

# High aboveground carbon stock of African tropical montane forests

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#### 1 High above-ground carbon stock of African tropical montane forests

2 3

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- 18 Ngute<sup>61,82</sup>, Douglas Sheil<sup>83</sup>, Demisse Sheleme<sup>84</sup>, Tibebu Y. Simegn<sup>85</sup>, Murielle Simo-Droissart<sup>22</sup>,
- 19 Bonaventure Sonké<sup>22</sup>, Teshome Soromessa<sup>16</sup>, Terry Sunderland<sup>86,87</sup>, Miroslav Svoboda<sup>88</sup>, Hermann
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- 22 Charlotte Wheeler<sup>64</sup>, Simon Willcock<sup>98,99</sup>, John T. Woods<sup>100</sup>, Etienne Zibera<sup>68</sup>
- 23

#### 24 Abstract

- 25 Tropical forests store 40-50% of terrestrial vegetation carbon<sup>1</sup>. Spatial variations in aboveground live
- tree biomass carbon (AGC) stocks remain poorly understood, in particular in tropical montane
- 27 forests<sup>2</sup>. Owing to climatic and soil changes with increasing elevation<sup>3</sup>, AGC stocks are lower in
- tropical montane compared to lowland forests<sup>2</sup>. Here we assemble and analyse a dataset of
- 29 structurally intact old-growth forests (AfriMont) spanning 44 montane sites in 12 African countries.
- 30 We find that montane sites in the AfriMont plot network have a mean AGC-stock of 149.4 Mg C ha<sup>-1</sup>
- 31 (95% CI 137.1-164.2), comparable to lowland forests in the African Tropical Rainforest Observation
- 32 Network<sup>4</sup> and about 70 per cent and 32 per cent higher than averages from plot networks in
- montane<sup>2,5,6</sup> and lowland<sup>7</sup> forests in the Neotropics, respectively. Notably, our results are two-thirds
- higher than the IPCC default values for these forests in Africa<sup>8</sup>. We find that the low stem density
- and high abundance of large trees of African lowland forests<sup>4</sup> is mirrored in the montane forests
- 36 sampled. This carbon store is endangered: we estimate that 0.8 million ha of old-growth African
- 37 montane forest have been lost since 2000. We provide country-specific montane forest AGC stock
- 38 estimates modelled from our plot network to help guide forest conservation and reforestation
- 39 interventions. Our findings highlight the need for conserving these biodiverse<sup>9,10</sup> and carbon-rich
- 40 ecosystems.
- 41
- 42

#### 43 Main text

- Tropical forests cover less than 10% of the global land area yet store 40–50% of terrestrial 44
- vegetation carbon<sup>1</sup> and contribute more than one third of primary productivity<sup>11</sup> so are a key 45
- component of the global carbon cycle<sup>12,13</sup>. There is substantial variation in carbon stocks across the 46
- biome, with lowland forests in Africa and Borneo storing more carbon per unit area than lowland 47
- 48 forests in the Neotropics<sup>4,7</sup>. This variation arises partly from structural differences: the signature
- 49 feature of African forests is their low stem density but relatively high abundance of large trees (>70
- 50 cm diameter) which store large quantities of carbon, while Bornean forests are characterised by high
- 51 stem density and basal area<sup>4,14,15</sup>.
- 52

53 Despite increased understanding of biogeographic differences in tropical lowland forests, patterns of 54 spatial variation in carbon stocks remain poorly understood in the 880,000 km<sup>2</sup> of tropical montane 55 forests located  $\geq$  1,000 m asl<sup>2</sup>. Montane forests are expected *a priori* to have lower aboveground live 56 tree biomass carbon (AGC) stocks than lowland forests because (1) temperature decreases with 57 increasing elevation, reducing net primary productivity and slowing nutrient recycling, (2) long 58 periods of cloud immersion in montane forests suppresses productivity, (3) soil waterlogging slows 59 nutrient recycling and (4) high epiphyte load, local wind exposure in crests and nutrient-limited soils 60 limit tree size and increase investment in roots over shoots<sup>3</sup>. While forest inventory plots provide 61 some support for these assumptions<sup>2</sup> data from African mountain regions are exceptionally sparse. 62 Indeed, in the most recent IPCC guidelines, there is no specific AGC default value for old-growth montane forests in Africa: the value given of 89.3 Mg C ha<sup>-1</sup> is simply a mean of secondary and old-63 growth forests found  $\geq$  1,000 m asl<sup>8</sup>. Mountain areas also pose special challenges for remote-sensing 64 65 approaches for estimating carbon stocks, as radar data are affected by geometric distortions<sup>16</sup> while steep slopes bias spaceborne LiDAR estimates towards overestimating canopy height<sup>17</sup>. These issues 66 67 are reflected in the limited correlation between estimates of AGC-stocks at mountain locations from 68 different recent remote-sensing derived carbon maps (Supplementary Information Table S1).

69

70 Better understanding of montane carbon stocks is important for many African countries, particularly 71 in eastern Africa where montane forests represent most of the extant evergreen old-growth forest 72 cover. Quantifying carbon stocks in these ecosystems is critical for estimating national carbon losses from deforestation and forest degradation<sup>18</sup>. Quantifying carbon stocks in old-growth montane 73 74 forests also serves to constrain potential carbon uptake by restored natural forests, given the high

75 commitment of most African nations to the Bonn Challenge effort to restore 150 million ha of

76 degraded and deforested lands by 2020 (see Table 1), and 350 million by 2030.

77

78 Here we measured, compiled and analysed a new dataset of 226 plot inventories spanning 44 sites 79 in 12 African countries, covering most major mountain regions on the continent (the "AfriMont" 80 dataset). Plots range from 800 to 3,900 m asl to include submontane forests (800-1,000 m asl) in smaller mountains closer to the ocean<sup>19,20</sup>. For all plots, stem diameter and species were recorded 81 82 for each tree  $\geq 10$  cm diameter at breast height (or above buttress) following standard methods<sup>21</sup>. 83 Tree height was sampled in 23 montane sites, allowing variation in height-diameter allometry to be 84 incorporated into the calculation of aboveground biomass. A total of 72,336 stems with diameter 85  $\geq$ 10 cm were measured. For each tree, we computed AGC (in Mg C ha<sup>-1</sup>) according to standard 86 procedures (see methods).

87

88 We find that the mean plot-level AGC-stock across sampled African tropical montane forests is 149.4 89 Mg C ha<sup>-1</sup> (95% confidence interval (CI) 137.1-164.2), two-thirds more than the IPCC default value of 90 89.3 Mg C ha<sup>-1</sup>. Our estimates are robust to subsampling our dataset (Extended Data Fig. 1) and 91 excluding small plots (Extended Data Fig. 2) and are not affected by the sampling strategy used to

- 92 establish plots in each study site (Extended Data Fig. 2). Comparing our dataset to previous
- 93 syntheses of montane<sup>2,5,6</sup> and lowland<sup>7</sup> forest plot networks reveals that tropical montane forests in

- 94 Africa have significantly higher AGC-stocks per unit area than both montane (95% CI = 50.4 – 71.9 95 Mg C ha<sup>-1</sup>) and lowland (95% CI = 124.0 – 147.9 Mg C ha<sup>-1</sup>) forests in the Neotropics, and that they 96 do not differ significantly from lowland forests in Africa (95% CI = -27.6 - 9.6 Mg C ha<sup>-1</sup>) (Fig. 1, Table 97 S2). The similar AGC-stocks in montane and lowland forests in Africa contrasts with the Neotropics 98 and Southeast Asia, where carbon stocks are lower in montane forests than lowland forests (albeit 99 not significantly different in Southeast Asia due to the small sample size, Fig. 1). These differences 100 are robust to accounting for differences in elevation among montane datasets: removing African 101 plots 800-1,000 m asl slightly reduces estimated montane forest AGC-stock to 145.0 Mg C ha<sup>-1</sup> (95% 102 Cl 129.6 – 163.2), but observed differences in AGC-stock among continents remain when plots are 103 restricted to elevations well represented in all continents (Extended Data Fig. 3). 105 The characteristic structural properties of lowland African forests (relatively low stem density and
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- greater importance of large trees compared to elsewhere in the tropics<sup>4</sup>) are also evident in the African montane forests we sampled. In these montane forests mean stem density is 483.3 stems ha <sup>1</sup> (± 177.7 s.d.) and mean basal area is 39 m<sup>2</sup>ha<sup>-1</sup> (± 14.8 s.d.). We find a high density of large stems  $(>70 \text{ cm diameter}, 19.1 \text{ stems ha}^{-1} \pm 15.4 \text{ s.d.})$  which contribute 35.3% (95% CI = 29.6 – 41.8 %) to plot-level AGC-stock (Fig. 2). The contribution of large trees to plot-level AGC-stock is also similar in montane and lowland Africa (95% CI of difference in square-root transformed proportional
- 112 contribution of large trees between lowland and montane forests = -0.100 - 0.075, P = 0.80). There
- 113 was no significant difference in the proportional contribution of any other size class to AGC-stocks
- 114 between our montane dataset and 132 lowland plots from the AfriTRON network ( $P \ge 0.24$ , Table S3),
- 115 although greater variation among plots is observed in montane forests (Fig. 2).
- 116

117 To investigate if elevation affected AGC or forest structure, we modelled these variables as functions 118 of elevation using random slopes mixed-effects models. This approach allows intercepts and 119 relationships to vary among sites, which would be expected as mountains can have very different 120 climate at the same elevation due to proximity to the ocean (generally the further, the drier) and because of the mass-elevation or telescopic effect<sup>22</sup> (larger mountains are better at warming the 121 atmosphere above them). We found that AGC, stem density or density of large stems (>70 cm 122 123 diameter) were not significantly related to elevation (Fig. 3, Table S4). Across sites these non-124 significant relationships were all negative, although there was some variation in strength and 125 direction amongst sites (Fig. 3). Similarly, in the Neotropics and Southeast Asia montane forest plot

126 datasets, AGC was not significantly correlated with elevation (Extended Data Fig. 4).

127

128 To assess potential environmental drivers of AGC-stock variation across the AfriMont plot network, 129 we related AGC to climate, soil and topography. We found that AGC-stocks increased with annual 130 precipitation (albeit not statistically significantly), decreased with soil fertility and were higher in 131 plots which were locally at higher elevation than their surroundings (Extended Data Fig. 5). 132 Relationships with other environmental variables were non-significant (Extended Data Fig. 5).

- 133 Although global datasets might not capture fine-scale variation in climate or soils in mountain
- regions<sup>23</sup>, leading to regression dilution<sup>24</sup>, the general absence of strong climate effects combined 134
- 135 with the lack of significant effect of elevation on AGC-stocks suggest that the high AGC-stock of 136 African montane forests is a pervasive phenomenon across a wide environmental gradient.
- 137

138 Although the AfriMont dataset covers most major mountain areas in tropical Africa (Fig. 4), some 139 areas remain under-sampled relative to forest extents (Extended Data Fig. 6), resulting in some

- 140 differences between the environmental conditions sampled by our plot network and the wider
- montane forest biome in Africa (Extended Data Fig. 7). Notably, the absence of plots from montane 141
- 142 forests of eastern Democratic Republic of the Congo (Fig. 4, Extended Data Fig. 6) means that the
- 143 AfriMont dataset samples forests are, on average, at higher elevations, and that are cooler and 144 cloudier than the wider montane forest biome in Africa (Extended Data Fig. 7). Using relationships

- 145 with environmental variables (Extended Data Fig. 5) to predict AGC-stocks in each 1-km grid cell 146 containing montane forest gives a mean (weighted by remaining forest cover) AGC-stock of 176.9 Mg C ha<sup>-1</sup> ( $\pm$  32.0 s.d.) for the tropical montane forest biome in Africa. This indicates that the 147
- 148 estimate we report based on our AfriMont plot network data (149.4 Mg C ha<sup>-1</sup>) is conservative.
- 149
- 150 Several mechanisms could explain the high AGC-stock of montane forests in the AfriMont plot 151 network. Firstly, large herbivores such as elephants (Loxodonta spp.) can have profound effects on 152 forest structure by consuming biomass, destroying small stems, dispersing seeds and transporting 153 nutrients<sup>25</sup>. Studies for lowland forests suggest that elephants can increase carbon stocks<sup>26,27</sup>. We tested if AfriMont plots with known elephant presence as of 2019 had significantly higher AGC-154 155 stocks, but found that they had significantly lower AGC-stocks, although significant differences were not observed in some countries (Extended Data Fig. 8). While the initial ecosystem response to 156 157 elephant removal might be greater AGC-stocks due to reduced biomass consumption and small-stem 158 destruction, the longer-term effects might differ. We were unable to fully disentangle such effects, 159 as we lacked details on both i) time since elephant extirpation, and ii) elephant abundance and its
- 160 determinants (see Table S5).
- 161

162 A second potential explanation is a relatively low frequency of large-scale abiotic disturbances,

- 163 allowing trees time to grow large and stands to self-thin, as is seen in lowland African forests<sup>4</sup>. For
- example, tropical cyclones are largely absent in mainland Africa (except in Mozambique<sup>28</sup>) and lava 164
- flows are limited even in the active volcano of Mt Cameroon<sup>29</sup>. Although fine-scale variability in 165
- landslide risk limits comparisons across large spatial scales, there are fewer areas with high landslide 166
- 167 susceptibility in mountains in tropical Africa than in the Andes and most mountain ranges in Southeast Asia<sup>30</sup>. If forests have been ecologically stable over evolutionary timescales, tree species 168
- 169 may be adapted to grow slowly but potentially reaching great sizes<sup>31</sup>. On Mt Kilimanjaro
- Entandrophragma individuals reach enormous heights and ages<sup>32</sup>. This low frequency of large-scale 170
- 171 abiotic disturbances contrasts with the Andes and several mountains in Southeast Asia (e.g. Barisian
- 172 mountains in western Sumatra), which are tectonically active, so the trees there are adapted to
- 173 sudden disturbance followed by intense competition to get established and grow. Future monitoring
- 174 of the AfriMont plot network will help determine the extent to which the high biomass of African 175 tropical montane forests results from them being dynamic and productive, or adapted to stability.
- 176 A third potential explanation could be the presence of conifers<sup>33</sup>. Mixed conifer/broad-leaved forests 177 178 tend to have greater basal area than purely broad-leaved forests due to a more effective use of light and other resources<sup>34</sup>. Podocarpaceae can be found in montane forests across the tropics<sup>35</sup>. Despite 179 having fewer species in Africa than in other continents<sup>36</sup>, these could be more abundant at the site-180 181 level. However, there is no pantropical comparative study on Podocarpaceae abundance in tropical 182 montane forests. In our dataset there was no significant correlation between plot-level AGC-stock
- 183 and conifer (Podocarpaceae) abundance (Extended Data Fig. 9). Other explanations could be
- 184 continental differences in mountain terrain (more gentle slopes or plateau regions in Africa) or types
- 185 of montane forests investigated (less cloud forest existing/sampled in Africa). Within our dataset,
- 186 slope did not have a significant effect on AGC-stocks (Extended Data Fig. 5). Contrary to the Neotropics<sup>37</sup>, there is no high-resolution map of cloud forests available for Africa, so while we found 187
- 188 no relationship between AGC-stock and cloud frequency (Extended Data Fig. 5), we were unable to
- 189 investigate differences in AGC-stock between cloud forest vs non-cloud forest plots.
- 190
- To understand the policy implications of our findings for African countries, we calculated montane 191
- (≥800 m asl) forest cover change between 2000 and 2018, using forest cover from ref.<sup>38</sup> clipped to 192
- 'primary humid forest' from ref.<sup>39</sup>. We show that tropical montane forests represent most -or all-193
- 194 evergreen old-growth forests found in ten African countries (Fig. 4), and that the Democratic
- 195 Republic of the Congo has two thirds of the remaining 16 million ha of montane forests in Africa.

Over 0.8 million ha (5%) have been lost in Africa since 2001, with the highest losses in the
Democratic Republic of the Congo (536,000 ha), Uganda (65,000 ha) and Ethiopia (62,000 ha) (Fig. 4,
Table 1) have a since 2001, with the highest losses in the

198Table 1). In terms of percentage, Mozambique and Côte d'Ivoire lost over 20% of their montane

199 forests over this period (Fig. 4, Table 1). In some sites, however, a larger proportion of montane

forests was lost before 2000, e.g. in Taita Hills in Kenya<sup>40</sup>. If absolute country-level deforestation
 rates continue, a further 0.5 million ha of tropical montane forests will be lost by 2030.

202

203 African tropical montane forests are not only carbon-rich, but they also harbour some of the highest concentrations of biodiversity and endemism in the world<sup>9,10</sup>. They are important 'water towers' as, 204 located at the headwaters of numerous river systems, including the Congo and the Nile, they 205 206 regulate timing and magnitude of runoff<sup>9</sup>. They also regulate local temperatures<sup>41</sup> and provide 207 numerous other services to people in the surrounding landscapes<sup>9</sup>. Clearly, more should be done to 208 avoid the destruction of these important ecosystems. Logging, mining and clearing land for farming, 209 but also political unrest and militia presence have affected -and continue to affect- these forests, e.g. 210 in Itombwe Mts in the Democratic Republic of the Congo<sup>42</sup>. Protected areas are known to help reduce deforestation in the tropics<sup>43</sup>. Beyond protected areas, other forest conservation 211 212 mechanisms could be implemented, including effective carbon finance. Previous IPCC AGC-stock 213 estimates for montane forests in Africa (89.3 Mg C ha<sup>-1</sup>) may have contributed to low incentives for 214 carbon finance mechanisms in these ecosystems. Our study shows the far greater carbon storage 215 potential in these tropical montane forests, which will be even higher if soil carbon stocks are considered (e.g. > 200 Mg C ha<sup>-1</sup> of organic carbon occurs in the top 0-30 cm soil on Mt Cameroon<sup>44</sup> 216

- and in the Usambara Mts, Tanzania<sup>45</sup>).
- 218

219 As well as conserving the remaining montane forests, efforts to restore them are critical. Forest 220 restoration at one of our sites, Kibale National Park in Uganda, indicates the potential for rapid AGC 221 accumulation<sup>46</sup>. Our study shows the high potential AGC-stock these montane forests can attain. The possible co-benefits of forest restoration, notably water regulation, control of soil erosion and 222 223 landslides and biodiversity conservation should also be considered. Most African nations are 224 committed to the Bonn Challenge; Ethiopia leading with 15 million ha committed (Table 1). We 225 provide country-specific estimates of potential AGC-stocks based on forests sampled in the AfriMont 226 dataset to help guide such interventions (Table 1, Extended Data Fig. 10). Caution is needed when 227 scaling-up our estimates to the landscape scale, as not all forests are closed-canopy old-growth and 228 structurally intact. Remote sensing or ancillary data (landcover maps, spatial environmental data) 229 could be used to identify e.g. exotic plantations, degraded or bamboo forests, and thus help create detailed AGC maps at different spatial scales<sup>18,47</sup>. A closer collaboration between air-borne, space-230 231 borne and ground approaches (such as the AfriMont and AfriTRON plot networks) is key for accurate 232 quantification and monitoring of landscape-scale tropical forest AGC-stocks, particularly in mountain 233 regions.

234

Our newly compiled dataset and analysis provides a large-scale quantification of AGC-stock in
African tropical montane forests, indicating it to be on average substantially higher than previously
thought. While there is variation around this mean AGC-stock within and across sites, it is not
systematically related to elevation. Apart from helping refine country-level estimates, IPCC
guidelines and ground-calibration of remote-sensing estimates, continued on-the-ground monitoring
of the AfriMont plot network will help determine ecosystem dynamics and carbon residence time in
these extraordinarily carbon-rich forests, as well as their responses to climatic changes.





Fig. 1 | Pantropical variation in aboveground carbon stocks sampled by plot networks in montane 247 248 (≥ 800 m asl) and lowland (< 800 m asl) tropical forests. Data from this study for African montane 249 forests (n = 226 plots, this study), montane forests in the Neotropics (n = 131) and Southeast Asia (n= 32) are from ref.<sup>2,5,6</sup>, lowland forests in Africa (n = 290), the Neotropics (n = 416) and Southeast 250 251 Asia (n = 60) arewfrom ref.<sup>7</sup>. Coloured points show the AGC-stock in each plot, with point size 252 proportional to square-root plot area. Black points show means for each continent-elevation 253 category estimated using linear mixed-effects models with site as a random effect, and lines show 254 95% confidence intervals around means. Letters indicate significant differences between continent 255 elevation category combinations (linear mixed-effects models with site as a random effect, P < 0.05). 256



258Size class259Fig. 2 | Proportion of plot-level aboveground carbon stock and stems accounted for by each size

class in montane and in lowland forests in Africa. Statistically significant differences in contribution

of each size class between montane and lowland forest plot networks are shown by asterisks (linear

262 mixed-effects model, P < 0.05). NS = non-significant difference. Montane (n = 226), lowland (n = 132). The thick line shows the median, and boxes cover the interquartile range (IQR). Values > 1.5

- times IQR away from the IQR are shown by points.
- 265
- 266







281 Fig. 4. | Old-growth evergreen humid forests in lowland and montane tropical Africa. Forest

extends circa 2018. Note that montane includes submontane forests (800-1,000 m asl, light purple).

283 Montane forests represent most (or all) evergreen humid old-growth forest in ten African nations:

284 Burundi, Ethiopia, Kenya, Rwanda, Tanzania, Uganda and Zimbabwe (included in AfriMont); and

285 Zambia, Malawi and South Sudan (no plot data available). Forest cover extracted from ref.<sup>38</sup> and

clipped to 'primary humid forest' using ref.<sup>39</sup>. See Table 1 for country-level absolute estimates.

288 Table 1 | Remaining forest area and aboveground carbon estimates for montane and lowland

289 tropical forests in Africa

290

Country	Montane (ha)	Montane lost (ha)	Montane AGC (MgC ha <sup>-1</sup> , 95% Cl)	Montane sites (plots)	Lowland (ha)	Lowland AGC (MgC ha <sup>-1</sup> , 95% Cl)	Lowland plots	Bonn Challenge by 2020 (ha)
Burundi	25,000	300	<b>94</b> (47-176)	1 (7)	0		0	2 million
Cameroon	840,000	30,200	<b>153</b> (121-195)	7 (37)	17.7 million	<b>166</b> (151-185)	72	12 million
DRC	10.2 million	536,500	<b>129</b> (84-202)	2 (37)	90 million	<b>158</b> (135-183)	48	8 million
Ethiopia	1.7 million	62,100	<b>165</b> (124-215)	8 (25)	145,000	а	0	15 million
Guinea	29,000	1,700	<b>314</b> (147-616) <sup>b</sup>	1 (2)	193,000	<b>157</b> (122 – 206) <sup>c</sup>	24	2 million
Kenya	568,000	44,100	<b>104</b> (79-136)	8 (38)	37,000		0	5.1 million
Mozambique	18,000	6,600 <sup>d</sup>	<b>226</b> (146-384) <sup>b</sup>	3 (4)	93,000	e	0	1 million
Nigeria	42,000	1,400	<b>120</b> (47-309) <sup>b</sup>	1 (1)	1.8 million	<b>161</b> (105-262)	2	4 million
Rwanda	53,000	300	<b>106</b> (65-168)	2 (11)	0		0	2 million
Tanzania	587,000	13,900	<b>175</b> (129-234)	6 (29)	130,000	<b>128</b> (101-163)	16	5.2 million
Uganda	427,000	64,600 <sup>d</sup>	<b>158</b> (111-209)	6 (23)	18,000		0	2.5 million
Zimbabwe	7,000	800 <sup>d</sup>	<b>203</b> (108-363)	1 (12)	<1,000		0	2 million

<sup>291</sup> 

292 Forest cover circa 2018 was extracted from ref.<sup>38</sup> and clipped to 'primary humid forest' using ref.<sup>39</sup>.

293 Montane forest lost covers the period 2000-2018. Mean aboveground carbon (AGC, in MgC ha<sup>-1</sup>)

estimates for montane (or lowland) forests were estimated from AfriMont and AfriTRON plot

295 network data. Mean AGC values are in boldface, 95% confidence intervals in parentheses. For details

on sites and plots used see Table S5. Bonn Challenge pledges for 2030 not yet available.

<sup>a</sup> Ref.<sup>48</sup> report 192 MgC ha<sup>-1</sup> for lowland.

<sup>b</sup> Few plots sampled, or very small plots sampled, AGC estimates may not be robust, see Extended

299 data Fig. 10.

300 <sup>c</sup> Data from neighbouring Liberia.

<sup>d</sup>Montane forest loss in Mozambique, Uganda and Zimbabwe represents 27%, 13% and 10% of the

302 existing montane forest in 2001, respectively. Montane forest loss in Côte d'Ivoire (no plot data

available) was estimated to be 21% for the same period.

<sup>e</sup> Ref.<sup>49</sup> report 132.2 MgC ha<sup>-1</sup> for lowland.

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#### 419 Methods

#### 420 AfriMont or montane Africa dataset

421 We compiled forest inventory plot data from the African Tropical Rainforest Observatory Network

- 422 (AfriTRON; <u>www.afritron.org</u>), with data curated at <u>www.ForestPlots.net</u><sup>50,51</sup> and the TEAM
- 423 network<sup>52</sup>, as well as from numerous site-specific publications detailed in Table S5 and mapped in
- 424 Fig. 4. Plots were selected for the analysis when conforming to the following criteria: ≥800 m asl,
- 425 closed-canopy evergreen wet or moist tropical forest, geo-referenced, old-growth and structurally
- 426 intact (not impacted by recent selective logging, fire or coffee cultivation), with no exotic species
- 427 present (e.g. *Eucalyptus* or *Pinus* spp.), all trees  $\geq$ 10 cm diameter measured and majority of stems 428 identified to encode *We* included plate from *Virunga* Massif in *Duranda* (Uganda even when not
- identified to species. We included plots from Virunga Massif in Rwanda/Uganda even when not
   100% closed-canopy due to high abundance of naturally-occurring bamboo. In all plots, tree
- 429 diameter was measured at 1.3 m along the stem from the ground, or above buttresses if present. In
- 431 23 sites tree height was sampled in the field for some stems, using a clinometer or a laser. Families
- 432 and species names follow the African Plant Database (<u>http://africanplantdatabase.ch</u>). The AfriMont
- dataset consists of 72,336 stems, of which 92.9% were identified to species, 98.4% to genus and
- 434 98.5% to family. This dataset represents a standardised safe long-term repository of valuable
- historical data (four sites initially considered could not be included because tree-level data hadalready been lost by data owners).
- 437

## 438 AfriTRON or lowland Africa dataset

The 132 lowland-forest plots are all from AfriTRON<sup>4,13,53</sup>. They were selected using the same criteria as above (but with elevation <800 m asl), restricted to countries for which we also had montane plots plus neighbouring countries where the mountains span international borders (e.g. Mt Nimba spans Guinea and Liberia). The dataset includes 51,305 stems, of which 89.6% were identified to species, 97.3% to genus and 97.7% to family. The plot data were retrieved from forestplot.net on 06/01/2019. The plot locations and details are in Table S6.

445

## 446 Literature dataset

- We compiled data on AGC-stocks in tropical lowland and montane forests to compare to the
   AfriMont data. Data for lowland forests came from ref.<sup>7</sup> and consisted of all multi- and single-census
- 449 plots that were <800 m asl. Data for montane forests were obtained from ref.<sup>2</sup>, with additional data
- 450 from Venezuela (ref.<sup>5</sup>) and Colombia (ref.<sup>6</sup>). Montane plots were defined as  $\geq$ 800 m asl; elevation
- 451 was not provided for the Colombian dataset so plots were selected based on the forest type, and
- 452 these plots were excluded from analyses requiring elevation. To avoid double counting plots,
- 453 Venezuelan and Colombian plots were removed from the ref.<sup>2</sup> dataset.
- 454

## 455 Aboveground carbon

- 456 For each tree in the montane dataset we used the published allometric equation by ref.<sup>54</sup> to
- 457 estimate aboveground biomass. This allometric equation was created using data from directly
- 458 harvested trees at 58 sites across the tropics, including eight sites with elevation ≥800m asl (range
- 459 900-3,000m asl including sites in Africa). We then converted this biomass to carbon, assuming that

460 aboveground carbon (AGC, in Mg C ha<sup>-1</sup>) is 45.6% of aboveground biomass<sup>55</sup>. AGC for each plot was

- 461 estimated as the sum of the AGC of each living stem, divided by planimetric plot area (in hectares). If
   462 field measurements of slope were unavailable, we converted surface to planimetric area extracting
- field measurements of slope were unavailable, we converted surface to planimetric area extracting
   slope from the NASA's Shuttle Radar Topography Mission (SRTM) product. We excluded tree ferns,
- bamboo and palms, as these were not measured in all plots. Ref.<sup>54</sup> includes tree diameter, wood
- 465 mass density and tree height. The best taxonomic match wood density of each stem was extracted
- 466 from a global database<sup>56,57</sup> following ref.<sup>53</sup>. For some sites, all trees in a plot had been sampled for
- height. If this was not the case, but some field measurements of height were available (typically ten
- stems per diameter class), we constructed a site-specific height-diameter model, using a Weibull
   equation following ref.<sup>58</sup>. If no field measurements of height were available, we constructed a
- 470 cluster-specific height-diameter model, using a Weibull equation, as explained in Table S7 in
- 471 Supplementary Information. The same approach was used to calculate aboveground biomass for
- 472 lowland forests. For these, height was estimated using a Weibull equation following ref.<sup>58</sup>.
- 473

## 474 Small plots and data subsampling

- For 22 sites where plots were small (<0.2 ha), we aggregated plots to groups of about 0.2 ha based 475 476 on their geographic proximity, elevation, environmental affinity and the co-authors' knowledge of 477 the site, to help reduce the variation among plots at site level. This is because the presence of an extremely large tree in a small plot can result in overestimates of AGC<sup>59</sup>. We investigated if using the 478 479 aggregated-plot approach affected AGC-stock estimates at the site level, and this was not the case 480 (Extended Data Fig. 2). We also investigated if including small plots affected the continental mean 481 AGC-stock estimates, as small plots have greater edge surface, and there is a tendency of some field 482 teams to include large trees inside plots when laying out the boundaries<sup>60</sup>. Including small plots did 483 not significantly affect our continental mean AGC-stock estimates (Extended Data Fig. 2). We also 484 explored the sensitivity of our continental mean AGC-stock estimates to data subsampling. Data 485 were resampled at different sample sizes either at plot level (sampling with replacement) or at site
- level (sampling without replacement). The number of plots (n=226) and the number of sites (n=44)
   we sampled indicate that our estimates of AGC-stock at the continental level are robust (Extended
- 488 Data Fig. 1). They are also not affected by the fact that we included plots 800-1,000 m asl (Extended 489 Data Fig. 3).
- 490

## 491 Size classes

For all plots, we computed the proportion of AGC which was distributed in each size-diameter class, using the classes of ref.<sup>15</sup>. We also computed stem density, basal area, density of large trees (>70 cm diameter, named SD<sub>70</sub> in stems ha<sup>-1</sup>) and Podocarpaceae abundance (in percentage of plot-level basal area).

496

# 497 Environmental variables and their effects

Climate variables (temperature annual mean and seasonality, and precipitation mean and 498 seasonality, i.e. Bio1, 4, 12 and 15) were extracted from WorldClimV2<sup>61</sup> at 30 arc-sec (~1-km) 499 resolution. Mean temperature values were adjusted for the difference in elevation between the plot 500 501 and the wider 1-km grid cell using the lapse rate of -0.005°C m<sup>-1</sup>. We obtained data on cloud cover 502 from ref.<sup>62</sup> and lightning frequency (0.1 degree, ~11 km) from the Lightning Imaging Sensor (LIS) very high resolution climatology<sup>63</sup>. Values for soil variables (cation exchange capacity, CEC, representing 503 soil fertility, and percentage clay representing soil texture) were extracted from SoilGrids<sup>64</sup> (~1-km 504 505 resolution) and a depth-weighted mean taken for values from 0 to 30 cm depth to give a single value 506 of each soil variable per plot. Elevation was obtained from SRTM (at 3 arc-second resolution, ~90 m). 507 Topographic metrics were calculated from elevation data using the terrain function in the raster R 508 package version 3.3-6. These were slope and topographic position index (TPI). TPI is the difference 509 between the elevation of the plot and the mean value of the eight surrounding grid cells – positive 510 values indicate locally high locations and negative values indicate locally low locations. Where small

- 511 plots were aggregated for analysis, environmental variables were extracted for the ungrouped plot
- 512 locations, and then an area-weighted mean taken to obtain a plot-level value.
- 513

#### 514 Elephant and conifer effects on AGC-stocks

- 515 For the current elephant presence in the AfriMont plots, we created a binary variable
- 516 (presence/absence) based on co-authors knowledge of elephant ranges and elevation distribution at
- each site as of 2019. Co-authors estimated that elephants were present in 2019 in 54 plots in 12
- 518 sites in five countries (see Table S5). For all plots which had at least one individual in the
- 519 Podocarpaceae family (47 plots, 16 sites, 7 countries), we computed the contribution of
- 520 Podocarpaceae to plot basal area and AGC-stock in terms of percentages.
- 521

#### 522 Estimating forest cover and loss

- 523 We obtained estimates of forest cover and loss in the years 2000 through to 2018, using the 'loss
- 524 year' dataset of the Global Forest Change database, version 1.6 (ref.<sup>38</sup>). To exclude plantation
- forests, 'dry' forests (e.g. miombo woodland) and degraded forests, we applied the 'primary humid
- 526 forest' mask developed by ref.<sup>39</sup>. We distinguished montane from lowland forests using an
- 527 elevational cut-off of 800-m elevation, using the SRTM v3 product at 1 arc-sec resolution (snapping
- to the ref.<sup>38</sup> grid of the same resolution). Where there were gaps in the 1 arc-sec SRTM product, we
- 529 filled these using a 1 arc-sec bilinear interpolation of the (gapless) 3 arc-sec SRTM product. Areal
- estimates of forest cover and loss were calculated at 30-m resolution using the Africa Sinusoidal
- projection. To estimate future forest loss by year 2030, we extrapolated absolute country-level
- 532 deforestation rates for the period 2000-2018 (in ha per year).
- 533

#### 534 Investigating AfriMont representativeness

- 535 To quantify AfriMont sampling effort within the montane forest biome in Africa, we used the map of 536 tropical montane forest extent (see above) and calculated the amount of remaining forest in each 1-
- degree grid-cell. By dividing the area sampled in the AfriMont dataset by the proportion of this
  biome in a grid-cell, we calculated the expected sampling intensity if sampling was proportional to
- remaining forest extent. To assess how representative our plot network was of the environmental
- 540 conditions of the wider tropical montane forest biome in Africa, we extracted the environmental
- 541 data (climate and soil variables presented above) at ~1-km resolution from grid-cells that contained
- 542 montane forest. We then visually compared the distribution of each variable in our dataset to its 543 distribution across the biome (Extended Data Fig. 7).
- 544

#### 545 AfriMont vs global AGC maps

- We extracted alternative AGC estimates for the AfriMont plots (unaggregated, n=666) from four
  different sources: Harris et al. (ref.<sup>65</sup>) (30-m resolution, dated 2000), the European Space Agency
  Climate Change Initiative Biomass map<sup>66</sup> (100-m resolution, 2017), Saatchi, et al. (ref.<sup>67</sup>) (1-km
  resolution, 2007/8) and Avitabile et al. (ref.<sup>68</sup>) (1-km resolution, circa 2000-2010). Most of the
- 550 AfriMont plots were sampled between 2000 and 2019 (Table S5). Where the plots were found within
- a single map pixel, we extracted that value. Where plots were larger than the pixel size, we averaged
- 52 the values from the surrounding pixels weighted according to the proportion of the pixel that was in
- 553 the plot.
- 554

#### 555 Statistical analysis

- 556 Data were analysed using linear mixed-effects models, with site as a random effect. Site was
- 557 included as a random intercept in all models, and as a random slope where relationships were
- assessed against elevation. Allowing the slope of the elevation effect to vary amongst sites in this
- 559 way captures the *a priori* expectation for slopes to differ among sites, for example due to mass
- 560 elevation effects. The effect of plot size on variation was accounted for by weighting observations by
- a power transformation of plot size; this was estimated during model fitting using the varPower

- 562 function in the nlme R package (ref.<sup>69</sup>), and then models refitted using the lme4 R package (ref.<sup>70</sup>)
- 563 using these estimated weights. Confidence intervals and *P*-values for mixed effects models
- parameters were estimated by bootstrapping models (1,000 iterations) using the
- bootstrap\_parameters function in the parameters R package (ref.<sup>71</sup>). AGC-stocks, stem density and
- 566 SD<sub>70</sub> were natural-log transformed (a small constant was added to SD<sub>70</sub> before log transforming to
- avoid log-transforming zeros) to meet assumptions of normality and avoid heteroscedacity. Likewise,
   the proportional contribution of each size class was square-root transformed. Differences in AGC-
- 569 stocks between all combinations of lowland and montane forests amongst continents were assessed
- 570 using Tukey post-hoc tests implemented in the multcomp R package (ref.<sup>72</sup>). Relationships between
- AGC-stocks and environmental variables were investigated by fitting all subsets of the full model
- 572 with all environmental covariates and averaging the best supported (difference in Akaike
- information criterion from the best supported model ΔAIC<4) models (using dredge and movel.avg</li>
   functions in the MuMIn R package (ref.<sup>73</sup>). We used these relationships with climate and soil to
- 575 predict AGC-stocks in each 1-km grid cell containing montane forests (holding topographic variables
- 576 at their dataset wide mean), and then took the forest-area weighted mean of these to obtain a
- 577 single mean for the tropical montane forest biome in Africa. Differences in AGC-stocks between
- 578 plots with and without elephants were tested using t-test with AGC-stocks natural-log transformed.
- 579 We investigated if Podocarpaceae abundance (in terms of basal area) and plot AGC-stocks were 580 significantly correlated using Spearman's rank correlation coefficient. To investigate if sampling
- design affected AfriMont AGC-stock estimates we used ANOVA to test whether site-level mean AGC stocks differed according to the sampling strategy used to establish plots at that site. To explore the
- relationship between AfriMont AGC-stock estimates and global maps, and among these global maps,
  we used Spearman's rank correlation test.
- 586 References Methods

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- A.C-S. conceived the study and assembled the AfriMont dataset. A.C-S. and M.J.P.S. analysed the
- plot data (with contributions from S.L.L.) and wrote the manuscript. P.J.P. analysed forest extents
- 713 and contributed to writing. S.L.L. conceived and managed the AfriTRON forest plot recensus

714 programme. E.T.A.M. and V.A. helped compare plot data with remote sensing carbon maps. All co-715 authors read and approved the manuscript. 716 717 Competing interests The authors declare no competing interests. 718 719 **Additional information** 720 Supplementary information is available for this paper at XX (to be added) 721 Correspondence and requests for materials should be addressed to A. C-S. 722 Reprints and permissions information are available at XX (to be added) 723 724 Data availability 725 Source data to generate figures and tables are available from: 726 https://doi.org/10.5521/forestplots.net/2021 5 727 728 **Code availability** 729 R code to generate figures and tables is available from: 730 https://doi.org/10.5521/forestplots.net/2021 5 731 732 Affiliations 733 <sup>1</sup>Department of Environment and Geography, University of York, York, UK. <sup>2</sup>Department of 734 International Environmental and Development Studies (NORAGRIC), Norwegian University of Life 735 Sciences, Ås, Norway. <sup>3</sup>Department of Natural Sciences, Manchester Metropolitan University, 736 Manchester, UK. <sup>4</sup>School of Geography, University of Leeds, Leeds, UK. <sup>5</sup>Climate Change Specialist Group, Species Survival Commission, International Union for Conservation of Nature, Gland, 737 738 Switzerland. <sup>6</sup>University College London, Department of Geography, London, UK. <sup>7</sup>Biology 739 Department, Université Officielle de Bukavu, Bukavu, DRC. <sup>8</sup>Service of Wood Biology, Royal Museum 740 for Central Africa, Tervuren, Belgium. <sup>9</sup>Department of Environment, Laboratory of Wood Technology (Woodlab), Ghent University, Ghent, Belgium. <sup>10</sup>University of Jos, Jos, Nigeria. <sup>11</sup>Nigerian Montane 741 742 Forest Project, Taraba State, Nigeria. <sup>12</sup>Department of Geosciences and Geography, University of 743 Helsinki, Finland. <sup>13</sup>Department of Zoology, Faculty of Science, Charles University, Prague, Czech Republic.<sup>14</sup>Institute of Vertebrate Biology, Czech Academy of Sciences, Brno, Czech Republic. 744 745 <sup>15</sup>Institute of Botany of the Czech Academy of Science, Třeboň, Czech Republic. <sup>16</sup>College of Natural and Computational Science, Addis Ababa University, Addis Ababa, Ethiopia. <sup>17</sup>Department of Natural 746 747 Resource Management, College of Agriculture and Natural Resource, Wolkite University, Wolkite, Ethiopia. <sup>18</sup>European Commission, Joint Research Centre, Ispra, Italy. <sup>19</sup>UK Centre for Ecology & 748 Hydrology, Edinburgh, UK.<sup>20</sup>Université du Cinquantenaire Lwiro, Département de sciences de 749 750 l'environnement, Kabare, Suk-Kivu, DRC.<sup>21</sup>Isotope Bioscience Laboratory (ISOFYS), Ghent University, 751 Ghent, Belgium. <sup>22</sup>Plant Systematic and Ecology Laboratory, Higher Teachers' Training College, 752 University of Yaoundé I, Yaoundé, Cameroon. <sup>23</sup>Institute of Tropical Forest Conservation, Mbarara University of Science and Technology, Uganda. <sup>24</sup>Biodiversity and Landscape Unit, Gembloux Agro-753 Bio Tech, Université de Liege, Liège, Belgium.<sup>25</sup>Institut for Geography, Friedrich-Alexander-754 Universität, Erlangen-Nürnberg, Germany.<sup>26</sup>Institut Supérieur d'Agroforesterie et de Gestion de 755 756 l'Environnement de Kahuzi-Biega (ISAGE-KB); Département de Eaux et Forêts, Kalehe, DRC. <sup>27</sup>UN 757 Environment World Conservation Monitoring Center (UNEP-WCMC), Cambridge, UK. 758 <sup>28</sup>Computational & Applied Vegetation Ecology (CAVElab), Faculty of Bioscience Engineering, Ghent 759 University, Ghent, Belgium.<sup>29</sup>Department of Anthropology, George Washington University, Washington DC, USA. <sup>30</sup>School of Life Sciences, University of KwaZulu-Natal, Scottsville, 760 Pietermaritzburg, South Africa. <sup>31</sup>Shaanxi Key Laboratory for Animal Conservation, Northwest 761 University, Xi'an, China. <sup>32</sup>International Centre of Biodiversity and Primate Conservation, Dali 762 University, Dali Yunnan, China. <sup>33</sup>University of Canterbury, New Zealand. <sup>34</sup>Inventory & Monitoring 763 764 Program, National Park Service, Fredericksburg, USA. <sup>35</sup>University of Ghent, Belgium. <sup>36</sup>World

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- 818 819 **Extended Data**
- 820





823 subsampling. AfriMont plot data were resampled at different sample sizes either at plot level

824 (sampling with replacement) or at site level (sampling without replacement). *N* = 1,000 resamples 825 for each sample size.



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828 Extended Data Fig. 2 | Effect of plot area, aggregation procedure and plot design on estimates of 829 aboveground carbon stocks across the AfriMont plot network. (a) Relationship between 830 aboveground carbon stocks and plot area of plots prior to aggregation. The red line shows the fit of a 831 locally weighted regression model (span = 0.75) relating these variables, with dashed lines showing 832 the standard errors. (b) Variation in aboveground carbon stocks using either all plots prior to 833 aggregation (unaggregated), plots prior to aggregation but excluding those < 0.2 ha (unaggregated, > 834 0.2 ha) or the aggregated plots used in the main analyses (aggregated). (c) Effects of plot design on 835 aboveground carbon stocks (each site represents one dot). Sampling strategies include random or 836 stratified random, plots positioned along transects, plots established within elevation bands, 837 subjective measures such as choosing an area of forest considered representative of the wider area, 838 and other strategies (one plot sampled per site or unclear strategy). Carbon stocks (log-transformed) 839 did not differ significantly between sites with different sampling strategies (ANOVA: F4,39 = 0.432, P 840 = 0.785). For specific site information see Table S5. 841





Extended Data Fig. 3 | Robustness of differences in tropical montane forest aboveground carbon (AGC) stocks among continents based on plot networks to differences in elevation. (a) Elevations of montane forests plots sampled in each continent. Violin plots show the distribution of data, with boxplots showing the median and interquartile range of elevation in each continent. (b) Effect of removing submontane plots (800-1,000 m asl) and high elevation plots (> 2,200 m asl, approximately the upper quartile of elevations for the African montane plot dataset) on AGC-stocks in montane forests sampled by plot networks in each continent. Mean AGC-stocks and 95% confidence intervals

are shown as estimated by models using i) all data, ii) excluding plots 800-1,000 m, and iii) restricting
plots to 1,000-2,200 m. Means for all plots differ from the analysis in Fig. 1 as literature plots without
elevation data (plots in Colombia) were excluded from this analysis. Point symbols are proportional
to square-root plot area. N = 324 plots.

855



858 Extended Data Fig. 4 | Relationship between aboveground carbon (AGC) stocks and elevation for

859 tropical montane forests in each continent based on plot networks. Dashed lines show

relationships from a linear mixed-effects model of log-transformed AGC-stocks as a function of
elevation, continent and their interaction. Site was included as a random effect, and AGC-stock –
elevation relationships allowed to vary among sites. Lines show fitted slopes across sites. Neither the
overall relationship between elevation and AGC-stocks (slope = -0.039 [95% CI = -0.127 - 0.057], *P* =
0.420) nor interactions between elevation and continent (Southeast Asia, change in slope = -0.074 [0.294 - 0.149], *P* = 0.503; Neotropics, change in slope = 0.006 [-0.132 - 0.149], *P* = 0.913) are

statistically significant. *N* = 324 plots.

867



#### Coefficient

869 Extended Data Fig. 5 | Environmental drivers of aboveground carbon stocks across the AfriMont 870 871 plot network. Coefficients are from a linear mixed-effects model with site as a random intercept. 872 Results are following all-subsets regression and model averaging, in which variables that do not 873 appear in well supported models are given coefficients of zero, leading to shrinkage in model coefficients. Statistically significant relationships (P < 0.05) are indicated with asterisks. TPI refers to 874 875 topographic position index (positive values indicate higher than surroundings, negative values 876 indicate lower than surroundings). T\_mean: annual mean temperature, T\_seasonality: temperature 877 seasonality, Precip total: annual precipitation, Precip seasonality: precipitation seasonality. 878



880 881 Extended Data Fig. 6 | Expected sampling effort if effort was distributed in proportion to the area 882 of tropical montane forest biome in Africa. Data are summarised at 1-degree resolution. Upward 883 triangles show grid-cells where AfriMont sampling effort is more than double expected effort, 884 downward triangles show grid-cells where AfriMont sampling effort is less than half expected effort. 885 Circles denote AfriMont sampling effort being between half and double expected effort. The extent of the tropical montane forest biome was defined as closed-canopy forests ≥ 800 m asl in December 886 887 2018, extracted from ref.<sup>38</sup> and clipped to 'primary humid forest' using ref.<sup>39</sup>. This grided map differs from Fig. 4 as numerous grids have very little tropical montane forest. 888 889 890



891 892 Extended Data Fig. 7 | Differences in the environmental conditions sampled by the AfriMont plot 893 network and the tropical montane forest biome in Africa. The extent of the biome was defined as closed-canopy forests  $\ge$  800m asl in December 2018, extracted from ref.<sup>38</sup> and clipped to 'primary' 894 humid forest' using ref.<sup>39</sup>. Environmental variables for the biome were extracted at ~1-km 895 896 resolution.





900 Extended Data Fig. 8 | Differences in aboveground carbon (AGC) stocks in AfriMont plots located in 901 montane forests with and without elephants. a) Differences across all plots. AGC-stocks are 902 statistically significantly lower in forests with elephants (*t*-test, t = 3.5, df=83.5, *P* = 0.001). The thick 903 line shows the median, and boxes cover the interquartile range (IQR). Values > 1.5 times IQR away 904 from the IQR are shown by points. **b**) Differences in countries where elephants are present in at least 905 one of the montane sites studied. Black squares show means in each country in forests with or 906 without elephants – solid lines denote statistically significant differences (*t*-tests, *P* < 0.05). Elephant 907 presence in 2019 was estimated by co-authors (see Table S5). 908 909



#### 910 911 Extended Data Fig. 9 | Relationship between aboveground carbon (AGC) stocks and

912 **Podocarpaceae.** (a) Relationship between AGC-stocks and Podocarpaceae basal area across plots in

913 the AfriMont network, expressed as a percentage of total plot basal area. These variables are not

significantly correlated ( $r_s = 0.083$ , n = 226, P = 0.212). (b) Distribution of plots with at least 20%

915 basal area of Podocarpaceae (black points) in relation to elevation and AGC-stocks. AGC-stocks are

916 not significantly related to elevation or Podocarpaceae basal area (Linear mixed effects model, *P* =

- 917 0.152 and 0.132 respectively).
- 918 919



922 Extended Data Fig. 10 | Within country variation in aboveground carbon stocks based on the

923 AfriMont plot network. Error bars show means and 95% confidence intervals estimated by linear

924 mixed-effects models. Modelled means not shown for countries with fewer than five plots. Point size

925 is proportional to plot area.

926