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Unsupervised Mapping of Urban Tree Diversity using Spatially-aware Visual Clustering

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Abstract

Urban tree biodiversity is critical for climate resilience, ecological stability, and livability in cities, yet most municipalities lack detailed knowledge of their canopies. Field-based inventories provide reliable estimates of Shannon and Simpson diversity but are costly and time-consuming, while supervised AI methods require labeled data that often fail to generalize across regions. We introduce an unsupervised clustering framework that integrates visual embeddings from street-level imagery with spatial planting patterns to estimate biodiversity without labels. Applied to eight North American cities, the method recovers genus-level diversity patterns with high fidelity, achieving low Wasserstein distances to ground truth for Shannon and Simpson indices and preserving spatial autocorrelation. This scalable, fine-grained approach enables biodiversity mapping in cities lacking detailed inventories and offers a pathway for continuous, low-cost monitoring to support equitable access to greenery and adaptive management of urban ecosystems.

33 1 Introduction

34 Quantifying biodiversity in urban forests is essential to assess ecosystem health and to
35 guide evidence-based management and policy decisions for urban green spaces. Widely
36 used metrics like Shannon entropy and Simpson's index integrate both richness (num-
37 ber of taxa) and evenness (distribution of abundances) into a single value, providing
38 interpretable indicators for comparing ecological conditions across locations and over
39 time [1, 2]. However, accurate estimation of these indices at the genus or species
40 level typically requires detailed and geo-referenced taxonomic inventories, most often
41 collected through arborist-led field surveys [3]. Although these surveys are reliable,
42 they are resource intensive. Urban street-tree programs may spend up to \$65 per tree
43 annually on inventory, monitoring, and management, which has limited large-scale
44 implementation to only a few well-funded cities [4, 5].

45 In the past 50 years, adapting to climate change has become a priority for cities
46 around the world. Seminal and contemporary studies [6–9] highlight that anthro-
47 pogenic changes in the past few decades have caused a rapid loss of urban biodiversity.
48 In addition to supporting biodiversity and urban ecosystems [10], trees play a critical
49 role in maintaining livability by contributing to equitable access to greenery [11–13],
50 improved air quality [14, 15], and thermal comfort [16, 17]. Yet, tree planting strategies
51 are often guided by aesthetic preferences rather than ecological function [18], resulting
52 in reduced diversity and increased vulnerability to pests, diseases, and climate stres-
53 sors [13]. A more diverse and ecologically informed urban canopy can buffer against
54 these risks and adapt over time [19–21].

55 Open access data sources, such as Street View Imagery (SVI), offer a scalable alter-
56 native for biodiversity monitoring due to their extensive spatial coverage [22]. However,
57 SVI datasets are unlabeled, meaning the images do not specify the tree species present
58 and require substantial pre-processing to extract ecological information suitable for
59 diversity estimations. Identifying individual taxa from imagery is particularly chal-
60 lenging: trees of the same species may look very different depending on season, age, or
61 management, while distinct species can appear deceptively similar. These difficulties
62 make automated classification highly sensitive to subtle morphological cues. Current
63 supervised ecological AI models [23, 24], while effective within their training domains,
64 often under-perform when applied to regions or taxa not well represented in training
65 data, largely due to geographic domain shift [25].

66 Self-supervised learning offers a promising alternative by eliminating the need for
67 labeled data [26]. However, when applied to limited or homogeneous datasets, these
68 methods often under-perform due to overfitting and poor generalization, as robust
69 representations typically require large and diverse training samples. Few clustering
70 approaches have been explicitly designed to preserve abundance distributions in a way
71 that enables accurate Shannon and Simpson estimation from unlabeled imagery. This

72 capability is essential for producing ecologically meaningful biodiversity metrics from
73 large-scale, unlabeled datasets.

74 In this work, we address the challenge of large-scale, label-free biodiversity moni-
75 toring by developing an unsupervised clustering approach that estimates genus-level
76 urban tree diversity directly from street-level imagery. We apply the framework across
77 eight North American cities with contrasting forest compositions and imbalance, and
78 evaluate its ability to recover biodiversity metrics such as Shannon and Simpson
79 indices as well as spatial distribution patterns. While our evaluation is necessarily con-
80 strained to cities with available ground-truth inventories, the results demonstrate that
81 the approach generalizes across diverse urban forest contexts and provides a scalable
82 pathway for integrating biodiversity insights into urban resilience planning.

83 2 Results

84 We evaluated our framework in eight North American cities spanning a wide range of
85 forest compositions and genera imbalances (see Section 4.1). All biodiversity metrics
86 are computed at the level of spatial grid cells, obtained by partitioning each city
87 into regular 500-m grids, with each cell representing a local urban tree community.
88 Accordingly, richness, Shannon, and Simpson are all treated as local (α -diversity)
89 measures, and $RMSE_\alpha$ quantifies error in cell-level richness rather than city-wide
90 (γ) diversity. The method consistently recovered meaningful biodiversity patterns,
91 demonstrating robustness even in cities dominated by a few genera. Table 1 reports
92 results for the best-performing configuration, selected through hyperparameter tuning
93 (see Appendix A) and ablation study (see Appendix B), which achieved a balanced
94 trade-off between richness accuracy, clustering quality, and diversity preservation.

Table 1: Clustering performance for our clustering framework across cities. Metrics include richness error ($RMSE_\alpha \downarrow$), clustering quality (V-score \uparrow), and distributional agreement with ground-truth Shannon (scaled) and Simpson diversity measured using the 1-Wasserstein distance ($W_1 \downarrow$). Values are reported as mean [95% CI] through bootstrapping [27].

City	$RMSE_\alpha \downarrow$	V-score \uparrow	$W_{1,Shannon} \downarrow$	$W_{1,Simpson} \downarrow$
Calgary	4.39 [4.20–4.59]	0.397 [0.387–0.406]	0.215 [0.208–0.222]	0.178 [0.169–0.186]
New York	3.42 [3.34–3.51]	0.476 [0.471–0.480]	0.073 [0.070–0.077]	0.064 [0.059–0.068]
Columbus	5.99 [5.76–6.22]	0.430 [0.422–0.437]	0.177 [0.172–0.183]	0.140 [0.134–0.147]
Denver	5.39 [5.15–5.62]	0.487 [0.480–0.494]	0.069 [0.065–0.073]	0.040 [0.036–0.045]
Los Angeles	3.74 [3.63–3.84]	0.477 [0.473–0.481]	0.070 [0.067–0.073]	0.067 [0.064–0.071]
Seattle	9.94 [9.56–10.3]	0.487 [0.481–0.493]	0.155 [0.150–0.160]	0.095 [0.089–0.101]
Washington	15.6 [14.7–16.3]	0.498 [0.491–0.505]	0.133 [0.127–0.140]	0.033 [0.029–0.039]
San Francisco	10.0 [9.17–10.8]	0.493 [0.485–0.501]	0.034 [0.028–0.041]	0.028 [0.023–0.034]

95 ***Richness and cluster quality results.***

96 As shown in Table 1, genera imbalance strongly influenced richness estimation error
 97 ($RMSE_\alpha$), though not uniformly across cities. In highly uneven forests such as
 98 Washington and Seattle, $RMSE_\alpha$ exceeded within-city variability, underscoring the
 99 difficulty of estimating richness under strong dominance patterns. By contrast, in more
 100 balanced systems such as Los Angeles, Denver, and New York, errors were smaller than
 101 or comparable to observed standard deviations, and in several cases outperformed a
 102 naïve mean-predictor baseline (see Appendix E). Despite these differences, the V-score
 103 remained consistent across all datasets (0.397–0.498), indicating that clustering pre-
 104 served the overall partitioning structure even when richness estimates deviated. These
 105 findings suggest that richness is not the primary strength of our approach, but per-
 106 formance is competitive in balanced forests and improves over statistical baselines.
 107 Richness estimation is inherently more sensitive to long-tail genus distributions, as
 108 rare taxa contribute little to the dominant visual and spatial signal regardless of learn-
 109 ing paradigm. This limitation is therefore not specific to unsupervised clustering, and
 110 could be mitigated in future work through targeted sampling [28], weak priors on rar-
 111 ity [29], or emerging long-tail-aware generative augmentation strategies [30], which
 112 remain challenging in ecological settings.

113 ***Diversity results.***

114 For Shannon and Simpson diversity, we assessed agreement with ground truth using
 115 the Wasserstein distance [31]. This metric quantifies the minimum "cost" of trans-
 116 forming one distribution into another, accounting not only for absolute differences but
 117 also for their positions across the value range. Unlike RMSE, which captures pointwise
 118 error, the Wasserstein distance is sensitive to systematic shifts in distributional shape
 119 and spread, making it particularly suitable for ecological indices. We focus on Shannon
 120 and Simpson entropies due to their long-standing theoretical foundation, widespread
 121 adoption, and interpretability in ecological research [1]. These classical metrics capture
 122 complementary aspects of community structure such as balanced weighting of common
 123 and uncommon taxa (Shannon) and dominance-driven structure (Simpson)[32].

124 In our datasets, Simpson diversity is bounded between 0 and 1, while Shannon
 125 diversity can reach values of about 3.5 (see Section 4.1). To enable direct comparison
 126 across indices and cities, Shannon values were normalized by their relative maximum,
 127 placing both metrics on a common [0,1] scale. In this context, Wasserstein distances
 128 below 0.05 indicate very strong alignment with ground truth distributions, while values
 129 near 0.1 represent moderate but still ecologically meaningful agreement [33]. Cities
 130 such as Washington (0.133 for Shannon, 0.033 for Simpson) and San Francisco (0.034
 131 for Shannon, 0.028 for Simpson) demonstrated strong alignment for both indices, with
 132 proportionally smaller errors for Simpson. Calgary, by contrast, exhibited the weakest
 133 alignment across metrics. We hypothesize that this may be linked to more aggressive
 134 visual obfuscation apparent in Canadian Street View imagery compared to the United
 135 States, where stronger privacy safeguards and stricter blurring protocols are enforced
 136 [34]. Fig. 1 illustrates this by comparing predicted and observed Shannon values in Los
 137 Angeles (Fig. 1a) and Simpson values in New York (Fig. 1b). A full set of city-level
 138 results, including baseline comparisons and variability relative to observed standard

139 deviations, is provided in Appendix E, which contextualizes how performance varies
140 across urban forest structures.

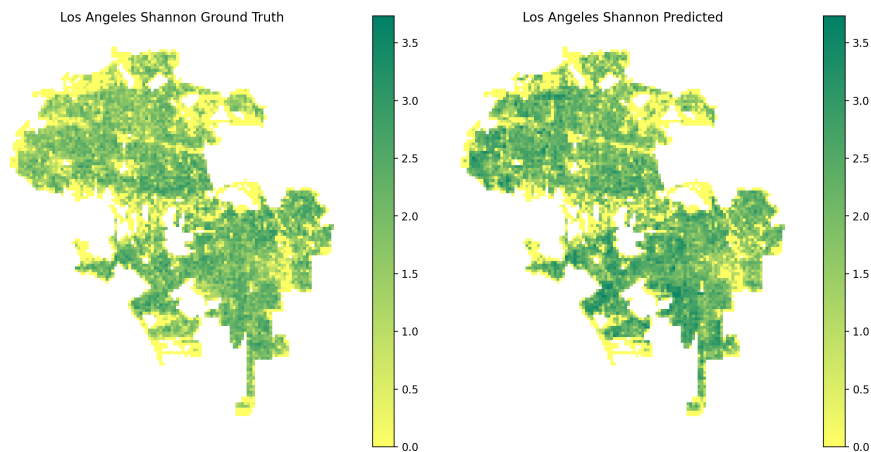
141 *Diagnosis of possible entropy inflation.*

142 One potential concern with unsupervised clustering is that lower distributional errors
143 may arise not from genuine recovery of community structure but from artificial entropy
144 inflation, e.g., through oversplitting or the proliferation of singleton clusters. To assess
145 this, we performed a dedicated diagnostic analysis (Appendix D) that examined over-
146 splitting factors, purity, evenness shifts, and rank–abundance correlations between
147 predicted and observed communities against a supervised model using linear probing
148 on BioCLIP [23] model. Across all cities, the singleton ratio was zero, oversplitting
149 factors remained close to one, and evenness shifts were modest, with New York even
150 showing a slight decrease. At the same time, rank–abundance correlations were con-
151 sistent high ($\rho \approx 0.9$), indicating that predicted clusters preserved the overall shape
152 of community distributions. These findings confirm that the improved alignment of
153 Shannon and Simpson diversity under the unsupervised approach is not an artifact
154 of inflated entropy, but rather reflects ecologically meaningful recovery of biodiversity
155 structure.

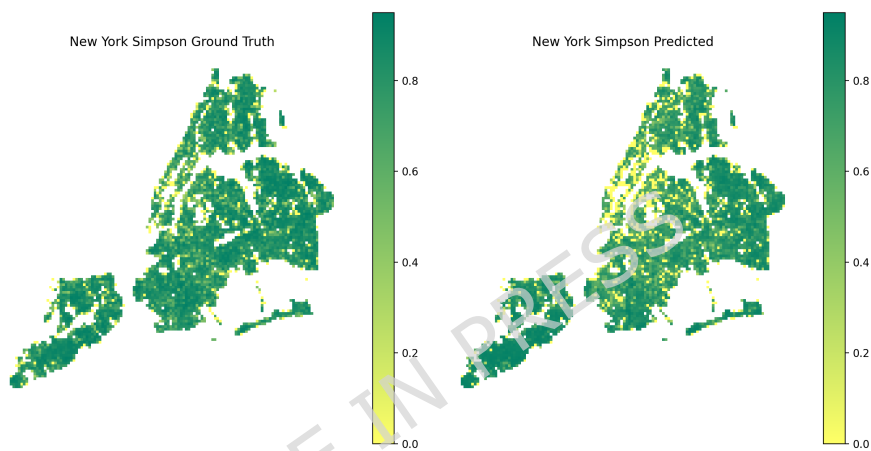
156 Overall, the method achieved low Wasserstein distances for both Shannon and
157 Simpson diversity across diverse urban forest conditions, including in highly imbal-
158 anced contexts. High Simpson accuracy suggests the framework effectively captured
159 dominant genera distributions, while strong Shannon performance indicates robustness
160 to abundance evenness patterns. Taken together, these results demonstrate that our
161 approach can accurately recover both abundance distributions and spatial diversity
162 patterns with strong spatial fidelity (see Appendix C) from entirely unlabeled street-
163 level imagery, directly addressing the lack of scalable, label-free biodiversity mapping
164 methods highlighted in the Introduction.

165 **3 Discussion**

166 Our results demonstrate that an unsupervised and spatially aware clustering frame-
167 work can recover both magnitude and spatially weighted biodiversity measures, where
168 diversity is computed locally and evaluated across spatial grid cells, from unlabeled
169 street-level imagery with high fidelity, even under markedly imbalanced genus distribu-
170 tions. Across diverse urban contexts, the method achieved low Wasserstein distances
171 for Shannon and Simpson diversity, while preserving spatial autocorrelation patterns
172 with minimal deviation from ground truth. These findings are consistent with previous
173 reports that dominant taxa patterns are easier to recover than local evenness structure
174 driven by the relative abundances of non-dominant genera [1, 22], but extend prior
175 work by showing that evenness-sensitive indices such as Shannon can also be estimated
176 reliably when spatial priors are incorporated. Crucially, by enabling robust, label-free
177 biodiversity mapping, this approach offers a scalable tool that could be integrated
178 into municipal monitoring systems, guide nature-based policy, and support adaptive
179 planning in cities with limited ecological data infrastructure.



(a) Los Angeles's Shannon Entropy Ground Truth and Predicted Results



(b) New York's Simpson Entropy Ground Truth and Predicted Results

Fig. 1: A comparison between ground truth and predicted values of Shannon entropy in Los Angeles and Simpson entropy in New York.

180 Our ablation analysis (Appendix B) showed that the joint use of outlier elimina-
 181 tion and cluster merging provided the best balance between clustering quality, richness
 182 accuracy, and preservation of diversity structure. This result indicates that addressing
 183 noise (through elimination) and fragmentation (through merging) in tandem is particu-
 184 larly effective for recovering meaningful ecological patterns from unlabeled imagery.
 185 More broadly, it suggests that unsupervised clustering in environmental applications
 186 benefits from strategies that explicitly mitigate both spurious assignments and over-
 187 segmentation. While prior studies have applied label-free clustering to infer urban

188 landforms from very high-resolution satellite data [35] or crop types from multispec-
189 tral time series [36], such approaches typically rely on costly or proprietary datasets.
190 By contrast, our framework demonstrates that freely available street-level imagery
191 can support scalable, label-free biodiversity assessment in urban forests, extending
192 applicability to regions where high-resolution remote sensing is not accessible.

193 From an applied perspective, the framework offers a scalable alternative to field-
194 based inventories for monitoring urban tree diversity and can be integrated into urban
195 forestry programs. Spatially explicit Shannon and Simpson maps at the city scale can
196 guide planting strategies, highlight areas at risk of monocultures, and support temporal
197 monitoring, while reducing the effort and cost of repeated ground surveys. Importantly,
198 this directly addresses the central gap identified in the Introduction in regards to the
199 inability of existing methods to recover abundance distributions and spatial diversity
200 patterns from unlabeled imagery. By overcoming this limitation, our approach provides
201 a pathway for continuous, cost-effective biodiversity monitoring that can inform urban
202 resilience planning. Here, “city-scale” refers to the aggregation and analysis of spatially
203 explicit grid-level diversity patterns, rather than to the direct estimation of total city-
204 wide (γ) richness. City-level insights are obtained by summarizing the distribution
205 and spatial organization of grid-level Shannon and Simpson diversity values, enabling
206 identification of dominance hotspots, monoculture risk, and spatial inequities without
207 explicitly inferring total genus counts.

208 Several constraints remain. Street-level imagery is updated at irregular intervals,
209 sometimes years apart, which complicates the interpretation of temporal change.
210 In practice, this means the framework is better suited to structural baselines and
211 cross-sectional monitoring than to precise year-to-year trend detection. Coverage is
212 uneven globally, with gaps in rural areas, private developments, and in certain coun-
213 tries; thus, results are most applicable in dense urban cores of cities with established
214 GSV infrastructure, and less transferable to places where coverage is patchy. Privacy-
215 driven blurring can obscure morphological features needed for genus discrimination,
216 potentially biasing genus detection toward larger or more distinctive trees, and under-
217 representing genera with finer-grained leaf or bark traits. Seasonal foliage changes,
218 variable lighting, and transient occlusions (e.g. vehicles or pedestrians) introduce visual
219 noise, which can reduce consistency across time and space and necessitate larger sam-
220 ple sizes to stabilize estimates. In such cases, phenological variation may cause visually
221 distinct seasonal appearances of the same genus to be assigned to separate pseudo-
222 taxa, potentially affecting cluster purity or inflating effective richness. Additionally,
223 the method only captures the visible streetscape; off-street vegetation is excluded,
224 which can underestimate diversity in areas where backyards, courtyards, or parks hold
225 a significant share of urban trees.

226 Looking ahead, performance could be enhanced by incorporating vision–language
227 models (VLMs) that leverage fine-grained textual descriptors of morphological traits
228 [37], improving discrimination between visually similar trees. As model efficiency
229 advances, large-scale VLM integration may become feasible. Further gains may come
230 from foundational models trained on ecologically diverse, high-resolution vegetation
231 datasets, coupled with domain-adaptive fine tuning to improve generalization to under-
232 represented regions. In parallel, prior work shows that satellite RGB imagery alone

233 is insufficient for reliable urban tree assessment [38, 39]; integrating complementary
234 modalities such as airborne LiDAR for structural detail and high-resolution multispec-
235 tral data or recent foundation models [40] for spectral cues would enrich representation
236 and enable a more holistic view of urban forest diversity. Beyond technical gains,
237 these enhancements would directly support cross-city generalization, allowing models
238 to adapt to ecological and urban form variability across different contexts. Moreover,
239 the fusion of tree diversity indicators with other environmental datasets such as air
240 quality, temperature, or land use would situate this work within the broader trend
241 of multimodal urban environmental monitoring, creating synergies for applications in
242 climate resilience, ecosystem services evaluation, and nature-based urban planning.

243 In sum, this work establishes that unsupervised clustering of spatial and visual
244 embeddings can be a practical, transferable, and data-efficient approach to quantifying
245 urban tree diversity. By recovering both the magnitude and spatial organization of
246 Shannon and Simpson diversity indices, the framework offers a pathway for continu-
247 ous, city-scale biodiversity monitoring and supports the development of more resilient,
248 ecologically informed urban forests. By enabling the recovery of both abundance distri-
249 butions and spatial diversity patterns from unlabeled imagery, our approach opens the
250 door to scalable, globally comparable biodiversity mapping for urban environments.

251 4 Methods

252 We approach urban tree biodiversity with an unsupervised clustering framework driven
253 by spatial priors, using neighborhood structure and planting patterns as the primary
254 signal. Rather than relying on datasets that label each tree by species or genus and
255 then training a supervised model on those labels, our method operates without labels
256 and instead groups trees into taxonomically coherent clusters using visual embeddings
257 extracted from street-level imagery. Each cluster functions as a pseudo-taxon, allowing
258 the computation of Shannon, Simpson, and related indices without the need for explicit
259 species identification. The end-to-end pipeline consists of five stages: data description,
260 spatial and visual embedding extraction, spatial-visual clustering, cluster refinement,
261 and biodiversity metric computation. An overview of the methodology is shown in
262 Fig. 2 and detailed later in this section.

263 4.1 Data Description

264 This study relies on the Auto Arborist dataset [38], a large-scale benchmark of urban
265 street trees derived from dense sampling of Google Street View (GSV) imagery. While
266 the dataset spans numerous cities, we selected eight cities that span a wide range of
267 urban form, genera composition, and spatial distribution, providing a robust testbed
268 for cross-city evaluation of the framework. Across these cities, the dataset contains
269 hundreds of thousands of geotagged tree instances annotated at the genus level.

270 To move from city-level coverage to local-scale analysis, each city is discretized into
271 regular 500-m grid cells, which serve as the fundamental spatial units of analysis. All
272 biodiversity metrics are computed at the grid-cell level, with each cell representing a

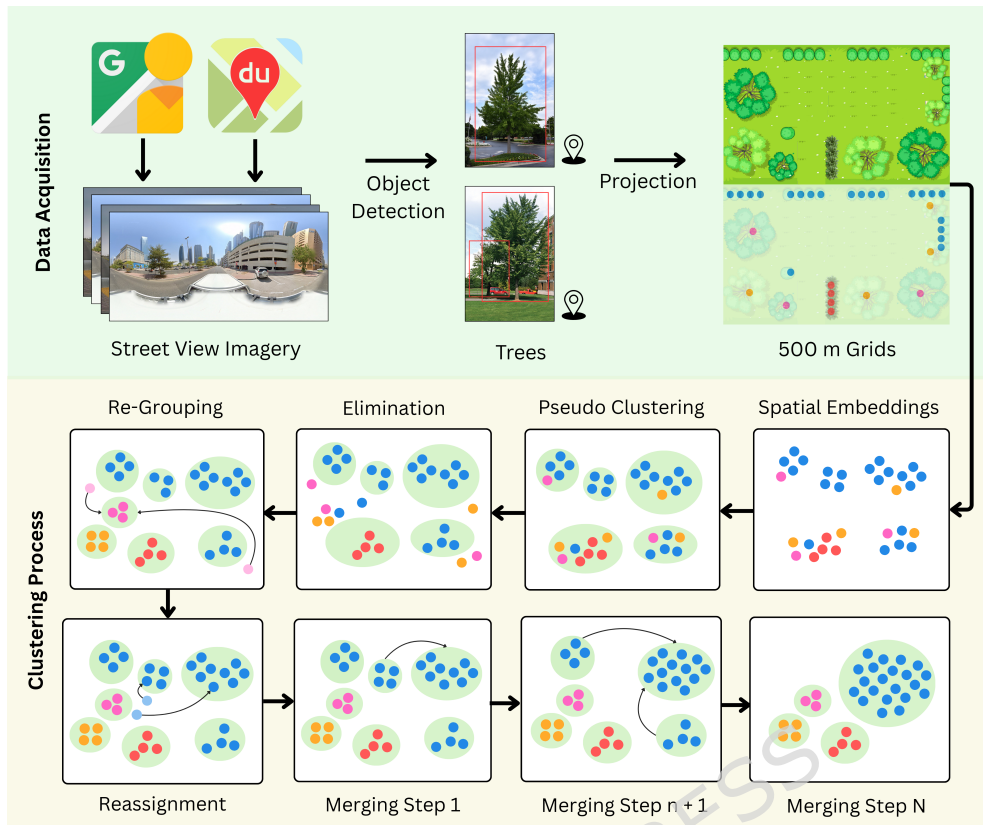


Fig. 2: Overview of the methodology. Street-level imagery (see Section 4.1) is processed using an object-detection model from prior work to extract tree instances, which are geo-located and mapped into 500 m grid cells (top row). The bottom rows illustrate the iterative clustering process applied to spatial embeddings: from left to right, the procedure consists of pseudo-clustering, elimination, and re-grouping (top sequence), followed by reassignment and successive merging steps (bottom sequence), culminating in stable genus-level clusters. Together, these stages yield coherent genus-level tree distributions across space.

273 local urban tree community. The dataset is no longer publicly hosted due to mainte-
 274 nance constraints, but remains available from the authors upon request. Full details
 275 of the acquisition and preprocessing pipeline are provided in [38].

276 All analyses are conducted at the genus level, which represents a well-established
 277 and pragmatic taxonomic resolution in large-scale urban tree studies. At this level,
 278 taxonomic groupings remain ecologically meaningful while avoiding the substantial
 279 ambiguity and visual overlap that often limit reliable species-level discrimination
 280 from street-level imagery. Many functional traits, environmental tolerances, and man-
 281 agement considerations are conserved within genera, making genus-level aggregation

282 suitable for biodiversity assessment and urban forestry applications [41–43]. This reso-
 283 lution also provides a stable basis for comparative analysis across cities while remaining
 284 compatible with the structure of widely used urban tree inventories. To ensure repro-
 285 ducibility and enable further research, we will release the spatial and visual embeddings
 286 of some cities generated in this study, providing a standardized, high-quality resource
 287 for consistent and cross-city evaluation.

Table 2: Summary of urban forest characteristics for the eight North American cities in the Auto Arborist dataset. Reported metrics include urban area, total tree counts, number of genera, genera imbalance ($\mu_{IR} \pm \sigma_{IR}$), and the maximum grid-cell-level Shannon diversity (H'_{\max}), shown as a reference for local α -diversity rather than city-wide entropy.

City	Urban Area (km ²)	#Trees	#Genera	$\mu_{IR} \pm \sigma_{IR}$	H'_{\max}
Calgary, AB	621.7	64,476	35	11.28 \pm 18.02	2.47
New York, NY	1,223.6	560,069	68	11.33 \pm 14.67	3.12
Columbus, OH	1,336.5	114,536	81	17.96 \pm 24.95	3.18
Denver, CO	1,760.0	175,438	97	18.40 \pm 20.25	3.27
Los Angeles, CA	5,910.0	391,788	202	20.98 \pm 27.90	3.50
Seattle, WA	2,546.0	150,983	142	34.06 \pm 37.07	3.44
Washington, DC	3,351.0	152,983	71	43.13 \pm 43.38	2.89
San Francisco, CA	1,330.0	154,698	195	57.96 \pm 77.78	3.48
Average	2,384.5	220,746	111	26.43 \pm 32.00	3.17

288 The dataset used encompasses eight major North American cities (Table 2), cap-
 289 turing substantial variation in urban extent, tree abundance, taxonomic richness, and
 290 genera imbalance. Urban area spans nearly an order of magnitude, from 621.7 km²
 291 in Calgary to 5,910 km² in Los Angeles. Total tree counts range from roughly 64,000
 292 in Calgary to over 560,000 in New York, reflecting differences in both city size and
 293 canopy coverage. Taxonomic richness, measured as the number of genera, peaks in Los
 294 Angeles (202) and San Francisco (195), indicating broad genera variety, and is lowest
 295 in Calgary (35), suggesting a more homogeneous composition.

296 The genera imbalance factor (IR), defined as the mean \pm standard deviation of the
 297 maximum-to-minimum (MM) genus count ratio within city grids, highlights marked
 298 contrasts in dominance patterns. New York exhibits the most even composition (11.33
 299 \pm 14.67), whereas San Francisco (57.96 \pm 77.78), Washington (43.13 \pm 43.38), and
 300 Seattle (34.06 \pm 37.07) display strong skew toward a few dominant genera, because
 301 richness is driven by the accurate detection of rare genera, which often appear as
 302 low-abundance or singleton occurrences and are therefore more susceptible to mis-
 303 assignment or merging errors in clustering-based approaches. Such high imbalance
 304 factors present greater challenges for biodiversity estimation, particularly for rich-
 305 ness metrics. Importantly, IR should be interpreted as a measure of dominance rather
 306 than a direct indicator of overall biodiversity: cities with high IR can still host many

307 genera, but those genera are unevenly distributed, with a few disproportionately rep-
308 presented. In our analysis, we found that while extreme IR values often coincided with
309 higher richness errors, they did not consistently predict lower Shannon or Simpson
310 diversity, since these indices emphasize evenness and relative abundance rather than
311 absolute richness. In particular, Simpson diversity and, to a lesser extent, Shannon
312 entropy largely reflect dominant and common genera, whereas rare genera contribute
313 disproportionately to richness but minimally to these entropy-based measures [32].

314 Shannon diversity, which in this dataset can reach values of approximately 3.5,
315 further distinguishes the cities. Larger and compositionally diverse cities such as Los
316 Angeles, San Francisco, and Seattle attain the highest maxima, reflecting both high
317 richness and evenness in certain localities. By contrast, cities with fewer genera or
318 pronounced dominance, such as Calgary and Washington, tend toward lower Shannon
319 maxima, consistent with reduced local evenness.

320 Collectively, these differences position the dataset used as a diverse and challenging
321 benchmark to test the generalizability of unsupervised clustering methods in urban
322 biodiversity assessment. A detailed pseudo-code of the pipeline is provided in Appendix
323 B (Algorithm ??), and the complete implementation, including hyperparameter tuning
324 and preprocessing scripts, is available on Github.

325 4.2 Clustering Framework

326 Clustering urban tree data from street-level imagery requires methods that can
327 accommodate both fine-grained visual differences between genera and spatial patterns
328 imposed by urban design. Our framework combines visual and spatial embeddings in
329 a multi-stage pipeline that progressively refines initial groupings, removing visual out-
330 liers, merging highly similar clusters, and reassigning ambiguous samples. This design
331 balances taxonomic accuracy with spatial representativeness, ensuring that diversity
332 metrics capture both the magnitude and spatial structure of urban forest biodiversity.

333 4.2.1 Pseudo Clustering

334 Urban trees are often arranged linearly along streets and boulevards to provide shade,
335 enhance aesthetic appeal, and define pedestrian zones [44]. However, many clustering
336 algorithms perform poorly on linear or anisotropic spatial patterns, as they often
337 assume compact clusters with uniform densities. Clustering methods like Ordering
338 Point to Identify the Clustering Structure (OPTICS) [45] or Fuzzy *c*-Means (FCM) [46]
339 rely on parameters or assumptions that are not well suited for elongated or non-convex
340 cluster geometries.

341 Spatial embeddings derived from TaxaBind [24], a multimodal foundation model
342 that incorporates geographic location and environmental features globally, provide a
343 powerful solution to the limitations of traditional clustering algorithms when applied
344 to linearly arranged urban tree data. Unlike centroid- or density-based methods, which
345 often fail to capture elongated or irregular structures [47], spatial embeddings trans-
346 forms raw geographic coordinates into a learned representation that encodes relative
347 spatial context and structural relationships. This allows linearly aligned trees to be
348 represented as coherent patterns, even in the presence of spacing variability, gaps, or

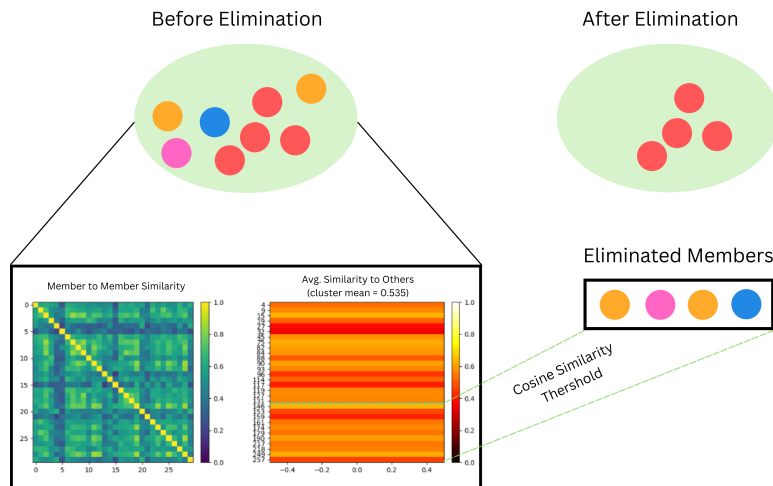


Fig. 3: Intra-cluster similarity filtering using a tuned threshold. Low-similarity members (highlighted) are excluded to increase cluster coherence.

349 missing data. By preserving and enhancing neighborhood relationships, spatial embed-
 350 dings help group trees that share local spatial context, making clustering more robust
 351 to urban morphology.

352 For visual features, we use BioCLIP [23] to extract fine-grained, taxonomically rel-
 353 evant representations from street-level imagery. These visual embeddings complement
 354 the spatial representation by capturing leaf, bark, and crown characteristics that are
 355 essential for distinguishing between visually similar genera.

356 As an initial step, we group TaxaBind spatial embeddings into pseudo-clusters
 357 using the HDBSCAN algorithm [48], leveraging the tendency of urban trees to be
 358 planted in structured, often linear arrangements. This approach typically results in
 359 taxonomically pure clusters due to the spatial regularity introduced by common urban
 360 planting strategies such as row planting along streets or uniform spacing in public
 361 parks.

362 4.2.2 Outlier Elimination

363 Outlier elimination based on intra-cluster visual similarities aims to improve the coher-
 364 ence and reliability of clustering by removing samples that visually deviate from the
 365 dominant patterns within their assigned group. After forming initial clusters based
 366 on spatial proximity, we compute pairwise cosine similarities [49] between all visual
 367 embeddings within each cluster. Cosine similarity is particularly suitable here because
 368 it measures the angular closeness of vectors rather than their raw magnitudes, which is
 369 advantageous in high-dimensional embedding spaces where differences in vector length
 370 are less informative than differences in orientation. This makes it effective for captu-
 371 ring whether samples share a common visual representation, even when the embedding
 372 space is large and sparse. Samples with consistently low similarity to other members
 373 are considered visually inconsistent and are eliminated as outliers.

374 Because visual differences among tree genera can be subtle and fine-grained, espe-
375 cially in urban settings with morphologically similar genera, directly detecting outliers
376 is challenging. To address this, we adopt a conservative strategy in which outliers are
377 identified as those samples whose average similarity to other cluster members falls
378 below a threshold determined through hyperparameter tuning (see Appendix A). This
379 thresholded filtering prioritizes consistency while avoiding premature exclusion of valid
380 but visually nuanced samples, resulting in more stable and interpretable clusters in
381 subsequent steps. This process is further illustrated in Fig. 3.

382 4.2.3 Re-grouping and Re-assignment

383 The re-grouping and re-assignment process targets samples eliminated during the out-
384 lier detection stage, which were initially clustered based primarily on spatial proximity
385 rather than visual coherence. Since such pseudo-clusters may group together visu-
386 ally dissimilar trees due to their physical arrangement, the eliminated members are
387 revisited to assess whether they form more semantically consistent groupings.

388 Visual embeddings of all eliminated samples are compared, and if a subset exhibits
389 mutual similarity above the tuned cosine similarity threshold (Appendix A), elimi-
390 nated samples are grouped into a new cluster, capturing visual structure that spatial
391 proximity alone may have missed. Remaining samples are evaluated for reassignment
392 to the most visually similar existing cluster, again using the tuned threshold. If no
393 such reassignment is possible, the sample is treated as a singleton cluster.

394 This step enhances visual coherence in the final clustering while improving the
395 model's ability to capture the long-tail distribution of urban tree genera where many
396 rare or underrepresented classes naturally appear as singletons.

397 4.2.4 Merging Process

398 Within each post-processing iteration, cluster merging serves as an adaptive consol-
399 idation step, reducing over-fragmentation as the clustering structure evolves. After
400 outlier elimination, some clusters become more visually coherent, and new clusters
401 may emerge from outlier grouping. These updates can reveal opportunities to combine
402 clusters that now exhibit strong visual similarity.

403 As shown in Fig. 4, we compute the average cosine similarity between the embed-
404 ding centroids of all cluster pairs. If the similarity between two clusters exceeds a
405 threshold determined through hyperparameter tuning (see Appendix A), they are
406 merged into a single group. Performing this merging iteratively, rather than in a single
407 final pass, ensures that decisions are made on the most up-to-date cluster compositions,
408 accounting for changes introduced by outlier removal, grouping, and reassignment.

409 This incremental approach prevents premature large-scale merges that risk col-
410 lapsing meaningful fine-grained distinctions, while still allowing the framework to
411 consolidate redundant clusters as they appear. Over successive iterations, the result
412 is a set of visually coherent, non-redundant clusters that better reflect the ecological
413 structure of the dataset. By reducing spurious fragmentation and redundancy, these
414 refinements directly improve the accuracy of abundance distributions, which in turn
415 leads to more reliable Shannon and Simpson diversity estimates.

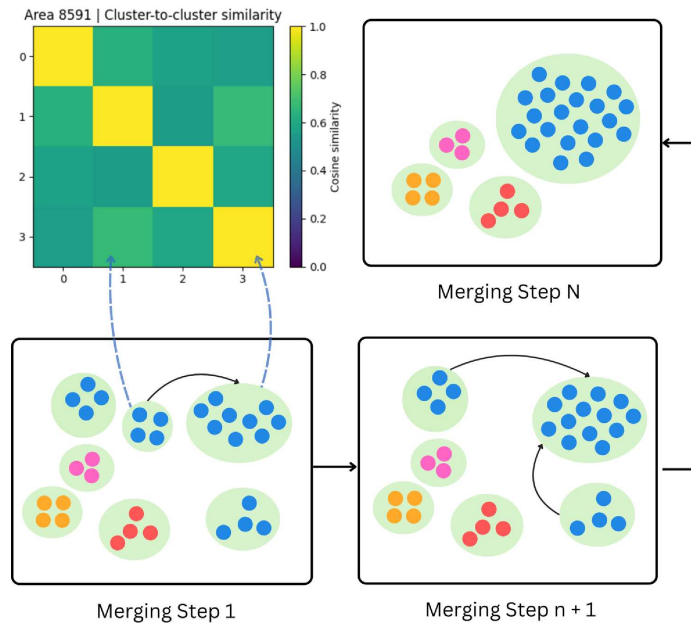


Fig. 4: Iterative merging of visually similar clusters, guided by centroid similarity, to consolidate over-fragmented groups.

416 5 Conclusion

417 This study introduced an unsupervised clustering framework for quantifying urban forest diversity directly from street level imagery, capturing both evenness and dominance based diversity indices. Extensive hyperparameter tuning and ablation experiments revealed that combining outlier elimination with cluster merging provided the best trade off between clustering quality, richness accuracy, and preservation of diversity structure, while additional reassignment steps offered only marginal benefits. Across diverse urban contexts, the method achieved low Wasserstein distances for both Shannon and Simpson diversity, indicating strong agreement with ground truth even in cities with highly imbalanced genera distributions.

426 Importantly, the framework preserved the spatial organization of diversity, as reflected by minimal differences in Moran's I values for Shannon and Simpson indices across cities. This ability to recover both the magnitude and spatial structure of key biodiversity metrics underscores the method's potential as a scalable monitoring tool for urban tree communities. While challenges remain including uneven imagery availability, occlusion from privacy blurring, and environmental variability, these results demonstrate the promise of computer vision based approaches for large scale, fine grained biodiversity assessment and their potential integration into urban forestry policy and planning.

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446 **5.2 Conflict of interest**

447 The authors declare no competing interests.

448 **5.3 Ethics approval and consent to participate**

449 Not applicable.

450 **5.4 Consent for publication**

451 Not applicable.

452 **Data availability**

453 All data supporting the findings of this study are available within the paper and its
454 Supplementary Information. The code used for analysis is available at <https://github.com/Diaa340/Releaf/>.

456 **5.5 Materials availability**

457 Not applicable.

458 **5.6 Author contribution**

459 M.M., F.D., I.Z., M.S., and D.A.A. conceptualized the study. D.A.A. and M.S.
460 performed the methodology development and data curation, conducted the formal
461 analysis, and wrote the original draft. M.M., F.D., and I.Z. supervised the research,
462 C.R. provided resources, and handled funding acquisition. All authors reviewed and
463 edited the manuscript.

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