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Root morphology, yield and must composition of grapevine subjected to application of urea methods and rates

This is the final peer-reviewed author's accepted manuscript (postprint) of the following publication:

*Published Version:*

Severo de Souza Kulmann, M., Stefanello, L.O., Schwalbert, R., Squizani Arruda, W., Sans, G.A., Fogliarini Parcianello, C., et al. (2023). Root morphology, yield and must composition of grapevine subjected to application of urea methods and rates. JOURNAL OF PLANT NUTRITION, 46(13), 2959-2976 [10.1080/01904167.2022.2160762].

*Availability:*

This version is available at: <https://hdl.handle.net/11585/912975> since: 2024-04-18

*Published:*

DOI: <http://doi.org/10.1080/01904167.2022.2160762>

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1 **Root morphology, yield and must composition of grapevine subjected**  
2 **to application of urea methods and rates**

3

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17

18 **Abstract**

19 Nitrogen (N) is often applied to soil surface during the grapevine cycle, increases N  
20 losses. One way to reduce N losses lies on its application method, however how N dose  
21 and application mode impact on grapevines remain poorly investigated. The aim of this  
22 study is to evaluate the root growth, as well as grape must yield and composition, in a  
23 sandy soil and subjected to different N application doses and modes under subtropical  
24 climate conditions. Grapevines were implanted in 2011 and, since 2014, they were  
25 subjected to surface (N-Surf), and fertigation (N-Fert) N application, at rates of 0, 40

26 and 100 kg N ha<sup>-1</sup> year<sup>-1</sup>. Leaf N concentration, yield, must quality parameters and root  
27 system morphology were evaluated in the 2016/2017 and 2017/2018 seasons. The  
28 application of N in both rates (40 and 100 kg N ha<sup>-1</sup> year<sup>-1</sup>), in the N-Surf and N-Fert  
29 methods, increased the surface area of root, number of living roots and root production.  
30 The application of N rates, in both N application modes, provided adequate N supply to  
31 the soil and increased root system development, which contributed to the increase of  
32 soil N uptake by plants, higher N concentration in leaves and greater grape yield.  
33 However, N supply decreased the quality of the must, as observed in decreased TSS and  
34 total anthocyanins values and in increased total titratable acidity values, which are not  
35 desirable for red wine making processes.

36

37 **Keywords:** Nitrogen fertilization; Sandy soil; Grapes quality; Root system;  
38 Minirhizotron.

39

## 40 **Introduction**

41 Grapevines are grown worldwide in sandy soils that often have low organic  
42 matter (OM) content, a fact that leads to low nitrogen (N) availability in the soil  
43 (Gustavo Brunetto et al., 2016, 2014; Lorensini et al., 2017). Thus, applications of N  
44 fertilizer are necessary to increase soil mineral N availability, *i.e.* nitrate (NO<sub>3</sub><sup>-</sup>) and  
45 ammonium (NH<sub>4</sub><sup>+</sup>) forms, which are the ones uptake by plants. Soil surface (without  
46 incorporation) N applications are often recommended, however this approach may  
47 increase N losses, mainly due to volatilization (Lorensini et al., 2012; Viero, Bayer,  
48 Mara Vieira Fontoura, Paulo de Moraes, & Professor, 2014), and decrease grapevines N  
49 uptake (Gustavo Brunetto et al., 2016; Gustavo Brunetto, Kaminski, Melo, Brunning, &  
50 Mallmann, 2006). Therefore, it is necessary adopting alternative ways of supplying N to

51 decrease losses and increase N use efficiency. Fertigation is one of these alternatives; it  
52 enables urea solubilization and hydrolysis reaction, decreases losses caused by  
53 volatilization, since it favors the migration of  $\text{NO}_3^-$  and  $\text{NH}_4^+$  in soil profile (Comas,  
54 Bauerle, & Eissenstat, 2010).

55 Increased mineral N contents in soil subsurface layers can contribute plant root  
56 system growth and development (Centinari, Heuvel, Goebel, Smith, & Bauerle, 2016;  
57 Comas et al., 2010; Zhou, Yang, Ren, Huang, & An, 2014). Increased  $\text{NO}_3^-$  and  $\text{NH}_4^+$   
58 concentrations in soil solution activates  $\text{NO}_3^-$  (NRTs) and  $\text{NH}_4^+$  (AMTs) transporters  
59 and, consequently, contributes for N uptake (Pii, Aldrighetti, Valentinuzzi, Mimmo, &  
60 Cesco, 2019; Tomasi, Monte, Varanini, Cesco, & Pinton, 2015; Xu, Fan, & Miller,  
61 2012) and for the increase of grapevine root emission and root surface area (Krouk et  
62 al., 2010; Remans et al., 2006). In addition, high  $\text{NO}_3^-$  concentrations can inhibit auxin  
63 transport in the root system due to competition at the NRT1.1 transporter site; thus,  
64 auxin accumulates in plant roots and leads to an increase of lateral root growth in sites  
65 presenting high  $\text{NO}_3^-$  concentrations (Mounier, Pervent, Ljung, Gojon, & Nacry, 2014).  
66 On the other hand, high N doses can decrease soil pH due to  $\text{NH}_4^+$  nitrification into  
67  $\text{NO}_3^-$ . This phenomenon, which is often seen in vineyards (Lorensini et al., 2017), can  
68 solubilize minerals in the soil and release aluminum ( $\text{Al}^{+3}$ ) into soil solution;  
69 consequently,  $\text{Al}^{+3}$  can be uptake by plant roots and lead to root length shortening and to  
70 root diameter increase (Inostroza-Blancheteau, Aquea, Reyes-Díaz, Alberdi, & Arce-  
71 Johnson, 2011; Riaz et al., 2018; Zhou et al., 2014). This outcome is undesirable  
72 because it reduces the volume of soil explored by the root system, as well as the  
73 absorption of water and nutrients (Riaz et al., 2018).

74 Mineral N provided by N fertilization can be uptake and transported to plant  
75 shoot, increasing N content in the leaves. This process occurs mainly in stages of

76 intense root growth, such as at flowering, when the increased root volume contributes to  
77 great N uptake (Radville et al., 2016). Increased leaf N content can lead to an increased  
78 photosynthetic rate and CO<sub>2</sub> fixation, since RuBisCo corresponds to 50% of soluble  
79 proteins in the leaves of C3 plants (Stefano, Agyei, Njoku, & Udenigwe, 2018). This  
80 outcome favors cell division and elongation conditions in plant tissues and increases  
81 vegetative growth (Blank, Tittmann, Ghozlen, & Stoll, 2018; Moriwaki et al., 2019) in  
82 grape clusters, resulting in greater crop yield (Gustavo Brunetto et al., 2009a; Yu et al.,  
83 2012).

84 In addition to increased crop yield, there may be changes in oenological  
85 parameters associated with wine quality; these changes may be due to total  
86 anthocyanins and total soluble solids (TSS) dilution, as well as to increase in total  
87 titratable acidity values (Gustavo Brunetto et al., 2009a; Janjanin et al., 2016; Kelly et  
88 al., 2017; Spayd, Nagel, & Edwards, 1995). Thus, it is recommended adopting  
89 strategies, such as N doses and application modes, to stimulate root system growth in  
90 order to increase water and nutrient absorption by plants, as well as to increase crop  
91 yield without decreasing the quality of must. The aim of the current study was to  
92 investigate whether N rates and modes of application can change root system  
93 morphology, yield and the composition of grape must in grapevines (*Vitis vinifera* L.)  
94 grown in sandy soil under subtropical climate conditions.

95

## 96 **Materials and Methods**

### 97 **Experimental site description**

98 The experiment was conducted in a vineyard established in 2011, in Santana do  
99 Livramento County, Rio Grande do Sul State (RS), Southern Brazil (30° 48' 31" S; 55°  
100 22' 33" W). Alicante Bouschet (*Vitis vinifera* L. grafted onto Paulsen 1103 rootstock)

101 was the adopted cultivar. Plant density was 2,976 hectare<sup>-1</sup> (2.8 m between rows × 1.2  
102 m between plants), based on the espalier system. Soil in the experimental site was  
103 classified as Typic Hapludalf (Soil Survey Staff, 2014); the main chemical properties of  
104 the 0.0-0.20 m layer before the experiment are shown in Table 1.

105         Based on Köppen classification, the climate in the region is subtropical humid  
106 (Cfa); it is featured by mild temperatures and rainfall rates, with little variation  
107 throughout the year. Mean annual rainfall is 1,600 mm, whereas mean temperature is  
108 23.8° C in the warmest month (January) and 12.4° C in the coldest month (July)  
109 (Alvares, Stape, Sentelhas, De Moraes Gonçalves, & Sparovek, 2013). Annual  
110 insolation reaches approximately 2,500 hours; the site presents gently undulating  
111 terrain. Two crop production cycles, 2016/17 and 2017/18, were evaluated. According  
112 to data from the National Institute of Meteorology (INMET), the experimental area  
113 recorded 513.2 mm and 189.6 mm of rainfall from November 2016 to February 2017  
114 and from November 2017 to February 2018 (period between treatment application and  
115 harvest), respectively (Fig. 1).

116         Weeds in the grapevine row were desiccated with two to four sequential  
117 applications of 2 L ha<sup>-1</sup> y<sup>-1</sup> of non-selective systemic herbicide (*i.e.*, glyphosate),  
118 depending on cover plant development. Inter-row cover plants were cut near the soil  
119 surface, five times throughout each crop season and the residues were deposited on soil  
120 surface between rows. Grapevines were subjected to four applications of 45 kg P<sub>2</sub>O<sub>5</sub> ha<sup>-1</sup>  
121 year<sup>-1</sup> and 45 kg K<sub>2</sub>O ha<sup>-1</sup> year<sup>-1</sup> prior to experiment installation (CQFS-RS/SC, 2016).  
122 Triple superphosphate (42% P) and KCl (60% K) were used as P and K sources,  
123 respectively. Nitrogen fertilizers were applied to the crop based on each treatment.

124

125

## 126 **Experimental design and treatments**

127 Starting from September 2014, two N application methods: surface (N-Surf) and  
128 fertigation (N-Fert) and three rates 0, 40 and 100 kg N ha<sup>-1</sup> year<sup>-1</sup> were evaluated. The  
129 experiment followed a complete randomized block experimental design with three  
130 repetitions (plot) with 2 x 3 factorial arrangement (*i.e.*, two N application modes and  
131 three N rates). Each plot consisted of five plants were only the three central plants were  
132 used for data collection. In both application methods, urea was employed as N source  
133 (45% N); in N-Surf treatment, N was applied on the surface of the soil, within plant  
134 canopy projection, without incorporation. Every year, N application was performed  
135 during flowering (Gustavo Brunetto et al., 2009a). In N-Fert treatment, N was applied  
136 via drip irrigation (Netafim™ 132 Dripnet PC AS 16250), at a flow rate of 1.6 L h<sup>-1</sup>;  
137 drippers were spaced 0.6 m a part along the tree row, within canopy projection. The  
138 applications were performed at four different times, at 7-day intervals, according to the  
139 following procedure: 10 minutes of water; 10 minutes of water + N; and 10 minutes of  
140 water. Applications N-Fert provided 6.4 L of water per plant per crop season.

141

## 142 **Minirhizotron installation**

143 Minirhizotron tubes (Supplementary material 1) were installed in the ground of  
144 four plants per treatment in May 2016 to enable scanning the grapevine root system *in*  
145 *situ*. For each assessed grapevine plant, a hole was made with a spiral drill (70 mm  
146 diameter and 1,500 mm length), pulled by motorized ground driller (Branco, Bps 52),  
147 0.5 m away from the trunk, inside the canopy projection. Each individual hole was  
148 made at an angle of 45° to soil surface and a clear acrylic tube (70 mm x 1050 mm) was  
149 inserted. Three images were taken along each tube, referring to the soil layers: 0-20;  
150 40; and 40-60 cm deep. Each image had fixed dimensions of 216 mm × 196 mm. A

151 flexible, lightweight, waterproof and non-toxic plastic polyethylene rod (CID Bio-  
152 Science, Inc. Camas, WA, USA) internally protected the tubes. PVC pipes (100 mm  
153 diameter) were used to protect the external part of the tubes from damage caused by  
154 weather, handling or pest attacks. Soil disturbance during the hole drilling process made  
155 it necessary waiting eight months for soil-plant-tube system stabilization, before the  
156 beginning of the scanning activities.

157

### 158 **Image collection and measurements of root morphological parameters**

159 Root images were taken at grapevine flowering stage (in October) in the 2016/17  
160 and 2017/18 crop seasons. This phenological stage was selected because it coincides  
161 with a period of intense root growth and nutrient absorption, as reported in several  
162 studies (Comas, Anderson, Dunst, Lakso, & Eissenstat, 2005; Radville et al., 2016).  
163 Images of the root system were generated *in situ* with a CI-600™ cylindrical scanner  
164 (CID Bio-Science, Inc. Camas, WA, USA).

165 Subsequently, images were captured at different soil layers (0-20, 20-40 and 40-  
166 60 cm deep) and individually analyzed. The CI-600 scanner was connected to a  
167 computer by USB cable. Images captured in TIFF format (600 dpi resolution) were  
168 analyzed in the RootSnap!® CI-690 software version 1.3.2.25 (CID Bio-Science, Inc.  
169 Camas, WA, USA), which enables seeing the fine roots ( $\leq 2$  mm). Parameters evaluated  
170 in each tube based on generated images were (i) root surface area ( $\text{mm}^2 \text{tube}^{-1}$ ); (ii)  
171 mean root diameter ( $\text{mm tube}^{-1}$ ); (iii) number of living roots; (iv) number of dead roots;  
172 and (v) root production as the difference between the total length of living and dead  
173 roots ( $\text{mm tube}^{-1}$ ).

174

### 175 **Leaf collection for nitrogen analysis**



176 Six fully expanded leaves located in the opposite direction of the grape clusters,  
177 in the middle third of the annual branches, were collected at flowering (October 2016  
178 and 2017) and at berry veraison (December 2016 and 2017). According to CQFS-  
179 RS/SC (2016), these leaves are the diagnostic organ used to assess the nutritional status  
180 of grapevines. Leaves were washed in distilled water and dried in forced-air-circulation  
181 oven at 65 °C, until they reached constant weight. Then, they were milled in Wiley mill  
182 and sieved through 2-mm mesh. Subsequently, the leaf tissue was prepared and  
183 subjected to sulfuric digestion (Tedesco, Gianello, Bissani, Bohnen, & Volkweiss,  
184 1995). Total N concentration was determined in Kjeldahl semi-micro steam distiller  
185 (Tecnal, TE-0363, Brazil).

186

### 187 **Grape yield**

188 The number of grape clusters per plant was counted in the 2016/17 and 2017/18  
189 grape harvests. All grape clusters were harvested and weighed in a portable electronic  
190 scale (WalMur, 50 K, Brazil), to measure grape yield ( $\text{Mg ha}^{-1}$ ). Berries were collected  
191 at the upper, middle and lower third of the grape clusters, thus totaling 100 berries per  
192 repetition and 9 repetitions per treatment. They were weighed in scale (Bel Engineering,  
193 Precision Scale L, Brazil) to determine the weight of 100 berries, before storing for  
194 chemical analysis (in triplicate) of the must.

195

### 196 **Chemical analysis of the must**

197 The stored berries were peeled and only the pulp and seeds were analyzed. A  
198 benchtop refractometer (Cosmo, Type K - 32) was used to determine the total soluble  
199 solids (TSS) based on standards set by AOAC (2005). Total titratable acidity was  
200 determined by transferring 10 mL of extract to a 100 mL volumetric flask, whose

201 volume was completed with distilled water. For the titration of the resulting solution 0.1  
 202 mol L<sup>-1</sup> of sodium hydroxide (NaOH) was used (IAL, 2008).

203 Berry peel was used to determine total anthocyanin contents through a contact  
 204 with a 70:30 acidified (1% HCl) ethanolic solution (Ju and Howard 2003) at  
 205 peel/solution ratio of 1:3 (m/v) and crushed in a blender (Arno, Clic Lav Top, Brazil) at  
 206 low (1500 RPM) and high speed (3500 RPM) for 40 seconds each. The solution  
 207 resulting from the aforementioned extraction was placed in a 250 mL beaker and left to  
 208 rest for 30 minutes at room temperature (20 ± 1 °C) before it was centrifuged at 3500  
 209 RPM for 5 minutes. Extract supernatant was removed and stored in amber bottle, which  
 210 was placed in refrigerator (4-10°C). Subsequently, triplicate readings of these extracts  
 211 were performed.

212 Total anthocyanin content was determined based on the pH difference method  
 213 (Giusti and Wrolstad 2001), according to which samples were dissolved in two buffer  
 214 systems: 0.025 mol L<sup>-1</sup> potassium chloride at pH 1.0 and 0.4 mol L<sup>-1</sup> sodium acetate at  
 215 pH 4.5. Maximum absorption readings were taken at 515 nm and 700 nm, with the aid  
 216 of a UV-visible spectrophotometer (FEMTO, 600 plus, Brazil). Absorbance was  
 217 calculated through equation (1):

$$218 \quad A = (A_{515vis} - A_{700nm})_{pH1.0} - (A_{515vis} - A_{700nm})_{pH4.5} \quad (1)$$

219 Pigment concentration in the extract was calculated and represented in cyanidine-3-  
 220 glycoside through equation (2).

$$221 \quad \text{Anthocyanins (mg 100 mL}^{-1}\text{)} = (A \times MM \times DF \times 10000) / (\varepsilon \times 1) \quad (2)$$

222 wherein: A = absorbance; MM = molecular mass; DF = dilution factor; and ε = molar  
 223 absorptivity.

224

225

## 226 **Statistical analysis**

227 Data were subjected to analysis of variance by the R Studio software (R Core  
228 Team, 2019); whenever the analysis of variance showed a significant effect, means  
229 were compared through Scott-Knott test, at 5% significance level ( $P < 0.05$ ). In  
230 addition, data were subjected to multivariate principal component analysis (PCA) in the  
231 R software (R Core Team, 2019) in order to investigate correlation effects between  
232 response variables and treatments' distribution along the evaluated crop seasons.

233

## 234 **Results**

### 235 **Root morphological parameters**

236 In the 2016/17 crop season, the highest root surface areas, recorded in the 0-20  
237 cm layer, were observed in grapevines subjected to the application of 40 kg N ha<sup>-1</sup> year<sup>-1</sup>  
238 <sup>1</sup> as N-Surf and of 100 kg N ha<sup>-1</sup> year<sup>-1</sup> as N-Fert (Fig. 2a). As for the 2017/18 crop  
239 season, the highest root surface areas were observed in grapevines subjected to the  
240 application of 40 kg N ha<sup>-1</sup> year<sup>-1</sup> as N-Surf and of 100 kg N ha<sup>-1</sup> year<sup>-1</sup> as N-Surf and  
241 N-Fert (Fig. 2b). In the 2016/17 crop season, the highest root surface area, recorded in  
242 the 20-40 cm and 40-60 cm layers was observed in grapevines subjected to the  
243 application of 40 kg N ha<sup>-1</sup> year<sup>-1</sup> as N-Surf (Fig. 2a); whereas in the 2017/18 crop  
244 season, the highest values recorded were observed in grapevines subjected to  
245 application of 100 kg N ha<sup>-1</sup> year<sup>-1</sup> and 40 kg N ha<sup>-1</sup> year<sup>-1</sup> as N-Fert and N-Surf,  
246 respectively (Fig. 2b).

247 In the 2016/17 crop season, the highest mean root diameters, recorded in the 0-  
248 20 cm layer were observed at rate of 0 kg N ha<sup>-1</sup> year<sup>-1</sup> and 100 kg N ha<sup>-1</sup> year<sup>-1</sup>, as N-  
249 Surf and N-Fert, respectively (Fig. 2c). On the other hand, the highest root diameters,  
250 recorded in the 20-40 cm and 40-60 cm layers in the 2016/17 crop season, were

251 observed in grapevines subjected to all N rates (0, 40 and 100 kg N ha<sup>-1</sup> year<sup>-1</sup>) applied  
252 as N-Surf. In the 2017/18 crop season, the highest root diameters in the three soil layers  
253 were observed in plants subjected to 40 kg N ha<sup>-1</sup> year<sup>-1</sup> in the N-Surf mode and to 100  
254 kg N ha<sup>-1</sup> year<sup>-1</sup> as N-Fert mode (Fig. 2d).

255

### 256 **Root production and mortality**

257 In 2016/17 and 2017/18 crop seasons, the highest number of living roots,  
258 recorded in the 0-20 cm layer, were observed in plants subjected to 40 kg N ha<sup>-1</sup> year<sup>-1</sup>  
259 as N-Surf and to 100 kg N ha<sup>-1</sup> year<sup>-1</sup> as N-Fert mode (Fig. 3a, b). In the 2016/17 crop  
260 season, the highest number of living roots, recorded in the 20-40 cm and 40-60 cm  
261 layers, were observed in grapevines subjected to 40 kg N ha<sup>-1</sup> year<sup>-1</sup> as N-Surf and N-  
262 Fert modes (Fig. 3a). In the 2017/18 crop season, the highest number of living roots,  
263 recorded in the 20-40 cm layer were observed in grapevines subjected to 100 kg N ha<sup>-1</sup>  
264 year<sup>-1</sup> as N-Surf and N-Fert (Fig. 3b). In 40-60 cm layer, the highest number of living  
265 roots was in the 100 kg N ha<sup>-1</sup> year<sup>-1</sup> as N-Fert (Fig. 3b).

266 In the 2016/17 crop season, the highest number of dead roots, recorded in the 0-  
267 20 cm layer were observed in grapevines subjected to 40 kg N ha<sup>-1</sup> year<sup>-1</sup> as N-Surf  
268 mode and to 100 kg N ha<sup>-1</sup> year<sup>-1</sup> as N-Fert mode (Fig. 3c). In the 2017/18 crop season,  
269 the highest number of dead roots, recorded in the 0-20 cm layer, were observed in  
270 grapevines subjected to 100 kg N ha<sup>-1</sup> year<sup>-1</sup> as N-Fert and to 0 kg N ha<sup>-1</sup> year<sup>-1</sup> and 40  
271 kg N ha<sup>-1</sup> year<sup>-1</sup> as N-Surf (Fig. 3d). In the 2016/17 crop season, the highest number of  
272 dead roots, recorded in the 20-40 cm layer were observed in plants subjected to 40 kg N  
273 ha<sup>-1</sup> year<sup>-1</sup> as N-Surf and N-Fert and to 0 kg N ha<sup>-1</sup> year<sup>-1</sup> and 100 kg N ha<sup>-1</sup> year<sup>-1</sup> as N-  
274 Surf mode (Fig. 3c). In the 2017/18 crop season, the highest number of dead roots,  
275 recorded in the 20-40 cm layer, were observed in grapevines grown in control soil (0 kg

276 N ha<sup>-1</sup> year<sup>-1</sup>) N-Fert and in soil subjected to the application of 40 kg N ha<sup>-1</sup> year<sup>-1</sup> as N-  
277 Surf mode (Fig. 3d). In the 2016/17 crop season, the largest total number of dead roots  
278 recorded in the 40-60 cm layer, was observed in grapevines grown in the control soil N-  
279 Fert and N-Surf (Fig. 3c). In the 2017/18 crop season, the highest number of dead roots  
280 recorded were observed in grapevines subjected to 100 kg N ha<sup>-1</sup> year<sup>-1</sup> as N-Fert and to  
281 40 kg N ha<sup>-1</sup> year<sup>-1</sup> as N-Surf (Fig. 3d).

282 In the 2016/17 crop season, the highest root production, recorded in the 0-20 cm  
283 layer, was observed in grapevines grown in soil subjected to the application of 40 kg N  
284 ha<sup>-1</sup> year<sup>-1</sup> as N-Surf and to 100 kg N ha<sup>-1</sup> year<sup>-1</sup> as N-Fert (Fig. 4a). In the 2017/18 crop  
285 season, the highest root production, recorded in the 0-20 cm, 20-40 cm and 40-60 cm  
286 layers, were observed in grapevines subjected to 100 kg N ha<sup>-1</sup> year<sup>-1</sup> as N-Surf and N-  
287 Fert (Fig. 4b).

288

### 289 **Leaf N concentration**

290 In the 2016/17 crop season, the highest N concentrations, in leaves collected at  
291 flowering and at berry veraison were observed in grapevines subjected to application of  
292 40 kg N ha<sup>-1</sup> year<sup>-1</sup> as N-Surf and N-Fert (Fig. 5a, c). In the 2016/17 crop season, N  
293 concentration in leaves, collected at flowering were significant difference between N  
294 supply methods, except for rate of 100 kg N ha<sup>-1</sup> year<sup>-1</sup>, and at veraison there was a  
295 significant difference between the N supply methods for 0 kg N ha<sup>-1</sup> year<sup>-1</sup>. In the  
296 2017/18 crop season, the highest N concentration in leaves collected at flowering and at  
297 veraison was observed in grapevines grown in soil subjected to 100 kg N ha<sup>-1</sup> year<sup>-1</sup> as  
298 N-Fert (Fig. 5b, d). In 2017/18 crop season, N concentrations in leaves collected at  
299 flowering and at berries veraison in grapevines subjected to N-Surf did not show  
300 significant difference among N rates (Fig. 5b, d).

301

**302 Yield and its components**

303 In the 2016/17 crop season, the lowest weight of 100 berries was observed in  
304 grapevines grown in soil subjected to the application of 0 and 100 kg N ha<sup>-1</sup> year<sup>-1</sup>, than  
305 40 kg N ha<sup>-1</sup> year<sup>-1</sup>, as N-Fert (Fig. 6a). The weight of 100 berries recorded for  
306 grapevines subjected to N-Surf in the 2016/17 crop season did not statistically differ  
307 with N doses (Supplementary material 2). In the 2017/18 crop season, the weight of 100  
308 berries did not show statistically significant difference among N doses and modes (Fig.  
309 6b).

310 In 2016/17 crop season, the largest number of grape clusters per plant fertilized  
311 as N-Fert was observed at the rate of 100 kg N ha<sup>-1</sup> year<sup>-1</sup> (Fig. 6c). The number of  
312 grape clusters per plant fertilized as N-Surf were not affected by N application rates  
313 (Supplementary material 2). In 2017/18 crop season, the smallest numbers of grape  
314 clusters per plant were observed in control grapevines (Fig. 6d). In the two crop seasons  
315 - 2016/17 and 2017/18, the number of grape clusters per plant did not statistically differ  
316 between N supply modes (Supplementary material 2).

317 In the 2016/17 crop season, the lowest yield of N-Fert plant was observed in  
318 control (12.49 Mg ha<sup>-1</sup>) (Fig. 6e), whereas yield in N-Surf plants was not affected by  
319 application rates. In the 2017/18 crop season, the highest yields recorded were observed  
320 in grapevines grown in soil subjected to the application of 100 kg N ha<sup>-1</sup> year<sup>-1</sup> as N-  
321 Surf and N-Fert (29.77 and 27.86 Mg ha<sup>-1</sup>, respectively) (Fig. 6f). In the 2016/17 and  
322 2017/18 seasons, crop yield was not statistically affected by N application modes  
323 (Supplementary material 2).

324

**325 Chemical parameters of the must**

326 In the 2016/17 crop season, the highest total titratable acidity was observed in  
327 grape musts subjected to the application of 100 kg N ha<sup>-1</sup> year<sup>-1</sup> as N-Fert (Fig. 7a),  
328 whereas in the 2017/18 crop season, the highest total titratable acidity was recorded in  
329 grape musts subjected to the application of the same N dose, in both N-Fert and N-Surf  
330 modes (Fig. 7b). In the 2016/17 and 2017/18 crop seasons, the highest TSS values in  
331 must were observed in grapevines that were not fertilized (0 kg N ha<sup>-1</sup> year<sup>-1</sup>) (Fig. 7c,  
332 d). In the 2016/17 crop season, the highest total anthocyanin were observed in grape  
333 that were not fertilized (Fig. 7e), whereas in the 2017/18 crop season, the highest total  
334 anthocyanin values recorded were observed in grapevines subjected to the application of  
335 0 kg N ha<sup>-1</sup> year<sup>-1</sup> and 40 kg N ha<sup>-1</sup> year<sup>-1</sup>, for both N application modes (N-Surf and N-  
336 Fert modes) (Fig. 7f).

337

### 338 **Principal Component Analysis (PCA)**

339 The sum of components PC1 and PC2 explained 54.18% of original data  
340 variability (Fig. 8). The PC1 explained 32.22% of data variability and enabled  
341 observing treatment trends to show repeated behavior throughout crop seasons, a fact  
342 that was evidenced by the constant position of the 0 kg N ha<sup>-1</sup> year<sup>-1</sup> dose (at the right  
343 side) in the spatial distribution. These results differed from those recorded for N doses  
344 of 40 kg N ha<sup>-1</sup> year<sup>-1</sup> and 100 kg N ha<sup>-1</sup> year<sup>-1</sup>, which alternated in the most productive  
345 position between crop seasons and always headed left in the spatial distribution.

346 The PC2 explained 21.96% data variability and it was efficient in separating  
347 grapevines' productive behavior in each crop season, which was delimited by different  
348 ellipses. Thus, plants subjected to the application of 40 kg N ha<sup>-1</sup> year<sup>-1</sup> and 100 kg N  
349 ha<sup>-1</sup> year<sup>-1</sup> were mostly influenced by production variables such as yield, number of  
350 grape clusters and weight of 100 berries; as well as by several root parameters such as

351 surface area, diameter, number of living roots and root production, which were  
352 positively correlated to each other. On the other hand, grapevines that were not  
353 subjected to N application mainly showed the influence of variables on quality  
354 parameters such as total anthocyanins and TSS. This variable of quality parameters also  
355 showed negative linear correlation to grape yield.

356

## 357 **Discussion**

358         The increased root surface area observed in plants subjected to N application  
359 may have happened due to increased content of mineral N forms (*i.e.*  $\text{NH}_4^+$  and  $\text{NO}_3^-$ ) in  
360 the soil which derived from the applied urea (Radville et al., 2016; Yan et al., 2017).  
361 Nitrogen application as fertigation probably promoted the movement of  $\text{NO}_3^-$ , a form of  
362 N common in subtropical soils (Barlow, Bond, Holzapfel, Smith, & Hutton, 2009; G.  
363 Brunetto et al., 2017), through to deeper soil layers and, consequently, reduced N losses  
364 due to volatilization. This outcome most likely resulted from the downward flow of  
365 water deriving from fertigation (Castellanosa et al., 2013), considering that the  
366 investigated soil was sandy with low OM content, a fact that facilitates the ion  
367 approximation through mass flow until the outer surface of the root system (Comas et  
368 al., 2010).

369         Increased  $\text{NO}_3^-$  concentration in soil solution stimulates the development of  
370 plant root system components, such as root diameter and surface area, two components  
371 sensitive to  $\text{NO}_3^-$  concentrations in the soil (Othman and Leskovar 2019). It also enables  
372 activating transporters, such as NRT1.1, in the root system, a fact that increases the  
373 amount of  $\text{NO}_3^-$  uptake by plants (Krouk et al., 2010; Pii et al., 2013; Remans et al.,  
374 2006). This  $\text{NO}_3^-$  root transporter is also capable of transporting auxin, under the control  
375 of  $\text{NO}_3^-$  in root-growth medium. Roots grown at high  $\text{NO}_3^-$  concentrations shows the



376 preference for transporting  $\text{NO}_3^-$  over auxin, which accumulates in the roots and favors  
377 root growth (Mounier et al., 2014). Increased  $\text{NO}_3^-$  concentrations in root tissues also  
378 increases cytokinins metabolism rates (Silva and Delatorre 2009); cytokinines are  
379 important cell division regulators that lead to increased root system surface area  
380 (Centinari et al., 2016; Comas et al., 2010).

381 In the present study, the increased mineral N availability in the soil contributed  
382 to increase the mean root diameter in grapevines grown in soil subjected to N  
383 application rate of 40 and 100 kg N ha<sup>-1</sup> year<sup>-1</sup>. It may have happened due to decreased  
384 soil pH values, which reduced cell division and elongation (Barlow et al., 2009; Yan et  
385 al., 2017). Such process often takes place because  $\text{H}^+$  is released during nitrification and  
386 stimulates the solubilization of  $\text{Al}^{+3}$  from soil native minerals (Miotto et al., 2019).  
387 Released  $\text{Al}^{+3}$  can bind to carboxylic groups and to pectic materials in the cell wall.  
388 This process increases the stiffness and decreases the extensibility of root cell walls;  
389 consequently, it can lead to disorganized cell division, which impairs root elongation  
390 and enables larger root diameter (Riaz et al., 2018; Zhou et al., 2014).

391 Nitrogen applications increased mineral N contents in the soil, a fact that  
392 stimulated root emission, as observed in the total number of living roots in grapevines  
393 subjected to N addition in both application modes. Roots preferentially grow in soil  
394 regions presenting the highest nutrient contents (Centinari et al., 2016), including  
395 mineral N (Lima, Kojima, Takahashi, & von Wirén, 2010). This because  $\text{NO}_3^-$  available  
396 in the soil acts as a signaling molecule that can regulate the expression of genes  
397 involved in plant hormonal balance (Lee et al., 2016; Xuan, Beeckman, & Xu, 2017).  
398 This process contributes to the production and development of new roots, mainly in soil  
399 regions where  $\text{NO}_3^-$  is available (Krouk et al., 2010; Remans et al., 2006).

400           The increased production and number of living roots in grapevines subjected to  
401 the highest N dose application ( $100 \text{ kg N ha}^{-1} \text{ year}^{-1}$ ) can also be associated with  
402 strategies adopted by plants to uptake and accumulate N, such as amino acids and  
403 proteins, in perennial organs (Ortiz-Lopez, Chang, & Bush, 2000). The increased N  
404 availability in the soil increases the likelihood of nutrient uptake; however, in addition  
405 to N supply, root emission and renewal processes also depend on the internal carbon (C)  
406 balance of grapevines (Jagodzinski and Kalucka 2011; Yan et al. 2017). In periods of  
407 higher metabolic activity, such as intense annual plant growth, part of N can be  
408 redistributed to shoots and leaves, in order to enable biochemical and physiological  
409 processes in grapevines (Gustavo Brunetto et al., 2016; Zambrosi et al., 2012).  
410 However, senescence is expected to happen in part of the roots when most of the C  
411 found in plants is allocated to the shoot, as it happens in vegetative growth periods  
412 (Kou, Chen, et al., 2015; Kou, Guo, Yang, Gao, & Li, 2015). This was observed in  
413 grapevines subjected to  $40 \text{ kg N ha}^{-1} \text{ year}^{-1}$  as N-Surf mode and to  $100 \text{ kg N ha}^{-1} \text{ year}^{-1}$   
414 as N-Fert mode, in the two evaluated crop seasons.

415           Increased N concentrations in leaves collected at flowering and at berry veraison  
416 can be attributed to increased mineral N content in the soil, in grapevines subjected to N  
417 applications. Part of N uptake by plants can be incorporated into C skeletons, such as  
418 amino acids, amines and proteins, which can be transported to leaves, which undergo  
419 intense cell division and elongation processes (Radville et al., 2016; Yu et al., 2012).  
420 This phenomenon, in association with the application of higher N rates such as  $100 \text{ kg}$   
421  $\text{N ha}^{-1} \text{ year}^{-1}$ , led to increase of N concentrations in leaves of plants subjected to this  
422 treatment. In addition, large rainfall volumes recorded in concomitance of N  
423 applications, as observed in the 2016/17 crop season (Fig. 1), have stimulated  $\text{NO}_3^-$   
424 movement in the soil solution, since  $\text{NO}_3^-$  links itself to reactive particles in the soil by

425 means of external sphere complex (Comas et al., 2010; Li, Wang, & Stewart, 2013;  
426 Lorensini et al., 2012). This process may have stimulated the emission of new in-depth  
427 roots and increased grapevine N uptake (Gustavo Brunetto et al., 2016).

428         The increased number of grape clusters per plant and yield of grapevines  
429 subjected to the highest N application (100 kg N ha<sup>-1</sup> year<sup>-1</sup>) might be associated to a  
430 higher mineral N availability in the soil (Brunetto et al., 2016; Steenwerth and Belina,  
431 2010), a fact that stimulated root growth to 60 cm. On the one hand, increased N  
432 concentrations in plant leaves stimulated photosynthetic processes, since higher  
433 concentrations of chlorophylls and chloroplast proteins, such as RuBisCO, enabled  
434 increased CO<sub>2</sub> fixation in leaf tissues, as well as increased plant growth (Blank et al.,  
435 2018; Moriwaki et al., 2019). On the other hand, plants needed to be photosynthetically  
436 active in order to enable NO<sub>3</sub><sup>-</sup> uptake and reduction. Therefore, higher rates of  
437 photosynthesis can also favor the uptake of N to be increased, since enzymes  
438 responsible for reducing N were regulated by light, whereas the energy used in this  
439 process derived from chloroplasts (Krouk et al., 2010).

440         The highest total titratable acidity in the must of grapevines grown in soil  
441 subjected to the highest N applications may be explained by the inverse relationship  
442 between the total titratable acidity in the must and the vegetative growth of grapevine  
443 shoots. Increased leaf yield also increases the shading of clusters inside plants, which, in  
444 its turn, delays the ripening of grapes and decreases organic acid degradation in berries  
445 (Centinari et al., 2016; Vilanova, Fandiño, Frutos-Puerto, & Cancela, 2019). Decreased  
446 TSS values observed in the must of grapevines grown in soil subjected to the  
447 application of higher N doses resulted from the increased number of grape clusters per  
448 plant and, mainly, from the increased grape yield, which contributed to dilute the sugars  
449 in the berries (Guilpart, Metay, & Gary, 2014). In addition, decreased TSS levels in the

450 must may have resulted from increased leaf area in grapevines subjected to higher N  
451 applications, since excessive leaf growth reduces direct sunlight incidence on berries, a  
452 fact that leads to lower sugars concentration (Sadras and Moran 2012). Nitrogen  
453 applications at the highest dose ( $100 \text{ kg N ha}^{-1} \text{ year}^{-1}$ ) and N supply modes (N-Surf and  
454 N-Fert) led to TSS values lower than 14 °Brix, which are not appropriate for  
455 winemaking processes (Chiarotti, Guerios, Cuquel, & Biasi, 2011). The application of  
456 40 and  $100 \text{ kg N ha}^{-1} \text{ year}^{-1}$ , in both N supply methods, also reduced total anthocyanin  
457 values in grape must. This outcome may be associated with increased number of grape  
458 clusters per plant and with grape yield, which is often associated with increased berry  
459 size and pulp/peel ratio. Thus, the concentration of important compounds, such as total  
460 anthocyanins in the must, which account for the reddish color of the must and,  
461 consequently, of the wine, gets diluted (Brunetto et al. 2009b; Sadras and Moran, 2012;  
462 Sofo et al. 2012). In addition, as plants' vegetative growth increases, part of the  
463 anthocyanins found in the clusters may have been redistributed to other growth sites and  
464 had their contents decreased in berries (Terrier, Poncet-Legrand, & Cheynier, 2009).

465         Results of the principal component analysis showed intrinsic climate variation in  
466 each crop season, which resulted in plants' productive response to the application of  
467 different treatments. It also showed annual variations in fruit yield and nutritional status.  
468 It happened because fruit plant species have a cyclical dynamics of C allocation to  
469 different organs that can favor yield in a given year and species maintenance in the  
470 following year, due to reserve accumulation in organs that require more energy for  
471 protein synthesis, mainly in plants subjected to N applications (Ceccon et al. 2016;  
472 Artacho and Bonomelli 2017). The PCA enabled a better understanding of the positive  
473 relationship among root system, plants yield, N application rates and modes. These  
474 results showed that as photoassimilates accumulated in grapevines roots, water and

475 nutrient intake increased, a fact that improved crop yield (Antolín et al., 2003; Barlow et  
476 al., 2009; Comas et al., 2010). In addition to this process, there was a decrease in the  
477 oenological quality of the must, which was diagnosed based on decreased total  
478 anthocyanins and total soluble solids values, as well as on increased total acidity values.  
479

## 480 **Conclusion**

481         The application of 40 kg N ha<sup>-1</sup> year<sup>-1</sup>, as N-Surf mode, provided adequate N  
482 supply to the soil and increased root system development, which enabled larger root  
483 surface area, larger number of living roots and greater root production. This process  
484 enabled an increase of soil N uptake by plants, higher N concentration in leaves and  
485 greater grape yield. However, N supply decreased the quality of the must, as observed  
486 in decreased TSS and total anthocyanins values and in increased total titratable acidity  
487 values, which are not desirable for red wine making processes.

**488 Author contribution statement**

489           Conceptualization, MSSK; data duration, MSSK, LOS and RS; formal analysis,  
490 MSSK, LOS and RS; funding acquisition, MT, EB and TLT; investigation, WSA, GAS,  
491 CFP, GS, JPJS and ROSS; project administration, GB; supervision, GB; writing—  
492 original draft, MSSK; writing—review and editing, LOS, RS, MT, EB, TLT and GB.

493

**494 Conflicts of interest**

495           The authors declare no financial or other competing conflicts of interest.

496

**497 Acknowledgements**

498           We are grateful to the Conselho Nacional de Desenvolvimento Científico e  
499 Tecnológico (Brazilian National Council for Scientific and Technological  
500 Development) – CNPq (Process number 301509/2015-8, 408318/2018 and  
501 423772/2018), Coordenação de Aperfeiçoamento de Pessoal de Nível Superior  
502 (Brazilian Federal Agency for Support and Evaluation of Graduate Education) –  
503 CAPES, Empresa Brasileira de Pesquisa Agropecuária (Brazilian Agricultural Research  
504 Corporation) – Embrapa (Call announcement 04/2016), and Fundação de Amparo a  
505 Pesquisa do Estado do Rio Grande do Sul (Foundation for Research Support of the State  
506 of Rio Grande do Sul) - FAPERGS (Term of grant 17/2551-0000925-8) for granting  
507 scholarship provided and the financial resources made available for this study.

508

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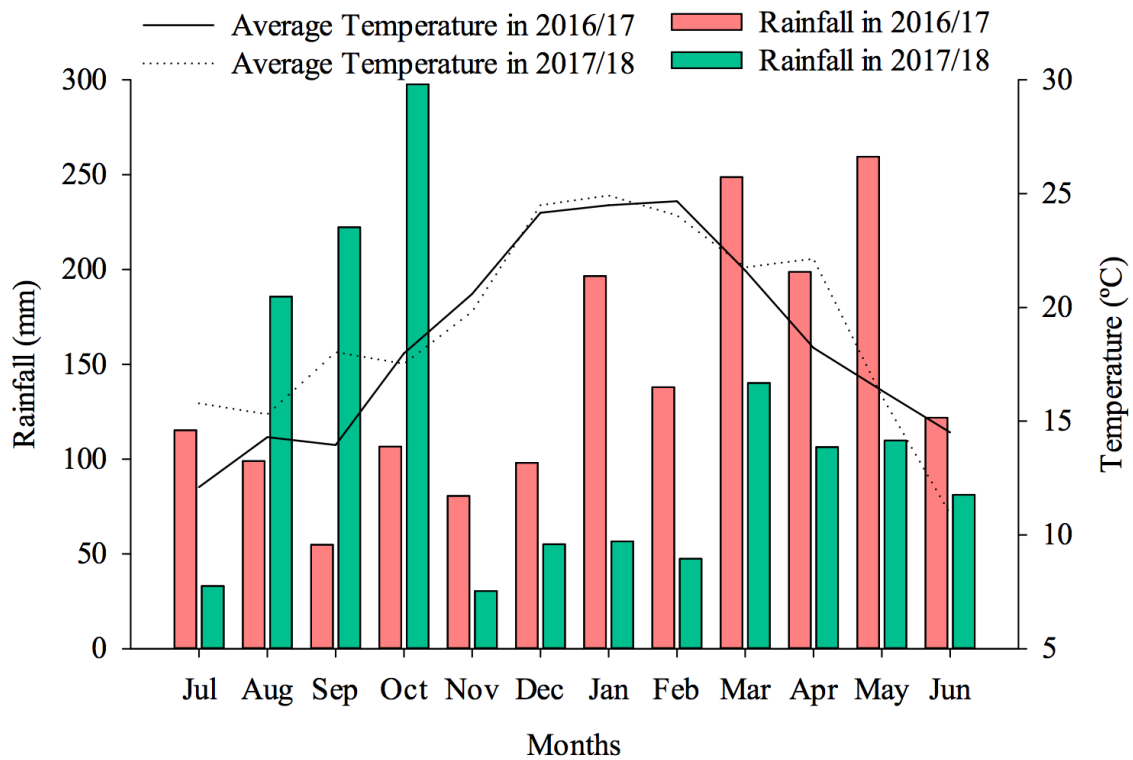


752 **Table 1** Main physical and chemical characteristics of the soil in the experimental site at 0.0-0.20 m soil  
753 layer

Soil characteristics	0.0-0.20 m
Clay (Pipette method) (g kg <sup>-1</sup> )	63
Silt (Pipette method) (g kg <sup>-1</sup> )	107
Sand (Pipette method) (g kg <sup>-1</sup> )	830
Organic matter (Walkley Black method) (g kg <sup>-1</sup> )	10.0
pH in water (1:1 ratio)	5.5
Available P (Extracted by Mehlich-1) (mg kg <sup>-1</sup> )	20.0
Exchangeable K (Extracted by Mehlich-1) (mg kg <sup>-1</sup> )	65.0
Exchangeable Ca (Extracted by KCl 1 mol L <sup>-1</sup> ) (cmol <sub>c</sub> kg <sup>-1</sup> )	1.7
Exchangeable Mg (Extracted by KCl 1 mol L <sup>-1</sup> ) (cmol <sub>c</sub> kg <sup>-1</sup> )	0.8
Exchangeable Al (Extracted by KCl 1 mol L <sup>-1</sup> ) (cmol <sub>c</sub> kg <sup>-1</sup> )	0.0

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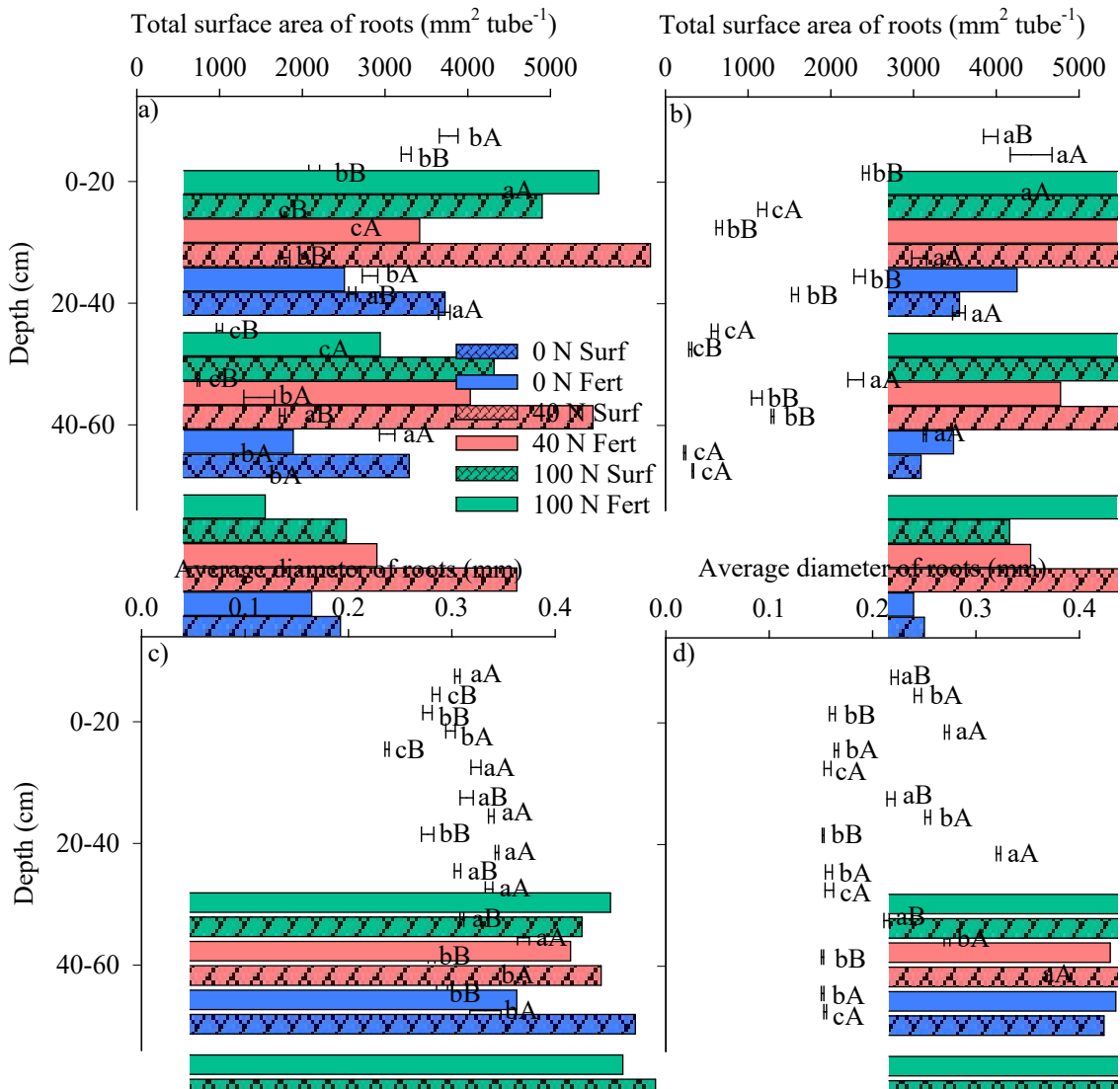
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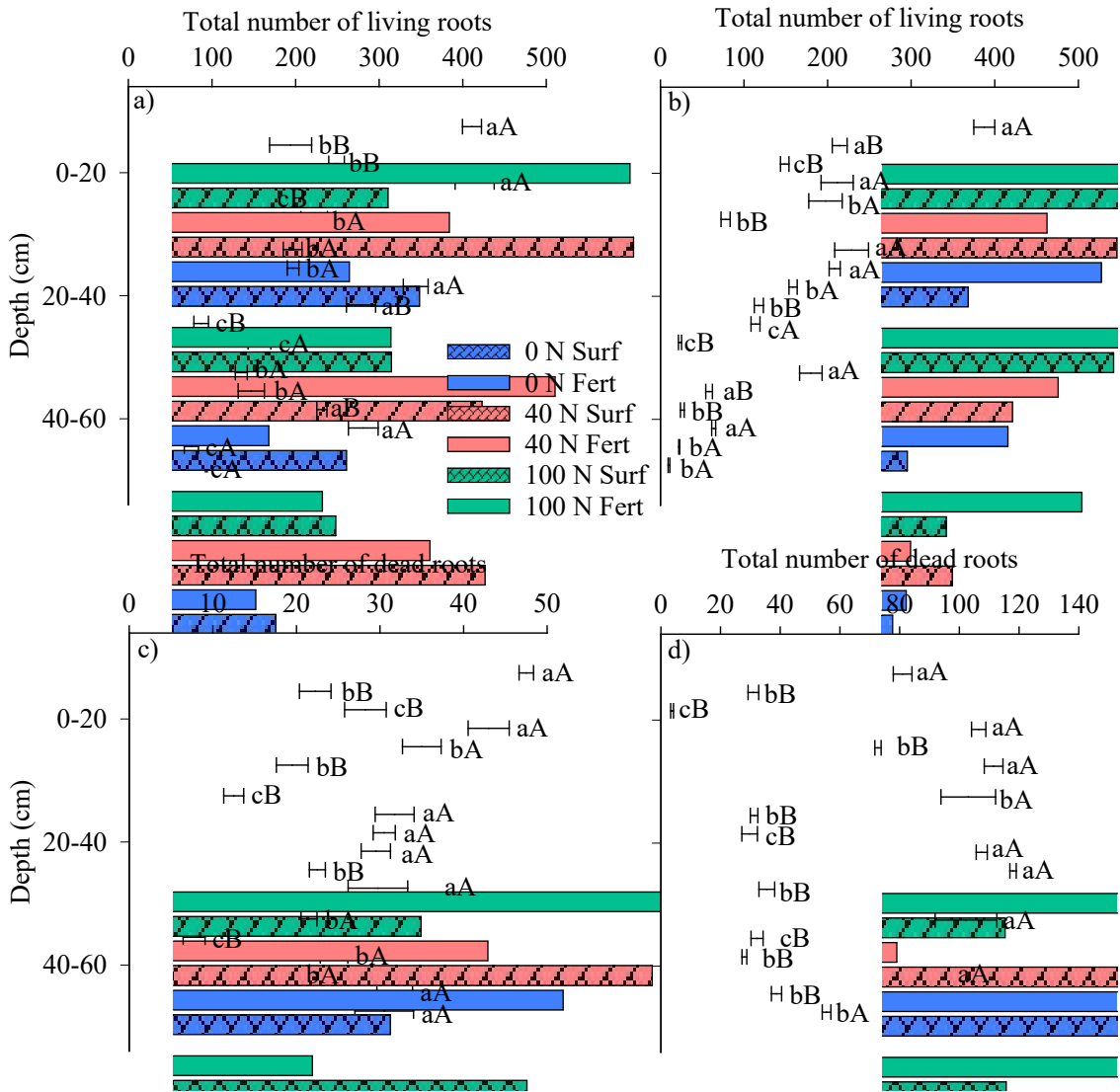
757 **Fig. 1** Monthly accumulated rainfall (mm) and average monthly temperature (°C) of the 2016/17 and  
 758 2017/18 crops season obtained at the experimental station in Santana do Livramento, state of Rio Grande  
 759 do Sul, Brazil

760



761

762 **Fig. 2** Effect of N application rate and method on root total surface area in the 2016/17 (a) and 2017/18  
 763 crop season (b), average diameter of roots in the 2016/17 (c) and 2017/18 crop season (d) in 0-20, 20-40  
 764 and 40-60 cm soil layers, at flowering of ‘Alicante Bouschet’ (*Vitis vinifera* L.) grapevine (N Surf -  
 765 Nitrogen surface; N Fert - Nitrogen fertigation; 0, 40 and 100 kg N ha<sup>-1</sup> year<sup>-1</sup>). Horizontal bars indicate  
 766 the standard error ( $n = 4$ ). Different lower-case letters indicate a significant difference among N doses in  
 767 the same application modes, and different upper-case letters indicate a significant difference among the N  
 768 application modes in the same N doses by the Scott-Knott test ( $p < 0.05$ )



770

771 **Fig. 3** Effect of N application rate and method on total number of living roots in the 2016/17 (a) and

772 2017/18 crop season (b); total number of dead roots in the 2016/17 (c) and 2017/18 crop season (d) in 0-

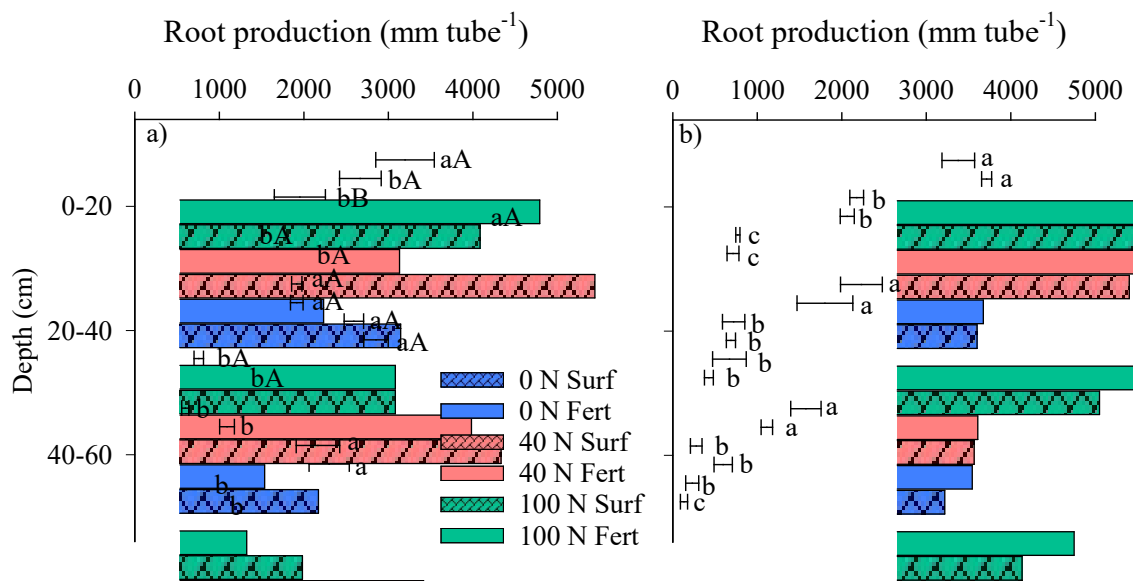
773 20, 20-40 and 40-60 cm soil layers, at flowering of ‘Alicante Bouschet’ (*Vitis vinifera* L.) grapevine (N

774 Surf - Nitrogen surface; N Fert - Nitrogen fertigation; 0, 40 and 100 kg N ha<sup>-1</sup> year<sup>-1</sup>). Horizontal bars

775 indicate the standard error ( $n = 4$ ). Different lower-case letters indicate a significant difference among N

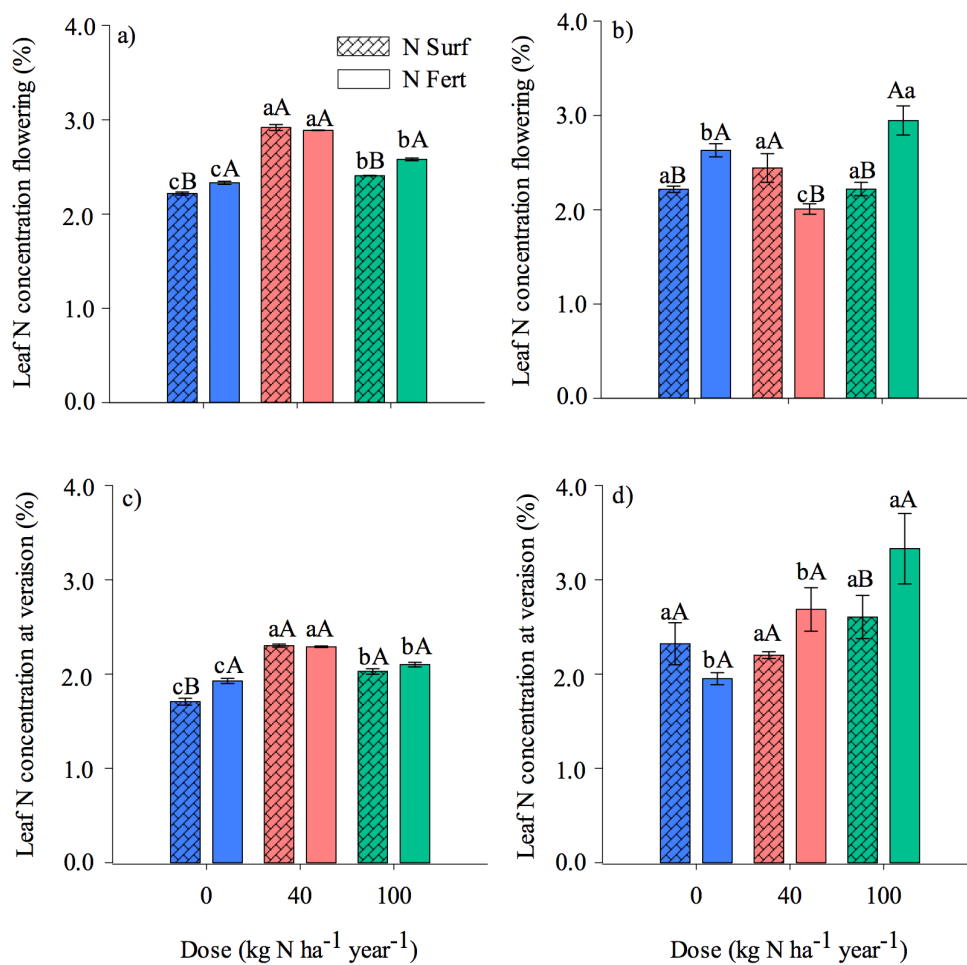
776 doses in the same application modes, and different upper-case letters indicate a significant difference

777 among the N application modes in the same N doses by the Scott-Knott test ( $p < 0.05$ )



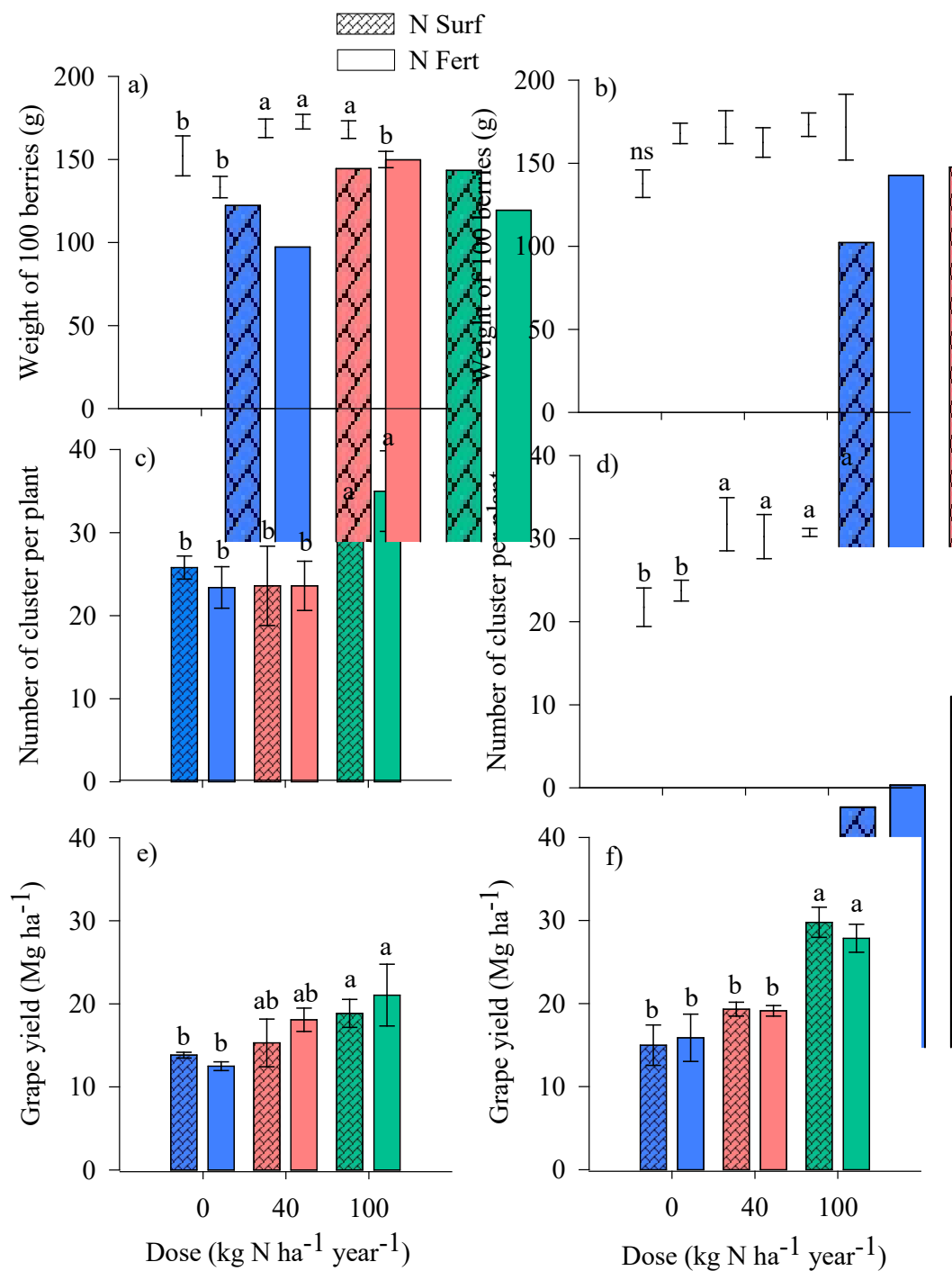
778

779 **Fig. 4** Effect of N application rate and method on root production (mm tube<sup>-1</sup>) in the 2016/17 (a) and  
 780 2017/18 crop season (b) at flowering of 'Alicante Bouschet' (*Vitis vinifera* L.) grapevine (N Surf -  
 781 Nitrogen surface; N Fert - Nitrogen fertigation; 0, 40 and 100 kg N ha<sup>-1</sup> year<sup>-1</sup>). Horizontal bars indicate  
 782 the standard error ( $n = 4$ ). Different lower-case letters indicate a significant difference among N doses in  
 783 the same application modes, and different upper-case letters indicate a significant difference among the N  
 784 application modes in the same N doses by the Scott-Knott test ( $p < 0.05$ )



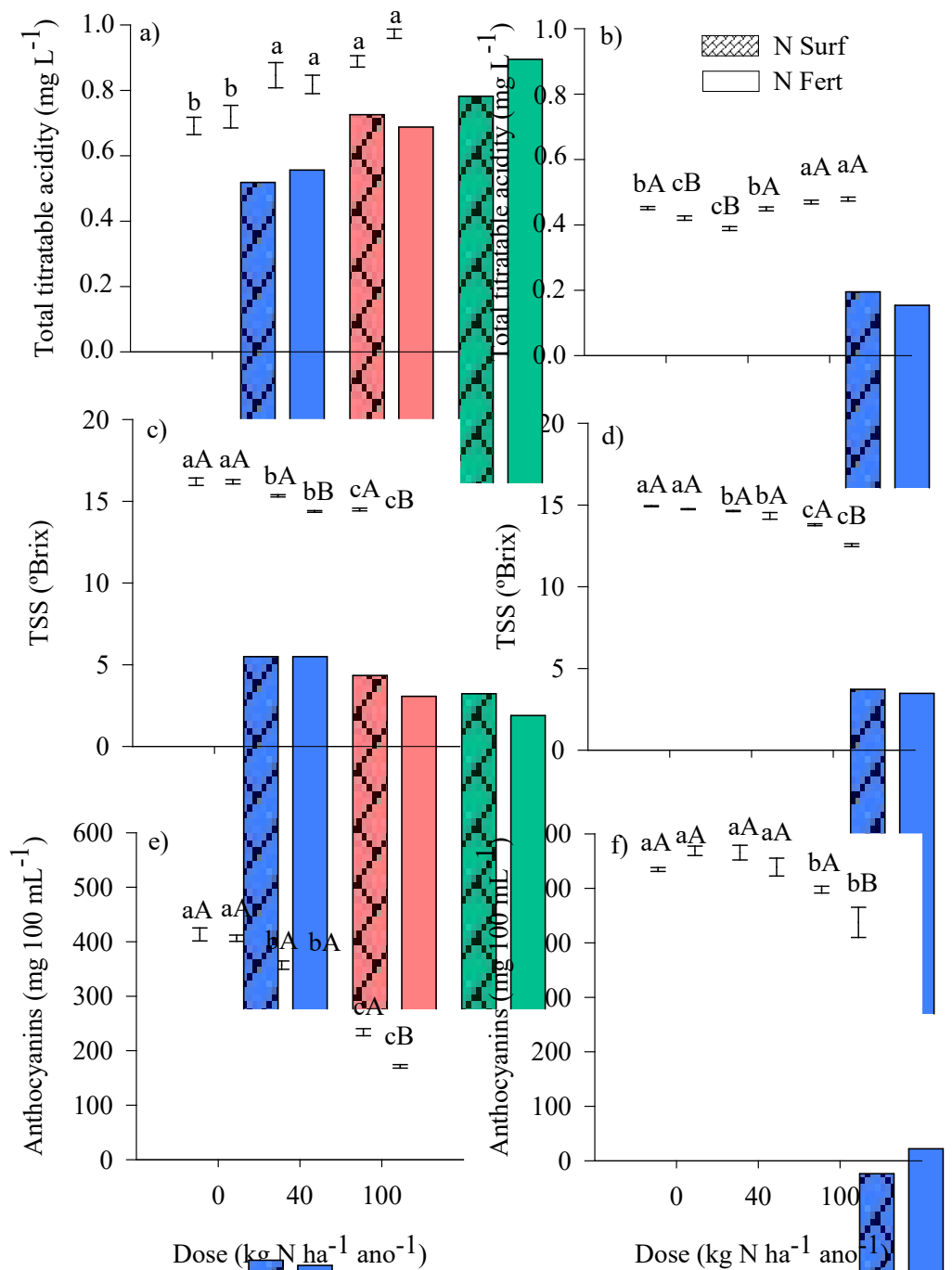
785

786 **Fig. 5** Effect of N application rate and method on leaf N concentration at flowering in 2016/17 (a) and  
 787 2017/18 crop season (b); leaf N concentration at veraison in 2016/17 (c) and 2017/18 crop season (d) of  
 788 'Alicante Bouschet' (*Vitis vinifera* L.) grapevine (N Surf - Nitrogen surface; N Fert - Nitrogen fertigation;  
 789 0, 40 and 100 kg N ha<sup>-1</sup> year<sup>-1</sup>). The vertical bars indicate the standard error ( $n = 4$ ). Different lower-case  
 790 letters indicate a significant difference among N doses in the same application modes, and different  
 791 upper-case letters indicate a significant difference among the N application modes in the same N doses by  
 792 the Scott-Knott test ( $p < 0.05$ )



793

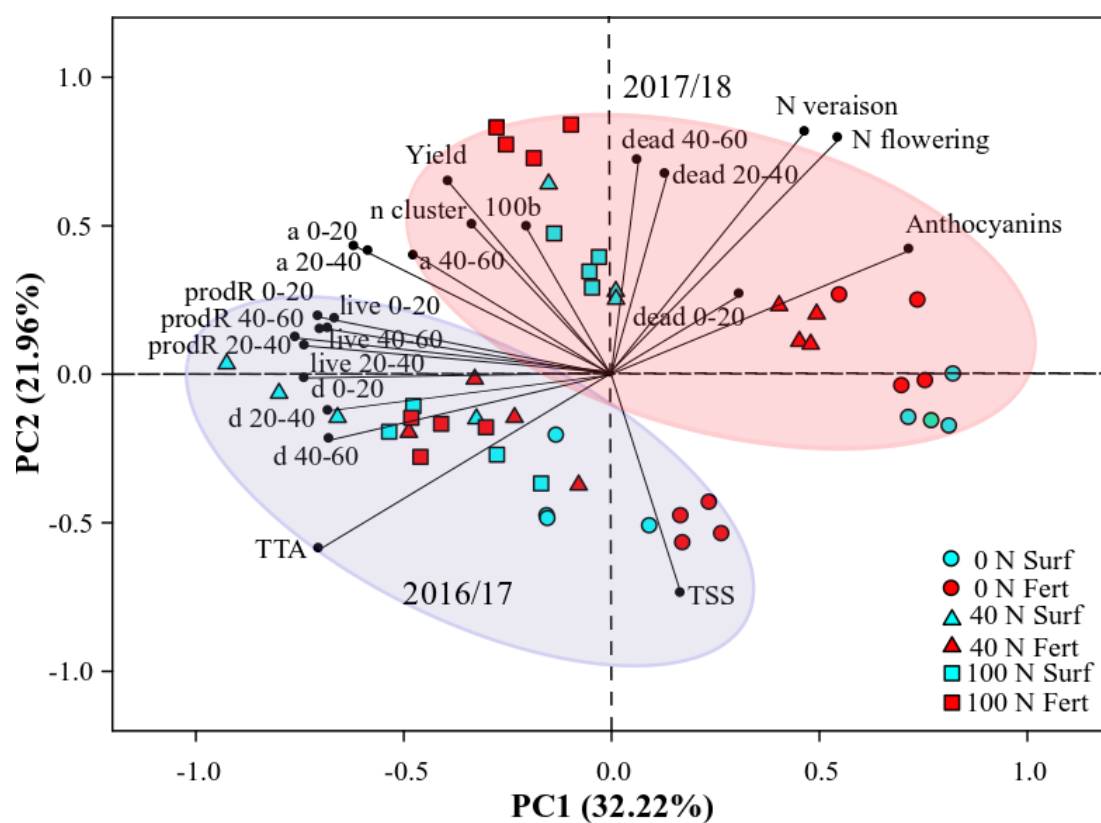
794 **Fig. 6** Effect of N application rate and method on weight of 100 berries in 2016/17 (a) and 2017/18 crop  
 795 season (b); number of cluster per plant in the 2016/17 (c) and 2017/18 crop season (d); grape yield in the  
 796 2016/17 (e) and 2017/18 crop season (f) of 'Alicante Bouschet' (*Vitis vinifera* L.) grapevine (N Surf -  
 797 Nitrogen surface; N Fert - Nitrogen fertigation; 0, 40 and 100 kg N ha<sup>-1</sup> year<sup>-1</sup>). The vertical bars indicate  
 798 the standard error ( $n = 4$ ). Different lower-case letters indicate a significant difference among treatments  
 799 by the Tukey test ( $p < 0.05$ )



800

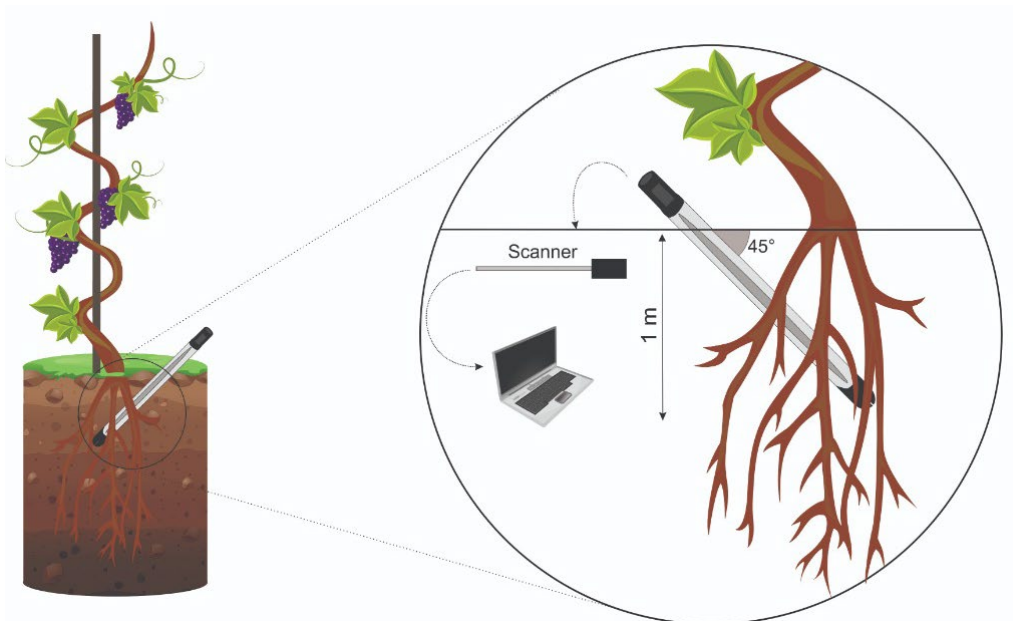
801 **Fig. 7** Effect of N application rate and method on total titratable acidity in 2016/17 (a) and 2017/18 crop  
 802 season (b); total soluble solids (TSS) in the 2016/17 (c) and 2017/18 crop season (d); total anthocyanins  
 803 (Anthocyanins) in the 2016/17 (e) and 2017/18 crop season (f) of the must of 'Alicante Bouschet' (*Vitis*  
 804 *vinifera* L.) grapevine (N Surf - Nitrogen surface; N Fert - Nitrogen fertigation; 0, 40 and 100 kg N ha<sup>-1</sup>  
 805 year<sup>-1</sup>). The vertical bars indicate the standard error ( $n = 4$ ). Different lower-case letters indicate a  
 806 significant difference among N doses in the same application modes, and different upper-case letters  
 807 indicate a significant difference among the N application modes in the same N doses by the Scott-Knott  
 808 test ( $p < 0.05$ )





809

810 **Fig. 8** Relationship between principal component 1 (PC1) and 2 (PC2) of the morphological parameters  
 811 of the root system (total surface area of roots (a), average diameter of roots (d), total number of living  
 812 roots (living), total number of dead roots (dead) and root production (prodR) in 0-20, 20-40 and 40-60 cm  
 813 soil layers), leaf N concentration (leaf N concentration at flowering (N flowering) and leaf N  
 814 concentration at veraison (N veraison)), grape yield parameters (grape yield (Yield), number of cluster  
 815 per plant (n cluster), weight of 100 berries (100b)) and chemical parameters of the must (total  
 816 anthocyanins (Anthocyanins), total soluble solids (TSS), total titratable acidity (TTA)) of 'Alicante  
 817 Bouschet' (*Vitis vinifera* L.) grapevine subjected to different N doses and application modes



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820 **Supplementary material 1.** Schematic representation of the imaging system from minirizotron tubes  
821 installed in the vineyard of the experimental area; *in situ* image capture high resolution (600 dpi) from a  
822 cylindrical scanner (Root Imager – CI600®).

823 **Supplementary material 2.** Analysis of variance of the roots parameters, leaf N concentration, grape  
 824 yield and chemical of the must of 'Alicante Bouschet' (*Vitis vinifera* L.) grapevine subjected to different  
 825 N doses and application modes.

Variables	Effect				
	Soil layer (cm)	Crop season	Mode	Dose	Interaction
Total surface area of roots (mm <sup>2</sup> tube <sup>-1</sup> )	0-20	2016/17	***	***	***
		2017/18	***	***	***
	20-40	2016/17	***	***	***
		2017/18	***	***	***
	40-60	2016/17	***	***	***
		2017/18	***	***	***
Average diameter of roots (mm)	0-20	2016/17	***	**	***
		2017/18	***	***	***
	20-40	2016/17	***	*	***
		2017/18	***	***	***
	40-60	2016/17	***	***	***
		2017/18	***	***	***
Total number of living roots	0-20	2016/17		***	***
		2017/18	***	***	***
	20-40	2016/17		***	***
		2017/18	***	***	**
	40-60	2016/17	**	***	***
		2017/18	***	***	***
Total number of dead roots	0-20	2016/17	***	*	***
		2017/18	***	***	***
	20-40	2016/17	***	*	***
		2017/18	***		***
	40-60	2016/17	***	***	*
		2017/18	*	***	***
Root production (mm tube <sup>-1</sup> )	0-20	2016/17	*	***	**
		2017/18		***	
	20-40	2016/17	**	***	
		2017/18		***	
	40-60	2016/17		***	
		2017/18		***	**
Leaf N concentration at flowering (%)	2016/17	***	***	***	
	2017/18	*	**	***	
Leaf N concentration at veraison (%)	2016/17	***	***	***	
	2017/18		*	*	
Grape yield (Mg ha <sup>-1</sup> )	2016/17		***		
	2017/18		***		
Number of cluster per plant	2016/17		**		
	2017/18		***		
Weight of 100 berries (g)	2016/17		***		
	2017/18				
Total anthocyanins (mg 100mL <sup>-1</sup> )	2016/17	***	***	**	
	2017/18		***	*	
Total soluble solids (°Brix)	2016/17	***	***	***	
	2017/18	***	***	***	
Total titratable acidity (mg L <sup>-1</sup> )	2016/17		***		
	2017/18	*	***	***	

826 \*0.05; \*\*0.01; \*\*\*<0.001