



1 **Quantifying vegetation indices using TLS: methodological**
2 **complexities and ecological insights from a Mediterranean**
3 **forest**

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10 **Abstract.** Accurate measurement of vegetation density metrics including plant, wood and leaf area indices (PAI,
11 WAI and LAI) is key to monitoring and modelling carbon storage and uptake in forests. Traditional passive sensor
12 approaches, such as Digital Hemispherical Photography (DHP), cannot separate leaf and wood material, nor
13 individual trees, and require many assumptions in processing. Terrestrial Laser Scanning (TLS) data offer new
14 opportunities to improve understanding of tree and canopy structure. Multiple methods have been developed to
15 derive PAI and LAI from TLS data, but there is little consensus on the best approach, nor are methods
16 benchmarked as standard.

17 Using TLS data collected in 33 plots containing 2472 trees of five species in Mediterranean forests, we compare
18 three TLS methods (*LiDAR Pulse*, *2D Intensity Image* and *Voxel-Based*) to derive PAI and compare with co-
19 located DHP. We then separate leaf and wood in individual tree point clouds to calculate wood to total plant area
20 (α), a metric to correct for non-photosynthetic material in LAI estimates. We use individual tree TLS point clouds
21 to estimate how α varies with species, tree height and stand density.

22 We find the *LiDAR Pulse* method agrees most closely with DHP, but is limited to single scan data so cannot
23 determine individual tree α . The *Voxel-Based* method shows promise for ecological studies as it can be applied to
24 individual tree point clouds. Using the *Voxel-Based* method, we show that species explain some variation in α ,
25 however, height and density were stronger predictors.

26 Our findings highlight the value of TLS data to improve fundamental understanding of tree form and function,
27 but also the importance of rigorous testing of TLS data processing methods at a time when new approaches are
28 being rapidly developed. New algorithms need to be compared against traditional methods, and existing
29 algorithms, using common reference data. Whilst promising, our results show that metrics derived from TLS data
30 are not yet reliably calibrated and validated to the extent they are ready to replace traditional approaches for large
31 scale monitoring of PAI and LAI.

32



33 **1 Introduction**

34 Terrestrial Laser Scanning (TLS) generates high-resolution 3D measurements of whole forests and individual
35 trees (Burt et al., 2018; Disney, 2018), leading to the development of completely new monitoring approaches to
36 understand the structure and function of ecosystems (Lines et al., 2022). Unlike traditional passive sensors, TLS
37 can estimate plant, wood and leaf area indices (PAI; WAI; LAI) for both whole plots and individual tree point
38 clouds (Calders et al., 2018), and is unaffected by illumination conditions. This has led to the development of
39 several methods for processing TLS data to extract the key metrics PAI, WAI and LAI (e.g. Hosoi and Omasa,
40 2006; Jupp et al., 2008; Zheng et al., 2013). However, intercomparison of algorithms and processing approaches
41 to derive the same metrics from different TLS methods are lacking.

42 Leaf Area Index (LAI), defined as half the amount of green leaf area per unit ground area (Chen and Black, 1992),
43 determines global evapotranspiration, phenological patterns and canopy photosynthesis, and is therefore an
44 essential climate variable (ECV), as well as a key input in dynamic global vegetation models (Sea et al., 2011;
45 Weiss et al., 2004). Accurate measurements of LAI, WAI and PAI have historically been derived from labour
46 intensive destructive sampling (Baret et al., 2013; Jonckheere et al., 2004), so over large spatial or temporal scales
47 these can only be measured indirectly, typically with remote sensing. Large-scale remote sensing, using
48 spaceborne and airborne instruments, has been widely used to estimate LAI over large areas (Pfeifer et al., 2012),
49 but requires calibration and validation using in situ measurements to constrain information retrieval (Calders et
50 al., 2018). Non-destructive in situ vegetation index estimates have historically been made by measuring light
51 transmission below the canopy and using simplifying assumptions about canopy structure to estimate the amount
52 of intercepting material (e.g. Beer-Lambert law; Monsi and Saeki, 1953). The most common method, Digital
53 Hemispherical photography (DHP; Figure 1a), requires both model assumptions and subjective user choices during
54 data acquisition and processing in order to estimate both PAI and LAI (Breda, 2003). DHP images are processed
55 by separating sky from canopy, but not photosynthetic from non-photosynthetic vegetative material, so additional
56 assumptions are needed to calculate either LAI or WAI (Jonckheere et al., 2004; Pfeifer et al., 2012). Separation
57 of LAI from PAI can be achieved by removing or masking branches and stems from hemispherical images (e.g.
58 Sea et al., 2011; Woodgate et al., 2016), but is not reliable when leaves are occluded by woody components
59 (Hardwick et al., 2015). An alternative approach is to take separate DHP measurements in both leaf on and leaf
60 off conditions, and derive empirical wood to plant ratios (WAI/PAI, α) (Leblanc and Chen, 2001), but this is not
61 always practical, for example in evergreen forests. The difficulty of separation means that studies often omit
62 correcting for the effect of WAI on optical PAI measurements altogether (Woodgate et al., 2016), but since woody
63 components in the forest canopy can account for more than 30% of PAI (Ma et al., 2016) this can introduce
64 overestimation. Further, although DHP estimates of LAI or PAI are valuable both for ecosystem monitoring and
65 developing satellite LAI products (Hardwick et al., 2015; Pfeifer et al., 2012), they are limited to sampling only
66 at a neighbourhood or plot level (Hardwick et al., 2015; Pfeifer et al., 2012, Weiss et al., 2004), and cannot be
67 used to measure individual tree LAI except for open grown trees (Béland et al., 2014).

68 The ratio of wood to total plant area, α , is known to be dynamic, changing in response to abiotic and biotic
69 conditions. For example, the Huber value (sapwood to leaf area ratio, a related measure to α) may vary according
70 to water availability (Carter and White 2009). Leaf area may therefore be indicative of the drought tolerance level
71 of a tree, with more drought tolerant species displaying a lower leaf area, reducing the hydraulic conductance of



72 the whole tree and therefore increasing its drought tolerance (Niinemets and Valladares, 2006). α has been
73 hypothesised to increase with the size of a tree in response to the increased hydraulic demand associated with
74 greater hydraulic resistance of tall trees (Magnani et al., 2000) and higher transpiration rates of larger LAI
75 (Battaglia et al., 1998; Phillips et al., 2003). Stand density may also impact α (Long and Smith, 1988; Whitehead,
76 1978), as increased stand level water use scales linearly with LAI (Battaglia et al., 1998; Specht and Specht, 1989),
77 reducing water availability to individual trees competing for the same resources (Jump et al., 2017). Large scale
78 quantification of α or Huber value, however, is difficult as studies usually rely on a small number of destructively
79 sampled trees (e.g. Carter and White, 2009; Magnani et al., 2000), litterfall traps (e.g. Phillips et al., 2003) or
80 masking hemispherical images (e.g. Sea et al., 2011; Woodgate et al., 2016). These approaches are only applicable
81 on a small to medium scale, and in the case of image masking, cannot differentiate between individuals. Variation
82 in α , for example by species and or stand structure, is therefore largely unknown.

83 1.2 TLS methods for calculating PAI, LAI and WAI

84 TLS methods for extracting PAI, LAI and WAI can be broadly categorised into two types: (1) LiDAR return
85 counting, using single scan data (e.g., the *LiDAR Pulse* method; Jupp et al., 2008, and *2D Intensity Image* method;
86 Zheng et al., 2013) and (2) point cloud voxelisation, usually using co-registered scans (e.g., the *Voxel-Based*
87 method; Hosoi and Omasa, 2006).

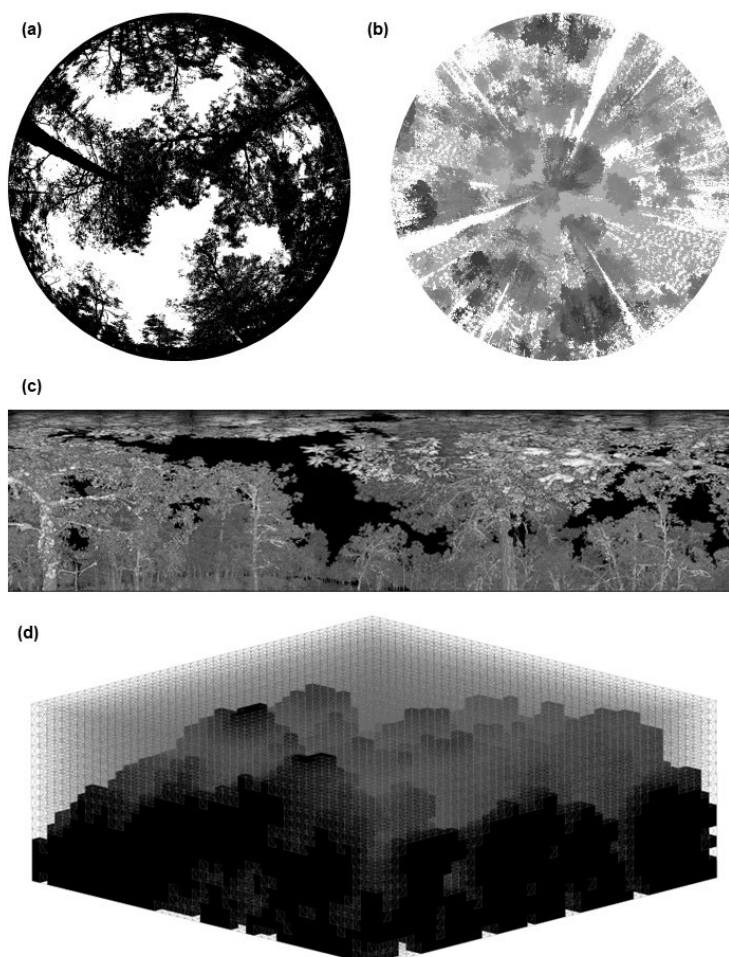
88 The *LiDAR Pulse* method (Jupp et al., 2008; Figure 1b) estimates gap fraction (P_{gap}) using single scan data, as a
89 function of the total number of outgoing LiDAR pulses from the sensor and the number of pulses that are
90 intercepted by the canopy. This method, which eliminates illumination impacts associated with the use of DHP
91 (Calders et al., 2014), has been implemented in the python module, *PyLidar* (www.pylidar.org) and the R package,
92 *rTLS* (Guzman, et al. 2021). Using the *LiDAR Pulse* method, Calders et al. (2018) compared TLS PAI estimates
93 from two ground-based passive sensors (LiCOR LAI-2000 and DHP) with TLS data collected with a RIEGL VZ-
94 400 TLS in a deciduous woodland, and found the two passive sensors underestimated PAI values compared to
95 TLS, with differences dependent on DHP processing and leaf on/off conditions.

96 The *2D Intensity Image* method (Zheng et al., 2013; Figure 1c), also uses raw single scan TLS point clouds, but
97 unlike the *LiDAR Pulse* method, this approach converts LiDAR returns into 2D panoramas where pixel values
98 represent intensity. PAI is estimated by classifying pixels as sky or vegetation, based on their intensity value, to
99 estimate P_{gap} , and then applying Beer-Lambert's law. As for the *LiDAR Pulse* method, this approach has been
100 shown to generate higher PAI estimates than DHP (Calders et al., 2018; Woodgate et al., 2015; Grotti et al., 2020),
101 with differences attributed to the greater pixel resolution and viewing distance of TLS resolving more small
102 canopy details (Grotti et al., 2020).

103 The *Voxel-Based* method (Figure 1d) estimates PAI by segmenting a point cloud into voxels and either simulating
104 radiative transfer within each cube (Béland et al., 2014; Kamoske et al., 2019), or classifying voxels as either
105 containing vegetation or not, and dividing vegetation voxels by the total number of voxels (Hosoi and Omasa,
106 2006; Itakura and Hosoi, 2019; Li et al., 2017). Crucially, this method may be applied to multiple co-registered
107 scan point clouds and so can be used to calculate PAI for both whole plots and individual, segmented TLS trees.



108 The *LiDAR Pulse* method and *2D Intensity Image* method both use single scan data. However, to generate robust
109 estimates of canopy properties that avoid errors from occlusion effects, multiple co-registered scans taken from
110 different locations are likely needed (Wilkes et al., 2017). Further, both these methods require raw unfiltered data
111 to accurately measure the ratio of pulses emitted from the scanner and number of pulses that are intercepted by
112 vegetation. This means “noisy” points caused by backscattered pulses (Wilkes et al., 2017) are included in
113 analyses, potentially leading to higher PAI estimates. However, the *LiDAR Pulse* and *2D Intensity Image* methods
114 may introduce fewer estimation errors compared DHP, which is influenced by differences in sky illumination
115 conditions and camera exposure (Weiss et al., 2004).



116

117

118 **Figure 1: Methods for PAI estimation applied in this study: (a) a binarised digital hemispherical photograph (DHP),**
119 **(b) TLS raw single scan point cloud, used within the LiDAR Pulse method (Jupp et al., 2008). Image shows a top-down**
120 **view of raw point cloud and greyscale represents low (grey) and high (black) Z values, (c) TLS 2D intensity image for**
121 **the 2D Intensity Image method (Zheng et al., 2013), (d) Voxelised co-registered whole plot point cloud for the Voxel-**
122 **Based method (Hosoi and Omasa, 2006), showing a representative schematic of cube voxels with edge length of 1m,**



123 voxelised using the *R* package *VoxR* (Lecigne et al., 2018). Solid black voxels are classified as containing vegetation
124 (filled) and voxels outlined with grey lines are voxels classified as empty.

125 1.3 Scope and aims

126 In this study we use a dataset of 528 co-located DHP and high-resolution TLS scans from 33 forest plots to
127 compare DHP derived PAI with estimates from three methods to estimate PAI from TLS data: the *LiDAR Pulse*
128 method; the *2D Intensity Image* method and the *Voxel-Based* method (Figure 1). We use a dataset collected from
129 a network of pine/oak forest plots in Spain (Owen et al., 2021) and ask (1) are the three TLS methods able to
130 reproduce DHP PAI estimates at single scan and whole plot level? (2) does α , calculated from the *Voxel-Based*
131 method on individual tree point clouds, vary with species and tolerance to drought; and (3) does α scale with
132 height and stand density?

133 2. Methods

134 2.1 Study site

135 We collected TLS and DHP data from 29 plots in Alto Tajo Natural Park ([40°41'N 02°03'W](#); FunDIV plots;
136 Baeten et al., 2013) and four plots in Cuellar ([41°23'N 4°21'W](#)) in June - July 2018 (see Owen et al., 2021 for full
137 details). Plots contained two oak species: semi-deciduous *Q. faginea* and evergreen *Q. ilex*, and three pine species:
138 *P. nigra*, *P. pinaster* and *P. sylvestris*. *P. sylvestris* is the least drought tolerant species, followed by *P. nigra*, *Q.*
139 *faginea*, *Q. ilex*; shade tolerance follows the same ranking (Niinemets and Valladares, 2006; Owen et al., 2021).
140 Although not quantitatively ranked, *P. pinaster* has been shown to be very drought tolerant, appearing in drier
141 areas than the other species (Madrigo-González et al., 2017). The area is characterised by a Mediterranean climate
142 (altitudinal gradient 840 – 1400 m.a.s.l.) (Jucker et al., 2014; Madrigo-González et al., 2017). In addition to the
143 five main canopy tree species, plots contained an understory of *Juniperus thurifera* and *Buxus sempervirens*
144 (Kuusk et al., 2018).

145 2.2 Field protocol

146 In each of the 33 30 x 30 m plots we collected TLS scans on a 10 m grid, making 16 scan locations following
147 Wilkes et al., (2017) to minimise occlusion effects associated with insufficient scans. We used a Leica HDS6200
148 TLS set to super high resolution (3.1 x 3.1 mm resolution at 10 m with a beam divergence of ≤ 5 mm at 50 m; scan
149 time 6m 44 s; see Owen et al., 2021). At each of the 528 scan locations and following the protocol in Pfeifer et
150 al., (2012), we captured co-located DHP images with three exposure settings (automatic and \pm one stop exposure
151 compensation), levelling a Canon EOS 6D full frame DSLR sensor with a Sigma EX DG F3.5 fisheye lens,
152 mounted on a Vanguard Alta Pro 263AT tripod.

153 2.3 Calculation of single scan and whole plot PAI using DHP data

154 For each of the red-green-blue (RGB) DHP images we extracted the blue band for image thresholding, as this best
155 represents sky/vegetation contrast (Pfeifer et al., 2012). For each plot, we picked the exposure setting that best
156 represented sky/vegetation difference based on pixel brightness histograms of four sample locations indicative of
157 the plot. We carried out automatic image thresholding using the Ridler and Calvard method (1978), to create a
158 binary image of sky and vegetation, avoiding subjective user pixel classification (Jonckheere et al., 2005). We
159 calculated PAI from the binary image, limiting the field of view to a 5° band centred on the hinge angle of 57.5°



160 (55° – 60°). The hinge angle has a path length through the canopy twice the canopy height, so the band around it
161 is an area of significant spatial averaging taken as representative of canopy structure of the area (Calders et al.,
162 2018; Jupp et al., 2008). From the binarised hinge angle band we calculated gap fraction as the number of sky
163 pixels divided by the total number of pixels and PAI using an inverse Beer-Lambert law equation (Monsi and
164 Saeki, 1953). We calculated whole plot PAI as the arithmetic mean within plot scan location PAI. As this value
165 does not correct for canopy clumping, it is better described as effective PAI, rather than true PAI (Woodgate et
166 al., 2015). However, as the TLS and DHP methods we apply here account for canopy clumping differently, we
167 compare effective values and here-on refer to effective PAI as PAI (Calders et al., 2018).

168 **2.4 Calculation of single scan and whole plot PAI from TLS data**

169 To calculate PAI using the *LiDAR Pulse* method (Jupp et al., 2008), we calculated the gap fraction (P_{gap}) for a
170 single scan (Figure 1b) by summing all returned laser pulses and dividing by the number of total outgoing pulses,
171 following Lovell et al. (2011; see Eq. 7 in that study), and then estimated PAI following Jupp et al. (2008; see Eq.
172 18 in that study), setting the sensor range to 5° around the hinge angle as before (55° – 60°). Single scan PAI was
173 taken as the cumulative sum of PAI values estimated by vertically dividing the hinge region into 25 cm intervals
174 (Calders et al., 2014). We implemented the *LiDAR Pulse* method using the open-source *R* (R Core Team, 2020)
175 package, *rTLS* (Guzmán and Hernandez, 2021).

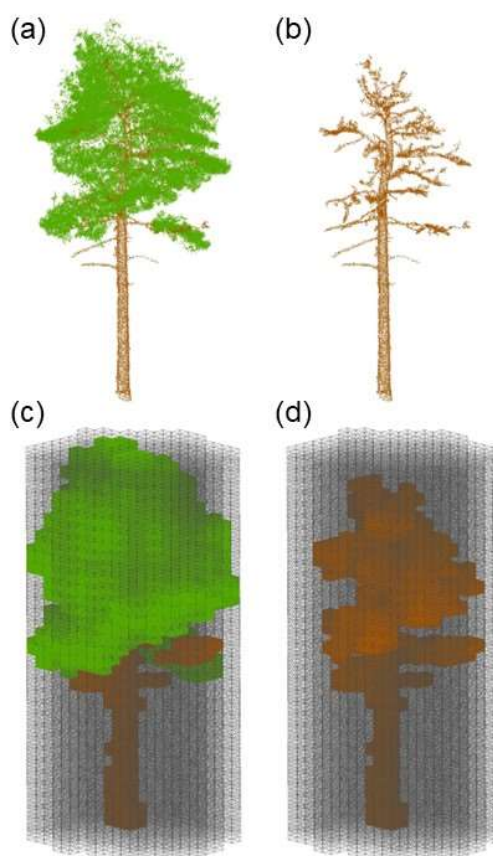
176 To calculate PAI using the *2D Intensity Image* method (Zheng et al., 2013) we converted 3D TLS point cloud
177 data from all 528 scan locations into polar coordinates and scaled intensity values to cover the full 0-255 range
178 (Figure 1c) and rasterised into a 2D intensity image using the open-source *R* package, *raster* (Hijmans, 2022). We
179 cut the 2D intensity image to a 5° band around the hinge angle (55° – 60°) and classified sky and vegetation pixels
180 in each image using the Ridler and Calvard method (1978). We calculated P_{gap} as the number of pixels classified
181 as sky divided by the total number of pixels and derived PAI with an inverse Beer-Lambert law equation (Monsi
182 and Saeki, 1953).

183 Following the same approach as applied to our DHP data, we calculated whole plot PAI for the *LiDAR Pulse* and
184 *2D Intensity Image* methods as the arithmetic mean of within plot single scan PAI estimates.

185 To calculate PAI using the *Voxel-Based* method, we followed a voxel classification approach (Hosoi and Omasa,
186 2006), matching the voxel size to the resolution of the point cloud (0.05 m), following Li et al., (2017), who
187 showed that matching the voxel size to the point cloud point to point minimum distance (resolution) increases
188 accuracy as small canopy gaps are not included in voxels classified as vegetation. We combined individually
189 segmented trees (see Owen et al., 2021) into whole plot point clouds and voxelised them using the open source *R*
190 package, *VoxR* (Lecigne et al., 2018), with a full grid covering the minimum to maximum XYZ ranges of the plot.
191 We classified any voxel containing > 0 points as vegetation (“filled”), and empty voxels as gaps. We then split
192 the voxelised point cloud into slices one voxel high. Within each slice, the contact frequency is calculated as the
193 fraction of filled to total number of voxels. We then multiplied the contact frequency by a correction factor for
194 leaf inclination, set at 1.1 (Li et al., 2017), and whole plot PAI was calculated as the sum of all slices’ contact
195 frequencies.



196 2.5 Calculation of individual tree PAI, WAI and α using the voxel-based method



197 **Figure 2: Visualisation of the workflow for applying the Voxel-Based method to estimate individual-tree PAI, WAI and**
198 **α . (a) Individual tree point cloud; (b) separated leaf off (wood) individual tree point cloud; (c) voxelised individual tree**
199 **point cloud; (d) voxelised wood cloud. Solid black voxels are filled voxels and grey lines are empty voxels. Empty voxels**
200 **occupy the space within the projected crown area of the tree. Image shows schematic of point cloud voxelised with cube**
201 **voxels with edge length of 0.5 m. Wood and leaf separation was carried out using *TLSeparation* (Vicari et al., 2019).**
202 **Point cloud voxelisation was carried out using modified functions from *R* package *VoxR* (Lecigne et al., 2018).**

203 As the only method using multiple co-registered scans, the *Voxel-Based* method is only method we found capable
204 of estimating individual tree leaf and wood properties. We estimated PAI and WAI for 2472 individual trees
205 segmented from co-registered point clouds following a similar method to the whole plot point cloud. We extracted
206 individual trees using the automated tree segmentation program *treeSeg* (Burt et al., 2019), implemented in C++,
207 see Owen et al., (2021) for full details, and Owen et al., (2022) for individual segmented tree data.

208 To estimate PAI, WAI and α for each tree, we first separated leaf from wood points in individual tree point clouds
209 using the open source Python library *TLSeparation* (Vicari et al., 2019), and then used the wood only point clouds
210 to calculate WAI. *TLSeparation* classifies points as leaf or wood, iteratively looking at a predetermined number
211 of nearest neighbours (*knn*). The *knn* of each iteration is directly dependent on point cloud density, since high



212 density point clouds will require higher a *knn* (Vicari et al., 2019). We used the utility package in *TLSeparation*
213 to automatically detect the optimum *knn* for each tree point cloud.

214 To voxelise individual tree complete (Figure 2a) and wood only (Figure 2b) point clouds, we used a modified
215 approach based on Lecigne et al., (2018), voxelising within the projected crown area of the whole tree point cloud
216 (Figure 2c) to calculate PAI. We calculated WAI within the projected crown area of the whole tree (Figure 2d;
217 using the whole crown and not just the wood point cloud), and derived α for each tree as WAI/PAI .

218 2.6 Statistical Analyses

219 We tested the relationships between TLS PAI and DHP PAI estimates using Standardised Major Axis (SMA)
220 using the open source *R* (R Core Team, 2020) package, *smatr* (Warton et al., 2012). SMA is an approach to
221 estimating a line of best fit where we are not able to predict one variable from another (Warton et al., 2006); we
222 chose SMA because we do not have a ‘true’ validation dataset, so avoid assuming either DHP or any of the TLS
223 methods produces the most accurate results. For each TLS method, we assessed the relationship with DHP using
224 the coefficient of determination and RMSE. To further understand observed drivers of variance in PAI, we tested
225 the relationship between PAI and TLS estimated whole plot crown area index, CAI, calculated as the sum of
226 projected crown area, divided by the plot area (Owen et al., 2021), and indicative measure of stand density, using
227 SMA.

228 To test if α differs by species, we used linear mixed models (LMMs) in the *R* package, *lme4* (Bates et al., 2015).
229 We included an intercept only random plot effect to account for local effects on α :

230

$$231 \alpha_{i,sj} = a_s + Plot_j \quad (1)$$

232

233 here, α_i is α of an individual of species s , in plot j , and a_s is the parameter to be fit. To test the effect of stand
234 structure and tree height on α we fit relationships separately for each species, again including a random plot effect:

235

$$236 \alpha_{i,sj} = a_s + b_s H_i + c_s CAI_j + Plot_{sj} \quad (2)$$

237

238 here H_i is the height of the tree, CAI_j is the crown area index for the plot, with other parameters as before.

239 For each species’ model (equation 2), we calculated the intra-class correlation coefficient (ICC). The ICC, similar
240 to coefficient of determination, quantifies the amount of variance explained by the random effect in a linear mixed
241 model (Nakagawa et al., 2017).

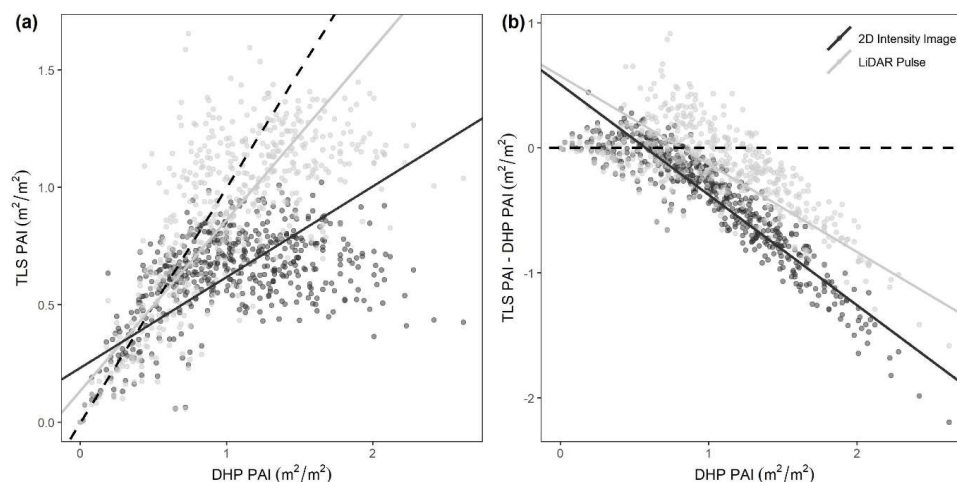


242 3. Results

243 3.1 Comparison of plant area index estimated by DHP and single scan TLS

244 Of the two single scan TLS methods tested (*LiDAR Pulse* method and *2D Intensity Image* method), we found that
245 PAI estimated using the *LiDAR Pulse* method more strongly agreed with DHP PAI, but there was also significant
246 correlation for the *2D Intensity Image* method (SMA; *LiDAR Pulse* method $R^2 = 0.50$, slope = 0.73, $p < 0.001$,
247 RMSE = 0.14, and *2D Intensity Image* method $R^2 = 0.22$, slope = 0.38, $p < 0.001$, RMSE = 0.39, respectively,
248 Figure 3a). At larger PAI values, both TLS methods underestimated PAI compared with DHP (Figure 3b). We
249 found statistically significant negative correlations between residuals and DHP for both methods (SMA; *2D*
250 *Intensity Image* method residuals $R^2 = 0.85$, slope = -0.88, $p < 0.01$; *LiDAR Pulse* method residuals $R^2 = 0.47$, slope
251 = -0.70, $p < 0.01$; Figure 3b). The *2D Intensity Image* method showed larger underestimation at higher DHP PAI
252 values, suggesting this method may saturate sooner than both DHP and the *LiDAR Pulse* method at higher PAI
253 values (Figure 3b).

254



255 **Figure 3: Comparison of single scan TLS PAI and DHP PAI estimates, for all 528 scan locations (16 per plot). (a) The**
256 **correlation between DHP derived PAI with PAI derived using the 2D Intensity Image method, and LiDAR Pulse**
257 **method. Dashed line represents 1:1 relationship. (b) The difference between TLS and DHP PAI estimates for the 2D**
258 **Intensity Image method, and LiDAR Pulse method (dashed line at 0). Lines show statistically significant relationships**
259 **fitted using SMA ($p < 0.01$).**

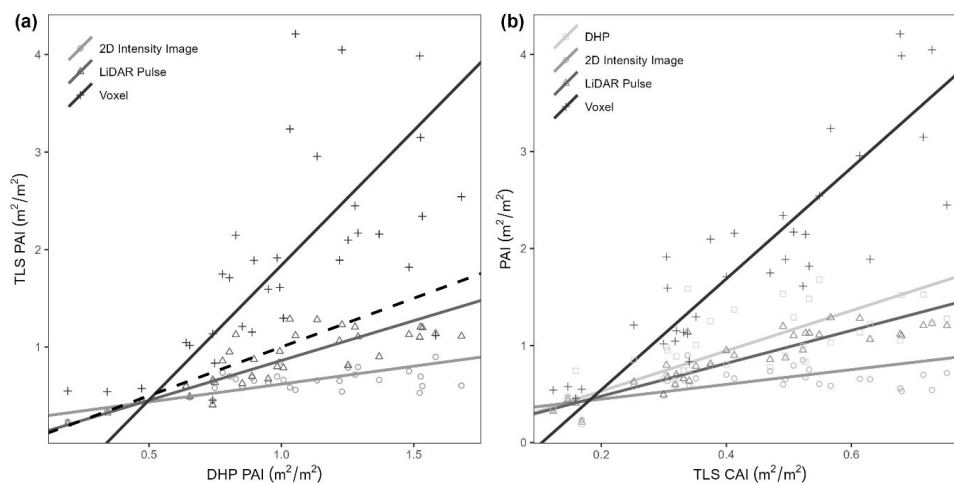
260 3.2 Comparison of whole plot plant area index estimated using TLS and DHP and the effect of plot structure 261 on PAI

262 We found statistically significant correlations between TLS whole plot PAI values and DHP PAI for all three TLS
263 methods. As for single scans (Figure 3), the *LiDAR Pulse* method showed the closest agreement to DHP PAI, here
264 compared to both the *Voxel-Based* and *2D Intensity Image* methods (SMA; *LiDAR Pulse* method $R^2 = 0.66$, slope
265 = 0.82, $p < 0.01$, RMSE = 0.14; *Voxel-Based* method $R^2 = 0.39$, slope = 2.76, $p < 0.01$, RMSE = 0.88; *2D Intensity*
266 *Image* method $R^2 = 0.35$, slope = 0.36, $p < 0.01$, RMSE = 0.39, respectively; Figure 4a). The *2D Intensity Image*
267 method and *LiDAR Pulse* method consistently underestimated PAI compared to DHP, whilst the *Voxel-Based*
268 method underestimated in plots with lower DHP PAI and overestimated in plots with higher DHP PAI. The *Voxel-*



269 *Based* method's high PAI values compared to other methods is likely due to its use of multiple co-registered scans
 270 reducing occlusion effects prevalent in single scan data.

271 To assess the effect of plot structure on variation in TLS derived PAI, we compared TLS PAI estimates to TLS
 272 estimated crown area index (CAI, m² projected crown area per m² ground area, Figure 4b). We found a significant
 273 positive relationship between CAI and PAI estimated using each of the *LiDAR Pulse* method, the *Voxel-Based*
 274 method, and DHP (SMA; *LiDAR Pulse* method R² = 0.79, slope = 1.69, p<0.01; *Voxel-Based* method R² = 0.76,
 275 slope = 5.72, p<0.01; *2D Intensity Image* method R² = 0.15, slope = 0.76; DHP R² = 0.46, slope = 2.07, p<0.01,
 276 respectively; Figure 4b), where the *2D Intensity Image* method appears to saturate at medium CAI values (Figure
 277 4b).



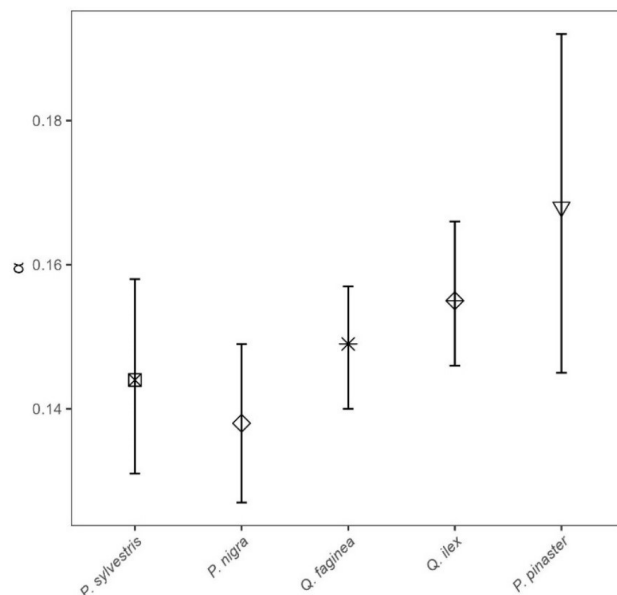
278 **Figure 4: Comparison of plot level TLS PAI and DHP PAI, and CAI vs PAI estimates for all 33 plots. (a) The correlation**
 279 **between DHP derived PAI and PAI derived using 2D Intensity Image (circle), LiDAR Pulse (triangle) and Voxel-Based**
 280 **(cross) methods (b) The correlation between TLS derived CAI and PAI derived using DHP (square), LiDAR Pulse**
 281 **(triangle) and Voxel-Based (cross) methods. Lines show statistically significant relationships fitted using SMA (p<0.01).**
 282 **Dashed line represents 1:1 relationship.**

283 3.4 Influence of species, tree height and CAI on α

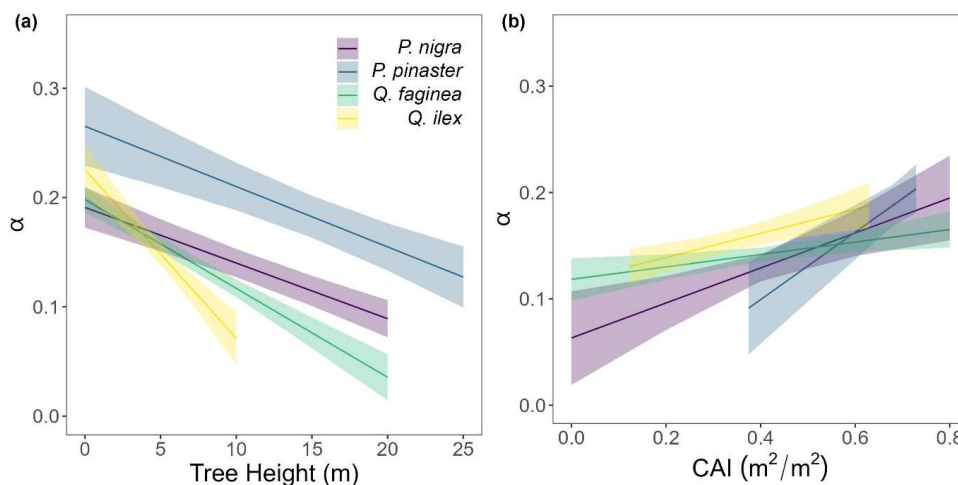
284 To understand drivers of variance in α , we used individual tree PAI and WAI, calculated using the *Voxel-Based*
 285 method to test the relationship between species and α , and height/ CAI and α . We found that more drought tolerant
 286 species generally had higher α values than less drought tolerant species (Table A1; Figure 5), however, confidence
 287 intervals were wide and overlapping, suggesting that species is not a strong predictor of variation in α . We found
 288 a statistically significant negative effect of height (p<0.001; Table A2; Figure 6a) and positive effect of CAI
 289 (p<0.01 – 0.05; Table A2; Figure 6b) on α for all species apart from *P. sylvestris*. α decreased more rapidly with
 290 height and increased less rapidly with CAI for oaks than pines. Statistically significant ICC values were higher
 291 for *P. nigra* (ICC = 0.211; Table A2) than *P. pinaster*, *Q. faginea* and *Q. ilex* (ICC = 0.036; 0.060; 0.070,



292 respectively), showing that more α variation is explained by the random plot effect in *P. nigra* than the other
 293 species. *P. pinaster* has a wider confidence interval (Figure 5), possibly explained by its lower sample size.



294 **Figure 5:** Linear mixed model derived α values (a, equation 1) for all 2472 individual trees of species *P. sylvestris*, *P.*
 295 *nigra*, *Q. faginea*, *Q. ilex* and *P. pinaster*. Error bars represent 95% confidence intervals. Species are listed from low –
 296 high drought tolerance, with the exception of *P. pinaster*, for which drought tolerance index has not been calculated in
 297 the literature.



298 **Figure 6:** Variation in α for each species: *Pinus nigra*, *P. pinaster*, *Q. faginea* and *Q. ilex* with (a) height and (b) plot
 299 CAI. Lines represent statistically significant linear mixed models (equation 2; $p < 0.001$). Ribbons represent 95%
 300 confidence intervals. The model for *P. sylvestris* was not statistically significant.

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304 **4. Discussion**

305 **4.1 Comparison of approaches to deriving PAI from remote sensed data**

306 We found substantial differences in PAI values estimated from TLS and DHP and from different TLS processing
307 methods (Figures 3 and 4). Further, differences between TLS methods varied across plot structure (CAI), with the
308 greatest differences between methods in plots with high CAI, and therefore high canopy density. Although
309 previous studies have presented TLS as an improvement over DHP due to its independence of illumination and
310 sky conditions during the data acquisition phase, and ability to resolve fine-scale canopy elements and gaps
311 (Calders et al., 2018; Grotti et al., 2020; Zhu et al., 2018), we have shown that there is large variability between
312 TLS processing methods in Mediterranean forests. Rigorous intercomparison of approaches, ideally using
313 standard benchmarking TLS datasets, and destructive sampling, would improve trust and reliability of TLS
314 algorithms.

315 **4.2 The *LiDAR Pulse* and *2D Intensity Image* method derived PAI estimates were lower than those derived
316 from DHP and the *Voxel-Based* method**

317 We found the *LiDAR Pulse* method (Jupp et al., 2008) to have the best agreement with DHP for both whole plot
318 and single scan PAI estimates. In contrast to previous TLS – DHP comparisons (Calders et al., 2018; Grotti et al.,
319 2020; Woodgate et al., 2015), we found that the *LiDAR Pulse* and *2D Intensity Image* methods underestimated
320 PAI compared to DHP, except at very low PAI values. Quantification of PAI from DHP may introduce additional
321 sources of error, for example, its relatively lower resolution compared to TLS could lead to mixed pixels that have
322 a greater chance of misclassification of sky as vegetation (Jonckheere et al., 2004). This effect could be enhanced
323 in a Mediterranean forest as trees in drier climates tend to have smaller leaves (Peppe et al., 2011), leading to
324 more small canopy gaps that TLS may resolve where DHP cannot. Further, although we took steps to reduce the
325 error introduced at DHP data acquisition and processing steps, including using automatic thresholding and
326 collecting images with multiple exposures, DHP processing requires both model and user assumptions that can
327 impact results. For example, DHP PAI estimates are highly sensitive to camera exposure; increasing one stop of
328 exposure can result in 3 – 28% difference in PAI and use of automatic exposure can result in up to 70% error
329 (Zhang et al., 2005).

330 We found the *Voxel-Based* method overestimated PAI values compared to the other methods at the whole plot
331 level. This is likely due to the method's use of co-registered scans, rather than averaged single scan PAI values,
332 since co-registered scans will reduce occlusion effects prevalent in single scan data that could lead to an
333 underestimation of PAI (Wilkes et al., 2017). The *Voxel-Based* method is, however, sensitive to voxel size (Li et
334 al., 2016), and larger voxels lead to larger PAI estimates as they fill small canopy gaps; we chose a voxel size of
335 0.05 m to match the minimum distance between points in our downsampled dataset. However, the *Voxel-Based*
336 method is a memory intensive approach to calculating PAI, and smaller voxels have higher memory requirements.
337 We picked this data resolution, and therefore voxel size, to balance the need to capture fine-scale canopy details
338 against memory requirements for running many large plots. Voxel size could have been chosen based on
339 estimates' match to DHP, but this would assume (1) that DHP estimates are most accurate, and (2) that DHP data
340 are always available, limiting the wider applicability of our findings. Understanding which method is over or
341 underestimating would require a destructively sampled dataset for validation, which was not possible for this
342 study (or most ecosystems). However, other studies using voxel approaches have found that although these



343 produce high LAI values for individual trees, these are underestimates compared with destructive samples (Li et
344 al., 2016). Regardless, PAI and LAI estimates using a *Voxel-Based* approach are highly dependent on voxel size
345 (Béland et al., 2014), and future work should test the influence of voxel size on PAI estimates, using destructive
346 samples in a range of environments.

347 **4.3 Relationship between PAI and CAI varied according to method and sensor**

348 The *LiDAR Pulse* method had the strongest relationship (defined as highest R^2) with TLS derived CAI,
349 demonstrating that the method is well suited to measuring PAI across the range of plot CAI values used in this
350 study. Although the *2D Intensity Image* method can tackle the significant challenges presented by edge effects
351 and partial beam interceptions, particularly present in phase-shift systems (Grotti et al., 2020), our results suggest
352 this method has a lower performance ability, with saturation occurring sooner than all other methods in dense
353 forests (Figures 3 and 4). The *2D Intensity Image* method uses the same raw single scan data as the *LiDAR Pulse*
354 method, so the better performance from the latter is likely due to the method's use of vertically resolved gap
355 fraction; both the *LiDAR Pulse* method and *Voxel-Based* method account for the vertical structure of the canopy
356 by summing vertical slices through the canopy.

357 **4.4 α variation between species and plot**

358 We used the *Voxel-Based* method to investigate individual tree α variation between species and across structure,
359 as this was the only approach we identified that could be applied to single tree point clouds. We found α values
360 obtained were within the range of values obtained from destructive approaches (0.1 – 0.6, Gower et al., 1997).
361 The drought and shade intolerant *P. nigra* showed stronger variability in α across plots (higher ICC value, Table
362 A2) than other species, suggesting its wood – leaf ratio may be more sensitive to site factors. However, as the
363 plots measured in this study vary in both abiotic conditions (altitude, aspect, slope, wetness) as well as species
364 composition, stem density and canopy cover, there may be other drivers of variation in α values.

365 We found some evidence that species with higher drought tolerance had higher α values (Figure 5; Table A1),
366 however, confidence intervals were wide, suggesting a weak relationship. There is evidence that trees that tolerate
367 water limited environments have a lower leaf area (Battaglia et al., 1998; Mencuccini and Grace, 1995), so higher
368 α values may reflect maintenance of homeostasis of leaf water use through adjustment of wood to leaf area ratio
369 (Carter and White, 2009; Gazal et al., 2006). The potential for a tree to lose water is mostly regulated through leaf
370 traits including stomatal conductance and leaf area, and both stand (Battaglia et al., 1998; Specht and Specht,
371 1989) and individual tree (Mencuccini, 2003) water use have been found to scale linearly with LAI, with drought
372 often mitigated through leaf shedding (López et al., 2021).

373 **4.5 Tree stature and stand density drives α variation**

374 Although species explain some variation in α , tree height and plot CAI were stronger predictors for all species,
375 showing the importance of local stand structure on leaf and woody allocation. We found that α scaled negatively
376 with height for all species apart from *P. sylvestris*, suggesting that in this environment, taller trees generally have
377 a lower proportion of wood to plant area index than shorter ones. *P. sylvestris*, which is at the edge of its
378 geographical range and physiological limits (Castro-Díez et al., 1997; Owen et al., 2021), showed no significant
379 relationship between height and α . We found that α scaled positively with plot level CAI for all species apart from
380 *P. sylvestris*, that is, trees growing in denser plots have a higher α . This supports theory that trees growing in dense



381 forests are competing for resources, reducing individual tree leaf area (Jump et al., 2017). The negative height –
382 α and positive CAI – α relationships in our model suggest that trees may initially invest in vertical growth to reach
383 the canopy level, and once there invest in lateral growth, with more leaf area, to increase light capture. This
384 supports theory that trees grow to outcompete neighbouring individuals for light capture (Purves and Pacala, 2008)
385 and evidence that both lateral growth and LAI are reduced beneath closed canopies (Beaudet and Messier, 1998;
386 Canham, 1988).

387 Wood may be harder to accurately classify than leaves in TLS data (Vicari et al., 2019), resulting in a higher
388 occurrence of false positives in wood clouds, potentially leading to an overestimation in WAI, and therefore
389 underestimation of α , especially in trees with small leaves which are prevalent in dry, Mediterranean environments
390 (Peppe et al., 2011). The problem of misclassification will increase in taller trees due to TLS beam divergence,
391 occlusion and larger beam footprint at further distances (Vicari et al., 2019), suggesting that WAI overestimation
392 could be more pronounced in tall trees. Although our dense scanning strategy (Owen et al., 2021) was designed
393 to mitigate some of these effects, it is possible our findings could underestimate the slope of the negative
394 relationship between α and tree height.

395 **4.6 Correcting for non-photosynthetic elements in LAI estimates using TLS**

396 The value of TLS data to estimate individual tree PAI, WAI and subsequently α , demonstrates their potential to
397 corrective factors for non-photosynthetic components in ground based remote sensing measurements of LAI.
398 Properly correcting for WAI in LAI estimates is of global importance as small errors in ground based
399 measurements propagate through to large scale satellite observations generating large errors in global vegetation
400 models (Calders et al., 2018). Our results echo others' in finding that the prevalence of woody material in the tree
401 canopy, and therefore α , is dynamic and varies by species as well as senescence, crown health and, in the case of
402 deciduous forests, leaf phenology (Gower et al., 1999). The use of single α value in a plot or region (Olivas et al.,
403 2013; Woodgate et al., 2016), invariant of species, size and forest structure, to convert PAI to LAI is therefore
404 problematic (Niu et al., 2021). Our study demonstrates the importance of taking species mix and structural
405 variation into account when correcting for non-photosynthetic material in ground-based LAI estimates.

406 **5. Conclusions**

407 We tested three methods for estimating PAI using Terrestrial Laser Scanning data and compared these against
408 traditional DHP measurements. We found large variation between PAI values estimated from each TLS method
409 and DHP, demonstrating that care should be taken when deriving PAI from ground based remote sensing methods.
410 Although the *LiDAR Pulse* method was found to have the best agreement with both single scan and whole plot
411 PAI values measured by DHP, the *Voxel-Based* method allowed separate analysis of the key metric used to correct
412 for the effect of WAI in LAI measurements, α , in individual trees. We recommend the *LiDAR Pulse* method as a
413 fast and effective method for PAI estimation independent of illumination conditions. Whilst the *Voxel-Based*
414 method may be used to analyse individual tree α and determine ecological drivers of variation, work remains to
415 determine the validity of these approaches, in particular correct voxel size choice. We found that α varies by
416 species, height and stand density, showing the importance of accurately correcting for WAI on the individual tree
417 level and the utility of TLS to do so.



418 The variation in our results for the different methods used to derive PAI from TLS data show that there is some
419 way to go before TLS derived vegetation indices can be interpreted as robust and reliable. Validation using
420 destructive samples and further intercomparison studies of methods are needed to demonstrate the advantages of
421 TLS, and use of benchmarking datasets should be standard. DHP is a faster, cheaper and more widely accessible
422 method for PAI estimation, and while TLS promises to alleviate potential bias in DHP estimates, results are highly
423 methods dependent. Our results demonstrate the challenges that stand in the way of large scale adoption of TLS
424 for vegetation indices monitoring.

425 **6. Code availability**

426 See https://github.com/will-flynn/tls_dhp_pai.git for all processing and modelling code.

427 **7. Data availability**

428 See Owen et al., (2022) for individual segmented tree data.

429 **8. Author contribution**

430 All authors designed the study. HJFO and WRMF collected and processed TLS and DHP data; WRMF performed
431 formal analysis with guidance from all authors. WRMF led the writing with input from all authors. All authors
432 contributed critically to drafts and gave final approval for publication.

433 **9. Competing interests**

434 The authors declare that they have no conflict of interest.

435 **7. Acknowledgements**

436 WRMF was funded through a London NERC DTP PhD studentship. ERL, HJFO and SWDG were funded through
437 the UKRI Future Leaders Fellowship awarded to ERL (MR/T019832/1).

438 **References**

- 439 Baeten, L., Verheyen, K., Wirth, C., Bruelheide, H., Bussotti, F., Finér, L., Jaroszewicz, B., Selvi, F.,
440 Valladares, F., Allan, E., Ampoorter, E., Auge, H., Avăcăriei, D., Barbaro, L., Bărnoaiea, I., Bastias, C. C.,
441 Bauhus, J., Beinhoff, C., Benavides, R., Benneter, A., Berger, S., Berthold, F., Boberg, J., Bonal, D.,
442 Brüggemann, W., Carnol, M., Castagneyrol, B., Charbonnier, Y., Češko, E., Coomes, D., Coppi, A., Dalmaris,
443 E., Dănilă, G., Dawud, S. M., de Vries, W., De Wandeler, H., Deconchat, M., Domisch, T., Duduman, G.,
444 Fischer, M., Fotelli, M., Gessler, A., Gimeno, T. E., Granier, A., Grossiord, C., Guyot, V., Hantsch, L.,
445 Hättenschwiler, S., Hector, A., Hermy, M., Holland, V., Jactel, H., Joly, F.-X., Jucker, T., Kolb, S., Koricheva,
446 J., Lexer, M. J., Liebergesell, M., Milligan, H., Müller, S., Muys, B., Nguyen, D., Nichiforel, L., Pollastrini, M.,
447 Proulx, R., Rabasa, S., Radoglou, K., Ratcliffé, S., Raulund-Rasmussen, K., Seiferling, I., Stenlid, J., Vesterdal,
448 L., von Wilpert, K., Zavala, M. A., Zielinski, D., and Scherer-Lorenzen, M.: A novel comparative research
449 platform designed to determine the functional significance of tree species diversity in European forests,
450 *Persepect. Plant. Ecol.*, 15, 281–291, <https://doi.org/10.1016/j.ppees.2013.07.002>, 2013.
- 451 Baret, F., Weiss, M., Lacaze, R., Camacho, F., Makhmara, H., Pacholczyk, P., and Smets, B.: GEOV1: LAI and
452 FAPAR essential climate variables and FCOVER global time series capitalizing over existing products. Part I:
453 Principles of development and production, *Remote Sens. Environ.*, 137, 299–309,
454 <https://doi.org/10.1016/j.rse.2012.12.027>, 2013.



- 455 Bates, D., Mächler, M., Bolker, B., and Walker, S.: Fitting Linear Mixed-Effects Models Using lme4, *J. Stat.*
456 *Softw.*, 67, <https://doi.org/10.18637/jss.v067.i01>, 2015.
- 457 Battaglia, M., Cherry, M. L., Beadle, C. L., Sands, P. J., and Hingston, A.: Prediction of leaf area index in
458 eucalypt plantations: effects of water stress and temperature, *Tree Physiol.*, 18, 521–528,
459 <https://doi.org/10.1093/treephys/18.8-9.521>, 1998.
- 460 Beaudet, M. and Messier, C.: Growth and morphological responses of yellow birch, sugar maple, and beech
461 seedlings growing under a natural light gradient, *Can. J. Forest Res.*, 28, 1007–1015,
462 <https://doi.org/10.1139/x98-077>, 1998.
- 463 Béland, M., Baldocchi, D. D., Widlowski, J.-L., Fournier, R. A., and Verstraete, M. M.: On seeing the wood
464 from the leaves and the role of voxel size in determining leaf area distribution of forests with terrestrial LiDAR,
465 *Agr. Forest Meteorol.*, 184, 82–97, <https://doi.org/10.1016/j.agrformet.2013.09.005>, 2014.
- 466 Breda, N. J. J.: Ground-based measurements of leaf area index: a review of methods, instruments and current
467 controversies, *J. Exp. Bot.*, 54, 2403–2417, <https://doi.org/10.1093/jxb/erg263>, 2003.
- 468 Burt, A., Disney, M., and Calders, K.: Extracting individual trees from lidar point clouds using treeseg, *Methods*
469 *Ecol. Evol.*, 10, 438–445, <https://doi.org/10.1111/2041-210X.13121>, 2019.
- 470 Calders, K., Armston, J., Newnham, G., Herold, M., and Goodwin, N.: Implications of sensor configuration and
471 topography on vertical plant profiles derived from terrestrial LiDAR, *Agr. Forest Meteorol.*, 194, 104–117,
472 <https://doi.org/10.1016/j.agrformet.2014.03.022>, 2014.
- 473 Calders, K., Origo, N., Disney, M., Nightingale, J., Woodgate, W., Armston, J., and Lewis, P.: Variability and
474 bias in active and passive ground-based measurements of effective plant, wood and leaf area index, *Agr. Forest*
475 *Meteorol.*, 252, 231–240, <https://doi.org/10.1016/j.agrformet.2018.01.029>, 2018.
- 476 Canham, C. D.: Growth and Canopy Architecture of Shade-Tolerant Trees: Response to Canopy Gaps, *Ecology*,
477 69, 786–795, <https://doi.org/10.2307/1941027>, 1988.
- 478 Carter, J. L. and White, D. A.: Plasticity in the Huber value contributes to homeostasis in leaf water relations of
479 a mallee Eucalypt with variation to groundwater depth, *Tree Physiol.*, 29, 1407–1418,
480 <https://doi.org/10.1093/treephys/tpp076>, 2009.
- 481 Castro-Díez, P., Villar-Salvador, P., Pérez-Rontomé, C., Maestro-Martínez, M., and Monserrat-Martí, G.: Leaf
482 morphology and leaf chemical composition in three *Quercus* (Fagaceae) species along a rainfall gradient in NE
483 Spain, *Trees*, 11, 127–134, <https://doi.org/10.1007/PL00009662>, 1997.
- 484 Chen, J. M. and Black, T. A.: Defining leaf area index for non-flat leaves, *Plant Cell Environ.*, 15, 421–429,
485 <https://doi.org/10.1111/j.1365-3040.1992.tb00992.x>, 1992.



- 486 Disney, M.: Terrestrial LiDAR: a three-dimensional revolution in how we look at trees, *New Phytol.*, 222,
487 1736–1741, <https://doi.org/10.1111/nph.15517>, 2018.
- 488 Gazal, R. M., Scott, R. L., Goodrich, D. C., and Williams, D. G.: Controls on transpiration in a semiarid riparian
489 cottonwood forest, *Agr. Forest Meteorol.*, 137, 56–67, <https://doi.org/10.1016/j.agrformet.2006.03.002>, 2006.
- 490 Gower, S. T., Vogel, J. G., Norman, J. M., Kucharik, C. J., Steele, S. J., and Stow, T. K.: Carbon distribution
491 and aboveground net primary production in aspen, jack pine, and black spruce stands in Saskatchewan and
492 Manitoba, Canada, *J. Geophys. Res.*, 102, 29029–29041, <https://doi.org/10.1029/97JD02317>, 1997.
- 493 Gower, S. T., Kucharik, C. J., and Norman, J. M.: Direct and Indirect Estimation of Leaf Area Index, fAPAR,
494 and Net Primary Production of Terrestrial Ecosystems, *Remote Sens. Environ.*, 70, 29–51,
495 [https://doi.org/10.1016/S0034-4257\(99\)00056-5](https://doi.org/10.1016/S0034-4257(99)00056-5), 1999.
- 496 Grotti, M., Calders, K., Origo, N., Puletti, N., Alivernini, A., Ferrara, C., and Chianucci, F.: An intensity, image-
497 based method to estimate gap fraction, canopy openness and effective leaf area index from phase-shift terrestrial
498 laser scanning, *Agr. Forest Meteorol.*, 280, 107766, <https://doi.org/10.1016/j.agrformet.2019.107766>, 2020.
- 499 Hardwick, S. R., Toumi, R., Pfeifer, M., Turner, E. C., Nilus, R., and Ewers, R. M.: The relationship between
500 leaf area index and microclimate in tropical forest and oil palm plantation: Forest disturbance drives changes in
501 microclimate, *Agr. Forest Meteorol.*, 201, 187–195, <https://doi.org/10.1016/j.agrformet.2014.11.010>, 2015.
- 502 Hijmans, R. J.: raster: Geographic Data Analysis and Modeling R package version 3.5-21, [https://CRAN.R-](https://CRAN.R-project.org/package=raster)
503 [project.org/package=raster.](https://CRAN.R-project.org/package=raster), 2022.
- 504 Hosoi, F. and Omasa, K.: Voxel-Based 3-D Modeling of Individual Trees for Estimating Leaf Area Density
505 Using High-Resolution Portable Scanning Lidar, *IEEE T. Geosci. Remote*, 44, 3610–3618,
506 <https://doi.org/10.1109/TGRS.2006.881743>, 2006.
- 507 Itakura, K. and Hosoi, F.: Voxel-based leaf area estimation from three-dimensional plant images, *J. Agric.*
508 *Meteorol.*, 75, 211–216, <https://doi.org/10.2480/agrmet.d-19-00013>, 2019.
- 509 Jonckheere, I., Fleck, S., Nackaerts, K., Muys, B., Coppin, P., Weiss, M., and Baret, F.: Review of methods for
510 in situ leaf area index determination, *Agr. Forest Meteorol.*, 121, 19–35,
511 <https://doi.org/10.1016/j.agrformet.2003.08.027>, 2004.
- 512 Jonckheere, I. G. C., Muys, B., and Coppin, P.: Allometry and evaluation of in situ optical LAI determination in
513 Scots pine: a case study in Belgium, *Tree Physiol.*, 25, 723–732, <https://doi.org/10.1093/treephys/25.6.723>,
514 2005.
- 515 Jucker, T., Bouriaud, O., Avacaritei, D., Dănilă, I., Duduman, G., Valladares, F., and Coomes, D. A.:
516 Competition for light and water play contrasting roles in driving diversity-productivity relationships in Iberian
517 forests, *J. Ecol.*, 102, 1202–1213, <https://doi.org/10.1111/1365-2745.12276>, 2014.



- 518 Jump, A. S., Ruiz-Benito, P., Greenwood, S., Allen, C. D., Kitzberger, T., Fensham, R., Martínez-Vilalta, J.,
519 and Lloret, F.: Structural overshoot of tree growth with climate variability and the global spectrum of drought-
520 induced forest dieback, *Glob. Change Biol.*, 23, 3742–3757, <https://doi.org/10.1111/gcb.13636>, 2017.
- 521 Jupp, D. L. B., Culvenor, D. S., Lovell, J. L., Newnham, G. J., Strahler, A. H., and Woodcock, C. E.: Estimating
522 forest LAI profiles and structural parameters using a ground-based laser called 'Echidna(R), *Tree Physiol.*, 29,
523 171–181, <https://doi.org/10.1093/treephys/tpn022>, 2008.
- 524 Kamoske, A. G., Dahlin, K. M., Stark, S. C., and Serbin, S. P.: Leaf area density from airborne LiDAR:
525 Comparing sensors and resolutions in a temperate broadleaf forest ecosystem, *Forest Ecol. Manag.*, 433, 364–
526 375, <https://doi.org/10.1016/j.foreco.2018.11.017>, 2019.
- 527 Kuusk, V., Niinemets, Ü., and Valladares, F.: A major trade-off between structural and photosynthetic
528 investments operative across plant and needle ages in three Mediterranean pines, *Tree Physiol.*, 38, 543–557,
529 <https://doi.org/10.1093/treephys/tpx139>, 2018.
- 530 Leblanc, S. G. and Chen, J. M.: A practical scheme for correcting multiple scattering effects on optical LAI
531 measurements, *Agr. Forest Meteorol.*, 110, 125–139, [https://doi.org/10.1016/S0168-1923\(01\)00284-2](https://doi.org/10.1016/S0168-1923(01)00284-2), 2001.
- 532 Lecigne, B., Delagrange, S., and Messier, C.: Exploring trees in three dimensions: VoxR, a novel voxel-based R
533 package dedicated to analysing the complex arrangement of tree crowns, *Ann. Bot-London*, 121, 589–601,
534 <https://doi.org/10.1093/aob/mcx095>, 2018.
- 535 Li, S., Dai, L., Wang, H., Wang, Y., He, Z., and Lin, S.: Estimating Leaf Area Density of Individual Trees
536 Using the Point Cloud Segmentation of Terrestrial LiDAR Data and a Voxel-Based Model, *Remote Sens-Basel*,
537 9, 1202, <https://doi.org/10.3390/rs9111202>, 2017.
- 538 Li, Y., Guo, Q., Tao, S., Zheng, G., Zhao, K., Xue, B., and Su, Y.: Derivation, Validation, and Sensitivity
539 Analysis of Terrestrial Laser Scanning-Based Leaf Area Index, *Can. J. Remote Sens.*, 42, 719–729,
540 <https://doi.org/10.1080/07038992.2016.1220829>, 2016.
- 541 Lines, E. R., Fischer, F. J., Owen, H. J. F., and Jucker, T.: The shape of trees: Reimagining forest ecology in
542 three dimensions with remote sensing, *J. Ecol.*, 110, 1730–1745, <https://doi.org/10.1111/1365-2745.13944>,
543 2022.
- 544 Long, J. N. and Smith, F. W.: Leaf area - sapwood area relations of lodgepole pine as influenced by stand
545 density and site index., *Can. J. Forest Res.*, 18, 247–250, 1988.
- 546 López, R., Cano, F. J., Martin-StPaul, N. K., Cochard, H., and Choat, B.: Coordination of stem and leaf traits
547 define different strategies to regulate water loss and tolerance ranges to aridity, *New Phytol.*, 230, 497–509,
548 <https://doi.org/10.1111/nph.17185>, 2021.
- 549 Lovell, J. L., Jupp, D. L. B., van Gorsel, E., Jimenez-Berni, J., Hopkinson, C., and Chasmer, L.: Foliage Profiles
550 from Ground Based Waveform and Discrete Point Lidar, *SilviLaser*, 1–9, 2011.



- 551 Ma, L., Zheng, G., Eitel, J. U. H., Magney, T. S., and Moskal, L. M.: Determining woody-to-total area ratio
552 using terrestrial laser scanning (TLS), *Agr. Forest Meteorol.*, 228–229, 217–228,
553 <https://doi.org/10.1016/j.agrformet.2016.06.021>, 2016.
- 554 Madrigal-González, J., Herrero, A., Ruiz-Benito, P., and Zavala, M. A.: Resilience to drought in a dry forest:
555 Insights from demographic rates, *Forest Ecol. Manag.*, 389, 167–175,
556 <https://doi.org/10.1016/j.foreco.2016.12.012>, 2017.
- 557 Magnani, F., Mencuccini, M., and Grace, J.: Age-related decline in stand productivity: the role of structural
558 acclimation under hydraulic constraints, *Plant Cell Environ.*, 23, 251–263, [https://doi.org/10.1046/j.1365-](https://doi.org/10.1046/j.1365-3040.2000.00537.x)
559 [3040.2000.00537.x](https://doi.org/10.1046/j.1365-3040.2000.00537.x), 2000.
- 560 Mencuccini, M.: The ecological significance of long-distance water transport: short-term regulation, long-term
561 acclimation and the hydraulic costs of stature across plant life forms, *Plant Cell Environ.*, 26, 163–182,
562 <https://doi.org/10.1046/j.1365-3040.2003.00991.x>, 2003.
- 563 Mencuccini, M. and Grace, J.: Climate influences the leaf area/sapwood area ratio in Scots pine, *Tree Physiol.*,
564 15, 1–10, <https://doi.org/10.1093/treephys/15.1.1>, 1995.
- 565 Monsi, M. and Saeki, T.: On the Factor Light in Plant Communities and its Importance for Matter Production,
566 *Ann. Bot-London*, 95, 549–567, <https://doi.org/10.1093/aob/mci052>, 1953.
- 567 Nakagawa, S., Johnson, P. C. D., and Schielzeth, H.: The coefficient of determination R² and intra-class
568 correlation coefficient from generalized linear mixed-effects models revisited and expanded, *J. R. Soc.*
569 *Interface*, 14, 20170213, <https://doi.org/10.1098/rsif.2017.0213>, 2017.
- 570 Niinemets, Ü. and Valladares, F.: Tolerance to shade, drought, and waterlogging of temperate northern
571 hemisphere trees and shrubs, *Ecol. Monogr.*, 76, 521–547, [https://doi.org/10.1890/0012-](https://doi.org/10.1890/0012-9615(2006)076[0521:TTSDAW]2.0.CO;2)
572 [9615\(2006\)076\[0521:TTSDAW\]2.0.CO;2](https://doi.org/10.1890/0012-9615(2006)076[0521:TTSDAW]2.0.CO;2), 2006.
- 573 Niu, X., Fan, J., Luo, R., Fu, W., Yuan, H., and Du, M.: Continuous estimation of leaf area index and the
574 woody-to-total area ratio of two deciduous shrub canopies using fisheye webcams in a semiarid loessial region
575 of China, *Ecol. Indic.*, 125, 107549, <https://doi.org/10.1016/j.ecolind.2021.107549>, 2021.
- 576 Olivas, P. C., Oberbauer, S. F., Clark, D. B., Clark, D. A., Ryan, M. G., O’Brien, J. J., and Ordoñez, H.:
577 Comparison of direct and indirect methods for assessing leaf area index across a tropical rain forest landscape,
578 *Agr. Forest Meteorol.*, 177, 110–116, <https://doi.org/10.1016/j.agrformet.2013.04.010>, 2013.
- 579 Owen, H. J. F., Flynn, W. R. M., and Lines, E. R.: Competitive drivers of inter-specific deviations of crown
580 morphology from theoretical predictions measured with Terrestrial Laser Scanning, *J. Ecol.*, 109, 2612–2628,
581 <https://doi.org/10.1111/1365-2745.13670>, 2021.
- 582 Owen, H. J. F., Flynn, W. R. M., and Lines, E. R.: Individual TLS tree clouds collected from both Alto Tajo and
583 Cuellar in Spain., 2022.



- 584 Peppe, D. J., Royer, D. L., Cariglino, B., Oliver, S. Y., Newman, S., Leight, E., Enikolopov, G., Fernandez-
585 Burgos, M., Herrera, F., Adams, J. M., Correa, E., Currano, E. D., Erickson, J. M., Hinojosa, L. F., Hoganson, J.
586 W., Iglesias, A., Jaramillo, C. A., Johnson, K. R., Jordan, G. J., Kraft, N. J. B., Lovelock, E. C., Lusk, C. H.,
587 Niinemets, Ü., Peñuelas, J., Rapson, G., Wing, S. L., and Wright, I. J.: Sensitivity of leaf size and shape to
588 climate: global patterns and paleoclimatic applications, *New Phytol.*, 190, 724–739,
589 <https://doi.org/10.1111/j.1469-8137.2010.03615.x>, 2011.
- 590 Pfeifer, M., Gonsamo, A., Disney, M., Pellikka, P., and Marchant, R.: Leaf area index for biomes of the Eastern
591 Arc Mountains: Landsat and SPOT observations along precipitation and altitude gradients, *Remote Sens.*
592 *Environ.*, 118, 103–115, <https://doi.org/10.1016/j.rse.2011.11.009>, 2012.
- 593 Phillips, N., Bond, B. J., McDowell, N. G., Ryan, M. G., and Schauer, A.: Leaf area compounds height-related
594 hydraulic costs of water transport in Oregon White Oak trees, *Funct. Ecol.*, 17, 832–840,
595 <https://doi.org/10.1111/j.1365-2435.2003.00791.x>, 2003.
- 596 Purves, D. and Pacala, S.: Predictive Models of Forest Dynamics, *Science*, 320, 1452–1453,
597 <https://doi.org/10.1126/science.1155359>, 2008.
- 598 Ridler, T. W. and Calvard, S.: Picture Thresholding Using an Iterative Selection Method, *IEEE T. Syst. Man.*
599 *Cyb.*, 8, 630–632, <https://doi.org/10.1109/TSMC.1978.4310039>, 1978.
- 600 Sea, W. B., Choler, P., Beringer, J., Weinmann, R. A., Hutley, L. B., and Leuning, R.: Documenting
601 improvement in leaf area index estimates from MODIS using hemispherical photos for Australian savannas,
602 *Agr. Forest Meteorol.*, 151, 1453–1461, <https://doi.org/10.1016/j.agrformet.2010.12.006>, 2011.
- 603 Specht, R. L. and Specht, A.: Canopy structure in Eucalyptus-dominated communities in Australia along
604 climatic gradients, *Canopy structure in Eucalyptus-dominated communities in Australia along climatic*
605 *gradients*, 10, 191–213, 1989.
- 606 Vicari, M. B., Disney, M., Wilkes, P., Burt, A., Calders, K., and Woodgate, W.: Leaf and wood classification
607 framework for terrestrial LiDAR point clouds, *Methods Ecol. Evol.*, 10, 680–694, [https://doi.org/10.1111/2041-](https://doi.org/10.1111/2041-210X.13144)
608 [210X.13144](https://doi.org/10.1111/2041-210X.13144), 2019.
- 609 Warton, D. I., Wright, I. J., Falster, D. S., and Westoby, M.: Bivariate line-fitting methods for allometry, *Biol.*
610 *Rev.*, 81, 259–291, <https://doi.org/10.1017/S1464793106007007>, 2006.
- 611 Warton, D. I., Duursma, R. A., Falster, D. S., and Taskinen, S.: smatr 3 - an R package for estimation and
612 inference about allometric lines: *The smatr 3 - an R package*, *Methods Ecol. Evol.*, 3, 257–259,
613 <https://doi.org/10.1111/j.2041-210X.2011.00153.x>, 2012.
- 614 Weiss, M., Baret, F., Smith, G. J., Jonckheere, I., and Coppin, P.: Review of methods for in situ leaf area index
615 (LAI) determination, *Agr. Forest Meteorol.*, 121, 37–53, <https://doi.org/10.1016/j.agrformet.2003.08.001>, 2004.



- 616 Whitehead, D.: The Estimation of Foliage Area from Sapwood Basal Area in Scots Pine, *Forestry*, 51, 137–149,
617 <https://doi.org/10.1093/forestry/51.2.137>, 1978.
- 618 Wilkes, P., Lau, A., Disney, M., Calders, K., Burt, A., Gonzalez de Tanago, J., Bartholomeus, H., Brede, B., and
619 Herold, M.: Data acquisition considerations for Terrestrial Laser Scanning of forest plots, *Remote Sensing of*
620 *Environment*, 196, 140–153, <https://doi.org/10.1016/j.rse.2017.04.030>, 2017.
- 621 Woodgate, W., Jones, S. D., Suarez, L., Hill, M. J., Armston, J. D., Wilkes, P., Soto-Berelov, M., Haywood, A.,
622 and Mellor, A.: Understanding the variability in ground-based methods for retrieving canopy openness, gap
623 fraction, and leaf area index in diverse forest systems, *Agr. Forest Meteorol.*, 205, 83–95,
624 <https://doi.org/10.1016/j.agrformet.2015.02.012>, 2015.
- 625 Woodgate, W., Armston, J. D., Disney, M., Jones, S. D., Suarez, L., Hill, M. J., Wilkes, P., and Soto-Berelov,
626 M.: Quantifying the impact of woody material on leaf area index estimation from hemispherical photography
627 using 3D canopy simulations, *Agr. Forest Meteorol.*, 226–227, 1–12,
628 <https://doi.org/10.1016/j.agrformet.2016.05.009>, 2016.
- 629 Zhang, Y., Chen, J. M., and Miller, J. R.: Determining digital hemispherical photograph exposure for leaf area
630 index estimation, *Agr. Forest Meteorol.*, 133, 166–181, <https://doi.org/10.1016/j.agrformet.2005.09.009>, 2005.
- 631 Zheng, G., Moskal, L. M., and Kim, S.-H.: Retrieval of Effective Leaf Area Index in Heterogeneous Forests
632 With Terrestrial Laser Scanning, *IEEE T. Geosci. Remote*, 51, 777–786,
633 <https://doi.org/10.1109/TGRS.2012.2205003>, 2013.
- 634 Zhu, X., Skidmore, A. K., Wang, T., Liu, J., Darvishzadeh, R., Shi, Y., Premier, J., and Heurich, M.: Improving
635 leaf area index (LAI) estimation by correcting for clumping and woody effects using terrestrial laser scanning,
636 *Agr. Forest Meteorol.*, 263, 276–286, <https://doi.org/10.1016/j.agrformet.2018.08.026>, 2018.
- 637