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Tree Species Traits Determine the Success of LiDAR-Based Crown Mapping in a Mixed Temperate Forest

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Abstract: The ability to automatically delineate individual tree crowns using remote sensing data opens the possibility to collect detailed tree information over large geographic regions. While individual tree crown delineation (ITCD) methods have proven successful in conifer-dominated forests using Light Detection and Ranging (LiDAR) data, it remains unclear how well these methods can be applied in deciduous broadleaf-dominated forests. We applied five automated LiDAR-based ITCD methods across fifteen plots ranging from conifer- to broadleaf-dominated forest stands at Harvard Forest in Petersham, MA, USA, and assessed accuracy against manual delineation of crowns from unmanned aerial vehicle (UAV) imagery. We then identified tree- and plot-level factors influencing the success of automated delineation techniques. There was relatively little difference in accuracy between automated crown delineation methods (51–59% aggregated plot accuracy) and, despite parameter tuning, none of the methods produced high accuracy across all plots (27—90% range in plot-level accuracy). The accuracy of all methods was significantly higher with increased plot conifer fraction, and individual conifer trees were identified with higher accuracy (mean 64%) than broadleaf trees (42%) across methods. Further, while tree-level factors (e.g., diameter at breast height, height and crown area) strongly influenced the success of crown delineations, the influence of plot-level factors varied. The most important plot-level factor was species evenness, a metric of relative species abundance that is related to both conifer fraction and the degree to which trees can fill canopy space. As species evenness decreased (e.g., high conifer fraction and less efficient filling of canopy space), the probability of successful delineation increased. Overall, our work suggests that the tested LiDAR-based ITCD methods perform equally well in a mixed temperate forest, but that delineation success is driven by forest characteristics like functional group, tree size, diversity, and crown architecture. While LiDAR-based ITCD methods are well suited for stands with distinct canopy structure, we suggest that future work explore the integration of phenology and spectral characteristics with existing LiDAR as an approach to improve crown delineation in broadleaf-dominated stands.

Keywords: LiDAR; individual tree crown delineation (ITCD); temperate forest; tree architecture

1. Introduction

Individual tree crown delineation (ITCD) via remote sensing platforms offers potential to obtain detailed tree inventory/information over large areas. ITCD has been used to map species [1], biodiversity [2], and carbon stocks [3], as well as to quantify tree structural [4] and spectral characteristics [5]. While manually delineating crowns from high resolution imagery provides accurate measurements for small scale studies [5,6], effective automated methods are necessary if efforts are to be scaled to larger geographic regions. An ideal crown delineation method would be broadly applicable across stands varying in structural and compositional complexity. Given that many forests across the globe are under increasing pressure from climate change [7], invasive pests [8], and land-use change [9], reliable methods for measuring and mapping forests takes on additional urgency. Yet, broad-scale application of automated ITCD techniques remains difficult and reliability is uncertain.

Considerable work has been done to develop and improve automated ITCD techniques [10–15]. Light Detection and Ranging (LiDAR) crown delineation methods tend to be favored over spectral methods because they are not impaired by shadow and illumination artifacts [16], and because of their ability to directly measure crown architecture [17]. However, reported accuracies of different LiDAR-based methods varies considerably [12].

The success of ITCD is directly linked to the architecture of individual tree crowns and their position relative to neighboring crowns [18]. Crown architecture controls leaf display [19], and trees must balance resource acquisition (e.g., light capture) with mechanical constraints (e.g., buckling under their own weight [20]). Tree crown shape is also plastic [21–23] and crown shape is affected by competitive interactions with neighboring trees [24,25], as well as site history and disturbance [23,26].

Despite the seemingly stochastic nature of crown and stand structural development, there are also predictable differences between needle-leaf evergreen (conifer) and deciduous broadleaf plant functional types that influence ITCD. Conifer and broadleaf species differ in their physiological traits and adaptations to acquire resources [27,28] that manifest in differences in crown shape and stand arrangement. LiDAR-based ITCD methods have been successfully applied in conifer-dominated systems [29–31], while broadleaf-dominated systems have proven to be more challenging [17,32]. Discrepancies in accuracy of ITCD methods between conifer and broadleaf systems is often attributed to the characteristic plagiotropic growth form (ellipsoidal or umbrella-shape) of broadleaf crowns that make it difficult to identify tree tops, differentiate neighboring crowns, and group split canopies of an individual crown, such as those arising from forked trees [12].

Despite the challenges, there is a need for ITCD in many regions dominated by broadleaf and mixed stands. The temperate forests of the northeastern United States are typically dense mixed species stands with closed canopies, where crowns often overlap and have irregular shape. It remains unclear the degree to which automated ITCD techniques can be employed in the region, or what factors influence the success of these automated techniques. Here, we applied five automated LiDAR-based ITCD methods in a mixed temperate forest across a gradient of plots ranging from conifer- to broadleaf-dominated. We identified tree-level factors related to tree size and crown shape and plot-level factors related to vertical and horizontal structural and compositional complexity that influenced the outcome of our delineation analysis. We discussed how the architecture of individual trees and stand arrangement influences automated crown delineation results. Finally, we commented on how understanding the ecology of conifer and broadleaf species might best be exploited to delineate trees in temperate forests.

2. Materials and Methods

2.1. Study Site Description

This study was conducted in a Smithsonian Forest Global Earth Observatory (ForestGEO) MegaPlot [33] at the Harvard Forest, in north-central Massachusetts (42°32′ N, 72°11′ W). Located within the Prospect Hill Tract of Harvard Forest, the 35-ha MegaPlot is structurally and compositionally

representative of mature central New England forests. It encompasses a continuous forest comprised of mature eastern hemlock (*Tsuga canadensis*) stands, mixed-broadleaf stands, remnant red pine (*Pinus resinosa*) plantations, and a 3-ha swamp [34]. The age structure is dominated by 75–125 year old second growth forest [35]. Dominant species include red oak (*Quercus rubra*), red maple (*Acer rubrum*), eastern hemlock, and white pine (*Pinus strobus*). Other common species include Norway spruce (*Picea abies*), American beech (*Fagus grandifolia*) and birch (*Betula* spp.). There is little terrain variation in the MegaPlot, and elevation ranges from 340.2 to 367.8 meters above sea level. Soil are primarily moderately- to well-drained loam formed from glacial till [36].

Between 2010 and 2014 a census of the MegaPlot was conducted, where all woody stems ≥ 1 cm diameter at breast height (DBH) were mapped, measured, and identified to species [34]. The height of all stems was estimated using site-specific allometric equations [37]. For the present study, we used fifteen 20 m radius subplots located across the MegaPlot (Figure S1). Plots were selected to capture a full range of tree functional composition from conifer-dominated to broadleaf-dominated.

2.2. Remote Sensing Data

LiDAR and hyperspectral data were collected over Prospect Hill by NASA's Goddard LiDAR, Hyperspectral and Thermal (G-LiHT) sensor package [38] between 19–21 June 2012. LiDAR point cloud, canopy height model (CHM), and hyperspectral data were downloaded on 22 October 2018 (https://glihtdata.gsfc.nasa.gov/), and clipped to a 10 m buffered extent of the MegaPlot. The LiDAR point cloud has an average density of 26.98 points per m^2 within the MegaPlot. The hyperspectral and LiDAR CHM data have a spatial resolution of 1 m. Although LiDAR data from NEON AOP [39] were collected in 2017, we opted to use the 2012 G-LiHT data set because of its higher point density and because changes in the canopy since 2012 were minimal.

Aerial surveys of the MegaPlot were conducted throughout the 2018 growing season by an RGB camera-equipped DJI Phantom 4 Pro unmanned aerial vehicle (UAV) equipped with a stock 20 megapixel sensor. The UAV followed a pre-programmed flight path 100 m above ground level at 20 km/h, obtaining a side overlap of 70% and a forward overlap of 80% of the images. The orthomosaic, generated using AgiSoft PhotoScan, had a spatial resolution of 0.025m but was down sampled to 0.1m for use in this analysis. Using the *georeferencer* plugin in QGIS (v. 2.18; [40]), UAV imagery was registered to the G-LiHT remote sensing data by identifying distinguishable features (e.g., emergent white pine crowns, MET towers, wetland boundaries) in both the UAV imagery and the hyperspectral imagery. Each UAV image was aligned with the G-LiHT imagery with 20 control points and transformed using a first-order polynomial. The resulting georeferenced UAV images were found to be in good visual agreement and tree crowns aligned with those visible in the G-LiHT hyperspectral and LiDAR imagery and field-measured stem locations. We further inspected the datasets and confirmed that there was no tree dieback or notable changes to the forest canopy between G-LiHT and UAV imagery collection.

2.3. Crown Delineation

All tree crowns visually distinguishable within the fifteen plots were manually delineated by onscreen digitizing of the UAV image collected on September 13th. This study excluded understory crowns not visible within UAV imagery. Manual delineation of individual tree crowns (M_{ITC}) was done with a stylus pen using the *FreehandEditing* plugin in QGIS. While crown digitization across all plots was performed on the September 13th image for consistency, multiple dates of imagery (September 13th, October 5th, October 12th, and November 4th) were used to help distinguish crowns and identify the species of each crown based on differences in shadow and phenology (Figure 1).

We manually delineated 650 tree crowns from 14 unique species. M_{ITC} species label and associated stem attributes (DBH and allometrically derived tree height) were manually assigned during the digitization process from the ForestGEO stem data. In rare cases where a crown could conceivably belong to one of multiple stems from either the same species or stems from different species that could not be distinguished using phenology and textural cues, the crown was assigned to the stem with the

taller allometrically derived tree height. Crown area and maximum CHM-derived crown height were calculated for each M~~r~~. Using M~~r~~ crowns, conifer fraction of each plot was calculated as the ratio of conifer crown area to broadleaf crown area.

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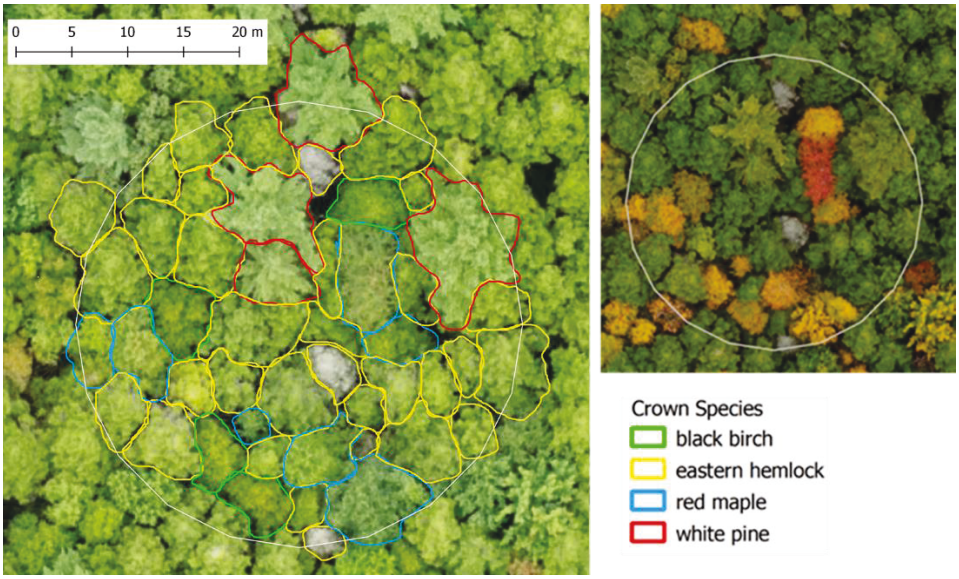

Figure 1. Manual crown delineation was performed using 2018 high resolution UAV imagery. All delineations were done on the September 13th image (left panel), but other dates of imagery were used to help differentiate crowns growing in close proximity. The right panel (October 12th) gives an example of phenologic differences between species that can be leveraged to help separate crowns that might otherwise be clumped during manual interpretation.

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We tested five automated individual tree crown (ITC) delineation techniques (Table 1) available in the R (v. 3.5.1; [41]) lidR package [42]. Four routines are surface-based methods applied to a rasterized CHM, and the fifth is a 3D method applied to a LiDAR point cloud. Dalponte2016 (DALPONTE) is a surface-based seed and region growing method [43]. Silva2016 (SILVA) is a surface-based seed and watershed segmentation method [30]. Simple Watershed (SWS) [44] is a surface-based watershed segmentation [44]. Marker-controlled Watershed (MCWS) is a watershed segmentation that relies on a priority seed map. Li2012 (LI) is a 3D region growing method applied to a point cloud [29]. All techniques were run using the *lastrees* function. Treetop priority seed points used in DALPONTE, SILVA, and MCWS were created with the *lmf* (local maximum filtering) algorithm [45]. SWS did not rely on a priority seed map and LI has a tree top detection built into the function. Final ITC polygons were generated using the *tree_hulls* function, by creating a 2D concave hull around the segmented point cloud. We chose not to smooth CHM data (e.g., Gaussian filtering) prior to crown delineation as our preliminary results showed, on certain cases, decreased overall accuracy of the methods.

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Table 1. Five automated LiDAR-based individual tree crown delineation routines were evaluated in this study. † Four routines are surface-based methods applied to rasterized canopy height models. ‡ The fifth routine is a 3D method applied to a point cloud. All routines were implemented in the R package lidR, developed by Roussel and Auty [42].

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Crown Delineation Routine	Reference
Marker-controlled Watershed (MCWS) †	[43]
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2.4. Parameter Tuning and Accuracy Assessment

To apply each crown delineation method, we optimized parameters against manually delineated crowns. Each automated delineation technique has different input parameters controlling how the algorithm searches and delineates the CHM or point cloud, and methods vary in input parameter complexity. Parameters include search window sizes, maximum height or radius values and drop-off thresholds. We first applied each automated delineation technique with default parameters. We specified a 3 × 3 moving window for the *lmf* tree top detection during default parameterization because a default parameter was not given.

We then tuned each technique's input parameters to find (1) the best plot-tuned parameters—potentially unique parameters that maximized accuracy for each individual plot and (2) the best generalized parameters—a single set of parameters that achieved the highest accuracy when evaluated across all 15 plots. Parameter tuning was done using a bootstrapping approach, where input parameters were randomly selected within a predefined range (Tables S1 and S2). Following each delineation iteration, accuracy was assessed by comparing the generated A_{ITC} polygons to the reference M_{ITC} polygons. Automated delineations were paired to manual delineations (details provided in supplemental material) so that any given M_{ITC} was labeled as either correctly or incorrectly delineated. A detection accuracy score (DA) was assigned to each iteration:

$$DA = \frac{n_{TP}}{N}, \quad (1)$$

where, n_{TP} is the number of correctly delineated A_{ITC} and N is the number of M_{ITC} [46]. A given A_{ITC} was considered correctly delineated (true positive) if ≥ 50% of the area of both A_{ITC} and M_{ITC} overlap (Figure 2; e.g., [47,48]). Each routine, except LI, was iterated 500 times. LI was only iterated 200 times because it was substantially slower than the surface-based methods and because maximum accuracy achieved did not improve beyond the first 100 iterations. We retained tuning iterations for the highest plot-tuned parameter accuracy and the highest generalized parameter accuracy (aggregated across the 15 plots) for each automated crown delineation technique.

To further understand how each method performed at the crown-level, we characterized the incorrect M_{ITC} delineations by type of error. Therefore, each crown was ultimately assigned to one of four categories based on their overlap with A_{ITC} (Figure 2):

1. Over-segmentation: The intersecting area between A_{ITC} and M_{ITC} is greater than or equal to 50% of the area of *only* A_{ITC}, indicating that the automated crown is smaller than manual crown.
2. True Positive: The intersecting area between A_{ITC} and M_{ITC} is greater than or equal to 50% of the area of *both* A_{ITC} and M_{ITC} (as defined above), indicating that the automated crown is well matched to the manual crown.
3. Under-segmentation: The intersecting area between A_{ITC} and M_{ITC} is greater than or equal to 50% of the area of *only* M_{ITC}, indicating that the automated crown is larger than the manual crown.
4. False Positive: The intersecting area between A_{ITC} and M_{ITC} is greater than or equal to 50% of the area of *neither* A_{ITC} and M_{ITC}, indicating poor matching of the automated and manual crowns.

Given that any M_{ITC} can only be linked to one A_{ITC}, in the case where multiple A_{ITC} fell within a single M_{ITC} (as can occur with over-segmentation), the M_{ITC} was assigned to the A_{ITC} that had the highest proportion of overlapping area with the particular M_{ITC}.

2.5. Statistical Analysis

To understand the factors that influenced automated crown delineation success, we evaluated tree-level variables (field-measured DBH, CHM-derived M_{ITC} crown height, and M_{ITC} crown area) and metrics of plot-level vertical and horizontal structural and compositional complexity (canopy complexity, uniformity of crown spacing, relative density, trees per plot, and species diversity). Plot-level metrics only included stem attributes associated with M_{ITC} data.

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Plot canopy complexity was estimated using the Rumple Index [49], a ratio of canopy surface area to projected ground area, describing the roughness of the canopy terrain surface. The horizontal spatial distribution of trees in each plot was calculated from Moran's *I* using an aggregation index (AGI) [50], which estimates the uniformity of spacing. Plot stand density was expressed using a mixed species relative density equation [51], as well as by using the number of trees per plot. We estimated the diversity of species on each plot using three diversity indices: Shannon's Diversity (*H*), Pielou's Evenness (*J*), and Species Richness [52]. All predictor variables were standardized to have a mean of zero and a standard deviation of one by subtracting the mean and dividing by one standard deviation [53].

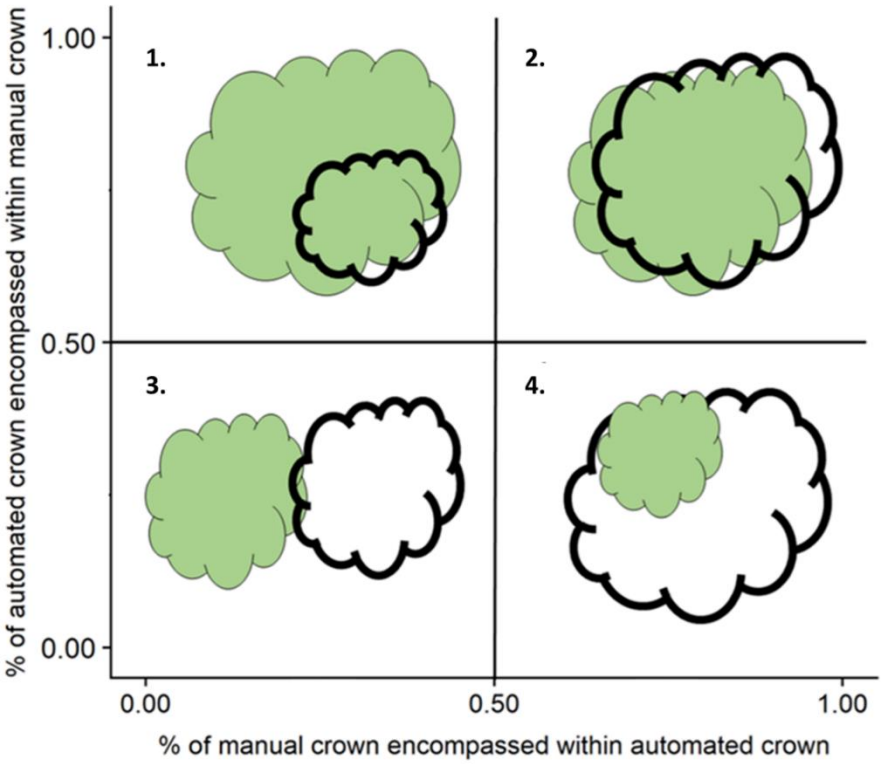

Figure 2. Automated crown delineations (A_{ITC}; shown with bold outline) were assessed against manual crown delineations (M_{ITC}; shown with green fill) and assigned into one of four categories based on overlapping area: (1) Over-segmentation: The intersecting area between A_{ITC} and M_{ITC} is greater than or equal to 50% of the area of only A_{ITC}. (2) True Positive: The intersecting area between A_{ITC} and M_{ITC} is greater than or equal to 50% of the area of both A_{ITC} and M_{ITC}. (3) Under-segmentation: The intersecting area between A_{ITC} and M_{ITC} is greater than or equal to 50% of the area of only M_{ITC}. (4) False Positive: The intersecting area between A_{ITC} and M_{ITC} is less than 50% of the area of both A_{ITC} and M_{ITC}.

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To identify important plot-level variables we performed univariate linear regressions between plot-level metrics and plot-tuned accuracy ($n = 15$) for all five crown delineation routines. We then built global multiple linear regression models including all significant ($\alpha < 5\%$) variables from the univariate analyses. Multicollinearity of the global model was evaluated using variance inflation factor (VIF), and we sequentially removed highly inter-correlated variables until VIF of all variables was <10 [54]. The best model for plot-level performance was chosen using a corrected Akaike Information Criterion (AICc) to account for small sample size [55].

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Finally, we built mixed-effect logistic regressions to help understand which of the tree- and plot-level factors influenced agreement of A_{ITC} with M_{ITC} as a linear function of covariates in a logistic regression [56]. Logit models were built in the R package lme4 [57]. Each global model included tree-level variables and the plot-level variables found to be significant during the univariate analyses described above. We controlled for plot-level variability by including plot as a random effect in each model. Model selection was performed by backward elimination from the global model, and the final

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Model accuracies were evaluated using a 10-fold cross validation, where the developed logistic relationships were each trained on 90% of the data and tested on the remaining 10%. Training and testing were performed on all 10 folds of data and the results were averaged to give an estimate of each model's accuracy.

3. Results

3.1. Manual Crown and Plot Characteristics

Of the 650 tree crowns we manually delineated, 379 were conifer crowns, and 271 were broadleaf crowns. The range in height, DBH, and crown area was comparable between conifer and broadleaf species. On average conifers were taller and had larger DBH (Error! Reference source not found.), while median conifer crown area was 27% smaller than broadleaf crowns.

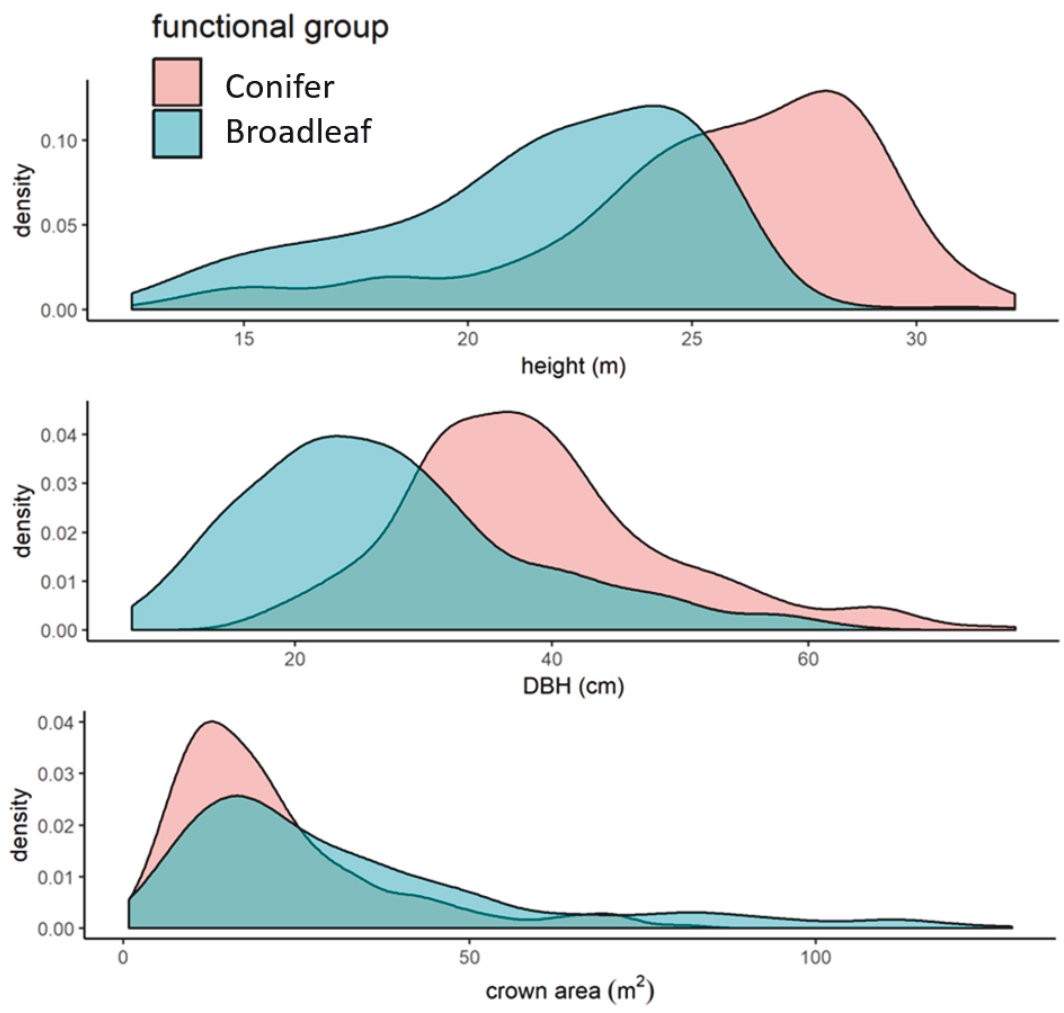

Figure 3. Density distribution of tree-level variables showing differences between conifer and broadleaf functional groups.

Trees per plot ranged from 25 to 66 across the 15 plots (mean 43), and structural complexity and composition varied substantially. Average plot canopy height ranged from 15.5 m to 28.9 m, and

average crown area ranged from 13.75 m^2 to 47 m^2. Conifer fraction, as estimated by crown area, ranged from 14% to 96%. Six of the 15 plots were characterized as conifer-dominated with ≥ 50% conifer fraction.

Plot-level characteristics showed varying relationships with conifer fraction (Figure S2). For example, there was a strong positive linear relationship between conifer fraction and rumple (r: 0.89; $p < 0.001$). Rumple ranged from 1.34 to 2.16, with conifer-dominated plots occupying the upper end of this range (1.69–2.16). Species evenness also had a strong linear relationship with conifer fraction (r: −0.84; $p < 0.001$), as did Shannon's Index (r: −0.76; $p < 0.001$). Conifer fraction showed a weak, but non-significant relationship with trees per plot (r: 0.41; $p = 0.12$) and AGI (r: 0.41; $p = 0.13$).

3.2. Automated Crown Delineation Accuracy

3.2.1. Influence of Parameter Tuning and Differences in Model Accuracies

The influence of generalized parameter tuning compared to default parameters varied by method (1–41%) (Table 2). While further plot-tuning of method parameters only marginally improved overall accuracy scores (+2–5%), we chose to continue the analyses using plot-tuned results because plot-level accuracy (Table S3) improved by as much as 36% (LI) and because we were interested in understanding the factors that influenced the highest quality delineations.

Following plot-tuning overall accuracy and plot-level accuracy did not vary substantially across delineation methods. Overall accuracy ranged from 51% by SWS to 59% by LI. Though LI was marginally more accurate (+4%) than the second highest overall accuracy (MCWS: 55%), it came at a substantial increase in processing time and complexity of input parameters (and necessarily required parameter tuning to achieve high accuracy).

Plot-level accuracy ranged from a low of 27% (DALPONTE and SWS; plot 11) to a high of 90% (MCWS; plot 14). The difference between the most- and least- accurately delineated plot was >40% for all methods. Accuracy of all methods varied similarly across all plots (Figure S3), and significantly related to conifer fraction ($p < 0.05$).

Table 2. Overall site accuracy (0-1 range) of five different automated crown delineation routines. The table included default, generalized, and plot-tuned parameters. ‡ Conifer and broadleaf accuracies are from plot-tuned model runs.

Routine	Default	Generalized	Plot-tuned	Conifer ‡	Broadleaf ‡
MCWS	0.49	0.53	0.55	0.65	0.40
SWS	0.08	0.49	0.51	0.54	0.46
DALPONTE	0.46	0.48	0.52	0.63	0.38
SILVA	0.48	0.49	0.54	0.67	0.39
LI	0.38	0.55	0.59	0.69	0.46

3.2.2. Differences in Accuracy across Species

All methods more accurately delineated conifer crowns (mean 64%) than broadleaf crowns (mean 42%). Each method had trade-offs in accuracy at the species level, and no single method stood out has having the highest accuracy across all species (Figure 4). For example, SILVA delineated red pine especially well (81%), but had consistently low broadleaf accuracy scores. Alternatively, SWS had the lowest red pine accuracy (53%) but excelled at delineating red oak in comparison to other methods (+9%).

3.3. Variables Influencing Accurate Automated Crown Delineation

3.3.1. Linear Regressions

Five plot-level variables (J, trees per plot, rumple, H, and AGI) were found to be significant ($p < 0.05$) in at least one univariate regression (Figure 5). All five variables were initially included in

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Five plot-level variables (J, trees per plot, rumple, H, and AGI) were found to be significant ($p < 0.05$) in at least one univariate regression (**Error! Reference source not found.**). All five variables were initially included in the multivariate global model. However, trees per plot and AGI were highly correlated ($r = 0.97$), and H and J were highly correlated ($r = 0.94$). To constrain VIF values to below 10, H and trees per plot were removed from the analysis. For each of the five multivariate regression analyses, the model including only J as an independent variable was selected as the best model. This relationship between accuracy and J was significant ($p < 0.05$) for all models (Table S4).

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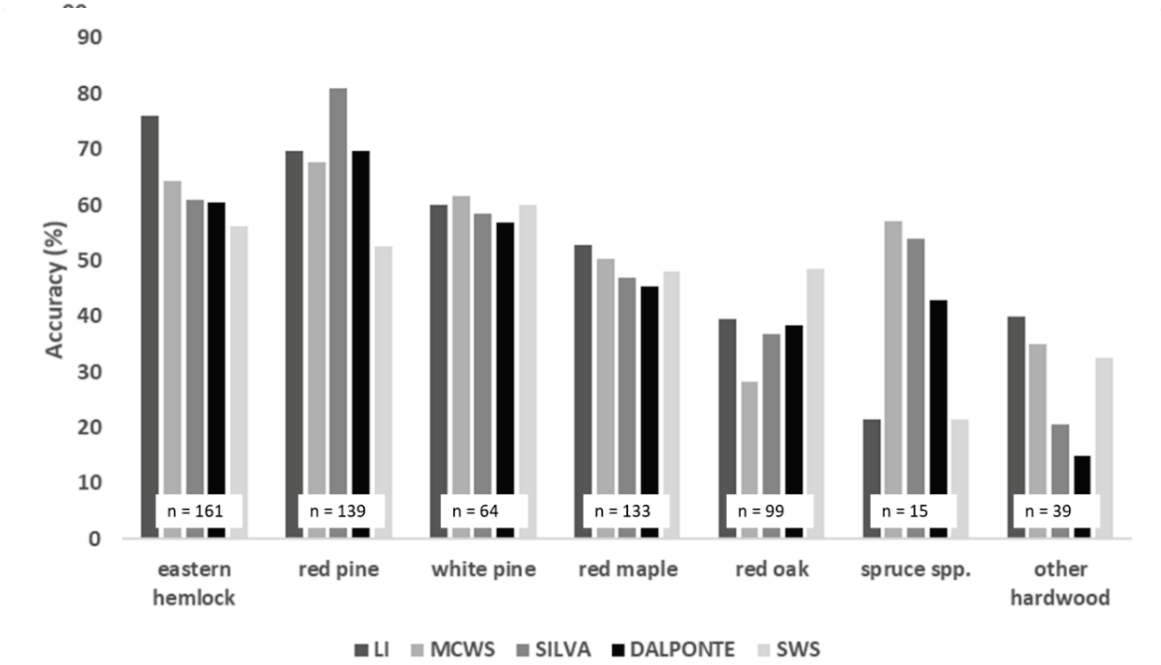

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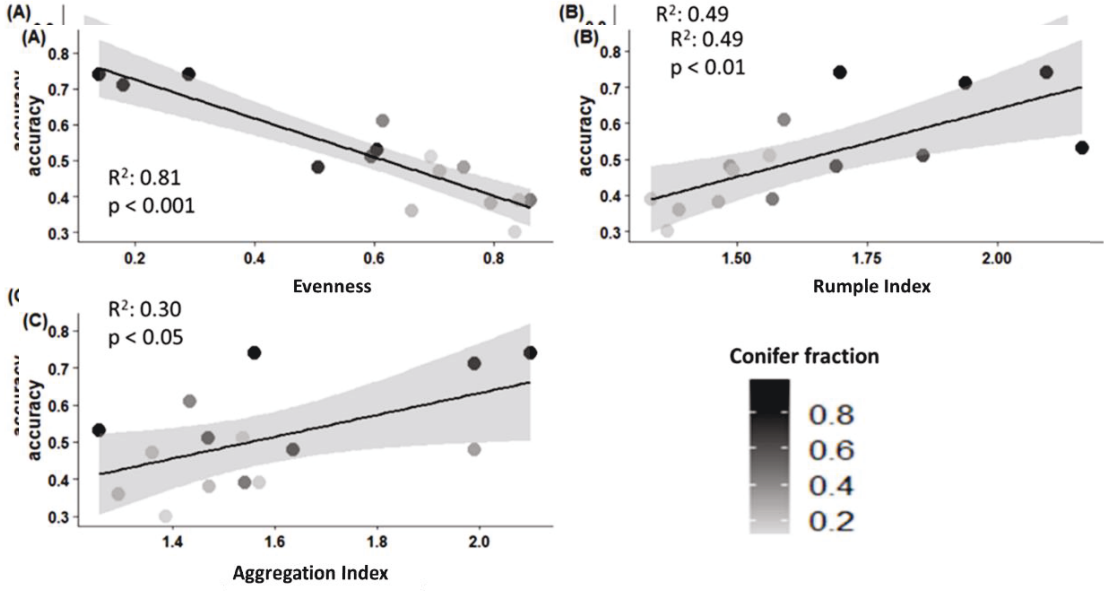

Figure 5. Prediction accuracy in relation to plot-level (**a**) evenness, (**b**) rumple index, and (**c**) aggregation index for one crown delineation method (DALPONTE). Points are colored to show fraction of conifer crown area per plot (conifer fraction). The relationship between accuracy and evenness was significant ($p < 0.05$) across all methods. However, the relationship between accuracy and rumple index was only significant for DALPONTE, SILVA and LI.

Figure 5. Prediction accuracy in relation to plot-level (**a**) evenness, (**b**) rumple index, and (**c**) aggregation index for one crown delineation method (DALPONTE). Points are colored to show fraction of conifer crown area per plot (conifer fraction). The relationship between accuracy and evenness was significant ($p < 0.05$) across all methods. However, the relationship between accuracy and rumple index was only significant for DALPONTE, SILVA and LI.

3.3.2. Logistic Regressions

Global logit models consisted of tree-level variables (DBH, height, and crown area) and the plot-level variables (rumple, J and AGI) identified in the linear regression analyses. Results of the final logit models are shown in Table 3. Cross validation model accuracy ranged from 61% (MCWS) to 70% (SWS), suggesting that while we captured the most impactful variables in predicting crown delineation, there are additional factors unaccounted for. There was no single variable in the logit models that was included across all five ITCD approaches. However, all but one model (SWS) consisted of at least one tree-level variable related to tree size and one plot-level variable related to tree arrangement.

Table 3. Results of the logit models assessing the important tree- and plot-level variables influencing the odds of successful individual tree crown delineation. All variables were standardized prior to analyses. The table includes the 10-fold cross-validation (CV %) model accuracy estimates, coefficients of variables in the models and the corresponding standard error (SE) (* $p < 0.05$, ** $p < 0.01$, *** $p < 0.001$).

Routine	Variable	CV %	Coefficient	SE (Coef)	Z Value	*p*-Value	
MCWS		60.77					
	Intercept		0.19	0.16	1.16	0.25	
	Crown Area		−0.38	0.11	−3.41	0	***
	DBH		0.42	0.14	3.06	0	**
	Height		0.39	0.28	1.39	0.16	
	J		−0.68	0.26	−2.64	0.01	**
	Rumple		−0.49	0.26	−1.89	0.06	.
SWS		70					
	Intercept		−0.1	0.33	−0.3	0.76	
	Crown Area		0.76	0.12	6.39	0	***
	Height		1.25	0.25	4.91	0	***
DALPONTE		62					
	Intercept		0.12	0.09	1.35	0.18	
	AGI		0.36	0.12	2.88	0	**
	DBH		0.6	0.11	5.51	0	***
	J		−0.25	0.12	−2.11	0.03	*
SILVA		65.32					
	Intercept		0.27	0.2	1.33	0.18	
	AGI		0.47	0.21	2.21	0.03	*
	Height		1.04	0.24	4.37	0	***
LI		61.54					
	Intercept		0.41	0.11	3.78	0	***
	Crown Area		−0.2	0.1	−1.97	0.05	*
	DBH		0.5	0.11	4.39	0	***
	J		−0.33	0.12	−2.75	0.01	**

All tree-level variables were important, each showing up in three of the five logit models, though not always together. Height and DBH always had a positive effect on successful crown delineation. Crown area had a negative effect in two models, but it had a positive effect on delineation for the SWS method.

Plot-level variables were not consistently important across methods. Species evenness was the most important, showing up in three models, and each time having a negative effect on crown delineation accuracy ($p < 0.05$). AGI was included as positively affecting delineation in two models ($p < 0.05$), while rumple was only included in one model and negatively affected delineation odds ($p = 0.06$).

4. Discussion

4.1. Differences between Segmentation Methods

Automated crown delineation shows promise but remains difficult to apply in closed canopy mixed species forests. Despite methodological differences between segmentation routines, we found similar patterns in accuracy and in which crowns were best delineated. Even with parameter tuning, none of the methods produced high overall accuracy (51–59%), and the variation in accuracy across plots was similar for each method (i.e., the most and least accurate plots identified by each method were generally the same). Furthermore, all methods better delineated conifer crowns than deciduous crowns.

We found that the point cloud segmentation method (LI) only marginally outperformed the CHM-based segmentation methods. This may be because our validation method (manual delineation of UAV imagery) allowed for accurate surveying of canopy trees but did not attempt to address understory trees. Point cloud-based methods outperform CHM-based methods in detection and delineation of understory and suppressed crowns [58], but there is less of an advantage for delineation of canopy trees.

Emerging point cloud segmentation methods have taken novel approaches (e.g., Multiclass Graph Cut [15]; Mean Shift [59]) that may or may not be better suited to delineate closed-canopy temperate forests with difficult deciduous crown architecture. However, superior performance of point cloud segmentation methods is likely to be largely limited by availability of LiDAR data with high measurement density [60]. An appeal of exploring CHM-based segmentation is the wide and growing availability of high-quality CHM data, which can make these techniques more broadly applicable than point cloud segmentation approaches.

One reason that methods that relied on an external tree detection algorithm (DALPONTE, SILVA, and MCWS) all performed similarly may be because we used the same tree detection algorithm (local maximum filtering) for each method. Given that we found such little variation between these CHM-based segmentation methods, this may suggest that the choice of tree detection method (e.g., [61]) is of equal or greater importance than the choice of segmentation method (e.g., region growing vs. watershed).

Similarities in crown delineation patterns indicate that the ability to delineate crowns is driven by physical canopy traits across plots rather than by methodological differences in crown delineation algorithms. Conifer and broadleaf species have evolved traits that distinguish their ability to compete for resources and respond to disturbance and competition. In turn these traits influence tree height, crown architecture (crown spreading and leaf-display), and how crowns interact with neighboring crowns.

4.2. Tree Architecture

4.2.1. Tree Size

Taller trees and larger DBH trees were more likely to be correctly delineated using automated methods. This is in part because large trees often hold dominant positions in the canopy and tend to have more symmetrical crown shape [21]. Yet, this is also because conifers in the canopy tended to be taller and have larger DBH (Figure 3). Conifer species identified in the plots have lower average wood density than the broadleaf species [51,62]. Lower wood density is energetically efficient for height growth [63,64], while resulting in lower stand density contributions of individual trees [51]. In our study region, in higher diameter size classes, conifer species diverge from broadleaf species, growing taller. This divergence is associated with both lower wood density and evergreen foliage [65].

Conifers, especially white pine, are larger (DBH and height) at Harvard Forest because of site history and growth strategy. Much of the northeastern United States landscape has been shaped by historical land use [66]. White pine are successful colonizers on disturbed sites, and many of the large white pines are old-field pines that invaded agricultural and pastoral fields following abandonment in the mid-1800s [67]. Low density wood, and relatively high photosynthetic rates [27,63] allow white

pine to achieve rapid vertical growth, and they continue to avoid direct competition by occupying a higher canopy stratum than broadleaf species. On a canopy height model, emergent white pines appear as hotspots (Figure S4) because they often stand five or more meters above the continuous canopy; thus, they are easily detected and delineated by automated crown delineation methods.

4.2.2. Crown Spread

Smaller diameter crowns were more likely to be successfully delineated which, as was the case with height, is likely related to differences between conifer and broadleaf traits. Conifer crowns tend to spread less than broadleaf crowns, though it is possible to find white pine or hemlock that are comparable in spread to broadleaf crowns. However, conifers maintain a more rigid, apically controlled, growth form and are less likely to exhibit crown plasticity [68]. This results in a singularly defined orthotropic bole and the characteristically conical crown shape, and it is far rarer to find conifers with forked trunks and split crowns. This may be because many conifers have evolved in resource poor conditions, and invest in long-lasting low-nitrogen (N) foliage [69]. Convergent leaf and canopy structural properties (conical crown shape, clumped foliage) promote light scattering and more even/diffuse light conditions throughout the canopy, which in turn increases radiation use efficiency. Ollinger [70] pointed out that plants grown (or adapted to grow) in resource poor conditions allocate fewer resources to wood compared to foliage, constraining crown spread.

In contrast to conifers, broadleaf species invest in high-N deciduous foliage, with higher photosynthetic capacities and shorter leaf lifespans. To benefit from this strategy, broadleaf species must maximize direct light interception. Intermediate- and shade-tolerant broadleaf species (e.g., red oak and red maple) achieve this by spreading their crowns to maximize foliage display on a more even plane. Thus, many broadleaf species have weak apical control that results in plagiotropic growth forms [71]. Weak apical control allows multiple stems to compete for a dominant terminal position, the result of which can be a broad and flat crown, often with forked trunks and multiple differentiated sections within a single crown (i.e., crown splitting).

The ability to spread branches laterally is also positively associated with wood density. Higher wood density is correlated with structural properties, including resistance to splitting, rupture stress, dynamic breakage, and increased elasticity [20]. While low density wood is a lower carbon-cost approach to attain vertical expansion, broadleaf species with denser wood can expand lateral branching without compromising structural integrity [63,64]. This is in agreement with crown radius - DBH allometric equations developed by Sullivan et al. [37] at the Harvard Forest. They found the slope of the crown radius - DBH relationship to be steeper for broadleaf than conifer species, and that this relationship was related to wood specific gravity.

Red oak, in particular, often have substantial crown spread and split crowns. This type of architecture presents two major challenges for automated tree crown delineation: (1) It is difficult to define a singular local maximum (tree detection) and (2) crowns either interdigitate with neighboring crowns—resulting in under-segmentation, or crowns split—resulting in over-segmentation. We found all methods most often over-segmented red oak (Figure 6). These results agree with other studies that found broadleaf canopies are often over-segmented [17].

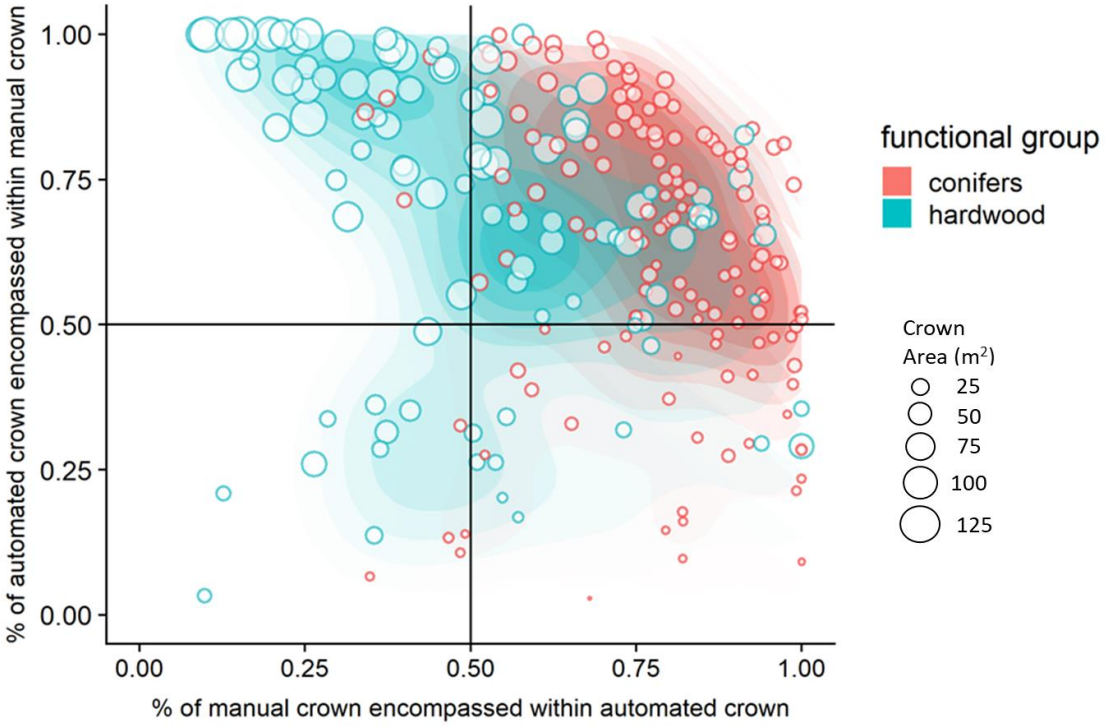

Figure 6. Two-dimensional density plot showing different patterns of crown delineation accuracy between conifer and broadleaf functional groups. The circles provide crown size comparison for two end member species: red pine (shown in red circles) and red oak (shown in blue circles). This figure corresponds with delineation categories described in **Error! Reference source not found.**: the top right quadrant signifies true delineations, the top left signifies over-segmentation, the bottom right signifies under-segmentation, and the bottom left signifies false positive. This figure shows data generated using the DALPONTE method, although all methods produce similar patterns.

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4.2.3. Mechanical Interactions

Mechanical interactions between neighboring crowns is another major dynamic controlling lateral branch expansion, perhaps even more than resource competition [72]. Crown shyness—gaps that form between adjacent crowns, often of the same species—can result from mechanical bud abrasion and branch damage during crown collisions [73]. While mechanical interactions occur between all adjacent crowns in closed-canopy stands, crown shyness is controlled by branch fragility and rates of regrowth following lateral branch damage [72].

Crown shyness is a common occurrence in even-aged conifer-dominated stands [74]. Crown shyness is especially visible in red pine dominated plots (Figure S5) located in an even-aged remnant pine plantation [75]. Shyness likely contributed not only to the high accuracy in these plots (as high as 80%), but also the fidelity of the delineations because gaps between adjacent crowns create defined borders for delineation (Figure 6). In comparison to broadleaf species with strong, dense branches (e.g., red oak), red pine is more susceptible to collision damage. High height:diameter ratios coupled with low wood density make the crowns of red pine susceptible to wind damage [76] through increased crown mobility and resulting high-impact crown collisions [77] that maintain crown shyness.

4.3. Species Evenness

We found that species evenness was the most important plot-level variable associated with crown delineation success. As species evenness decreased, the likelihood of successful delineation increased. Evenness was likely important because of (1) its negative relation to conifer fraction, and (2) a relationship between evenness and canopy space filling efficiency.

There was a strong relationship between species evenness and conifer fraction (Figure S2); the least even plots had the highest conifer fraction, while the most even plots tended to have the lowest conifer fraction. It is important to note that two of the low evenness conifer plots were artificial in the sense that they are remnant red pine plantation [75], though red pine can grow naturally in monocultures. However, the other low evenness conifer plot was in a natural mature hemlock stand, a common occurrence in northeastern temperate forests. Hemlock stands often have low species evenness because of deep shade cast that precludes the establishment of less shade tolerant species

[78]. Further, conifer needles have high carbon to nitrogen ratios, resulting in lower soil fertility which can deter broadleaf growth [27,79].

In addition to the influence of conifer fraction, the evenness—accuracy relationship may also be reflective of increased efficiency of canopy space filling (i.e., crown packing) in higher diversity plots [22,80], which makes successful delineation more challenging. In low diversity stands, trees from the same species compete similarly for growing space [81], while in higher diversity stands niche partitioning and complementarity of crown architecture promote partitioning of resources [82,83] allowing for more efficient use of available canopy space [84,85]. As plot diversity increases crown packing increases, and it becomes increasingly difficult to differentiate neighboring crowns (**Error! Reference source not found.Error! Reference source not found.**).

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To further investigate the potential relationship between species evenness and crown packing, we calculated average plot NDVI [86] from the G-LiHT hyperspectral data as a proxy estimate of leaf area index and foliar density [87], assuming increased crown packing would be related to higher LAI. We found evenness is strongly related to NDVI ($p < 0.001$; r: 0.9). However, because NDVI is also related to conifer fraction [88], we performed a partial correlation test. After accounting for conifer fraction, NDVI was still positively correlated (r: 0.58) with species evenness, lending support to the idea that the evenness–accuracy relationship is both a result of conifer fraction and increased crown packing in higher diversity plots. Future work would benefit from a more detailed analysis exploring how crown delineation results are expressly affected by local neighborhood tree species diversity (e.g., [89]), and the interaction of neighboring crowns.

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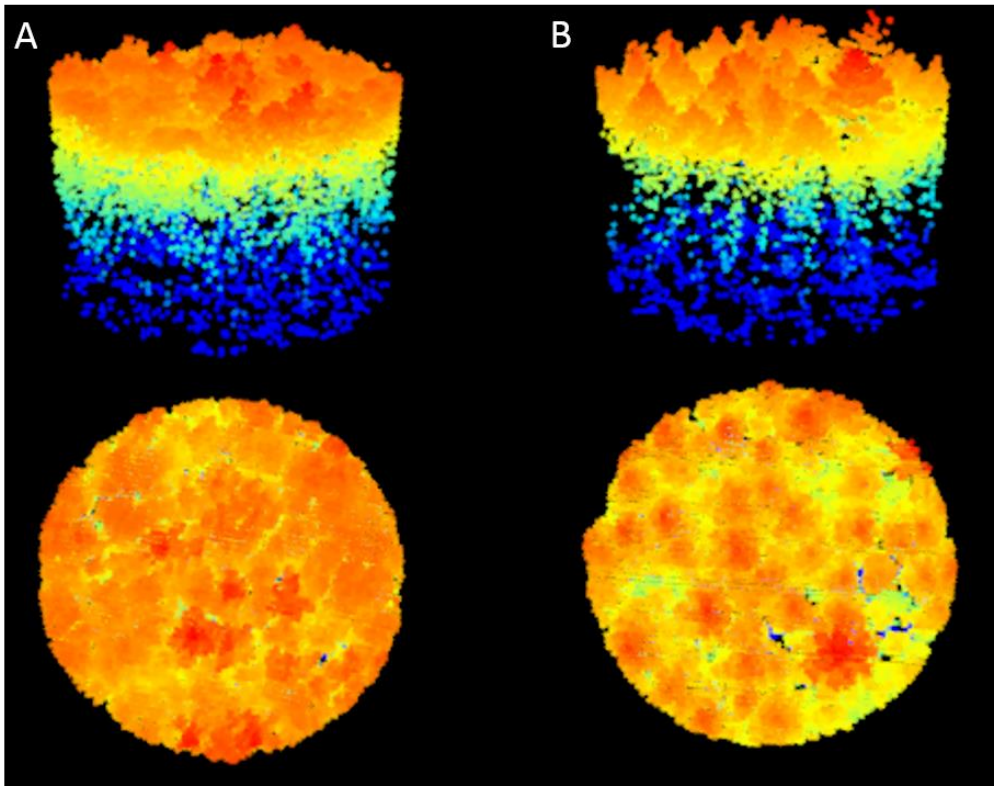

Figure 7. G-LiHT LiDAR point cloud comparison highlighting the differences in structure between a high-evenness broadleaf-dominated stand (**A**) and a low-evenness conifer-dominated stand (**B**). Warmer colors represent higher points in the canopy. The conifer-dominated stand exhibits higher canopy rumple, and uniformity of crown shape. The conical, less-plastic shape of conifer crowns may also reduce canopy space filling efficiency.

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4.4. Silver Linings: Where Do Automated Crown Delineation Methods Work Well?

Our findings show that automated LiDAR-based ITCD methods have substantial promise for delineation of large trees. Despite lower accuracy for smaller size trees, these results are encouraging

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4.4. A Silver Lining: Where Do Automated Crown Delineation Methods Work Well?

Our findings show that automated LiDAR-based ITCD methods have substantial promise for delineation of large trees. Despite lower accuracy for smaller size trees, these results are encouraging given the important role large trees play in terrestrial ecosystems [90], especially in terms of carbon accumulation [91,92]. We were able to successfully delineate 62–70% of all trees ≥ 40 cm DBH, which is promising for the prospect of tree-centric carbon mapping [3,43].

We also found these methods to perform especially well in conifer-dominated stands. High accuracy mapping of conifers has important implications for forest management and conservation. In particular, the ability to delineate mature eastern hemlock could provide important data for monitoring and mapping hemlock woolly adelgid (HWA) infestations [93]. Given the impact HWA has on the structure and composition of infested forests [94], existing crown delineation methods could aid in measuring and mapping HWA impacts.

Much of the northeast United States is comprised of aggrading second growth forest [66]. However, while our plots cover a range of structure and composition, they are undoubtedly still just a sample of the different forest types found across the northeast. LiDAR-crown delineation methods are likely to show varying degrees of accuracy based on additional factors influencing structure, such as stage of forest succession [95]. Relatively young stands in stem-exclusion stage (*sensu* [81]) are likely to be especially difficult to delineate because of high-stem density and intense competition. Given that tree size [96], stand structural complexity [97], and canopy surface complexity [98] often increase with stand age our results suggest that mature- and old-growth stands could be delineated with high accuracy.

4.5. Moving forward

While structurally distinct conifers stands are ideally suited for LiDAR-based crown delineation, these methods may not be ideal for delineating broadleaf canopies due the tendency of broadleaf crowns to be irregularly shaped, have forked trunks, and overlap with neighboring crowns. Broadleaf species often lack the distinctive crown architecture of conifers that allows for higher accuracy LiDAR-based crown delineation.

An alternate path for improving crown mapping of mixed- and deciduous-dominated stands may lie within the species-specific phenological and spectral traits of broadleaf species. Indeed, much of the information we relied upon to manually delineate tree crowns—subtle differences in hue and texture from RGB images—is lost in a LiDAR dataset. Even more information may be available from hyperspectral or multi-temporal RGB imagery (e.g., [99]). Many studies have shown great success for spectrally distinguishing canopy species using hyperspectral (e.g., [1]) and multi-temporal imagery (e.g., [6]), while fewer studies have made use of this wealth of information available to delineate mixed- and broadleaf-dominated forests [100–102].

Of course, the use of high resolution imagery in crown delineation has been explored (e.g., [47,48]). However, many of the studies have relied on panchromatic [4] or single band imagery [103]. Future work could benefit from development of spectral or integrated LiDAR—spectral delineation methods for distinguishing broadleaf tree crowns. The increasing wide-spread availability of spectral platforms—including airborne sensor packages that acquire co-aligned hyperspectral and LiDAR data (e.g., G-LiHT [38] and NEON AOP [39])—adds incentive to develop effective methods because of the potential to apply methods broadly.

5. Conclusions

The ability to automatically delineate individual tree crowns in all types of forests would be a major step forward for applications of remote sensing in ecology and land management. We found that LiDAR-based crown delineation methods worked well in conifer-dominated stands but were less reliable in broadleaf-dominated stands. Overall, discrepancies in accuracy appears to be driven by differences in underlying traits controlling tree architecture and how trees interact with one another rather than differences between different LiDAR-based delineation methods. While broadleaf crowns often lack the same level of structural distinction as conifers, they have unique phenology and spectral characteristics that may be exploited to improve delineation techniques. Our work demonstrates a need to develop crown delineation techniques that integrate both structural and spectral characteristics to effectively delineate mixed species stands.

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Author Contributions: Conceptualization, J.H.H., S.V.O., A.P.O., and R.S.-D; methodology, J.H.H., S.V.O., A.P.O., M.W.P, and M.J.D.; software, J.H.H. and F.B.S.; validation, J.H.H.; formal analysis, J.H.H.; investigation, J.H.H., S.V.O., M.J.D., M.W.P., A.P.O., R.S.-D., F.B.S., D.B., and D.A.O.; resources, D.A.O, D.B., and S.V.O; data curation, J.H.H.; writing—original draft preparation, J.H.H.; writing—review and editing, J.H.H., S.V.O., M.J.D., M.W.P., A.P.O., R.S-D., F.B.S., D.B., D.A.O.; visualization, J.H.H.; supervision, S.V.O.; project administration, J.H.H., S.V.O.; funding acquisition, S.V.O. All authors have read and agreed to the published version of the manuscript.

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References

1. Shi, Y.; Skidmore, A.K.; Wang, T.; Holzwarth, S.; Heiden, U.; Pinnel, N.; Zhu, X.; Heurich, M. Tree species classification using plant functional traits from LiDAR and hyperspectral data. *Int. J. Appl. Earth Obs. Geoinf.* **2018**, *73*, 207–219. [CrossRef]
2. Zhao, Y.; Zeng, Y.; Zheng, Z.; Dong, W.; Zhao, D.; Wu, B.; Zhao, Q. Forest species diversity mapping using airborne LiDAR and hyperspectral data in a subtropical forest in China. *Remote Sens. Environ.* **2018**, *213*, 104–114. [CrossRef]
3. Coomes, D.A.; Dalponte, M.; Jucker, T.; Asner, G.P.; Banin, L.F.; Burslem, D.F.R.P.; Lewis, S.L.; Nilus, R.; Phillips, O.L.; Phua, M.-H.; et al. Area-based vs tree-centric approaches to mapping forest carbon in Southeast Asian forests from airborne laser scanning data. *Remote Sens. Environ.* **2017**, *194*, 77–88. [CrossRef]
4. Palace, M.; Keller, M.; Asner, G.P.; Hagen, S.; Braswell, B. Amazon forest structure from IKONOS satellite data and the automated characterization of forest canopy properties. *Biotropica* **2008**, *40*, 141–150. [CrossRef]
5. Clark, M.L.; Roberts, D.A.; Clark, D.B. Hyperspectral discrimination of tropical rain forest tree species at leaf to crown scales. *Remote Sens. Environ.* **2005**, *96*, 375–398. [CrossRef]
6. Fang, F.; McNeil, B.E.; Warner, T.A.; Maxwell, A.E. Combining high spatial resolution multi-temporal satellite data with leaf-on LiDAR to enhance tree species discrimination at the crown level. *Int. J. Remote Sens.* **2018**, *39*, 9054–9072. [CrossRef]
7. Zhang, T.; Niinemets, Ü.; Sheffield, J.; Lichstein, J.W. Shifts in tree functional composition amplify the response of forest biomass to climate. *Nature* **2018**, *556*, 99–102. [CrossRef]
8. Seebens, H.; Blackburn, T.M.; Dyer, E.E.; Genovesi, P.; Hulme, P.E.; Jeschke, J.M.; Pagad, S.; Pyšek, P.; Winter, M.; Arianoutsou, M.; et al. No saturation in the accumulation of alien species worldwide. *Nat. Commun.* **2017**, *8*, 1–9. [CrossRef] [PubMed]
9. Houghton, R.A. Land-use change and the carbon cycle. *Glob. Chang. Biol.* **1995**, *1*, 275–287. [CrossRef]
10. Ayrey, E.; Fraver, S.; Kershaw, J.A.; Kenefic, L.S.; Hayes, D.; Weiskittel, A.R.; Roth, B.E. Layer Stacking: A Novel Algorithm for Individual Forest Tree Segmentation from LiDAR Point Clouds. *Can. J. Remote Sens.* **2017**, *43*, 16–27. [CrossRef]

11. Jing, L.; Hu, B.; Noland, T.; Li, J. An individual tree crown delineation method based on multi-scale segmentation of imagery. *ISPRS J. Photogramm. Remote Sens.* **2012**, *70*, 88–98. [CrossRef]
12. Lu, X.; Guo, Q.; Li, W.; Flanagan, J. A bottom-up approach to segment individual deciduous trees using leaf-off lidar point cloud data. *ISPRS J. Photogramm. Remote Sens.* **2014**, *94*, 1–12. [CrossRef]
13. Wan Mohd Jaafar, W.; Woodhouse, I.; Silva, C.; Omar, H.; Abdul Maulud, K.; Hudak, A.; Klauberg, C.; Cardil, A.; Mohan, M. Improving Individual Tree Crown Delineation and Attributes Estimation of Tropical Forests Using Airborne LiDAR Data. *Forests* **2018**, *9*, 759. [CrossRef]
14. Zhen, Z.; Quackenbush, L.J.; Stehman, S.V.; Zhang, L. Agent-based region growing for individual tree crown delineation from airborne laser scanning (ALS) data. *Int. J. Remote Sens.* **2015**, *36*, 1965–1993. [CrossRef]
15. Williams, J.; Schonlieb, C.-B.; Swinfield, T.; Lee, J.; Cai, X.; Qie, L.; Coomes, D.A. 3D Segmentation of Trees Through a Flexible Multiclass Graph Cut Algorithm. *IEEE Trans. Geosci. Remote Sens.* **2019**. [CrossRef]
16. Dalponte, M.; Reyes, F.; Kandare, K.; Gianelle, D. Delineation of individual tree crowns from ALS and hyperspectral data: A comparison among four methods. *Eur. J. Remote Sens.* **2015**, *48*, 365–382. [CrossRef]
17. Zhen, Z.; Quackenbush, L.J.; Zhang, L. Trends in automatic individual tree crown detection and delineation-evolution of LiDAR data. *Remote Sens.* **2016**, *8*, 333. [CrossRef]
18. Vauhkonen, J.; Ene, L.; Gupta, S.; Heinzel, J.; Holmgren, J.; Pitkänen, J.; Solberg, S.; Wang, Y.; Weinacker, H.; Hauglin, K.M.; et al. Comparative testing of single-tree detection algorithms under different types of forest. *Forestry* **2012**, *85*, 27–40. [CrossRef]
19. Valladares, F.; Niinemets, U. The Architecture of Plant Crowns: From Design Rules to Light Capture and Performance. In *Functional Plant Ecology*; CRC Press: Boca Raton, FL, USA, 2007; pp. 101–150.
20. Chave, J.; Coomes, D.; Jansen, S.; Lewis, S.L.; Swenson, N.G.; Zanne, A.E. Towards a worldwide wood economics spectrum. *Ecol. Lett.* **2009**, *12*, 351–366. [CrossRef]
21. Muth, C.C.; Bazzaz, F.A. Tree canopy displacement and neighborhood interactions. *Can. J. For. Res.* **2003**, *33*, 1323–1330. [CrossRef]
22. Pretzsch, H. Canopy space filling and tree crown morphology in mixed-species stands compared with monocultures. *For. Ecol. Manag.* **2014**, *327*, 251–264. [CrossRef]
23. Forrester, D.I.; Benneter, A.; Bouriaud, O.; Bauhus, J. Diversity and competition influence tree allometric relationships—developing functions for mixed-species forests. *J. Ecol.* **2017**, *105*, 761–774. [CrossRef]
24. Fichtner, A.; Härdtle, W.; Li, Y.; Bruelheide, H.; Kunz, M.; von Oheimb, G. From competition to facilitation: How tree species respond to neighbourhood diversity. *Ecol. Lett.* **2017**, *20*, 892–900. [CrossRef] [PubMed]
25. Givnish, T.J. Ecological Constraints on the Evolution of Breeding Systems in Seed Plants: Dioecy and Dispersal in Gymnosperms. *Evolution* **2002**, *34*, 959. [CrossRef]
26. Oliver, C.D.; Stephens, E.P. Reconstruction of a Mixed-Species Forest in Central New England. *Ecology* **1977**, *58*, 562–572. [CrossRef]
27. Brodribb, T.J.; Pittermann, J.; Coomes, D.A. Elegance versus Speed: Examining the Competition between Conifer and Angiosperm Trees. *Int. J. Plant. Sci.* **2012**, *173*, 673–694. [CrossRef]
28. Augusto, L.; Davies, T.J.; Delzon, S.; de Schrijver, A. The enigma of the rise of angiosperms: Can we untie the knot? *Ecol. Lett.* **2014**, *17*, 1326–1338. [CrossRef]
29. Li, W.; Guo, Q.; Jakubowski, M.K.; Kelly, M. A New Method for Segmenting Individual Trees from the Lidar Point Cloud. *Photogramm. Eng. Remote Sens.* **2012**, *78*, 75–84. [CrossRef]
30. Wang, Y.; Hyyppa, J.; Liang, X.; Kaartinen, H.; Yu, X.; Lindberg, E.; Holmgren, J.; Qin, Y.; Mallet, C.; Ferraz, A.; et al. International Benchmarking of the Individual Tree Detection Methods for Modeling 3-D Canopy Structure for Silviculture and Forest Ecology Using Airborne Laser Scanning. *IEEE Trans. Geosci. Remote Sens.* **2016**, *54*, 5011–5027. [CrossRef]
31. Silva, C.A.; Hudak, A.T.; Vierling, L.A.; Loudermilk, E.L.; O'Brien, J.J.; Hiers, J.K.; Jack, S.B.; Gonzalez-Benecke, C.; Lee, H.; Falkowski, M.J.; et al. Imputation of Individual Longleaf Pine (Pinus palustris Mill.) Tree Attributes from Field and LiDAR Data. *Can. J. Remote Sens.* **2016**, *42*, 554–573. [CrossRef]
32. Broadbent, E.N.; Asner, G.P.; Peña-Claros, M.; Palace, M.; Soriano, M. Spatial partitioning of biomass and diversity in a lowland Bolivian forest: Linking field and remote sensing measurements. *For. Ecol. Manag.* **2008**, *255*, 2602–2616. [CrossRef]

33. Anderson-Teixeira, K.J.; Davies, S.J.; Bennett, A.C.; Gonzalez-Akre, E.B.; Muller-Landau, H.C.; Joseph Wright, S.; Abu Salim, K.; Almeyda Zambrano, A.M.; Alonso, A.; Baltzer, J.L.; et al. CTFS-ForestGEO: A worldwide network monitoring forests in an era of global change. *Glob. Chang. Biol.* **2015**, *21*, 528–549. [CrossRef] [PubMed]
34. Orwig, D.; Ellison, A. Harvard Forest CTFS-ForestGEO Mapped Forest Plot since 2014. *Environ. Data Initiat.* **2015**. [CrossRef]
35. Plotkins, A.B.; O'Keefe, J.; Foster, D.R. Harvard University Forest, Massachusetts, United States of America. In *Forest Plans of North America*; Siry, J.P., Bettinger, P., Merry, K., Grebner, D.L., Boston, K., Cieszewski, C., Eds.; Academic Press: Waltham, MA, USA, 2015; pp. 69–77. ISBN 9780127999364.
36. Orwig, D.A.; Boucher, P.; Paynter, I.; Saenz, E.; Li, Z.; Schaaf, C. The potential to characterize ecological data with terrestrial laser scanning in Harvard Forest, MA. *Interface Focus* **2018**, *8*, 20170044. [CrossRef]
37. Sullivan, F.B.; Ducey, M.J.; Orwig, D.A.; Cook, B.; Palace, M.W. Forest Ecology and Management Comparison of lidar- and allometry-derived canopy height models in an eastern deciduous forest. *For. Ecol. Manag.* **2017**, *406*, 83–94. [CrossRef]
38. Cook, B.D.; Corp, L.A.; Nelson, R.F.; Middleton, E.M.; Morton, D.C.; Mccorkel, J.T.; Masek, J.G.; Ranson, K.J.; Ly, V.; Montesano, P.M. NASA Goddard's LiDAR, Hyperspectral and Thermal (G-LiHT) Airborne Imager. *Remote Sens.* **2013**, *5*, 4045–4066. [CrossRef]
39. Kampe, T.U. NEON: The first continental-scale ecological observatory with airborne remote sensing of vegetation canopy biochemistry and structure. *J. Appl. Remote Sens.* **2010**, *4*, 043510. [CrossRef]
40. Quantum GIS Development Team. *Quantum GIS Geographic Information System*. Open Source Geospatial Foundation Project. 2018. Available online: http://qgis.osgeo.org (accessed on 10 October 2019).
41. R Core Team. *R: A Language and Environment for Statistical Computing*; R Foundation for Statistical Computing: Vienna, Austria, 2018.
42. Roussel, J.R.; Auty, D. lidR: Airborne LiDAR Data Manipulation and Visualization for Forestry Application. 2019. Available online: https://rdrr.io/cran/lidR/ (accessed on 10 October 2019).
43. Dalponte, M.; Coomes, D.A. Tree-centric mapping of forest carbon density from airborne laser scanning and hyperspectral data. *Methods Ecol. Evol.* **2016**, *7*, 1236–1245. [CrossRef]
44. Vincent, L.; Soille, P. Watersheds in Digital Spaces: An Efficient Algorithm Based on Immersion Simulations. *IEEE Trans. Pattern Anal. Mach. Intell.* **1991**, *13*, 583–598. [CrossRef]
45. Popescu, S.C.; Wynne, R.H. Seeing the Trees in the Forest. *Photogramm. Eng. Remote Sens.* **2013**, *70*, 589–604. [CrossRef]
46. Yin, D.; Wang, L. How to assess the accuracy of the individual tree-based forest inventory derived from remotely sensed data: A review. *Int. J. Remote Sens.* **2016**, *37*, 4521–4553. [CrossRef]
47. Lamar, W.R.; McGraw, J.B.; Warner, T.A. Multitemporal censusing of a population of eastern hemlock (Tsuga canadensis L.) from remotely sensed imagery using an automated segmentation and reconciliation procedure. *Remote Sens. Environ.* **2005**, *94*, 133–143. [CrossRef]
48. Leckie, D.G.; Jay, C.; Gougeon, F.A.; Sturrock, R.N.; Paradine, D. Detection and assessment of trees with Phellinus weirii (laminated root rot) using high resolution multi-spectral imagery. *Int. J. Remote Sens.* **2004**, *25*, 793–818. [CrossRef]
49. Kane, V.R.; Gillespie, A.R.; McGaughey, R.; Lutz, J.A.; Ceder, K.; Franklin, J.F. Interpretation and topographic compensation of conifer canopy self-shadowing. *Remote Sens. Environ.* **2008**, *112*, 3820–3832. [CrossRef]
50. Pommerening, A. Approaches to quantifying forest structures. *Forestry* **2002**, *75*, 305–324. [CrossRef]
51. Ducey, M.J.; Knapp, R.A. A stand density index for complex mixed species forests in the northeastern United States. *For. Ecol. Manag.* **2010**, *260*, 1613–1622. [CrossRef]
52. Heip, C.H.R.; Herman, P.M.J.; Soetaert, K. Indices of diversity and evenness. *Océanis* **1998**, *24*, 61–87.
53. McCune, B.; Grace, J.B. *Analysis of Ecological Communities*; MjM Sotware Design: Gleneden Beach, OR, USA, 2002.
54. Hair, J.F.; Anderson, R.E.; Tatham, R.L.; Black, W.C. *Multivariate Data Analysis*, 3rd ed.; MacMillian: New York, NY, USA, 1995.
55. Burnham, K.P.; Anderson, D.R. *Model Selection and Multimodel Inference: A Practical Information-Theoretic Approach*, 2nd ed.; Library of Congress Cataloging-in-Publication Data; Springer: New York, NY, USA, 2002; Volume 172, ISBN 978-0-387-22456-5.

56. Oberle, B.; Ogle, K.; Zanne, A.E.; Woodall, C.W. When a tree falls: Controls on wood decay predict standing dead tree fall and new risks in changing forests. *PLoS ONE* **2018**, *13*, 1–22. [CrossRef]
57. Bates, D.; Maechler, M.; Bolker, B.; Walker, S. Fitting Linear Mixed-Effects Models Using lme4. *J. Stat. Softw.* **2015**, *67*, 1–48. [CrossRef]
58. Aubry-Kientz, M.; Dutrieux, R.; Ferraz, A.; Saatchi, S.; Hamraz, H.; Williams, J.; Coomes, D.; Piboule, A.; Vincent, G. A comparative assessment of the performance of individual tree crowns delineation algorithms from ALS data in tropical forests. *Remote Sens.* **2019**, *11*, 1086. [CrossRef]
59. Xiao, W.; Zaforemska, A.; Smigaj, M.; Wang, Y.; Gaulton, R. Mean shift segmentation assessment for individual forest tree delineation from airborne lidar data. *Remote Sens.* **2019**, *11*, 1263. [CrossRef]
60. Lindberg, E.; Holmgren, J. Individual Tree Crown Methods for 3D Data from Remote Sensing. *Curr. For. Rep.* **2017**, *3*, 19–31. [CrossRef]
61. Eysn, L.; Hollaus, M.; Lindberg, E.; Berger, F.; Monnet, J.M.; Dalponte, M.; Kobal, M.; Pellegrini, M.; Lingua, E.; Mongus, D.; et al. A benchmark of lidar-based single tree detection methods using heterogeneous forest data from the Alpine Space. *Forests* **2015**, *6*, 1721–1747. [CrossRef]
62. Miles, P.D.; Smith, W.B. Specific Gravity and Other Properties of Wood and Bark for 156 Tree Species Found in North America. In *Research Note Nrs-38*; United States Forest Service: Newtown Square, PA, USA, 2009; Volume 35.
63. Anten, N.P.R.; Schieving, F. The Role of Wood Mass Density and Mechanical Constraints in the Economy of Tree Architecture. *Am. Nat.* **2010**, *175*, 250–260. [CrossRef] [PubMed]
64. Horn, H.S. *The Adapative Geometry of Trees*; Princeton University Press: Princeton, NJ, USA, 1971.
65. Ducey, M.J. Evergreenness and wood density predict height-diameter scaling in trees of the northeastern United States. *For. Ecol. Manag.* **2012**, *279*, 21–26. [CrossRef]
66. Thompson, J.R.; Carpenter, D.N.; Cogbill, C.V.; Foster, D.R. Four Centuries of Change in Northeastern United States Forests. *PLoS ONE* **2013**, *8*, e72540. [CrossRef]
67. Abrams, M.D. Eastern White Pine Versatility in the Presettlement Forest. *Bioscience* **2001**, *51*, 967–979. [CrossRef]
68. Strigul, N.; Pristinski, D.; Purves, D.; Dushoff, J.; Pacala, S. Scaling from trees to forests: Tractable macroscopic equations for forest dynamics. *Ecol. Monogr.* **2008**, *78*, 523–545. [CrossRef]
69. Gower, S.T.; Isebrands, J.G.; Sheriff, D.W. Carbon Allocation and Accumulation in Conifers. In *Resource Physiology of Conifers*; Academic Press: New York, NY, USA, 1995; pp. 217–254.
70. Ollinger, S.V. Sources of variability in canopy reflectance and the convergent properties of plants. *New Phytol.* **2011**, *189*, 375–394. [CrossRef]
71. Pretzsch, H.; Rais, A. Wood quality in complex forests versus even-aged monocultures: Review and perspectives. *Wood Sci. Technol.* **2016**, *50*, 845–880. [CrossRef]
72. Hajek, P.; Seidel, D.; Leuschner, C. Mechanical abrasion, and not competition for light, is the dominant canopy interaction in a temperate mixed forest. *For. Ecol. Manag.* **2015**, *348*, 108–116. [CrossRef]
73. Putz, F.E.; Parker, G.G.; Archibald, R. Mechanical Abrasion and Intercrown Spacing. *Am. Midl. Nat.* **1984**, *112*, 24–28. [CrossRef]
74. Goudie, J.W.; Polsson, K.R.; Ott, P.K. An empirical model of crown shyness for lodgepole pine (Pinus contorta var. latifolia [Engl.] Critch.) in British Columbia. *For. Ecol. Manag.* **2009**, *257*, 321–331. [CrossRef]
75. Rainey, S.M.; Nadelhoffer, K.J.; Silver, W.L.; Martha, R. Effects of Chronic Nitrogen Additions on Understory Species in a Red Pine Plantation. *Ecol. Appl.* **1999**, *9*, 949–957. [CrossRef]
76. Wonn, H.T.; O'Hara, K.L. Height:diameter ratios and stability relationships for four northern Rocky Mountain tree species. *West. J. Appl. For.* **2001**, *16*, 87–94. [CrossRef]
77. Rudnicki, M.; Silins, U.; Lieffers, V.J.; Josi, G. Measure of simultaneous tree sways and estimation of crown interactions among a group of trees. *Trees Struct. Funct.* **2001**, *15*, 83–90. [CrossRef]
78. Orwig, D.A.; Foster, D.R. Forest Response to the Introduced Hemlock Woolly Adelgid in Southern New England, USA. *J. Torrey Bot. Soc.* **1998**, *125*, 60–73. [CrossRef]
79. Foster, D.R.; Baiser, B.; Plotkins, A.B.; D'Amato, A.; Ellison, A.M.; Orwig, D.; Oswald, W.; Thompson, J. *Hemlock: A Forest Giant on the Edge*; Foster, D.R., Ed.; Yale University Press: New Haven, CT, USA, 2014.
80. Jucker, T.; Bouriaud, O.; Coomes, D.A. Crown plasticity enables trees to optimize canopy packing in mixed-species forests. *Funct. Ecol.* **2015**, *29*, 1078–1086. [CrossRef]

81. Oliver, C.D.; Larson, B.C. *Forest Stand. Dynamics*; John Wiley and Sons Inc.: New York, NY, USA, 1996; ISBN 0471138339.
82. Sapijanskas, J.; Paquette, A.; Potvin, C.; Kunert, N.; Loreau, M. Tropical tree diversity enhances light capture through plastic architectural changes and spatial and temporal niche differences. *Ecology* **2012**, *95*, 1–32.
83. Morin, X.; Fahse, L.; Scherer-Lorenzen, M.; Bugmann, H. Tree species richness promotes productivity in temperate forests through strong complementarity between species. *Ecol. Lett.* **2011**, *14*, 1211–1219. [CrossRef]
84. Pretzsch, H.; Schütze, G. Effect of tree species mixing on the size structure, density, and yield of forest stands. *Eur. J. For. Res.* **2016**, *135*, 1–22. [CrossRef]
85. Williams, L.J.; Paquette, A.; Cavender-Bares, J.; Messier, C.; Reich, P.B. Spatial complementarity in tree crowns explains overyielding in species mixtures. *Nat. Ecol. Evol.* **2017**, *1*, 0063. [CrossRef] [PubMed]
86. Roberts, D.; Roth, K.; Perroy, R. Hyperspectral Vegetation Indices. In *Hyperspectral Remote Sensing of Vegetation*; CRC Press: Boca Raton, FL, USA, 2011; ISBN 978-1-4398-4537-0.
87. Qiao, K.; Zhu, W.; Xie, Z.; Li, P. Estimating the Seasonal Dynamics of the Leaf Area Index Using Piecewise LAI-VI Relationships Based on Phenophases. *Remote Sens.* **2019**, *11*, 689. [CrossRef]
88. Waring, R.H.; Law, B.E.; Goulden, M.L.; Bassow, S.L.; McCreight, R.W.; Wofsy, S.C.; Bazzaz, F.A. Scaling gross ecosystem production at Harvard Forest with remote sensing: A comparison of estimates from a constrained quantum-use efficiency model and eddy correlation. *Plant. Cell Environ.* **1995**, *18*, 1201–1213. [CrossRef]
89. Hui, G.; Zhao, X.; Zhao, Z.; Gadow, K. Von Evaluating Tree Species Diversity Based on Neighborhood Relationships. *For. Sci.* **2011**, *57*, 292–300.
90. Freckleton, R.P.; Watkinson, A.R. Asymmetric competition between plant species. *Funct. Ecol.* **2001**, *15*, 615–623. [CrossRef]
91. Stephenson, N.L.; Das, A.J.; Condit, R.; Russo, S.E.; Baker, P.J.; Beckman, N.G.; Coomes, D.A.; Lines, E.R.; Morris, W.K.; Rüger, N.; et al. Rate of tree carbon accumulation increases continuously with tree size. *Nature* **2014**, *507*, 90–93. [CrossRef]
92. Lutz, J.A.; Furniss, T.J.; Johnson, D.J.; Davies, S.J.; Allen, D.; Alonso, A.; Anderson-Teixeira, K.J.; Andrade, A.; Baltzer, J.; Becker, K.M.L.; et al. Global importance of large-diameter trees. *Glob. Ecol. Biogeogr.* **2018**, *27*, 849–864. [CrossRef]
93. Orwig, D.A.; Thompson, J.R.; Povak, N.A.; Manner, M.; Niebyl, D.; Foster, D.R. A foundation tree at the precipice: Tsuga canadensis health after the arrival of Adelges tsugae in central New England. *Ecosphere* **2012**, *3*, 1–16. [CrossRef]
94. Small, M.J.; Small, C.J.; Dreyer, G.D.; The, S.; Society, B.; Sep, N.J.; Small, M.J.; Small, C.J.; Dreyer, G.D. Changes in a Hemlock-Dominated Forest following Woolly Adelgid Infestation in Southern New England. *J. Torrey Bot. Soc.* **2005**, *132*, 458–470. [CrossRef]
95. van Ewijk, K.Y.; Treitz, P.M.; Scott, N.A. Characterizing Forest Succession in Central Ontario using Lidar-derived Indices. *Photogramm. Eng. Remote Sens.* **2013**, *77*, 261–269. [CrossRef]
96. Lorimer, C.; Frelich, L.E. A structural Alternative to Chronosequence Analysis for Uneven-Aged Northern Hardwood Forests. *J. Sustain. For.* **1998**, *6*, 347–365. [CrossRef]
97. Bradford, J.B.; Kastendick, D.N. Age-related patterns of forest complexity and carbon storage in pine and aspen–birch ecosystems of northern Minnesota, USA. *Can. J. For. Res.* **2010**, *40*, 401–409. [CrossRef]
98. Ogunjemiyo, S.; Parker, G.; Roberts, D. Reflections in bumpy terrain: Implications of canopy surface variations for the radiation balance of vegetation. *IEEE Geosci. Remote Sens. Lett.* **2005**, *2*, 90–93. [CrossRef]
99. Klosterman, S.; Melaas, E.; Wang, J.; Martinez, A.; Frederick, S.; O'Keefe, J.; Orwig, D.A.; Wang, Z.; Sun, Q.; Schaaf, C.; et al. Fine-scale perspectives on landscape phenology from unmanned aerial vehicle (UAV) photography. *Agric. For. Meteorol.* **2018**, *248*, 397–407. [CrossRef]
100. Yang, J.; He, Y.; Caspersen, J.P.; Jones, T.A. Delineating Individual Tree Crowns in an Uneven-Aged, Mixed Broadleaf Forest Using Multispectral Watershed Segmentation and Multiscale Fitting. *IEEE J. Sel. Top. Appl. Earth Obs. Remote Sens.* **2017**, *10*, 1390–1401. [CrossRef]
101. Maschler, J.; Atzberger, C.; Immitzer, M. Individual tree crown segmentation and classification of 13 tree species using Airborne hyperspectral data. *Remote Sens.* **2018**, *10*, 1218. [CrossRef]

102. Huang, H.; Li, X.; Chen, C. Individual tree crown detection and delineation from very-high-resolution UAV images based on bias field and marker-controlled watershed segmentation algorithms. *IEEE J. Sel. Top. Appl. Earth Obs. Remote Sens.* **2018**, *11*, 2253–2262. [CrossRef]
103. Ke, Y.; Quackenbush, L.J. A comparison of three methods for automatic tree crown detection and delineation from high spatial resolution imagery. *Int. J. Remote Sens.* **2011**, *32*, 3625–3647. [CrossRef]