

1 Title: Developmental plasticity of *Brachypodium distachyon* in response to P deficiency: modulation by
2 inoculation with phosphate-solubilizing bacteria

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23 Abstract

24 Background Mineral P fertilisers must be used wisely in order to preserve rock phosphate, a limited and non-
25 renewable resource. The use of bio-inoculants to improve soil nutrient availability and trigger an efficient plant
26 response to nutrient deficiency is one potential strategy in the attempt to decrease P inputs in agriculture.

27 Method A gnotobiotic co-cultivation system was used to study the response of *Brachypodium distachyon* to
28 contrasted P supplies (soluble and poorly soluble forms of P) and inoculation with P solubilizing bacteria.
29 *Brachypodium's* responses to P conditions and inoculation with bacteria were studied in terms of developmental
30 plasticity and P use efficiency.

31 Results *Brachypodium* showed plasticity in its biomass allocation pattern in response to variable P conditions,
32 specifically by prioritizing root development over shoot productivity under poorly soluble P conditions. Despite
33 the ability of the bacteria to solubilize P, shoot productivity was depressed in plants inoculated with bacteria,
34 although the root system development was maintained. The negative impact of bacteria on biomass production in
35 *Brachypodium* might be attributed to inadequate C supply to bacteria, an increased competition for P between
36 both organisms under P-limiting conditions, or an accumulation of toxic bacterial metabolites in our cultivation
37 system. Both P and inoculation treatments impacted root system morphology. The modulation of
38 *Brachypodium's* developmental response to P supplies by P solubilizing bacteria did not lead to improved P use
39 efficiency.

40 Conclusion Our results support the hypothesis that plastic responses of *Brachypodium* cultivated under P-limited
41 conditions are modulated by P solubilizing bacteria. The considered experimental context impacts plant–bacteria
42 interactions. Choosing experimental conditions as close as possible to real ones is important in the selection of P
43 solubilizing bacteria. Both persistent homology and allometric analyses proved to be useful tools that should be
44 considered when studying the impact of bio-inoculants on plant development in response to varying nutritional
45 context.

46

47 Keywords

48 biomass allocation, root system morphology, P use efficiency, bio-inoculants, P solubilizing bacteria

49

50 Abbreviations

51 HA hydroxyapatite

52 PPUE physiological P use efficiency

53 PSB phosphate-solubilizing bacteria

54 PUpE P uptake efficiency

55 PUtE P utilization efficiency

56 PUE P use efficiency

57 RMF root mass fraction

58 SMA standardized major axis

59 TCP tricalcium phosphate

60 TRL total root length

61 **1 Introduction**

62 An important challenge for this century is to implement sustainable cropping systems that preserve the
63 environment and non-renewable resources. This is particularly true concerning phosphate rock, a finite resource
64 mined from only a few countries (Cooper et al. 2011). Phosphate rock constitutes the main source of phosphorus
65 inorganic fertiliser and has been extensively used by farmers in industrialized countries during the past century.
66 Given the growing demand for food, the increasing demand for P fertilisers is predicted to continue (mainly in the
67 developing countries). Although this is a controversial issue, it is estimated that global commercial P reserves
68 could be depleted within 50 to 100 years (Cordell et al. 2009). In the foreseeable future, a peak in P production is
69 expected in 2033. This implies that the growing demand for P will overtake the economically available supply due
70 to decreasing quality and accessibility of the remaining phosphate rock reserves (Cordell and White 2011). As the
71 European Union is strongly dependent on P imports (van Dijk et al. 2016), phosphate rock has been classified as
72 a “critical raw material” by the European Commission since 2014 (European Commission 2014). These
73 considerations emphasize the necessity of adapting fertilisation strategies.

74 In industrialized countries, the excessive use of P fertilisers has led to an accumulation of P in agricultural soils,
75 constituting a new source of P reserves known as “legacy soil P” (van Dijk et al. 2016; Menezes-Blackburn et al.
76 2018; Rowe et al. 2016). In some regions, this accumulation has reached levels that generate an environmental
77 risk of watercourse contamination and subsequent eutrophication (Haygarth et al. 2014). It has been estimated that
78 the total soil P stock for arable and grassland soils represents 352 ± 26 years of agronomic P use, with
79 orthophosphate and monoester (organic) phosphate accounting for the greatest proportion (study based on 258
80 different soils collected in Europe, Oceania and North America; Menezes-Blackburn et al. 2018). However, this
81 soil P reserve is unavailable to plants due to the capacity of many soils to fix P (Shen et al. 2011). Soil P reserve
82 mobilizing technologies must be developed in order to reduce the use of inorganic P fertilisers. The requirement
83 for inorganic P fertilisers could be reduced by 50% if legacy soil P was included in nutrient management practices
84 (Sattari et al. 2012).

85 Plants have developed strategies to cope with P deficiency and enhance their P use efficiency (PUE), including
86 alteration of the root morphology and architecture, as well as exudation of carboxylates and hydrolytic enzymes
87 for P solubilization. Micro-organisms can also be useful in mobilizing soil P reserves, by directly increasing the P
88 availability in soils (through solubilization) or enhancing plant P nutrition processes (through hormonal stimulation
89 of root growth, for example) (Richardson et al. 2011). The use of micro-organisms, broadly called biological
90 inoculants (“bio-inoculants”), is considered a technology to improve soil P use by crops and pastures (Owen et al.
91 2015). Those products belong to the biostimulants category as defined by du Jardin (2015). Biostimulants
92 including substances and/or microorganisms act in addition to fertilisers, with the aim of optimising the efficiency
93 of those fertilisers and reducing nutrient application rates (European Parliament and Council of the EU 2019).

94 Plants exhibit modular growth, potentially allowing them to add new branches to their body. New meristems are
95 exposed to the direct influence of the environment, making plant growth a flexible process (Schmid 1992).
96 Phenotypic plasticity, i.e. the environment-driven alteration of a phenotype, gives the plant a great potential to
97 respond to fluctuating environments (Nicotra et al. 2010; Schmid 1992). It is, at least in part, genetically controlled
98 and heritable (Nicotra et al. 2010). Breeding programs have traditionally opted for phenotypic stability over
99 plasticity, ensuring high yield in constant agricultural systems with high inputs. However, the uncertainty of the

100 future environment and climate requires us to reconsider the place of phenotypic plasticity in breeding strategies.
101 Indeed, phenotypic plasticity can be an advantage to plants living in changing or heterogeneous environments by
102 increasing plant fitness (Lobet et al. 2019; Nicotra et al. 2010). The plastic response of plants to abiotic factors and
103 to the presence of microorganisms still needs clarification. Additionally, the microbial-triggered change in plants'
104 plastic response to P deficiency deserves greater attention in order to optimize plant P nutrition and reduce the use
105 of fertilisers (Goh et al. 2013).

106 This study aimed to characterize *Brachypodium distachyon*'s response to contrasted P supplies (soluble and poorly
107 soluble forms of P), as well as the impact of plant inoculation with single strains of phosphate-solubilizing bacteria
108 (PSB) on this response in terms of developmental plasticity. The following hypotheses were tested: (i) biomass
109 allocation and root system development in *Brachypodium* show plasticity in response to contrasted P conditions;
110 (ii) inoculation with PSB modulates the plant's plastic response to contrasted P supplies; and (iii) this modulation
111 induces changes in plant PUE. Biomass accumulation and allocation, shoot P concentration and PUE, as well as
112 root architectural traits, were considered. *Brachypodium*'s developmental plasticity was assessed using tools
113 including allometry analysis for the biomass allocation and persistent homology analysis for the root system
114 architecture (RSA). It is the first time, to our knowledge, that these tools have been used to precisely evaluate the
115 impact of biostimulants on a plant's response to nutrient limitation.

116 **2 Material and Methods**

117 **2.1 Plant and bacterial material**

118 *Brachypodium distachyon* (L.) P. Beauv. (Bd21 line) caryopses were kindly provided by Dr Philippe Vain from
119 the John Innes Centre (Norwich, UK) and propagated under greenhouse conditions.

120 Four bacterial strains were selected for their potential plant growth promotion and phosphorus solubilization
121 capacities: *Bacillus velezensis* GB03 (BveGB03), *Bacillus velezensis* FZB42 (BveFZB42), *Pseudomonas*
122 *fluorescens* 29ARP (Pfl29ARP) and *Azotobacter vinelandii* F0819 (AviF0819). The *Bacillus* strains were selected
123 for their plant growth promotion activities on *Poaceae* (Delaplace et al. 2015; Myresiotis et al. 2015; Zhang et al.
124 2014), as well as their ability to solubilize different forms of P (Giles et al. 2014; Idris et al. 2007; Idriss et al.
125 2002; Liu et al. 2015). *Pseudomonas fluorescens* also exhibited P solubilizing activities and promoted wheat
126 (Shaharouna et al. 2008) and maize growth (Li et al. 2017). *Azotobacter vinelandii*, a free diazotrophic bacteria,
127 exhibited P solubilization activity (Nosrati et al. 2014) and PGP traits (Taller and Wong 1989). *Escherichia coli*
128 DH5 α 99B829 (Eco99B829), was selected as a negative control for plant growth promotion (Delaplace et al. 2015;
129 Wu et al. 2016; Zhou et al. 2016). The strains BveGB03 and Eco99B829 were kindly provided by Dr Paul W. Paré
130 and Dr John McInroy (Texas Tech University, Lubbock, TX, USA), Pfl29ARP by Dr Alain Sarniguet (Institut
131 National de la Recherche Agronomique, Rennes, France), AviF0819 by the Katholieke Universiteit Leuven
132 (Leuven, Belgium), and BveFZB42 by Pr Rainer Borriss (Nord Reet UG, Greifswald, Germany). The bacterial
133 strains were stored at 80°C in LB medium containing 20% v/v glycerol before plating.

134 **2.2 In vitro P solubilization assay**

135 One week before the experiment, the bacteria were plated on LB agar plates (2.5% w/v LB broth, Prod. No. L3152;
136 1.5% w/v agar, Prod. No. 05039, Sigma-Aldrich Co., St. Louis, USA) and incubated at 28°C. The day before the
137 experiment, the bacteria were suspended in 40 ml of LB (2.5% w/v LB broth) and incubated overnight at 150 rpm
138 and 30°C (Innova 4340, New Brunswick Scientific Co. Inc., Edison, USA). The concentration of the bacterial

139 suspensions was derived from the optical density, measured at 540 nm. The tubes were centrifuged (20 min at
140 4000 rpm) and the LB medium was removed. The bacterial pellets were rinsed with 25 ml of 10 mM MgSO₄ in
141 order to avoid P contaminations. The tubes were centrifuged again (20 min at 4000 rpm) and the MgSO₄ solution
142 was removed. The bacteria were suspended in an adequate volume of NBRIP medium (National Botanical
143 Research Institute's phosphate growth medium, Nautiyal 1999) containing tricalcium phosphate (Ca₃(PO₄)₂,
144 hereafter named "TCP", Prod. No. C0506.1000, Duchefa Biochemie, Haarlem, The Netherlands) or
145 hydroxyapatite (Ca₅(PO₄)₃OH, hereafter named "HA", Prod. No. 8450.1, Carl Roth GmbH + Co. KG, Karlsruhe,
146 Germany) at a concentration of 5 g/l (pH 7) in order to obtain a bacterial concentration of 10⁷ CFU/ml. Bottles
147 containing 90 ml of NBRIP medium were successively inoculated with 10 ml of the prepared suspensions to obtain
148 a final concentration of 10⁶ CFU/ml, and incubated for 3 days at 30°C and 150 rpm.

149 10 ml were sampled daily for subsequent analysis. 1 ml was subsampled for serial dilution and plating on LB agar
150 plates in order to monitor bacterial growth. The remaining samples were centrifuged and the supernatant was filter-
151 sterilized (pore size 0.2 µm) for pH and soluble P content measurements. The P content in the solution (as soluble
152 phosphate) was measured according to the phosphomolybdate blue colorimetric method (Murphy and Riley 1962)
153 (Prod. No. 69888, molybdate reagent solution, Fluka Sigma-Aldrich Co., St. Louis, USA).

154 **2.3 *Brachypodium*-bacteria co-cultivation in axenic conditions**

155 One week before the experiment, the bacteria were plated on LB agar plates and incubated at 28°C. The day before
156 the experiment, the bacteria were suspended in 40 ml of LB and incubated overnight at 150 rpm and 30°C. The
157 tubes were centrifuged (20 min at 4000 rpm) and the LB medium was removed. Inoculums at 10⁸ CFU/ml were
158 finally prepared in 10 mM MgSO₄ for subsequent inoculation of the plantlets.

159 *Brachypodium distachyon* Bd21 seeds were surface sterilized (30 s in 70% v/v ethanol, rinsed once with sterile
160 water, 10 min in sodium hypochlorite 5% v/v, rinsed three times with sterile water) and stratified for 2 days at 4°C
161 on Hoagland agar plates (0.125% w/v Hoagland, Prod. No. DU1201, Duchefa Biochemie, Haarlem, The
162 Netherlands; 0.094% w/v Ca(NO₃)₂·4H₂O; 0.8% w/v Plant agar, Prod. No. P1001, Duchefa Biochemie, Haarlem,
163 The Netherlands). The seeds were then incubated for 24 hours in a growth chamber (23°C, 16h/8h day light, PPF
164 140 µmol.m⁻².s⁻¹) for germination.

165 Homogeneous 24 hour-old plantlets were selected and inoculated with bacteria by dipping them into 10 mM
166 MgSO₄ containing a bacterial strain at 10⁸ CFU/ml for 10 minutes (control plantlets were dipped into 10 mM
167 MgSO₄). The plantlets were then transferred into Magenta® boxes (GA-7 Magenta vessel, Magenta LLC,
168 Lockport, USA) filled with 180 g of sterilized black gravel (rinsed three times with tap water and autoclaved; 1-3
169 mm quartz gravel, prod. no. 400723, Flamingo, Geel, Belgium) and 50 ml of sterile nutrient solution. One plantlet
170 was placed into each Magenta® box. Three modified Hoagland nutrient solutions and a reference solution,
171 corresponding to the contrasting P treatments, were used: a P-limiting supply containing 25µM of KH₂PO₄ ("P-"),
172 a P-limiting supply supplemented with 1 g/l TCP ("P-/TCP") or 1 g/l HA ("P-/HA"), and a P-sufficient supply
173 containing 1mM KH₂PO₄ ("P+"). The boxes were sealed with Leukopor® tape (prod.no. 02454-00, BSN medical
174 GmbH, Hamburg, Germany) and incubated in the growth chamber for four weeks (23°C, 16h/8h day light, PPF
175 140 µmol.m⁻².s⁻¹). Six independent experiments were performed (three with P-, P-/TCP and P+ treatments; three
176 with P-, P-/HA and P+ treatments) and five plants were cultivated for each treatment (90 plants per experiment).
177 Four week-old plants were harvested and cut to measure fresh biomass accumulation in shoot and roots. The

178 presence of bacteria was assessed by scratching agar plates with the root system. The root system was scanned for
179 three plants per treatment (1200 dpi, Epson Perfection V800 Photo, Epson America Inc., Long Beach, USA) in
180 order to perform RSA analyses. Shoots were stored at -80°C before P content measurements. Total biomass and
181 root mass fraction (hereafter named “RMF”, mg root biomass/mg total biomass) were computed from the measured
182 biomasses. RMF was recorded in order to analyse biomass allocation in *Brachypodium*, considering allocation as
183 a partitioning process. According to this perspective, plants divide a given amount of resources among structures
184 according to their developmental priorities (Weiner 2004).

185 **2.4 P concentration in plant tissues**

186 P content in *Brachypodium* shoots was measured by ICP-OES on frozen samples (C.A.R.A.H. ASBL, Ath,
187 Belgium). The samples were calcinated overnight at 450°C. The ashes were then suspended in nitric acid for
188 digestion. The P concentration was measured by ICP-OES (Thermo Fisher iCAP 7600, Thermo Fisher Scientific,
189 Waltham, USA). The five replicates of each treatment in each independent experiment were pooled. Three pooled
190 samples were analysed for the P-/TCP and P-/HA treatments. Six pooled samples were analysed for the P- and P+
191 treatments. The results were expressed as total shoot P concentration (µg P/ mg fresh weight).

192 **2.5 Root system architecture measurement**

193 An automated evaluation of the total root length (“TRL”) was performed for all scanned root systems using the
194 ImageJ macro IJ_Rhizo (Pierret et al. 2013; Schneider et al. 2012). For each image, the TRL was estimated
195 using the Kimura method as it provides more accurate length estimates than the other methods available in
196 IJ_Rhizo (Delory et al. 2017).

197 In addition, more detailed root system architecture analyses were performed using SmartRoot (Lobet et al. 2011).
198 Only the 1st and 2nd order roots were analysed because the thinner, higher order, roots break easily at harvest. These
199 manual analyses were performed for the control treatment and the two strains found to impact plant development
200 the most. The RSML (Root System Markup Language) outputs were then processed with the archiDART package
201 for morphological analysis using persistent homology (R 3.5.2, R core Team 2018; archiDART package version
202 3.3, Delory *et al.*, 2016; Delory et al. 2018). A geodesic distance function was used to compute a persistence
203 barcode for each root system. The degree of dissimilarity between barcodes (i.e. root systems) was assessed by
204 computing a pairwise distance matrix containing dissimilarities calculated using a bottleneck distance method.
205 Morphological differences between root systems were then visualized using multidimensional scaling (R 3.5.2, R
206 Core Team 2018).

207 **2.6 P use efficiency**

208 The P use efficiency analysis was performed by considering three different parameters: (i) the P uptake efficiency
209 (PU_pE, µg P/mg P applied), corresponding to the shoot P content per unit of soluble P applied; (ii) the P utilization
210 efficiency (PU_tE, mg FW/µg P), corresponding to the biomass produced by unit of shoot P; and (iii) the
211 physiological P use efficiency (PPUE, mg² FW/µg P), corresponding to the produced biomass divided by the tissue
212 P concentration (Neto et al. 2016).

$$213 \quad PU_{pE} = \frac{[P] \times FW}{P_{applied}}$$

$$214 \quad PU_{tE} = \frac{FW}{[P] \times FW}$$

215
$$PPUE = \frac{FW}{[P]}$$

216 **2.7 Statistical analyses**

217 The relationship between P solubilization and pH variation in the NBRIP medium was studied by performing
218 regression analyses (lm function, R 3.5.2, R Core Team 2018). The model order was increased until there was not
219 significant difference with the higher order model (anova function, R 3.5.2, R Core Team 2018).

220 Three-way ANOVAs were performed to study the impact of P supply, bacteria inoculation, independent temporal
221 repetitions and the interaction between P supply and bacteria inoculation on the following factors: shoot, root and
222 total biomass parameters; RMF; TRL; shoot P concentration; and the three components of PUE. A model with
223 crossed fixed factors was applied (lm, glm and anova functions, R 3.5.2, R Core Team 2018; Gamma family
224 distribution with a log-link function was used for GLM models). Dunnett's post-hoc tests were performed to
225 compare the treatments to the control situation (non-inoculated plants for inoculation treatment, P+ for P treatment)
226 (R 3.5.2, R Core Team 2018; multcomp package version 1.4-8, Hothorn et al. 2008).

227 Allometry analyses were performed on shoot and root biomass in order to study the biomass allocation pattern.
228 The "smatr" package (R 3.5.2, "smatr" version 3.4-8) was used for estimation, inference and plotting of allometric
229 lines as well as for checking assumptions (Warton et al. 2012). The standardized major axis ("SMA") analysis was
230 used and all variables were log-transformed. In brief, this analysis consists of a model II regression, estimating
231 how one variable scales against another. The obtained allometric trajectories depict the relative development of
232 the shoot and root compartments, i.e. how the root system growth impacts the shoot development. Inference
233 statistics compare coefficients of the regression lines (slope and elevation) between the populations (Warton et al.
234 2006). Firstly, differences in slope between groups were tested. If there was no difference in slope between groups,
235 differences in elevation were tested using a common slope for all groups. When significant differences between
236 groups were highlighted, pairwise multiple comparisons were performed in order to identify which populations
237 differed from each other. Differences in slope (i.e. investment in shoot biomass per additional unit of root biomass)
238 or elevation (i.e. shoot productivity for similar root biomass) among treatments led to different allometric
239 trajectories. Change in allometric trajectory due to different treatments revealed plasticity in the biomass allocation
240 process (Weiner 2004; Xie et al. 2015). The analysis of allometric trajectories is complementary to the analysis of
241 RMF for the study of biomass allocation plasticity.

242 Differences in root system architecture were investigated using permutational multivariate analysis of variance
243 (PERMANOVA) (R 3.5.2, R Core Team 2018; vegan package version 2.5-4, Oksanen et al. 2019). The
244 dissimilarity matrix used in the model formula was the pairwise distance matrix returned by the persistent
245 homology analysis of plant root systems. Bacterial strain, P treatment and their interaction were used as
246 independent variables in the model. For each fixed factor, a post-hoc test was performed by running a separate
247 PERMANOVA for each pairwise comparison. *P* values were adjusted for multiple comparisons using the
248 Bonferroni method.

249 All figures shown in this study were generated using the "ggplot2" package (R 3.5.2, "ggplot2" version 3.1.0).

250 Data and R scripts are accessible on Zenodo repository at <https://doi.org/10.5281/zenodo.3555566>

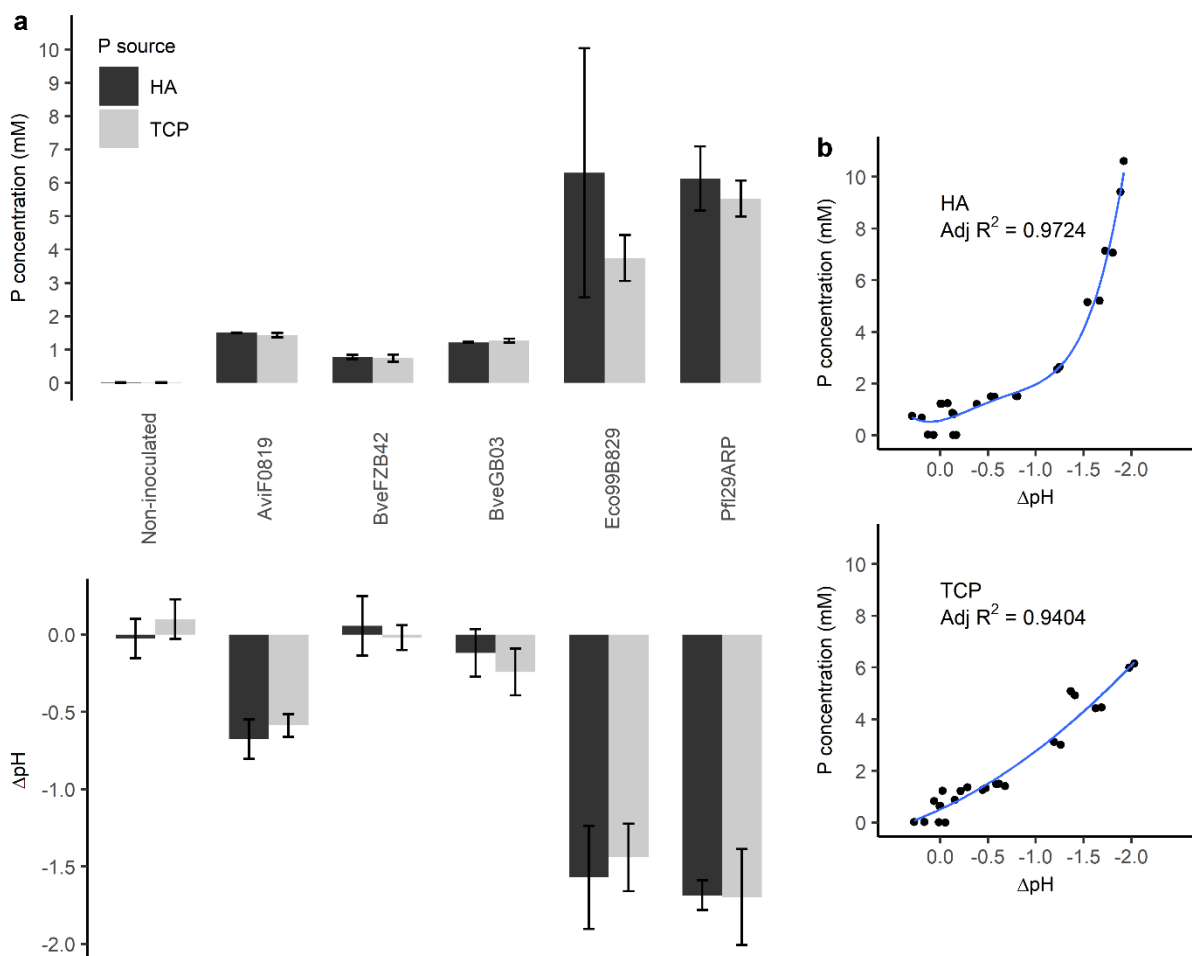
251

252 3 Results

253 3.1 The selected bacterial strains solubilized tricalcium phosphate and hydroxyapatite, while acidifying 254 their growth medium

255 The bacteria's ability to solubilize poorly available forms of P was assessed using TCP and HA in a modified
256 NBRIP medium. After three days, all the selected strains were able to solubilize both forms of P to some extent
257 (Fig. 1a) compared to the non-inoculated control treatment. For both forms of P, the best performing bacterial
258 strains were Eco99B829 and Pfl29ARP. The solubilization of TCP and HA were similar for all bacterial strains
259 with the exception of Eco99B829, which exhibited a stronger solubilization ability for HA despite a greater
260 variability between independent replicates. All the strains were able to maintain stationary populations during the
261 duration of the experiment (data not shown). BveGB03, AviF0819, Pfl29ARP and Eco99B829 generated a pH
262 drop during the experiment for both forms of P (Fig. 1a). As for the P concentration, Eco99B829 and Pfl29ARP
263 induced the strongest acidification.

264 Regarding HA solubilization, the relationship between the soluble P concentration and Δ pH in the growing
265 medium was best fitted by a 4th order polynomial model (Fig. 1b). The HA solubilization activity clearly intensified
266 as the acidification became stronger. The regression between soluble P concentration and Δ pH for TCP
267 solubilization was best fitted by a 2nd order polynomial model (Fig. 1b). As for HA, the TCP solubilization activity
268 intensified with increasing pH variation, but to a lesser extent.



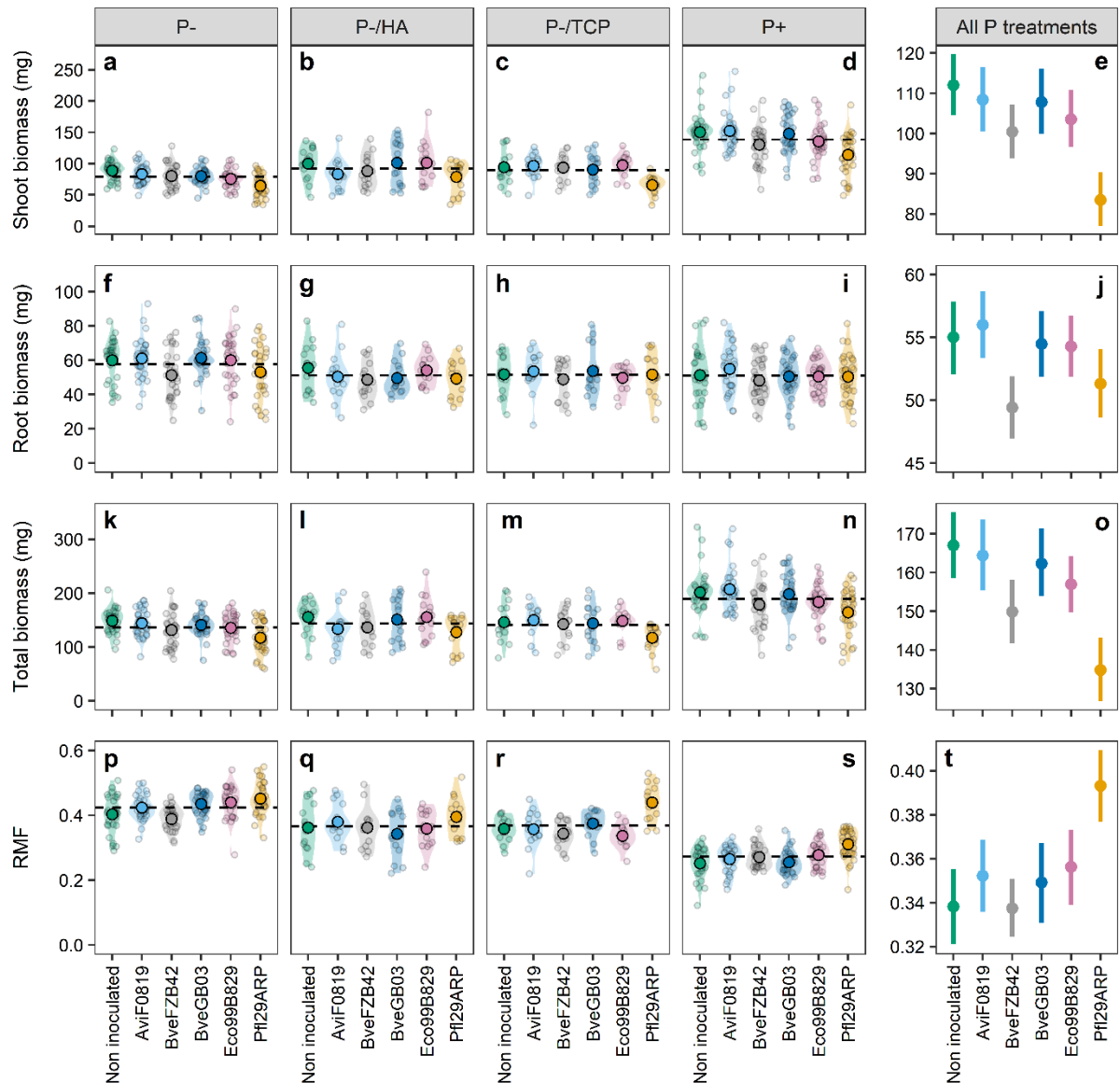
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270 **Fig. 1** (a) Soluble P concentration and pH variation in NBRIP medium after three days of bacteria cultivation in
271 the presence of either TCP or HA as poorly soluble forms of P (n=4, mean \pm SD). (b) Regression curves linking

272 the observed P concentration and the ΔpH in the growing medium after three days of incubation. For each
273 regression model, adjusted R^2 values are displayed on the graphs. Regression coefficients for HA solubilization: y
274 $= 0.571 - 0.814x + 2.7675x^2 + 4.219x^3 + 2.039x^4$; regression coefficients for TCP solubilization: $y = 0.513 -$
275 $1.727x + 0.530x^2$

276 **3.2 Biomass accumulation in *Brachypodium* was altered by soluble P deficiency and inoculation with P** 277 **solubilizing bacteria**

278 Shoot biomass production was lower in plants grown under P-, P-/TCP and P-/HA conditions, with a diminution
279 of 43.1%, 35.2% and 33.4% compared to the P+ treatment, respectively ($P < 0.001$, Fig. 2a-d). Plant inoculation
280 with PSB strains had either no impact or induced a lower shoot biomass accumulation (Fig. 2e). Inoculation with
281 BveFZB42 and Pfl29ARP led to a significantly lower shoot biomass, with up to 13.2% reduction in plants
282 inoculated with BveFZB42 under the P+ treatment and 30.3% reduction in plants inoculated with Pfl29ARP under
283 the P-/TCP treatment ($P < 0.001$). The impact of P conditions on the accumulation of biomass in roots was more
284 limited, with only plants grown under the P- treatment having a significantly greater root biomass (+13.3%)
285 compared to the plants exposed to P+ conditions ($P < 0.001$, Fig. 2f-i). Inoculation had either no impact or a negative
286 impact on the accumulation of biomass in roots ($P = 0.003$, Fig. 2j). Indeed, plants inoculated with BveFZB42
287 exhibited a significant reduction of the root biomass of up to 14.5% under the P- treatment. The total biomass
288 decreased by 27.8%, 25.2% and 24.1% under the P-, P-/TCP and P-/HA treatments respectively ($P < 0.001$ Fig. 2k-
289 n). Plant inoculation with PSB strains led to a repression of the biomass accumulation at the whole plant level
290 ($P < 0.001$, Fig. 2o). Inoculation with BveFZB42 and Pfl29ARP induced a significantly lower total biomass in
291 comparison with the non-inoculated control. The growth reduction reached 12.2% with BveFZB42 under P-/HA
292 conditions and 21.1% with Pfl29ARP under P- conditions. A table comprising mean values per treatment, standard
293 deviation and coefficients of ANOVAs is available in Online Resource 1.



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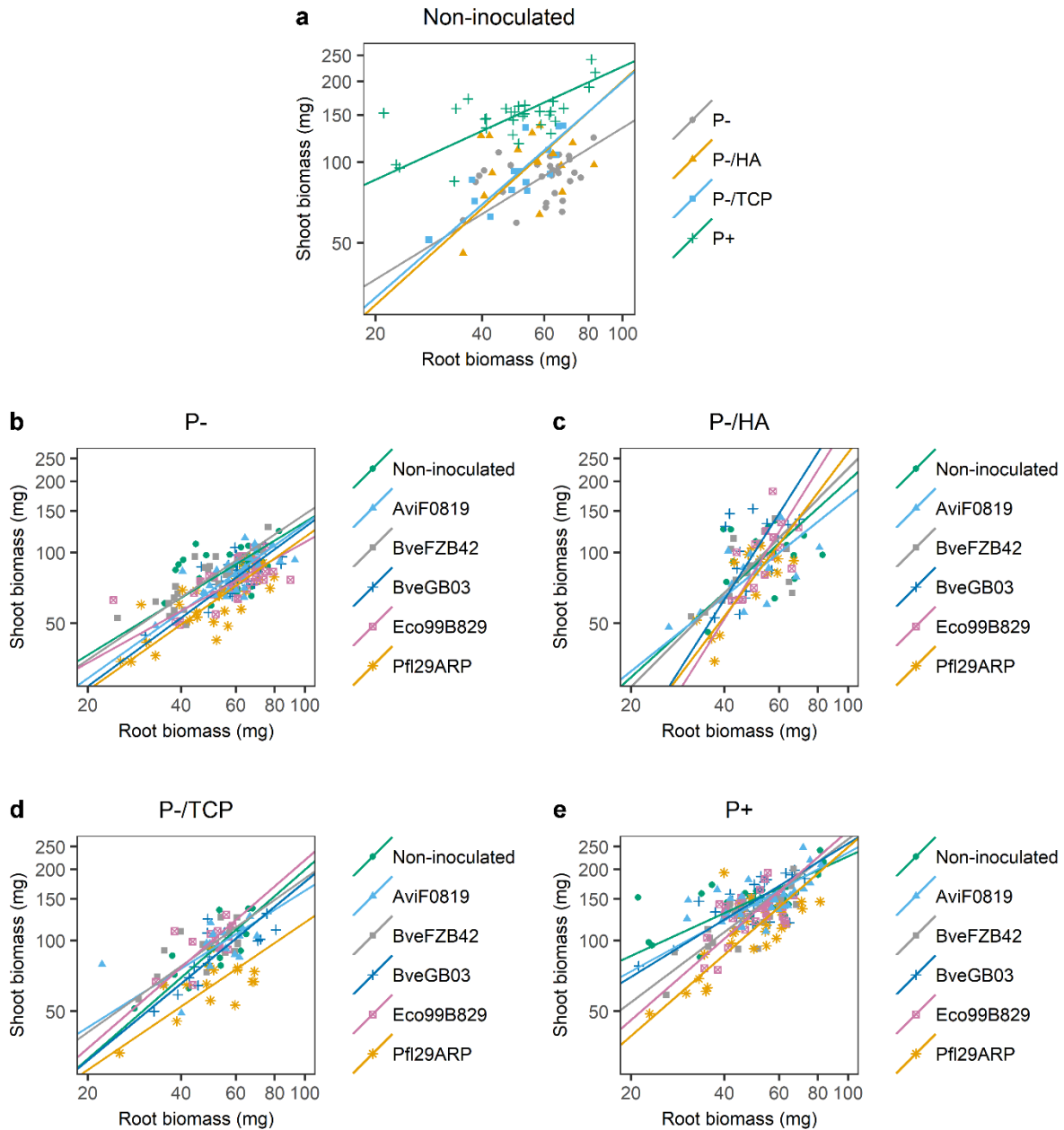
295 **Fig. 2** Average shoot biomass (a-e), root biomass (f-j), total biomass (k-o) and root mass fraction (p-t) of four-
 296 week-old *Brachypodium* plantlets exposed to contrasted P supplies and either inoculated or not inoculated with
 297 bacteria. n = 30 for the P- and P+ treatments, n=15 for the P-/HA and P-/TCP treatments. For each P treatment,
 298 the grand mean is shown by a dashed horizontal line. For each inoculation treatment, large black-circled dots
 299 represent mean values, and shaded areas show the density distribution of each population. Individuals are displayed
 300 as small grey-circled dots in the graphs. In panels e, j, o and t, values are means +/- 95% confidence intervals
 301 calculated across P treatments

302 **3.3 Shifts in biomass partitioning and allometric trajectories of *Brachypodium* were observed when**
 303 **exposed to contrasted P supplies and inoculated with P solubilizing bacteria**

304 Exposure of *Brachypodium* to soluble P limitation (P-, P-/HA and P-/TCP) increased RMF by 55.8%, 34.9% and
 305 35.7% compared to the P+ treatment, respectively ($P < 0.001$, Fig. 2p-s). The impact of *Brachypodium* inoculation
 306 with PSB was dependent on the P environment, as there was a significant interaction between these two variables
 307 ($P < 0.001$). Plants inoculated with Pf29ARP exhibited the greatest RMF under all treatments (Fig. 2t) and this
 308 effect was significant for the P-, P-/TCP and P+ treatments (12.1%, 22.7% and 23.4% increase, respectively).

309 Under P- conditions, plants inoculated with BveGB03 and Eco99B829 also had a significantly greater RMF
310 compared to non-inoculated plants (Fig. 2p-t). Mean values per treatment, standard deviation and coefficients of
311 ANOVAs are available in Online Resource 1.

312 The allocation pattern between shoots and roots was further analysed using SMA regression models (Fig. 3). In
313 non-inoculated plants grown under P-/TCP and P-/HA conditions, the shoot biomass increase per unit of root
314 biomass was greater than that of non-inoculated plants grown under P- and P+ conditions (slopes: 1.15, 1.19, 0.81
315 and 0.60, respectively; $P=0.021$; Fig. 3a). Non-inoculated plants grown under P+ conditions exhibited the greatest
316 shoot productivity, but invested the lowest amount of biomass into the shoot per unit of root production.
317 Inoculation of plants grown under P- conditions did not induce a significant difference in slope ($P=0.757$, Fig. 3b).
318 Significant differences in elevation were observed ($P<0.001$), with non-inoculated plants and plants inoculated
319 with BveFZB42 showing the greatest shoot productivity and plants inoculated with Pfl29ARP the lowest, for
320 similar root biomass. Under the P-/HA treatment, slope and elevation did not significantly vary among groups
321 ($P=0.174$ and 0.433 , respectively; Fig. 3c), even if plants inoculated with BveGB03 and Eco99B829 showed
322 greater shoot biomass increase per unit of root biomass, when considering the graphical trends. When plants were
323 grown in the presence of TCP, the inoculation with PSB did not significantly affect the slope ($P=0.835$, Fig. 3d).
324 Elevation was significantly altered when plants were inoculated with Pfl29ARP, leading to the lowest shoot
325 productivity for similar root biomass (lowest elevation, $P<0.001$). Significant differences in slope were observed
326 under the P+ treatment ($P=0.008$), with the greatest production of shoot biomass per unit of root production in
327 plants inoculated with Pfl29ARP, Eco99B829 and BveFZB42 (Fig. 3e). SMA coefficients and results of
328 covariance analysis are available in Online Resource 2.



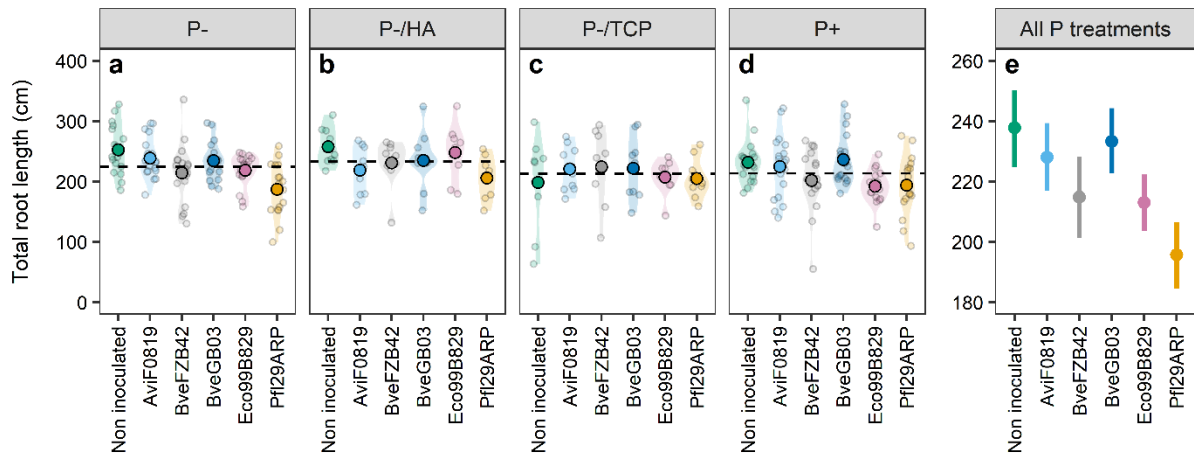
329

330 **Fig. 3** Allometric relationship between shoot biomass and root biomass of four-week-old *Brachypodium* plantlets
 331 exposed to contrasted P supplies and grown with or without bacterial inoculation. X and Y axes are log-scaled.
 332 Symbols represent individuals. Lines represent SMA regression lines. (a) Non-inoculated plants exposed to
 333 contrasted P supplies. Inoculated and non-inoculated plants grown under (b) P-, (c) P-/HA, (d) P-/TCP and (e) P+
 334 conditions

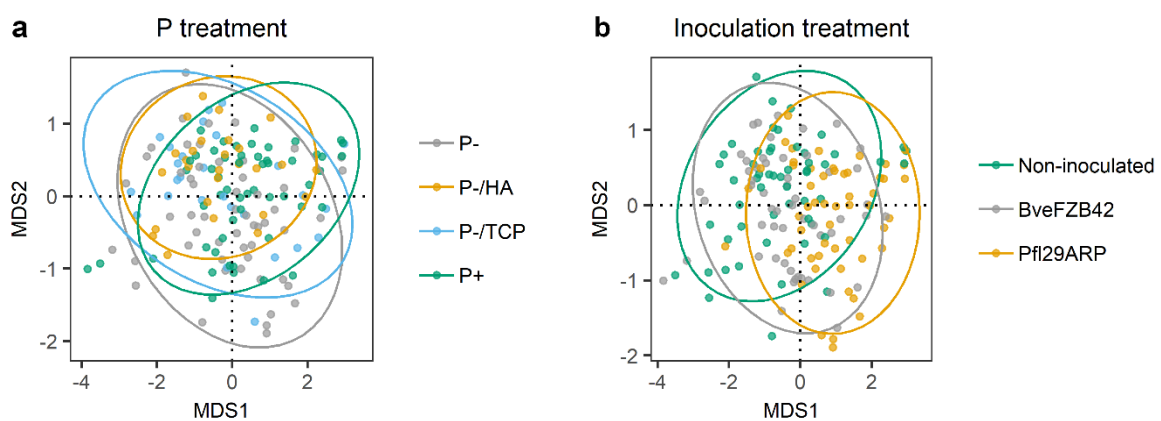
335 **3.4 *Brachypodium* total root length and root system morphology were impacted by P supply and**
 336 **inoculation with P solubilizing bacteria**

337 *Brachypodium* TRL increased by 8.97% when plants were exposed to the P-/HA treatment compared to the P+
 338 treatment ($P=0.023$, Fig. 4a-d). In comparison with the TRL measured in non-inoculated plants, the TRL of plants
 339 inoculated with BveFZB42, Eco99B829 or Pfl29ARP decreased by 9.64%, 11.61% and 16.67% respectively,
 340 whatever the nutritional context ($P < 0.001$, Fig. 4e). Mean values per treatment, standard deviation and coefficients
 341 of ANOVAs are available in Online Resource 3.

342 The persistent homology analysis of the root systems was performed on 1st and 2nd order roots of non-inoculated
 343 plants and plants inoculated with Pfl29ARP or BveFZB42, as those strains showed a strong impact on root biomass
 344 accumulation (Fig. 5). Both PSB inoculation and P treatment had a significant impact on root system morphology
 345 ($P < 0.001$ and $= 0.006$ respectively). Pairwise comparisons revealed that, on average, the morphology of plant root
 346 systems inoculated with Pfl29ARP was different from those of non-inoculated plants and plants inoculated with
 347 BveFZB42. Despite a significant impact of P treatment on root system morphology, pairwise comparisons did not
 348 highlight the P treatments that differed from one another. The coefficients of the statistical analysis are available
 349 in Online Resource 4.



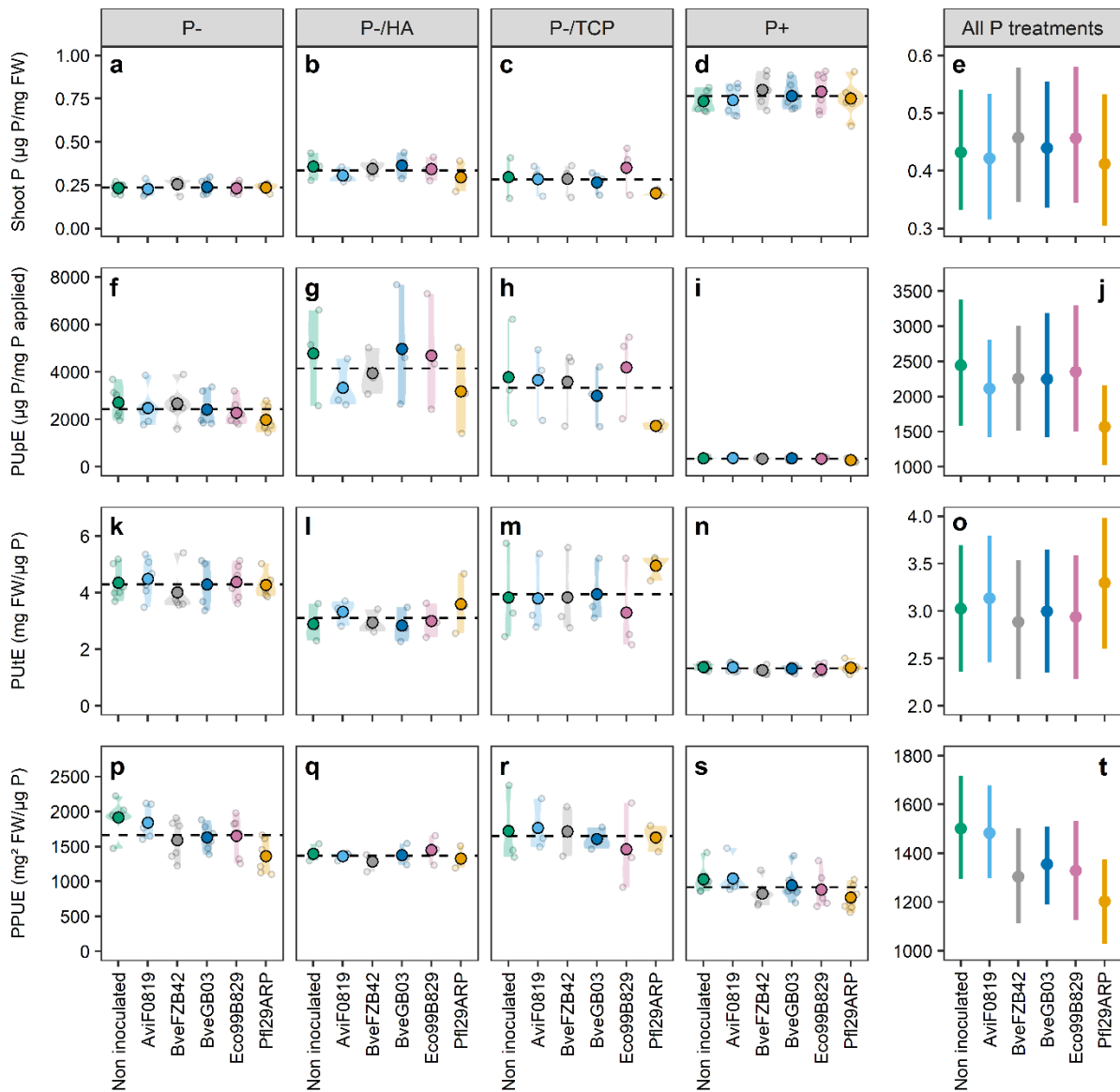
350
 351 **Fig. 4** Average total root length of four-week-old *Brachypodium* plantlets exposed to contrasted P supplies and
 352 either inoculated or not inoculated with bacteria. $n = 18$ for the P- and P+ treatments, $n = 9$ for the P-/HA and P-
 353 /TCP treatments. For each P treatment, the grand mean is shown by a dashed horizontal line. For each inoculation
 354 treatment, large black-circled dots represent mean values, and shaded areas show the density distribution of each
 355 population. Individuals are displayed as small grey-circled dots in the graphs. In panel e, values are means +/- 95%
 356 confidence intervals calculated across P treatments



357
 358 **Fig. 5** Multidimensional scaling plots displaying morphological differences between root systems, induced by P
 359 (a) and inoculation (b) treatments. The Euclidean distance separating two branching structures (dots) on the plot
 360 is a close representation of the true dissimilarity between these structures. 95% confidence ellipses for the centroids
 361 are plotted for each treatment

362 **3.5 Low P availability induced lower shoot P concentration, even in the presence of P solubilizing**
 363 **bacteria**

364 P concentration in the shoot of plants exposed to the P-, P-/HA and P-/TCP treatments was lower than in plants
 365 exposed to P+ (-68.9%, -56.2% and -63.2% respectively; $P < 0.001$; Fig. 6a-d). Plants grown under these three
 366 treatments showed P deficiency symptoms, such as necrosis starting from the apex of mature leaves (Arvalis,
 367 Institut du végétal). Inoculation with bacteria did not help the plants to increase the shoot P concentration, even in
 368 the presence of the potentially mobilizable P sources TCP or HA (Fig. 6e). Mean values per treatment, standard
 369 deviation and coefficients of ANOVAs are available in Online Resource 5.



370
 371 **Fig. 6** Average shoot P concentration (a-e), P uptake efficiency “PUpE” (f-j), P utilization efficiency “PUtE” (k-
 372 o) and physiological P use efficiency “PPUE” (p-t), of four-week-old *Brachypodium* plants grown under
 373 contrasted P supplies and either inoculated or not inoculated with bacterial strains. $n=6$ for the P- and P+
 374 treatments, $n=3$ for the P-/HA and P-/TCP treatments. For each P treatment, the grand mean is shown by a
 375 dashed horizontal line. For each inoculation treatment, large black-circled dots represent mean values, and

376 shaded areas show the density distribution of each population. Individual data points (pool of 5 plantlets) are
377 displayed as small grey-circled dots in the graphs. In panels e, j, o and t, values are means +/- 95% confidence
378 intervals calculated across P treatments

379 **3.6 P supply and inoculation with Pfl29ARP impacted P use efficiency components in *Brachypodium***

380 Regarding the PUpE (i.e. the ratio between shoot P content and applied soluble P), plants exposed to soluble P
381 deficiency had a greater uptake efficiency ($P < 0.001$), with the greatest values measured on plants grown in the
382 presence of TCP and HA (883.5% and 1128.8% increase, respectively, compared to plants exposed to P+; Fig. 6f-
383 i). *Brachypodium* acquired and accumulated a greater amount of P in shoots when TCP or HA were added to the
384 nutrient solution, in comparison with the P- treatment and regardless of the bacterial treatment. Plants inoculation
385 with Pfl29ARP led to lower PUpE values under all P treatments compared to non-inoculated plants (average
386 decrease of 35.8% across all P treatments, $P = 0.011$, Fig. 6j).

387 Plants grown under soluble P deficiency were more efficient at utilizing P for biomass accumulation (PUtE,
388 biomass produced by unit of plant P content; $P < 0.001$; Fig. 6k-n). These plants accumulated more shoot biomass
389 per unit of shoot P content compared to plants exposed to P+ condition. Plants grown under the P- treatment were
390 globally the most efficient. Inoculation of *Brachypodium* by any of the bacterial strains had no significant impact
391 on PUtE ($P = 0.436$, Fig. 6o).

392 The PPUE (i.e. shoot biomass divided by shoot P concentration), was significantly higher in plants grown under
393 P-, P-/HA and P-/TCP conditions compared to plants exposed to sufficient P supply (81.8%, 49.1% and 80.1%
394 increase respectively compared to P+ condition, $P < 0.001$, Fig. 6p-s). Plants exposed to a deficiency in soluble P
395 produced shoot biomass more efficiently at lower shoot P concentration. The inoculation with Pfl29ARP induced
396 a 19.9% reduction in PPUE compared to non-inoculated plants ($P = 0.008$, Fig. 6t). Mean values per treatment,
397 standard deviation and coefficients of ANOVAs are available in Online Resource 5.

398 **4 Discussion**

399 This study aimed to explore the impact of PSB inoculation on the response of *Brachypodium distachyon* Bd21 to
400 contrasted P conditions. *Brachypodium* and the PSB were co-cultivated over four weeks in an *in vitro* gnotobiotic
401 system and exposed to four different nutritional conditions: a low level of soluble P (P-); a low level of soluble P
402 supplemented with poorly soluble forms of P (P-/TCP and P-/HA); and a high level of soluble P (P+). The plant
403 biomass production and allocation, the root system architecture and the P use efficiency were studied.

404 **4.1 *Brachypodium* shows developmental plasticity in response to contrasted P conditions**

405 Our study demonstrated that *Brachypodium* biomass accumulation is highly responsive to P supply, with lower
406 shoot biomass but stable or greater root biomass accumulation under soluble P deficiency compared to high soluble
407 P levels. The reduction in shoot biomass under soluble P deficiency was also reported in *Brachypodium* (Bd21-3)
408 by Poiré et al. (2014) and *Dactylis glomerata* by Haling et al. (2016). Interestingly, a reduction in root biomass
409 was observed in their studies, while our results showed no impact or even an increase in root biomass accumulation
410 when plants were exposed to soluble P deficiency. The younger growth stage obtained in our confined
411 experimental system could explain these results, as the stress was not as intense as it would have been under, for
412 example, greenhouse conditions. Similarly to our results, Giles et al., (2017) showed a decrease in shoot dry weight
413 and an increase in root dry weight in hydroponically grown barley under P deficiency. The stimulation of root
414 development under low P conditions is a common reported response, facilitating the plant to explore the substrate

415 and take up P (Lynch et al., 2012). Nonetheless, it seems that both the growing conditions and the plant growth
416 stage are important factors affecting biomass accumulation in response to P deficiency.

417 *Brachypodium* displayed different allometric trajectories under contrasted P conditions, showing responsiveness
418 of the allocation pattern to the P supply. Plants grown in the presence of TCP or HA exhibited a higher shoot
419 development per unit of root biomass than plants grown under the P- and P+ treatments. From this we can infer
420 that the presence of unavailable but potentially mobilizable P sources induced a reduction of investment into the
421 root compartment, in comparison with plants grown under P- conditions. Nevertheless, for similar root biomass,
422 the shoot biomass was the highest in plants supplied with the P+ treatment compared to the three other treatments.
423 We can hypothesize that stressed plants (P- conditions) maintained root development at the expense of the shoot
424 compartment. This is confirmed by the greater RMF observed under soluble P limitation. On the contrary, when
425 there was no nutrient limitation, there was no need for the plants to prioritize extension of their root systems and
426 the plants maintained the biomass accumulation into the above-ground compartment. These results are in
427 accordance with the “functional equilibrium model”, which states that a plant shifts allocation towards the organ
428 involved in the acquisition of the most limiting resources (Brouwer 1963), and reveal a true plasticity in response
429 to P supply. Contrasted results were found in previous studies about the allocation pattern in response to P nutrition.
430 Some of them concluded in a “conservative response” of the plants adjusting their size rather than their allocation
431 pattern (apparent plasticity; Müller et al. 2000). Others described an impact on the allocation pattern, but only
432 under severe P stress (Rubio et al. 2013) or in interaction with nitrogen fertilisation (Sims et al. 2012). Plasticity
433 of biomass allocation was also demonstrated, with a strong impact from the nutritional context (Poorter et al. 2012;
434 Poorter and Nagel 2000; Shipley and Meziane 2002).

435 Regarding the root system, plants exposed to the P-/HA treatment exhibited a greater TRL. The observed root
436 system lengthening was associated with greater root biomass and RMF for plants grown under P- conditions. These
437 results are consistent with those of a hydroponics experiment on several barley varieties, which revealed a general
438 trend towards root lengthening in response to P deficiency (Giles et al. 2017). On the other hand, Shen et al. (2018)
439 reported that under moderate P stress, wheat plants maintained root length and reduced root biomass whereas
440 under severe P stress both TRL and root biomass were reduced.

441 **4.2 Despite their ability to solubilize tricalcium phosphate and hydroxyapatite, the bacterial inoculants** 442 **did not alleviate P deficiency stress in *Brachypodium* under the experimental growing conditions**

443 All the selected bacteria were able to solubilize the poorly available forms of P (TCP and HA) in NBRIP medium
444 (Nautiyal 1999). HA, despite being reported as less soluble than TCP (Bashan et al. 2013, Havlin et al. 2014), was
445 as easily solubilized as TCP. Some acidification of the medium was observed, with the best solubilizer strains
446 acidifying the most. Medium acidification by proton release is the most straightforward P solubilization process
447 (Bashan et al. 2013) and numerous studies have reported an acidification-associated P solubilization (Collavino et
448 al. 2010; Fernández et al. 2012; Pereira and Castro 2014; Yu et al. 2011). The relationship between soluble P
449 concentration and pH variation tended towards an intensification of P solubilization activity as the pH variation
450 became stronger. This was more pronounced for HA than for TCP solubilization. This raises the hypothesis that
451 HA solubilization mechanisms other than acidification are involved, such as complexing or chelating reactions
452 (Bashan et al. 2013).

453 The use of PSB as bio-inoculants is increasingly reported in the literature, with interesting effects of microbial P
454 mobilization on plant development and yield (Bakhshandeh et al. 2015; Li et al. 2017; Oteino et al. 2015; Pereira
455 and Castro 2014), but few results have reported the inefficiency of *in vitro*-selected PSB to promote plant growth
456 in the presence of poorly soluble forms of P (Collavino et al. 2010; Yu et al. 2011). In our study, the biomass
457 accumulated in shoots and roots was reduced when plants were grown in the presence of bacteria. The strains
458 Pfl29ARP and BveFZB42 had the strongest impact on plant development. Despite their ability to solubilize TCP
459 and HA in NBRIP medium, the selected strains were not able to mobilize these poorly soluble forms of P under
460 co-cultivation conditions and by this way alleviate P-starvation stress in *Brachypodium*. The soluble P
461 concentration in the Hoagland solution at the end of the cultivation was below the detection limit of our analytical
462 method for the P-, P-/TCP and P-/HA treatments (data not shown). A slight acidification of the nutrient solution
463 was observed at the end of the co-cultivation in the presence of bacteria, but the pH remained within an acceptable
464 range for plant development (data not shown). The available carbon source is of great importance for P
465 solubilization by bacteria. Nico *et al.* (2012) reported a reduced P solubilization by bacteria unless glucose was
466 added to the growing medium of rice. Soil experiments also resulted in the absence of beneficial effects of
467 microbial consortia products on maize grown in a substrate with low organic matter content (Bradácová et al.
468 2019). Glucose is the most abundant sugar detected in *Brachypodium* exudates (Kawasaki et al. 2016). In our
469 study, the bacterial strains were tested for P solubilization with glucose as the sole C source in NBRIP medium
470 (Nautiyal 1999). During the co-cultivation experiment, the concentration of glucose provided through root
471 exudates may have been too low to sustain the bacteria solubilization activity. As our gnotobiotic co-cultivation
472 system was closed during the entire experiment duration, toxic bacterial metabolites may have accumulated in the
473 system, leading to a repression of plant growth. Some studies have revealed a deleterious impact of inoculation
474 with bacterial strains on plant growth under gnotobiotic conditions (Rybakova et al. 2016; Timmusk et al. 2015).
475 The efficacy of a system to test for PSB activity in the presence of a host plant appears to be highly dependent on
476 the considered organisms, but also on the co-cultivation conditions.

477 **4.3 The plastic response of *Brachypodium* to P deficiency was modulated by inoculation with P** 478 **solubilizing bacteria**

479 Regarding the biomass allocation pattern, inoculation with PSB revealed an alteration of the plant's response to P
480 conditions, except in the presence of HA. Under P- conditions, inoculation with PSB (except with BveFZB42) led
481 to a reduced shoot productivity for similar root biomass. The same observation was made under the P-/TCP
482 treatment, mainly with Pfl29ARP. The depletion in shoot growth benefited the root system, the development of
483 which was either unaffected or less impacted than the shoot. This resulted in an increase in RMF. Under the P+
484 treatment, investment into the root compartment was reduced in inoculated plants, except with AviF0819 and
485 BveGB03. The RMF was still increased for the same reason as before: a repression of shoot biomass but a steady
486 root biomass accumulation. As the root system is the place where the interaction with the bacteria occurs, it appears
487 that the plant modulated the development of this interface of interaction depending on the nutritional context.
488 These contrasted behaviours in *Brachypodium* should be explored more deeply. The complementarity between
489 biomass partitioning (RMF) and allometric trajectories appears clearly here for the analysis of biomass allocation
490 patterns under environmental variation. Both approaches should be considered when studying the impact of
491 biostimulants on plant biomass allocation in response to environmental constraints.

492 The total root length of *Brachypodium* was significantly impacted by the P supply and inoculation with PSB.
493 Regardless the P treatment, inoculation with BveFZB42, Eco99B829 and Pfl29ARP led to a reduction in TRL.
494 These results contrast with others reported in the literature. Indeed, Talboys et al. (2014) demonstrated a root
495 elongation promotion effect of BveFZB42 inoculation on wheat (through auxin production), in both low and high
496 P-level soils. In a soil experiment, *Pseudomonas fluorescens* strains also exhibited a positive impact on wheat root
497 elongation under contrasted P fertilisation (Zabihi et al. 2011). The persistent homology analysis performed in our
498 study revealed that inoculation with Pfl29ARP impacted the morphology of the plant root system (considering 1st
499 and 2nd order roots) in comparison with non-inoculated plants and plants inoculated with BveFZB42. The P
500 conditions also induced changes in root system morphology, but these were less easily characterized. According
501 to our results, *Brachypodium* showed a modification of root development, triggered by contrasted P supply and
502 inoculation with bacteria. Geometrical and topological aspects of the root system architecture are important for
503 nutrient foraging in soils. Both aspects are covered by the persistent homology analysis of the root system
504 morphology. Thus, the methodological approaches used in this study appear suitable in seeking to characterize the
505 plant's response to P supply and inoculation with PSB. This study did not consider root hairs, yet they constitute
506 an important strategy for P nutrition (Lynch 2011) and should be further investigated.

507 **4.4 Inoculation with P solubilizing bacteria did not improve *Brachypodium* P use efficiency under the** 508 **experimental growth conditions**

509 The shoot P concentration and PUE in *Brachypodium* were mainly affected by the P supply, but also by PSB
510 inoculation to some extent. The shoot P concentration was the lowest in plants grown under P- conditions,
511 confirming the P-deficient status of those plants. Despite the demonstrated ability of the bacterial strains to
512 solubilize TCP and HA, they did not alleviate P deficiency in the plants. The soluble P concentration in the
513 Hoagland solution at the end of the cultivation was null for the P-, P-/TCP and P-/HA treatments (data not shown).
514 This result reinforces the above-mentioned hypothesis that the PSB did not extensively solubilize TCP and HA in
515 our gnotobiotic conditions. On the other hand, the P+ solution contained enough soluble P after four weeks for
516 avoiding nutritional stress in the plants (data not shown). Considering the slightly higher shoot P concentration in
517 the presence of TCP and HA regardless the inoculation treatment, we assume that *Brachypodium* was able to partly
518 solubilize those poorly soluble forms of P. Indeed, plants are able to acidify the rhizosphere and release organic
519 anions, mobilizing poorly available P sources (Hinsinger et al. 2003; Wang and Lambers 2019). Citrate, malate,
520 succinate, fumarate and oxalate were detected in *Brachypodium* root exudates (Kawasaki et al. 2016) and may be
521 implied in P solubilization processes. The PUpE was significantly higher in plants exposed to soluble P deficiency
522 compared to plants grown under the P+ treatment, as the stressed plants took up all the available soluble P and
523 partly used it to build their shoots. The highest PUpE values were obtained in the presence of TCP and HA. This
524 observation is consistent with the higher shoot P concentration observed under these treatments and reinforces the
525 above-mentioned hypothesis of partial P solubilization by *Brachypodium*. The PUpE reduction in plants inoculated
526 with Pfl29ARP is consistent with the observed decrease in shoot biomass accumulation, which impairs their P
527 accumulation ability. The PUE was significantly higher under soluble P deficiency than under the P+ treatment,
528 with the highest efficiency under the P- treatment. Therefore, stressed plants produced the largest biomass per unit
529 of accumulated P. The inoculation of *Brachypodium* with bacteria did not impact the PUE, as expected from their
530 poor P solubilization activity during the co-cultivation experiment. As observed for PUpE and PUE, the PPUE
531 values were higher under soluble P deficiency, meaning that for similar shoot P concentration the stressed plants
532 produced more shoot biomass. The inoculation with Pfl29ARP induced a reduction in PPUE. Indeed, shoot P

533 concentration was similar in non-inoculated plants and in plants inoculated with Pfl29ARP, but shoot biomass
534 accumulation was reduced in inoculated plants.

535 **5 Conclusion**

536 The selected PSB efficiently solubilized TCP and HA in an *in vitro* liquid cultivation system. However, they did
537 not alleviate P deficiency in *Brachypodium* under gnotobiotic co-cultivation conditions. Some negative impact of
538 the PSB on plant biomass accumulation was even observed, probably due to inadequate carbon supply through
539 root exudates or to the accumulation of bacterial toxic metabolites in the system. *Brachypodium* showed
540 developmental plasticity in response to contrasted P conditions, prioritizing the development of the root
541 compartment upon P starvation. Despite their inability to alleviate P deficiency, the selected PSB modulated
542 *Brachypodium*'s response to P conditions by altering the plant allocation pattern and the root system development.
543 Nevertheless, this modulation did not improve PUE in *Brachypodium* under our experimental conditions. This
544 study highlights the necessity to select experimental conditions as close as possible to realistic conditions in the
545 perspective of screening PSB for the purpose of using them as plant inoculants. Co-cultivation experiments are
546 mandatory in order to confirm a beneficial interaction and test the related hypothesis. To our knowledge, this study
547 represents the first time that allometry and persistent homology analyses were used to assess the impact of
548 biostimulants on plant development under nutritional deficiency. They revealed to be convenient tools to study
549 potential plasticity in biomass allocation or change in root system morphology. The plasticity in biomass allocation
550 could be explored more deeply by considering a temporal perspective of the biomass allocation patterns; this would
551 allow the experiment to cover a broader range of plant sizes and clearly assess the interaction between the use of
552 biostimulants and varying nutrient supply. As root hairs are an important trait in nutrient acquisition, they deserve
553 consideration in addition to root system architecture parameters, providing a more precise insight into root system
554 plasticity in response to P supply and PSB inoculation. Integrating the proposed analyses and tools in future
555 research would provide a better understanding of the impact of biostimulants on plant plasticity in a changing
556 environment.

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562

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- 735

736 **Online Resource 1** Biomass accumulation and RMF of four-week-old *Brachypodium* plantlets grown in Magenta
 737 boxes, exposed to contrasted P supplies and either inoculated or not inoculated with bacterial strains (n=30 for the
 738 P- and P+ treatments, n=15 for the P-/TCP and P-/HA treatments). Results of 3-way ANOVAs (degree of freedom
 739 “df”, *P* and *F* values) and Dunnett’s *post hoc* tests (annotated with stars; P+ and non-inoculated treatments used
 740 as references)

	Shoot biomass (mg)		Root biomass (mg)		Total biomass (mg)		RMF	
	mean	sd	mean	Sd	mean	sd	mean	sd
<i>P treatment</i>								
P-	78.65 *	17.53	57.78 *	13.43	136.43 *	27.98	0.42	0.05
P-/HA	92.09 *	29.20	51.25	11.08	143.34 *	35.52	0.37	0.06
P-/TCP	89.56 *	23.15	51.59	11.75	141.25 *	31.30	0.37	0.06
P+	138.25	33.14	51.00	12.81	188.90	41.99	0.27	0.04
ANOVA	df=3, <i>P</i> <2.2e-16, <i>F</i> =188.9276		df=3, <i>P</i> =4.278e-07, <i>F</i> =11.1453		df=3, <i>P</i> <2.2e-16, <i>F</i> =84.2410		df= 3, <i>P</i> <2.2e-16, <i>F</i> =313.3280	
<i>Inoculation treatment</i>								
Non-inoculated	112.05	36.78	55.02	14.04	167.07	41.82	0.34	0.08
AviF0819	108.42	39.08	56.00	13.06	164.42	44.44	0.35	0.08
BveFZB42	100.46 *	32.72	49.42 *	12.10	149.88 *	39.84	0.34	0.06
BveGB03	107.84	38.83	54.49	12.45	162.33	42.21	0.35	0.09
Eco99B829	103.56	34.44	54.28	11.58	156.97	35.83	0.36	0.08
Pfl29ARP	83.48 *	32.72	51.33	13.27	134.81 *	39.88	0.39	0.08
ANOVA	df=5, <i>P</i> =8.956e-16, <i>F</i> =17.2350		df=5, <i>P</i> =0.002956, <i>F</i> =3.6519		df=5, <i>P</i> =3.476e-11, <i>F</i> =12.2019		df=5, <i>P</i> =2.356e-15, <i>F</i> =16.7725	
<i>Repetition</i>								
ANOVA	df=1, <i>P</i> =0.2585, <i>F</i> =1.2795		df=1, <i>P</i> =0.013298, <i>F</i> =6.1721		df=1, <i>P</i> =0.1091, <i>F</i> =2.5765		df=1, <i>P</i> =0.2085544, <i>F</i> =1.5855	
<i>Interaction</i>								
P- non-inoculated	88.84	15.04	60.01	11.97	148.86	22.02	0.40	0.06
P- AviF0819	83.13	15.56	61.05	12.06	144.18	25.07	0.42	0.04
P- BveFZB42	80.21	19.12	51.28	13.53	131.49	31.51	0.39	0.03
P- BveGB03	79.81	13.45	61.35	10.44	141.16	21.41	0.43 *	0.04
P- Eco99B829	75.62	14.65	59.99	14.23	135.61	26.00	0.44 *	0.05
P- Pfl29ARP	64.34	17.60	53.11	15.08	117.44	30.64	0.45 *	0.05
P-/HA non-inoculated	100.09	25.94	55.54	13.70	155.62	31.15	0.36	0.08
P-/HA AviF0819	83.17	23.63	50.40	13.22	133.57	32.78	0.38	0.06
P-/HA BveFZB42	87.99	26.38	48.61	10.86	136.61	32.44	0.36	0.06
P-/HA BveGB03	101.44	35.68	49.55	9.45	150.99	40.80	0.34	0.07
P-/HA Eco99B829	101.27	32.89	54.15	8.14	155.43	37.93	0.36	0.06
P-/HA Pfl29ARP	78.59	24.21	49.23	9.98	127.83	32.08	0.40	0.06
P-/TCP non-inoculated	94.26	26.26	51.83	11.63	146.09	36.12	0.36	0.04
P-/TCP AviF0819	96.39	19.21	53.53	11.80	149.92	27.51	0.36	0.06
P-/TCP BveFZB42	93.72	20.98	48.94	11.43	142.65	29.81	0.34	0.04
P-/TCP BveGB03	90.40	24.71	53.77	14.86	144.16	37.03	0.37	0.04
P-/TCP Eco99B829	97.48	17.70	49.70	7.81	148.43	23.06	0.34	0.04
P-/TCP Pfl29ARP	65.7	14.42	51.70	12.92	117.40	23.81	0.44 *	0.05
P+ non-inoculated	150.13	31.20	51.36	16.15	201.49	43.82	0.25	0.05
P+ AviF0819	152.36	30.56	54.98	13.41	207.34	40.21	0.26	0.04
P+ BveFZB42	130.30	30.35	48.21	11.87	178.51	40.32	0.27	0.03
P+ BveGB03	147.60	30.07	50.53	11.90	198.13	38.98	0.25	0.04
P+ Eco99B829	135.48	28.99	50.50	9.14	183.69	32.12	0.28	0.04
P+ Pfl29ARP	113.97	32.82	50.40	13.33	164.36	42.14	0.31 *	0.05
ANOVA	df=15, <i>P</i> =0.1269, <i>F</i> =1.4330		df=15, <i>P</i> =0.735267, <i>F</i> =0.7482		df=15, <i>P</i> =0.6874, <i>F</i> =0.7917		df=15, <i>P</i> =0.0004245, <i>F</i> =2.7517	

741

742 **Online Resource 2** Coefficients, R² and P value of SMA lines (n=30 for the P- and P+ treatments, n=15 for the
 743 P-/HA and P-/TCP treatments), results of covariance analysis for differences among SMA lines coefficients
 744 (degree of freedom “df”, P and likelihood ratio test “LR” values). If no significant difference was noticed between
 745 slopes, a common slope was used to test for difference in elevation. Treatments without any common letter are
 746 significantly different from each other (pairwise comparison)

	elevation	slope	R ²	P
<i>Non-inoculated</i>				
P-				
Non-inoculated	0.51	0.81 <i>ab</i>	0.090	0.10762
AviF0819	-0.08	1.19 <i>a</i>	0.072	0.33277
BveFZB42	0.00	1.15 <i>a</i>	0.668	0.00019
BveGB03	1.15	0.60 <i>b</i>	0.399	0.00018
Eco99B829				
Pfl29ARP				
Covariance analysis	/	df= 3, P=0.020606,		
<i>P-</i>				
Non-inoculated	0.38 <i>ab</i>		0.090	0.10762
AviF0819	0.34 <i>bc</i>		0.468	3.0804e-5
BveFZB42	0.39 <i>a</i>	0.88	0.718	3.5339e-9
BveGB03	0.32 <i>cd</i>		0.460	5.3114e-5
Eco99B829	0.31 <i>cd</i>		0.411	0.00013
Pfl29ARP	0.28 <i>d</i>		0.604	4.3258e-7
Covariance analysis	df=5, P=2.4697e-9, LR=48.77	df=5, P=0.75661, LR=2.631		
<i>P-/HA</i>				
Non-inoculated	-0.71		0.072	0.33277
AviF0819	-0.73		0.394	0.01228
BveFZB42	-0.69	1.56	0.264	0.05028
BveGB03	-0.65		0.227	0.07242
Eco99B829	-0.71		0.384	0.01371
Pfl29ARP	-0.75		0.582	0.00093
Covariance analysis	df=5, P=0.43264, LR=4.865	df=5, P=0.17355, LR=7.7		
<i>P-/TCP</i>				
Non-inoculated	0.23 <i>ab</i>		0.668	0.00019
AviF0819	0.23 <i>ab</i>		0.249	0.05839
BveFZB42	0.25 <i>ab</i>	1.02	0.513	0.00269
BveGB03	0.19 <i>b</i>		0.630	0.00069
Eco99B829	0.27 <i>a</i>		0.339	0.03689
Pfl29ARP	0.08 <i>c</i>		0.419	0.00905
Covariance analysis	df=5, P=4.4342e-7, LR=37.65	df=5, P=0.83515, LR=2.1		
<i>P+</i>				
Non-inoculated	1.15	0.60 <i>c</i>	0.399	0.00018
AviF0819	0.93	0.72 <i>bc</i>	0.312	0.00132
BveFZB42	0.45	0.99 <i>ab</i>	0.621	2.3175e-7
BveGB03	0.79	0.81 <i>bc</i>	0.512	1.2825e-5
Eco99B829	0.17	1.15 <i>a</i>	0.312	0.00165
Pfl29ARP	0.10	1.15 <i>a</i>	0.529	5.3439e-6
Covariance analysis	/	df=5, P=0.00842, LR=15.5		

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749 **Online Resource 3** TRL of four-week-old *Brachypodium* plantlets grown in Magenta boxes, exposed to
 750 contrasted P supplies and either inoculated or not inoculated with bacterial strains (n=18 for the P- and P+
 751 treatments, n=9 for the P-/HA and P-/TCP treatments). Results of 3-way ANOVAs (degree of freedom “df”, *P*
 752 and *F* values) and Dunnett’s *post hoc* tests (annotated with stars; P+ and non-inoculated treatments used as
 753 references)

	TRL (cm)	
	mean	sd
<i>P treatment</i>		
P-	224.58	42.78
P-/HA	233.06*	42.37
P-/TCP	213.26	49.83
P+	213.87	46.39
ANOVA	df=3, <i>P</i> =0.02126, <i>F</i> =3.2821	
<i>Inoculation treatment</i>		
Non-inoculated	237.85	48.83
AviF0819	228.11	42.30
BveFZB42	214.92*	50.62
BveGB03	233.41	40.92
Eco99B829	213.11*	35.64
Pfl29ARP	195.82*	41.59
ANOVA	df=5, <i>P</i> =1.99e-06, <i>F</i> =7.2513	
<i>Repetition</i>		
ANOVA	df=1, <i>P</i> =0.07758, <i>F</i> =3.1363	
<i>Interaction</i>		
P- non-inoculated	252.83	40.23
P- AviF0819	239.12	33.89
P- BveFZB42	214.76	49.38
P- BveGB03	234.62	31.18
P- Eco99B829	218.86	26.65
P- Pfl29ARP	187.32	42.97
P-/HA non-inoculated	257.96	31.15
P-/HA AviF0819	219.12	39.79
P-/HA BveFZB42	231.37	40.44
P-/HA BveGB03	235.36	49.72
P-/HA Eco99B829	248.26	45.64
P-/HA Pfl29ARP	206.27	33.97
P-/TCP non-inoculated	198.82	76.65
P-/TCP AviF0819	221.07	38.39
P-/TCP BveFZB42	224.41	61.96
P-/TCP BveGB03	222.32	51.99
P-/TCP Eco99B829	207.66	28.38
P-/TCP Pfl29ARP	205.30	33.29
P+ non-inoculated	232.32	36.08
P+ AviF0819	225.11	52.78
P+ BveFZB42	202.13	51.09
P+ BveGB03	236.76	41.63
P+ Eco99B829	192.52	27.13
P+ Pfl29ARP	194.36	47.95
ANOVA	df=15, <i>P</i> =0.15092, <i>F</i> =1.3891	

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756 **Online Resource 4** Results of PERMANOVA performed on the persistent homology analysis output of plant
757 root systems. n=18 for the P- and P+ treatments, n=9 for the P-/HA and P-/TCP treatments. Post-hoc tests were
758 performed by running a PERMANOVA for each pairwise comparison and *P* values were adjusted for multiple
759 comparisons using the Benferroni method.

760

	Df	<i>F</i> model	<i>P</i>
Inoculation treatment	2	11.1650	0.000999
P treatment	3	2.8237	0.005994
Interaction	6	1.2461	0.217782
Residuals	150		

761

762 Post-hoc tests:

	<i>F</i> model	<i>P</i>
P- vs P-/TCP	1.8911	0.68931
P- vs P+	2.1396	0.60539
P- vs P-/HA	2.5906	0.28771
P-/TCP vs P+	3.1768	0.19780
P-/TCP vs P-/HA	1.4693	1.00000
P+ vs P-/HA	3.5882	0.17982
Pfl29ARP vs non-inoculated	18.7287	0.00099
Pfl29ARP vs BveFZB42	13.9209	0.00099
Non-inoculated vs BveFZB42	1.0035	0.38462

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765 **Online Resource 5** Shoot P concentration and PUE parameters of four-week-old *Brachypodium* plantlets grown
 766 in Magenta boxes, exposed to contrasted P supplies and either inoculated or not inoculated with bacterial strains
 767 (n=6 for the P- and P+ treatments, n=3 for the P-/HA and P-/TCP treatments). Results of 3-way ANOVAs
 768 (degree of freedom “df”, *P* and *F* values) and Dunnett’s *post hoc* tests (annotated with stars; P+ and non-
 769 inoculated treatments used as references)

	Shoot P concentration ($\mu\text{g P/mg FW}$)		PU _p E		PU _t E		PPUE	
	mean	sd	mean	sd	Mean	Sd	mean	sd
<i>P treatment</i>								
P-	0.237*	0.032	2418.692*	662.318	4.296*	0.603	1664.549*	294.805
P-/HA	0.335*	0.064	4146.142*	1768.202	3.099*	0.626	1365.267*	135.433
P-/TCP	0.281*	0.092	3318.446*	1565.915	3.940*	1.267	1649.241*	361.100
P+	0.764	0.086	337.427	60.022	1.325	0.151	915.617	222.696
ANOVA	df=3, $P<2\text{e-}16$, $F=292.0433$		df=3, $P<2\text{e-}16$, $F=306.9495$		df=3, $P<2\text{e-}16$, $F=217.6723$		df=3, $P<2.2\text{e-}16$, $F=60.9819$	
<i>Inoculation treatment</i>								
Non-inoculated	0.432	0.233	2444.236	2012.003	3.026	1.488	1501.401	467.108
AviF0819	0.421	0.243	2110.551	1542.716	3.136	1.493	1482.431	419.206
BveFZB42	0.457	0.260	2255.018	1662.922	2.886	1.410	1304.184	436.268
BveGB03	0.440	0.248	2247.915	1942.801	2.997	1.443	1354.915	360.533
Eco99B829	0.457	0.261	2350.842	2048.284	2.937	1.485	1329.075	453.353
Pfl29ARP	0.412	0.256	1568.265*	1258.735	3.298	1.543	1202.833*	381.678
ANOVA	df=5, $P=0.2825$, $F=1.2751$		df=5, $P=0.01069$, $F=3.2079$		df=5, $P=0.4357$, $F=0.9788$		df=5, $P=0.007738$, $F=3.3921$	
<i>Repetition</i>								
ANOVA	df=1, $P=0.6384$, $F=0.2225$		df=1, $P=0.83696$, $F=0.0426$		df=1, $P=0.4312$, $F=0.6257$		df=1, $P=0.648586$, $F=0.2092$	
<i>Interaction</i>								
P- non-inoculated	0.234	0.031	2704.936	682.528	4.350	0.608	1913.742	245.850
P- AviF0819	0.228	0.037	2476.670	739.706	4.487	0.686	1841.606	221.698
P- BveFZB42	0.255	0.037	2664.388	746.187	4.009	0.710	1588.721	289.616
P- BveGB03	0.239	0.040	2412.522	692.725	4.284	0.701	1631.159	186.833
P- Eco99B829	0.232	0.033	2275.638	540.169	4.378	0.607	1649.714	296.266
P- Pfl29ARP	0.236	0.022	1977.999	544.750	4.268	0.430	1362.351	248.987
P-/HA non-inoculated	0.357	0.079	4775.731	2043.682	2.896	0.664	1395.929	126.322
P-/HA AviF0819	0.305	0.044	3327.955	1067.802	3.323	0.456	1361.300	58.025
P-/HA BveFZB42	0.344	0.046	3942.647	989.521	2.944	0.416	1283.964	129.002
P-/HA BveGB03	0.363	0.076	4969.660	2542.434	2.836	0.608	1374.750	154.550
P-/HA Eco99B829	0.343	0.069	4684.583	2454.629	2.998	0.602	1449.523	212.164
P-/HA Pfl29ARP	0.295	0.089	3176.275	1807.334	3.594	1.055	1326.138	163.137
P-/TCP non-inoculated	0.296	0.117	3770.126	2229.912	3.826	1.715	1722.268	566.133
P-/TCP AviF0819	0.285	0.089	3654.221	1536.643	3.792	1.388	1765.498	368.753
P-/TCP BveFZB42	0.286	0.095	3586.993	1625.492	3.835	1.530	1714.052	354.350
P-/TCP BveGB03	0.266	0.067	2989.873	1264.261	3.944	1.112	1606.530	157.781
P-/TCP Eco99B829	0.351	0.141	4186.080	1884.872	3.296	1.666	1460.423	611.666
P-/TCP Pfl29ARP	0.203	0.020	1723.384	157.770	4.947	0.453	1626.672	188.967
P+ non-inoculated	0.736	0.062	354.844	41.169	1.368	0.114	1031.362	204.255
P+ AviF0819	0.741	0.088	363.894	57.738	1.365	0.163	1042.288	220.108
P+ BveFZB42	0.802	0.086	335.845	48.824	1.260	0.138	824.824	180.071
P+ BveGB03	0.765	0.078	351.455	70.629	1.318	0.129	942.947	232.530
P+ Eco99B829	0.791	0.111	341.557	38.802	1.286	0.186	882.537	248.321
P+ Pfl29ARP	0.751	0.102	276.967	73.901	1.354	0.193	769.744	189.371
ANOVA	df=15, $P=0.5820$, $F=0.8858$		df=15, $P=0.72824$, $F=0.7489$		df=15, $P=0.9355$, $F=0.4976$		df=15, $P=0.937471$, $F=0.4938$	

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