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RESEARCH ARTICLE

Using multiscale lidar to determine variation in canopy structure from African forest elephant trails

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Abstract

Recently classified as a unique species by the IUCN, African forest elephants (*Loxodonta cyclotis*) are critically endangered due to severe poaching. With limited knowledge about their ecological role due to the dense tropical forests they inhabit in central Africa, it is unclear how the Afrotropics are influenced by elephants. Although their role as seed dispersers is well known, they may also drive large-scale processes that determine forest structure through the creation of elephant trails and browsing the understory, allowing larger, carbon-dense trees to succeed. Multiple scales of lidar were collected by NASA in Lopé National Park, Gabon from 2015 to 2022. Utilizing two airborne lidar datasets in an African forest elephant stronghold, detailed canopy structural information was used in conjunction with elephant trail data to determine how forest structure varies on and off trails. Forest along elephant trails displayed different structural characteristics than forested areas off trails, with lower canopy height, canopy cover, and different vertical distribution of plant density. Less plant area density was found on trails at 1 m in height, while more vegetation was found at 12 m, compared to off trail locations. Trails in forest areas with previous logging history had lower plant area in the top of the canopy. Forest elephants can be considered as "logging light" ecosystem engineers, affecting canopy structure through browsing and movement. Both airborne lidar scales were able to capture elephant impact along trails, with the high-resolution discrete return lidar performing higher than waveform lidar.

Introduction

African forest elephants (*Loxodonta cyclotis*) are critically endangered (Gobush et al., 2021), facing severe threats from poaching and deforestation with estimated population losses of 62% in central Africa from 2002 to 2011 (Maisels et al., 2013). Despite their recent, broad-scale decline and listing as a separate species in 2021 (Hart et al., 2021), forest elephants are understudied due to poor visibility in African tropical forests. Specifically, the forest elephant's ecological role is severely understudied in comparison to the African savanna elephants (*Loxodonta africana*), whose role as a keystone species has been studied for decades (Asner & Levick, 2012; Bond, 1994; Coverdale et al., 2016). Previous work identified forest elephants as important seed and lateral nutrient dispersers

through plant digestion, defecation, urination and the ingestion of mineral rich water and soil (Blake et al., 2009; Campos-Arceiz & Blake, 2011; Doughty et al., 2016; Metsio Sienne et al., 2014). However, their influence on forest structure is less clear.

It is hypothesized that forest elephants impact forest structure through their feeding preferences. As generalist browsers, forest elephants modify their environment by trampling, ingesting and shaking vegetation to access preferred fruits (Maisels et al., 2002). It is proposed that through browsing in the understory, small saplings are consumed which allows large, woody trees to succeed (Terborgh et al., 2016). For example, Berzaghi et al. (2023) collated >100 000 records of elephant feeding preferences in central Africa and showed that low wood density vegetation was consumed at significantly higher rates than high wood density tree species. At the same time, fruits from larger trees were preferred by forest elephants. By feeding on low wood density plants but fruit from larger trees, elephants may promote forests with higher aboveground biomass. It is estimated that if tree species preferentially dispersed by elephants were replaced by other species, above ground carbon could decrease by up to 12% in central Africa (Berzaghi et al., 2023).

One of the most obvious ways forest elephants impact their environment is by the creation of trails through the forest (Blake & Inkamba-Nkulu, 2004; Vanleeuwé & Gautier-Hion, 1998). These paths are usually 0.5–0.9 m wide on average (Blake & Inkamba-Nkulu, 2004; Vanleeuwé & Gautier-Hion, 1998), allowing elephants to access important resources such as streams, mineral-rich areas (bais), and high-priority fruit trees (Benitez & Queenborough, 2021; Blake & Inkamba-Nkulu, 2004). Through consistent use, structural changes in the understory have been found along elephant trails in the Republic of the Congo (Blake & Inkamba-Nkulu, 2004; Vanleeuwé & Gautier-Hion, 1998), the DRC (Inogwabini et al., 2013), and Uganda (Benitez & Queenborough, 2021). Vanleeuwé and Gautier-Hion (1998) found that forest elephants created larger "boulevard trails" for long-distance movement which cut through all forest types and "foraging trails" which wound through the thick Marantaceae and Zingiberaceae (terrestrial herbaceous vegetation). Elephant trail networks can be dense (e.g., 9.82 trails/km in Ndoki National Park; Blake & Inkamba-Nkulu, 2004) and probably last for hundreds of years as elephants pass on their mental maps of the forest to their offspring (Haynes, 2006).

Additional fauna—including humans, maintain elephant trail systems and utilize them to access rivers, bais, and hard-to-access regions of the forest (Blake, 2002; Remis & Jost Robinson, 2020). Indigenous tribes, such as the BaAka, have used elephant trails for hunting and travel for centuries (Remis & Jost Robinson, 2020). Elephant trails in Gabon were found to act as natural firebreaks along forest-savanna boundaries, assisting in the protection of forest interiors (Cardoso et al., 2020). To gain access to forest interiors, logging companies have used elephant trails and further widened them with bulldozers in some forests. However, the effects that elephants and humans have had on the African tropics through the creation and maintenance of these trails is undetermined. Understanding the complexities of vegetation structure surrounding these trails could shed light on forest elephant feeding behaviors, seed dispersal patterns, space use, and role as ecosystem engineers—all of which are still relatively unknown.

Lidar (light detection and ranging) sensors are used to measure forest structural properties, such as cover and canopy height (Dubayah & Drake, 2000). Energy pulses returned from the ground and vertical sub-canopy vegetation structure are digitized and converted into a 3D representation of topography and plant biomass distribution. Lidar has proven effective in modeling habitat preferences and understanding predator–prey dynamics, movement behavior, and species richness in relation to both vertical and horizontal structure (Davies & Asner, 2014; Goetz et al., 2010; McLean et al., 2016). To our knowledge, lidar has never been used to determine forest elephants' impact on the structure of forests in the Afrotropics.

Here, we use lidar to investigate vegetation structure at broad spatial scales along forest elephant trails in Lopé National Park, Gabon. This study aims to characterize the role forest elephants play as ecosystem engineers through the creation of trails, specifically by using lidar to quantify their impact on canopy structure. Multiple lidar collections have taken place over the last 8 years in Lopé National Park, Gabon, a protected area with one of the highest elephant densities in central Africa (0.93 elephants/km^2; Bezangoye & Maisels, 2010). Utilizing elephant trail geolocation data and two scales of lidar (2× aircraft) from NASA, questions regarding how elephants affect canopy structure along trails can be addressed. Specifically, our research questions are:

1. How do lidar-assessed canopy structure metrics (e.g., canopy height, canopy cover, vegetation area index, and height of median energy) vary with distance from elephant trails?
2. Can changes in forest structure associated with elephant trails be detected with lidar? If so, which resolution of lidar sensors can detect structural variation?

Materials and Methods

Study area

Lopé National Park covers 4960 km^2 near the center of Gabon (0° 10′S 11° 35′ E; Fig. 1) and was established as a

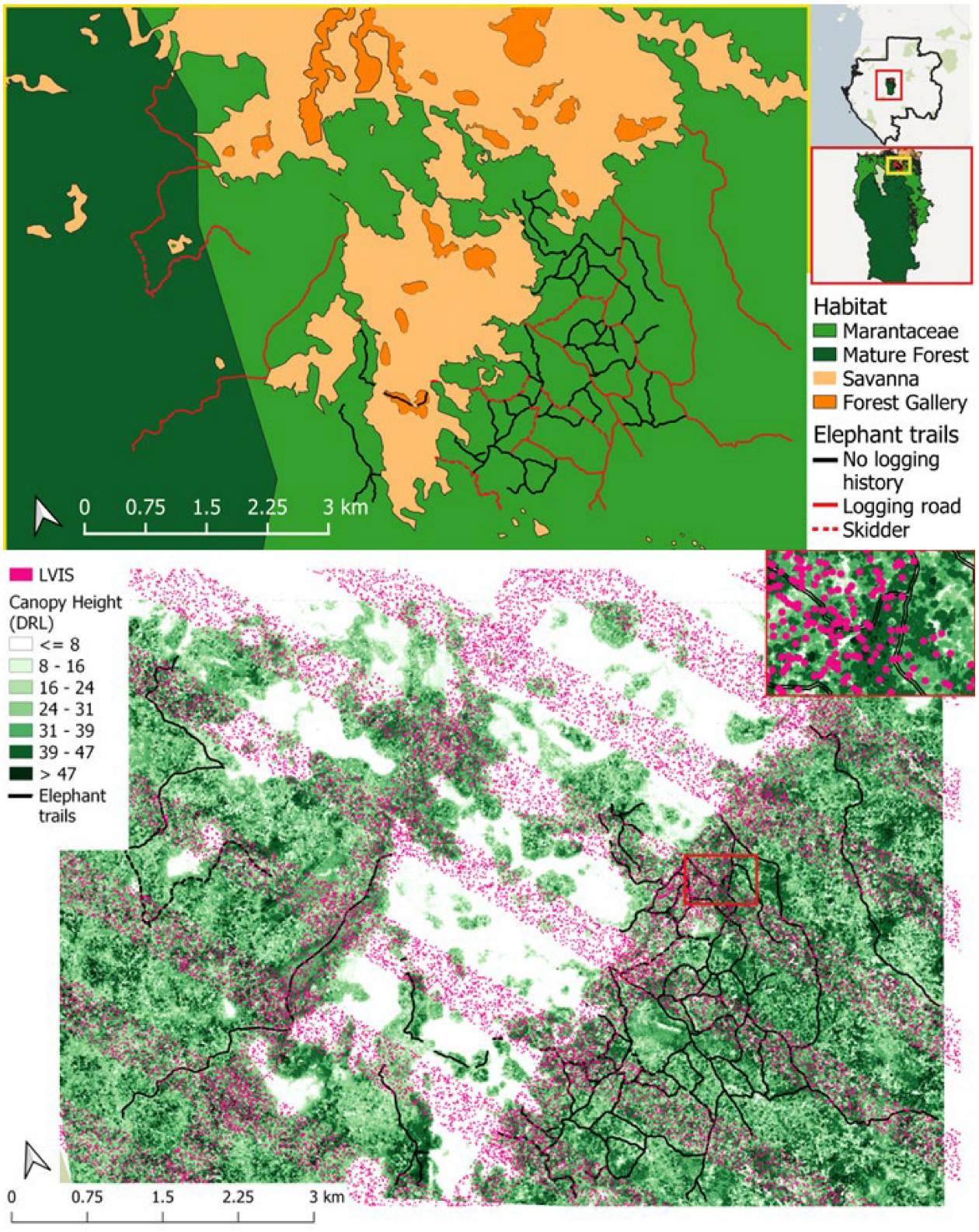

Figure 1. Study area within Lopé National Park, Gabon. Habitat types and elephant trail history are indicated in the top panel, while the two lidar datasets are shown in the bottom panel with overlapping elephant trails (black).

nature reserve in 1946 and as a UNESCO World Heritage Site in 2007 for its natural and archeological richness. The park is dominated by closed canopy tropical forest with a forest-savanna mosaic located in the northern section. Forest types include mature, Marantaceae, gallery, and young forest (due to savanna colonization). The most dominant vegetation type in Lopé is mature forest (63.7% according to Gabon's national park agency), which comprises medium to large trees with very dense canopy cover and a limited understory (White & Abernethy, 1997). Marantaceae forest is the second most prevalent forest type characterized by dense understory dominated by plants in the Marantaceae (arrowroot) family and other herbaceous vegetation preferred by elephants and gorillas, with large trees dominating the canopy (White & Abernethy, 1997). Gallery forests are located along water courses; small woodland patches ("bosquets") are located in the savannas. Both typically have shorter canopies and sparser understories than mature or Marantaceae forests. Lopé Faunal Reserve was selectively logged from the 1960s to the early 2000s for Okoumé (*Aucoumea klaineana*) trees and over 50 other hardwood species with removal rates <2 trees/ha, prior to becoming a national park (White, 1994). Since the gazettement of the area as a National Park in 2002, there has been limited human influence in the park due to a low density of human settlement in this region of Gabon. Eight settlements occur along the northern and north-eastern park boundaries (<4000 people in total) and some indigenous communities use the interior of the southern region of the park (Rakotonarivo et al., 2021).

Elephant trail data

Fifteen kilometers of elephant trails were surveyed and geolocated in Lopé National Park in January 2022. Trail data were collected with a handheld GPS (Garmin 66i), which has a standard 3-m accuracy. Tracklogs were then downloaded onto Garmin BaseCamp and converted to shapefiles in QGIS for analysis. Width of trails at breast height (~1.4 m) and on the ground were measured and recorded every 200 m along trails. Trail edges on the ground were determined by the presence of bare ground, while trail edges at breast height were recorded where the vegetation subsided on either side. These data were supplemented by an existing dataset with 62 km of georeferenced elephant trails from Gabon's National Park Agency (ANPN), collected between 1986 and 2010 and regularly refined as path locations changed and GPS technology advanced (White, 1995; SEGC unpublished).

Elephant trail history was verified by K. Abernethy (Associate Researcher for the National Centre for Research in Science and Technology in Gabon) and L. Makaga (Lopé Research Station Manager) as either: (a) elephant-engineered, (b) previous logging road, and (c) previous skidder trail. All trails are now elephant trails, while some were previously created or used by the logging industry to access the forest more easily. Although all logging in the study area of the park ceased 40–50 years ago, it is vital to take the effect of logging into account due to its known legacy in affecting forest structure (Hall et al., 2003). Skidder trails were used to extract logs, with some small trees removed but not clear-cut or bulldozed. Previous logging road trails were clear-cut and bulldozed to achieve an understory opening of >5 m, allowing heavy equipment to enter the forest. Elephant-engineered trails were never anthropogenically widened or logged but have been maintained by elephants. To simplify these classifications, trails that were once logging roads were deemed "elephant-and-human engineered," while trails maintained by elephants only are "elephant-engineered." Skidder trails were removed from the analysis due to a small sample size.

Lidar data

We used lidar data acquired from two different sensors, with spatial resolution ranging from ~1 to 20 m (Table 1). The first dataset was acquired in July 2015 as a discrete return point cloud using a helicopter low-flying rotary platform, providing 54 km^2 of wall-to-wall coverage (Discrete Return Lidar or DRL; Silva et al., 2018). The second dataset was acquired in March 2016 as waveform lidar using NASA's Land, Vegetation, and Ice Sensor (LVIS) mounted on a King Air B-200 airplane flown at 7.3 km in altitude (LVIS; Blair

Table 1. Lidar data specifications from both airborne sensors. Full coverage of DRL is found across the study area. LVIS shot locations by habitat type are found in Figure 1.

Name	Type	Platform + Instrument	Footprint Width	Vertical/Horizontal Geolocation Accuracy	Pulse Density	Acquisition Date(s)
DRL	Airborne discrete return point cloud	EC 135 Helicopter + Riegl VQ480U (1550 nm)	<0.1 m	<1 m/<1 m	78 000 pulses/ha	July 2015
LVIS	Airborne full waveform	NASA Langley King Air B-200 Airplane + LVIS laser (1064 nm)	20 m	~1 m/0.1 m	4.6 points/ha	March 2016

et al., 1999). LVIS data return individual "shots" instead of providing wall-to-wall coverage, with shots 20 m in diameter. Figure 1 shows the location of the two lidar datasets in relation to known elephant trails surveyed in Lopé National Park. Table 1 describes the spatial resolution, instrument specifications, and data collection timeline for each lidar acquisition. With the highest spatial resolution of all lidar in the study at 1 m, the DRL dataset provides wall-to-wall coverage of the study area (Fig. 1). The DRL and LVIS data were validated with field plots and compared in 2018 for structural metrics (Silva et al., 2018). LVIS canopy cover and vertical profile products (plant area volume density or PAVD) were downloaded from Oak Ridge National Laboratory Distributed Active Archive Center (ORNL DAAC; Tang et al., 2018).

Data analysis

Elephant trail data

Elephant-engineered and elephant-and-human-engineered trails in Marantaceae forests were used in the lidar analysis. These were selected as Marantaceae is the dominant habitat type in our study area, and skidder trails had a small sample size (mature forest and forest gallery results are found in the supplementary material). All elephant trails were clipped to exclude savannas. A 2 m diameter buffer (1 m from trail center) was used for "on-trail" analyses to account for handheld GPS positional uncertainty. While GPS uncertainty fluctuates, elephant trails are on average 0.5–0.9 m in width. Therefore, in order to detect any understory changes in vegetation, a 2 m buffer was chosen. While 2 m does not account for high estimates of GPS uncertainties, it captures the trail center where the highest vegetation changes should occur. Each trail was geolocated multiple times by Gabon's ANPN, allowing for a higher accuracy from repetitive sampling. Additional 10 m buffers were created from 10 to 60 m from trails and used to compare on and off trail vegetation. A DTM was used to compare the environmental conditions of on and off trail forest. On-trail buffers had an average DTM value of 291.7 m while the off-trail buffers averaged 288.4 m. Waterways were removed from all trail shapefiles using a 50 m diameter (25 m from center) stream buffer (Fig. 2). There were 39 elephant-engineered "on trail" buffers for each marked trail, totaling 0.037 km^2 in area coverage. The subsequent 39 elephant-engineered "off trail" buffers totaled 0.263 km^2 in area coverage (10 m in width for each side of the trail). Elephant-and-human-engineered "on trail" buffers had a count of 12, with 0.037 km^2 in area coverage with 12 corresponding "off trail" buffers totaling 0.328 km^2.

Lidar processing

The discrete return lidar were clipped to both on and off trail buffers (segment tool) and filtered using the

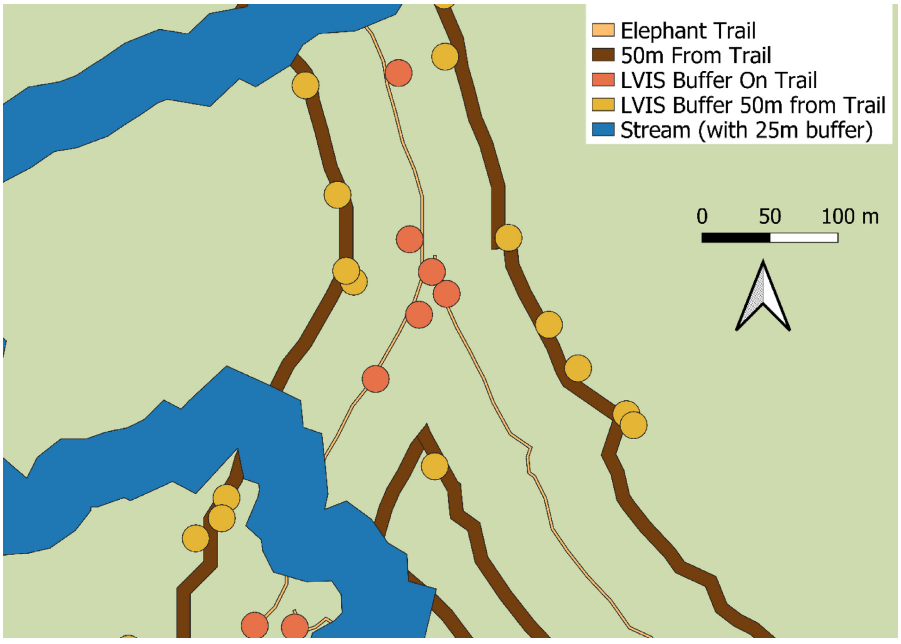

Figure 2. Visualization of buffered trail and lidar data used in the structural analyses.

statistical outlier filter with 2 standard deviations in CloudCompare (v.2.13alpha). Any additional erroneous points located above top of canopy height or below the ground were manually removed with the segment tool. Within R (version 4.2.1; R Core Team, 2022), the lidR, sp., and lidRmetrics packages were used to process and manipulate the point cloud (Roussel et al., 2020; Roussel & Auty, 2023). A DTM was created from the cleaned point cloud using the kriging method (k = 10 L), which was then used to normalize the height of all lidar returns. An additional manual filter of the normalized point cloud was performed to ensure a clean point cloud without erroneous points ($0 < Z < 65$). LVIS point data were buffered by their respective footprint widths (20 m in diameter for LVIS; Fig. 2). These buffers were then used to clip the wall-to-wall DRL pointcloud for the multiscale lidar comparisons (DRL vs. LVIS) to ensure direct overlap between all lidar scales for forest structure comparisons. LVIS footprints that had at least 50% overlap with elephant trail buffers were selected for subsequent canopy structural analyses. There were 528 LVIS shots on trails in marantaceae habitat, 70 in mature forest, and 25 in forest galleries. Off trail 50–60 m buffers totaled 908 LVIS in marantaceae, 148 in mature, and 52 in forest galleries.

"On Trail" and "Off Trail" forest structural comparisons

We used a variety of canopy metrics to compare vegetation structure on and off elephant trails (Table 2). First, leaf area density (LAD) and plant area volume density (PAVD; Dubayah et al., 2021) were used to understand how the vertical structure profile differed on and off trails. LAD and PAVD quantify the distribution of both woody and foliar material across discrete height bins (Table 2). In this analysis, LAD is including not only leaves but other plant and woody material and therefore representing the plant area density. Although LAD and PAVD are calculated using different equations, they are both indicators of the vertical distribution of plant area in the forest. We are confident in using them to compare the plant area distribution across lidar types in Lope National Park, as these two lidars were compared in other published work (Silva et al., 2018). LVIS PAVD is discretized into 1 m height bins. LAD was calculated in the subsequent DRL using the lidR package (Roussel et al., 2020) from the cleaned and normalized point cloud into 1 m height bins using a standard extinction coefficient of foliage of 0.5 and a clumping index of 1 (also used for LVIS). Within each category of trail (elephant-engineered or elephant-and-human engineered), the LAD at each 1 m height bin was calculated for each on or off trail buffer shapefile. To compare LAD between the on and off trail forest, Wilcoxon tests were run for each height bin to account for the non-normal data. For example, 39 LAD values were calculated at 1 m in height for the 39 elephant-engineered "on trail" shapefiles, which we were then compared to the 39 LAD values at 1 m for the subsequent "off trail" shapefiles in a Wilcoxon test. The P-values from each Wilcoxon test from 1 to 50 m in height were then stored as a data table, and significant P-values were displayed on the LAD plots at their corresponding height. Finally, standard error bars were generated for each LAD and PAVD values and displayed on the plots at their subsequent height bins.

Table 2. Canopy structural metrics descriptions and references.

Metric	Description	References
Leaf area density (LAD/PAVD)	Leaf area per unit of volume	Bouvier et al. (2015)
Max canopy height (CH)	Maximum of canopy height	Drake (2001)
Height of median energy (HOME)	Median of returned energy	Drake (2001)
Vertical distribution ratio (VDR)	VDR = [CH − HOME]/CH An index ranging from 0 to 1 depicting the vertical distribution of plant matter	Goetz et al. (2007)
Total vegetation area index (VAI)	Sum of LAD for all height bins	Bouvier et al. (2015)
Canopy cover fraction (CCF)	1- Deep Gap Fraction	LaRue et al. (2020)

The maximum canopy height is equated with the LVIS RH98 metric – the height (relative to the ground) at which 98% returned energy is reached (Table 2). RH98 represents the top of the canopy and is not as sensitive to atmospheric noise such as fog or very low clouds in comparison to RH100, making it a more stable indicator. Canopy height is an indicator of forest successional stage and above ground biomass (Drake, 2001). RH50 is also referred to as the height of median energy (HOME), which indicates where the bulk of vegetation is located vertically. The vertical distribution ratio (VDR; Goetz et al., 2007) equals (RH98−RH50)/RH98. VDR is normalized by RH98 to indicate the relative distribution of biomass within the vertical profile. High VDR values (closer to 1) are associated with bottom-heavy vertical profiles (mid successional) while lower values (closer to 0) are associated with top heavy vertical profiles (young and old forests). Finally, canopy cover fraction (CCF) was calculated to determine the amount of forest cover within each buffer. Canopy cover fraction is ecologically important as it influences light availability, water interception, and temperature regulation. Total vegetation area index (VAI) represents the total plant or vegetation area within the vertical column.

Violin plots for all canopy metrics were created and Wilcoxon tests were used to compare each metric for on and off trail forests. Finally, LAD vertical bins were tested for normality using Kruskal Wallace tests. As the data were not normally distributed, Wilcoxon tests were used to determine differences in LAD/PAVD for each vertical height bin (either 1 m for DRL or 5 m for LVIS).

Results

Canopy structural metrics

Elephant-engineered trails and elephant-and-human trails had a mean ground width of 88.8 ± 32.4 SD cm and 123.7 ± 44.1 SD cm, respectively (Table S1). Average width at breast height increased for both trail types, with elephant-engineered at 218.9 ± 60.3 SD cm and elephant-and-human at 287.1 ± 86.8 cm (Table S1). When averaging LAD from 1 to 4 m in height with the DRL, less LAD is found on trails than off (Fig. 3). However, this increase is relatively small with LAD values generally around 0.6 m^2m^{-3}. Furthermore, both elephant-engineered and elephant-and-human trails had significantly less LAD on trails than off at the 1 m vertical height bin (Fig. 3).

All "on-trail" and "off-trail" buffers were analyzed for forest structure using the DRL, as it provided full coverage of the study area. From these analyses, max canopy height was significantly higher 50 m from trails than on both elephant-engineered ($P < 0.001$) and elephant-and-human trails ($P < 0.01$) (Fig. 4). No difference was found between HOME or VDR on elephant-engineered trails; however, elephant-and-human trails had higher HOME values off trail ($P < 0.01$) and lower VDR values off trail than on ($P < 0.05$). Total VAI was higher off trails than on elephant-and-human trails ($P < 0.01$), with no difference observed for elephant-engineered trails.

Elephant-engineered trails showed significantly higher canopy cover 50 m from trails than on them ($P < 0.001$), and no significant difference was found in cover between on and off elephant-and-human trails. When comparing the vertical profiles (average LAD) of for on and off trail forest, elephant-engineered trails had higher LAD at the 1, 12, 47, 48, and 49 m height bins (Fig. 5). Elephant-and-human trails had less LAD at 1 m and 30–49 m heights and more LAD at 16, 17, and 18 m heights than off trail forest.

Comparing multiple lidar scales

To determine how different scales of lidar detect changes in forest structure on elephant trails, overlapping lidar data were compared on and off trails. The DRL and LVIS

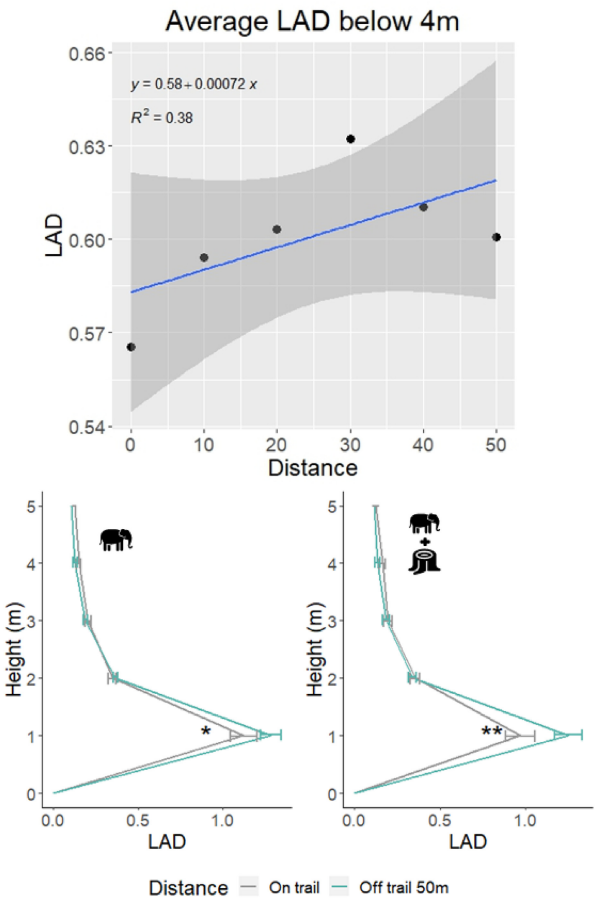

Figure 3. Top: Discrete return lidar (DRL) average leaf area density from 1 to 4 m on elephant-engineered trails in Marantaceae forests fitted with a linear trend line with standard error. The trend line accounts for 38% of the variation. Bottom left: Average leaf area density (LAD) from 0 to 5 m in vertical height on elephant-engineered trails. Bottom right: Average leaf area density (LAD) from 0 to 5 m in vertical height on elephant-and-human-engineered trails. Statistical comparisons of average LAD at each 1 m vertical height bin using Wilcoxon tests between on and off trail are indicated by one asterisk ($P < 0.001$), two asterisks ($P < 0.01$), or three asterisks ($P < 0.001$).

were used to compare canopy structure on and off elephant-engineered trails (Fig. 5). Significant differences were observed in plant area density from DRL height bins at 22–36 and 48 m (Fig. 5). Almost all height bins were significantly different between on and off elephant-engineered trails from LVIS, except from 10–20 to 35–40 m (Fig. 5). A similar pattern is seen in the DRL versus LVIS comparison for the elephant-and-human-engineered trails (Fig. S1).

Due to smaller lidar overlap in mature forests and forest galleries, limited canopy structural analyses were completed for those habitat types. Using the two lidar datasets (DRL and LVIS), elephant-and-human trails in

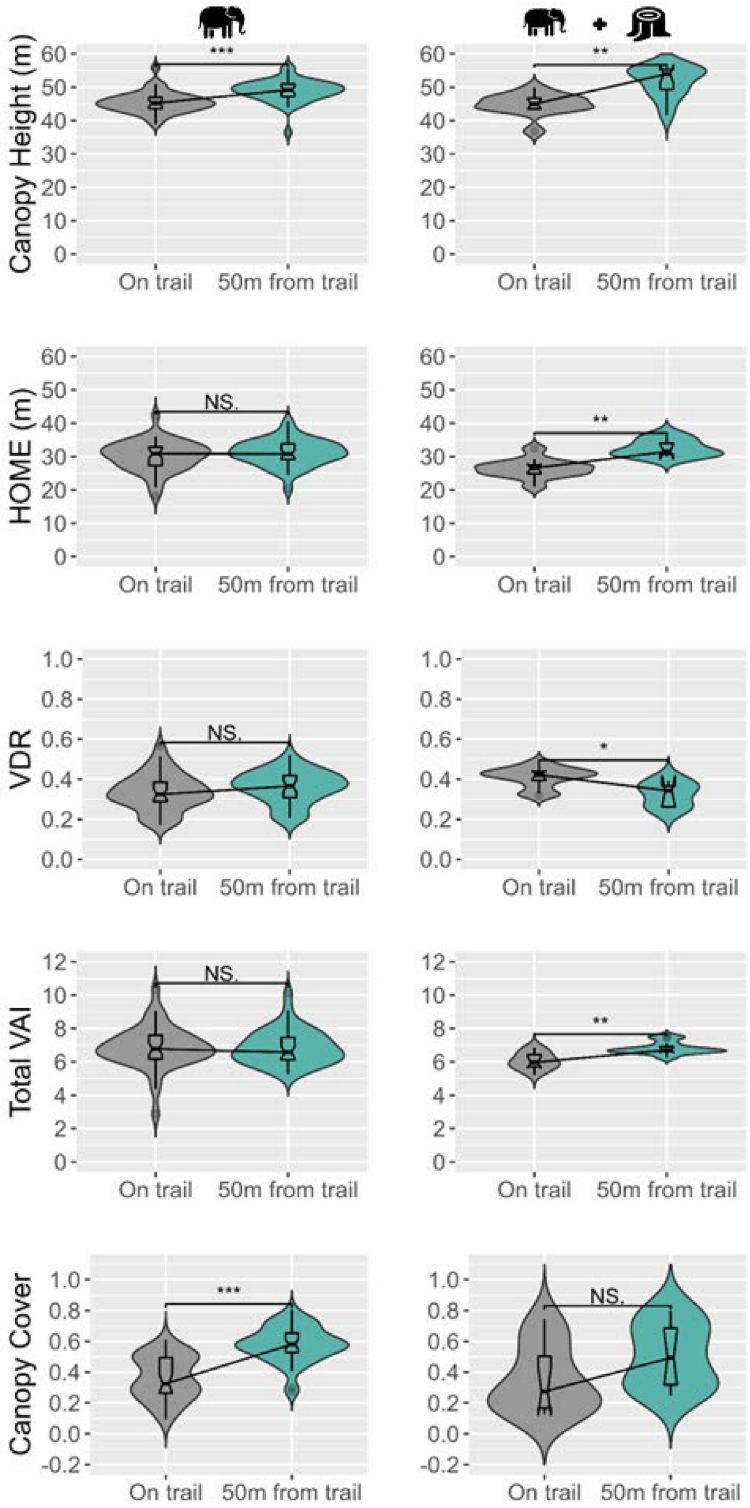

Figure 4. Maximum canopy height, height of median energy (HOME), vertical distribution ratio (VDR), total vegetation area index (VAI), and canopy cover violin plots from DRL on elephant trails and 50 m from elephant trails. Comparisons using elephant-engineered trails are indicated in the left panels, while elephant-and-human-engineered trails are indicated in the right panels. Notched box plots are within each violin plot with trend lines connecting median values between on and off trail. Statistical comparisons using the Wilcoxon test between on and off trail are indicated as not significant (NS), one asterisk ($P < 0.001$), two asterisks ($P < 0.01$), or three asterisks ($P < 0.001$).

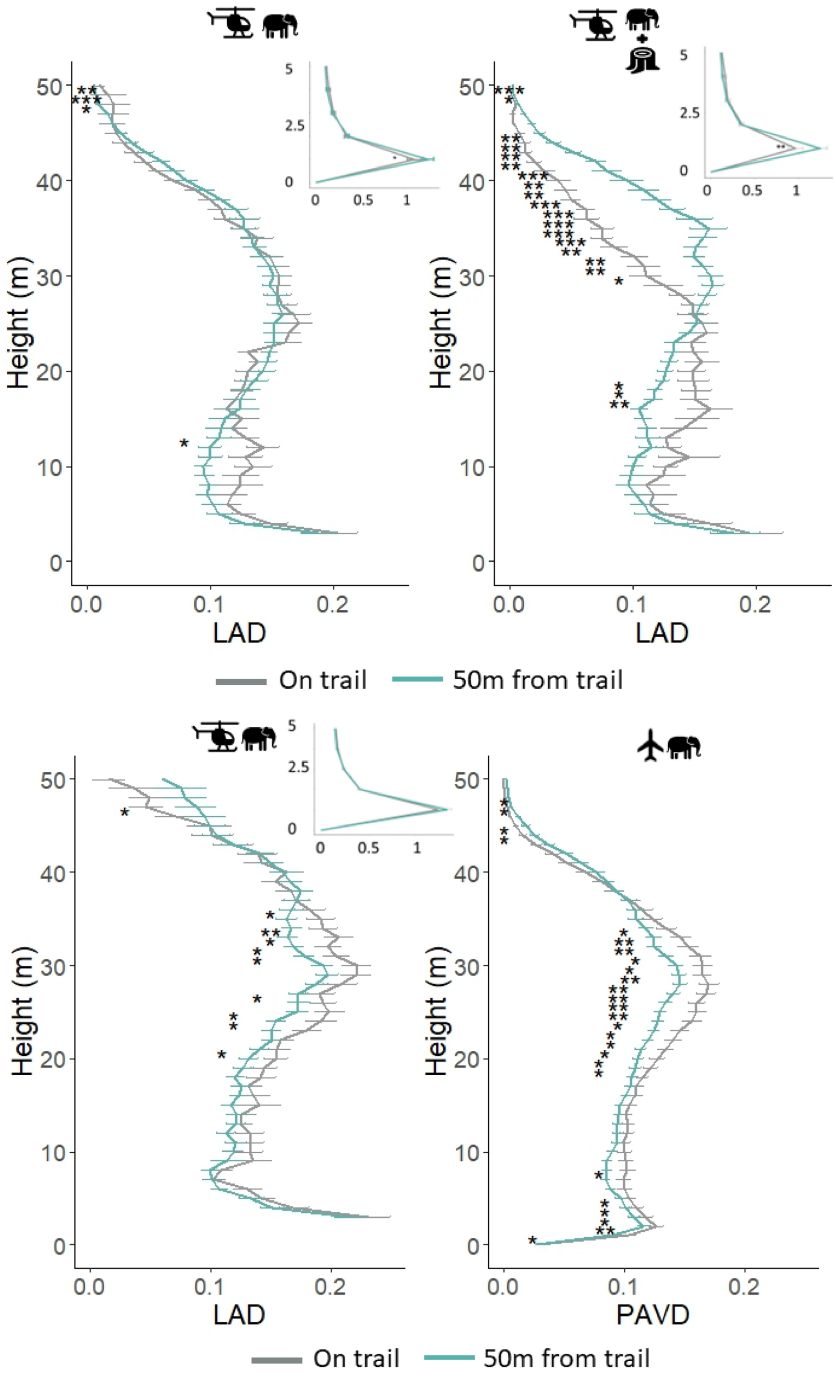

Figure 5. (Top) Average leaf area density (LAD) profiles from discrete return lidar (DRL) for elephant-engineered trails and elephant-and-human-engineered trails with standard error bars. (Bottom) Average LAD and PAVD profiles with standard error bars from overlapping DRL and LVIS shots on elephant-engineered trails (N = 219 on trail, N = 355 off trail). Statistically significant differences from Wilcoxon tests for each height bin are indicated by one ($P < 0.05$), two ($P < 0.01$), or three ($P < 0.001$) asterisks. Inset figures display DRL LAD values from 0 to 5 m.

mature forests depicted similar canopy structure to those in Marantaceae forests with higher LAD on trails in from 1 to 15 m and lower leaf area on trails from 30 to 45 m (Fig. S2). Elephant-engineered trails in forest galleries also showed similar trends to those in Marantaceae forests (Fig. S3).

Discussion

African forest elephant trails impact canopy structure

Elephant trails in the Marantaceae forests of Lopé National Park, Gabon appear to impact forest structure by reducing canopy cover and canopy height, while changing the vertical LAD. Less leaf area was found 1 m above the ground for both trail types, while more leaf area was found at 12 m above the ground for elephant-engineered trails, and at 16, 17, and 18 m in height for elephant-and-human trails. Leaf area is significantly less on elephant-and-human-engineered trails from 30 to 49 m in height. With similar HOME and VDR metrics, the vertical distribution of plant area is similar both on and off of elephant-engineered trails; however, canopy height and cover are smaller on trails (Fig. 4). Through the creation and maintenance of trails, it appears that African forest elephants impact canopy structure by removing canopy cover and lowering canopy height of vegetation while keeping total VAI the same. Total VAI was similar both on and off elephant-engineered trails indicating that the higher plant area in the lower vertical bins made up for a shorter canopy.

When comparing forest structural changes from elephant-engineered trails to trails altered by the logging industry (through clear-cutting and bulldozing), considerably more impact on the canopy by the latter is observed. The legacy of selective logging is noticeable in our study, showing that after ~50 years since logging subsided, the overall vertical structure remains affected. The widening and compacting of trails during tree harvesting produced long lasting effects by lowering canopy height, total VAI, HOME, and canopy cover (Fig. 4). These structural changes are in line with previous research, showing that logged forests in central Africa have altered ecosystem composition from the loss of larger trees and increased diversity in the understory from higher light penetration (Hall et al., 2003; Sullivan et al., 2022). Yet, the intertwining influences of both elephant and logging disturbance to vegetation on trails are novel. While the effects of logging in the Afrotropics are well documented (Cazzolla Gatti et al., 2015), our study suggests that African forest elephants create a lighter disturbance in the canopy—acting as a "logging light" ecosystem engineer. To understand these effects in more depth, we identified two hypotheses on how elephant trail vegetation structure is established.

To understand our findings of higher plant area (PAVD; Fig. 4) from 5 to 30 m above the ground, we hypothesize that elephants create trails in the thickest Marantaceae to gain access to better browse, while simultaneously promoting understory growth through increased canopy gaps. While some trails could be direct paths moving toward a water source or fruiting tree, most others could be created to access the Marantaceae roots—a preferred browse by forest elephants (White et al., 1993). In a similar manner to logging, additional gaps in the forest caused by forest elephant browsing and soil compaction would allow more light to penetrate the canopy and reach the ground (Kamga et al., 2022; Struhsaker et al., 1996), which increases growth of understory vegetation. Added light to the lower and mid canopy promotes the expansion of light seeking fast-growing saplings. Consequently, a feedback effect consisting of elephants consuming vegetation along trails while also promoting more growth is possible. Mount Cameroon National Park has a natural elephant exclosure area (from lava flows), allowing researchers to compare forest structure in elephant and non-elephant areas of the park. In this unique environment, forest elephants promoted canopy gaps and fostered a more heterogeneous forest, which in turn increased the diversity of insects and birds (Kamga et al., 2022; Maicher et al., 2020). Our findings support this hypothesis by showing that both elephant-and-human and elephant-engineered trails have lower canopy cover (Fig. 4) and higher plant area in the understory (Fig. 5) than areas without trails.

Our second hypothesis is that increased soil fertility along trails from animal use encourages more fast-growing vegetation to thrive along elephant trails. The addition of seed deposition and disturbance from forest elephants might promote more fertile growing conditions near elephant paths. Megaherbivores are known to disproportionately affect nutrient availability and forest fertility through the deposition of feces and urine (Doughty et al., 2016; Enquist et al., 2020; Stanbrook, 2018; Wolf et al., 2013). Little is known about forest elephants' use of space in regard to departing from trails to enter the forest interior. However, Inogwabini et al. (2013) found less elephant dung with increasing distance from trails, suggesting they typically stay along the established paths, although the sample size was small. Increased nutrient deposition along elephant trails could therefore promote increased fast-growing vegetation from higher soil fertility. Finally, the effects of higher soil nutrients could contribute to lower canopy height on trails, which is in line with previous findings associating lower aboveground biomass with higher N and P concentrations in the tropics (Unger et al., 2012).

These findings suggest that African forest elephants alter the structure of central African rainforests along their trails. However, further studies are needed on an ecosystem scale to understand their impact on structure more broadly. It is likely that tropical forests with elephants will have less canopy cover and a more heterogenous vertical structure. These ecosystem changes may

provide environmental conditions that support higher insect biodiversity (Kamga et al., 2022; Maicher et al., 2020). Our study found shorter canopy heights along elephant trails, but to understand how overall forest height is affected by elephant density, a larger study must be completed with varying densities of elephants. Forest elephants are the sole disperser of many tall, carbon-dense tree species in the African rainforests; therefore, it is possible that they increase carbon content and canopy height of forests at regional scales as suggested by Berzaghi and others (2022). As our study took place within a park with a high number of elephants, there is no way to determine density related changes. However, with the release of spaceborne lidar from the Global Ecosystem Dynamics Investigation (GEDI; Dubayah et al., 2020), these questions can be addressed on a larger scale.

Determining what scale of lidar is needed to detect elephant trails

It is apparent that lidar can distinguish changes in forest structure from elephant paths, but how are these differences characterized by lidar collected at different scales? As expected, our multiscale lidar analysis found the highest resolution airborne lidar (DRL) detected fine-scale changes in structure. Even though DLR is more sensitive to elephant related structure differences than LVIS, LVIS still detects variation and will be available over larger areas. Furthermore, it may help us link with spaceborne lidar from GEDI.

Limitations and uncertainties

Additional limitations to this study include anthropogenic influences, collection time differences, and geolocation uncertainties for the trail and lidar data. With mean widths of 88 cm on elephant-engineered trails, a 3-m GPS uncertainty could produce changes in the lidar results, making it difficult to find the understory vegetation gap along the path. Each lidar dataset has accompanying GPS errors which could impact our results as well. Silva et al. (2018) compared the DRL and LVIS collected in Lopé and found some large differences in Z values on a footprint scale, mainly due to complex topography. When compared across larger scales, variation decreased and no significant differences were found. UAV lidar could be highly useful when studying small trails in thick Marantaceae forests and should be explored in future research.

Conclusion

Using lidar, we were able to determine canopy height, canopy cover, and vertical structural differences between forest along elephant trails and the surrounding areas in Lopé National Park, Gabon. We also showed that a structural signature exists on logging roads now used as elephant trails, even 50 years beyond the abandonment of logging activity. These findings are novel, as lidar has never been used to quantify how large animals such as African forest elephants influence canopy and vegetation structure through the creation and maintenance of trails (to our knowledge). In addition, the persistence of the structural impacts from selective logging has not been followed on such a long time scale. Although the highest resolution lidar detected more variation in forest structure than LVIS, increased coverage of lidar in the African tropics is imperative to broaden our understanding of elephants' ecological role. Future studies on forest elephant trails in these habitat types would benefit from measuring light availability, characterizing Marantaceae thickness, and classifying trail use with direct measurement using camera traps. Additional research is needed to understand elephant preferences and motivations (e.g. accessing fruiting trees, water courses, bais) to create their trail networks in a variety of habitat types. The full effect elephants have as engineers and trailblazers of Gabonese forests is still under investigation, but it is clear they play a role in influencing canopy height, canopy gaps, and the vertical distribution of plant mass.

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References

Asner, G.P. & Levick, S.R. (2012) Landscape-scale effects of herbivores on treefall in African savannas. *Ecology Letters*, **15** (11), 1211–1217.

Benitez, L. & Queenborough, S.A. (2021) Fruit trees drive small-scale movement of elephants in Kibale National Park, Uganda. *Biotropica*, **53**(6), 1620–1630.

Berzaghi, F., Bretagnolle, F., Durand-Bessart, C. & Blake, S. (2023) Megaherbivores modify forest structure and increase carbon stocks through multiple pathways. *Proceedings of the*

National Academy of Sciences of the United States of America, **120**(5), e2201832120.

Bezangoye, A. & Maisels, F. (2010) Great ape and human impact monitoring in the Lopé-Waka Exceptional Priority Area, Gabon. Part 1: Lope National Park. GACF Agreement: 98210-8-G529. Final performance report to USFWS. In, p. 62. WCS.

Blair, J.B., Rabine, D.L. & Hofton, M.A. (1999) The laser vegetation imaging sensor: a medium-altitude, digitisation-only, airborne laser altimeter for mapping vegetation and topography. *ISPRS Journal of Photogrammetry and Remote Sensing*, **54**(2–3), 115–122.

Blake, S. (2002) The ecology of forest elephant distribution and its implications for conservation.

Blake, S., Deem, S.L., Mossimbo, E., Maisels, F. & Walsh, P. (2009) Forest elephants: tree planters of The Congo. *Biotropica*, **41**(4), 459–468.

Blake, S. & Inkamba-Nkulu, C. (2004) Fruit, minerals, and forest elephant trails: do all roads lead to Rome? *Biotropica*, **36**(3), 392–401.

Bond, W.J. (1994) Keystone species. In: Schulze, E.D. & Mooney, H.A. (Eds.) *Biodiversity and ecosystem function*. Berlin, Heidelberg: Springer Berlin Heidelberg, pp. 237–253.

Bouvier, M., Durrieu, S., Fournier, R.A. & Renaud, J.P. (2015) Generalizing predictive models of forest inventory attributes using an area-based approach with airborne LiDAR data. *Remote Sensing of Environment*, **156**, 322–334.

Campos-Arceiz, A. & Blake, S. (2011) Megagardeners of the forest–the role of elephants in seed dispersal. *Acta Oecologica*, **37**(6), 542–553.

Cardoso, A.W., Malhi, Y., Oliveras, I., Lehmann, D., Ndong, J.E., Dimoto, E. et al. (2020) The role of forest elephants in shaping tropical forest–savanna coexistence. *Ecosystems*, **23**, 602–616.

Cazzolla Gatti, R., Castaldi, S., Lindsell, J.A., Coomes, D.A., Marchetti, M., Maesano, M. et al. (2015) The impact of selective logging and clearcutting on forest structure, tree diversity and above-ground biomass of African tropical forests. *Ecological Research*, **30**, 119–132.

Coverdale, T.C., Kartzinel, T.R., Grabowski, K.L., Shriver, R.K., Hassan, A.A., Goheen, J.R. et al. (2016) Elephants in the understory: opposing direct and indirect effects of consumption and ecosystem engineering by megaherbivores. *Ecology*, **97**(11), 3219–3230.

Davies, A.B. & Asner, G.P. (2014) Advances in animal ecology from 3D-LiDAR ecosystem mapping. *Trends in Ecology & Evolution*, **29**(12), 681–691.

Doughty, C.E., Roman, J., Faurby, S., Wolf, A., Haque, A., Bakker, E.S. et al. (2016) Global nutrient transport in a world of giants. *Proceedings of the National Academy of Sciences of the United States of America*, **113**(4), 868–873.

Drake, J.B. (2001) *Estimation of tropical forest aboveground biomass using large-footprint lidar*. College Park: University of Maryland.

Dubayah, R., Blair, J.B., Goetz, S., Fatoyinbo, L., Hansen, M., Healey, S. et al. (2020) The global ecosystem dynamics investigation: high-resolution laser ranging of the Earth's forests and topography. *Science of Remote Sensing*, **1**, 100002.

Dubayah, R., Tang, H., Armston, J., Luthcke, S., Hofton, M. & Blair, J. (2021) *GEDI L2B canopy cover and vertical profile metrics data global footprint level V002* [data set]. Sioux Falls, SD: NASA EOSDIS Land Processes DAAC.

Dubayah, R.O. & Drake, J.B. (2000) Lidar remote sensing for forestry. *Journal of Forestry*, **98**(6), 44–46.

Enquist, B.J., Abraham, A.J., Harfoot, M.B., Malhi, Y. & Doughty, C.E. (2020) The megabiota are disproportionately important for biosphere functioning. *Nature Communications*, **11**(1), 699.

Gobush, K.S., Edwards, C.T.T., Maisels, F., Wittemyer, G., Balfour, D. & Taylor, R.D. (2021) Loxodonta cyclotis (errata version published in 2021). The IUCN Red List of Threatened Species 2021, e.T181007989A204404464.

Goetz, S., Steinberg, D., Dubayah, R. & Blair, B. (2007) Laser remote sensing of canopy habitat heterogeneity as a predictor of bird species richness in an eastern temperate forest, USA. *Remote Sensing of Environment*, **108**(3), 254–263.

Goetz, S.J., Steinberg, D., Betts, M.G., Holmes, R.T., Doran, P.J., Dubayah, R. et al. (2010) Lidar remote sensing variables predict breeding habitat of a neotropical migrant bird. *Ecology*, **91**(6), 1569–1576.

Hall, J.S., Harris, D.J., Medjibe, V. & Ashton, P.M.S. (2003) The effects of selective logging on forest structure and tree species composition in a central African forest: implications for management of conservation areas. *Forest Ecology and Management*, **183**(1–3), 249–264.

Hart, J., Gobush, K., Maisels, F., Wasser, S., Okita-Ouma, B. & Slotow, R. (2021) African forest and savannah elephants treated as separate species. *Oryx*, **55**(2), 170–171.

Haynes, G. (2006) Mammoth landscapes: good country for hunter-gatherers. *Quaternary International*, **142**, 20–29.

Inogwabini, B.I., Ngama-Nkosi, M., Wema-Wema, L. & Longwango, M. (2013) Elephant effect on forest physical structure and plant species composition in Salonga and Malebo (lac Tumba landscape), Democratic Republic of Congo. *Pachyderm*, **53**, 28–37.

Kamga, S.M., Tamungang, S.A., Awa, T., Ewome, F.L., Motombi, F.N., Hořák, D. et al. (2022) The importance of Forest elephants for vegetation structure modification and its influence on the bird Community of a mid-Elevation Forest on Mount Cameroon, west-Central Africa. *Diversity*, **14**(3), 227.

LaRue, E.A., Wagner, F.W., Fei, S., Atkins, J.W., Fahey, R.T., Gough, C.M. et al. (2020) Compatibility of aerial and terrestrial LiDAR for quantifying forest structural diversity. *Remote Sensing*, **12**(9), 1407.

Maicher, V., Delabye, S., Murkwe, M., Doležal, J., Altman, J., Kobe, I.N. et al. (2020) Effects of disturbances by forest

elephants on diversity of trees and insects in tropical rainforests on Mount Cameroon. *Scientific Reports*, **10**(1), 21618.
Maisels, F., Blake, S. & Turkalo, A.K. (2002) Wild forest elephants shake down fruit and leaves from trees. *Pachyderm*, **33**, 88–90.
Maisels, F., Strindberg, S., Blake, S., Wittemyer, G., Hart, J., Williamson, E.A. et al. (2013) Devastating decline of forest elephants in Central Africa. *PLoS One*, **8**(3), e59469.
McLean, K.A., Trainor, A.M., Asner, G.P., Crofoot, M.C., Hopkins, M.E., Campbell, C.J. et al. (2016) Movement patterns of three arboreal primates in a neotropical moist forest explained by LiDAR-estimated canopy structure. *Landscape Ecology*, **31**, 1849–1862.
Metsio Sienne, J., Buchwald, R. & Wittemyer, G. (2014) Differentiation in mineral constituents in elephant selected versus unselected water and soil resources at central African bais (forest clearings). *European Journal of Wildlife Research*, **60**, 377–382.
R Core Team. (2022) *R: a language and environment for statistical computing*. Vienna, Austria: R Foundation for Statistical Computing. https://www.R-project.org/
Rakotonarivo, S.O., Bell, A.R., Abernethy, K., Minderman, J., Duthie, A.B., Redpath, S. et al. (2021) The role of incentive-based instruments and social equity in conservation conflict interventions. *Ecology and Society*, **26**(2), 8.
Remis, M.J. & Jost Robinson, C.A. (2020) Elephants, hunters, and others: integrating biological anthropology and multispecies ethnography in a conservation zone. *American Anthropologist*, **122**(3), 459–472.
Roussel, J. & Auty, D. (2023) Airborne LiDAR data manipulation and visualization for forestry applications. R package version 4.0.3. https://cran.r-project.org/package=lidR
Roussel, J.R., Auty, D., Coops, N.C., Tompalski, P., Goodbody, T.R., Meador, A.S. et al. (2020) lidR: an R package for analysis of airborne laser scanning (ALS) data. *Remote Sensing of Environment*, **251**, 112061.
Silva, C.A., Saatchi, S., Garcia, M., Labriere, N., Klauberg, C., Ferraz, A. et al. (2018) Comparison of small-and large-footprint lidar characterization of tropical forest aboveground structure and biomass: a case study from Central Gabon. *IEEE Journal of Selected Topics in Applied Earth Observations and Remote Sensing*, **11**(10), 3512–3526.
Stanbrook, R. (2018) Assessing the nutrient status of elephant dung in the Aberdare National Park, Kenya. *Pachyderm*, **59**, 86–90.
Struhsaker, T.T., Lwanga, J.S. & Kasenene, J.M. (1996) Elephants, selective logging and forest regeneration in the Kibale Forest, Uganda. *Journal of Tropical Ecology*, **12**(1), 45–64.
Sullivan, M.K., Biessiemou, P.A.M., Niangadouma, R., Abernethy, K., Queenborough, S.A. & Comita, L. (2022) A decade of diversity and forest structure: post-logging patterns across life stages in an Afrotropical forest. *Forest Ecology and Management*, **513**, 120169.
Tang, H., Armston, J., Hancock, S., Hofton, M., Blair, J.B., Fatoyinbo, T. et al. (2018) *AfriSAR: canopy cover and vertical profile metrics derived from LVIS, Gabon, 2016*. Oak Ridge, Tennessee, USA: ORNL DAAC. Available from: https://doi.org/10.3334/ORNLDAAC/1591
Terborgh, J., Davenport, L.C., Niangadouma, R., Dimoto, E., Mouandza, J.C., Scholtz, O. et al. (2016) Megafaunal influences on tree recruitment in African equatorial forests. *Ecography*, **39**(2), 180–186.
Unger, M., Homeier, J. & Leuschner, C. (2012) Effects of soil chemistry on tropical forest biomass and productivity at different elevations in the equatorial Andes. *Oecologia*, **170**, 263–274.
Vanleeuwé, H. & Gautier-Hion, A. (1998) Forest elephant paths and movements at the Odzala National Park, Congo: the role of clearings and Marantaceae forests. *African Journal of Ecology*, **36**(2), 174–182.
White, L. & Abernethy, K. (1997) *A guide to the vegetation of the Lopé reserve, Gabon*. New York, NY: Wildlife Conservation Society.
White, L.J. (1994) The effects of commercial mechanised selective logging on a transect in lowland rainforest in the Lope Reserve, Gabon. *Journal of Tropical Ecology*, **10**(3), 313–322.
White, L.J. (1995) Factors affecting the duration of elephant dung piles in rain forest in the Lopé Reserve, Gabon. *African Journal of Ecology*, **33**(2), 142–150.
White, L.J., Tutin, C.E. & Fernandez, M. (1993) Group composition and diet of forest elephants, Loxodonta africana cyclotis Matschie 1900, in the Lopé Reserve, Gabon. *African Journal of Ecology*, **31**(3), 181–199.
Wolf, A., Doughty, C.E. & Malhi, Y. (2013) Lateral diffusion of nutrients by mammalian herbivores in terrestrial ecosystems. *PLoS One*, **8**(8), e71352.

Supporting Information

Additional supporting information may be found online in the Supporting Information section at the end of the article.

Table S1. Field measurements of elephant trails in Marantaceae forests collected in Lopé National Park, Gabon, in January 2022.
Figure S1. DRL and LVIS leaf area density comparison on elephant-and-human engineered trails in Marantaceae forests.
Figure S2. DRL and LVIS leaf area density comparison on elephant-and-human engineered forests in mature forests.
Figure S3. DRL and LVIS leaf area density comparison in forest galleries on elephant-engineered trails.

