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1 **Implications of hornbeam and beech root systems on slope stability: from field and**
2 **laboratory measurements to modelling methods**

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17 Abstract

18 *Purpose* This study investigated root distribution and root reinforcement estimated by field and laboratory
19 measurements and modelling methods, in function of species, trees diameter at breast height (DBH), slope position,
20 altitude, vertical and horizontal distances from tree in Hyrcanian temperate ecoregions of Iran.

21 *Method* 1080 profile trenches with maximum 1 m depth were excavated on upslope and downslope from trunks of
22 *Carpinus betulus* and *Fagus orientalis* with the DBH of 7.5-32.5, 32.5-57.5, and 57.5-82.5 cm at three altitudes (400,
23 950, and 1300 m a.s.l.).

24 *Results* Root distribution results indicated that: (i) frequency of small roots (2-5mm of diameter) of *C. betulus* and fine
25 roots (0-2 mm) of *F. orientalis* are the highest, whereas the frequency of large roots (>10mm) of both species is the
26 lowest, (ii) the Root Area Ratio (RAR) of *C. betulus* is always higher than *F. orientalis*, (iii) the trees with larger DBH
27 have more roots than those with a smaller DBH, (iv) the RAR of *F. orientalis* in upslope is higher than in downslope;
28 however, the RAR of *C. betulus* for both slopes are similar, (v) the RAR in the 1300 m altitude is the highest, and (vi)
29 the RAR decreases with increasing distance from tree trunk and from soil surface. Furthermore, it is evident that: (i)
30 root reinforcement of *C. betulus* is higher than *F. orientalis*, (ii) altitude has a significant effect on root reinforcement
31 of *C. betulus*, (iii) root reinforcement of large trees is the highest, and (iv) root reinforcement decreases with increasing
32 distance from tree trunks.

33 *Conclusion* *C. betulus* is preferable to *F. orientalis* for increasing slope stability. Forest managers should consider this
34 outcome when developing strategies for silvicultural treatment and reforestation projects in mountainous areas.

35 **Keywords:** Hillslope stabilization, Root mechanical properties, Root Bundle Model Weibull, Root distribution, Root
36 reinforcement, Soil bioengineering, Iran.

37 1. Introduction

38 Forests play a significant role in preventing and mitigating hydrogeomorphic hazards such as shallow landslides,
39 rockfalls, and avalanches. Trees generally provide more protective functions than shrubs and herbs through some
40 hydrological and mechanical processes: rainfall interception (Sadeghi et al. 2020; Lin et al. 2020), soil coverage by
41 rainfall splash (Lin et al. 2020; Williams et al. 2020), buttressing and arching (Gray and Sotir 1996), soil reinforcement
42 by roots in shallower and deeper soil layers (Morgan and Rickson 1995). In detail, trees contribute to regulate water
43 cycle: intercepting rainfall, altering hydraulic conductivity through physical transformation of the soil by roots, and
44 transpiring stored water (Stokes et al. 2014; Vergani et al. 2017a). Moreover, roots contrast the triggering mechanisms
45 of shallow landslides up to 2 m extending and penetrating the soil mantle and sometimes crossing the failure plane,
46 and increasing tensile and shear soil resistance (Norris et al. 2008). On hillslopes, thick roots can act as piles to reinforce
47 the soil, while fine roots act in tension, cross the slip surface and strengthen the soil by adding cohesion. Quantifying
48 the mechanical effects of vegetation on hillslope stabilization remains an unsolved issue and has been frequently
49 investigated since the Sixties (e.g., Endo and Tsuruta 1969; O'Loughlin 1974; Genet et al. 2008). Root systems
50 stabilize the shallower soils by three different mechanisms, called in general root reinforcement: basal root

51 reinforcement, lateral root reinforcement, and root reinforcement under compression (stiffening and buttressing of
52 sliding mass under compression (Giadrossich et al. 2019). When roots cross the shear plane, basal root reinforcement
53 acts as an anchor inside the stable soil layer or bedrock. Lateral root reinforcement is the most effective mechanism in
54 stabilizing landslide-prone slopes, although increasing the landslide size, such contribution declines (Milledge et al.
55 2014). Finally, the contribution of roots under compression consists in mobilizing an additional resistance across the
56 shear plane, which leads to a complex bending-tensioning of rooted-soil (Schwarz et al. 2015).

57 Estimating root reinforcement depends by four main factors: root density (i.e. the number of roots into the soil), root
58 spatial distribution into the soil, root diameter (i.e. larger roots are more resistant than fine ones), and root
59 biomechanical properties (i.e., tensile resistance, elasticity, etc.) (Mao 2022). Several parameters individually influence
60 root distribution, including forest stand characteristics (such as tree spatial distribution), diameter at the breast height
61 (DBH), tree age, tree species composition, distance from the trunk, position on the slope, growth conditions (including
62 soil temperature, soil depth, nutrient), and moisture content (Genet et al. 2010; Mao et al. 2012; Cislighi et al. 2021).
63 Moreover, it is worth noting that the spatial root distribution depends on species morphology and environmental
64 conditions, soil type, soil depth, and availability of water and nutrients (Phillips et al. 2014; Vergani et al. 2017a). For
65 these reasons, investigating the spatial heterogeneity of roots at stand scale and modelling the lateral root distribution
66 is very challenging (Vergani et al. 2017a). A proxy parameter of the spatial distribution of roots is the Root Area Ratio
67 (RAR), computed as the total cross-sectional area of all roots divided by the total soil area, and largely used in the
68 scientific literature (Bischetti et al. 2009; Mao et al. 2012; Arnone et al. 2016). Conversely, most studies have been
69 exclusively focused on fine roots (<2 mm), whereas there is less information on coarse roots, especially roots larger
70 than 10 mm (Giadrossich et al. 2020). In addition, few studies about root distribution as a function of DBH and distance
71 from the tree trunk have allowed modelling the root distribution (Schwarz et al. 2010; Cislighi et al. 2021). Roots
72 biomechanical properties have been extensively studied compared to root distribution.

73 Observing data on biomechanical properties and root distribution allowed to develop and improve the use of
74 numerical models that quantify root reinforcement values to be included into slope stability analysis (Ekanayake and
75 Phillips 1999). Numerical models have been generally used since the development of the pioneering model Wu and
76 Waldron (W&W) based on the assumption that roots are elastic fibers extending perpendicular to a shear surface,
77 moreover all roots break at the same time (Wu, 1976; Waldron, 1977). Pollen and Simon (2005) developed the Fiber
78 Bundle Model (FBM) due to the significant overestimation of root reinforcement by W&W model. According to the
79 FBM, all roots assumed to be parallel and have similar elastic properties. When each root breaks, the load is
80 continuously redistributed over the remaining roots until the entire bundle are broken. Besides the already mentioned,
81 the Root Bundle Model Weibull (RBMw) was an interesting improvement since it provides the root reinforcement in
82 function of the observed root distribution at hillslope scales and of the displacement due to a potential triggering
83 mechanism at hillslope scales (Schwarz et al. 2013). RBMw has been used in different environments across the world,
84 including temperate forests of Norway spruce (*Picea abies* (L.) H. Karst.; Bischetti et al. 2007; Schwarz et al. 2013,
85 2015; Vergani et al. 2014; Moos et al. 2016; Cohen and Schwarz 2017; Cislighi et al. 2021), jolcham oak (*Quercus*
86 *serrate* Murray; Yamase et al. 2021), Monterey Pine (*Pinus radiata* D. Don; Giadrossich et al. 2020), black locust
87 (*Robinia pseudoacacia* L.; Zydroń et al. 2019; Zydroń and Gruchot 2021), black poplar (*Populus nigr* L.; Zydroń et

88 al. 2019), common hornbeam (*Carpinus betulus* L.; Zydroń and Gruchot 2021), green alder (*Alnus viridis* (Chaix)
89 D.C.), red willow (*Salix purpurea* L.), goat willow (*Salix caprea* L.), hazel (*Corylus avellana* L.), European ash
90 (*Fraxinus excelsior* L.), and European larch (*Larix decidua* Mill.; Bischetti et al. 2007), silver birch (*Betula pendula*
91 Roth), small-leaved lime (*Tilia cordata* Mill.), English oak (*Quercus robur* L.), and Sweet Cherry (*Prunus avium* L.;
92 Zydroń and Gruchot 2021), sweet chestnut (*Castanea sativa* Mill.; Dazio et al. 2018; Cislaghi et al. 2021), European
93 beech (*Fagus sylvatica* L.; Bischetti et al. 2007; Gehring et al. 2019; Cislaghi et al. 2021), Scots pine (*Pinus sylvestris*
94 L.; Vergani et al. 2017b), and subtropical forest for the white mangrove (*Avicennia marina* (Forssk.) Vierh.; Karimi et
95 al. 2022).

96 In this context, although it is evident the efforts of scientific community to investigate the implications of forests
97 on slope stability around the world, several scientific gaps remain for some mountainous areas, often prone to
98 landslides, such as the Hyrcanian temperate forests. In this environment, few studies provided useful advances for
99 implementing nature-based solutions such as the forests, to reduce the negative impacts of the shallow landslides. Abdi
100 et al. (2010a) observed that RAR of oriental beech (*Fagus orientalis* Lipsky.), Persian ironwood (*Parrotia persica*
101 (DC.) C.A.Mey.) and *Carpinus betulus* trees decreased with the depth and the maximum RAR values in the upper soil
102 layers. For three pioneer species Caucasian alder (*Alnus subcordata* C.A.Mey), velvet maple (*Acer velutinum* Boiss.),
103 and *Parrotia persica*, Abdi and Deljouei (2019) showed that in shallower depths of the soil, RAR was higher, and in
104 profiles nearer trees, they showed significantly higher RAR than in far trenches. Deljouei et al. (2020) explored the
105 most important parameters that affect fine roots resistance of two common temperate species (*F. orientalis* and *C.*
106 *betulus*) in the Hyrcanian forest, finding that tree species and DBH make a significant difference in fine roots
107 resistance. Despite, the effect of variability in soil profiles, tree DBH, altitude, and slope positions on the spatial
108 variability of roots and root reinforcement is still underexplored, with only a couple of studies on the topic (Moos et
109 al. 2016; Cislaghi et al. 2021). For these reasons, the present study focuses on spatial and mechanical characteristics
110 of the root systems of the dominant species (*F. orientalis* and *C. betulus*) in the Hyrcanian forest. In such environment,
111 *F. orientalis* forests account for approximately 30% of the standing volume and 23.6% of the stem number, covered
112 large areas at altitude from 300 to 2000 m a.s.l (Sagheb-Talebi et al. 2014). Meanwhile, *C. betulus* species accounts
113 for 30.5% of the standing volume and 30% of the stem number and can be found at altitude from 100 to 1500 m a.s.l
114 in the Hyrcanian forests (Sagheb-Talebi et al. 2014). Both species are representative and valuable study cases not only
115 they cover a vast area of temperate forests in the world but also these species are often located on slopes of mountainous
116 areas where the protection function of forests is significantly important to reduce the landslide susceptibility in
117 proximity of infrastructures and villages.

118 Recently, a lot of importance has been given to the implications of slope stability and evaluating the potential of trees
119 in stabilizing steep slopes. Some critical challenges in setting a safety factor are understanding how vegetation
120 enhances slope stability and quantifying vegetation's contribution to soil shear strength. The ability to evaluate
121 different areas in terms of their susceptibility to shallow landslides as well as the potential of local trees in stabilizing
122 the hillslopes enables optimization in forestry practices. To the authors' best knowledge, no studies modelling root
123 reinforcement of *C. betulus* and *F. orientalis* in function of DBH, slope position, altitude, soil depth, and distance from
124 the tree trunk. Hence, the main aims of the study consist in: (i) investigating the spatial (root distribution) and
125 mechanical variability of root systems, and (ii) modelling root reinforcement by Root Bundle Model Weibull (RBMw)

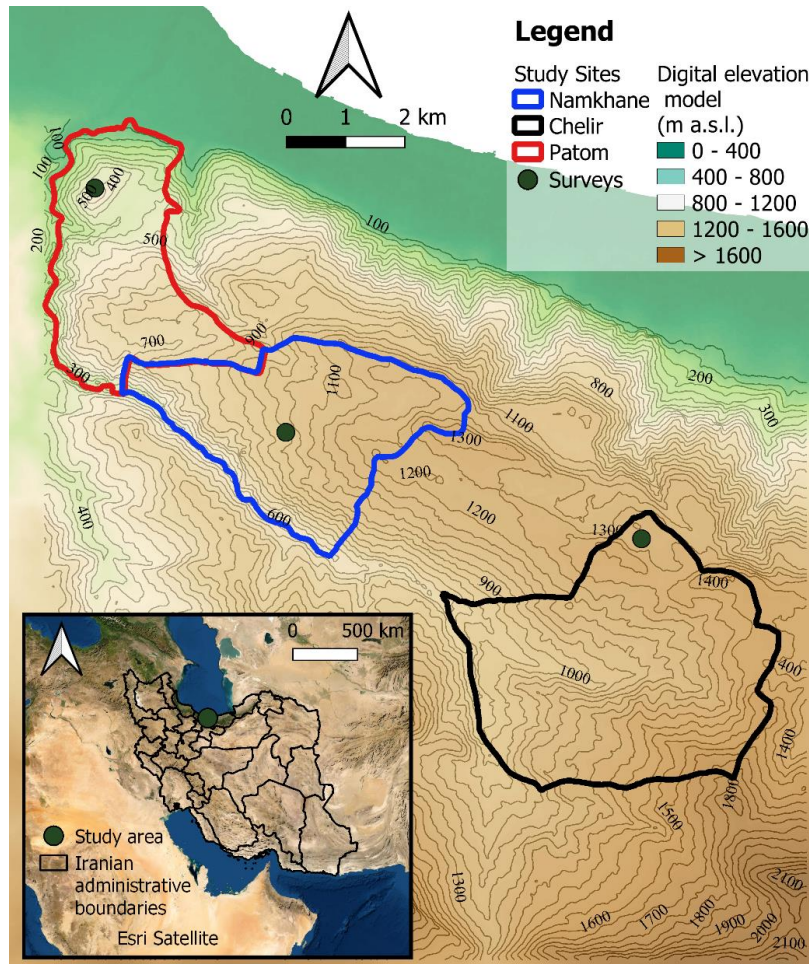
126 of two hardwood species in the Hyrcanian forest; (iii) providing a simplified framework for evaluating the effects of
127 trees in terms of slope stabilization. In addition, the study conducted a statistical analysis of the similarities and
128 differences between two main species of Hyrcanian temperate forests and several dendrometric characteristics,
129 altitude, and spatial position on root reinforcement, which forest managers must consider to mitigate shallow
130 landslides.

131 2. Material and Methods

132 2.1. Study site

133 Iran is one of the landslide-prone countries due to its specific geologic, morphologic, climatic, and tectonic conditions,
134 where most landslides are concentrated on the rim of Alborz Mountains. Approximately, 2600 landslides occurred in
135 the year 2000 in the country, which caused 162 deaths, destruction of 176 houses, and damages to 170 roads
136 (Abbaszadeh Shahri and Maghsoudi 2021). One of the regions of Iran in which landslides occur frequently is the
137 Hyrcanian ecoregion, particularly in such circumstances in which the vegetation has been cleared to construct roads.
138 Hyrcanian forests are classified as hilly and mountainous temperate forests, forming a green belt over the northern
139 slopes of the Alborz Mountain. They cover the southern coast of the Caspian Sea spreading on about 1.9 million
140 hectares. The area is abundant in hardwood species, including approximately 50 trees and 80 shrub species. Dominant
141 species are the *Carpinus betulus*, *Fagus orientalis*, *Parrotia persica*, Cappadocian maple (*Acer cappadocicum* Gled.),
142 *Acer velutinum*, common alder (*Alnus glutinosa* (L.) Gaertn), wych elm (*Ulmus glabra* Huds.), and the chestnut-leaved
143 oak (*Quercus castaneifolia* C.A.Mey.). Hyrcanian forests are used for wood production, tourism, environmental
144 protection and supportive services. These forests play an important role in conserving soil and water resources on steep
145 mountain slopes that are vulnerable to landslide. One of the severe problems in Hyrcanian forests is slope failure and
146 shallow landslides, because bare soils are vulnerable during intensive rainstorms, specifically where the trees have
147 been clear cut to make space for forest roads (Abdi et al. 2010a). In this ecoregion, landslides often cause economic
148 losses, property damages and high maintenance costs, as well as injuries or mortality (Pourghasemi et al. 2012).

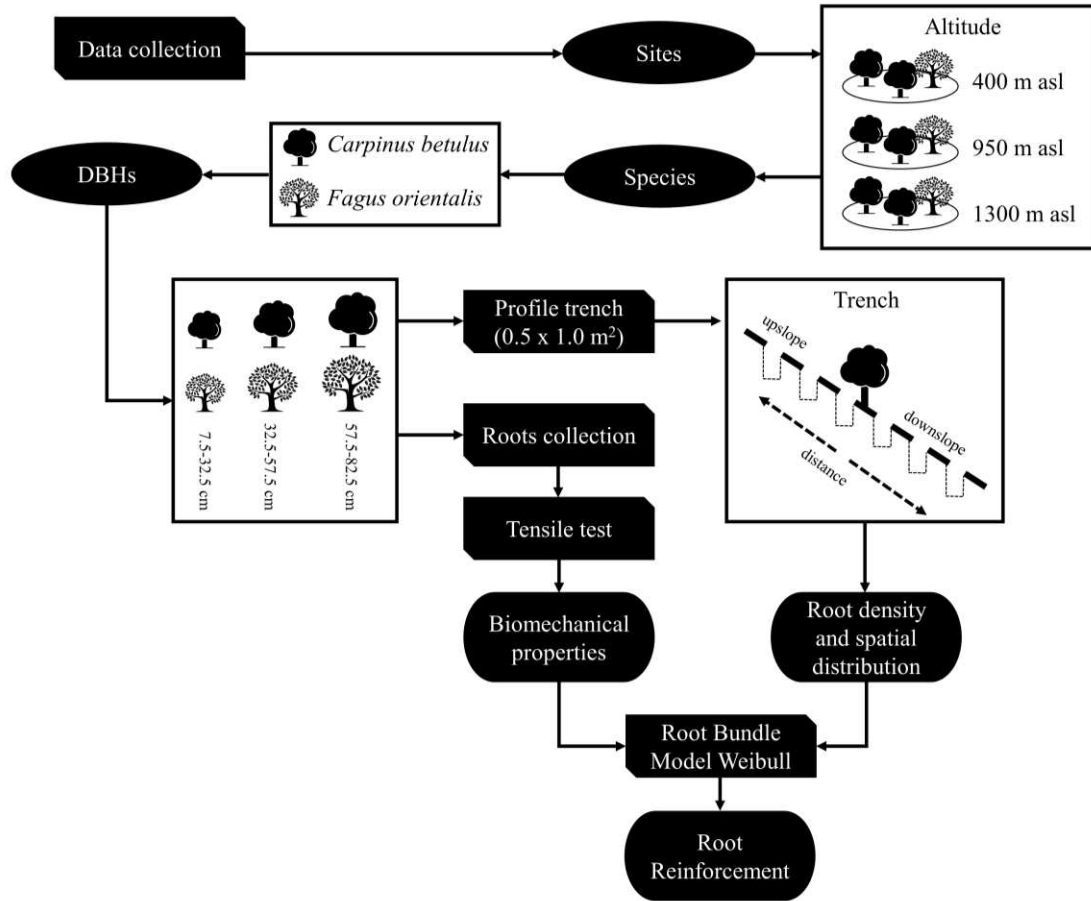
149 Kheyroud Forest, which covers an area of ~8000 ha, was selected as the study location (Fig. 1). The climate of the area
150 is humid, and the temperature fluctuations are relatively limited. The average annual precipitation is 1300 mm, falling
151 mainly as rain. The mean summer and winter temperatures are estimated to be 25.1 and 7.1 °C, respectively. Field
152 sampling was carried out in three districts of the Kheyroud Forest (Fig. 1), namely Patom, Namkhane, and Chelir.
153 Altitude of the study sites ranges from 400 m a.s.l. in Patom, with the highest mean temperature and lowest annual
154 precipitation, to 950 m a.s.l. in Namkhane, and 1300 m a.s.l. in Chelir, the latter exhibiting the lowest mean temperature
155 and the highest yearly precipitation. According to the unified soil classification system, the soils on the three study
156 sites were clays with high plasticity (i.e., CH). The mean values (\pm SD) of the Atterberg limits of the soils (soil liquid
157 limit (Casagrande cup method), soil plastic limit (rolling and thread method), and soil plasticity index) from the study
158 sites were estimated at 65% (\pm 6.2%), 26.4% (\pm 3.1%), 38.6% (\pm 3.8%) in Patom district, 88.5% (\pm 7.4%), 38.3% (\pm
159 4.9%), 50.2% (\pm 4.6%) in Namkhane district, and 85.7% (\pm 6.9%), 37.7% (\pm 3.7%), 48.0% (\pm 5.0%) in Chelir district,
160 respectively.



161
 162 **Fig. 1** Study location: map of Iran showing the general location of the study and districts taken into study

163 **2.2. Measuring root distribution**

164 Root distribution was measured for 5 sample trees of each of the two investigated species (*C. betulus* and *F. orientalis*),
 165 for each study site (Patom, Namkhane, and Chelir) at altitude of 400, 950, and 1300 m a.s.l., respectively, and for each
 166 DBH class (small = 7.5-32.5 cm, medium = 32.5-57.5 cm, and large = 57.5-82.5 cm). Hence, 90 trees were randomly
 167 selected (3 altitudes × 3 DBH classes × 2 species × 5 trees) and used as a sampling reference in this study. Six trenches
 168 with a width of 0.5 m and a length of 1 m were excavated manually to the maximum rooting depth (1 m soil depth);
 169 located on the downslope and upslope at distances of 1, 1.5, 2, 2.5, 3.5, and 4 m from the tree (Fig. S1). The profile
 170 trenching method was used to characterize the root distribution (Böhm 1979; Fig. S1). Layers of 10 cm were marked
 171 on the vertical profile walls using pins and string (Fig. S1). The number of roots, diameter, and maximum depth were
 172 measured in both downslope and upslope trenches. The diameters of roots intersecting the soil profile were measured
 173 with a digital calliper. Based on their diameter, the roots were included in four classes, namely fine roots (0-2 mm),
 174 small roots (2-5 mm), medium roots (5-10 mm), and large roots (>10 mm). The field measurements lasted between
 175 August and October 2016. All the steps of collecting data are shown in Fig. 2.



176

177 **Fig. 2** Flowchart describing the steps for quantifying root reinforcement: selecting study sites at different altitudes (400
 178 m, 950 m, 1300 m a.s.l), selecting *Carpinus betulus* and *Fagus orientalis* samples with different DBH classes (small:
 179 7.5-32.5 cm, medium: 32.5-57.5 cm, large: 57.5-82.5), evaluating root density and spatial distribution at downslope
 180 and upslope at six distances from tree trunk (1, 1.5, 2, 2.5, 3.5, and 4 m), investigating roots biomechanical properties
 181 and quantifying root reinforcement by Roor Bundle Model Weibull

182 **2.3. Measuring biomechanical properties**

183 Root samples were collected randomly from downslope and upslope at a depth of about 30 cm from the surface (Mao
 184 et al. 2012). To prevent mould and microbial degradation, a 15% alcohol solution was sprayed on the roots, then the
 185 treated roots were placed into plastic bags and refrigerated (4 °C) until tested (time between sampling and testing in
 186 the laboratory was of about 48 h) (Vergani et al. 2012). Roots with a length of 15 cm were placed in the clamps of the
 187 Universal Testing Machine (SANTAM Co./SMT-5, Tehran, Iran), and mechanical tests were conducted at a speed of
 188 10 mm min⁻¹ until rupture occurred. **Only specimens that broke near the middle of the root segment were considered.**
 189 Then, the relationships between the root diameter and biomechanical properties (i.e., maximum tensile force, Young's
 190 modulus, maximum elongation) were calculated through power-law functions, as follows:

$$F_{\max} = F_0 \phi^\xi \tag{1}$$

$$E = E_0 \phi^\beta \quad (2)$$

$$L = L_0 \phi^\alpha \quad (3)$$

191 where F_{max} is maximum tensile force (in N), ϕ is root diameter (in mm), E is root elasticity (MPa), and L is root
 192 elongation (in mm). F_0 , E_0 , and L_0 are constant coefficients (in N, MPa, and mm, respectively), ξ , β , and α are exponents
 193 (dimensionless).

194 2.4. Root Bundle Model weibull (RBMw)

195 Root reinforcement was calculated using the Root Bundle Model Weibull (RBMw; Schwarz et al. 2013). RBMw is a
 196 strain-step fiber bundle model, developed to include the failure probability of roots due to variability in root mechanical
 197 properties. RBMw calculates force-displacement behaviour of a root bundle based on root distribution in diameter
 198 classes and on a series of power-distributed relationships (Eqs. 1-3). Root reinforcement c_r (in kPa) is calculated by
 199 summing up the force contributions F (in N) for each root per unit of area (m^2) multiplied by the Weibull survival
 200 function S (dimensionless), as follows:

$$c_r = \sum_{i=1}^N F(\phi_i, \Delta x) S(\Delta x^*) \quad (4)$$

201 Where Δx is the displacement unit in mm and S is a function of the normalized displacement Δx^* (dimensionless). The
 202 following equation calculates the $S(\Delta x^*)$:

$$S(\Delta x^*) = \exp \left[- \left(\frac{\Delta x^*}{\lambda} \right)^\omega \right] \quad (5)$$

203 In equation (5), λ is the scale Weibull parameter (dimensionless) and ω is the shape Weibull parameter (dimensionless).
 204 The ratio between the displacements is estimated by each single tensile tests and the corresponding displacement values
 205 are calculated using fitted values of tensile forces.

$$F(\phi_i, \Delta x) = \frac{\pi E_0}{4 L_0} \phi_i^{2+\beta-\alpha} \quad F(\phi_i, \Delta x) < F_{max}(\phi_i) \quad (6)$$

206 All input parameters (F_0 , E_0 , L_0 , ξ , β , and α) were calculated from the tensile tests.

207 2.5. Evaluation of the hillslopes stability using a Monte Carlo approach

208 The majority of distributed slope-stability models that predict shallow landslides overlook the forces acting on
 209 landslide scars. Based on back calculations and observations in the field, most shallow landslides are controlled by
 210 lateral root strength (Casadei et al. 2003). The infinite slope analysis is a rather simplistic and well-known method,
 211 being used to assess slope stability; however, it assumes that the plane of failure is parallel to the slope (Taylor 1948).

212 In comparison, the physical-based 3D slope stability approach overcomes the limitations of infinite slope analysis and
 213 incorporates the root reinforcement effectively. By applying the limit equilibrium theory, slope stability can be
 214 expressed using the Factor of Safety (FoS) as proposed by Casadei et al. (2003) and modified by Chiaradia et al.
 215 (2016), equation 7:

$$FoS = \frac{S}{T} = \frac{c_s + c_{r-b} + \frac{A_l}{A_b} c_s + \frac{A_l}{A_b} c_{r-l} + \{[(D - D_w)\gamma_s + (\gamma_{sat} - \gamma_w)D_w + q_0] \cos \alpha\} \tan \varphi'}{[(D - D_w)\gamma_s + \gamma_{sat}D_w + q_0] \sin \alpha} \quad (7)$$

216 where S is the resisting force, T is the shear force, c_s is the soil cohesion (kPa), c_{r-b} and c_{r-l} are the contributions of plant
 217 roots to slope stability (root reinforcement) along the basal and lateral surfaces (in kPa), A_b and A_l are basal and lateral
 218 area of the sliding volume (in m²), D is the average depth of the sliding surface (in m), γ_s is the unit weight of dry soil,
 219 D_w is the average height of the seepage in respect to the sliding surface, γ_w is the unit weight of water, q_0 is the tree
 220 surcharge per unit area (in kPa), α is the slope steepness, φ' is the effective friction angle (°), and γ_{sat} is the unit weight
 221 of saturated soil.

222 Based on a mathematical formula, Monte Carlo Simulation is an effective method for calculating distributions for the
 223 FoS. It identifies independent and random sets of possible values for each input parameter to determine the FoS for
 224 each pass. The Monte Carlo approach was applied to overcome the variability and uncertainties of each input
 225 parameter. The slope angle varies from 20 ° to 60 °. The soil bulk density is normally distributed around 13.5 kNm⁻³
 226 (Abdi and Deljouei 2019). The internal friction angle is uniformly distributed between 26 ° and 35 °, and the soil
 227 cohesion is normally set around the average value of 15.3 kPa (Abdi and Deljouei 2019). The A_l/A_b ratio can be
 228 approximated by a value of 0.50 (Milledge et al. 2014). Trees surcharge for different DBH classes was considered as
 229 140 Nm⁻², 220 Nm⁻², and 320 Nm⁻² for small, medium and large *C. betulus* trees, and corresponding values for *F.*
 230 *orientalis* were 100 Nm⁻², 230 Nm⁻², and 400 Nm⁻², respectively (Chiaradia et al. 2016; Hayati et al. 2017). C_{r-l} is the
 231 RBMw-calculated root reinforcement in function of the most critical conditions i.e. distance of 4 m and upslope
 232 position. The mean, minimum, and maximum values of C_{r-l} are 3.46, 0.06, and 53.24 kPa, respectively. C_{r-b} is a
 233 percentage of C_{r-l} in function of RAR and soil depth which depends on slip failure. To summarise, the parameters for
 234 Monte Carlo Simulation are reported in Table 1. For this analysis, Monte Carlo simulation is replicated 1000 times;
 235 furthermore, to reduce the effects of random selection (Hammond et al. 1992).

236 Table 1. The parameters for Monte Carlo Simulation in function of species, altitude and DBH classes, considering the
 237 upslope position and the far distance from the trunk (i.e. 4 m). The parameters were specified using a range and the
 238 distribution function.

Parameter	Unit	Range	Distribution function	Reference
A_b	m ²	$\mu=104$ $\sigma=2.63$	Normal	Milledge et al. 2014
L/W^*	-	$\mu=1.42$ $\sigma=0.20$	Normal	Milledge et al. 2014
D	m	[0.5; 1.5]	Uniform	Hammond et al. 1992
D_w/D	-	$\mu=1.42$ $\sigma=0.20$	Normal	Schwarz et al. 2010

γ_s	kN/m ³	$\mu=13.5$ $\sigma=1.00$	Normal	Abdi and Deljouei 2019
ϕ'	°	[26; 35]	Uniform	Abdi and Deljouei 2019
c_s	kPa	$\mu=13.5$ $\sigma=2.00$	Normal	Abdi and Deljouei 2019
ω_c **	-	$\mu=0.20$ $\sigma=0.01$	Normal	Abdi and Deljouei 2019

239 $w = \sqrt{A_b \cdot \frac{w}{L}}$

240 $\gamma_{sat} = \gamma_s(1 + \omega_c)$

241 2.6. Statistical analysis

242 Statistical analysis was used to check the differences in root distribution and root reinforcement by considering the
 243 species, DBH, slope position, **altitude**, and distance from the trees. The Shapiro–Wilk and Levene’s tests were used to
 244 check the normality and homogeneity of the data, respectively. Since the datasets were found to violate the normality
 245 and homogeneity assumptions, a nonparametric Kruskal-Wallis test (H) was used to compare the RAR and soil
 246 reinforcement of different root diameter classes within DBH classes, slope positions, and study sites for *C.*
 247 *betulus* and *F. orientalis* trees (Tables S1-S5). When the residuals were normal and variance was heterogeneous,
 248 parametric Welch t-tests were used to compare between two independent groups. Finally, when the residuals were
 249 normal and variance was homogenous, One-way ANOVA (analysis of variance) was used to compare the means of
 250 two or more groups for one dependent variable. All statistical analyses were implemented using the R software
 251 (<https://www.r-project.org>). Confidence intervals were set for a probability level of 0.05.

252 3. Results

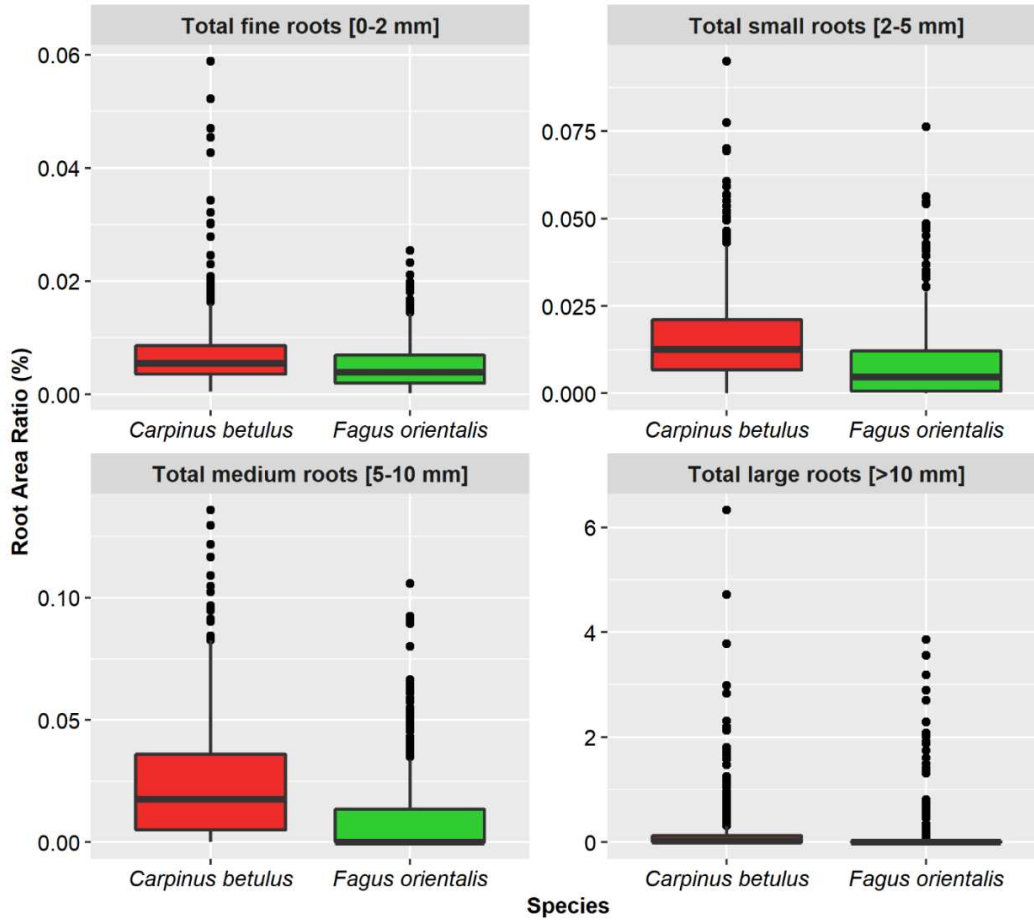
253 3.1. Variability of root distribution

254 3.1.1. Root distribution as a function of species

255 RAR was measured by considering a number of 1080 profile trenches. Figure 3 showed the total RAR for fine (0-2
 256 mm), small (2-5mm), medium (5-10mm), and large roots (>10mm) as a function of species. Root distributions were
 257 remarkably different in regard to the diameter classes. As a fact, for *C. betulus*, small roots had the highest RAR value
 258 whereas the value of large was the lowest among all the root diameter classes (Fig. 3). Furthermore, fine roots of *F.*
 259 *orientalis* had a higher frequency compared with that from the rest of diameter classes (Fig. 3). The Kruskal-Wallis
 260 test indicated that RAR values of *C. betulus* was significantly higher than those of *F. orientalis* ($H_1 = 65.13$, $H_1 =$
 261 140.65 , $H_1 = 177.01$, $H_1 = 117.44$; $p < 2.2e-16$ to $p < 1e-15$; for fine, small, medium, and large roots, respectively; Fig.
 262 3 and Table S2). The total RAR ($H_1 = 191.37$, $p < 2.2e-16$; Fig. 4 and Table S2) and **total roots per unit area** ($H_1 =$
 263 99.60 , $p < 2.2e-16$; Fig. 4 and Table S2) of *C. betulus* and *F. orientalis* demonstrated a significant difference, in which
 264 *F. orientalis* had fewer roots. The minimum, maximum and mean value of RAR for *C. betulus* were of 0.0010%,
 265 0.0040%, and 0.0020%, respectively, while for *F. orientalis* the same statistics accounted for 0.0005%, 0.0020%, and

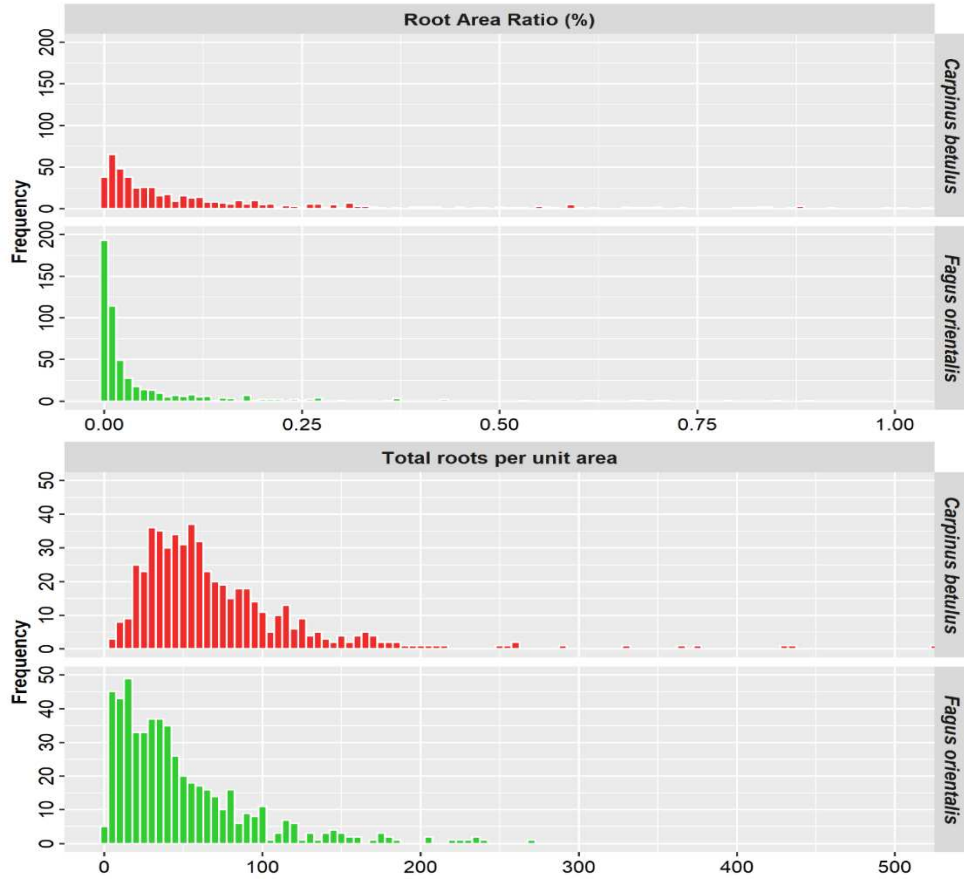
266 0.0010%, respectively. The results showed that *C. betulus* had more roots in all root diameter classes, in which the
267 total number of roots [per unit area](#) for *C. betulus* was 38987 and for *F. orientalis* was 26079 (Fig. 4).

268



269

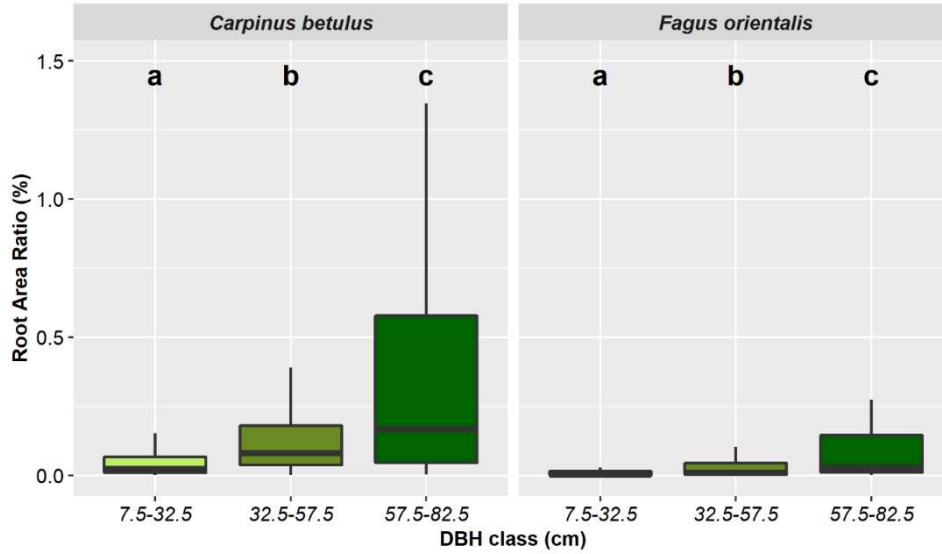
270 **Fig. 3** Root Area Ratio (RAR, in %) of fine roots (0-2 mm), small roots (2-5 mm), medium roots (5-10 mm), and
271 large roots (>10 mm) as a function of species: *Carpinus betulus* and *Fagus orientalis*



272
 273 **Fig. 4** Total Root Area Ratio (RAR, in %) and total roots per unit area as a function of tree species: *Carpinus betulus*
 274 and *Fagus orientalis*

275 **3.1.2. Root distribution as a function of DBH**

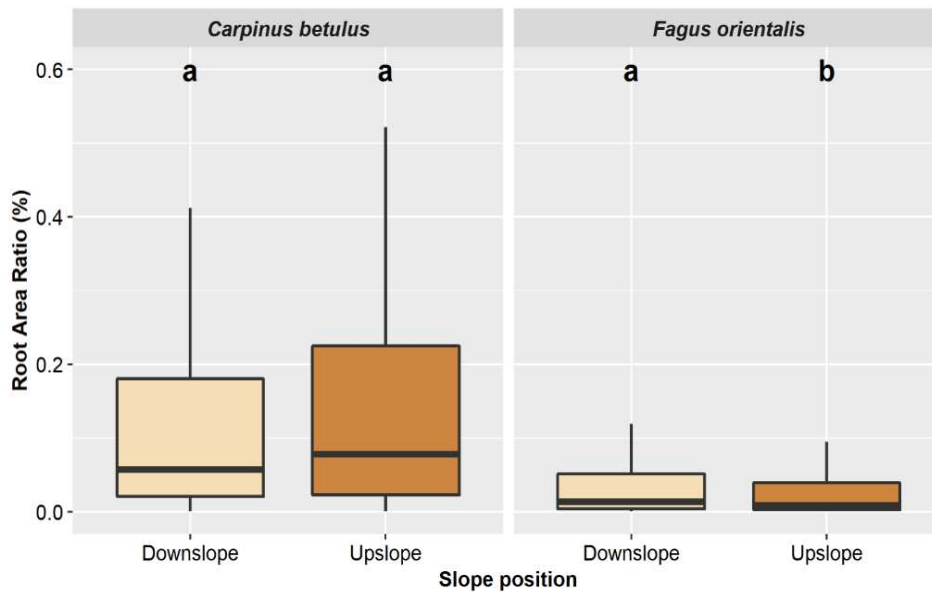
276 Kruskal-Wallis test was conducted to verify the differences in RAR as a function of DBH for *C. betulus* ($H_2 = 111.01$)
 277 and *F. orientalis* ($H_2 = 112.80$) in which significant differences were found to be caused by DBH classes of both
 278 species ($p < 2.2e-16$; Table S3). On average, the largest DBH tree class of both species had the highest RAR value
 279 compared to the other DBH classes (Fig. 5). The mean values of RAR for large trees was 0.0050% and 0.0030%, for
 280 *C. betulus* and *F. orientalis*, respectively (Fig. 5). Mean values of RAR for medium and small trees of *C. betulus* and
 281 *F. orientalis* were of 0.0010% and 0.0004%, 0.0006% and 0.0001%, respectively (Fig. 5).



282
 283 **Fig. 5** Root Area Ratio (RAR in %) for *Carpinus betulus* and *Fagus orientalis* as a function of DBH class: Small (7.5-
 284 32.5 cm), Medium (32.5-57.5 cm), and Large (57.5-82.5 cm). Boxes with the similar lowercase letters are not
 285 significantly different

286 **3.1.3. Root distribution as a function of slope position**

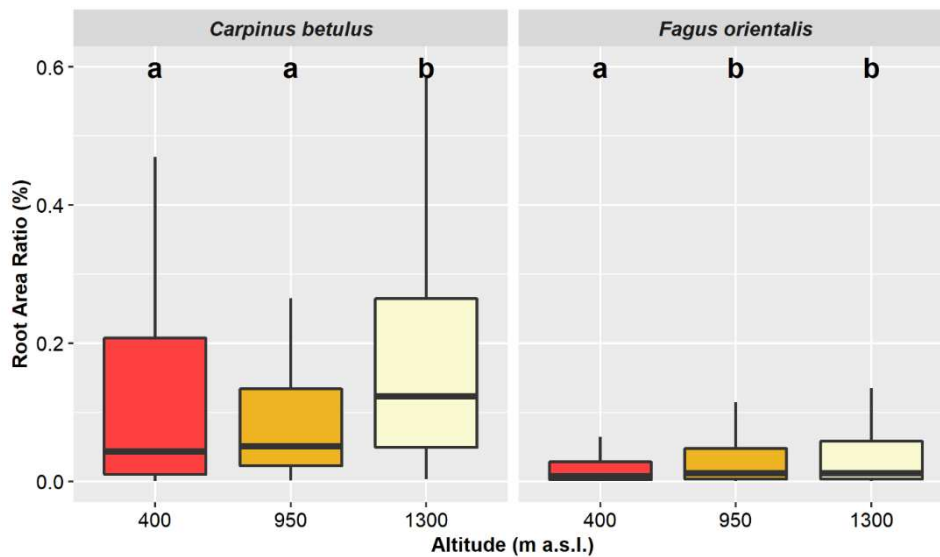
287 A synthesis of the RAR as a function of slope position is shown in Figure 5. RAR values of downslope and upslope
 288 were not significantly different for *C. betulus* ($H_1 = 2.86$, $p = 0.09$; Table S4). However, while the RAR values of *F.*
 289 *orientalis* were significantly higher for downslope ($H_1 = 7.01$, $p = 0.01$; Table S4). The mean value of RAR for
 290 downslope and upslope of *C. betulus* was estimated to be 0.0020%, while the mean value of RAR for *F. orientalis* was
 291 of 0.0020% and 0.0007% for downslope and upslope, respectively (Fig. 6).



293 **Fig. 6** Root Area Ratio (RAR in %) for *Carpinus betulus* and *Fagus orientalis* in function of slope position: Downslope
294 and Upslope. Boxes with the similar lowercase letters are not significantly different

295 3.1.4. Root distribution as a function of altitude

296 RAR values varied among altitudes: 400, 950, and 1300 m of *C. betulus* ($H_2 = 32.73$, $p < 1e-7$; Table S5) and *F.*
297 *orientalis* ($H_2 = 6.46$, $p = 0.04$; Table S5). Figure 7 showed the statistical differences brought by the altitude on the RAR
298 of the two species. For *C. betulus*, the highest mean value of RAR was found in 1300 m (0.0020%) and it was followed
299 by 950 m (0.0010%) and 400 m (0.0030%). For *F. orientalis*, RAR values in 1300 and 950 m were larger than in 400
300 m (Fig. 7). According to the altitude, mean RAR for *F. orientalis* was reported as 0.0010% for 1300 m and 950 m;
301 furthermore, it was recorded as 0.0009% for 400 m (Fig. 7).

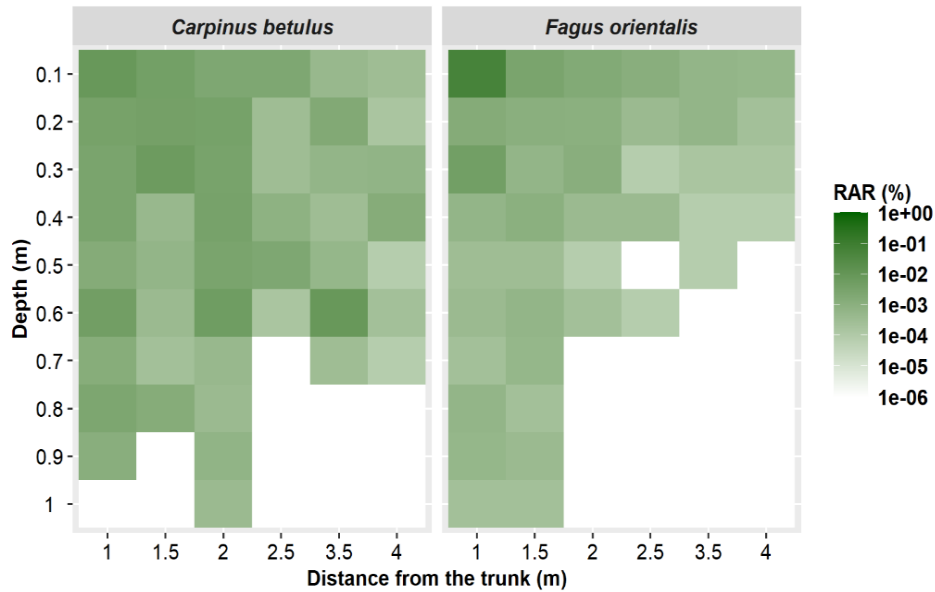


302
303 **Fig. 7** Root Area Ratio (RAR in %) for *Carpinus betulus* and *Fagus orientalis* as a function of altitude: 400, 950, and
304 1300 m. Boxes with the similar lowercase letters are not significantly different

305 3.1.5. Root distribution as a function of vertical and horizontal directions

306 In the case of both species, the predominant number of roots was found in the subsurface soil layers to a depth of up
307 to 30 cm after which the root frequency decreased (Fig. 8). For both *C. betulus* and *F. orientalis*, RAR decreased
308 monotonically with distance to tree beyond a 2 m (Fig. 8). The highest RAR values were found closest to the tree trunk
309 with greater concentrations at 1-1.5 m from the tree trunk for both species (Fig. 8). The mean RAR at 1 m distance
310 was calculated 0.07% and 0.06% for *C. betulus* and *F. orientalis*, respectively. It was recorded that mean RAR at 1.5
311 m distance was 0.03% for *C. betulus* and 0.01% for *F. orientalis* (Fig. 8). Indeed, roots were detected for the 49.21%
312 at the 1 m distance for *C. betulus* and 82.93% for *F. orientalis*. Only 20.80% and 10.76% of roots were distributed at
313 1.5 m distance for *C. betulus* and *F. orientalis*. It was noted that mean values of RAR at the 0-10 cm soil depth were
314 0.11 and 0.08% for *C. betulus* and *F. orientalis*, respectively (Fig. 8). In the case of 10-20 cm soil depth, it was

315 estimated 0.06% for *C. betulus* and 0.02% for *F. orientalis* (Fig. 8). Approximately, in the case of *C. betulus*, 50% of
 316 roots distributed at the 0-10 cm soil depth and 69% for *F. orientalis*. Furthermore, at a depth of 10-20 cm, root
 317 distribution was reported 25.55% and 19.77% for *C. betulus* and *F. orientalis*, respectively. Overall, root distribution
 318 decreases with increasing soil depth and distance from the tree trunk. Moreover, root distribution was found higher on
 319 *C. betulus* than *F. orientalis*.

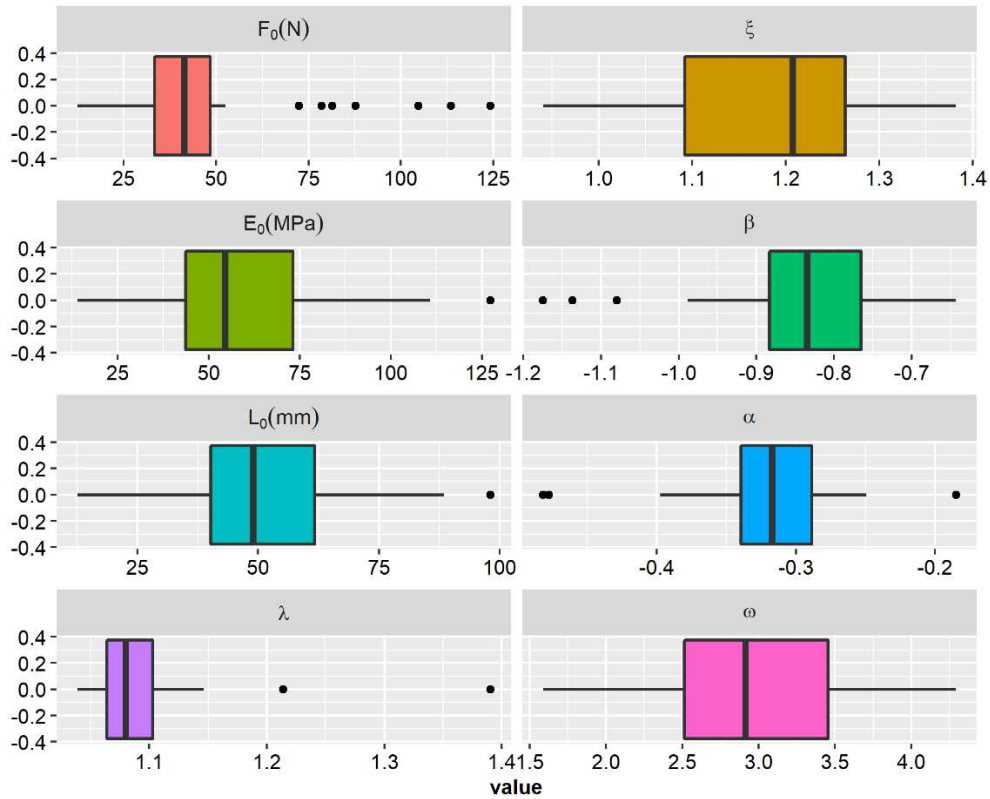


320
 321 **Fig. 8** Spatial distribution of Root Area Ratio (RAR in %) as a function of vertical and horizontal direction of *Carpinus*
 322 *betulus* and *Fagus orientalis*

323 **3.2. Variability of the mechanical properties of roots**

324 **3.2.1. Variability of RBMw parameters**

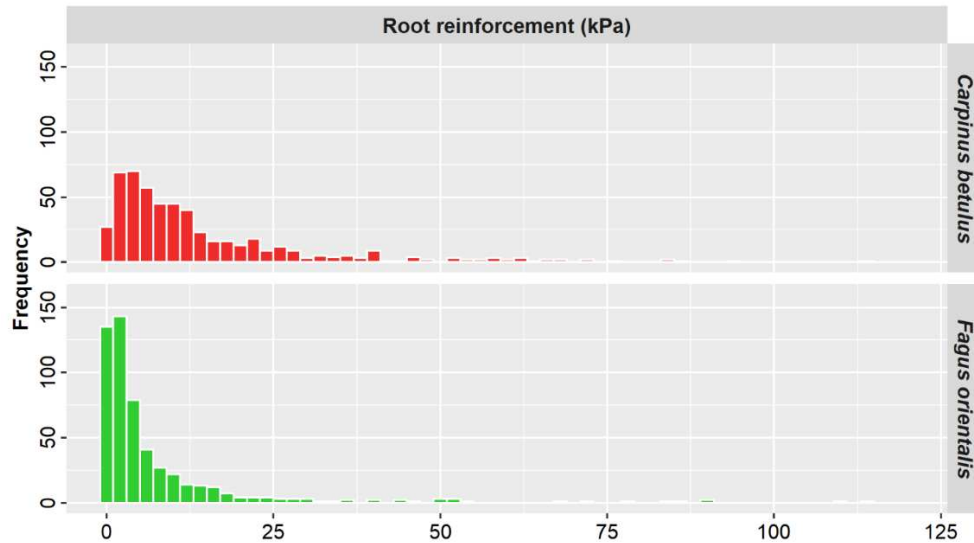
325 The main input parameters, including the relationship between root tensile force and root diameter, the regression
 326 coefficients for Young's modulus, root elongation, and the Weibull survival function are shown in Figures S2, S3, S4,
 327 S5, S6, S7, and Table S7. The results of the root tensile tests indicated strong relations between the mechanical and
 328 geometrical characteristics of the roots (root diameter) by power-law regression (Figures S2- S7). **Variability of Root**
 329 **Bundle Model Weibull (RBMw) parameters including constant coefficients F_0 , E_0 , L_0 and exponents ξ , β , α , λ , ω**
 330 **coefficients were shown in Fig. 9. F_0 ranged between 12.41 and 124.18 N, E_0 ranged between 1.39E+07 and 1.27E+08**
 331 **MPa, and L_0 ranged between 12.56 and 98.07 mm (Fig. 9). The minimum values of ξ , β , α , λ , ω were 0.94, -1.17, -**
 332 **0.48, 1.04, and 1.59, respectively. The maximum values of these exponents were 1.38, -0.64, -0.18, 1.39, and 4.29,**
 333 **respectively (Fig. 9).**



334
 335 **Fig. 9** Variability of Root Bundle Model Weibull (RBMw) parameters: constant coefficients F_0 , E_0 , L_0 and exponents
 336 ξ , β , α , λ , ω coefficients (dimensionless)

337 **3.2.2. Species-specific root reinforcement using RBMw model**

338 The mean values of root reinforcement for *C. betulus* and *F. orientalis* were of 16.08 and 7.69 kPa, respectively. Figure
 339 10 showed that root reinforcement of *C. betulus* distributed more and higher than *F. orientalis*, whereas the Kruskal-
 340 Wallis test showed a statistically significant difference in root reinforcement between the species ($H_1 = 168.22$, $p <$
 341 $2.2e-16$; Table S6). The minimum and maximum values of root reinforcement for *C. betulus* calculated as 0.23 and
 342 216.95 kPa, respectively which is concentrated in shallower soil layers and nearer to tree trunk. Root reinforcement
 343 for *F. orientalis* varied from 0.07 to 145.39 kPa.

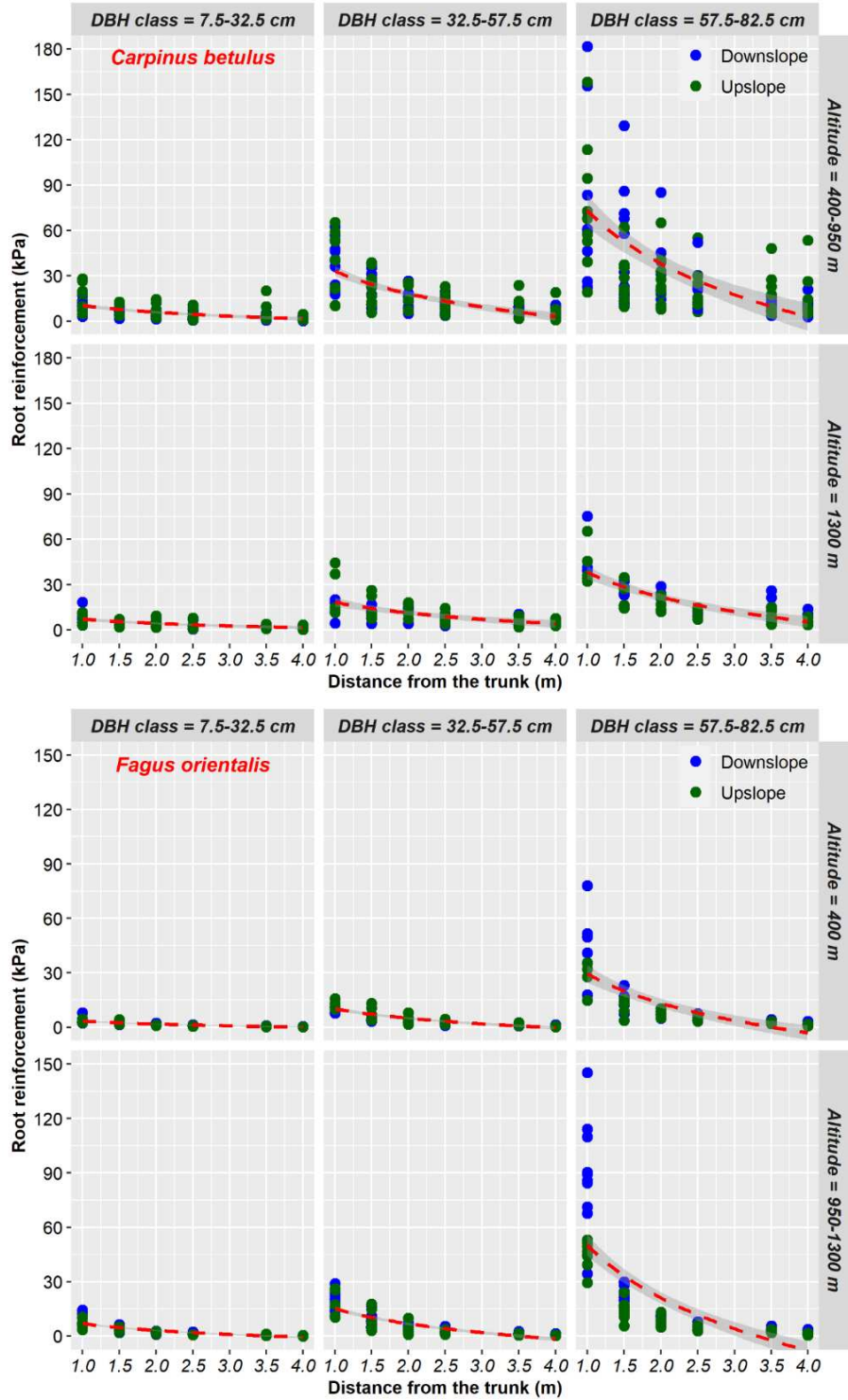


344

345 **Fig. 10** Root reinforcement in function of species (*Carpinus betulus* and *Fagus orientalis*)

346 **3.2.3. Impacts of environmental factors on RBMw-calculated root reinforcement**

347 Applying RBMw, the root reinforcement of *C. betulus* was the highest in 400 and 950 m compared with 1300 m. The
 348 mean values of c_r showed larger variation, including 23.89, 12.95, and 11.41 kPa at 400, 950, and 1300 m, respectively.
 349 Conversely, the mean value of c_r for *F. orientalis* reported 5.65, 9.58, and 7.85 kPa at the same altitudes, respectively.
 350 The results showed that c_r declined by decreasing DBH for both species (Fig. 11). Mean values of c_r reached 29.12,
 351 14.29, and 4.82 kPa for large, medium and small trees of *C. betulus*, respectively; however, for the same DBH classes,
 352 the mean values of c_r for *F. orientalis* were of 15.52, 5.30, and 2.26 kPa, respectively. Furthermore, the c_r decreased
 353 with increasing distance from the tree trunk (Fig. 11). The highest mean value of c_r for *C. betulus* was of 38.55 kPa at
 354 1 m distance from the tree, whereas it was of 26.99 kPa for *F. orientalis* at the same distance (Fig. 11). The lowest
 355 mean values of c_r were of 5.85 and 0.78 kPa, and they were found at distances of 4 m for *C. betulus* and *F. orientalis*,
 356 respectively (Fig. 11).

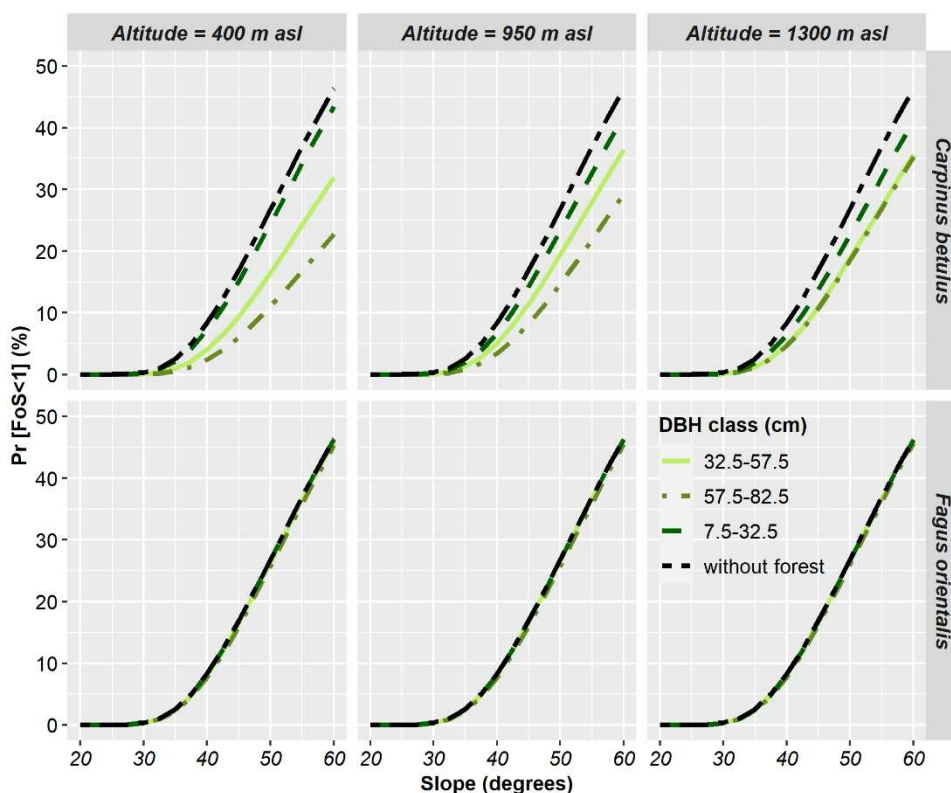


357

358 **Fig. 11** Root reinforcement of *Carpinus betulus* and *Fagus orientalis* as a function of DBH class (Small: 7.5-32.5 cm,
 359 Medium: 32.5-57.5 cm, and Large: 57.5-82.5 cm), altitudes (400, 950, and 1300 m), and distance from tree (1, 1.5, 2,
 360 2.5, 3.5, and 4 m)

361 **3.3. Slope stability analysis**

362 Including both basal and lateral reinforcement, the results of slope stability analysis (Eq. 7) suggested that the most
 363 stabilizing species is *C. betulus* in a mature growth (DBH= 57.5-82.5 cm) at the distance of 4 m. In fact, at 400, 950,
 364 and 1300 m a.s.l trees of *C. betulus* maintained an instability probability near 22.7%, 29.2%, and 35.3% for very steep
 365 conditions, respectively (Fig. 11). Instability probability for small trees at same altitudes varied from 40.9% to 43.3%.
 366 These values for medium trees of *C. betulus* reported between 31.9% and 35.6% (Fig. 12). In contrast, the instability
 367 probabilities of the *F. orientalis* was higher, reaching up to approximately 46.0% for all DBH classes at 400, 950, and
 368 1300 m a.s.l (Fig. 12). The instability probabilities was less than 10% until 40 ° for *F. orientalis* at the distance of 4 m
 369 (Fig. 12). In the areas without forest with very steep conditions instability probability for *C. betulus* and *F.*
 370 *orientalis* was 46.17% (Fig. 12).



371 **Fig. 12** Factor of safety (FoS), probability of failure (Pr(FoS < 1)), in function of species (*Carpinus betulus* and *Fagus*
 372 *orientalis*), altitude (400, 950, and 1300 m), and DBH classes (small: 7.5-32.5 cm, medium: 32.5-57.5 cm, and large:
 373 57.5-82.5 cm) considering the upslope position and the far distance from the trunk (i.e. 4 m). In all the case, the
 374 contribution to slope stability by the vegetation is evaluated as the sum of both basal and lateral root reinforcement
 375 values
 376 values

377 **4. Discussion**

378 **4.1. Root distribution**

379 The scientific literature often used RAR to quantify, first, root distribution into the soil and, second, the root
380 reinforcement. In this study, the root density, observed though total roots per unit area and RAR, was significantly
381 different in of the two investigated species. In fact, it was evident that root density of *C. betulus* was higher than that
382 of *F. orientalis*. In parallel to our findings, previous literature showed that root density is affected by tree species
383 (Bischetti et al. 2007; Phillips et al. 2014; Vergani et al. 2017a; Moresi et al. 2019; Gholami-Derami et al. 2021). For
384 example, Bischetti et al. (2007) found differences among European alpine species, including *Alnus viridis*, *Fagus*
385 *sylvatica*, *Salix purpurea*, *Salix caprea*, *Corylus avellana*, *Fraxinus excelsio*, *Picea abies*, and *Larix decidua* in which
386 root density varies significantly for the same species within a same locality. Gholami-Derami et al. (2021) compared
387 RAR of *C. betulus* with that of *Alnus subcordata* and among two non-native tree species (*Pinus sylvestris* and *Robinia*
388 *pseudoacacia*) with similar habitat conditions in Northern Iran. They found that RAR value in exotic species is higher
389 than native species. A study in New Zealand reported that willow roots were more numerous than poplar (Phillips et
390 al. 2014). Several reasons can relate to differences between species, above all else is heterogeneity in environmental
391 factors (Burylo et al. 2011), such as soil bulk density (Goodman and Ennos 1999), and soil moisture and fertility (Taub
392 and Goldberg 1996; Hodge 2004).

393 For both studied species, our results revealed that a tree with larger DBH has more roots than a tree with a smaller
394 diameter, and this finding is in line with previous studies (Bischetti et al. 2009; Schwarz et al. 2010; Mehtab et al.
395 2021), and is consistent with Abdi et al. (2010b) who showed that a significant effect of DBH on total RAR in three
396 hardwood species in Hyrcanian forests (*F. orientalis*, *C. betulus*, and *Parrotia persica*). In accordance, John et al.
397 (2001), comparing three stands of 6, 15, and 23 years old, showed that root distribution increases by tree age.
398 Additionally, it was reported that in the older stands, fine roots (0-2mm) declined as they were converted into coarse
399 roots (>10 mm) to provide more structural support. As a result, coarse root mass increased substantially compared to
400 fine roots (John et al. 2001). McQueen (1968) reported that fine roots peak at the early ages of the stands and are
401 relatively maintained constant after that. Our findings indicate that larger trees maintain their anchorage, with
402 increasing root numbers and RAR values. In fact, nutrient availability increases lateral root extension as well as causes
403 changes in lateral roots anatomy by supplying nutrients (Goss et al. 1993). Also, Ford and Deans (1977) reported a
404 higher concentration of fine roots because of increased nutrient availability. Furthermore, in this study, moisture
405 content has been highlighted as a key factor in the distribution of roots. So, greater root mass could be attributed to the
406 higher nutrient content and better aeration of the surface soil (Ford and Deans 1977). Overall, root growth can be
407 affected by a variety of environmental factors such as soil texture, soil structure, aeration, moisture, temperature, and
408 competition with other plants (Kramer and Boyer 1995).

409 The slope influences root distribution with the roots-oriented upslope, assisting soil anchorage (Vergani et al. 2017a),
410 as observed in this study for *F. orientalis* species. RAR distribution of *F. orientalis* in downslope is greater than
411 upslope; however, in the case of *C. betulus*, the RAR values of both slopes were similar. Mechanical function of the
412 root system in downslope and upslope is different based on the specific type of root system architecture (i.e.,
413 asymmetry of the cross-sectional area; Chiatante et al. 2003). It was stated that mechanical stress is one reason for
414 large cross-sectional areas (Di Iorio et al. 2005), therefore, the higher RAR value of *F. orientalis* in upslope is a kind
415 of adaptability in response to the environment. Burylo et al. (2011) reported that the heterogeneity in soil properties,
416 natural obstacles, and interactions between vegetation such as competition makes a high variability of RAR. Stokes et

417 al. (2008) found that this variability was related to the interactions of genetic and environmental factors. In the case of
418 *C. betulus*, RAR values of downslope and upslope were not significantly different, probably implying that trees are
419 thickening the roots instead of increasing the number in the reaction of the mechanical stresses (Abdi et al. 2010a).
420 Our results highlighted that RAR values fluctuate among different altitudes, and RAR in the 1300 m is higher than 950
421 and 400 m. One possible reason is the variation among the meteorological parameter and altitude above sea level.
422 These altitudes range from low altitude (i.e., 400 m a.s.l.) with the highest mean temperature and lowest annual
423 precipitation to mid- altitude (i.e., 950 m a.s.l) to the highest altitude (i.e., 1300 m a.s.l.) having the lowest mean air
424 temperature and greatest annual precipitation with the most proportion of snowfall (Azaryan et al. 2015; Deljouei et
425 al. 2020). Bischetti et al. (2009) showed that for *F. sylvatica*, RAR distributions were statistically more in a site with
426 highest altitude than lowest altitude (altitude ranges from 1100 to 1454 m a.s.l.). Mao et al. (2012) found similar results
427 for *P. abies*, *Abies alba*, and *F. sylvatica* growing at 1400 m a.s.l. and 1700 m a.s.l. The use of altitude gradients is
428 considered an excellent way to examine vegetation responses to environmental change (Sundqvist et al. 2013;
429 Weemstra et al. 2021). Higher altitudes in temperate regions typically have longer growing seasons and more
430 seasonality, and their vegetation is adapted to the extreme variations in climate they can experience (Körner 1999;
431 Sundqvist et al. 2013). In most cases, high-altitude soils are more heterogeneous in terms of soil nutrient availability
432 (Holtmeier and Broll, 2005) and less fertile (Sveinbjörnsson et al. 1995), since the cooler temperatures slow down
433 microbial activity (Loomis et al. 2006; Mayor et al. 2017), leaf and root litter decomposition rates (Moore 1986;
434 Loomis et al. 2006; See et al. 2019), and mineralization rates (Sveinbjörnsson et al. 1995). This means that changes in
435 climate and soil properties along an altitude gradient can profoundly influence intraspecific root trait variation
436 (Weemstra et al. 2021). Another significant factor is the distance from the trunk. In fact, root distribution decreases
437 with increasing distance from tree trunk and soil depth for both species. Seventy percent of *C. betulus* roots and 94%
438 *F. orientalis* roots were distributed at 1.5 m of distance from the tree trunk. Furthermore, the maximum RAR values
439 (50% of *C. betulus* roots and 69% of *F. orientalis* roots) were situated in the surface soil, i.e. 1-10 cm of soil depth and
440 approximately were smaller in deeper soil. Species or genetics, climate characteristics determined root distribution
441 throughout the soil profiles, and soil type (Bischetti et al. 2005); for instance, changes in nutrient content, water
442 availability and aeration will affect root distribution, whereas the most available nutrients are detected in the topsoil
443 and cause a reduction in vertical root distribution (Bischetti et al. 2005; Abdi et al. 2010a; Mao et al. 2012; Bordoni et
444 al. 2019; Moresi et al. 2019). It was pointed out that root development might depend mainly on the quantity of organic
445 matter in the soil (Bordoni et al. 2019). In addition, deeper layers due to compacted soil layers and bedrock caused the
446 roots to grow horizontally (Coppin and Richards 1990; Zydroń et al. 2019). The decreasing pattern of RAR values in
447 this study is similar to the values reported by other researchers for various species and in other locations such as the
448 Mediterranean (Moresi et al. 2019), subtropical (Genet et al. 2008), temperate (Bischetti et al. 2005; Abdi et al. 2010a;
449 Abdi and Deljouei 2019), arid (Abdi et al. 2019) climate zones.

450 4.2. Root reinforcement

451 Root reinforcement is fundamentally determined by combining root distribution and mechanical properties (Schwarz
452 et al. 2013). Mechanical properties of roots are well-known by root maximum tensile force and stiffness. Several

453 researchers indicated that different parameters affect root resistance, including species (Abdi and Deljouei 2019), root
454 size and age (Loades et al. 2013; Gilardelli et al. 2017; Boldrin et al. 2017), tree age (Genet et al. 2008), root length
455 (Zhang et al. 2012), DBH (Deljouei et al. 2018), cellulose and lignin content (Hales et al. 2009; Abdi et al. 2014), root
456 moisture content (Moresi et al. 2019), root dehydration (Ekeoma et al. 2021), season (Makarova et al. 1998; Abdi and
457 Deljouei 2019), living or decaying roots (Vergani et al. 2014), **altitude** (Genet et al. 2011), slope position (Stokes 2002;
458 Abdi et al. 2010a), soil moisture content (Tsige et al. 2020), and elastic modulus as a root reinforcement input parameter
459 (Cislaghi 2021).

460 The root system may be adapted according to the environmental conditions and tree location, where roots resistance
461 will be varied by different **altitudes** (Genet et al. 2011). If **altitude** causes a variation in root cellulose content, as a
462 result, we can expect that root resistance will be altered. Consequently, environmental conditions might help
463 researchers to understand the mechanisms of root reinforcement better and use these data in modelling slope stability,
464 however, such information about the effect of **altitude** on root systems remains largely unsolved and further
465 investigation on root morphological adaptation is required. Root reinforcement of *C. betulus* at 400 m a.s.l. higher than
466 1300 m a.s.l. i.e. root reinforcement decreased significantly with increasing **altitude**, which means it was required more
467 force for root failure. These findings are similar to other studies performed on root reinforcement of different **altitudes**
468 (Hales et al. 2009; Genet et al. 2011; Vergani et al. 2012). It was investigated that the differences in site interactions
469 for various species result from several conditions of growing sites and environment (Vergani et al. 2012). It is clarified
470 that soil's chemical and physical properties are a consequence of changes in root resistance and root reinforcement
471 with increasing **altitude** (Genet et al. 2011).

472 The quantity of root reinforcement assessed by the contribution of different species might indicate the impact of roots
473 in reducing shallow landslides and slope instabilities. Root numbers of *C. betulus* were more **numerous** than *F.*
474 *orientalis*, so root reinforcement of *C. betulus* is higher than *F. orientalis*. Also, root **resistance** has differed between
475 **altitudes**, and roots of *C. betulus* have shown **higher resistance in terms of force** than *F. orientalis* (see Deljouei et al.
476 2020).

477 Root reinforcement of large trees is the highest among all the tree DBH classes, concurs with Cohen and Schwarz
478 (2017). The roots of the oldest trees were the most resistant in tension compared to the middle age and young trees
479 (Genet et al. 2006). It could be defined by the differences in the root structure of trees in the early growth stage and
480 older trees, as large DBH trees may possess a higher amount of cellulose (Genet et al. 2006). As a result, investigating
481 cellulose content in roots of large, medium, and small trees would be interesting in future research.

482 Root reinforcement decreases with increasing distance from tree trunks and varies considerably, even at the same
483 distance from trees, and similar results were found in past research (Moos et al. 2016). Therefore, we highlighted that
484 considering uniform cohesion value for vegetation in landslide models may not be appropriate to represent the effect
485 of trees on slope stability.

486 **4.3. The implications on slope stability and possible countermeasures of forest management**

487 According to our results, species, trees DBH, slope position, distance from the tree trunk, and **altitude** strongly
488 influence tree roots' stabilizing effect. Hence, the addition of a uniform cohesion term for vegetation in physically-

489 based landslide models (e.g., CHASM model (Wilkinson et al. 2002), TRIGRS model (Baum et al. 2008), SOSlope
490 (Cohen and Schwarz 2017), PRIMULA model (Cislaghi et al. 2018), and SLIP model (Montrasio and Valentino 2008))
491 may not be appropriate to represent the trees impacts on slope stability. The variable value for root reinforcement based
492 on species, trees DBH, slope position, distance from the tree trunk, and **altitude** shows to be a good approximation for
493 root reinforcement models. These findings indicated that specific root reinforcement should be considered due to
494 various species, tree DBH, etc. Due to the high cost and time involved in investigating root reinforcement may be of
495 great relevance to the large-scale model application. Based on a large database about root properties and root
496 reinforcement in beech and hornbeam species in the temperate forests of Europe and Iran (Bischetti et al. 2005, 2007,
497 2009; Abdi et al. 2010b; Chiaradia et al. 2016; Abdi and Deljouei 2019; Deljouei et al. 2020; Cislaghi, 2021; Cislaghi
498 et al. 2021), modelling root reinforcement can be applied **for various environmental conditions**.
499 **By comparing the performance of various species in terms of additional root reinforcement, the species most likely to**
500 **increase slope stability were identified. Also, our result can be used for nature-based solutions targeting root**
501 **reinforcement, like the effect of different forest stand structures on slope stability (Moos et al. 2016; Dazio et al. 2018),**
502 **and forest management scenarios (Kumar et al. 2021).** Over a long period, *C. betulus* is preferable to *F. orientalis* when
503 promoting one species over another to increase (or maintain) slope stability. Forest managers should consider this
504 outcome when developing strategies for large forests in mountainous areas. Certainly, factors including inter-and
505 intraspecific competition affect the performance of species concerning slope stability (Chiaradia et al. 2016). As shown
506 in this paper, this performance can be considered by using a large, widely distributed dataset and evaluating the
507 probabilistic function used to describe the values in the field survey. By clarifying the higher prevention power's
508 behaviour from large DBH *C. betulus* trees, it is possible to propose forest management that keep this tree species with
509 a large diameter in landslide-prone areas. **Using both basal and lateral reinforcement, the results suggest that the most**
510 **stabilizing species is *C. betulus*.** Importantly, our findings imply tree diameter and **species** may be appropriate for
511 assessing FoS by common tree species in Hyrcanian temperate forests. **Future studies can be conducted on the available**
512 **landslide inventory data with back analysis.**

513 **5. Conclusions**

514 This study conducted a total of 1080 profile trenches for widely species in temperate forests of Iran and European
515 countries (i.e., *C. betulus* and *F. orientalis*) at various **altitudes** (i.e., 400, 940, and 1350 m a.s.l). Root distribution and
516 mechanical properties of trees with 7.5-32.5, 32.5-57.5, and 57.5-82.5 cm DBH were used as input parameters of
517 RBMw. This study highlighted quantitative information about the impact of roots mechanical properties and
518 distribution in DBH classes, slope position, **altitude**, and vertical and horizontal distance from tree trunk of *C.*
519 *betulus* and *F. orientalis* on root reinforcement and slope stability is the practical conjunction for forest management.
520 Our results reported that **altitude**, DBH classes, and slope position significantly affect root reinforcement in Hyrcanian
521 temperate forests. Slope stabilization is decreased in further distances (more than 1 m) from the tree trunk; furthermore,
522 it is entirely different for both **species** in which *C. betulus* shows better **root reinforcement** than *F. orientalis*.
523 Furthermore, the most stabilizing species is mature *C. betulus* **at all altitudes**, which maintained an instability
524 probability near 29% for very steep conditions. Overall, we could highlight that vegetation's impact on soil slope

525 stability depends on various parameters, and considering the constant value of soil reinforcement via tree roots in soil
526 stabilization modelling is incorrect. As far as slope stabilization is concerned, there are no specific species suitable for
527 all ecological conditions since its suitability depends not only on its root reinforcement characteristics but also on the
528 species ability to grow and support healthy forest cover. Protecting slopes from long-term instability is dependent on
529 the forest's ability to withstand disturbances and its ability to recover after disturbances.

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