The Global Canopy Atlas: analysis-ready maps of 3D structure for the world's woody ecosystems

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1 Abstract

Woody canopies regulate exchanges of energy, water and carbon, and their three-dimensional

(3D) structure supports much of terrestrial biodiversity. Remote sensing technologies such as

airborne laser scanning (ALS) now enable the 3D mapping of entire landscapes. However, we

lack the large, harmonized and geographically representative ALS collections needed to build

a global picture of woody ecosystem structure. To address this challenge, we developed the

Global Canopy Atlas (GCA): 3,458 standardized and analysis-ready maps of canopy height

and elevation at 1 m^2 resolution derived from high-resolution ALS acquisitions. The GCA

covers 56,554 km^2 across all major biomes. Of its acquisitions 19% of have been scanned

multiple times and 87% are publicly available, covering 95% of the total area. To showcase its

wide range of applications, we applied the GCA in three case studies. First, we validated three

global satellite-derived canopy height maps, finding poor performance at native resolution (1-

30 m, $R^2 < 0.38$) and moderate performance at 250 m resolution ($R^2 < 0.65$). Second,

analyzing global patterns in canopy gap size frequency we discovered an unexpectedly large

variation of power law exponents from branch to stand level (α = 1.52 to 2.38), pointing to

fundamental scale-dependencies of forest structure. Third, we developed a framework to

standardize forest turnover quantification from multi-temporal ALS. Applied to repeat

acquisitions in a temperate forest in North America, it revealed that 21% of canopy gaps closed

within 12 years of opening and would thus have been missed by infrequent monitoring. As

demonstrated by these case studies, the GCA provides a novel data source for ecologists,

foresters, remote sensing scientists and the ecosystem modelling community that

substantially advances our ability to understand the structure and dynamics of woody

ecosystems at global scales.

Key words: airborne laser scanning, ecosystem structure, digital terrain model, canopy

height, satellite remote sensing, forest gaps, scaling law, disturbance, canopy closure

1. Introduction

Over the past two decades, airborne laser scanning (ALS) has become integral to the environmental sciences (Beland et al., 2019; Eitel et al., 2016). Laser pulses emitted from aircraft penetrate deep into vegetation canopies, and – by reflecting off leaves, branches and trunks – create a three-dimensional (3D) image of vegetation structure. This information is now routinely used for mapping forest resources (Næsset et al., 2004; Nilsson et al., 2017), microclimates (Gril et al., 2023) and biodiversity (Davies & Asner, 2014; Toivonen et al., 2023). It is also central to the upscaling of vegetation properties such as aboveground carbon stocks (Asner et al., 2014; Coops et al., 2021; Ometto et al., 2023; Xu et al., 2017). At submetric accuracy, ALS reveals an ecological complexity that we are only just finding a language for (Atkins et al., 2023; Lines et al., 2022), and, with flight automation and frequent surveying on the horizon (Besson et al., 2022), it can help uncover the fundamental processes of how organisms, communities and their biomass develop in space and time (Battison et al., 2024; Jackson et al., 2024; Næsset et al., 2013). These insights will enable us to train a new generation of satellite monitoring tools (Kellogg et al., 2020; Quegan et al., 2019), calibrate high-resolution vegetation models (Fischer et al., 2020; Shugart et al., 2015), and shape conservation policies (Gonzalez et al., 2023). However, to build a comprehensive picture of the 3D structure and dynamics of ecosystems, we must move beyond single-time, single-site studies and harmonize ALS data and derivatives at global scales (Valbuena et al., 2020).

ALS data are increasingly available for this purpose. Most countries in Europe and North America now have substantial ALS coverage, with wall-to-wall acquisitions in many areas at regional scale or below (Moudrý et al., 2024; White et al., 2025), and facilities such as OpenTopography (https://opentopography.org), 3DEP (https://www.usgs.gov/3d-elevation-program) or ELVIS (https://elevation.fsdf.org.au) consolidate ALS data from a wide range of sources in the United States, New Zealand and Australia. As an alternative to wall-to-wall coverage, targeted ALS campaigns have been employed to collect representative samples over large biomes (Wulder, White, Nelson, et al., 2012). Examples include the Amazon

(Ometto et al., 2023), the Congo Basin (Xu et al., 2017), Borneo (Melendy et al., 2018), Peru

(Asner et al., 2014), and Canada's boreal forests (Wulder, White, Bater, et al., 2012), as well

as data-rich ecological "supersites" (Kampe, 2010; Karan et al., 2016; White et al., 2019).

However, despite these advances, there is still a distinct lack of harmonization at global scales

(Stereńczak et al., 2020). Three issues stand out in particular. First, the acquisition of ALS

varies considerably across locations and projects. They are acquired in different seasons, with

different instruments, at different altitudes, and often without consistent reporting standards –

all of which can introduce substantial uncertainties into assessments of ecosystem 3D

structure (Almeida et al., 2019; Fischer et al., 2024; Joerg et al., 2012; Næsset, 2005, 2009;

Okyay et al., 2019; Riofrío et al., 2022; Roussel et al., 2017). Second, the processing of ALS

data varies widely. It lacks minimum quality specifications and is carried out with different

algorithms, for different aims, and often with little documentation. Ecologically relevant

quantities such as canopy height, cover, and density can therefore come with errors that are

difficult to correct for and make robust comparisons across studies difficult if not impossible

(Fischer et al., 2024; Kissling & Shi, 2023; LaRue et al., 2022; Quan et al., 2021; Vincent et

al., 2023; Zhang et al., 2024). Third, the accessibility of ALS data and derived products is

inconsistent, with data provided in different formats, with varying data sharing policies and

access protocols. It requires specialist knowledge and considerable time to find, download

and harmonise ALS data, which substantially slows progress towards global syntheses.

To tackle these challenges, we here present the Global Canopy Atlas (GCA), a

harmonised database of 3,458 ALS-derived maps of canopy height and elevation at 1 m^2

resolution, covering more than 56,000 km^2 of the world's woody ecosystems. Its ALS samples

have been processed using a harmonised pipeline designed to enable robust comparisons

across acquisitions (Fischer et al., 2024). Products include several digital terrain models

(DTMs, ground elevation), digital surface models (DSMs, total surface elevation), and canopy

height models (CHMs, top canopy height), all delivered in an easy-to-share GeoTIFF format.

Products are supplemented with a range of ancillary spatialized data quality layers and

comprehensive metadata recording everything from acquisition dates, instrumentation, flight

specifications and geolocation. Around 87% of the GCA products have been derived from

openly available ALS acquisitions, and to facilitate their use we will make them freely available

through the European Space Agency's Multi-Mission Algorithm and Analysis Platform

(MAAP), including thorough documentation and links to the original ALS data sources. In doing

so, the GCA provides a novel resource for ecological research, global change studies, and

the environmental sciences more broadly. Below, we describe its assembly and contents

before demonstrating its range of applications through three case studies that focus on (1) the

validation of three global satellite-derived canopy height models, (2) global patterns in canopy

gap size frequency, and (3) forest turnover quantification from multi-temporal ALS data.

92 2. Database construction

93 2.1 Data aggregation

To assemble the GCA, we searched and documented ALS acquisitions across the globe that

could be converted into high-quality maps of terrain and canopy height. We employed a broad

definition of ALS data, including acquisitions from airplanes and drones (fixed-wing and rotor-

based). We did not include any form of terrestrial laser scanning (TLS) or mobile laser

scanning (MLS).

We followed a number of guidelines during the assembly of the GCA. First, we put a

focus on landscapes with natural woody vegetation cover, i.e., areas without large-scale

human interventions such as urban and agricultural zones or plantations of introduced

species. Second, we prioritised representativity and balanced sampling in geographical and

environmental space. This means, when downloading data from publicly available ALS

collections, we selected subsets covering areas of at least 1 km^2, ideally 5-10 km^2, a wide

geographic range, and a diversity of ecosystem types. Third, we only included acquisitions

where raw ALS data – so-called "point clouds" – were available, so that a standardized

processing pipeline could be applied (Fischer et al., 2024). Fourth, we filtered data according

to a set of minimum quality standards. These included a target sampling density of at least 2

pulses m^{-2} (based on metadata information), compliance with the LAS format standard

(https://www.asprs.org/wp-content/uploads/2019/07/LAS_1_4_r15.pdf, last accessed on 26

August 2025), and the availability of metadata for flight configuration and acquisition dates.

We also prioritised "leaf-on" over "leaf-off" acquisitions for ecosystems that are deciduous.

This meant not including a wide range of openly available wall-to-wall ALS data, such as

available through governmental agencies, as these are often insufficiently documented, low in

pulse density or from winter acquisitions. We note, however, that these were broad guidelines.

We made exceptions to improve geographic coverage, or to sample sites of particular

ecological interest. Therefore, the GCA also includes acquisitions from heavily

anthropogenically modified landscapes, with densities <2 pulses m^{-2}, or with imprecise

metadata on flight dates. Areal coverage can reach several 100 km^2 in national parks, but also

go down to a few hectares for drone acquisitions.

Using these guidelines, we assembled the GCA from a wide range of sources. An

overview over all datasets that we included is provided in Table S1. We downloaded open

collections from dedicated ecosystem observatories such as the Brazilian Sustainable

Landscapes program (https://www.paisagenslidar.cnptia.embrapa.br/, this and all following

links accessed on 26 August 2025), the United States' National Ecological Observatory

Network (Kampe, 2010) and Australia's Terrestrial Ecosystem Research Network (TERN,

http://portal.tern.org.au/). We supplemented them with ALS campaigns such as the Estimativa

de Biomassa na Amazônia (EBA) initiative (https://www.ccst.inpe.br/projetos/eba-estimativa-

de-biomassa-na-amazonia/), the World Wildlife Fund's (WWF) acquisitions over the

Democratic Republic of Congo (Xu et al., 2017) and individual research projects. We

complemented targeted ALS acquisitions with openly available data from country-specific

websites (e.g., https://geoservices.ign.fr/lidarhd in France) and collections such as

OpenTopography, with a focus on protected areas and those with a low human footprint.

Additionally, we contacted research groups who had acquired ALS data over large

environmental gradients or at specific research sites (Stereńczak et al., 2020) and invited them

to contribute to the database.

2.2 Data processing and quality checks

All data processing was carried out with the same standardized pipeline, which is described in detail in Fischer et al. (2024). The pipeline is fully automatic and based on a few key software tools, mainly the laser scanning software LAStools and the R packages data.table (Dowle & Srinivasan, 2025), terra (Hijmans, 2025), sf (Pebesma, 2018), and lidR (Roussel et al., 2020). It carries out several pre- and post-processing steps, such as the splitting of ALS data into coherent spatial units as well as noise filtering, deduplication, and ground classification. For each individual scan, the output is a set of DTMs, DSMs, and CHMs, as well as several types of ancillary layers for quality control. For most applications, we recommend the default DTM and a CHM derived either from a triangulated irregular network (TIN) or a locally adaptive version of the spikefree algorithm (Khosravipour et al., 2016), both of which performed well in a robustness assessment (Fischer et al., 2024). An overview of the different types of layers generated by the pipeline is provided in Table 1, with further information in Table S2.

Throughout processing, we kept all parameters constant across ALS acquisitions, with only few manual adjustments. For example, pre-classified noise points (LAS class 7 and 18) were always excluded, but for some datasets we also excluded custom noise classes defined by the data providers. These noise classes reflected manual corrections of scan artefacts or dense noise that would not be captured by automatic noise detection. In terms of geolocation, our default approach was to retain any provided metric coordinate system to ensure consistency with metadata. However, if acquisitions were provided in imperial units (ft), we converted them into SI units and Universal Transverse Mercator (UTM) coordinates to ensure global comparability. The same approach – retaining the original data configuration where possible – was applied to vertical geo-referencing, so acquisitions were generally processed with the provided ellipsoid or geoid reference. While this does not affect woody ecosystem structure quantification (CHMs), there may be a location-dependent offset between ellipsoidal and geoidal datum up to around 100m (DTMs and DSMs).

Map type	Products	Description	Example
Digital surface model (DSM)	• dsm_lspikefree.tif • dsm_tin.tif • dsm_highest.tif	Three maps of combined ground elevation and vegetation height at 1 m resolution. They reflect varying methods of interpolating top height. The default is a "spikefree" model (dsm_lspikefree.tif). The raster of highest laser returns (dsm_highest.tif) should be used cautiously, as it underestimates height in low-pulse density areas.	
Digital terrain model (DTM)	• dtm.tif • dtm_lasdef.tif • dtm_supplied.tif • dtm_lasfine.tif • dtm_highest.tif	Five maps of ground elevation at 1 m resolution, interpolated from ground returns. They reflect varying methods of classifying returns as ground and interpolating between them. The default (dtm.tif) builds on the standard ground classification algorithm of the LAStools software but refines it on steep slopes (>= 40°).	
Canopy height model (CHM)	• chm_lspikefree.tif • chm_tin.tif • chm_highest.tif	Three maps of vegetation height at 1 m resolution. Each CHM is derived from the corresponding DSM by subtracting the default DTM (dtm.tif).	
Pulse density	• pulsedensity.tif • pulsedensity_last return.tif • pulsedensity_sca nangle20.tif	Three maps of laser pulse density at 1 m resolution. They reflect different methods for pulse density inference (first returns, last returns, first returns with scan angles < 20°). The default is first return density (pulsedensity.tif), which also includes single returns. Best aggregated to coarser resolution (5-10 m) to reduce small-scale noise.	
Scan angle	• scanangle_abs.tif	One map of absolute scan angle at 1 m resolution. Reflects the mean scan angle of pulses that sampled a particular 1 m² pixel.	
Classification	• classification_su pplied.tif	One map of land classification at 1 m resolution. Only available if laser scans have been pre-classified by data provider. Reflects default lidar classes as defined in the LAS standard. If present, can be used to differentiate buildings (class 6) and rail or road surfaces (classes 10 and 11) from vegetation (classes 3 to 5).	

Table 1: Overview of types of spatial layers provided as part of the Global Canopy Atlas. For several of the derived layers (e.g., CHMs and DTMs) we provide a range of alternative options generated using different algorithms which are described in Table S2 and in Fischer et al. (2024). We briefly summarise these in the description provided above and highlight the suggested default maps for most purposes. Sample data come from a 2020 scan at the Sepilok Forest Reserve in Malaysian Borneo and show a 1 km² tile of where the reserve (right) borders an oil palm plantation (left).

Data quality standards varied across data providers and instrumentation, with errors ranging from negligible (isolated noise points) to considerable but repairable (faulty geolocation, flight line misalignment) to substantial and beyond repair (insufficient canopy penetration or flight line overlap). To process large numbers of ALS acquisitions, we developed automatic

procedures for quality assurance and control. The vast majority of these quality checks are

directly embedded in the processing pipeline and provided as ancillary layers alongside the

DTMs and CHMs (Table 1, Table S2). This gives users flexibility in choosing appropriate

quality levels before and during data analysis. General quality layers include pulse density

rasters that enable users to identify areas of poor sampling (pulsedensity.tif), derived pulse

density masks (mask_pd02.tif and mask_pd04.tif, subsetting to regions with >2 or 4 pulses m$^-$

^{2}, respectively), rasterizations of scan angles (scanangle_abs.tif), which enable identification

of flightline patterns, as well as masks for potential high noise or cloud points (mask_cloud.tif).

In addition, we provide several alternative DTM layers to assess robustness of ground

detection, which is a major source of uncertainty on steep slopes and under dense forest cover

(Stereńczak et al., 2016), as it requires both dense sampling and high laser power. Users can

investigate how and where DTMs disagree or use pre-computed DTM masks that highlight

areas with high uncertainty (mask_unstabledtm.tif) and those on steep slopes where features

such as cliffs hamper robust ground classification (mask_steep.tif).

2.3 Processing reports

Each processed scan is accompanied by an automatic two-page PDF processing report (Fig.

S1, also provided as separate pdf in the Supplementary). The first page provides an informal

quality check, showing the scan outline overlaid on local maps from OpenStreetMap and

optical imagery from ESRI, downloaded via the R basemaps package (Schwalb-Willmann,

2024). The second page provides a more detailed and quantitative overview. It shows a pulse

density rasterization where areas with pulse densities <2 m^{-2} are flagged to provide a rapid

assessment of lower-quality scan areas, as well as a DTM reliability mask

(mask_unstabledtm.tif). It also shows the timing of the scan with respect to vegetation

greenness/phenology (NDVI and EVI) and snow cover obtained from the MODIS Aqua and

Terra sensors (Didan, 2025). Finally, the second page of the processing report also compares

ALS-derived products with three satellite-derived layers: the Copernicus World DSM at 90 m

resolution (European Space Agency, 2024), top canopy height (RH98) estimated from

GEDI/ICESat-2 at 100 m resolution (Saatchi & Favrichon, 2023), and global canopy height

derived from Landsat at 30 m resolution (Potapov et al., 2022). ALS data provide more

accurate representations of terrain and vegetation structure than satellites, but we have found

the comparison useful for practical purposes. First, it helps identify potential geo-location

errors. For example, correlations between ALS DSMs and the Copernicus World DSM are

generally high ($R^2 > 0.9$), so low correlations can indicate assignment to the wrong UTM zone

or constant horizontal offsets. Second, the comparison also provides a quick indicator of how

well common satellite products represent local vegetation structure, including temporal offsets,

and thus offers an insight into upscaling opportunities, a key application of ALS (Coops et al.,

2021).

3. Database overview

The GCA database consists of 3,458 ALS acquisitions converted into high-resolution maps of

vegetation height and terrain elevation (Fig. 1a). These vary considerably in size from 0.1 to

203.7 km^2 (95% interval). The vast majority (85%) cover 1 km^2 or more, and the overall median

area is 8.5 km^2. All acquisitions together cover 56,554 km^2 of land area spread across 56

countries and 224 ecoregions (Dinerstein et al., 2017). About 57% of this area (31,554 km^2)

is in intact forest landscapes (Potapov et al., 2017) or has some form of protection designation

(based on World Database of Protected Areas). About 95% of the area – represented by 87%

of the acquisitions – is accessible through openly available data. The total file size of GCA

derived products is 3.2 TB.

Geographic coverage of the GCA extends to all biomes and biogeographic realms (Fig.

1). The total number of acquisitions in the tropics is nearly twice as large (1,950) as in

temperate regions (1,102) and six times as large as in the boreal zone (336). However,

temperate and boreal acquisitions tend to cover larger areas and consequently the GCA

comprises a total scan area of 20,271 km^2 (36% of the total) in the tropics, 27,543 km^2 in the

temperate zone (49%, including Mediterranean forests), and 6,898 km^2 in boreal regions

(12%). A few acquisitions cover other biomes, such as mangroves and highly xeric systems

with little woody canopy cover (1,842 km^2, 3%).

In terms of bioclimatic space, the GCA spans a large gradient of mean annual

temperatures, going from as low as -10 °C up to 30 °C, and ranges in precipitation from less

than 100 mm a year to over 6,000 mm (Fig. 1b). The woody ecosystem structures sampled in

the GCA are also highly diverse, from open systems with canopy cover of less than 10% and

trees that never exceed 10-20 m, to dense, tall forests where canopy cover reaches close to

100% and trees can exceed 100 m in height (Fig. 1c).

Fig. 1: The Global Canopy Atlas. The distribution of each of the 3,458 individual ALS acquisitions is shown (a)

on a map of the world, (b) in the environmental space, and (c) in terms of woody ecosystem structure. The top and

bottom rows show sample canopy height models (CHMs, 200×200 m) for 12 sample sites that span a wide variety

of ecosystems across the globe. Colours, ranging from dark blue to yellow with increasing height, are rescaled to

fit the range of heights observed in each system, with the maximum height of the vegetation patch shown in the

bottom right corner. The location of these 12 sites is also shown in geographic, environmental and structural space

in panels (a–c) using numeric labels. Closed and open canopy ecosystem layers shown in (a) were derived by

aggregating biomes in the Ecoregions2017 dataset (Dinerstein et al., 2017), for details cf. Table S3. Climatic

variables in panel (b) were extracted from 1-km CHELSA climatologies for the years 1981-2010 (Karger et al.,

2017) and averaged across the entire scan area. Maximum canopy height in panel (c) is defined as the 99th

percentile of canopy height across the entire scan area. Canopy cover at 2 m is calculated as the percentage of

CHM pixels $\geq$ 2 m.

In addition to the extensive sampling of geography, environment and woody ecosystem

structure, the GCA covers large tracts of land that have been scanned more than once (Fig.

S2). The area covered by repeated ALS acquisitions is 10,532 km^2 (19% of the total), but

because several of these sites were scanned on multiple occasions the cumulative area of

repeat acquisitions is 33,954 km^2. This is more than 50% of the single-scan area coverage in

the GCA, and brings the total scan area to 90,508 km^2. Repeat acquisitions are available for

all biomes, with the majority of the area covered by acquisitions on NEON sites in the United

States of America (62%). The median interval between acquisitions is 1.2 years (95% range

from 0.6 to 10.0 years).

In terms of sampling density and data quality, 93% of GCA acquisitions have an

average pulse density >2 m^{-2}, and the median density across all acquisitions is 5.1 m^{-2}. There

is, however, a large spread around these values, with a 95% interval ranging from 1.2 to 123.1

m^{-2}. Temperate acquisitions in the GCA generally have higher pulse densities (median of 8.6

m^{-2}) than their tropical (4.9 m^{-2}) or boreal (3.5 m^{-2}) counterparts, and drone acquisitions have

much higher densities (up to 1,000 m^{-2}) than airplane and helicopter-based ones, but these

high pulse density scans only account for a small fraction of the database (49 acquisitions >

300 pulses m^{-2}).

In terms of temporal coverage, acquisition dates range from 2006 to 2024, with 90%

acquired between 2012 and 2022. More than 74% of acquisitions were completed within a 1-

week period (>94% within three months and >99% within six months). Most acquisitions have

been acquired near or at peak EVI (87% of acquisitions) and NDVI (92%), with no clear

geographic biases in acquisition timing. Uncertainty in ground classification is generally low.

The average scan has topographic uncertainties over less than 2% of its area, and only 51

acquisitions (~1.5%) have topographic uncertainties over more than 20% of the scanned area.

4. Case studies

All three case studies are based on analyses of CHMs generated using the locally adaptive spikefree algorithm (chm_lspikefree.tif) described in Fischer et al. (2024). R code and ancillary data to replicate them are archived on Zenodo (https://doi.org/10.5281/zenodo.16987211).

4.1 Case study 1: How accurate are satellite-derived global canopy height models?

Large-scale models of canopy height derived from satellite data are increasingly being used to map the structure and aboveground carbon storage of woody ecosystems (Liu et al., 2025; Schwartz et al., 2023; Wagner et al., 2025). At global scale, height maps are mostly created by combining wall-to-wall optical satellite imagery with laser scanning data acquired from space-borne or airborne sensors using machine learning. Three global examples include: (1) Potapov et al. (2021, 2022), who used Landsat imagery together with GEDI and ICESat-2 shots to map canopy height at 30 m resolution, (2) Lang et al. (2023), who fused GEDI data and Sentinel-2 imagery to map canopy height at 10 m resolution; and (3) Tolan et al. (2024), who trained models on ALS data and predicted height from Maxar imagery at 1 m. However, while these models generally replicate global-scale gradients in canopy height, questions persist about their accuracy at landscape scale (Besic et al., 2025; Moudrý et al., 2024; Wagner et al., 2024). Optical satellite data have limitations both in open systems, where they underestimate tree cover (Brandt et al., 2020), and in closed canopy forests due to saturation effects (Mutanga et al., 2023). To address these concerns, we assessed the landscape scale accuracy of all three global canopy height models against ALS data from the GCA.

We compared the three satellite-derived height maps against ALS-derived CHMs from across 20 landscapes that had not been used to train the models. These ALS acquisitions span forest types with distinct 3D structures and were collected within 1-2 years of the satellite data used to generate the global models (Table S1.1). Agreement between ALS and the global maps was quantified by calculating the coefficient of determination (R^2) and the relative root mean square error (RMSE), i.e., the RMSE standardized by the mean canopy height. Since

the global canopy height models differed in resolution, we first converted them to 1-m resolution via nearest-neighbour resampling and then calculated the R^2 and RMSE across a wide range of resolutions between 1-250 m. By default, we averaged canopy height at each scale, but global models may represent different aspects of height (Besic et al., 2025), so we also calculated the R^2 and RMSE for maximum height (99th percentile) at each resolution.

Fig. 2. Accuracy of global canopy height models. Panels (a-b) show the agreement between the three global canopy height models and ALS data across 20 landscapes at different spatial resolutions on the basis of R^2 and relative RMSE. Rasters are aggregated to different resolutions via averaging, with ALS data assumed to provide the true value. The distributions at each resolution are summarized via median values (dots) and 25th and 75th

percentiles (error bars). Note that some data points exceeded 250% in relative RMSE, so have been excluded from

the visualized range in panel (b). The panels below show sample canopy height models (2×2 km) from 3

landscapes, including the reference ALS product at 1 m resolution. The height range of these maps is provided by

the scale bars on the right. Examples of landscapes with particularly large errors for each of the three models can

be found in Figs S1.4-1.6, an alternative assessment based on maximum canopy height in Fig. S1.1.

The three global canopy height models generally showed poor agreement with the

reference ALS data (Fig. 4). At their nominal resolutions, R^2 ranged from 0.28 (Lang, 10 m) to

0.31 (Tolan, 1 m) to 0.38 (Potapov, 30 m). The corresponding errors were lowest for Tolan

(relative RMSE = 95%), higher for Potapov (112%) and highest by some margin for Lang

(189%). However, agreement depended strongly on resolution and tended to increase at

coarser scales (Fig. 4a-b). When assessed at a common resolution of 30 m, the Tolan model

clearly outperformed the other two models (R^2 = 0.54 and RMSE = 58% compared to 0.38 and

112% for Potapov, and 0.39 and 181% for Lang). The same was true at 250 m, where the

Tolan model had an R^2 of 0.65 and an RMSE of 43%, compared to the poorer 0.56 and 93%

of Potapov and 0.56 and 170% of Lang. When comparing maximum heights instead of means,

the differences in RMSE between models narrowed (57-69% at 30 m, 40-56% at 250 m; Fig.

S1.1), but the accuracy of the predictions worsened (R2 = 0.38-0.55 at 250 m).

Global scale canopy height products are expected to be less accurate than costly high-

resolution airborne laser scans, but our results highlight strong deficiencies in currently

available global canopy height maps. While these products broadly mirror global height trends

(Fig. S1.3), they miss most of the variation in canopy height within landscapes at their nominal

resolutions (1–30 m). Even at 250 m resolution and for the best performing model (Tolan), the

mean relative error was >40% of the reference height and reached as high as 170% in certain

landscapes. This makes it near-impossible to capture within-landscape variation in canopy

height with reasonable accuracy (Moudrý et al., 2024). We found that some models, such as

the Lang model, predicted maximum canopy height better than mean height (Besic et al.,

2025), but errors still remained above 40%, and R^2 values were even lower than for mean

height. Across the vast majority of sites, the high-resolution approach of Tolan et al. (2024)

was clearly superior to other global maps tested here, with higher R^2 and lower RMSE (Fig.

4), and better representation of landscape variability (Fig. S1.3). This may be due to an

improved detection of trees from high-resolution imagery (Brandt et al., 2020) as well as the

training on ALS data, which offer better resolution and geolocation than spaceborne lasers.

That being said, the Tolan height map also contained notable errors and uncertainties,

underpredicting height at a tropical site in Amazonia by more than 20 m and consistently

misrepresenting cloud cover as canopy gaps in tropical regions (Figs. S1.4-6). Global canopy

height modelling is not even 15 years old (Simard et al., 2011) and we expect it to improve

massively in the coming years, especially with the integration of new data sources (Quegan

et al., 2019). It seems, however, that we are still a long way from operational products for

global forest structure monitoring at medium to high resolution (1-30 m). Combining high-

resolution satellite imagery with ALS for model training – as Tolan et al. (2024) have done –

seems a promising approach. But it would require a more careful screening of input data (e.g.,

removing cloud artefacts) and a wider range of ALS data for calibration. A recent canopy height

model for Amazonia shows considerable potential in this respect (Wagner et al., 2025). We

anticipate that the GCA, with its robust canopy height products available across all biomes,

can serve as a benchmark for the calibration and validation of such models.

4.2 Case study 2: Does the frequency of canopy gap sizes follow power laws?

Treefall gaps are an essential component of forest structure and dynamics (Franklin et al.,

1987). They reinitiate succession, create diverse forest microclimates and habitats (Ritter et

al., 2005), and shape species composition (Denslow, 1987). As legacies of forest disturbance,

they are key proxies for measuring canopy and carbon turnover (Dalagnol et al., 2021; Hunter

et al., 2015; Jucker, 2022) and indices of how humans reshape ecosystem structure (Zhang

et al., 2023). Over the last two decades, large-scale ALS surveys have revealed a fundamental

pattern: small gaps are legion, and large gaps are rare. Regional-scale studies have often

found that these heavy-tailed gap size frequency distributions (GSFD) exhibit power law

scaling (Goodbody et al., 2020; Kellner & Asner, 2009), possibly with a narrow range of

exponents (Asner et al., 2013). If true at global scale, this would imply an astonishing level of

generalization: (1) that forest dynamics are governed by multiplicative processes, (2) that

these processes show little variation from the boreal to the tropics and (3) that they are scale-

invariant, i.e., small-scale disturbances such as branchfalls follow similar rules as large-scale

disturbances such as those associated with wildfires and storms. However, not all studies

show power law scaling (Araujo et al., 2021), and the required scale invariance has never

been systematically tested at global scales. Comparisons between regional studies are

complicated if not impossible, as the inferred exponents depend on how CHMs are derived

(Fischer et al., 2024; White et al., 2018), how gaps are defined (Asner et al., 2013; Lobo &

Dalling, 2014), and how power laws are fitted (Hanel et al., 2017; Wedeux & Coomes, 2015).

Since forest dynamics and structure are scale dependent (Fischer & Jucker, 2024; Shugart et

al., 2010) and power laws difficult to distinguish from other heavy-tailed distributions (Newman,

2005; Taubert et al., 2013), it is an open question whether GSFDs follow power laws across

forest types, spatial scales, and the range of gap sizes .

To systematically tackle this question, we chose a subset of high-quality CHMs from

the GCA with a minimum of 70% canopy cover at 2 m and divided them into 1 km^2 cells using

a flexible Voronoi tessellation (n = 13,300, details on selection in Supplementary S2.1). For

each 1 km^2 cell we calculated the mean canopy height (μ_{CHM}) and defined gaps as contiguous

areas (rook neighborhood) below a height threshold of δ_{CHM} = 0.5 × μ_{CHM} and no larger than

100,000 m^2 (10% of the reference area). This excluded some larger gaps, but was not

expected to alter power law assessments below 100,000 m^2. Gap size was calculated as the

number of 1 m^2 pixels in a gap (Kellner & Asner, 2009), and discrete power laws were fitted

with the poweRlaw R package (Gillespie, 2015). Tests for power law behaviour exist, but they

reduce a continuous spectrum of deviations to binary hypothesis testing and are difficult to

interpret (Hanel et al., 2017). We therefore used an alternative heuristic that quantifies

deviations from scale invariance (Fischer & Jucker, 2024). Under ideal power law behaviour,

the scaling exponent α is scale-invariant. If a power law is fit to the full range of gap sizes and

to a subset (e.g., only large gaps), α should be approximately identical. By contrast, if α shifts

systematically, the distribution deviates from power-law scaling. We calculated this deviation

Δ_{PL} as the difference between α calculated over three different scale ranges. First, we

calculated α for all gaps from small branch falls $\geq 1\ m^2$ up to 100,000 m^2 (α_{branch}). This is the

lower cutoff used in Kellner & Asner (2009). The mean area of all gaps within that range was

32.8 m^2 (95% range: 9.4 to 91.8 m^2). Second, we calculated α only for single treefall gaps or

larger (α_{crown}). Crown area scales with tree height (Jucker et al., 2022), so we approximated

the typical tree crown area as circle with a diameter equivalent to the height cutoff δ_{CHM} in each

1 km^2 cell ($0.25 \times \delta_{CHM}^2 \times \pi$). The corresponding mean gap area was 420.4 m^2 (60.3 to 1,446.7

m^2). Finally, we calculated α only for the tail end of the gap size frequency distribution (α_{stand}).

For this, we calculated the cutoff as the midpoint between the typical tree crown size and the

size of the largest disturbances (95th percentile of gap sizes). The mean gap area was 1,830.9

m^2 (210.0 to 5691.1 m^2). Deviations were then quantified as $\Delta_{PL,\ branch} = \alpha_{crown} - \alpha_{branch}$ and $\Delta_{PL,}$

$_{crown} = \alpha_{stand} - \alpha_{crown}$ and calculated separately for boreal forests (n = 501), temperate forests

(n = 3,903) and tropical forests (n = 8,896). The robustness of results was tested through

several sensitivity analyses (Supplementary S2.2).

Fig. 3: Deviations from power law scaling across major biomes. Panel (a) shows the distribution
of power law exponents of gap size frequency distributions (α) for 1 km² cells across tropical, temperate
and boreal biomes and coloured by the size ranges over which they are calculated (α_{crown}, α_{branch} and
α_{stand}). Coloured points represent the median values of the distributions. Under true power law
behaviour, α should not depend on the size range over which it is calculated, so all distributions should
be indistinguishable, with a near-identical median or mean ($\Delta_{PL} \approx 0$). The righthand column (b–d) shows
results from a sample 1 km² cell in each of the three biomes to illustrate how α varies across scales.
Each black point represents the contribution of a single gap to the the observed complementary
cumulative distribution function (CCDF) of gap sizes, plotted on double-logarithmic scales. Coloured
lines show the fitted power laws. We chose a CCDF representation over size frequency plots, as it
avoids binning into size classes and noise in the tail end (Newman, 2005). Sample data are from a
tropical site in Indonesian Borneo (b), a temperate site in Northwestern United States (c), and a boreal
site in Québec, Canada (d) and were selected to be close to each biome's median in α_{branch} and α_{crown}.

We found that the power law exponent α was highly scale dependent, indicating that GSFDs

do not follow power law behaviour (Fig. 2). When small gaps were included, α was low and

narrowly distributed within and across biomes (mean α_{branch} = 1.52; 95% interval = 1.42-1.70).

By contrast, when focussing on gaps from treefalls or larger disturbances, exponents were

considerably higher and much more variable (α_{crown} = 2.22; 1.73-3.28), even though the gaps

excluded in the α_{crown} calculation only made up 2.59% (0.74-5.21) of the total gap area. Across

biomes, tropical forests generally had the largest α_{crown} and the largest variation around the

mean (2.29; 1.73-3.45), compared to boreal (2.02; 1.76-2.38) and temperate forests (2.08;

1.71-2.61). From branch to crown scale, the typical deviation from power law scaling $\Delta_{PL, branch}$

was 0.70 (0.16-1.73) and was highest in the tropics (0.77) and smallest in boreal landscapes

(0.49). Exponents increased again, albeit much less, when excluding treefall gaps and only

fitting power laws to the largest disturbances (α_{stand} = 2.38; 1.73-3.63 and $\Delta_{PL, crown}$ = 0.25; -

0.23-0.91). Qualitatively, our results were robust to methodological choices, with α changing

across scales and biomes also for different CHM types (Figs. S2.2–2.5) and when applying

fixed thresholds to define gaps (< 2 m) and minimum gap size ($\geq$ 9 m^2, Figs. S2.6–2.7).

Overall, our results provide strong evidence that GSFDs do not follow power laws. The

clear violation of scaling behaviour points to fundamentally different processes shaping forest

structure at small and at large scales (Shugart et al., 2010). When GSFDs include small,

branch-level gaps, inferred power law coefficients tend to be small and highly consistent

across systems (Fig. 2), but at larger scales they exhibit tremendous variability that likely

reflects the "fingerprints" of different disturbance regimes (Jucker, 2022). Violations of

fundamental scaling assumptions likely explain why previous studies have come to different

conclusions about the typical values and variability of power law coefficients (Asner et al.,

2013; Goodbody et al., 2020; Reis et al., 2022), with differences exacerbated by CHM

construction (Fig. 2 vs. Fig. S2.2, White et al., 2018). Based on our results ($\Delta_{PL, crown}$ << $\Delta_{PL,}$

$_{branch}$) and in line with other heavy-tailed distributions (Newman, 2005), it is possible that

GSFDs converge to power law behaviour at the tail end of the distribution, but our case study

reveals a fundamental scale dependence across a large size range. A wider range of functions

to model GSFDs (Araujo et al., 2021) is thus required to build a comprehensive picture of

global forest structure and how it is shaped by natural and anthropogenic disturbances.

4.3 Case study 3: How dynamic are forest canopies?

Accurate monitoring of forest growth and disturbance through time is an essential first step in understanding drivers of forest dynamics and forest responses to global change. Collections such as the GCA bring together ALS data from many sources and years. Some of these acquisitions overlap spatially, which creates the opportunity for large-scale repeat surveying of forests, in particular the monitoring of growth (Yu et al., 2004), disturbance (Huertas et al., 2022), recovery (Krüger et al., 2024), biomass change (Cao et al., 2016; Næsset et al., 2013), structural complexity (Rosen et al., 2024), and tree demographics (Battison et al., 2024). However, estimating change rates from multi-temporal and multi-source ALS comes with its own set of challenges and depends on instrument differences and methodological decisions (Jackson et al., 2024; Leitold et al., 2018). When intervals between acquisitions are short (1-5 years), small biological changes are easily confused with measurement errors (Heiskanen et al., 2024; Riofrío et al., 2022). By contrast, as intervals between acquisitions get longer, disturbances can be missed or underestimated in size due to rapid recovery between acquisitions or repeat disturbances in the same area. These transient disturbances and recovery events parallel the problems of unobserved tree mortality and growth in field inventories with long remeasurement intervals (Kohyama et al., 2019), but have not been explored for repeat ALS surveys. We here leverage a unique timeseries of 8 ALS acquisitions conducted over a period of 12 years at Harvard Forest in the Northeastern United States and propose a new approach to robustly estimate disturbance and recovery rates, instrumentation noise, and transient dynamics.

Harvard Forest is part of the National Ecological Observatory Network (NEON). It is a mixed temperate forest dominated by deciduous broadleaf trees (Quercus rubra, Acer rubrum) and white pine (Pinus *strobus*), with a mean canopy height of ~17 m and canopy cover of ~80%. The GCA contains 12 ALS acquisitions for Harvard Forest, but for the purposes of this analysis we focussed on 8 summer acquisitions conducted by NEON (2012, 2014, 2016, 2017, 2018, 2019, 2022, and 2024), as these cover a large area of overlap (~145 km^2, all pulse densities > 2 pulses m^{-2}). Each scan's CHM was split into tiles of 1 km^2. For each cell, we

defined gaps as contiguous areas (rook neighborhood) below 10 m canopy height. Non-forest

areas were defined as those lower than 10 m in canopy height which changed by less than 2

m over 12 years, and masked out. We calculated dynamics between each pair of NEON

acquisitions (28 combinations), differentiating disturbance (newly formed gap area as

proportion of intact canopy area), and recovery (closed gap area as a proportion of previous

gap area; Jackson et al., 2024). Assuming a typical yearly disturbance rate d and recovery

rate r, expressed as proportion of initial canopy and gap area, we derived an analytic

relationship between those rates and the observed disturbance and recovery for a given time

interval between surveys (Supplementary Material S3.1). This allowed us to separate

between-scan noise (d_0 and r_0 for disturbance and recovery) from biological signal and to

estimate unobserved dynamics by comparing observed rates to the theoretical rates that

would apply if there were no transient events (no short-term recovery after gap formation, no

immediate re-disturbance after closure). We fitted the models via the *brms* R package

(Bürkner, 2018, Supplementary Material S3.2) and explored the sensitivity of results to

modelling decisions (Supplementary Material S3.3-3.4).

We found near steady-state dynamics at Harvard Forest, with disturbance slightly

exceeding recovery (Fig. 3). We estimated a 1-year gap creation rate of d = 0.45 ha km^{-2} (95%

interval = 0.36-0.64) or slightly less than 0.5% of the intact canopy area (~90 ha km^{-2}), and a

1-year gap recovery rate r = 0.41 ha km^{-2} (0.07-1.11), or slightly less than 5% of the disturbed

area (~10 ha km^{-2}). After accounting for biases due to between-scan noise (d_0 = 1.09 ha km^{-2},

0.64-1.54, and r_0 = 0.93 ha km^{-2}, 0.68-1.17), we inferred a total of 5.30 ha km^{-2} (4.26-7.34)

of newly created gaps and 3.86 ha km^{-2} (0.85-7.26) of recovered canopy area over 12 years.

Of the new gaps, an estimated 1.11 ha km^{-2} (~21%) had closed again by 2024, and 0.1 ha km^{-2}

(~3%) of closed gaps had re-opened after closure. In canopies with higher turnover, we

estimated that such transient dynamics could account for up to 57% (short-term recovery of

gaps) and 7% (re-disturbance of recovered area), respectively (d = r = 2.0 ha km^{-2}, Fig. S3.1).

Our estimates varied slightly, but not fundamentally, when introducing a minimum gap size (d

= 0.45 ha km^{-2}, r = 0.34 ha km^{-2}, Fig. S3.2) or subsetting to a smaller area where additional

acquisitions were available (d = 0.58 ha km^{-2}, r = 0.38 ha km^{-2}, Fig. S3.3). Inferences from

single pairs of acquisitions were more uncertain (d ~ 0.24-1.11 ha km^{-2} and r ~ 0.05-1.04 ha

km^{-2}), but the overall averages were consistent (d = 0.45 ha km^{-2}, r = 0.40 ha km^{-2}, Fig. S3.4).

Estimates of biases due to between-scan noise were confirmed independently via temporally

coincident acquisitions (d_0 = 0.89 ha km^{-2}, 0.19-3.19, r_0 = 0.94 ha km^{-2}, 0.28-2.82, Fig. S3.5).

Fig. 4. Short-term dynamics at Harvard Forest. Panels show observed rates of (a) disturbance d and (b) recovery

r as a function of the time interval (years) between successive ALS surveys (black points). As the intervals between

observations get longer (a), an increasing proportion of disturbances recover before being re-surveyed, leading to

an underestimation of true disturbance rates (and equivalently for recovery in panel b). Rates were inferred as

proportion of existing canopy and gap area and then converted into area rates per grid cell (ha km^{-2}). Black lines

are fitted relationships, while coloured lines reflect the theoretical changes that would be expected if there was no

short-term recovery of disturbed area (a) and no rapid re-disturbance of recovered area (b). They are calculated

by extending d and r into the future, discounting only for loss of canopy and gap area, respectively. The intercepts

d_0 and r_0 are non-zero, as a proportion of pixels is expected to switch randomly between gap and non-gap states

between two ALS acquisitions (e.g., due to instrument differences or movement of leaves) and thus introduce a

baseline bias in disturbance and recovery rates. The five panels below show a sample canopy height model at

Harvard Forest (150×150 m) over a 12-year period. White contours indicate gaps or parts of gaps that formed between 2012 and 2014 and rapidly closed in the years afterwards. These would have gone undetected in a direct comparison between 2012 and 2024. The panels show that initial gap recovery stems predominantly from lateral ingrowth around gaps, and within-gap regeneration appears only in 2024 as lighter blue areas within gaps.

Our results show a moderately dynamic canopy at Harvard Forest, with close to 1% of the area (1 ha km^{-2}) either disturbed or recovering year-to-year, and more than 20% of the gap area created by branch or treefalls closing again over 12 years (Fig. 3, bottom panels). This transient turnover, which we cannot observe when ALS acquisitions are repeated infrequently, is expected to be even larger in forest canopies with higher disturbance and recovery rates (Fig. S3.1). To quantify it, we need for medium-to-high frequency monitoring programs (e.g., Cushman et al., 2022) and robust methods that can account for scanning noise. We found that nearly 2% of the pixels were expected to switch from canopy to gap status (~1%) or vice versa (~1%) simply due to measurement noise between ALS acquisitions. This was more than twice the yearly biological changes at Harvard Forest, and if not accounted for, would greatly inflate estimates of canopy dynamics. Single pairs of acquisitions may be particularly affected by these issues (Heiskanen et al., 2024), which can be further compounded by ecological variability, changes in acquisition timing (cf. "winter" outliers in Fig. S3.3a) or instable CHM algorithms (Fischer et al., 2024). Some instrumentation noise may be avoided by careful re-calibration (Riofrío et al., 2022) or by subsetting to larger or more severe disturbances (Fig. S3.2). However, the former is challenging, and the latter does not make biological sense when tracking recovery which is more gradual in nature (Körner, 2003). A more robust approach will consist in the pooling of information across multiple sites and acquisitions per site, ideally via Bayesian hierarchical models (Kohyama et al., 2018). Long-term ecological observatories such as NEON and harmonised collections like the GCA will provide an ideal backbone for such an approach and enable us to sketch a global picture of canopy turnover.

5. Outlook

5.1 In Humboldt's and Bonpland's tailwinds

Ever since Alexander von Humboldt and Aimé Bonpland set out to travel the Americas, comparing ecosystems and communities across large geographic gradients has been a cornerstone of modern Earth system science. On the one hand, ecosystems display a large degree of local idiosyncrasy, shaped as much by their unique geological and climatic past as the accidents of natural and human history. On the other hand, there are commonalities in how ecosystems develop that cut across biomes, continents and civilizations. These are encoded in vegetation structural patterns, such as how height and cover depend on water availability (Staver et al., 2011; Tao et al., 2016), how growing season length limits the elevation range of trees (Paulsen & Körner, 2014), and the successional mosaics left behind by natural and anthropogenic disturbance (Frelich et al., 2018; Jucker, 2022). Describing these patterns, identifying new ones, and linking them to the biophysical processes that underpin them not only enhances our understanding of the natural world, but also improves our ability to predict ecosystem responses to global change in decades to come (Blois et al., 2013). Remote sensing technologies such as ALS are uniquely poised to carry this project forward (Senf, 2022). Much of global ecology so far has been informed by a combination of local field samples and optical satellite layers – providing a "flat" view of the world as a tapestry of landcovers, punctuated by occasional data-rich information spikes grounded in field observations. ALS inserts 3D contours into this 2D world, converting discrete vegetation classes (e.g., forest and non-forest) into continuous vegetation metrics (e.g., vegetation height, cover and density). In doing so it helps us build a fuller and more nuanced picture of woody ecosystem structure and how it reflects current as well as past dynamics (Lenoir et al., 2022).

As we showed here, ALS can be a particularly powerful macroecological tool when acquisitions from different sources, sites and times are harmonized in large databases such as the GCA. High-resolution derived products such as CHMs and DTMs are comparatively

small in size, robust, and easily analysed and interpreted with a wide range of software

(Fischer et al., 2024). These detailed but accessible ecosystem maps shift the focus from

resource-intensive data processing to ecological analysis, model building and technological

innovation (White et al., 2019). As highlighted by our case studies, it requires comparatively

little effort to use 3D maps to validate predictions from satellite-based sensors (case study 1).

carry out robust global comparisons of complex structural features (case study 2) and

precisely quantify ecosystem dynamics through time (case study 3). Naturally, our case

studies offer only a glimpse into the opportunities provided by regional to global ALS

harmonization. Comparisons of ALS-derived maps across and within biomes may

fundamentally shift our conceptualization of ecosystem dynamics (Lines et al., 2022) and

usher in a new generation of ecosystem models (Shugart et al., 2015). Future versions of the

GCA and other similar projects could provide harmonised ALS collections covering even larger

spatial extents, such as wall-to-wall maps of entire countries or continents (Moudrý et al.,

2024). They could also be used to derive additional information from point clouds that goes

beyond rasterized maps of height, such as leaf area densities (Vincent et al., 2023) and

subcanopy structure (Jarron et al., 2020), further enhancing our picture of the 3D structure of

the world's terrestrial ecosystems.

5.2 Flying close to the ground

The new picture of woody ecosystem structure will take time to develop. When Humboldt and

Bonpland arrived in Venezuela, they were so enthralled by the flora and fauna that they ran

from one organism to the next like "madmen" (Humboldt, 1801). It then took them decades to

systematize their knowledge. The characterisation of ecosystem structure from remote

sensing and ALS is likely in a similar situation. Laser scanning has come a long way since its

first applications in forestry, which date back to the 1970s and 1980s (Nelson, 2013). We now

possess vast amounts of data and a wide range of tools to analyse them (Coops et al., 2021;

Næsset, 2002; Roussel et al., 2020), but, in our characterization of global ecosystems, we are

likely still in the initial exploration phase and will require years of consolidation to arrive at

robust categorizations and predictions (Atkins et al., 2018; Knapp et al., 2020; Loke &

Chisholm, 2022; Rosen et al., 2024; Valbuena et al., 2020). The GCA provides a strong

foundation for this enterprise. Its ecosystem maps provide samples from a wide range of

environments (Wulder, White, Nelson, et al., 2012), have been developed to be maximally

robust to differences between acquisitions, and are accompanied by extensive metadata,

quality layers and masks that can be used to identify areas where robustness may be

compromised (Fischer et al., 2024).

Automated data processing and quality checks have their limitations, and for certain

applications (e.g., measurements of within-canopy structure) additional corrections or data

filtering may be needed, including targeted ground corrections (Riofrío et al., 2022), manual

re-classification, or adjustments to account for instrument properties (Roussel et al., 2017;

Vincent et al., 2023). Furthermore, as our case studies show, the need for careful analysis

goes beyond these initial data selection and processing steps and extends to ecological model

building itself. Just because we can predict canopy height from ALS-trained satellite data (case

study 1), fit power laws to gap sizes (case study 2) or calculate structural change between

successive acquisitions (case study 3) does not mean the results are always reliable or useful.

Robust global ecosystem ecology needs to be guided by a biophysical understanding of the

underlying dynamics, a careful selection of metrics to monitor them (Fischer et al., 2024;

Zhang et al., 2024), models that can account and correct for biases, and standardized

methods to assess their predictive power (Ploton et al., 2020). A key challenge going forward

is to link ALS observations back to field data and fundamental ecological units such as trees

and species (White et al., 2016). This will be facilitated by a wide range of methods, such as

manual and automatic tree segmentation (Aubry-Kientz et al., 2021; Weinstein et al., 2021),

model inversion (Shugart et al., 2015), or algorithms that attempt to reconstruct inventories

bottom-up (Fischer et al., 2020; Spriggs et al., 2015; Taubert et al., 2015)

5.3 Flying together

Science is a social enterprise, and global databases such as the GCA are community efforts made possible by the collective investments, work and ideas of researchers across nations, generations and disciplines. While this creates a unique opportunity for collaborative research and data sharing, it also brings a responsibility to promote fair use. Access to and utilization of data are shaped by country-specific procurement models and regulatory openness, but also by historical power asymmetries, such as those that often divide the Global South and Global North (de Angeli Dutra et al., 2025) or countries in and outside the tropics (de Lima et al., 2022). Such asymmetries are a particular concern with regard to remote sensing technologies such as ALS, which rely on expensive instruments and can be employed by domestic or foreign actors without meaningful involvement of local communities or researchers. The open release of such data is key as it creates opportunities for local experts and a wider public to actively contribute to research. Equally important are clear licensing and documentation that acknowledge local data contributors and institutions, ongoing support for acquisitions in undersampled or difficult-to-access areas and further investments in capacity building. Care is also required to avoid misuse of data. Easily accessible remote sensing data could be used for the surveillance of communities (Millner et al., 2024) or to reveal the location of environmental assets (e.g. logging and mining opportunities) and archaeological sites (Fisher et al., 2021), potentially exacerbating disparities where access and safeguards are uneven. Moreover, external actors may violate data-ownership laws or license terms (e.g. non-commercial clauses), or repackage publicly available data into commercial products without proper attribution.

The GCA project cannot solve these problems by itself, but it can take steps to mitigate risks. First, by provisioning standardized ALS-derived products, but not the raw data, the GCA provides added value, but does not supplant the original data sources, which are thoroughly documented and linked to (cf. Table S1). Second, with its focus on simple, light-weight raster products, the GCA reduces the need for expensive computing infrastructure to process and

analyse data. Third, it brings visibility to acquisitions missing from large open data repositories,

which we hope will increase collaboration opportunities for local experts. Finally, the GCA has

a relatively equal distribution of data in space, with representative coverage across the tropics

(Fig. 1), ensuring that new products – often trained and tested in high-latitude temperate and

boreal zones – become more representative of ecosystems and research needs globally. On

balance, we believe that the benefits of making the majority of the GCA's products openly

available outweigh the risks. Most raw data in the GCA are already open, so the publication

of harmonised, easy-to-use derived products should mostly benefit local researchers and

practitioners who do not have the means to find, store and process large volumes of raw data.

Past experience has shown that pre-processed openly available global datasets, such as

meteorological and climate layers (Fick & Hijmans, 2017) and global trait compilations (Zanne

et al., 2009), can greatly benefit researchers world-wide and fill gaps in areas where

specialized local products and data are not as readily accessible.

To ensure the GCA promotes scientific discovery in a fair and equitable manner, we

aim to implement a number of measures to further widen participation in the maintenance and

governance of the database. This includes appointing an advisory committee to guide how

best to continue growing the GCA in a way that does not unintentionally reinforce historical

power imbalances. Similarly, we aim to invite local experts to join the project to help assemble

ancillary datasets for the GCA landscapes so that we can build a richer picture of their ecology

and history. Further, we plan to organise targeted online and in-person training courses

designed to lower the barrier to entry for working with ALS data and facilitate their integration

into existing research programs underway across GCA landscapes. Above all, we hope that

we can encourage users of the database, particularly in the Global North, to contribute to these

goals by involving local experts in their studies and investing in the maintenance and growth

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Competing interests

A Burt and M Demol are employees and/or shareowners of Sylvera Ltd. No other competing

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Author contributions

T Jucker conceived of the idea of the project. FJ Fischer and T Jucker led the data assembly

and construction of the database, with assistance from J Chave, D Coomes, KC Cushman, R

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FJ Fischer led the development of processing workflows and data processing, with assistance

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of the manuscript, with assistance from T Jucker, T Jackson and B Morgan. All co-authors

contributed substantially through data collection, data assembly and critical revisions of the

manuscript.

Data availability

All openly available GCA products (87% of the database, covering 95% of the area) will be

made freely available after review through the European Space Agency's Multi-Mission

Algorithm and Analysis Platform (MAAP). All code and materials necessary to reproduce the

analyses in this manuscript will be made publicly available on Zenodo

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