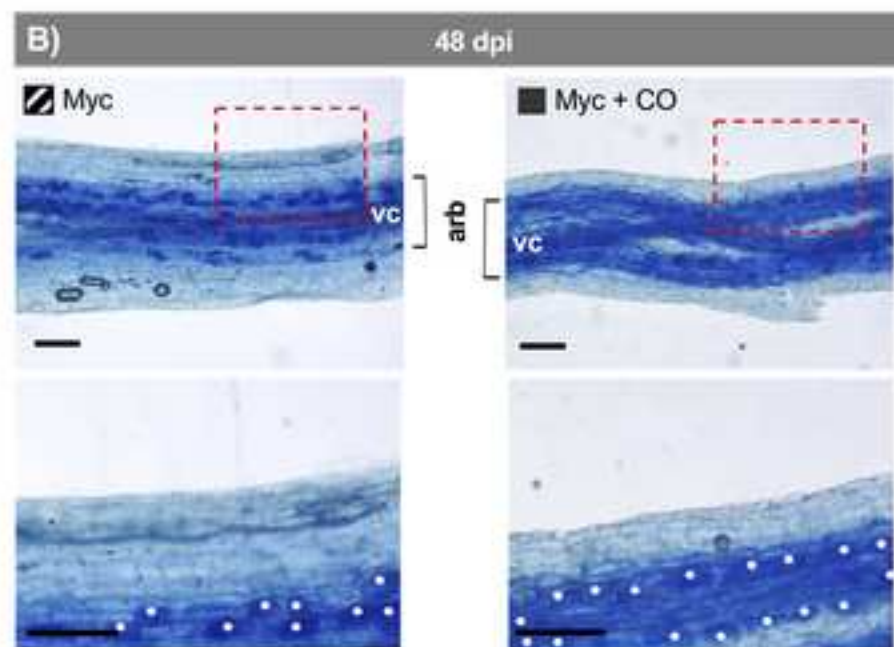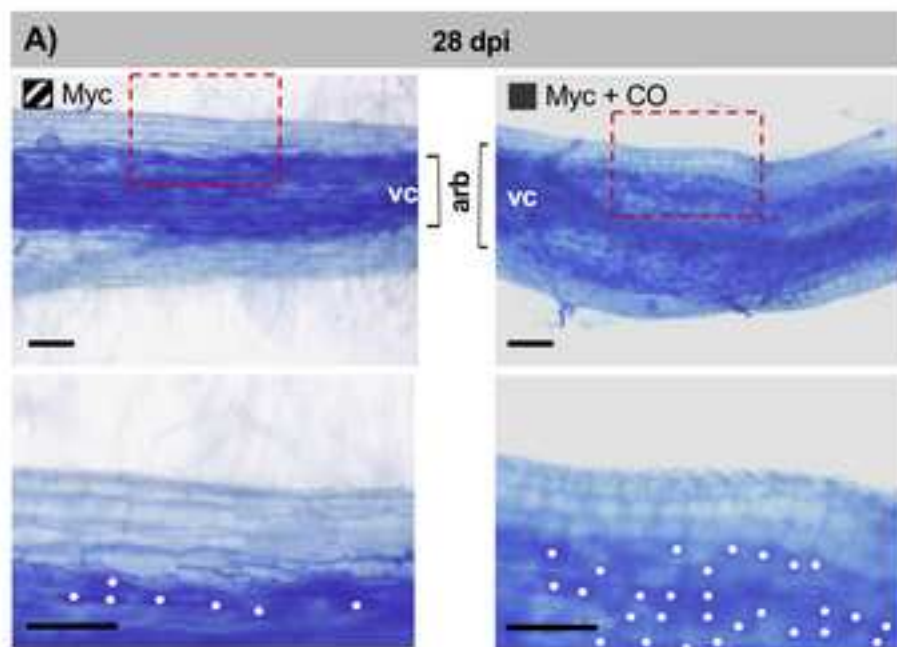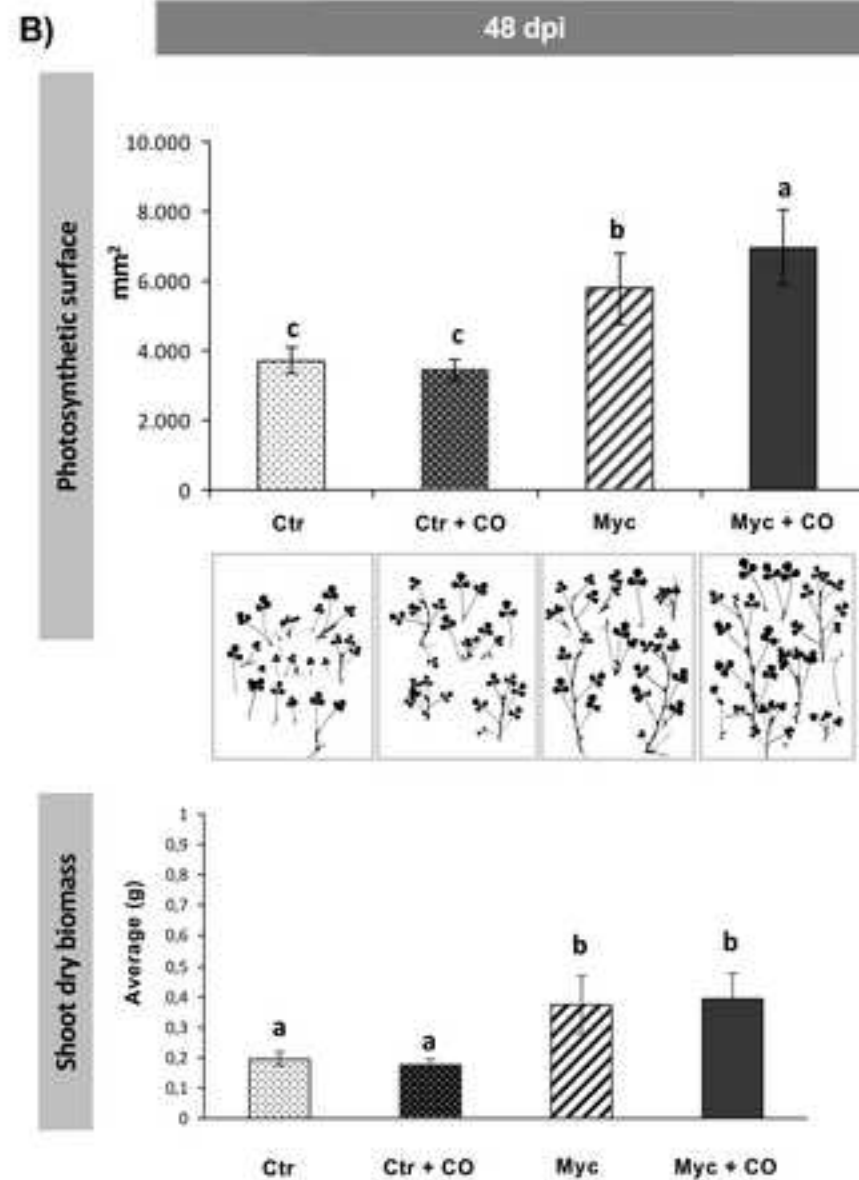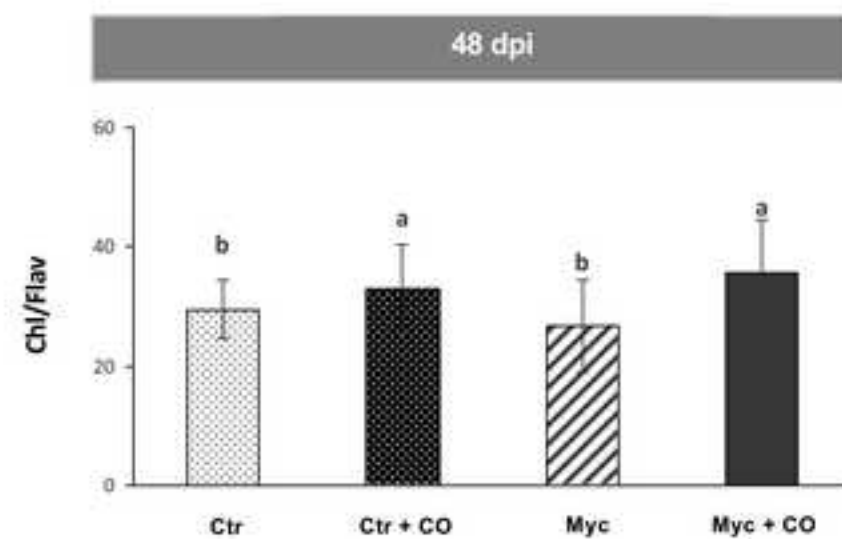
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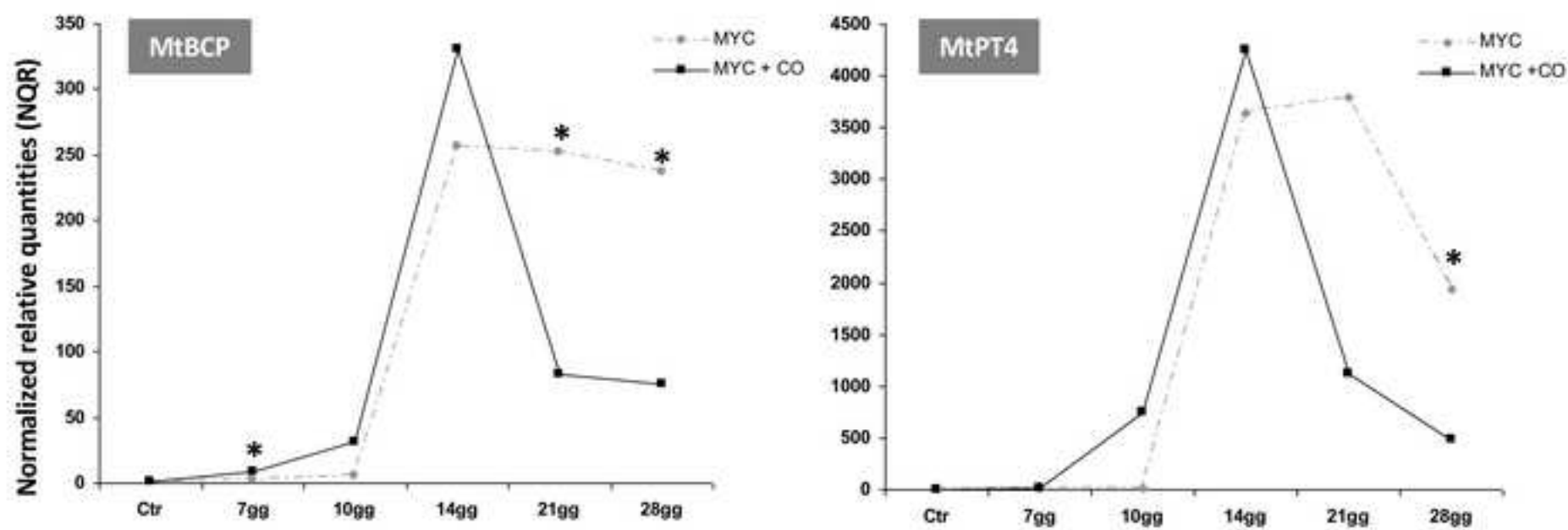
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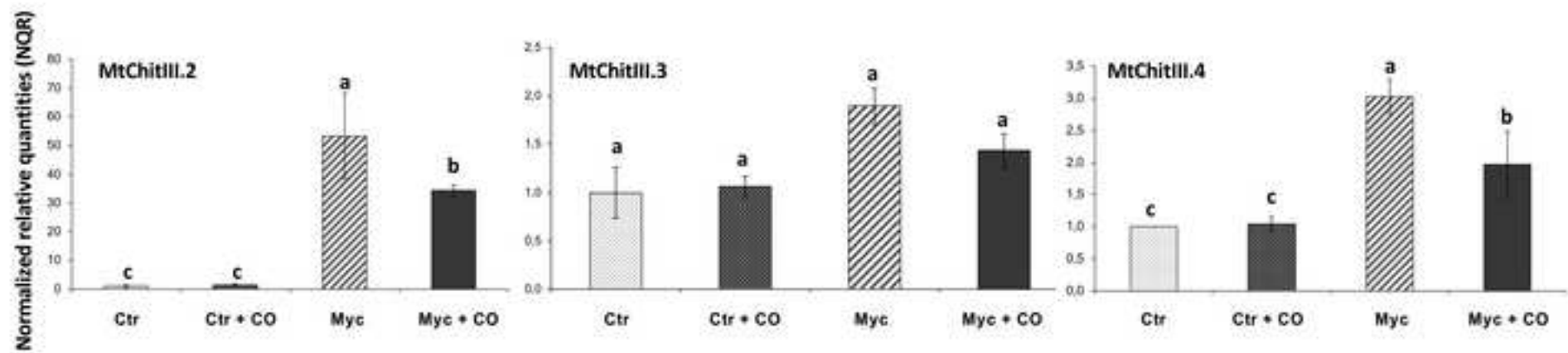
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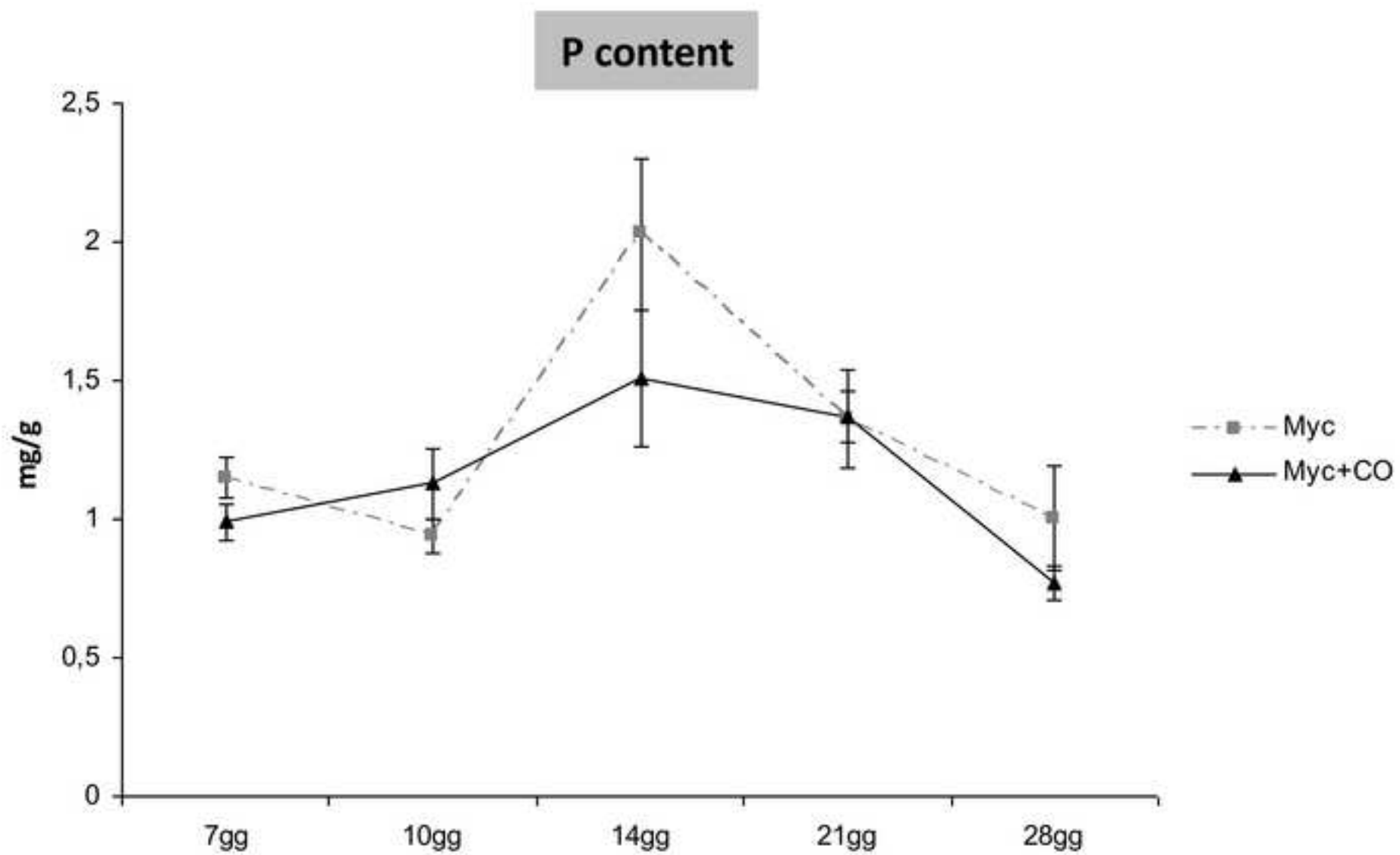
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Table S1

Table S1

Supplementary figures

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