

# 1 **Systematic losses in tree-canopy cover over three decades revealed by** 2 **integrating complementary data sources**

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## 12 **ABSTRACT**

13 Losses and gains in canopy cover of the world's tree canopies affect carbon stocks, species habitats,  
14 water cycles, and human livelihoods. Consistent and multi-decadal global data on tree-canopy cover  
15 dynamics are needed for modelling climate scenarios, tracking progress towards restoration targets, and  
16 diverse other research, management and policy applications. However, most data only map binary  
17 'forest'/'non forest' distinctions that are regionally restricted or biased by data gaps, and those mapping  
18 tree-canopy cover are limited to the 21st century, leaving longer-term dynamic in tree-canopy cover  
19 largely unknown. Here, we present an annual and global time-series of tree-canopy cover between 1992  
20 and 2018. To develop these data, we integrated complementary products, using their respective  
21 strengths to compensate for weaknesses, and exploiting path dependencies in change processes to derive  
22 predictions into the data-sparse 1990s. Our model validation indicates we can accurately map tree-  
23 canopy cover ( $r^2=0.95$  [ $\pm 0.01$ ], RMSE=6.75% [ $\pm 0.08$ ], F1-score=0.97 [ $< \pm 0.01$ ]) and our time-series  
24 show plausible broad-scale spatiotemporal patterns, as indicated by high correlations with national  
25 statistics ( $r^2=0.94$  [ $< \pm 0.01$ ]). Our analysis of these data revealed systematic global losses in tree-canopy  
26 cover that, area-wise, substantially exceed concurrent losses or gains of treescape extents. Our analysis  
27 and data provide novel insights into global dynamics of tree cover, and can support modelling and

28 reporting in the scope of the Kunming-Montréal Global Biodiversity Framework, the Paris Agreement,  
29 and other forest-related policies.

## 30 **INTRODUCTION**

31 Global dynamics in tree-canopy cover, resulting from changes in the densities, extents, and  
32 fragmentation of tree stands, have diverse impacts on Earth's biota and earth-system processes.  
33 Thinning tree canopies affect biogeochemical cycles directly by reducing carbon stores (Baul et al.,  
34 2021; L. Xu et al., 2018), and indirectly by increasing tree exposure to wind and solar radiation, thereby  
35 increasing their susceptibility to drying, fire, and subsequent carbon losses (Walker et al., 2019). Tree-  
36 canopy cover dynamics also affect biodiversity by altering resources available to different species (e.g.,  
37 shelter, food) and the microclimatic conditions to which they are exposed (Kašpar et al., 2021).  
38 Simultaneously, tree-canopy cover changes variously affect humans via effects on surface-water run-  
39 off (Selbig et al., 2022), soil erosion (Mohammad and Adam, 2010), crop pollination (Krishnan et al.,  
40 2020), urban heat islands (Schwaab et al., 2021), mental health (Lee et al., 2023), and real estate values  
41 (Kovacs et al., 2022).

42 Correspondingly, consistent historical data on the spatiotemporal dynamics of tree-canopy cover are  
43 needed to inform actions towards multiple goals and targets under the 2030 Agenda for Sustainable  
44 Development (Estoque et al., 2021) For example, long time-series of tree-canopy cover can support  
45 climate policy by enabling the modelling of future scenarios (Allen et al., 2017) and definitions of  
46 Nationally Determined Contributions to the Paris Agreement (UNFCCC, 2015). Similarly, long time-  
47 series help assess changes in forests and other wooded ecosystems since 1990, which is a globally  
48 adopted reference year for cutting greenhouse gas emissions (Kuyper et al., 2018).

49 Thanks to technical advances and capacity-building investments over recent decades, remote sensing  
50 increasingly offers cost-effective solutions to support spatiotemporal data requirements on key  
51 properties of global woodlands (Cameron et al., 2019), including on tree-canopy cover. Yet, existing  
52 global data on long-term tree-canopy cover dynamics are available only at coarse spatial resolutions  
53 (Song et al., 2018), with intermediate- and high-resolution products limited to post-2000 and post-2015

54 periods, respectively, or single years before then (Dimiceli et al., 2015; Sexton et al., 2013, Hansen et  
55 al., 2013; Tang et al., 2019).

56 In turn, global remote sensing products extending to earlier years map binary forest/non-forest  
57 distinctions. Those products rely on “forest” definitions based on thresholds of tree-canopy cover (e.g.,  
58 between 10 and 70 percent, alongside other criteria; (Chazdon et al., 2016)), which hinders the  
59 comparability of change reports enabled by different products. Moreover, there are risks of  
60 misinterpreting change magnitudes when using categorical products to analyse forest changes, which  
61 are often slow and progressive, as any change detection depends on crossing a specific threshold in  
62 either direction (Montibeller et al., 2020).

63 Deriving a global and dense time-series of tree-canopy cover requires surpassing satellite data  
64 limitations. For half a century, Landsat became a popular data source for historical forest-mapping  
65 thanks to its global coverage and high spatial and temporal resolution (Hansen et al., 2013; Kim et al.,  
66 2014; Vancutsem et al., 2021). However, the Landsat archive suffers from multi-year data gaps that  
67 limit consistent, global environmental monitoring (Remelgado et al., 2023). Consequently, existing  
68 products extending into the 1990s (all of which, to the best of our knowledge, tackle forest cover and  
69 not tree-canopy cover) are incomplete (Kim et al., 2014), focused on specific countries (Cohen et al.,  
70 2010; Lehmann et al., 2015; MapBiomass, 2021; Stibig et al., 2014; Wulder et al., 2020) or biomes  
71 (Vancutsem et al., 2021), or provide biased perceptions of change due to gradual improvements in  
72 satellite data coverages (Remelgado et al., 2023). Alternatively, to Landsat, AVHRR data offers a  
73 mostly consistent temporal and spatial coverage since the 1980s that enabled mapping long-term tree-  
74 cover changes (Song et al., 2018). Yet, its 5-km spatial resolution is too coarse to monitor forest changes  
75 reliably, given that those commonly occur at finer scales (Montibeller et al., 2020).

76 Predictive models that integrate heterogeneous data with support of covariates provide a unique  
77 opportunity to tackle these data limitations. Such models can capitalise on the ability of different  
78 remote-sensor systems and mapping algorithms to perceive different facets of forest dynamics (Wang  
79 et al., 2020). Model-based data integration can borrow information across spatial and temporal scales,  
80 building on the facts that local forest dynamics are characterised by path-dependent processes  
81 (Montibeller et al., 2020; Taubert et al., 2018), and that change dynamics are influenced by cross-scale

82 interactions (Soranno et al., 2014, **Fig. 1**). For example, although gaps in satellite imagery hamper direct  
83 fine-scale mapping of forest-change processes pre-2000, data for later years may still bear signals of  
84 earlier dynamics, and thus help reveal longer-term changes. Similarly, consistent coarser-resolution  
85 observations over longer periods capturing historical changes across larger landscapes may help  
86 determine the timing, direction, and magnitude of fine-scale changes. Yet, the potential of model-based  
87 data integration for global time-series mapping remains underutilised. Correspondingly, global, long-  
88 term dynamics of tree-canopy cover at sub-kilometre resolutions remain largely unknown.

89 Here, we leveraged these concepts by integrating state-of-the-art and high-resolution gridded data  
90 on tree-canopy cover and tree-cover losses (Hansen et al., 2013) with that on tree-canopy cover change  
91 - and drivers of that change - mapped at coarser spatial resolutions. The result is the Global Tree-Canopy  
92 Cover Change dataset (GTCCC), a global time-series of tree-canopy cover and change between 1992  
93 and 2018 mapped annually at a 300-m resolution. We use this synthesis product to analyse longer-term  
94 historical dynamics of tree-canopy cover, providing novel insights into their timing and speed, as well  
95 as into the magnitude, direction, and generality of net changes.

## 96 **MATERIALS AND METHODS**

### 97 Overview of modelling strategy

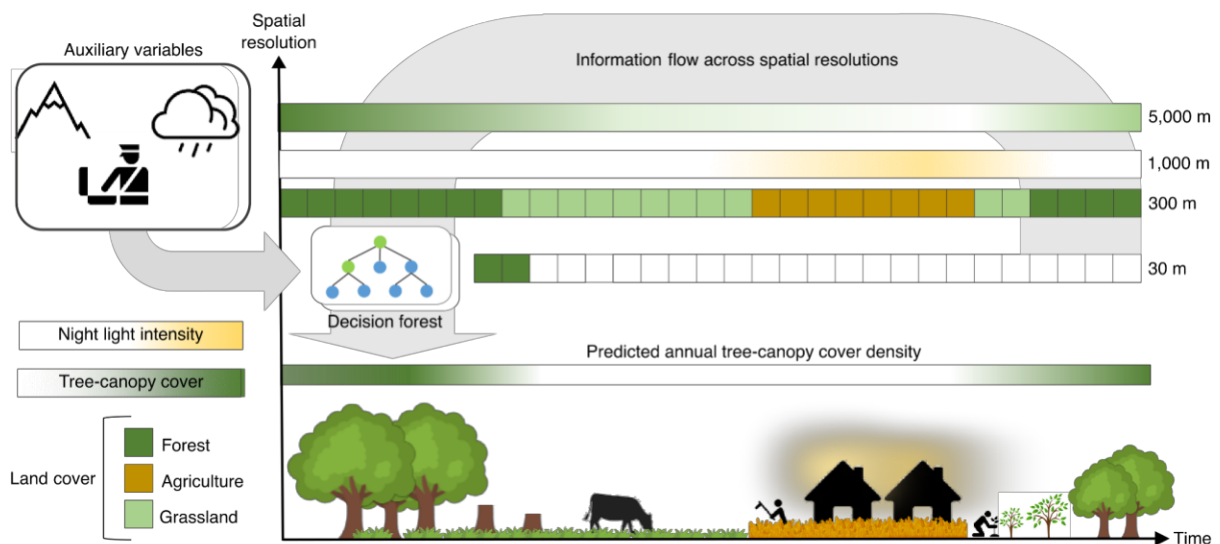
98 We mapped global tree-canopy cover using a machine-learning framework (**Fig. 1**) that exploits the  
99 fact that trajectories of forest persistence and change are commonly path-dependent (Montibeller et al.,  
100 2020; Taubert et al., 2018). This allows us to predict tree-canopy cover in the 1990s when high-  
101 resolution data is lacking, and consistently thereafter.

102 Our modelling framework uses higher-resolution data on tree-canopy cover as both a source of  
103 samples and as a predictor. These data inform on the outcome of long-term tree-canopy cover changes  
104 in a future reference year (T), and are obtained from a dataset available only since the year 2000 (see  
105 **Fine-scale tree-canopy cover**). For each mapping year (t), which span from 1992 to 2018, we instead  
106 used coarser-resolution data on environmental drivers of tree-canopy cover change (**Table 1 of SI**).  
107 This includes data on land-cover (see **Land cover**), vegetation continuous fields (see **Coarse-**

108 **resolution vegetation dynamics**), human settlements and political boundaries (see **Human factors**),  
 109 and climate and topography (see **Environmental drivers of tree growth**).

110 Furthermore, our modelling framework explicitly accounts for change dynamics. For this, we  
 111 included data on the direction and magnitude of changes between years  $t$  and  $T$  (see **Coarse-resolution**  
 112 **vegetation dynamics**), as well as data on fine-scale tree-canopy cover gains and losses by between  
 113 2000 and 2019 (the period for which our reference data is available, see **Fine-scale tree-canopy cover**).  
 114 In addition, we included a dummy variable on the number of years between the years  $t$  and  $T$  to control  
 115 for fixed temporal effects and temporal path-dependencies.

116 Finally, our modelling framework includes data on the (dis)agreement between inputs that measure  
 117 (or are sensitive to) tree-canopy cover (see **Cross-scale perceptions of tree-canopy cover**). This to  
 118 synthesise perceptions of tree-canopy cover, allowing us to exploit the strengths and weaknesses of  
 119 individual data products.



120  
 121 **Figure 1 – Conceptual overview of our modelling framework.** The model aims to reconstruct historical dynamics of  
 122 continuous tree-canopy cover at 300-m resolution. To capture signals of the true change trajectories (e.g., the iconized  
 123 example), the model integrates high-resolution, but temporally incomplete data on tree-canopy cover with coarser-resolution  
 124 data on categorical land cover, continuous canopy cover dynamics, night-lights, and auxiliary data (topography, climate,  
 125 political boundaries). The model learns about interactions among variables across spatial grains, spatiotemporally varying  
 126 strengths and weaknesses of coarser-grain products, and temporal path-dependencies of fine-grain change processes captured  
 127 only in later years, and uses the learned information for its annual predictions.

## 128 Modelling framework

129 We modelled tree-canopy cover with Random Forest Regression (RFReg, Breiman, 2001) as  
130 implemented in Python's Scikit-learn module (Pedregosa et al., 2011). The algorithm constructs  
131 multiple decision trees (i.e., a "forest"), each of which with its own decision path to the tree-canopy  
132 cover values reported in the training data. When applying the RFReg model to each pixel in a regular  
133 spatial grid, and in each prediction year, the different decision trees estimate a tree-canopy cover value,  
134 and the estimates of all trees are averaged into a final prediction.

135 We trained our model with 180 trees (hereafter referred to as "forest size"). To determine this  
136 number, for each of 10 runs, we extracted a random subset of 10,000 samples to test how forest sizes  
137 between 40 and 500 trees, in intervals of 20, led to gains in model performance (**Fig. 1a of SI**). At each  
138 iteration, we built a RFReg model and recorded its Out-Of-Bag (OOB) score, which corresponds to the  
139 coefficient of determination ( $r^2$ ) between the in-bag and out-of-bag samples. After all iterations, we  
140 calculated the mean OOB value at each forest size, and used Python's Kneed module (Satopaa et al.,  
141 2011) to find the start of a plateau in performances where increases in forest sizes did not improve  
142 accuracies noticeably (i.e., the 'knee' of the distribution).

## 143 Prediction uncertainties

144 We estimated the 95% Confidence Intervals (CI) across the predictions of all decision trees for each  
145 pixel and year combination. This was calculated as  $1.96 * SE$ , where the Standard Error (SE) is calculated  
146 as the standard deviation of the predictions, divided by the square root of the forest size. We used the  
147 CI to derive upper and lower confidence bounds around the average estimate for each pixel and year.  
148 We used these bounds to associate our subsequent assessments with uncertainty bounds (see **Change**  
149 **analysis** and **Comparison with national statistics**).

## 150 Sampling

151 We informed our model with samples from a high-resolution dataset that offers the most reliable  
152 global reference on tree-canopy cover (see **Fine-scale tree-canopy cover**). We sampled on a country-  
153 by-country basis, thereby accounting for likely differences in forest-management regimes across  
154 political boundaries (Herrera et al., 2019). Here, we treated any non-contiguous land masses of a given

155 country separately, resulting in 114,020 sampling units. For example, we treated the conterminous USA  
156 as a single unit, and addressed Alaska and each Hawaiian island separately, therefore ensuring sampling  
157 across unique environmental/management contexts.

158 For each country (referred hereafter as a ‘region of interest’, or roi), we stratified our sampling based  
159 on land-cover. Doing so ensured the collection of samples from every category, tackling the inability  
160 of RFReg to predict out-of-sample. In a given roi, the number of samples per class was estimated as  
161  $n_{roi} \sum(p_j = c)/n_c$ , where  $n_c$  is the total number of pixels of the target class,  $p_j$  is the target pixel at position  
162  $j$ , and  $n_{roi}$  is the total number of samples to be extracted within the roi. The value  $n_{roi}$  was estimated as  
163 being at least one sample per 150x150 pixels (i.e., 50-km<sup>2</sup> at the equator). This value was determined  
164 empirically as a compromise between model accuracy and computational intensity.

165 In a given roi, for each class  $c$ , we sampled half of the estimated  $n_c$  randomly over pixels where our  
166 reference data did not detect changes in tree-canopy cover between 2000 and 2018. We collected the  
167 remaining half over pixels with changes, where we applied a skewness-adjusted random-sampling  
168 approach. Here, we calculated the skewness of the change distribution. Then, we used the inverse of the  
169 shape parameter to build a skewed sampling distribution to guide the random selection of samples. We  
170 calculated the skewed distribution with NumPy’s skewnorm function (Harris et al., 2020).

171 This skewness-adjusted sampling tackles the potential rarity of changes at the upper and lower edges  
172 of the change distribution. For example, in a roi where smaller changes are dominant, the distribution  
173 will become left-skewed. Here, non-probabilistic random sampling would privilege smaller changes,  
174 creating an imbalanced sample set. This would negatively impact RFReg by restricting its predictive  
175 capabilities to a narrow range of outcomes.

176 Note that the sampling based on tree-canopy cover changes disregards pixels experiencing gains.  
177 This is because gains are mapped in the reference dataset as a mask of occurrences between 2000 and  
178 2012. Therefore, it is not possible to infer the true year of change. Nonetheless, our goal is to map year-  
179 specific values of tree-canopy cover and not change percentages. This filtering merely controls for  
180 sample quality, and does not limit our ability to predict gains thereafter.

181 We collected 78,135 sampling locations (**Fig. 1b of SI**). At each location, we sampled tree-canopy  
182 cover on an annual basis from 2000 to 2008, resulting in a training set of 625,080 samples, and  
183 accordingly extracted all predictor variables. We then used these data to train a model predicting annual  
184 tree-canopy cover densities (see **Overview of modelling strategy**).

185 Predictors

#### 186 *Fine-scale tree-canopy cover*

187 For each year for which we mapped tree-canopy cover, our modelling framework uses a later  
188 reference on that variable that informs on the outcome of long-term changes. Here, we constructed  
189 annual layers on tree-canopy cover, from 2000 to 2019, using the Global Forest Change dataset (GFC,  
190 version 1.8, Hansen et al., 2013). Specifically, we used 30-m data on tree-canopy cover mapped for  
191 2000, and extended it annually by disaggregating associated data on the year of tree-canopy cover  
192 losses. For each pixel and year combination, we preserved the tree-canopy cover value of year 2000  
193 until the loss year, at which point we updated the baseline value to 0% following the definition of a loss  
194 given by the authors. Then, we discarded pixels with < 10% tree-canopy cover due to reportedly high  
195 likelihoods of false detections of sparse tree-cover with Landsat imagery (Achard et al., 2014). Finally,  
196 we aggregated the annual layers via averaging to our target mapping resolution of 300-m.

197 In addition to data on tree-canopy cover changes, we accounted for changes in this variable at  
198 different spatial scales to account for spatial contagions (Ferrer Velasco et al., 2020; Rosa et al., 2013).  
199 Here, we derived two predictors. Firstly, we calculated the density of 30-m pixels nested within a 300-  
200 m pixel that experienced a loss at any point during the 2000-2019 period. Secondly, we calculated the  
201 average temporal distance of loss years relative to 2000. We produced both variables at our mapping  
202 resolution of 300-m and, additionally, at a 5-km resolution with respect to the coarsest data used in this  
203 study (see **Coarse-resolution vegetation dynamics**).

#### 204 *Coarse-resolution vegetation dynamics*

205 We used AVHRR-based Vegetation Continuous Fields data (AVHRR-VCF, Song et al., 2018),  
206 which map per-pixel percentages of tree cover (taller than 5 m), short vegetation, and non-vegetated



207 surfaces between 1982 and 2016. Due to its relatively high global and temporal consistency, these data  
208 allow us to describe regional vegetation dynamics, and provide our modelling framework with  
209 information on temporal path-dependencies.

210 The AVHRR-VCF data end in 2016, and we could not extend them with the original methods due  
211 to the degradation of AVHRR sensors (Dech et al., 2021). Instead, we extended these data with that of  
212 the Copernicus land cover dataset (CLC, Buchhorn et al., 2020), which maps per-pixel proportions of  
213 9 land-cover classes between 2015 and 2019 at a 100-m resolution. To match these data to the lower  
214 thematic and spatial resolution of the AVHRR-VCF data, we aggregated the CLC data by summing  
215 proportions of land-cover classes into AVHRR-VCF categories and averaged them to a 5-km resolution.  
216 A description of the modelling framework and the corresponding outputs can be found in a dedicated  
217 repository (<https://doi.org/10.5281/zenodo.8217455>).

218 We used the extended AVHRR-VCF time-series to derive predictor variables on percent changes.  
219 For each pixel and year, we estimated changes in relation to the nearest future year with available GFC  
220 data. For example, changes from 1992 onwards were calculated in comparison to the year 2000. In turn,  
221 changes from the year 2000 were estimated in comparison to data from the year 2001.

## 222 *Land cover*

223 We integrated annual data on the dominant land cover in 300-m-resolution pixels (distinguishing 38  
224 land-cover/use classes following the UN Land Cover Classification System; **Table 2 of SI**), as mapped  
225 by the Climate Change Initiative Land Cover dataset (CCILC, Santoro et al., 2017).

226 Categorical land-cover data aids predictions of tree-canopy cover in several ways. First, the class  
227 distinctions indicate whether or not tree-cover is perceived as the dominant land cover. Second, even if  
228 land-cover classifications face persistent regional issues, they can still carry relevant signals that help a  
229 model make more accurate predictions. This is because the spatial configurations of (correctly or  
230 falsely) perceived tree-cover classes, and also those of other land-cover classes that are easily confused  
231 with tree-cover (e.g., shrubland), often follow true gradients of either tree-canopy cover or of canopy-  
232 cover-affecting environmental conditions or management regimes.

233 *Cross-scale perceptions of tree-canopy cover*

234 We mapped tree-canopy cover at a 300-m resolution, but we lack such fine scale data throughout  
235 our mapping period. To tackle this issue, we included several other datasets that either directly map or  
236 are sensitive to this variable, and that provide consistent, annual information. Given these data result  
237 from different modelling strategies, they show disagreements. Yet, these disagreements provide varying  
238 perceptions that help complement the weaknesses of individual products.

239 We derived predictors on the differences in mapped tree-canopy cover densities between the CCILC,  
240 AVHRR-VCF, and GFC datasets. We computed the percent differences of each combination of the  
241 aforementioned datasets at a 5-km resolution, with respect for the lowest resolution dataset (i.e.,  
242 AVHRR-VCF). For the comparison between the AVRR-VCF and CCILC datasets, we estimated annual  
243 differences from 1992 and 2018. For the comparison of both AVHRR-VCF and CCILC with the GFC,  
244 we estimated annual differences between 2000 and 2019. Because the CCILC dataset is categorical, we  
245 translated land-cover classes into tree-canopy cover Following similar past experiments (Song et al.,  
246 2014), we estimated that classes occupy the central percentage value of the range of possible values  
247 indicated in their respective labels (**Table 2 of SI**).

248 *Environmental drivers of tree growth*

249 We integrated predictor variables to account for the driving influence of topography and climate on  
250 global tree abundances (Crowther et al., 2015). Firstly, elevation drives spatial variation in forest  
251 structure (Mazón et al., 2020), whereas slope influences both natural (e.g., landslides, Maraun et al.,  
252 2022) and human drivers of tree-canopy cover changes (e.g., feasibility of industrial logging, FAO,  
253 1998). Therefore, we obtained layers on elevation and slope from a 90-m-resolution digital elevation  
254 model (Yamazaki et al., 2017), and averaged them to our 300-m mapping resolution. Secondly, tree-  
255 growth is strongly determined by water availability and its seasonality (X. Xu et al., 2018). Therefore,  
256 we derived data on the annual total precipitation (P) and Aridity Index (AI, Barrow, 1992) based on  
257 monthly precipitation (P) and potential evapotranspiration (PET) data from the 1-km CHELSA dataset  
258 (version 2.0, Karger et al., 2017). Whereas P indicates the presence and abundance of available water,  
259 the AI, estimated for each year  $t$  as  $\sum_{t=1}^n P_t / \sum_{t=1}^n PET_t$ , serves as a proxy for water seasonality and

260 scarcity. When the AI is 0, this indicates the absence of water, or its rapid evaporation due to high  
261 temperatures. In turn, when the AI is high (generally above 1), precipitation is higher than PET,  
262 indicating moist conditions. In addition, when a higher P contrasts with a low AI, precipitation may  
263 contrast with seasonally high temperatures, such as in drylands.

#### 264 *Human factors*

265 The presence and intensity of human activities drives forest changes (IPCC, 2022). To account for  
266 these factors, we included annual 1-km-resolution data on night-time lights (NTL; Li et al., 2020).  
267 Additionally, we accounted for the fact that forest-management regimes and related human drivers often  
268 vary between countries (Herrera et al., 2019) by assigning unique country identifiers to each pixel  
269 (based on the Global Administrative boundary dataset; GADM, UCB, 2012). This served as a fixed  
270 spatial effect in our model.

#### 271 Model quality assessment

272 We tested our model using spatial block cross-validation (Roberts et al., 2017). We iterated through  
273 each level-1 administrative unit in the GADM dataset (i.e., district- or state-level), withholding the  
274 corresponding samples to validate a model trained with the remainder. We then recorded the predictions  
275 for the withheld sample locations to subsequently evaluate model performances. Through this process,  
276 we evaluated our ability to predict tree-canopy cover in pixels beyond an area occupied by reference  
277 data. We chose level-1 administrative units instead of countries for the block validation, because we  
278 included a country-level fixed effect, and decision-tree algorithms cannot predict out-of-sample.

279 We compared the pooled predictions for all administrative units against the respective reference  
280 values to derive the coefficient of determination ( $r^2$ ), the Root Mean Square Error (RMSE), and the F1-  
281 score (Taha and Hanbury, 2015) The  $r^2$  and the RMSE measure the quality of fit. The former shows  
282 how well we capture spatial gradients in the reference data, whereas the latter quantifies the magnitude  
283 of prediction errors. In turn, the F1-score evaluates the predicted presences and absences of tree-canopy  
284 cover, controlling for high commission errors hidden by low prediction errors.

285 We mapped tree-canopy cover by exploring temporal path-dependencies in tree cover change. This  
286 enabled us to reconstruct historical changes since 1992, although the earliest instance of our reference

287 data on tree-canopy cover is available for the year 2000. To test our model's reconstruction ability, we  
288 evaluated our quality metrics separately for pixels that, according to the GFC dataset, experienced no  
289 (0 ha), small ( $>0$  -  $<1$  ha), medium (1 -  $<4$  ha), or large ( $\geq 4$  ha) losses. We focused on losses because  
290 gains as depicted in the GFC dataset are estimated as a mask, and provided solely for the 2000-2012  
291 period. Despite this, it is important to reiterate that we do not predict changes in tree-canopy cover, but  
292 rather year-specific percentages, changes in which are mostly driven by losses.

293 Additionally, we tested our model's ability to predict tree-canopy cover with different numbers of  
294 years separating the prediction year (t) and the future reference year (T). We anticipated that larger  
295 temporal gaps would be easier to predict given our model exploits long-term change processes. To this  
296 end, we tested the performance of our model for each year for the 2000-2007 period (8 years). This  
297 emulates backward-predictions for the 1990s, when high-resolution reference data on forest dynamics  
298 is missing. Temporal gaps between mapping and reference years vary (from 1 year for 1999 predictions  
299 with a 2000 reference to 8 years for 1992 predictions with a 2000 reference).

300 We ran our quality assessments for each region and continent distinguished by the United Nations  
301 Food and Agriculture Organisation (FAO) when compiling Forest Resource Assessments (FRA's), as  
302 to inform on the potential value of our data product in supporting these assessments.

### 303 Comparison with national statistics

304 Forest Resource Assessments (FRAs) provide the most authoritative national-scale information on  
305 forest extents across multiple decades. Because our data on tree-canopy cover dynamics may be viewed  
306 as complementary to the assessment of forest extents, we compared them to FRAs to assess the  
307 plausibility of the mapped national, regional, and global patterns.

308 Tree-canopy cover is conceptually distinct from forest cover as reported in FRAs, primarily because  
309 the latter not only accounts for areas immediately covered by canopies, but also for open areas in  
310 between trees. In addition, FRAs limit forest accountancy to tree stands  $\geq 5$ m tall that cover  $\geq 0.5$ -1.0  
311 ha with a tree-canopy cover of  $\geq 10\%$ , and excludes tree-covered lands under non-forest uses (e.g.,  
312 agricultural tree plantations, corresponding to  $\sim 2.3\%$  of the global tree cover in 2015; FAO, 2016). In  
313 comparison, we based our model on data that aims to map tree occurrences with a height of  $\geq 5$ m, but

314 at a higher spatial resolution (0.09 ha). This is likely to make our modelled outputs more sensitive to  
315 the presence of trees compared to FRAs.

316 These conceptual differences prevent a direct comparison of mapped areas. Instead, we correlated  
317 national forest areas from FRAs with those estimated from the combined area of pixels for which we  
318 mapped some tree-canopy cover (describe tree covered landscapes, hereafter “treescapes”). The  
319 comparison between these variables follows the premise that, despite conceptual differences, treescape  
320 and forest extents should show similar cross-country variations.

321 We made comparisons against national forest-area statistics available for 1990, 2000, 2010, 2015,  
322 and 2020, and estimated their correlation ( $r^2$ ) against national treescapes calculated from our product  
323 for the nearest mapping years (i.e., 1992, 2000, 2010, 2015, and 2018). In addition, we compared the  
324 FRA of 2000 to treescapes derived from GFC’s data. This assessment informed on our ability to  
325 generate regional predictions of a comparable quality to the GFC, which was derived with optical  
326 remote sensing data, and is therefore informed by visual evidence of tree occurrences.

327 We correlated national treescape areas with national accountings of forest areas using weighted least-  
328 squares regression. The weights are country-specific and based on the reliability of FRA data based on  
329 national remote-sensing and forest-inventory capabilities (Nesha et al., 2021). Although monitoring  
330 capabilities are graded on individual FRA years, we used a static weight because grades were only  
331 available since 2005. For each country and FRA year, we calculated the minimum value across FRA  
332 years, preventing countries with evolving capabilities to bias earlier FRA comparisons.

333 We used our correlation analysis across multiple FRA years to evaluate the stability of our  
334 predictions. Yet, not every country contributed to FRAs in every assessment year, and the data for some  
335 countries may be unreliable due to poor monitoring capabilities in some years. Therefore, we excluded  
336 two countries without data in every target year, 93 with weights of 0 (i.e., very poor-quality reports),  
337 and eleven where the GFC reported no forest cover despite contrary indications in the FRAs due to a  
338 lack of usable Landsat acquisitions. Overall, we included 132 out of 238 territories (28/48 in Asia, 33/56  
339 in the Americas, 40/59 in Africa, 23/50 in Europe, 8/24 in Oceania).

340 From the remaining data pool, treescapes are unequally distributed across countries. In particular,  
341 countries with advanced monitoring capabilities and large forest extents (e.g., Brazil) received higher

342 weights, and thus had a stronger influence over the direction of the regression line. This results in  
343 relatively low residuals that inflate correlations. To tackle this issue, we log-transformed both forest-  
344 areas and treescape extents prior to the correlation analysis to give a ‘bigger voice’ to countries with  
345 relatively small forest extents.

#### 346 Change analysis

347 Tree-canopy cover influences multiple goals and targets of the 2030 Agenda for Sustainable  
348 Development (Estoque et al., 2021). Consequently, knowledge on historical patterns and directions of  
349 change is essential to refine post-2020 political action. Following this premise, we explored our data  
350 product to provide insights into long-term patterns of change in tree-canopy cover.

351 To achieve this, we first applied the Mann-Kendall test to the 1992-2018 time-series of each pixel  
352 to categorise significant negative and positive slopes as “losses” or “gains”, respectively. We defined a  
353 trend as “significant” if the p-value reported by the Mann-Kendall test was  $\leq 0.005$ , following recent  
354 recommendations for such assessments (Ioannidis, 2018). Then, for each trend category, we derived  
355 statistics within each combination of continent and forested biomes distinguished by the FAO (i.e.,  
356 ‘boreal’, ‘temperate’, ‘tropical dry’, and ‘tropical moist’).

357 For each change category, biome, and continent, (**Fig. 4a-d**), we summed the area of the  
358 corresponding pixels to inform on the extent of the affected area (hereafter referred to as ‘treescape  
359 extent’, **Fig. 4b-c**). Then, for the pixels accounted for in each change category, we calculated the  
360 difference in tree-canopy cover between the first and last years of our time-series, informing on the  
361 change in the openness of the canopies (**Fig. 4e-f**). Whereas the first metric informs on the spatial  
362 dispersal of change dynamics, the second informs on the magnitude of those changes.

## 363 RESULTS

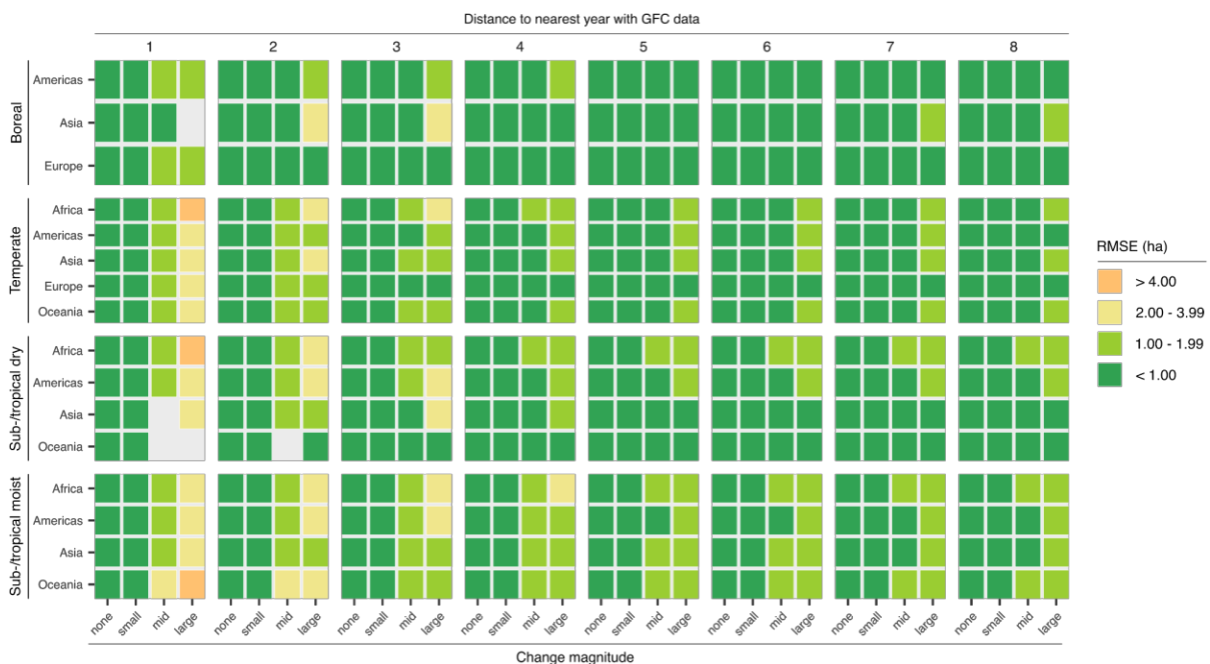
### 364 Quality assessment

365 Focusing on stable pixels (i.e., those not experiencing losses in tree-canopy cover), performances  
366 were higher across temporal gap sizes between the reference and target years. Within this group,  $r^2$ ,

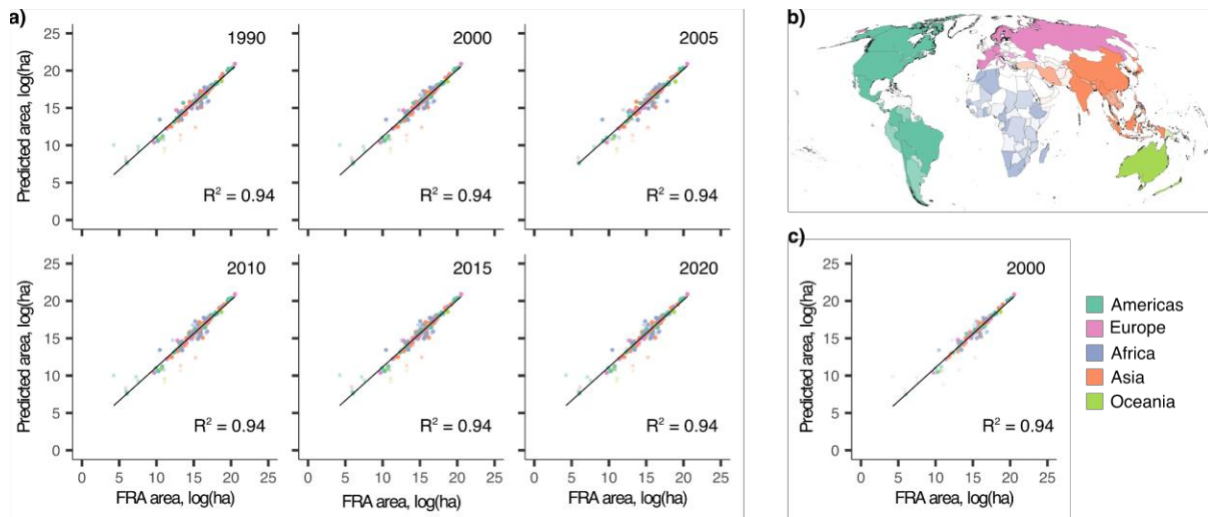
367 RMSE, and F1-score values were, respectively, 0.99 ( $< \pm 0.01$ ), 0.06 ha ( $\pm 0.01$ ) and 0.98 ( $\pm 0.05$ ) for  
 368 gap sizes of 8 years, and 0.98 ( $< \pm 0.01$ ), 0.16 ( $\pm 0.03$ ), and 0.95 ( $< \pm 0.01$ ) for gap sizes of 1 year.

369 For changing pixels, correlations between predicted and reference values varied between 0.83  
 370 ( $\pm 0.01$ ) for temporal gaps of 1 year, and 0.93 ( $< \pm 0.01$ ) for gaps of 8 years. When disaggregating these  
 371 values per change-size class, we recorded a correlation of 0.98 ( $\pm 0.01$ ) for small to mid-sized changes  
 372 with both 1 and 8 gap years, which for large changes dropped to 0.89 ( $\pm 0.01$ ) for a gap size of 8 years,  
 373 and 0.73 ( $\pm 0.01$ ) for a gap size of 1 year. Similarly, RMSE values across small and mid-sized changes  
 374 were lower for gap sizes of 8 years (0.40 ha,  $\pm 0.01$ ) compared to gap sizes of 1 year (0.59,  $\pm 0.01$ ), and  
 375 differences increased for large changes (3.26 ha [ $\pm 0.31$ ] for 1 year, and 1.32 ha [ $\pm 0.15$ ] for 8 years). In  
 376 contrast, F1-scores were persistent across gap sizes and change magnitudes (0.99 [ $\pm 0.01$ ]), suggesting  
 377 our ability to correctly perceive presences and absences of tree cover.

378 When comparing our predicted treescape extents against national forest extents reported by FRAs,  
 379 we recorded an  $r^2$  of 0.94 in every assessment year ( $\pm 0.00$ , **Fig. 3a-b**, see **Fig. 4 of SI** for plotted  
 380 uncertainties). Moreover, the GTCCC-FRA correlation for the year 2000 was as strong as the GFC-  
 381 FRA correlation, both globally (**Fig. 3c**) and regionally (**Table 3 of SI**).



382  
 383 **Figure 2 – Model quality.** RMSE of predicted tree-canopy cover for different time intervals between the focal mapping year  
 384 and the nearest year with reference data (shown in the header of each block), and for different continents, biomes, and tree-  
 385 canopy cover change sizes. The uncertainties surrounding the values in this plot are displayed by **Fig. 3 of SI**.



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### 393 Changes in treescape extents

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**Figure 3 – Alignment of global patterns of treescape extents with FRA data on forest extents.** a) Cross-country correlations of national extents of treescapes derived from the GTCCC with forest extents reported in FRA statistics for each reporting year (see **Fig. 4 of SI** for plots on upper/lower confidence bounds). The colour of points distinguishes geographical regions (**b**); countries shown in white were excluded from this comparison due to filtering. Correlations were weighted by the quality of national reporting capabilities, which are indicated in **b**) by varying levels of transparency. Solid colours indicate high quality data. c) As in **a**), but comparing FRA data for 2000 to the GFC.

We estimated global net losses in treescape extents of 8.8 million ha (Mha,  $\pm 0.4$ ) based on pixels with significant long-term trends (**Fig. 4a-c, Table 1**). The largest net losses were visible across the tropics (-11.5 Mha,  $\pm 0.6$ ), 60.0% and 40.0% of which occurred in the tropical moist and tropical dry biomes, respectively (**Table 1**). Although the tropical moist biome experienced larger global losses in treescape extents, tropical dry losses were larger in Africa (-1.0 Mha [ $\pm 0.0$ ] vs -0.6 Mha [ $\pm 0.0$ ]).

At higher latitudes, treescape gains within the temperate biome become more common than losses, minor negative balances in North America (-0.4 Mha,  $\pm 0.0$ ). Globally, temperate treescapes increase by 3.6 Mha ( $\pm 0.1$ ), 90.5% of which occurred in Europe (3.2 Mha,  $\pm 0.0$ ). Gains in Europe correspond to +1.2% relative to 1992 ( $\pm 0.0$ ), second only to Australia (+2.3%,  $\pm 0.0$ ).

Despite these gains, we again observed net losses in treescape extents at higher latitudes. The boreal biome experienced global net losses (-0.9 Mha,  $\pm 0.5$ ), with nearly twice as much area lost in Asia as was gained in Europe (-1.0 Mha [ $\pm 0.3$ ] vs. +0.6 Mha [ $\pm 0.0$ ]).



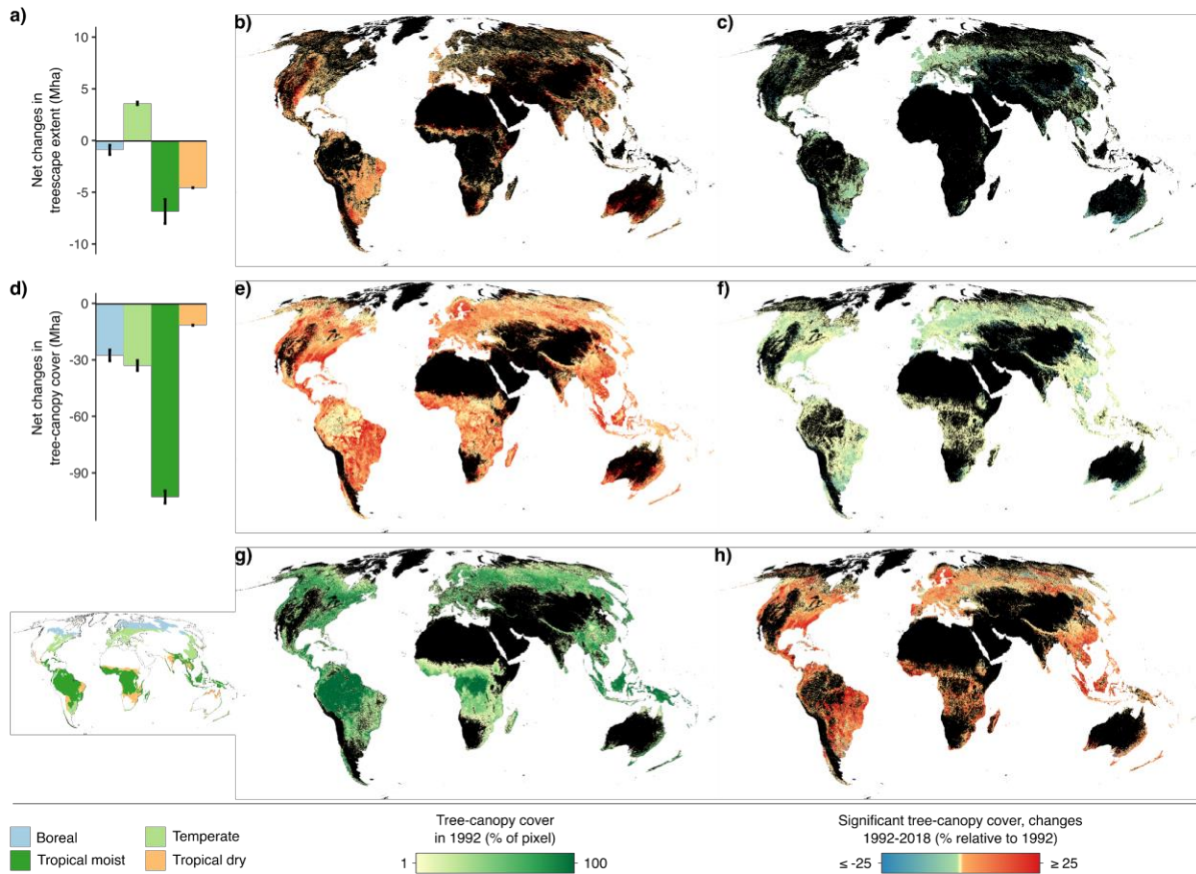
406 Changes in tree-canopy cover

407 Whereas our analysis of changes in treescape extents revealed regional contrasts, we found  
408 systematic losses in tree-canopy cover across all biomes and continents (-175.1 Mha [ $\pm 1.0$ ], **Fig. 4d-f**,  
409 **Table 1**). The bulk of these losses (58.7%) occurred in the tropical moist biome (-102.9 Mha,  $\pm 2.5$ ),  
410 and was 27 times greater than the simultaneous loss in the extent of treescapes.

411 Losses within the tropical moist biome predominantly occurred in South America (43.7% of net  
412 losses, -45.0 Mha [ $\pm 0.9$ ]). By comparison, the tree-canopy cover over the tropical dry biome decreased  
413 by -11.6 Mha ( $\pm 0.0$ ), nearly 9 times smaller than the losses in the tropical moist biome. However, this  
414 change is larger if viewed as a net percentage loss relative to the 1992 baseline (-7.5% [ $\pm 0.0$ ] vs -10.4%  
415 [ $\pm 0.1$ ] for tropical moist and tropical dry, respectively).

416 In the temperate biome, treescape and tree-canopy cover dynamics showed contrasting directions of  
417 change. The net losses in tree-canopy cover (-33.0 Mha,  $\pm 1.0$ ) were nine times larger than the areal net  
418 gains in treescape extents (corresponding to a loss of -6.3% [ $\pm 1.0$ ] relative to the 1992 baseline).

419 Even Europe's gains in temperate treescape extents were superseded nearly 1.7-fold by losses in  
420 tree-canopy cover (**Table 1**). In fact, net losses consistently dwarfed absolute net changes in treescape  
421 extents in all biomes and continents, typically by an order of magnitude or more (e.g., corresponding to  
422 a 93 times larger area in the tropical moist biome in Asia).



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**Figure 4 – Global changes in tree-canopy cover and treescape extents.** Global net changes of treescape extents (a) and tree-canopy cover (d) for different biomes over the 1992-2018 period, based on statistically significant trends (gains or losses) detected at a 300-m resolution (biomes indicated by bar colours, and mapped below), and aggregated for mapping to a 10-km resolution. b/c) Global percent losses (b) and gains (c) in treescape extents, relative to the 1992 baseline (shown in g). e/f) Global percent losses (e) and gains (f) in tree-canopy cover (same baseline). h) Net changes (i.e., losses or gains) in tree-canopy cover evaluated at the 10-km mapping resolution, highlighting a global dominance of net losses.

**Table 1 – Global change in treescape extents and tree-canopy cover.** Net 1992-2018 changes in treescape extents and tree-canopy across biomes and regions. Expressed in millions of ha (Mha) and as percentages (%) of the 1992 baseline.

Biome	Continent	<i>Δ Treescape extent (1992-2018)</i>		<i>Δ Tree-canopy cover (1992-2018)</i>	
		<i>Δ (Mha)</i>	<i>Δ (% of 1992 extent)</i>	<i>Δ (Mha)</i>	<i>Δ (% of 1992 cover)</i>
All	World	<b>-8.8 (±0.4)</b>	<b>-8.8 (±0.4)</b>	<b>-8.8 (±0.4)</b>	<b>-7.3 (±0.0)</b>
	Africa	-1.6 (±0.0)	-0.2 (±0.0)	-29.1 (±0.4)	-6.8 (±0.0)
	Asia	-1.6 (±0.4)	-0.2 (±0.0)	-43.4 (±1.4)	-8.1 (±0.0)
	Australia	0.5 (±0.0)	0.9 (±0.0)	-1.1 (±0.0)	-6.9 (±0.0)
	Europe	3.8 (±0.1)	0.9 (±0.0)	-15.2 (±0.4)	-7.2 (±0.0)
	North America	-0.9 (±0.3)	-0.2 (±0.1)	-31.6 (±0.8)	-10.4 (±0.0)
	Oceania	< 0.1	0.1 (±0.0)	-0.7 (±0.0)	-4.2 (±0.0)

	South America	-8.9 ( $\pm 1.0$ )	-0.8 ( $\pm 0.1$ )	-54.0 ( $\pm 1.1$ )	-7.2 ( $\pm 0.0$ )
<i>Boreal</i>	<b>World</b>	<b>-0.9 (<math>\pm 0.5</math>)</b>	<b>-0.1 (<math>\pm 0.1</math>)</b>	<b>-27.6 (<math>\pm 0.7</math>)</b>	<b>-7.5 (<math>\pm 0.0</math>)</b>
	Asia	-1.0 ( $\pm 0.3$ )	-0.3 ( $\pm 0.1$ )	-10.6 ( $\pm 0.2$ )	-6.8 ( $\pm 0.1$ )
	Europe	0.6 ( $\pm 0.0$ )	0.3 ( $\pm 0.0$ )	-7.3 ( $\pm 0.2$ )	-7.2 ( $\pm 0.0$ )
	North America	-0.5 ( $\pm 0.1$ )	-0.3 ( $\pm 0.1$ )	-9.7 ( $\pm 0.2$ )	-8.9 ( $\pm 0.0$ )
<i>Temperate</i>	<b>World</b>	<b>3.6 (<math>\pm 0.1</math>)</b>	<b>0.4 (<math>\pm 0.0</math>)</b>	<b>-33.0 (<math>\pm 1.0</math>)</b>	<b>-8.1 (<math>\pm 0.0</math>)</b>
	Africa	0.0 ( $\pm 0.0$ )	0.2 ( $\pm 0.0$ )	-0.4 ( $\pm 0.0$ )	-10.3 ( $\pm 0.1$ )
	Asia	0.1 ( $\pm 0.0$ )	0.0 ( $\pm 0.0$ )	-6.1 ( $\pm 0.1$ )	-5.5 ( $\pm 0.0$ )
	Australia	0.5 ( $\pm 0.0$ )	2.3 ( $\pm 0.0$ )	-1.0 ( $\pm 0.0$ )	-9.9 ( $\pm 0.1$ )
	Europe	3.2 ( $\pm 0.0$ )	1.2 ( $\pm 0.0$ )	-7.9 ( $\pm 0.2$ )	-7.2 ( $\pm 0.0$ )
	North America	-0.4 ( $\pm 0.1$ )	-0.2 ( $\pm 0.0$ )	-15.1 ( $\pm 0.5$ )	-11.0 ( $\pm 0.0$ )
	Oceania	< 0.1	0.2 ( $\pm 0.1$ )	-0.3 ( $\pm 0.0$ )	-5.0 ( $\pm 0.0$ )
	South America	0.1 ( $\pm 0.0$ )	0.1 ( $\pm 0.0$ )	-2.2 ( $\pm 0.1$ )	-7.9 ( $\pm 0.1$ )
<i>Tropical dry</i>	<b>World</b>	<b>-4.6 (<math>\pm 0.0</math>)</b>	<b>-1.0 (<math>\pm 0.0</math>)</b>	<b>-11.6 (<math>\pm 0.0</math>)</b>	<b>-10.4 (<math>\pm 0.1</math>)</b>
	Africa	-1.0 ( $\pm 0.0$ )	-0.4 ( $\pm 0.0$ )	-2.8 ( $\pm 0.0$ )	-6.7 ( $\pm 0.0$ )
	Asia	-0.4 ( $\pm 0.0$ )	-0.7 ( $\pm 0.0$ )	-1.5 ( $\pm 0.0$ )	-10.4 ( $\pm 0.1$ )
	Australia	- < 0.1	- < 0.1	- < 0.1	-1.0 ( $\pm 0.0$ )
	North America	-0.1 ( $\pm 0.0$ )	-0.5 ( $\pm 0.1$ )	-0.5 ( $\pm 0.0$ )	-6.3 ( $\pm 0.0$ )
	South America	-3.0 ( $\pm 0.0$ )	-2.6 ( $\pm 0.0$ )	-6.8 ( $\pm 0.0$ )	-15.6 ( $\pm 0.1$ )
<i>Tropical moist</i>	<b>World</b>	<b>-6.9 (<math>\pm 1.2</math>)</b>	<b>-0.3 (<math>\pm 0.1</math>)</b>	<b>-102.9 (<math>\pm 2.5</math>)</b>	<b>-7.5 (<math>\pm 0.0</math>)</b>
	Africa	-0.6 ( $\pm 0.0$ )	-0.1 ( $\pm 0.0$ )	-26.0 ( $\pm 0.5$ )	-6.8 ( $\pm 0.0$ )
	Asia	-0.3 ( $\pm 0.0$ )	-0.1 ( $\pm 0.0$ )	-25.2 ( $\pm 1.0$ )	-9.9 ( $\pm 0.0$ )
	Australia	< 0.1	0.2 ( $\pm 0.0$ )	- < 0.1	-2.6 ( $\pm 0.0$ )
	North America	0.1 ( $\pm 0.1$ )	0.1 ( $\pm 0.1$ )	-6.2 ( $\pm 0.1$ )	-13.1 ( $\pm 0.1$ )
	Oceania	- < 0.1	- < 0.1	-0.4 ( $\pm 0.0$ )	-3.6 ( $\pm 0.0$ )
	South America	-6.1 ( $\pm 1.0$ )	-0.7 ( $\pm 0.1$ )	-45.0 ( $\pm 0.9$ )	-6.7 ( $\pm 0.0$ )

## 432 DISCUSSION

433 We integrated remotely sensed information with complementary strengths to reconstruct global tree-  
434 canopy cover dynamics over nearly three decades. This offers new opportunities for assessing long-  
435 term changes in forests and other wooded ecosystems, analysing their impacts on biodiversity, training  
436 climate models, prioritising sites for ecosystem restoration, and various other applications supporting  
437 the 2030 Agenda for Sustainable Development. In turn, our analyses of the GTCCC offer new insights  
438 into the magnitude, direction, timing, and speed of global and regional tree-canopy cover changes,  
439 adding important nuance to previous forest change assessments.

440 Our analyses indicate that, since the early 1990s, wooded landscapes experienced systematic losses  
441 in tree-canopy cover at global, continental, and biome scales. These canopy-cover losses substantially  
442 exceed concurrent losses or gains of treescape extents. These results are in line with contrasts between  
443 trends in deforestation and forest disturbance reported from various regions (Aryal et al., 2021; Chen et  
444 al., 2021; Qin et al., 2021; Shapiro et al., 2021), and imply that common practices of estimating carbon-  
445 budget trends based on categorical land cover (e.g., Alkama and Cescatti, 2016; Baccini et al., 2017;  
446 Friedlingstein et al., 2022) may underestimate carbon losses.

447 Even in Europe, where forests are reportedly larger today than in the 20th century (Estoque et al.,  
448 2022), net gains in temperate and boreal treescape extents are dwarfed by concurrent losses in tree-  
449 canopy cover. Surprisingly, canopy-cover losses are noticeable across Eastern Europe (**Fig. 4e**), despite  
450 this being the European subregion with the largest treescape gains (**Fig. 4c**). These results are consistent  
451 with previous reports of increasing patch sizes and frequencies of forest disturbances (Senf and Seidl,  
452 2021), which have been linked to increases in wood harvests, and natural disturbances such as insects,  
453 windthrow and fires (McDowell et al., 2020; Seidl et al., 2014; Senf et al., 2018).

454 We note that our data neither indicate specific causes nor mechanisms of canopy-cover losses, which  
455 can affect carbon retention and other forest functions (Luyssaert et al., 2008; Thom and Seidl, 2016).  
456 Irrespective, the relative magnitude of tree-canopy cover losses relative to those in treescapes indicates  
457 that ignoring the former likely misses changes in those functions. Our results corroborate recent calls  
458 for better accountings of forest disturbances in European environmental policies (Maes et al., 2023),  
459 which currently aim solely for net increases in forest extents under the Green Deal, and which do not  
460 explicitly promote non-urban canopy cover gains (European Commission, 2022).

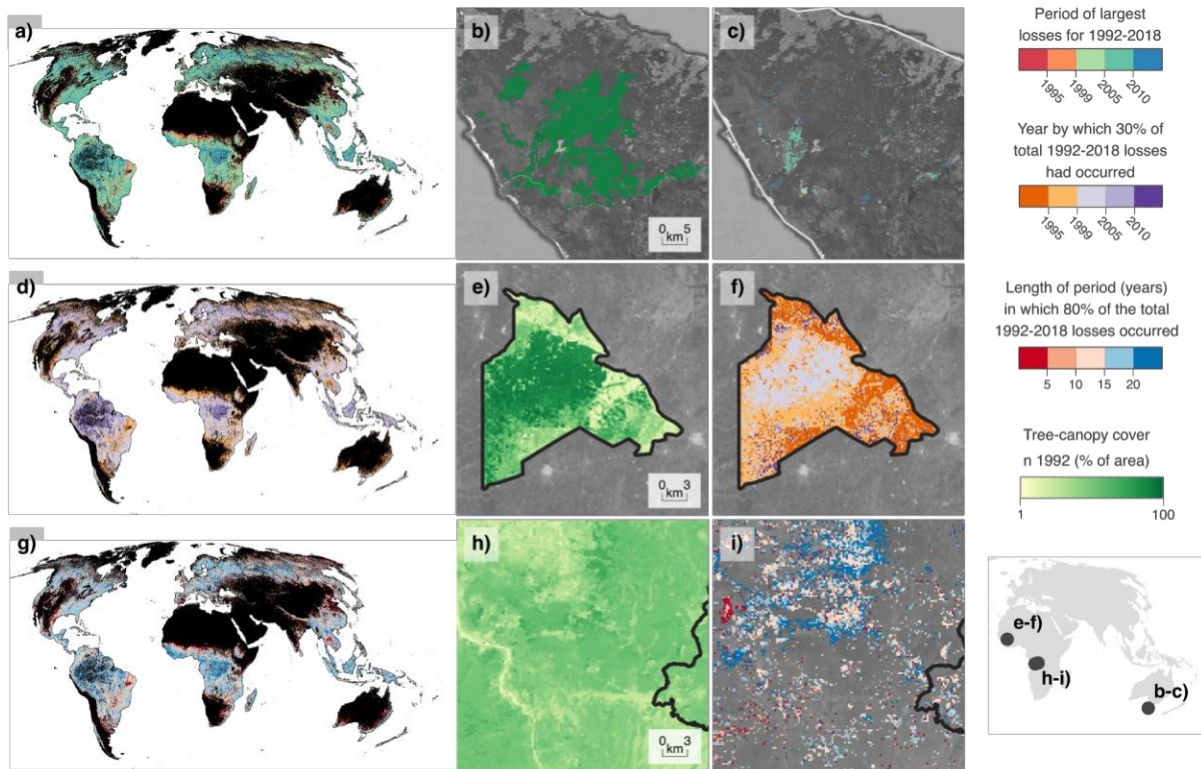
461 Similarly, our results indicate that even in countries with high national forest-monitoring capacities,  
462 the use of categorical data can grossly misrepresent structural changes in forest. For example,  
463 authoritative categorical data on canopy cover developed for Australia (Lucas et al., 2019) reveals net  
464 gains in temperate rainforests of +2.8% (-6.3%/+9.0%) between 1996 and 2016, whereas the Forest  
465 Practice Authority of Tasmania reported a loss of -3.7% during the same period (FPA, 2017). Our data  
466 integration framework enabled the detection of similar losses (-3.2%,  $\pm 0.2$ ). Literature relates these  
467 changes to growing human pressures in the form of logging and mining, and to the increasing risk of

468 fires driven by climate change (Mackey et al., 2017). The latter, in particular, lead to rainforest losses  
469 in recent years, such as seen over the Tarkine region of Tasmania (**Fig. 5b-c**).

470 Especially in tropical moist forest regions, where the historical Landsat archive has extensive  
471 temporal gaps during the 1990s (Wulder et al., 2016), integrating complementary data sources provided  
472 novel perceptions of the onset of major forest disturbances. For example, Côte d'Ivoire's Marahoué  
473 National Park (**Fig. 5e**) experienced extensive human settlement and the near-complete conversion of  
474 its forest area into cocoa plantations, both of which contributed to steep population declines in Western  
475 chimpanzees and other threatened primate species (Bitty et al., 2015; Campbell et al., 2008; Kühl et al.,  
476 2017). Landsat-based categorical time-series (Vancutsem et al., 2021) depict stable forest cover  
477 throughout the 1990s, with extensive forest degradation and deforestation thereafter. In contrast, by  
478 integrating data available during the 1990s only at coarse resolutions, we detected finer-resolution  
479 changes during this decade that make up for ~1/3 of 1992-2018 tree-canopy-cover losses (**Fig. 5f**, -  
480 8,085 ha vs. -17,006 ha), consistent with regional reports (**Fig. 5f**, -8,085 ha vs. -17,006 ha), consistent  
481 with regional reports (Hauhouot, 1992; Kouakou et al., 2018).

482 As we learned about these changes, we noted contrasts between tropical moist and dry biomes. In  
483 line with national statistics, we found the largest tree-canopy cover losses over the tropical moist biome  
484 (FAO, 2020) that threaten millions of species (Pillay et al., 2022) and historical carbon sinks (Achard  
485 et al., 2014). These risks earned the tropical moist biome disproportionate attention in literature  
486 compared to the tropical dry biome (Schröder et al., 2021). However, we found that the tropical dry  
487 biome experienced larger percent losses to its naturally sparser tree-canopy cover. This result supports  
488 calls for attention towards tropical dry woodlands (Schröder et al., 2021), on which millions of people  
489 depend as a source of fuelwood (Blackie et al., 2014).

490 Within tropical dry forests, change patterns and speeds are often heterogeneous (Buchadas et al.,  
491 2022). For example, in the surroundings of the Cangandala National Park in Angola (**Fig. 5h-i**, World  
492 Bank, 2019), the emergence of fast-paced changes associated with industrial farming (such as near the  
493 town of Cangandala, **Fig. 5i**) contrasts with slow-paced changes plausibly explained by household  
494 consumption of woodlands for fuelwood that is typical in the region (Michael Mills, 2017).



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**Figure 5. Temporal dynamics in tree-canopy cover.** Multi-temporal change metrics describing the timing and speed of tree-canopy cover changes over the 1992-2018 period, computed based on 300-m pixels experiencing significant losses (slope < 0; p-value < 0.0005). Panels **a/d/g** show global patterns of change dynamics (means per 3-km pixel); **c/f/i** show local examples, with **b/e/h** showing the initial (1992) tree-canopy cover for reference. **a/c/e** Period experiencing the largest losses in tree-canopy cover. **b** The Tarkine in north-eastern Tasmania is Australia’s largest old-growth temperate rainforest, known for its high integrity (note the continuous dense canopy cover in 1992; rainforest extent delineated by authoritative data (DPIPWE, 2020)). **c** Forest disturbances within the Tarkine, mainly after 2005. **d/f** Year by which 30% of the total 1992-2018 net tree-canopy cover losses had occurred. **e/f** Cote d’Ivoire’s Marahoué National Park was in 1992 still covered by tropical moist forest and savanna, which by the late 2000s had been nearly completely converted. The GTCCC perceives losses within Cote Ìvoire’s Marahoué National Park already in the 1990s (**e**, orange tones), with continuous losses into densely tree-covered portions of the park in the 2000s. **g** Speed of tree-canopy losses, expressed as the length of the shortest possible contiguous period (in years) within which 80% of the total 1992-2018 net losses occurred. Outside of Angola’s Cangandala National Park (**h-i**, outlined in black), faster miombo losses (i.e., <5 years, in bright red) coincide with the emergence of industrial farming, which contrasts with slow changes (i.e., > 20 years, shown in blue) likely associated with small-scale farming and charcoal-collection.

### Opportunities for more reliable monitoring and & scenario-based assessments

We found a strong correlation between FRAs and our data when equating the quality of national reporting capabilities (Nesha et al., 2021). This suggests our data is aligned with national views on the extent of forest resources. While tree-canopy cover is not directly comparable to ‘forest’ as a land-use

514 class as referred to in FRAs, the GTCCC can be combined with ancillary data to support more reliable  
515 assessments of the drivers of long-term losses and gains in forest extents. Moreso, it may support  
516 complementary assessments of the conditions of forests and other wooded ecosystems (e.g.,  
517 disturbance-succession dynamics due to climate change, Lantz et al., 2022; or forest-management  
518 regimes, Ankomah et al., 2019) , which is highly relevant for multiple goals and targets of the 2030  
519 Agenda for Sustainable Development (Estoque et al., 2021).

520 Note, however, that we reported treescape losses much smaller than those for forest cover (e.g., -  
521 0.45% vs. -17% in Vancutsem et al., 2020 for the tropical moist region). This is to be expected due to  
522 differences in mapping concepts (i.e., irrespective of additional influences of different mapping  
523 resolutions and possible data errors). Whereas categorical forest-cover data will map losses as soon as  
524 tree-cover is no longer perceived as dominant in the respective pixels, treescape extents will not show  
525 losses for as long as the underlying continuous tree-canopy cover layers perceive any trees.  
526 Correspondingly, perceived canopy-cover losses may reflect diverse processes, including woodland  
527 fragmentation or reduction of patch sizes (e.g., moving deforestation frontiers), disturbances (e.g.,  
528 landslides, windthrow), canopy thinning via selective harvesting or natural dying of individual trees,  
529 removal of small tree stands from largely treeless landscapes, as well as transitions from higher- to  
530 lower-density wooded ecosystems (e.g., forest to savanna). The GTCCC may thus support assessments  
531 of changes in extent or condition of different wooded ecosystem types, particularly if combined with  
532 other information distinguishing these processes.

533 In particular, studies of wooded ecosystem disturbances or fragmentation and related biological  
534 processes should consider whether expected canopy gaps in their study areas will be large enough to  
535 cause detectable breaks in canopy-cover gradients among neighbouring pixels. The GTCCC's pixel size  
536 (~9 ha at the equator) is at the lower ends of reported ranges of average deforestation patch sizes in  
537 Amazonia (10.6–24.7 ha; Trancoso, 2021), and average disturbance-patch sizes in moderately disturbed  
538 forest landscapes sampled across the temperate-forest biome (6.96–41.47 ha; Sommerfeld et al., 2018).  
539 However, disturbance patches are much smaller in low-disturbance temperate-forest landscapes (0.46–  
540 0.85 ha; Sommerfeld et al., 2018), and generally in European forests (mean size of 1.09 ha; Senf and  
541 Seidl, 2021). Similarly, the GTCCC data may not support meaningful assessments of habitat

542 fragmentation for species whose maximum dispersal abilities fall well below the ~300-m pixel size  
543 (e.g., many species with body sizes smaller than ~150g/~15 cm; Stevens et al., 2014).

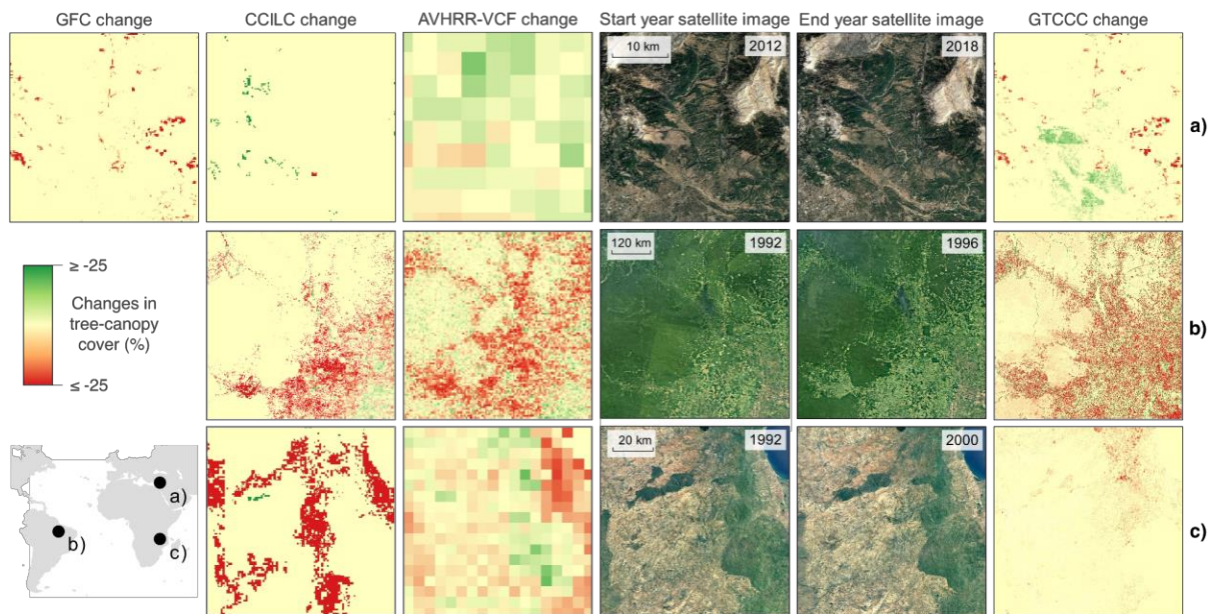
544 Besides applications in monitoring, the GTCCC can support climate modelling. Tree-canopy cover  
545 is an essential climate variable (WMO, 2022), and plays an important role in global and regional climate  
546 modelling (Gou et al., 2019). These data can more accurately inform translations of land-cover classes  
547 into fractional coverages of plant functional types, which is critical to link climate and ecosystem  
548 models (Bonan et al., 2002; Lawrence and Chase, 2007).

549 The derived climate models can then be used to infer climate normals against which to measure  
550 climate change (WMO, 2017), for which 30 years of observations are recommended to detect anomalies  
551 (WMO, 2017). For such long-time scales, however, long-term and continuous canopy-cover  
552 information had thus far only been available at coarse spatial resolutions (i.e., 5-km; Song et al., 2018)  
553 that may not meet spatial-resolution requirements of climate models (e.g.,  $\leq 1$ -km for carbon modelling;  
554 Zhao et al., 2010). With 27 years at 300-m resolution, the GTCCC thus offers improved support for  
555 climate change assessments.

#### 556 Combining strengths to overcome individual weaknesses

557 We built our data product with a novel modelling framework that explores complementary strengths  
558 of multiple existing products. For example, the use of data on land-cover allowed our model to learn  
559 how class distinctions relate to both averages and spreads of associated tree-canopy-cover values. This  
560 enabled our model to predict more plausible spatial patterns of flat vs. steep tree-canopy gradients,  
561 benefitting from a categorical product's ability to perceive transitions in tree-canopy cover dominance.  
562 In addition, by using data on coarse-scale (but temporally complete) vegetation changes compared to a  
563 future, reference year, and by relating those changes to the spatial distribution of high-resolution tree-  
564 canopy cover in that year, our model learned from the outcomes of longer-term change processes, which  
565 are co-informed by change drivers in the mapping years (i.e., climate, topography, human activity).





566

567 **Figure 6 – Mapping new tree-canopy cover changes.** Each row of figures (labelled as **a**, **b** and **c**) shows a regional example  
 568 (locations in inset map) of tree-canopy cover losses and gains as perceived by our new product (GTCCC; right) and by three  
 569 of its inputs (GFC, CCILC, AVHRR-VCFI; left to centre) during different time periods. Those periods' start/end years are  
 570 indicated in columns 4/5, which show Landsat-based true-colour images for the respective years for visual confirmation of  
 571 true change patterns. The three examples show different ways in which the GTCCC improves change depictions compared to  
 572 those of each input product taken individually. **a**) In southern Turkey, visible post-fire forest succession during the mid-2010s  
 573 is realistically captured as tree-canopy-cover gains in the GTCCC. By contrast, the CCILC only partially captures these but  
 574 instead maps large patches of gains elsewhere that are unsupported by visual inspection, as does the coarser-resolution  
 575 AVHRR-VCF, whereas the GFC by design only depicts losses. **b**) Regional patterns of tree-canopy-cover losses across a  
 576 north-eastern Amazonian deforestation frontier during the 1990s are broadly captured by both the CCILC and AVHRR-VCF  
 577 products. Yet, the new GTCCC product shows these patterns more completely than the CCILC and more refined than the  
 578 AVHRR-VCF. **c**) In a part of southern Malawi, the CCILC depicts extensive losses between 1992 and 2000, while the  
 579 AVHRR-VCF shows extensive mixtures of losses and gains. By contrast, the GTCCC depicts only minor losses and gains,  
 580 more consistent with Landsat true-colour images showing that tree-canopy cover has remained largely unchanged during that  
 581 period. The GFC is not included in b/c, as it only maps post-2000 losses.

582 Through this data integration process, our model was able to generate temporal patterns with fewer  
 583 outliers and back-and-forth changes compared to our input data (**Fig. 5 of SI**), as well as more plausible  
 584 regional patterns of tree-canopy cover changes compared to those perceived by individual inputs (**Fig.**  
 585 **6**). For example, the GTCCC better captures tree-canopy cover gains than either of the two input  
 586 products that are sensitive to tree-canopy cover (i.e., AVHRR-VCF, CCILC; note that the GFC does

587 not attempt to map gains past 2012; **Fig. 6a**). Simultaneously, the GTCCC could capture more complete  
588 patterns of gains and losses during the 1990s than the CCILC and AVHRR-VCF products (**Fig. 6b**).  
589 The model-based data-integration process also allowed correcting inaccuracies in individual products.  
590 For example, the CCILC product mapped losses of >30% (Radwan et al., 2021) during the 1990s in  
591 areas of southern Malawi that, according to visual evidence, had already been largely treeless by the  
592 beginning of that period (**Fig. 6c**). In turn, the GTCCC mapped negligible losses (-0.04% [ $\pm 0.03$ ]).

### 593 Data quality and caveats

594 The quality assessment of the GTCC yielded a high  $r^2$  ( $\geq 0.95$ ), and a relatively low RMSE ( $< 1$  ha).  
595 Yet, we found that the RMSE decreased with increasing temporal gap sizes (between the mapping year  
596 and the nearest year with reference data on tree-canopy cover), and with increasing tree-canopy covers  
597 to be reconstructed. In addition, our modelling framework had difficulties reconstructing mid-sized and  
598 large changes (2-4 ha and  $\geq 4$  ha, respectively) over periods of 1-3 years.

599 By contrast, our data performed consistently across continents and biomes when predicting small  
600 changes, and when predicting mid-sized and larger changes over  $\geq 4$  years. This indicates that the  
601 GTCCC is most suitable for studying mid- to long-term change dynamics across 4-year epochs, and is  
602 not generally suitable to analyse the timing of fine-scale changes.

603 Literature shows that most changes in tree-covered landscapes are path-dependent, multi-year  
604 processes (Taubert et al., 2018), whereas changes that are simultaneously extensive and fast are rare  
605 (Montibeller et al., 2020). However, like most global products, the GTCC is designed for broad-scale  
606 applications where specific, pixel-level errors may be expected to average out. Any smaller-scale  
607 studies that depend on highly accurate pixel-level patterns should use the GTCCC cautiously and ideally  
608 after assessing its specific fitness-for-purpose (e.g., via a local validation).

609 We validated our model's ability to predict tree-canopy cover as perceived by the GFC, a state-of-  
610 the-art product which we used as a source of samples and whose quality we aimed at replicating. In  
611 addition, we tested the plausibility of our predicted global treescape patterns via comparison against the  
612 most authoritative global information on the distribution of forest resources. Yet, it was not feasible to

613 validate tree-canopy cover values independently due to a lack of globally representative and  
614 independent validation data at relevant spatial resolutions (i.e., 300-m or coarser).

615 We caution users that although GFC's tree-canopy cover data was developed using very-high-  
616 resolution imagery, and tree-cover losses were carefully validated against independent data (Hansen et  
617 al., 2013), our random-forest model borrows its strengths along with its weaknesses. In particular, our  
618 data cannot correct cases where the GFCCC (near-)completely misses tree-canopy cover over large  
619 regions, even if trees are better captured by other products (e.g., in north-eastern Somaliland, the  
620 GTCCC underestimates the extents of montane woodlands, whereas the CCILC depicts their presence  
621 more accurately). In addition, we are unable to predict tree-canopy cover over 0.54% of the land surface,  
622 a proportion composed mainly by pacific islands and desertic lands. This is related to the fact that  
623 Landsat imagery in these areas is scarce during the early 2000s (Remelgado et al., 2023), which  
624 prevented the GFC from predicting tree-canopy cover. Since changes are inferred from the 2000  
625 baseline, changes thereafter are missing.

626 Going forward, such caveats could be addressed by integrating trusted regional products, both as  
627 source of samples and predictor data. Furthermore, forcing our machine-learning framework to consider  
628 prior theoretical understanding as exemplified in other disciplines (Seckler and Metzler, 2022; Wang et  
629 al., 2021) could help refine predictions by better acknowledging distinctions of biogeographical regions  
630 that, due to different plant adaptations, may result in specific relationships between tree-canopy cover  
631 and environmental predictors (Nicolson, 1996). Such future versions would ideally be able to rely on  
632 independent samples for validation (which are anyway a prerequisite for the existence of trustworthy  
633 regional products). Thus far, systematic efforts to derive standardised random validation samples from  
634 very-high-resolution imagery over broad spatial extents have been customised to serve modern satellite  
635 monitoring systems with higher spatial resolutions (e.g., up to a 100-m resolution; Tsendbazar et al.,  
636 2021). In the future, such systematic sampling designs could be extended to derive coarser-resolution  
637 reference data at least for recent years. Additionally, the opening of spy satellite archives holds promise  
638 for generating change-validation samples for many regions worldwide (e.g., as shown by Rizayeva et  
639 al., 2023). However, technical advances are still needed to reduce the manual effort required for  
640 generating analysis-ready images from these archives, a precondition for applying them at scale.

641 Concluding remarks

642 Model-based integration of existing remote-sensing products as showcased here cannot substitute  
643 continuous investments in higher-quality remote-sensing products based on intensive sampling and  
644 best-available algorithms for satellite-data interpretation. However, as hundreds of new remote-sensing  
645 products become available every year for different regions, scales, and application domains, there are  
646 additional opportunities in synthesising information in ways that enable additional insights or better  
647 meet the demands of certain user groups. We hope that our showcase may inspire further developments  
648 of modelling frameworks that effectively use the complementary strengths of different products to  
649 derive synthesis products that are more than the sum of their parts.

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## 655 **DATA AND CODE AVAILABILITY**

656 The GTCCC and the code needed to produce it are available through Zenodo (  
657 <https://doi.org/10.5281/zenodo.7901290>). Its predictors, however, are provided through a separate  
658 Zenodo repository (  
659 <https://doi.org/10.5281/zenodo.8217237>).

## 660 **AUTHOR CONTRIBUTIONS**

661 RR and CM designed the study. RR developed the modelling framework with support from CM. RR  
662 developed the data products reported in this paper, including the underlying code. RR and CM  
663 interpreted the results. RR and CM wrote the paper.

## 664 **CONFLICT OF INTEREST STATEMENT**

665 We declare no conflict of interest.

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1 **SUPPLEMENTARY INFORMATION**

2 **Table 1. Model predictors.** Listing of variables used to predict canopy densities. as well as the respective data sources,  
 3 temporal coverage, and spatial and temporal resolution. The column “reasoning” summarises the thought process behind the  
 4 selection of each variable.

<b>predictor</b>	<b>dataset</b>	<b>source</b>	<b>period</b>	<b>spatial res.</b>	<b>temporal res.</b>	<b>reasoning</b>
<i>Country</i>	GADM <sup>52</sup>					Controls for the influence of national governance
<i>Forest density</i>					annual	Expected future outcome of persistent gains/losses
<i>Mean year of change</i>				300-m		Long-term trends inform on the tendency for persistent loss
<i>Density of changes</i>	GFC <sup>32</sup>	Hansen et al.	2000-2020		multi-year aggregate	
<i>Mean year of change</i>				5-km		Controls for spillover effects
<i>Density of changes</i>						
<i>Land cover</i>	CCILC <sup>44</sup>		1992-2020	300-m		e.g. higher deforestation are more likely under agriculture than under dense tree cover
<i>Light intensity</i>	NTL <sup>51</sup>		1992-2018	1-km	annual	The existence and intensity of human activity drive deforestation
<i>Tree cover density</i>						
<i>Short vegetation density</i>						Year-specific data on vegetation cover occurrence and density
<i>Non-vegetated surface density</i>						
<i>Tree cover density change</i>	AVHRR-VCF <sup>45</sup>	Song et al.	1982-2016	5-km	annual	
<i>Short vegetation density change</i>						Direct information on sub-regional, long-term trends
<i>Non-vegetated surface density change</i>						
<i>Elevation</i>	MERIT DEM <sup>47</sup>	Yamazaki et al			snapshot	Changes are more common at low altitudes and over flat terrain
<i>Slope</i>						
<i>Aridity</i>	CHELSA <sup>48</sup>	-	1979-2018	1-km	multi-year aggregate	e.g. moist regions motivate extensive deforestation in favor of agricultural expansion
<i>Precipitation</i>						
<i>Tree cover mismatch (1)</i>	GFC and AVHRR-VCF		2000-2016	5-km		
<i>Tree cover mismatch (2)</i>	GFC and CCILC	-		5-km	annual	Controls for competing perceptions on the extent and density of forest cover given by different datasets
<i>Tree cover mismatch (3)</i>	CCILC and AVHRR-VCF		1992-2016	5-km		
<i>Temporal distance</i>	-	-	-	300-m	-	Time-fixed effect, controlling for the temporal distance between the mapping year and the nearest GFC data point

5

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7



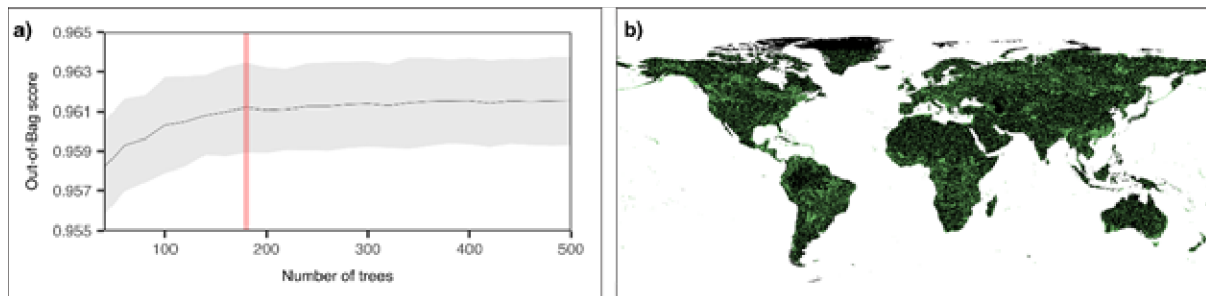
8 **Table 2. CCI land cover tree-canopy fractions.** Tree-canopy cover for each land cover class in the CCILC dataset, assuming  
 9 the central percentage value of the range of possible values indicated in the class label (cf ref (Song et al., 2014)).

Land cover class	Forest cover (%)
<i>Cropland, rainfed</i>	
<i>Herbaceous cover</i>	
<i>Tree or shrub cover</i>	
<i>Cropland, irrigated or post-flooding</i>	
<i>Mosaic cropland (&gt;50%) / natural vegetation (tree, shrub, herbaceous cover) (&lt;50%)</i>	8.3
<i>Mosaic natural vegetation (tree, shrub, herbaceous cover) (&gt;50%) / cropland (&lt;50%)</i>	25
<i>Tree cover, broadleaved, evergreen, closed to open (&gt;15%)</i>	100
<i>Tree cover, broadleaved, deciduous, closed to open (&gt;15%)</i>	100
<i>Tree cover, broadleaved, deciduous, closed (&gt;40%)</i>	100
<i>Tree cover, broadleaved, deciduous, open (15-40%)</i>	100
<i>Tree cover, needleleaved, evergreen, closed to open (&gt;15%)</i>	100
<i>Tree cover, needleleaved, evergreen, closed (&gt;40%)</i>	100
<i>Tree cover, needleleaved, evergreen, open (15-40%)</i>	100
<i>Tree cover, needleleaved, deciduous, closed to open (&gt;15%)</i>	100
<i>Tree cover, needleleaved, deciduous, closed (&gt;40%)</i>	100
<i>Tree cover, needleleaved, deciduous, open (15-40%)</i>	100
<i>Tree cover, mixed leaf type (broadleaved and needleleaved)</i>	100
<i>Mosaic tree and shrub (&gt;50%) / herbaceous cover (&lt;50%)</i>	37.5
<i>Mosaic herbaceous cover (&gt;50%) / tree and shrub (&lt;50%)</i>	12.5
<i>Shrubland</i>	
<i>Shrubland evergreen</i>	
<i>Shrubland deciduous</i>	
<i>Grassland</i>	
<i>Lichens and mosses</i>	
<i>Sparse vegetation (tree, shrub, herbaceous cover) (&lt;15%)</i>	
<i>Sparse tree (&lt;15%)</i>	
<i>Sparse shrub (&lt;15%)</i>	
<i>Sparse herbaceous cover (&lt;15%)</i>	
<i>Tree cover, flooded, fresh or brakish water</i>	100
<i>Tree cover, flooded, saline water</i>	100
<i>Shrub or herbaceous cover, flooded, fresh/saline/brakish water</i>	
<i>Urban areas</i>	
<i>Bare areas</i>	
<i>Consolidated bare areas</i>	
<i>Unconsolidated bare areas</i>	
<i>Water bodies</i>	
<i>Permanent snow and ice</i>	

11 **Table 3. Continental correlations compared to national statistics.** RMSE (in 1Mha) and  $r^2$  for the year 2000, comparing  
 12 treescape extents mapped by the GFC and by our data product against forest extents reported in national FRAs. The values  
 13 derived with our data product are accompanied by uncertainty bounds based on the upper and lower confidence intervals  
 14 surrounding the mean model predictions. We derived these metrics for each continent. In addition, we estimated the proportion  
 15 of the total FRA area that the estimated RMSE corresponds to.

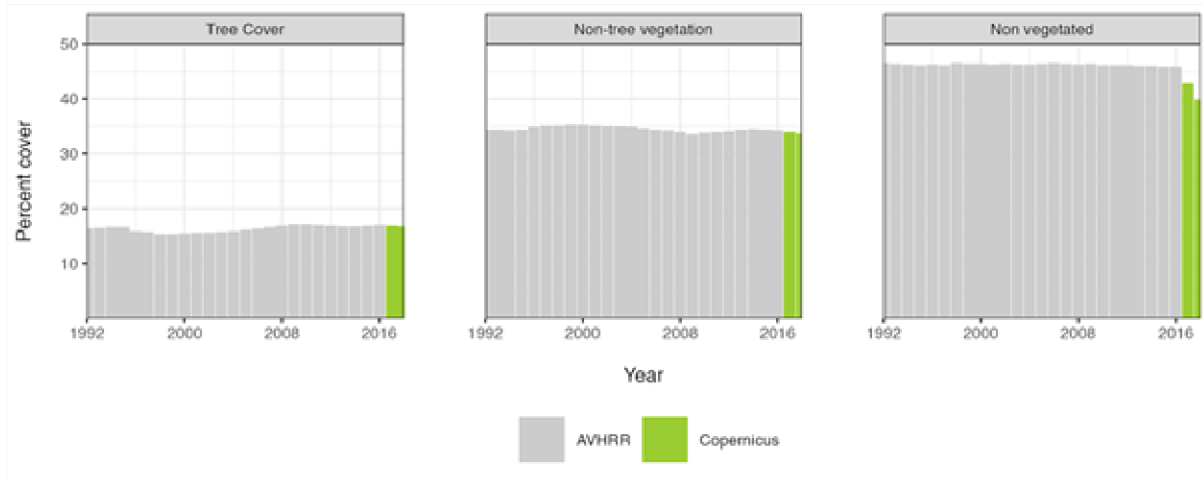
	<i>FRA</i>	$r^2$		<i>RMSE</i>		<i>RMSE as a % of the FRA</i>	
	<i>forest</i>					<i>forest area</i>	
	<i>area</i>	GTCCC	GFC	TCCC	GFC	TCCC	GFC
<i>Africa</i>	658.18	0.80 ( $\pm 0.00$ )	0.80	22.61 ( $\pm 0.01$ )	11.43	3.46 ( $\pm 0.41$ )	1.67
<i>Americas</i>	1615.36	0.97 ( $\pm 0.00$ )	0.97	79.28 ( $\pm 0.28$ )	29.88	4.92 ( $\pm 0.32$ )	1.82
<i>Asia</i>	570.46	0.89 ( $\pm 0.00$ )	0.89	33.91 ( $\pm 0.07$ )	19.63	5.96 ( $\pm 0.70$ )	3.55
<i>Europe</i>	956.7	0.98 ( $\pm 0.00$ )	0.98	92.07 ( $\pm 0.20$ )	72.97	9.63 ( $\pm 0.18$ )	7.67
<i>Oceania</i>	180.81	0.95 ( $\pm 0.00$ )	0.95	10.42 ( $\pm 0.18$ )	47.53	5.76 ( $\pm 0.25$ )	26.36

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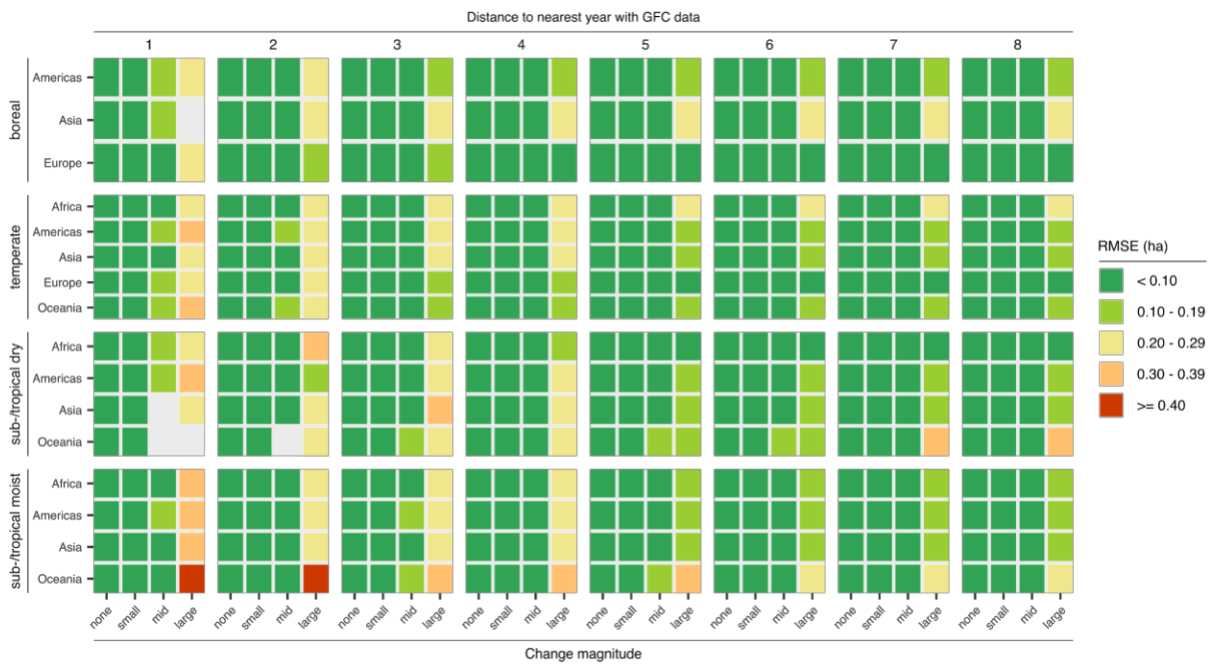


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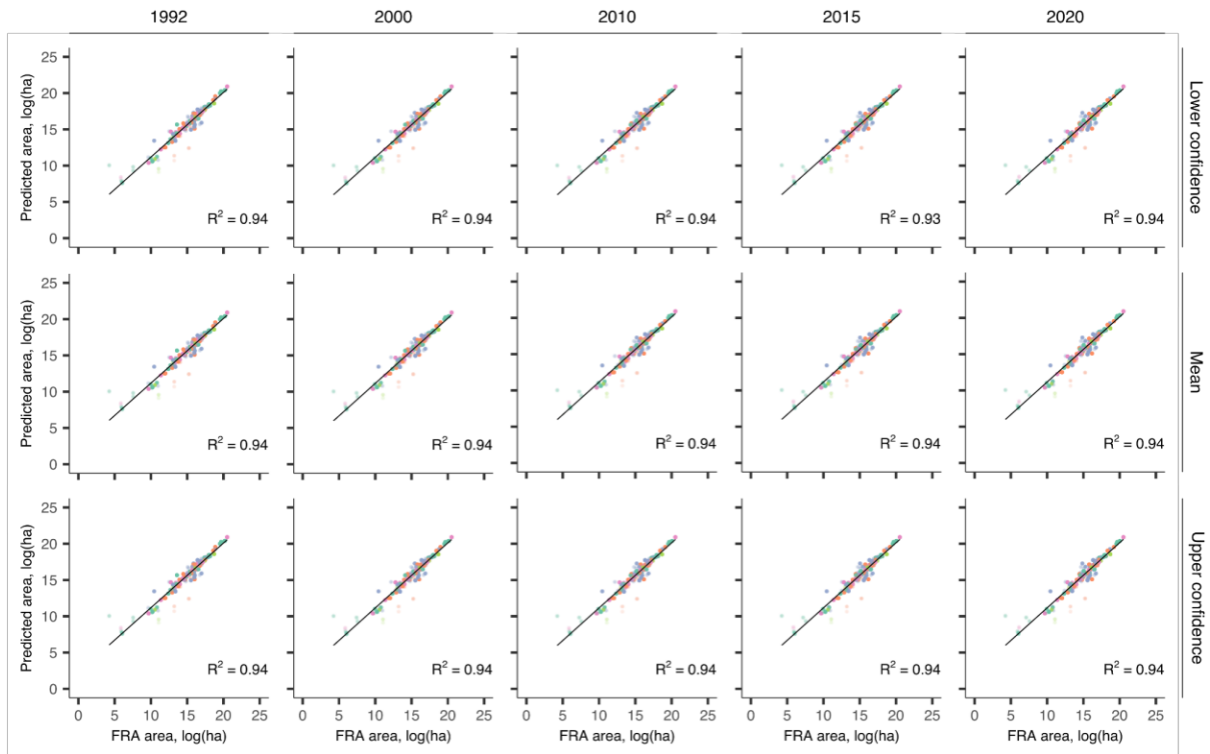
18 **Figure 1 – Model training.** Panel a) illustrates the selection of an optimal number of trees with which to train our Random  
 19 Forest Regression model (RFReg). The centre dark line shows the mean Out-of-Bag (OOB) performance for each tested forest  
 20 size, and the circumventing grey areas depict the standard deviation around the mean. The red vertical line shows the optimal  
 21 forest size of 180 trees which we selected using knee detection. Note that the knee occurred when OOB performances stopped  
 22 increasing notably. Panel b) shows the distribution of samples used to train our RFReg model. The green dots represent 78,135  
 23 unique sample locations within which we collected 625,080 samples across an 8-year period.



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 25 **Figure 2 – Temporal stability of VCF predictions.** Global percent cover of each VCF class between 1992 and 2018. Grey bars depict  
 26 values obtained through the AVHRR-VCF, while green bars represent values obtained from the predicted VCF data. We see how the  
 27 predictions of tree cover and non-tree vegetation are in line with global trends. However, predictions of non-vegetated cover deviate visibly.



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 29 **Supplementary Figure 3 – Uncertainty of quality assessment.** Confidence interval of the RMSE for each continent, biome,  
 30 and change magnitude (discriminated below), for different temporal gap sizes between the focal mapping year and the nearest  
 31 year with reference data on tree-canopy cover (shown above). This plot is supplementary to Fig. 2, which displays the  
 32 corresponding RMSE values.

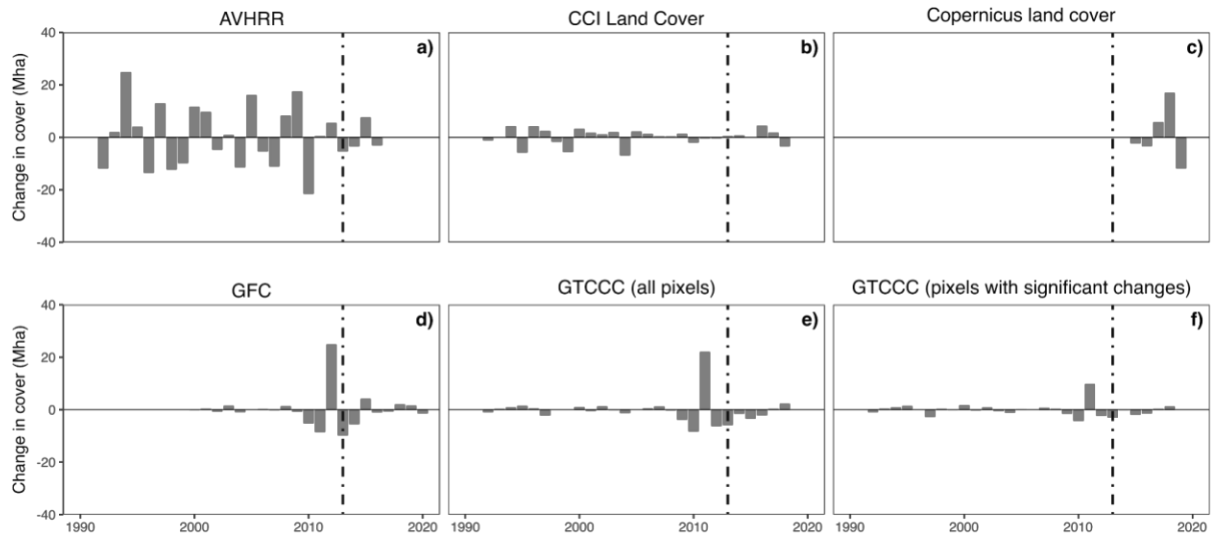


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34 **Figure 4 – Extended comparison with national statistics.** Each plot shows the correlation between forest area reported in  
 35 the Forest Resource Assessments (FRAs) and tree-canopy cover predicted by the presented GTCCC product. We ran this  
 36 analysis for each FRA assessment year (each column, distinguished above), and repeated it for the mean pixel predictions and  
 37 the upper and lower bounds of its 95% confidence interval (each row, distinguished on the right). The colours of each point  
 38 indicate the continent each country belongs to (see in Fig. 3), and faded colours indicate uncertainties regarding the quality of  
 39 national statistics. Our results show our predictions are consistent across FRA years and that correlations are robust to  
 40 uncertainty in the GTCCC predictions.  $R^2$  values were 0.94 for every iteration.

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**Figure 5. Time series consistency.** Positive/negative outliers in year-to-year changes in cover (expressed in Mha) relative to the smoothed trend (constructed with LOWESS and a span of 0.5). Each panel shows this variable for each of the datasets contributing to our data product (a-d), namely AVHRR (a), CCI land cover (b), Copernicus land cover (c, used to extend the AVHRR time-series past 2016), and GFC (d). In addition, we applied the same analysis to our data before (e) and after (f) detecting significant changes using the Mann-Kendall test. The dashed vertical line highlights 2013, when Landsat 8 was launched, and which marks a substantial improvement in the frequency and quality of the Landsat data the GFC is based on (i.e., our reference data and sample source). The plot shows that our product (e) preserves anomalies visible in the GFC between 2010 and 2013 (d), corresponding to the period when Landsat 5 was decommissioned, leaving Landsat 7 as the only source of data despite its heavy degradation. However, the magnitude of these anomalies appears smaller in our data. When limiting our focus to pixels with significant changes ( $p$ -value  $< 0.0005$ ) identified through the Mann-Kendall test, the prevalence of anomalies is reduced further. Note that despite integrating coarse-resolution (a), categorical (b), and contemporary data (e) to build our predictive model, their year-to-year anomalies are not replicated in our data.

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