# Genetic diversity and population structure of three tree species across an urban to remnant landscape gradient 

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#### Abstract

Trees in urban areas offer ecosystem services like carbon sequestration, storm water attenuation, reduction of energy used in buildings and wildlife habitat. Cities invest substantial funds and resources to maintain a healthy urban forest, and much research has been done to improve its resiliency and sustainability. Studies have been done to determine the quantity, density and variety of tree species in urban areas, but little is known about their genetic diversity. In this study the genetic diversity and population structure of three tree species (Quercus rubra (red oak), Acer saccharum (sugar maple) and Fraxinus pennsylvanica (green ash)) were compared on an urban to remnant gradient in the Chicago area. Microsatellites were used to compare neutral diversity. Genetic diversity did not differ significantly among land use types for any of the three species as measured by numbers of alleles, private alleles and heterozygosity. There was no genetic structure based on land use type among red oaks. However, genetic structure among sugar maple and green ash is evident with significant genetic differentiation between urban and remnant sites. The variance in genetic structure is due to the variable cultivation history of the three tree species. There are few red oak cultivars, and this species has remained genetically indistinguishable from remnant forests. Sugar maple and green ash have been cultivated over the last 50 years, leading to these species becoming genetically differentiated from remnant populations. Careful sourcing practices will be necessary to ensure the future genetic diversity of the urban forest.


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## Introduction

The urban forest is defined as all of the trees that grow in urban areas (e.g. street and park trees, spontaneous trees in brown fields or in parking lots, those that grow in managed and unmanaged forests, and trees planted in private yards) (Nowak et al. 2010), and it provides a myriad of ecosystem services. In the face of climate change trees are championed for their ability to remove and store carbon dioxide and other pollutants, which is especially important in urban areas where both are produced in large quantities (Akbari and Pomerantz 2001; Nowak et al. 2010). The urban tree canopy can also offset the heat island effect in several ways. It cools urban areas through evapotranspiration and altering air flow (Sawka et al. 2013). Trees also shade buildings, reducing the amount of solar radiation stored in impermeable surfaces, which would otherwise be redirected as heat (Sawka et al. 2013). This shading also reduces the energy necessary to cool buildings (Akbari and Pomerantz 2001). The urban forest absorbs water runoff, which can prevent it from reaching the sewage system or polluting nearby bodies of water (Walsh et al. 2012). Finally, trees increase land value, improve aesthetics, reduce erosion, interrupt wind gusts and provide habitat for wildlife (Donovan et al. 2010, Nowak et al. 2010).

The urban forest is valuable, but it is also expensive to maintain and can be susceptible to pests that cause tremendous damage (Roman and Scatena 2011). Large cities budget millions of dollars annually to prune and maintain trees (Nowak et al. 2002). On top of this regular maintenance cities occasionally contend with massive tree failures: when a pest, disease or environmental condition (such as drought or severe weather) kills a large number of individuals in a brief time period (Jim et al. 1997). One of the most devastating occurrences of massive tree failure in the urban forest was caused by Dutch elm disease. The American elm (Ulmus americana L.) was once the most commonly planted street and shade tree in eastern North

America (Biggerstaff et al. 1999). Beginning in the 1920's the exotic disease swept through the American elms, eventually killing close to $95 \%$ of the population (Biggerstaff et al. 1999). Because American elm was planted so heavily, the death of these trees left many cities with a large percentage of trees needing removal; in some cases entire blocks were denuded of shade trees (Biggerstaff et al. 1999). Today, emerald ash borer (Agrilus planipennis, Fairmaire) is wreaking similar havoc on ash trees (Fraxinus sp.). By 2019 the emerald ash borer is expected to destroy 38 million trees in the Midwestern United States; 17 million of which are street and park trees and will need to be removed lest they cause a safety hazard, and in many cases replaced, at an estimated cost of $\$ 10.7$ billion (Kovacs et al. 2010). Additionally, the health of other species may be adversely affected by other stresses such as climate change, the gypsy moth or the Asian longhorn beetle; any one of which will cost cities millions of dollars (Ball et al. 2007; Nowak et al. 2010).

In the aftermath of Dutch elm disease, urban foresters have learned the importance of planting a variety of tree species (Raup et al. 2006; Ball et al. 2007). By planting many different species, genera and families the city can help ensure that a single pest will not destroy a large percentage of its trees (Schoene et al. 2011; Lacan et al. 2008). Much attention and research has gone into ensuring that the urban forest is species rich, and many urban forests strive to contain no more than $10 \%$ of a given species, $20 \%$ of a genus and $30 \%$ of a family (Santamour 1990). However, little is known about the genetic diversity within species of the urban forest. It is assumed that the urban forest is not genetically diverse (Fissore et al. 2012). Most trees that are planted by the city are sourced from a handful of growers, and these growers obtain their seed stock from a limited number of trees (Worfolk, interview 2012). Additionally, many of the trees that are planted in urban areas are cultivars, some of which are grafted clones (Santamour 1990).

Foresters seek out uniformity within species: individuals with the same origin will have similar survivorship, adaptability, morphological characteristics and performance (Santamour 1990). In short, these trees are predictable. However, morphological uniformity often comes at the expense of genetic uniformity, and a lack of genetic diversity may prove to be deleterious (Raupp et al. 2006, Sork et al. 2013, Reed and Frankham 2003).

Genetic diversity, and in particular levels of heterozygosity, have been found to be related to population health in a variety of studies (see Reed and Frankham (2003) and Leimu et al. (2006) for reviews). Correlations between genetic diversity and fitness show that low genetic diversity will result in increased homozygosity as closely related individuals interbreed through generations. This inextricably leads to the expression of deleterious alleles and inbreeding depression (Reed and Frankham 2003). However, many trees in the urban forest do not breed naturally. All trees that are intentionally planted in urban areas are sourced from nurseries. Lack of genetic diversity in these instances may be caused by the excessive planting of a single cultivar or related individuals, but because new individuals are created in nurseries and not from the existing population this does not lead to inbreeding depression among intentionally planted trees. There are trees, however, that grow spontaneously in the urban forest. Unmanaged properties and city forests have a host of trees that reproduce naturally (Nowak 2012). Many of these trees are remnants from historic forests, but when they are enveloped by urbanization the trees may be influenced by planted trees. In these areas cultivars may interbreed with the autochthonous trees and cause genetic swamping (Lesica and Allendorf 1999). Over generations the spontaneous trees in the urban forest may experience deleterious inbreeding effects.

Even in forests where natural breeding does not occur (and the threat of inbreeding depression is nonexistent), genetic diversity is an important predictor of forest sustainability. If,
for example, all of the maples in a city are genetic clones of one another, a single pest or disease could rapidly damage or kill them all, whereas, if they are genetically diverse some individuals may be resistant to that pest or disease, limiting overall losses (Sork et al. 2013). American elms that are resistant to Dutch elm disease have recently been discovered (Whittemore, 2011). Ashes have proven to be variably tolerant of the emerald ash borer, with some trees still surviving after many years of infection (Marshall et al. 2013). Trees have long generation times, meaning that they are poor at rapidly adapting to new pests, disease or changing climatic conditions (Aitken et al. 2008; Sork et al. 2013). Current genetic diversity may be one of the best predictors of a tree population's ability to survive disasters (Sork et al. 2013). Even in the case of catastrophic disease a few individuals in a diverse forest may persist. Knowledge about the genetic diversity of trees in the urban forest will foster a more advanced understanding of how the trees may react to massive tree failures and help guide city planning.

## Research objectives

The objective of the study was to compare neutral genetic diversity and structure in three tree species across a land use gradient: planted trees in urban areas, trees that occur spontaneously in urban areas, and remnant forests. While neutral genetic diversity does not necessarily correlate with genetic fitness, it can be used to identify where there is a dearth of diversity (Holderegger et al. 2006). The study species were: red oak (Quercus rubra L.), sugar maple (Acer saccharum Marsh.), and green ash (Fraxinus pennsylvanica Marsh.). These species were chosen because they are easily found in both remnant and urban settings and are native to the Chicago region. Sugar maple and red oak are currently planted by the City of Chicago and although green ash is no longer planted, it is currently one of the most abundant street trees
(Nowak et al. 2010). Using microsatellite markers, the following hypotheses were tested: 1) There will be a decrease in genetic diversity across a land use gradient from remnant to spontaneous to planted land use for all three species, 2) There will be distinct genetic structure between the planted trees and the remnant forest, while spontaneous trees will have genetic structure that is a mix of the other two land use types.

## Methods

## Study site

This study focuses on the trees in and around the City of Chicago, a city that has made urban greening a priority in recent years. In the past 20 years canopy cover has increased from $11 \%$ to $17 \%$ largely through the planting of street and park trees (Nowak et al. 2010). However, this forest is under considerable stress right now. Emerald ash borer is predicted to affect $11 \%$ of the Chicago forest, around 3,000,000 trees (Nowak et al. 2010). Many of these trees will need to be removed and replaced. Chicago foresters are already planning on how best to replace these trees with the goal of making a sustainable urban forest. Developing an understanding of intraspecies genetic diversity could help achieve this goal.

## Study species

Three species were analyzed in this study: red oak (Quercus rubra L.), sugar maple (Acer saccharum Marsh.), and green ash (Fraxinus pennsylvanica Marsh.) (henceforth all species will be referred to by their common names). All of these species are native and are common in the Chicago urban forest and in the surrounding remnant woodlands. Additionally, they are all planted intentionally and occur spontaneously in the City of Chicago (Nowak 2012). Pollination
strategies can affect the ability of a species to breed with distant populations. If the study species have varying pollination syndromes it could have an effect on the landscape genetics. These three genera were chosen because they are all wind pollinated and should have a similar capacity for gene flow through pollination (Dow and Ashley 1996; Fore et al. 1992; Heuertz 2003). All three species also have methods for seed dispersal. Green ash and sugar maples seeds are both dispersed by wind while red oak seeds are moved by squirrels. None of these methods tend to move seeds over great distances, but do allow for additional movement beyond what gravity alone would provide (Fore et al. 1992; Kennedy 1990; Sork 1984).

## Field methods

For each individual tree included in the study, a leaf was taken for genetic analysis. This leaf was stored in a cooler immediately after collection, and then frozen at $-20^{\circ} \mathrm{C}$ within three hours. Additional data taken included: the diameter at breast height (DBH) and GPS coordinates. This studied endeavored to look at trees across an urban to remnant gradient. To do this, trees were sampled in three land use categories. The first category was trees that were intentionally planted in the city (hereafter known as planted). These trees were located along streets and in parks. Next, trees that had seeded spontaneously within the urban matrix were sampled (hereafter known as spontaneous). Planted and spontaneous trees were differentiated using strategies described by Nowak (2012). Remnant forests were found outside of the City of Chicago, and are areas that have been continuously forested since before European colonization. In the planted category, 59 red oak, 75 sugar maple and 69 green ash samples were collected: in the spontaneous category, 57 red oak, 42 sugar maple and 48 green ash samples were collected; and in the remnant category, 72 red oak, 80 sugar maple and 66 green ash samples were collected.

Planted trees were sampled differently from the other two land use types. Planted sites were selected using a grid system. Fifteen locations were chosen using a random number generator that corresponded with city blocks. Trees were sampled within a half mile by half mile square surrounding this point. Every sugar maple and red oak was sampled within these plots; while every fifth green ash was sampled, because ash trees were much more prevalent. Additionally, sugar maples and red oaks are generally less commonly planted as street trees, but are frequently found in city parks. For eight of the city blocks a nearby park was also identified and sampled (Figure 1).

Remnant and spontaneous areas were sampled in an identical manner to each other. First, an assessment of the forest composition was completed. In forests that had abundant maples and few oaks and ashes, every fifth maple encountered would be sampled and every other oak and ash. If oaks were encountered more frequently they were sampled less regularly and ashes and maples more regularly. Four urban forests were sampled: Catherine Chevalier Woods, Dan Ryan Woods, Labagh Woods, and the Montrose Point Bird Sanctuary. The remnant areas sampled were: Busse Woods, Morton Arboretum, and Somme Woods (Figure 1). These sites are all within the Chicago metropolitan area, are relatively pristine and have been wooded since before European colonization.


Figure 1: Sampling locations for study. Each point represents a single tree and the colors represent different land use types.

## Molecular methods

A modified cetyltrimethyl ammonium bromide method was used to extract DNA (Doyle and Doyle 1987). Microsatellite primers measured neutral diversity in all species. DNA was amplified using the polymerase chain reaction (PCR) (methods for each species follow), and genotypes were obtained for nuclear microsatellite loci. All forward primers were modified with the addition of an M13 sequence to the $5^{\prime}$ end ( $5^{\prime}$-CACGACGTTGTAAAACGAC-3') and
labeled with Wellred Dye D2, D3 or D4 (Sigma-Proligo, St. Louis, Missouri, USA) for visualization (Schuelke 2000).

Primers and PCR conditions varied for each species. For red oak an initial $10 \mu \mathrm{~L}$ reaction was conducted with $5 \mu \mathrm{~L}$ of PCR MasterMix (Promega, Madison, Wisconsin, USA; final concentration of $0.025 \mu \mathrm{~g} / \mu \mathrm{L}$ Taq DNA polymerase in a proprietary reaction buffer [ pH 8.5 ], $200 \mu \mathrm{M}$ of each dNTP, and 1.5 mM MgCl 2 ), along with 5 ng template DNA, and $0.25 \mu \mathrm{M}$ of both forward and reverse primers. The conditions for this PCR were: 1 min of denaturation at $94{ }^{\circ} \mathrm{C}$; then 15 cycles of $94{ }^{\circ} \mathrm{C}$ for $30 \mathrm{~s}, 45^{\circ} \mathrm{C}$ for 45 s , and $72{ }^{\circ} \mathrm{C}$ for 30 s ; and a final extension of 72 ${ }^{\circ} \mathrm{C}$ for 9 min . The product was labeled through a second $15-\mu \mathrm{L}$ PCR reaction containing $10 \mu \mathrm{~L}$ of the original PCR product, with an additional $2.5 \mu \mathrm{LMasterMix}$ (Promega) and $0.25 \mu \mathrm{M}$ of M13 primer labeled with WellRed D2, D3, or D4 Dye. The conditions for this PCR were: 1 min of denaturation at $94^{\circ} \mathrm{C}$; then 30 cycles of $94{ }^{\circ} \mathrm{C}$ for $30 \mathrm{~s}, 45^{\circ} \mathrm{C}$ for 45 s , and $72{ }^{\circ} \mathrm{C}$ for 30 s ; and a final extension of $72^{\circ} \mathrm{C}$ for 9 min .

The same two-step reaction was used for both sugar maple and green ash. For the sugar maples the initial $10-\mu \mathrm{L}$ reaction contained $5 \mu \mathrm{~L}$ PCR MasterMix (Promega), along with 5 ng template DNA, $0.5 \mathrm{ng} / \mu \mathrm{L}$ BSA, $0.25 \mathrm{ng} / \mu \mathrm{L} \mathrm{MgCl} 2_{2}$ and $0.25 \mu \mathrm{M}$ of both forward and reverse primers. The conditions of the PCR were: 15 min of denaturation at $95^{\circ} \mathrm{C}$; then 15 cycles of 94 ${ }^{\circ} \mathrm{C}$ for $45 \mathrm{~s}, 56^{\circ} \mathrm{C}$ for 1 min , and $72{ }^{\circ} \mathrm{C}$ for 45 sec ; and a final extension of $72{ }^{\circ} \mathrm{C}$ for 10 min . The labeling $15-\mu \mathrm{L}$ PCR reaction contained $10 \mu \mathrm{~L}$ of the original PCR product, with an additional 2.5 $\mu \mathrm{L}$ of PCR MasterMix (Promega), $0.5 \mathrm{ng} / \mu \mathrm{L}$ BSA, $0.25 \mathrm{ng} / \mu \mathrm{L} \mathrm{MgCl}_{2}$ and $0.25 \mu \mathrm{M}$ of M13 primer labeled with the Wellred Dye. The conditions for this PCR were: 15 min of denaturation at $95^{\circ} \mathrm{C}$; then 35 cycles of $94^{\circ} \mathrm{C}$ for $45 \mathrm{~s}, 56^{\circ} \mathrm{C}$ for 1 min , and $72^{\circ} \mathrm{C}$ for 45 sec ; and a final extension of $72{ }^{\circ} \mathrm{C}$ for 10 min .

The initial $10 \mu \mathrm{~L}$ reaction for green ash contained $5 \mu \mathrm{~L}$ PCR MasterMix (Promega), along with 5 ng template DNA, $0.5 \mathrm{ng} / \mu \mathrm{L} \mathrm{BSA}, 0.25 \mathrm{ng} / \mu \mathrm{L} \mathrm{MgCl}_{2}$, and $0.25 \mu \mathrm{M}$ of both forward and reverse primers. The conditions of the PCR were: 5 min at $96^{\circ} \mathrm{C}$; then 15 cycles of $94{ }^{\circ} \mathrm{C}$ for 1 $\min , 52^{\circ} \mathrm{C}$ for 1 min , and $72^{\circ} \mathrm{C}$ for 30 s ; and a final extension of $72^{\circ} \mathrm{C}$ for 9 min . The labeling $15-\mu \mathrm{L}$ PCR reaction contained $10 \mu \mathrm{~L}$ of the original PCR product, with an additional $2.5 \mu \mathrm{~L}$ of PCR $2 \times$ MasterMix (Promega), $0.5 \mathrm{ng} / \mu \mathrm{L}$ BSA, $0.25 \mathrm{ng} / \mu \mathrm{L} \mathrm{MgCl}_{2}$ and $0.25 \mu \mathrm{M}$ of M13 primer labeled with Wellred Dye. The conditions for this PCR were: 5 min at $96^{\circ} \mathrm{C}$; then 35 cycles of $94{ }^{\circ} \mathrm{C}$ for $1 \mathrm{~min}, 52^{\circ} \mathrm{C}$ for 1 min , and $72^{\circ} \mathrm{C}$ for 30 s ; and a final extension of $72^{\circ} \mathrm{C}$ for 9 min .

PCR products were analyzed and scored using a CEQ 8000 Genetic Analysis System version 9.0 (Beckman Coulter, Brea, California, USA). Each well included $30 \mu \mathrm{~L}$ of HiDi formamide (Azco Biotech., San Diego, CA), $3.3 \mu \mathrm{~L}$ of 400 bp ladder (Beckman Coulter, Brea, CA) and the following amounts of PCR product: $0.5 \mu \mathrm{~L}$ of Wellred Dye D4 labeled product, 1 $\mu \mathrm{L}$ of Wellred Dye D3 labeled product and $2.5 \mu \mathrm{~L}$ of Wellred D2 labeled product.

Primers used for each species are listed in Table 1. Primers used in the red oak analysis were originally described by Aldrich et al. (2002) and Aldrich and Jagtop (2003) Primers for green ash were originally developed for European ash (Fraxinus excelsior L.) and were described by Lefort and Frascaria-Lacoste (1999). Sugar maple primers were originally developed for sycamore leaf maple (Acer pseudoplatinus L.) by Pandey et al. (2004).

Table 1:Primers for red oak were developed by Aldrich et al. (2002) and Aldrich and Jagtop (2003). Primers used for sugar maple were developed by Pandey et al. (2004). Green ash primers were developed by Lefort and Frascaria-Lacoste (1999).

| Quercus rubra (Red oak) |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: |
| Locus | Repeat | GenBank accession | Primer sequence | Size range (bp) |
| quru-GA-0A01 | $(\mathrm{GA})_{11}$ | AF523851 | F: СTCTCGCTCTGCACGTGACTCA R:TTTGATTGATATAATTGATCGCT | 132-170 |
| quru-GA-1G13 | $(\mathrm{GA})_{14}$ | AF523862 | F: AAAACTCACACAGCCGATTACTA R:GATTCCATTGTCAACTGCGAAGA | 179-215 |
| quru-GA-1C06 | $(\mathrm{GA})_{29}$ | AF523858 | F: CAAATAAATATTGTGGGGTTCA R:GGAGGGGATCCGGAAAA | 247-305 |
| quru-GA-0A03 | $(\mathrm{GA})_{17}$ |  | F: ATTTTATATTAGCATAAGGGTG R:GGCTTCACATTGAGAACGTTG | 187-245 |
| quru-GA-1F02 | $(\mathrm{GA})_{15}$ | AF523860 | F: ССААТССАСССТТССААGTTCC R:TGGTTGTTTTGCTTTATTCAGCC | 165-215 |
| quru-GA-0I21 | $(\mathrm{GA})_{16}$ | AF523855 | F: ATATGGTCCCGATTAATTC <br> R:GGGCAACATTCAAATGTATCTA | 173-213 |
| quru-GA-2G07 | $(\mathrm{GA})_{23}$ |  | F: GCCAACAAATTTAACTATCCAT R:TAACTGGGCTAGATAATCAG | 218-258 |
| quru-GA-1C08 | $(\mathrm{GA})_{29}$ | AF523859 | F: TCCCAATCGATGTTTGATAAGG R:GGGCTCTTGAGAGGATGTAGG | 272-328 |
| quru-GA-2H14 | $(\mathrm{GA})_{18}$ |  | F: ATTACGCGAGCGTGCAGT <br> R:GTGCTCCACGAATGCTCTAGCCA | 281-347 |
| quru-GA-0E09 | $(\mathrm{GA})_{16}$ | AF523854 | F: TGCCATCCCTATACACAACCA R: ССТССАТСАСАAAGTTGCC | 183-251 |
| Acer saccharum (Sugar maple) |  |  |  |  |
| Locus | Repeat | GenBank accession | Primer sequence | Size range (bp) |
| MAP2 | $(\mathrm{GT})_{23}$ | AJ620722 | F: CATTAAACACATTTAAGCAAAACAAG <br> R: ATCGGTTTGACATTGAGTGG | 152-186 |
| MAP9 | $(\mathrm{GA})_{8}$ | AJ620723 | F: ACAATAAAAGAGCCCACATAGATAG R:TCTCTTCAATTGCAAGGCTTC | 110-126 |
| MAP33 | $(\mathrm{GT})_{18}$ | AJ620726 | F:GCAATGAACACATATACAAACAAGAG <br> R:GCAACAAATGCCCTCTCAAG | 132-168 |
| MAP34 | $(\mathrm{CA})_{21}$ | AJ620727 | F: АССАТТСТСАССССТССАТС R:TAAGTGGGAACATGGCAAGG | 142-186 |
| MAP46 | $(\mathrm{GT})_{8} \mathrm{GAT}$ | AJ620729 | F: CATAATGTAGGGACACATATGAATG | 169-177 |

$\left.\left.\begin{array}{|lllll|} & (\mathrm{GT})_{8} & & \text { R:GAGCGTCAAAGATTGACTTGG }\end{array}\right] \begin{array}{l}\text { Size } \\ \text { range } \\ \text { (bp) }\end{array}\right)$

## Statistical methods

Each primer was tested for possible null alleles and deviations from Hardy Weinberg equilibrium using Microchecker (van Oosterhout et al. 2004). In order to determine if there was a decrease in genetic diversity from remnant to planted sites average number of alleles per loci $\left(N_{a}\right)$, number of effective alleles $\left(N_{e}\right)$, Shannon's information index (I), observed and expected heterozygosity ( $\mathrm{H}_{\mathrm{o}}$ and $\mathrm{H}_{\mathrm{e}}$ ) F statistics (inbreeding coefficient) and average number of private alleles per loci ( $\mathrm{P}_{\mathrm{a}}$ ) were calculated using GenAlEx v.6.5 (Peakall and Smouse, 2012). F statistics used in this analysis are described by Hartl and Clark (1997). Significance of variation among across land use types and size classes was determined using ANOVA. If significant differences were found Tukey's honestly significant difference (HSD) test was used to determine the source of differences. Diversity was compared using two variables: land use and DBH. DBH has been shown to be a good predictor of tree age in these three species (Jones et al. 2006). DBH size classes are described in Table 2. It should, however, be noted that trees do increase in diameter much more quickly in urban areas than in natural forests. The correlation used in this
study was found in natural areas, and will overestimate the age of urban trees (Bowles, personal communication). DBH groupings were selected in such a way to create roughly equal numbers of individuals in each class.

Table 2: Tree size classes. Size classes using DBH and approximate tree age for each class is given for each species (calculated using data from M. Bowles (unpublished data)).

|  | Small | Medium | Large |
| :--- | :---: | :---: | :---: |
| Red oak | $2.5-20 \mathrm{~cm}$ | $20-60 \mathrm{~cm}$ | $60+\mathrm{cm}$ |
|  | $(10-60$ years $)$ | $(60-165$ years $)$ | $(165+$ years $)$ |
| Sugar maple | $2.5-15 \mathrm{~cm}$ | $15-30 \mathrm{~cm}$ | $30+\mathrm{cm}$ |
|  | $(10-50$ years $)$ | $(50-110$ years $)$ | $(110+$ years $)$ |
| Green Ash | $2.5-20 \mathrm{~cm}$ | $20-35 \mathrm{~cm}$ | $35+\mathrm{cm}$ |
|  | $(10-60$ years $)$ | $(60-100$ years $)$ | $(100+$ years $)$ |

A suite of analyses were used to detect structure and differentiation among land use types. First, AMOVA and pairwise $\mathrm{F}_{\text {st }}$ values were calculated using GenAlEx. Methods used to calculate $\mathrm{F}_{\text {st }}$ are described by Nei (1977). Isolation by distance can explain the origin of genetic differentiation. To determine if geographic distance could explain differentiation in these populations $\mathrm{F}_{\text {st }}$ was compared to the distance between sites using Genepop (Raymond and Rousset 1995; Rousset 2008). Distance between sites was determined by measuring the distance of the centroid of each site.

To identify population structure that may not be apparent with standard population genetic approaches, a Bayesian approach was taken using the program STRUCTURE (Pritchard et al. 2000). STRUCTURE identifies genetic clusters (K) that correspond to population structure. For each species, independent runs were carried out for each K . Twenty iterations were run using a 10,000 burn in period and data for 50,000 iterations were collected. The most likely K was selected from 1-25 theoretical populations using the procedure detailed in Evanno et al. (2005).

An unweighted pair group method for arithmetic mean (UPGMA) was also used to visualize relationships among individual trees. To do this, genetic distance was first determined using GenAlEx. This matrix was used to create a UPGMA tree with the program PHYLIP v. 3.6 (Felsenstein 2005). This tree was visualized using FigTree v. 1.4 (Rambaut 2012).

## Results

Red oak
No diversity measures varied significantly for red oak across land use types. Table 3 shows the raw values for each diversity measure across all sites and land use types. Table 4 shows the results of an ANOVA that compared each variable across the land use types. Both of these tables show no variation among sites nor land use types. Diversity measures did vary significantly among DBH size classes, showing that diversity has changed over time (Tables 5 and 6). Post hoc Tukey's HSD on $\mathrm{N}_{\mathrm{e}}$, I and F (the three diversity values that showed significant variation) all showed that there was no significant difference between small and medium trees, but small and medium trees were both significantly different from large trees ( $\mathrm{p}<0.01$ ).

Table 3：Summary of diversity data from all tree species in all sites and land use types． N is the number of individuals，Na the average number of alleles per loci，Ne the number of effective alleles per loci，I the Shannon information index，Ho the number of observed heterozygotes and He the expected， F is the difference of these two values $((\mathrm{He}-\mathrm{Ho}) / \mathrm{He})$ and Pa is the average number of private alleles in each population．

|  |  | Site | N | $\mathrm{N}_{\mathrm{a}}$ | $\mathbf{N e}^{\text {e }}$ | I | $\mathbf{H}_{0}$ | $\mathbf{H}_{\mathbf{e}}$ | F | $\mathbf{P}_{\text {a }}$ |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | $\begin{aligned} & \text { 粷 } \\ & \text { 而 } \end{aligned}$ | Northwest | 9 | 8.20 | 5.655 | 1.824 | 0.767 | 0.778 | 0.039 | 0.40 |
|  |  | Northeast | 17 | 11.10 | 7.297 | 2.040 | 0.665 | 0.816 | 0.184 | 1.40 |
|  |  | Southwest | 10 | 9.90 | 7.335 | 2.096 | 0.730 | 0.850 | 0.146 | 1.30 |
|  |  | Southeast | 23 | 12.70 | 7.170 | 2.142 | 0.717 | 0.830 | 0.138 | 2.30 |
|  | $\begin{gathered} \tilde{0} \\ \text { ش } \end{gathered}$ | Labagh | 18 | 11.60 | 7.364 | 2.105 | 0.739 | 0.834 | 0.119 | 1.90 |
|  |  | Chavelier | 26 | 13.20 | 8.118 | 2.165 | 0.735 | 0.823 | 0.103 | 2.90 |
|  |  | Dan Ryan | 13 | 9.50 | 6.392 | 1.964 | 0.731 | 0.815 | 0.108 | 1.80 |
|  | 药 | Somme | 32 | 15.20 | 8.101 | 2.251 | 0.720 | 0.840 | 0.151 | 3.40 |
|  |  | Morton | 12 | 10.60 | 7.510 | 2.120 | 0.708 | 0.846 | 0.168 | 1.00 |
|  |  | Busse | 28 | 13.10 | 7.686 | 2.090 | 0.718 | 0.794 | 0.111 | 1.60 |
|  | $\begin{aligned} & \text { 苐 } \\ & \text { 荷 } \end{aligned}$ | Northeast | 18 | 4.80 | 2.855 | 1.020 | 0.411 | 0.492 | 0.193 | 0.80 |
|  |  | Northwest | 21 | 5.20 | 2.521 | 0.963 | 0.360 | 0.468 | 0.487 | 1.40 |
|  |  | South central | 20 | 4.80 | 2.741 | 0.936 | 0.488 | 0.443 | －0．083 | 1.20 |
|  |  | South | 16 | 4.00 | 2.477 | 0.823 | 0.343 | 0.413 | 0.302 | 0.80 |
|  | $\begin{gathered} \text { E } \\ \text { شे } \end{gathered}$ | Montrose | 12 | 4.40 | 2.799 | 0.984 | 0.374 | 0.474 | 0.361 | 1.20 |
|  |  | Chavelier | 12 | 4.60 | 3.290 | 0.985 | 0.367 | 0.461 | 0.140 | 1.20 |
|  |  | Labagh | 11 | 4.20 | 2.351 | 0.932 | 0.373 | 0.475 | 0.280 | 0.80 |
|  |  | Dan Ryan | 7 | 2.80 | 1.826 | 0.575 | 0.286 | 0.292 | －0．013 | 0.00 |
|  | $\begin{aligned} & E \\ & \text { En } \end{aligned}$ | Somme | 27 | 5.60 | 2.218 | 0.944 | 0.322 | 0.451 | 0.474 | 1.60 |
|  |  | Morton | 26 | 4.60 | 2.353 | 0.929 | 0.346 | 0.470 | 0.326 | 1.00 |
|  |  | Busse | 27 | 5.00 | 2.378 | 0.962 | 0.320 | 0.485 | 0.395 | 1.20 |
| $\begin{aligned} & \tilde{\pi} \\ & \pi \\ & \tilde{U} \\ & \tilde{U} \end{aligned}$ | $\begin{aligned} & \text { 鸦 } \\ & \vec{U} \end{aligned}$ | Northeast | 15 | 6.50 | 3.325 | 1.415 | 0.517 | 0.671 | 0.220 | 0.25 |
|  |  | Northwest | 20 | 6.75 | 3.059 | 1.247 | 0.500 | 0.596 | 0.179 | 0.25 |
|  |  | Southeast | 18 | 6.00 | 3.094 | 1.249 | 0.520 | 0.614 | 0.170 | 0.50 |
|  |  | Southwest | 16 | 5.00 | 2.732 | 1.118 | 0.484 | 0.574 | 0.213 | 0.25 |
|  | $\begin{gathered} \tilde{0} \\ \text { 合 } \end{gathered}$ | Montrose | 9 | 5.25 | 3.519 | 1.333 | 0.594 | 0.668 | 0.118 | 0.25 |
|  |  | Chavelier | 17 | 6.75 | 3.784 | 1.499 | 0.568 | 0.699 | 0.202 | 0.75 |
|  |  | Labagh | 16 | 6.75 | 3.40 | 1.436 | 0.656 | 0.684 | 0.024 | 0.25 |
|  |  | Dan Ryan | 24 | 7.25 | 2.754 | 1.325 | 0.536 | 0.631 | 0.140 | 0.50 |
|  | $\begin{aligned} & \text { EI } \\ & \text { In } \end{aligned}$ | Somme | 23 | 7.50 | 3.778 | 1.538 | 0.512 | 0.720 | 0.289 | 0.75 |
|  |  | Busse | 25 | 7.25 | 3.722 | 1.539 | 0.617 | 0.723 | 0.151 | 0.25 |

Table 4：Comparison of diversity values from Table 3 using ANOVA across land use types．This table shows the $p$ values of each ANOVA result．There are no significant variances in any measure for any species（ $p>0.05$ ）．See Table 3 for definitions of terms．

|  | $\mathbf{N}_{\mathbf{a}}$ | $\mathbf{N}_{\mathbf{e}}$ | $\mathbf{I}$ | $\mathbf{H}_{\mathbf{0}}$ | $\mathbf{H}_{\mathbf{e}}$ | $\mathbf{F}$ | $\mathbf{P}_{\mathbf{a}}$ |
| :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- |
| Red oak | 0.327 | 0.321 | 0.404 | 0.674 | 0.065 | 0.289 | 0.460 |
| Sugar maple | 0.137 | 0.563 | 0.701 | 0.187 | 0.627 | 0.341 | 0.387 |
| Green ash | 0.209 | 0.110 | 0.158 | 0.857 | 0.181 | 0.184 | 0.588 |

Table 5: Summary of diversity data for each species in all land use types, divided by size classes. See Table 3 for definitions of terms.

|  |  | Size class | N |  | $\mathrm{Na}_{\mathrm{a}}$ | $\mathbf{N}_{\mathrm{e}}$ | I | $\mathrm{H}_{0}$ | $\mathbf{H}_{\mathbf{e}}$ | F | $\mathbf{P a}_{\text {a }}$ |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  | Small |  | 26 | 13.6 | 8.015 | 2.197 | 0.750 | 0.838 | 0.105 | 3.6 |
|  |  | Medium |  | 17 | 10.9 | 7.447 | 2.056 | 0.706 | 0.814 | 0.133 | 1.3 |
|  |  | Large |  | 13 | 9.5 | 6.363 | 1.954 | 0.746 | 0.810 | 0.084 | 1.6 |
|  | $\begin{aligned} & \text { E } \\ & \text { شे } \end{aligned}$ | Small |  | 24 | 12.8 | 7.766 | 2.143 | 0.696 | 0.823 | 0.163 | 2.4 |
|  |  | Medium |  | 21 | 12.6 | 7.779 | 2.194 | 0.686 | 0.844 | 0.192 | 2.8 |
|  |  | Large |  | 13 | 9.9 | 6.413 | 1.991 | 0.792 | 0.814 | 0.024 | 1.4 |
|  | $\underset{\sim}{E}$ | Small |  | 25 | 14.1 | 7.629 | 2.199 | 0.738 | 0.832 | 0.12 | 2.8 |
|  |  | Medium |  | 32 | 14.7 | 8.560 | 2.259 | 0.688 | 0.840 | 0.192 | 2.7 |
|  |  | Large |  | 15 | 9.9 | 6.672 | 1.936 | 0.748 | 0.788 | 0.074 | 0.5 |
|  | $\frac{\vec{H}}{\underset{A}{4}}$ | Small |  | 23 | 5.2 | 2.895 | 1.034 | 0.383 | 0.505 | 0.381 | 0.8 |
|  |  | Medium |  | 31 | 6.8 | 2.923 | 1.086 | 0.430 | 0.488 | 0.421 | 2.4 |
|  |  | Large |  | 21 | 4.4 | 2.726 | 0.886 | 0.378 | 0.451 | 0.232 | 1.0 |
|  | $\begin{gathered} \tilde{\partial} \\ \text { के } \end{gathered}$ | Small |  | 15 | 5.2 | 3.285 | 1.074 | 0.339 | 0.505 | 0.438 | 1.6 |
|  |  | Medium |  | 16 | 4.6 | 2.581 | 0.917 | 0.383 | 0.462 | 0.073 | 1.2 |
|  |  | Large |  | 11 | 4.2 | 2.364 | 0.902 | 0.345 | 0.460 | 0.309 | 0.8 |
|  | E | Small |  | 45 | 6.2 | 2.250 | 0.960 | 0.326 | 0.454 | 0.483 | 2.6 |
|  |  | Medium |  | 30 | 4.8 | 2.260 | 0.903 | 0.330 | 0.476 | 0.345 | 1.2 |
|  |  | Large |  | 5 | 3.2 | 2.649 | 0.902 | 0.350 | 0.547 | 0.402 | 0.4 |
| $\begin{aligned} & \tilde{\tilde{n}} \\ & \tilde{ت} \\ & \tilde{U} \\ & 0 \end{aligned}$ | $\frac{\vec{H}}{\frac{\pi}{a}}$ | Small |  | 2 | 2.5 | 2.317 | 0.747 | 0.500 | 0.438 | -0.156 | 0.0 |
|  |  | Medium |  | 29 | 7.75 | 3.365 | 1.380 | 0.500 | 0.636 | 0.219 | 0.75 |
|  |  | Large |  | 38 | 8.0 | 3.131 | 1.325 | 0.509 | 0.629 | 0.216 | 0.75 |
|  | $\begin{aligned} & \text { E } \\ & \text { شे } \end{aligned}$ | Small |  | 34 | 9.75 | 3.700 | 1.576 | 0.628 | 0.706 | 0.105 | 1.25 |
|  |  | Medium |  | 13 | 5.5 | 3.552 | 1.426 | 0.545 | 0.704 | 0.220 | 0.0 |
|  |  | Large |  | 19 | 6.0 | 2.679 | 1.255 | 0.518 | 0.620 | 0.166 | 0.5 |
|  | $\begin{aligned} & E \\ & \text { تِ } \end{aligned}$ | Small |  | 31 | 8.5 | 3.649 | 1.571 | 0.549 | 0.720 | 0.233 | 1.0 |
|  |  | Medium |  | 11 | 4.75 | 3.501 | 1.341 | 0.568 | 0.687 | 0.157 | 0.0 |
|  |  | Large |  | 6 | 4.75 | 3.664 | 1.378 | 0.650 | 0.712 | 0.092 | 0.0 |

Table 6: Comparison of diversity values from Table 5 using ANOVA across size classes in each land use type. Only red oak values varied beyond the null expectation ( $\mathrm{p}<0.05$ ). See Table 3 for definitions.

|  | $\mathbf{N}_{\mathbf{a}}$ | $\mathbf{N}_{\mathbf{e}}$ | $\mathbf{I}$ | $\mathbf{H}_{\mathbf{0}}$ | $\mathbf{H}_{\mathbf{e}}$ | $\mathbf{F}$ | $\mathbf{P}_{\mathbf{a}}$ |
| :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- |
| Red oak | $0.018^{*}$ | $0.005^{* *}$ | $0.009^{* *}$ | 0.358 | 0.064 | $0.015^{*}$ | 0.052 |
| Sugar maple | 0.114 | 0.708 | 0.409 | 0.554 | 0.899 | 0.325 | 0.25 |
| Green ash | 0.906 | 0.76 | 0.929 | 0.891 | 0.8 | 0.417 | 0.509 |

AMOVA gave little indication of differentiation among land use types: $99 \%$ of variation was within land use types and only $1 \%$ among. Comparison of pairwise $\mathrm{F}_{\mathrm{st}}$ values for red oak also revealed little differentiation among sites, let alone among land use types (Table 7). Only a handful of the sites varied significantly from one another, and these differences were evenly
spread out among land use types. That is, planted sites were as varied from one another as they were from the remnant and spontaneous sites, and there is no evidence for variation among land use types.

Table 7: Pair wise comparison of $\mathrm{F}_{\text {st }}$ values for red oak. Each site is separated into its associated land use: planted, spontaneous and remnant.

|  | Planted |  |  |  | Spontaneous |  |  | Remnant |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | Northwest | Northeast | Southwest | Southeast | Labagh | Chavelier | Dan <br> Ryan | Somme | Busse |
| Planted |  |  |  |  |  |  |  |  |  |
| Northeast | 0.030* |  |  |  |  |  | * | p $<0.05$ |  |
| Southwest | 0.021 | 0.018 |  |  |  |  | ** | $0.05<\mathrm{p}<0.001$$\mathrm{D}<0.001$ |  |
| Southeast | 0.029 | 0.024 | 0.018 |  |  |  | *** |  |  |
| Spontan. |  |  |  |  |  |  |  |  |  |
| Labagh | 0.024 | 0.021 | 0.016 | 0.021 |  |  |  |  |  |
| Chavelier | 0.023 | 0.020 | 0.015 | 0.016 | 0.016 |  |  |  |  |
| Dan Ryan | 0.042** | 0.021 | 0.025 | 0.031* | 0.029* | 0.027 |  |  |  |
| Remnant |  |  |  |  |  |  |  |  |  |
| Somme | 0.024* | 0.014 | 0.012 | 0.017 | 0.016 | 0.013 | 0.020 |  |  |
| Busse | 0.026* | 0.012 | 0.013 | 0.016 | 0.019* | 0.015 | 0.020 | 0.011 |  |
| Morton | 0.026 | 0.019 | 0.018 | 0.024 | 0.020 | 0.019 | 0.031* | 0.019 | 0.016 |

Often times genetic differences among sites is correlated with distance. However, this does not seem to be the case for red oaks. Comparison of genetic distance to geographic distance showed no significant correlation (Figure 2).


Figure 2: Isolation by distance for red oak. There is not significant relationship between $\mathrm{F}_{\text {st }}$ and geographic distance in any land use type ( $\mathrm{p}>0.05$ ).

STRUCTURE analysis provided further evidence that the red oaks from different land use types have little to no genetic structure. No clear K value could be determined using the Evanno (2005) procedure. The best theoretical K value should be selected where the line plateaus when the theoretical $K$ is plotted against average $L(K)$, however, no such point exists (Figure 3A). Additionally, a peak should be evident at the best K value when theoretical K is plotted against mean $L(K)$, again, no such peak is displayed (Figure 3B). This suggests that there is no genetic structure and that oaks across all land use types are essentially interbreeding.


Figure 3: Determination of theoretical K for red oak using the Evanno method (2005). In Figure 2A a plateau should form at the correct value of $K$. In 2B a peak should form at the same value. Neither of these features is observed, indicating that the red oaks in the Chicago region are one interbreeding population with little genetic structure.

Because there is no genetic structure in red oaks an UPGMA was used to visualize relationships among individuals (Figure 4). Each individual is color coded to a specific site. If individuals from different sites or land use types clustered together on the dendogram there
would be evidence for genetic boundaries in the landscape. However, red oak individuals from different sites and land use types appear evenly spread throughout the dendogram, indicating no structure. There is one indication of differentiation in the UPGMA. In the lower right hand portion of the figure there are four individuals that are closely related both geographically and genetically (Planted049 - Planted053). These trees also have similar morphological qualities: all had a DBH of around 5 cm and identical leaf shape and branching structure.


Figure 4: UPGMA tree for red oaks. The planted sites are all in green tones, the spontaneous sites are red tones and the remnant sites are in tones of blue. Further information including the size and geographic location of each individual can be found in the appendix.

## Sugar maple

As with red oak, there was no significant variation of any diversity measure among land use types (Table 3and 4). There was not, however, significant variation among DBH classes, indicating that diversity has not changed over time (Table 5 and 6). AMOVA for sugar maple shows a little evidence of differentiation of populations in sugar maple: $96 \%$ of variation occurred within sites and $4 \%$ among sites. Pairwise $\mathrm{F}_{\text {st }}$ did reveal differences in genetic distances among sites (Table 8). Nearly all of the planted sites varied from the spontaneous and remnant sites, and only one of the remnant and spontaneous site pairs did not vary significantly. However, these differences did not seem to be based on distances between the sites (Figure 5). Comparison of genetic and geographic distance showed no correlation.

Table 8: Pairwise comparison of $\mathrm{F}_{\text {st }}$ values for sugar maple.

|  | Planted |  |  |  | Spontaneous |  |  |  | Remnant |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | Northwest | Northeast | Southwest | Southeast | Montrose | Labagh | Chavelier | Dan Ryan |  |  |
| Planted |  |  |  |  |  |  |  |  |  |  |
| Northeast | 0.030* |  |  |  |  |  |  | * | p < 0.05 |  |
| Southwest | 0.028 | 0.021 |  |  |  |  |  | ** | $0.05<p$ | < 0.001 |
| Southeast | 0.039* | 0.041* | 0.027 |  |  |  |  | *** | p $<0.00$ |  |
| Spontan. |  |  |  |  |  |  |  |  |  |  |
| Montrose | 0.057** | 0.019 | 0.027 | 0.050* |  |  |  |  |  |  |
| Labagh | 0.043* | 0.023 | 0.030 | 0.051* | 0.028 |  |  |  |  |  |
| Chavelier | 0.069** | 0.070*** | 0.050* | 0.019 | 0.069* | 0.072* |  |  |  |  |
| Dan Ryan | 0.119*** | 0.086* | 0.094** | 0.144** | 0.073 | 0.041 | 0.168** |  |  |  |
| Remnant |  |  |  |  |  |  |  |  |  |  |
| Somme | 0.027* | 0.040* | 0.026 | 0.045** | 0.043* | 0.026 | 0.068*** | 0.075* |  |  |
| Busse | 0.028* | 0.056*** | 0.041** | 0.050*** | 0.079*** | 0.063*** | 0.073*** | 0.141*** | 0.020* |  |
| Morton | 0.023 | 0.042** | 0.027* | 0.037** | 0.049** | 0.043*** | 0.063*** | 0.121*** | 0.016 | 0.017 |

## Sugar maple isolation by distance



Figure 5: Isolation by distance for sugar maple. There is not significant relationship between $\mathrm{F}_{\text {st }}$ and distance for any land use type ( $p>0.05$ ).

Structure analysis confirmed that there was some variation among land use types. It was determined that there were five theoretical populations using the procedure described by Evanno (2005). The curve in Figure 6A levels out at K equals five, and in Figure 6B there is a small peak at the same value. There is some room for interpretation in these results, as the structure is fairly weak, but the visual readout for $\mathrm{K}=5$ does highlight some interesting features about the sugar maple population.


Figure 6: Determining theoretical K for sugar maple. Figure 6 A shows a plateau at $\mathrm{K}=5$. This is further affirmed by Figure 6B, which shows a small peak at $\mathrm{K}=5$.

The structure analysis for $\mathrm{K}=5$ shows a large percentage of cluster three in the planted population while clusters one and two dominate the remnant population (Figure 7). Planted individuals are differentiated from remnant sites, and spontaneous individuals have intermediate characteristics (Figure 7B). The spontaneous sites have intermediate levels of clusters one two and three and these sites are dominated by clusters four and five. These clusters are less common
in the remnant and planted sites. It seems that the spontaneous sites do have some qualities that bridge between remnant and planted sites, but also have some unique characteristics (Figure 8). There is little indication that there is structure among sites. While there are many individuals in remnant land use types that are dominated by cluster one or two, these individuals are not separated by site.


Figure 7: Sugar maple structure for $K=5$. Figure 7A shows results for each individual, sorted by land use type and geographic proximity. Large gaps denote breaks in land use types and small gaps denote sites. 7B illustrates the average composition of each land use type. The planted trees have a large percentage of cluster three, showing that it is genetically distinct from the new and remnant populations.


Figure 8: Detail of structure results for sugar maples spontaneous sites.
There is no indication of structure by DBH (Figure 9). This is especially interesting for the planted sites. I expected that smaller trees would be very similar to one another (because of an abundance of cultivars) than the older trees which might have more remnant individuals. There is not strong evidence that this is the case.


Figure 9: STRUCTURE results for sugar maple sorted by DBH across land use types. DBH is sorted by small to large, with smallest trees on the left.

An UPGMA further illustrates the structure in the sugar maple populations (Figure 10). Many of the planted individuals are clustered into two sections of the analysis. Remnant trees are also loosely clustered together in two sections. Spontaneous trees do not cluster out discretely, but are sprinkled throughout the dendogram and are often in between the remnant and planted clusters. Additionally, individuals are clustered more by land use types than by sites. The clusters of planted and remnant trees are composed of a mix of sites.


Figure 10: UPGMA for sugar maple. The planted sites are all in green tones, the spontaneous sites are red tones and the remnant sites are in tones of blue. Further information including the size and geographic location of each individual can be found in the appendix.

## Green ash

Average number of alleles per locus did not vary significantly among land use types, nor did average number of private alleles, heterozygosity, F statistics, and Shannon's information
index. As with sugar maple and red oak, genetic diversity among all three land use types and DBH classes does not vary significantly (Table 3-5).

AMOVA for green ash was very similar to sugar maple: $95 \%$ of diversity existed within land use types and only $5 \%$ among land use types. Pairwise $\mathrm{F}_{\mathrm{st}}$ shows little variation within land use types (Table 9). However, 62.5\% of planted sites varied from spontaneous sites and $100 \%$ varied from remnant sites. Most spontaneous sites also varied from remnant sites.

Table 9: Pairwise comparison of $\mathrm{F}_{\mathrm{st}}$ values for green ash.


There is not a significant correlation between genetic distance and geographic distance for green ash (Figure 11). There is a positive correlation for spontaneous sites, but it is not significant $(\mathrm{p}=0.209)$.


Figure 11: Isolation by distance for green ash. There is not a significant relationship between $\mathrm{F}_{\text {st }}$ and distance in any land use type ( $\mathrm{p}>0.05$ ).

Structure analysis further illustrates that the city trees were differentiated from the remnant forests, although the differentiation was again weak. The determination of the number of theoretical populations is not clear, and there is room for interpretation. In Figure 12A the curve peak begins to level out at $K=5$, but there is a small peak in Figure 12B at $K=6$. For this analysis $K=6$ was selected, but the structure results for $K=5$ were very similar and the same conclusions would be made using that number of theoretical populations.


Figure 12: Determination of theoretical K for green ash. The correct K value is not readily apparent. The curve begins to level out at $K=5$ in $6 A$, but there is a slight peak in $K=6$ in $6 B$. $\mathrm{K}=6$ was selected for structure analysis.

Spontaneous and remnant forests appear to have nearly identical genetic structure to one another, but the planted trees look quite different from the forested areas (Figure 13A and B). The planted trees have a much higher percentage of clusters one and five and lower percentages of the other clusters. There is no indication of genetic structure among sites.


Figure 13: Green ash structure for $\mathrm{K}=6$. In 12A large gaps denote breaks in land use types and smaller gaps denote sites. There is little difference between spontaneous and remnant trees. However, the planted population has much more of clusters one and five. This is further illustrated by 12B.

Structure results that were sorted by DBH of individuals show an interesting pattern in the planted sites (Figure 14). Cluster five is has a much higher abundance in small planted trees than in larger ones. This indicates that genetically similar individuals (likely cultivars) are being planted more heavily in recent years. This was not shown to correlate with a drop in genetic
diversity (Table 5). However, DBH size classes for these analyses were chosen to create a roughly even distribution of individuals in each size class across all land use types. This resulted in the presence of only two small individuals in the planted sites. When size classes are redistributed in the planted sites a drop in all diversity measures in small trees is observed (Table 10). There are not enough individuals in this analysis to determine if the differences in diversity are significant.


Figure 14: Green ash structure results sorted by DBH and land use type.
Table 10: Diversity measures for planted green ash across size classes. Size classes have been redefined from Table 5 in order to make roughly even numbers of individuals in each size class.

|  | $\mathbf{N}$ | $\mathbf{N}_{\mathbf{a}}$ | $\mathbf{N}_{\mathbf{e}}$ | $\mathbf{I}$ | $\mathbf{H}_{\mathbf{0}}$ | $\mathbf{H}_{\mathbf{e}}$ | $\mathbf{F}$ | $\mathbf{P}_{\mathbf{a}}$ |
| :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- |
| Small | 22 | 5.250 | 2.312 | 0.954 | 0.489 | 0.487 | 0.030 | 0.250 |
| Medium | 22 | 7.500 | 3.395 | 1.414 | 0.527 | 0.662 | 0.224 | 0.408 |
| Large | 24 | 8.500 | 3.577 | 1.498 | 0.500 | 0.678 | 0.263 | 0.500 |

The majority of planted individuals are again clustered into two sections in the UPGMA
(Figure 15). Most of the trees that are in these planted clusters are small to medium individuals, again indicating that planting of genetically similar individuals has increased in recent years. However, unlike with the sugar maples the remnant individuals do not cluster at all. Instead, the remnant and spontaneous sections look more like red oak: entirely intermixed.


Figure 15: UPGMA for green ash. The planted sites are all in green tones, the spontaneous sites are red tones and the remnant sites are in tones of blue. Further information including the size and geographic location of each individual can be found in the appendix.

## Discussion

Genetic diversity across all three species did not vary across the land use gradient. No diversity measure varied significantly. In short, planted trees were not less diverse than spontaneous trees nor than remnant forests. While red oaks had identical genetic structure across
all three land use types, sugar maple and green ash city trees were differentiated from the remnant forests. The differences in population structure among these three species can be explained by the history of cultivation of each species.

## Red oak

The lack of genetic structure in red oak is likely due to two factors: a dearth of red oak cultivars and oak's ability to spread pollen over great distances. There are only two red oak cultivars currently in production, and both of them are uncommon (University of Connecticut Horticulture Website 2013). Red oaks are difficult to graft, and the majority of red oaks planted in urban areas are grown from seed. These seeds usually come from a small subset of parents that are known to produce reliable seedlings (Worfolk interview 2012). This practice could reduce genetic diversity in the city trees, except that oaks broadcast pollen over great distances, which allows for gene flow (Dow and Ashley 1996). There is sufficient gene flow into planted trees to not only maintain genetic diversity, but also to keep the red oak city trees from becoming genetically distinct.

There is one portion of the UPGMA results that indicates the potential for a decrease in red oak genetic diversity in the City of Chicago. Four planted oak trees were side by side along a city block (Figure 4). They had similar DBH (they were around 20 years old) and physical appearance (branching structure, bark, and tree health were identical). These four individuals were likely planted at the same time. These oaks were not clones of one another, but were very closely related. It is likely that they are full siblings or even the result of several generations of inbreeding. If these sorts of breeding practices become more widespread genetic diversity in the planted trees will decrease and its genetic structure will become distinct from remnant forests.

Even though there is potential for decreases in diversity with the planting of related individuals, it does not seem to be the current trend. Almost all of the measures of diversity in red oak showed significant variances across size classes (Table 6). Tukey’s HSD indicated that small and medium trees were significantly varied from large trees, and Table 5 shows that smaller trees have higher diversity measures than larger ones. The UPGMA results showed that planted trees are overall not different from remnant sites, so it is not likely that the increase in diversity in coming from this source. The Chicago urban forest (like most developed areas) has become fragmented through urbanization. One would expect this to cause barriers to breeding among disparate populations, which would likely lead to decreased diversity in populations, and eventually lead to inbreeding and decreased population fitness. That does not seem to be the case with these red oaks, and very few other studies have documented a drop in diversity in fragmented populations of long lived, wind dispersed species (see Kramer et al. (2008) for a review). Some studies have even shown an increase in genetic diversity in fragmented forests (Fore et al. 1992; Muir et al. 2004; Young et al. 1993). The species in each of these studies were self incompatible, like red oak (Cottam et al. 1982). It is suspected that fragmentation encourages self incompatible species to accept pollen from other fragmented populations, leading to overall higher outcrossing rates than would be seen in a continuous forest (Kramer et al. 2008). This may be the case with the oaks in this study.

## Sugar maple

Diversity measures among land use types for sugar maple were not significantly different, but AMOVA, $\mathrm{F}_{\mathrm{st}}$, STRUCTURE and UPGMA suggested differences among the three populations. The structure results did not indicate strong differentiation across land use types, but
there were some key difference between the sites. City trees have far more of the cluster three, while the remnant forest is dominated by clusters one and two (Figure 7). UPGMA also showed clustering of planted trees in two different sections (Figure 10). Sugar maple has been widely planted in urban areas for over a century (Worfolk interview 2012), and unlike red oak, there are a number of maple cultivars, some of which have been available since the 1960's (Flemer 1962). It is likely that heavy cultivation and wide planting of cultivars in the city has led to a genetic divergence from the sugar maples in remnant forests.

The presence of a large number of sugar maple cultivars (which are often clonal) in the city was expected to reduce the genetic diversity in city trees when compared to remnant trees. However, even though planted individuals are differentiated from other forests, there were no significant differences in diversity measures between these land use types (Tables 3 and 4). It is possible that a number of different cultivars are being planted in the Chicago region, and this serves to keep diversity comparable to remnant forests, even though the populations are differentiated. UPGMA results indicate that this is the case. There are two distinct clusters of sugar maples that are genetically similar to one another but distinct from the rest of the sampled individuals. These trees could be different sugar maple cultivars. Planted trees in the urban forest seem to be made up of several cultivars and a number of autochthonous or wild type trees. This combination keeps diversity as high as in remnant forests.

I predicted that spontaneous trees would have elements of genetic structure from both planted trees and remnant forests. Sugar maples have been shown to have high capacity for gene flow (Fore et al. 1992), allowing city trees to interact with nearby forests, which could alter their genetic identity. To some extent, the STRUCTURE results show that this may be the case (Figure 7). Clusters one and two dominated in the remnant forests, while cluster three (and to a
lesser four and five) were the most abundant in the city trees. Some of the spontaneous population did look similar to remnant forests, but other parts were more similar to the planted trees. UPGMA results further illustrate these relationships (Figure 10). There are two clusters of planted individuals and two clusters of remnant individuals in the sugar maple dendogram. Spontaneous trees are not entirely intermixed with either land use type, but instead are scattered in the areas between with planted and remnant clusters: some spontaneous trees are more like planted trees and others are more like remnant forests. The reason for this differentiation is not immediately evident. It seemed likely that younger trees in spontaneous sites would have more in common genetically with planted trees than older trees. Over time more cultivars have been developed and planted and urbanization has continued to push closer to wild areas. It would stand to reason that these planted trees would have an increasing influence on spontaneous trees in more recent years. To see if this is the case I sorted STRUCTURE results by DBH (Figure 9). The clusters associated with city trees (cluster three and to a lesser extent four) were not more prevalent in the smaller trees in the spontaneous sites. Age of trees cannot explain the apparent influence of planted trees on the spontaneous sites.

Instead differences within the spontaneous trees fell by site (Figure 8). Clusters three and five were mostly present in Catherine Chevalier Woods, cluster four was largely found in Montrose Point Bird Sanctuary and Dan Ryan Forest Preserve, while Labagh Woods was dominated by clusters one and two (Figure 8). Aerial maps from 1939 show that Labagh Woods was largely unchanged from current forest cover. Chevalier Woods was forested in the early $20^{\text {th }}$ century but much of the surrounding land was being farmed, while the areas sampled in Dan Ryan Preserve and Montrose Point were largely denuded of trees (Figure 16). Structure results show that the continuously forested Labagh Woods was not differentiated from remnant forests
(see Figure 7 for comparison). The disturbance and reforestation that has occurred in Chevalier Woods, Montrose Point and Dan Ryan Preserve caused a change in their genetic structure. These sites are now largely composed of clusters that are more associated with city trees than remnant forests. These results indicate that while sugar maple is able to interbreed among land use types, the city trees do not seem to influence the spontaneous trees unless disturbance of some sort has occurred.




Figure 16: Comparison of 1939 aerial maps to satellite images from 2010-2013 of trees sampled in spontaneous sites. The black and white images are from 1939 and the colored ones are current. The dots on the current maps note locations of samples trees.

## Green ash

Green ash and sugar maple results were very similar, as are the species’ cultivation histories. Genetic diversity did not vary among land use types for green ash, but AMOVA, pairwise $\mathrm{F}_{\text {st }}$, STRUCTURE analysis and UPGMA suggest weak population differentiation among land use types. In the city trees there were two dominant clusters: one and five (Figure 13). There are also two groupings of planted individuals in the UPGMA (Figure 15). These results could indicate that there are two different cultivars that are prevalent in the city, neither of which is present in the other land use types. Genotyping of known cultivars would be necessary to confirm this hypothesis. Green ash cultivars have been developed since the 1970's (Klehm
and Klehm 1973), and a variety of them are currently planted in the City of Chicago (Scott unpublished data). Planted green ash in the City of Chicago seem to be composed of several different cultivars and a number of autochthonous trees, and this combination keeps genetic diversity in the urban forest on the same level as what exists in remnant areas.

There were several indications that green ash diversity is changing over time. STRUCTURE results sorted by the size of the trees showed large percentage of a single cluster in the smallest trees (Figure 14). These trees were genetically distinct from the older planted trees. Furthermore, the trees that clustered together in the UPGMA results had relatively small DBHs. I predicted that an abundance of cultivars would result in a decrease in genetic diversity. When size classes were redefined to have roughly equal numbers of individuals in each class it became apparent that this was the case (Table 10). An abundance of a few cultivars without a large number of autochthonous trees can result in reduced genetic diversity in the urban forest.

Unlike sugar maple, there was no structure between spontaneous and remnant sites. The STRUCTURE readout for green ash was nearly identical for spontaneous and remnant forests (Figure 13). In the UPGMA results individuals from spontaneous and remnant sites were completely intermixed (Figure 15). The spontaneous areas bore no similarity to the city trees, indicating that there is no gene flow between these land use types. This is even true at disturbed sites where the sugar maples were differentiated from the remnant forests. This is at first surprising, as green ash has nearly identical reproductive strategies as sugar maple. They are wind pollinated and have seeds that are wind dispersed. However, green ash pollen has been shown to spread less far than sugar maple pollen (Kennedy et al. 1990, Fore et al. 1992). This could reduce its ability to interact with spontaneous areas. Additionally, the majority of green ash cultivars that are intentionally planted in urban areas are either male or seedless varieties
(University of Connecticut Horticulture Website 2013). This means that green ash cultivars are only able to interact with spontaneous areas through pollen transfer, and because green ash has been shown to only move pollen from 60-80 m (Kennedy et al. 1990), its ability to breed with spontaneous trees is also reduced. Only green ash cultivars that are planted very close to spontaneous areas would have any ability to interbreed in these sites, and even then they may only be able to interact with trees on the perimeter of the spontaneous site.

Diversity results for green ash are especially interesting given that ash trees are under eminent threat from the emerald ash borer. It has been shown that some ash individuals are more able to cope with the borer than others. Marshall et al. (2013) showed that susceptibility to the borer varied with the roughness of the trunk, but more data is required to determine if these morphological variations have a genetic basis.

## Comparisons of genetic diversity to other regions

None of these species had significant changes in diversity across an urban to remnant gradient. However, it is uncertain whether this is caused by urban areas having relatively high diversity or remnant areas having low diversity. To determine which is the case, this data set needs to be compared with data from other remnant forests. This is possible for red oaks. Aldrich et al. quantified red oak diversity in a large remnant forest in northern Indiana (2003). They found diversity levels that were very similar to those found in this study using nearly identical microsatellite primers. This shows that planted red oaks in the City of Chicago are not only as genetically diverse as trees in surrounding forests, but in healthy, large forests. There are no comparable studies for sugar maple or green ash. The levels of diversity found in this study are similar to what was reported in the primer notes (Pandey et al. 2004; Lefort and Frascaria-

Lacoste 1999). However, these measurements were for different species (A. psuedoplatanus and F. excelsior). We can make some inferences on the possible state of genetic diversity of these species in remnant forests based on the ecological history of the area. Both sugar maple and green ash were somewhat rare before mesophication of Illinois forests (Nowacki et al. 2008). The area surrounding the City of Chicago was historically fire prone, which selected for oak species and against more mesophytic species like ashes and maples. This could mean that all of the ash and maples in the region have come from a fairly small founder population, and could have comparatively low genetic diversity. To be certain if this is the case additional individuals should be sampled at the heart of green ash and sugar maple diversity.

## Past and future changes in genetic diversity

Only red oaks had significant variation across size classes (Tables 5 and 6). Although smaller red oaks were found to have higher neutral diversity than larger ones, there are several indications that genetic diversity in the urban forest may decrease in coming years. STRUCTURE and UPGMA analyses showed that sugar maple and green ash city trees were genetically distinct from the remnant areas. City trees from these species are dominated by cultivars, and if a reduced number of cultivars are planted diversity could decrease and the city trees will become further differentiated. This already seems to be happening with green ash. The smallest green ashes in the planted trees are less diverse than larger trees and spontaneous and remnant sites (Table 10). This is also possible with red oaks. The current diversity in red oaks is high, and there is no differentiation from planted trees to remnant forests. However, UPGMA results show that some planted individuals are very closely related, and with the increased
planting of closely related individuals diversity in the city trees could plummet, causing this land use type to become genetically differentiated from remnant forests.

In order to maintain or increase genetic diversity in the urban forest managers need to plant trees from multiple sources. This could be achieved by planting a variety of cultivars and by planting trees from seed sources from multiple areas. Adding diversity to the urban forest can be done in a way to maximize sustainability. In the context of climate change it may be advantageous to start planting cultivars (or source trees from other regions) that are known to be better adapted to drier and warmer conditions. By making small changes in sourcing practices the urban forest could be made more robust and potentially better able to withstand diseases, pests and climate change.

## Application for other species and sites

The results from this study can be applied to other tree species with caution. This study showed that the genetic diversity in city trees is dependent upon the trees cultivation history and its interactions with the surrounding forests. The species in this study were chosen because they are common in forests surrounding the City of Chicago. This is not the case for most trees in the urban forest. Many of the trees planted in Chicago are not native to North America (e.g. little leaf linden (Tilia cordata Mill.) and Norway maple (Acer platanoides L.)) while others are native but are not common (e.g. catalpa (Catalpa speciosa Englem), and Kentucky coffee tree (Gymnocladus dioicus K. Koch)). These species would have no influence from the surrounding forest, and their diversity would be completely dependent upon what was planted anthropogenically. Their genetic diversity would, like the trees in this study, be influenced by the cultivation history of the species; cultivars could reduce diversity but with the presence of a
variety of cultivars high diversity can be achieved. Some trees like hackberry (Celtis occidentalis L.) and ironwood (Ostrya virginiana K. Koch) could have very similar results to the species in this study. They are native, common and have similar cultivation history. Species in other cities would likely have similar diversity results, given that the cultivation history and influences from surrounding forests were the same.

## Conclusion

Red oak, sugar maple and green ash in the Chicago region do not appear to have reduced genetic diversity, as measured by standard diversity measures, across an urban to remnant gradient. However, the genetic structure of green ash and sugar maple from city streets and parks is distinct from remnant forests. This pattern was not observed in red oak, which had no genetic structure across land use types. Although the urban forest currently has comparable genetic diversity to remnant forests, there is indication that the diversity might change in the future. There was one cluster of red oak street trees that were nearly genetically identical to one another. Red oak cultivars are currently rare, but these trees indicate that the planting of full siblings or inbred individuals can reduce diversity. On the other hand, green ash and sugar maple cultivars are very common, and STRUCTURE and UPGMA results indicate that they are frequently planted. There are currently several different cultivars planted along with autochthonous, which maintains genetic diversity that is comparable to remnant forests. But if a single cultivar is planted in mass it could greatly reduce genetic diversity of a species in the urban forest.

The state of genetic diversity in the urban forest is dictated by decisions made by city foresters. Currently trees are sourced from nurseries with little concern for their genetic makeup, and although efforts are made to increase species diversity, little thought is put into intraspecies
diversity (Santamour1990). The genetic diversity in the urban forests could be one of the major predictors of the forests ability to withstand stressors (Sork et al. 2013). A sustainable, resilient forest will need to have a mixture of genotypes. To attain this, foresters should not only increase species diversity but intraspecies diversity by planting an increased number of cultivars and seedlings from a variety of parents.

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Appendix: List of all individuals sampled. Data includes identification number and species; the land use, site and geographic location of the individual; also the tree's DBH and health.

| ID | Species | Landuse | Site | Lat | Long | DBH | Health |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| AM01 | A. saccharum | Spontaneous | Montrose sanctuary | 41.96249 | -87.63448 | 29.5 | ood |
| AM02 | A. saccharum | Spontaneous | Montrose sanctuary | 41.96232 | -87.63453 | 35.5 | poor |
| AM03 | A. saccharum | Spontaneous | Montrose sanctuary | 41.96273 | -87.63432 | 29.7 | good |
| AM04 | A. saccharum | Spontaneous | Montrose sanctuary | 41.96257 | -87.63433 | 34.9 | good |
| AM05 | A. saccharum | Spontaneous | Montrose sanctuary | 41.96251 | -87.63386 | 53.0 | ood |
| AM06 | A. saccharum | Spontaneous | Montrose sanctuary | 41.9625 | -87.63408 | 2.7 | air |
| AM07 | A. saccharum | Spontaneous | Montrose sanctuary | 41.96251 | -87.63411 | 23.8 | fair |
| AM08 | A. saccharum | Spontaneous | Montrose sanctuary | 41.96235 | -87.63366 | 51.7 | good |
| AM09 | A. saccharum | Spontaneous | Montrose sanctuary | 41.96248 | -87.63359 | 37.2 | good |
| AM10 | A. saccharum | Spontaneous | Montrose sanctuary | 41.9625 | -87.63357 | 40.6 | good |
| AM101 | A. saccharum | Spontaneous | Catherine Chevalier | 41.97671 | -87.84969 | 17.5 | good |
| AM102 | A. saccharum | Spontaneous | Catherine Chevalier | 41.97605 | -87.84944 | 16.5 | good |
| AM103 | A. saccharum | Spontaneous | Catherine Chevalier | 41.97594 | -87.84936 | 11.2 | air |
| AM104 | A. saccharum | Spontaneous | Catherine Chevalier | 41.97598 | -87.84913 | 11.9 | poor |
| AM105 | A. saccharum | Spontaneous | Catherine Chevalier | 41.97601 | -87.84899 | 21.2 | poor |
| AM106 | A. saccharum | Spontaneous | Catherine Chevalier | 41.97597 | -87.84853 | 8.7 | good |
| AM107 | A. saccharum | Spontaneous | Catherine Chevalier | 41.97582 | -87.84836 | 12.2 | good |
| AM109 | A. saccharum | Spontaneous | Catherine Chevalier | 41.97574 | -87.84839 | 36.1 | good |
| AM109 | A. saccharum | Spontaneous | Catherine Chevalier | 41.97585 | -87.84916 | 11.3 | fair |
| AM11 | A. saccharum | Spontaneous | Montrose sanctuary | 41.96247 | -87.63351 | 63.5 | good |
| AM110 | A. saccharum | Spontaneous | Catherine Chevalier | 41.97563 | -87.84925 | 17.6 | fair |
| AM111 | A. saccharum | Spontaneous | Catherine Chevalier | 41.97972 | -87.84942 | 31.5 | fair |
| AM112 | A. saccharum | Spontaneous | Catherine Chevalier | 41.97989 | -87.84685 | 37.6 | fair |
| AM12 | A. saccharum | Spontaneous | Montrose sanctuary | 41.96269 | -87.63281 | 19.8 | good |
| AM13 | A. saccharum | Spontaneous | Labagh woods | 41.97943 | -87.74465 | 5.2 | good |
| AM14 | A. saccharum | Spontaneous | Labagh woods | 41.97956 | -87.74481 | 2.8 | good |
| AM15 | A. saccharum | Spontaneous | Labagh woods | 41.97958 | -87.74422 | 2.5 | good |
| AM16 | A. saccharum | Spontaneous | Labagh woods | 41.97979 | -87.74422 | 9.3 | good |
| AM17 | A. saccharum | Spontaneous | Labagh woods | 41.98023 | -87.74369 | 10.4 | good |
| AM18 | A. saccharum | Spontaneous | Labagh woods | 41.98204 | -87.7454 | 23.1 | good |
| AM19 | A. saccharum | Spontaneous | Labagh woods | 41.98193 | -87.74559 | 2.5 | good |
| AM20 | A. saccharum | Spontaneous | Labagh woods | 41.9819 | -87.74573 | 20.4 | good |
| AM21 | A. saccharum | Spontaneous | Labagh woods | 41.98232 | -87.74322 | 8.3 | good |
| AM22 | A. saccharum | Spontaneous | Labagh woods | 41.98324 | -87.743 | 7.4 | fair |
| AM23 | A. saccharum | Spontaneous | Labagh woods | 41.98238 | -87.74287 | 18.0 | good |
| AM51 | A. saccharum | Spontaneous | Dan Ryan Preserve | 41.7387 | -87.67645 | 10.1 | good |
| AM52 | A. saccharum | Spontaneous | Dan Ryan Preserve | 41.7387 | -87.67654 | 18.8 | fair |
| AM53 | A. saccharum | Spontaneous | Dan Ryan Preserve | 41.73879 | -87.67657 | 23.8 | fair |
| AM54 | A. saccharum | Spontaneous | Dan Ryan Preserve | 41.73678 | -87.67423 | 16.9 | good |
| AM55 | A. saccharum | Spontaneous | Dan Ryan Preserve | 41.73668 | -87.67449 | 15.5 | good |
| AM56 | A. saccharum | Spontaneous | Dan Ryan Preserve | 41.73679 | -87.67469 | 16.0 | fair |
| AM57 | A. saccharum | Spontaneous | Dan Ryan Preserve | 41.73697 | -87.67485 | 30.1 | fair |
| AN01 | A. saccharum | Remnant | Somme | 42.13849 | -87.82129 | 16.4 | good |
| AN02 | A. saccharum | Remnant | Somme | 42.13851 | -87.82137 | 26.3 | good |
| AN03 | A. saccharum | Remnant | Somme | 42.13882 | -87.82144 | 12.1 | good |


| AN04 | A. saccharum | Remnant | Somme | 42.13953 | -87.81117 | 25.2 | good |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| AN05 | A. saccharum | Remnant | Somme | 42.14058 | -87.8134 | 13.0 | fair |
| AN06 | A. saccharum | Remnant | Somme | 42.14047 | -87.81436 | 32.4 | good |
| AN07 | A. saccharum | Remnant | Somme | 42.14069 | -87.81599 | 6.2 | fair |
| AN08 | A. saccharum | Remnant | Somme | 42.13978 | -87.81801 | 66.5 | good |
| AN09 | A. saccharum | Remnant | Somme | 42.13928 | -87.81909 | 6.0 | good |
| AN10 | A. saccharum | Remnant | Somme | 42.13921 | -87.81917 | 6.2 | good |
| AN101 | A. saccharum | Remnant | Morton | 41.81663 | -88.04152 | 15.7 | poor |
| AN102 | A. saccharum | Remnant | Morton | 41.81559 | -88.04128 | 6.4 | good |
| AN103 | A. saccharum | Remnant | Morton | 41.81633 | -88.04162 | 4.3 | good |
| AN104 | A. saccharum | Remnant | Morton | 41.81602 | -88.04077 | 7.2 | good |
| AN105 | A. saccharum | Remnant | Morton | 41.81532 | -88.04066 | 21.4 | fair |
| AN106 | A. saccharum | Remnant | Morton | 41.81527 | -88.04089 | 12.3 | fair |
| AN107 | A. saccharum | Remnant | Morton | 41.81432 | -88.04031 | 24.4 | good |
| AN108 | A. saccharum | Remnant | Morton | 41.81373 | -88.04079 | 25.8 | fair |
| AN109 | A. saccharum | Remnant | Morton | 41.81545 | -88.04658 | 2.5 | good |
| AN109 | A. saccharum | Remnant | Morton | 41.81359 | -88.0406 | 22.2 | good |
| AN11 | A. saccharum | Remnant | Somme | 42.13942 | -87.81988 | 74.0 | good |
| AN110 | A. saccharum | Remnant | Morton | 41.81311 | -88.04043 | 8.8 | good |
| AN111 | A. saccharum | Remnant | Morton | 41.81258 | -88.04056 | 14.4 | good |
| AN112 | A. saccharum | Remnant | Morton | 41.81272 | -88.04158 | 19.8 | good |
| AN113 | A. saccharum | Remnant | Morton | 41.81288 | -88.0415 | 25.9 | good |
| AN114 | A. saccharum | Remnant | Morton | 41.81284 | -88.04189 | 15.1 | good |
| AN115 | A. saccharum | Remnant | Morton | 41.81377 | -88.0456 | 7.1 | poor |
| AN116 | A. saccharum | Remnant | Morton | 41.81343 | -88.04312 | 21.0 | good |
| AN117 | A. saccharum | Remnant | Morton | 41.81351 | -88.04454 | 18.5 | fair |
| AN118 | A. saccharum | Remnant | Morton | 41.81341 | -88.04707 | 24.0 | good |
| AN119 | A. saccharum | Remnant | Morton | 41.8144 | -88.04827 | 10.1 | good |
| AN12 | A. saccharum | Remnant | Somme | 42.14078 | -87.82669 | 11.3 | fair |
| AN120 | A. saccharum | Remnant | Morton | 41.81467 | -88.04798 | 6.4 | fair |
| AN121 | A. saccharum | Remnant | Morton | 41.81522 | -88.04723 | 17.4 | good |
| AN122 | A. saccharum | Remnant | Morton | 41.81547 | -88.04645 | 4.0 | good |
| AN123 | A. saccharum | Remnant | Morton | 41.81573 | -88.04644 | 14.3 | good |
| AN124 | A. saccharum | Remnant | Morton | 41.81527 | -88.04624 | 19.9 | good |
| AN125 | A. saccharum | Remnant | Morton | 41.81618 | -88.04552 | 8.4 | fair |
| AN126 | A. saccharum | Remnant | Morton | 41.81778 | -88.04269 | 29.0 | good |
| AN127 | A. saccharum | Remnant | Morton | 41.81808 | -88.04159 | 19.8 | fair |
| AN128 | A. saccharum | Remnant | Morton | 41.81731 | -88.04244 | 11.1 | good |
| AN129 | A. saccharum | Remnant | Morton | 41.81715 | -88.0434 | 14.3 | fair |
| AN13 | A. saccharum | Remnant | Somme | 42.14264 | -87.8243 | 11.5 | poor |
| AN14 | A. saccharum | Remnant | Somme | 42.14265 | -87.82423 | 6.3 | fair |
| AN14b | A. saccharum | Remnant | Somme | 42.1398 | -87.81991 | 5.6 | good |
| AN15 | A. saccharum | Remnant | Somme | 42.13988 | -87.81982 | 55.0 | fair |
| AN16 | A. saccharum | Remnant | Somme | 42.13982 | -87.81991 | 16.2 | good |
| AN17 | A. saccharum | Remnant | Somme | 42.14028 | -87.8168 | 7.4 | poor |
| AN18 | A. saccharum | Remnant | Somme | 42.14034 | -87.81781 | 5.2 | good |
| AN19 | A. saccharum | Remnant | Somme | 42.14038 | -87.81764 | 16.2 | good |
| AN20 | A. saccharum | Remnant | Somme | 42.14255 | -87.81698 | 5.0 | good |
| AN21 | A. saccharum | Remnant | Somme | 42.14259 | -87.81746 | 6.3 | fair |


| AN23 | A. saccharum | Remnant | Somme | 42.14414 | -87.82191 | 15.9 | good |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| AN24 | A. saccharum | Remnant | Somme | 42.14392 | -87.82186 | 11.0 | good |
| AN25 | A. saccharum | Remnant | Somme | 42.14391 | -87.82172 | 6.3 | good |
| AN26 | A. saccharum | Remnant | Somme | 42.14382 | -87.82161 | 23.8 | good |
| AN27 | A. saccharum | Remnant | Somme | 42.14101 | -87.82134 | 21.8 | good |
| AN28 | A. saccharum | Remnant | Somme | 42.1408 | -87.82123 | 19.3 | good |
| AN50 | A. saccharum | Remnant | Busse | 42.0383 | -88.00255 | 15.3 | good |
| AN50 | A. saccharum | Remnant | Busse | 42.03795 | -88.00245 | 83.0 | fair |
| AN51 | A. saccharum | Remnant | Busse | 42.03621 | -88.00266 | 11.1 | good |
| AN52 | A. saccharum | Remnant | Busse | 42.03916 | -88.00269 | 7.6 | good |
| AN53 | A. saccharum | Remnant | Busse | 42.03941 | -88.00274 | 16.0 | good |
| AN54 | A. saccharum | Remnant | Busse | 42.03967 | -88.00292 | 7.1 | good |
| AN55 | A. saccharum | Remnant | Busse | 42.03971 | -88.00396 | 5.0 | good |
| AN56 | A. saccharum | Remnant | Busse | 42.0397 | -88.00404 | 12.4 | good |
| AN57 | A. saccharum | Remnant | Busse | 42.03966 | -88.00411 | 7.5 | good |
| AN58 | A. saccharum | Remnant | Busse | 42.03934 | -88.00505 | 9.3 | fair |
| AN59 | A. saccharum | Remnant | Busse | 42.03801 | -88.00518 | 29.0 | good |
| AN60 | A. saccharum | Remnant | Busse | 42.038 | -88.00543 | 77.3 | good |
| AN61 | A. saccharum | Remnant | Busse | 42.03812 | -88.00619 | 13.7 | good |
| AN62 | A. saccharum | Remnant | Busse | 42.03816 | -88.00634 | 20.5 | fair |
| AN63 | A. saccharum | Remnant | Busse | 42.03741 | -88.00644 | 6.6 | good |
| AN64 | A. saccharum | Remnant | Busse | 42.03706 | -88.00718 | 17.8 | good |
| AN65 | A. saccharum | Remnant | Busse | 42.03642 | -88.00922 | 6.1 | good |
| AN66 | A. saccharum | Remnant | Busse | 42.03601 | -88.00803 | 8.1 | good |
| AN67 | A. saccharum | Remnant | Busse | 42.037 | -88.006 | 16.3 | fair |
| AN68 | A. saccharum | Remnant | Busse | 42.03543 | -88.0046 | 12.1 | good |
| AN69 | A. saccharum | Remnant | Busse | 42.03512 | -88.00434 | 12.1 | good |
| AN70 | A. saccharum | Remnant | Busse | 42.03508 | -88.00405 | 16.8 | good |
| AN71 | A. saccharum | Remnant | Busse | 42.03495 | -88.00393 | 15.7 | good |
| AN72 | A. saccharum | Remnant | Busse | 42.03526 | -88.00331 | 6.6 | good |
| AN74 | A. saccharum | Remnant | Busse | 42.03553 | -88.00291 | 7.3 | good |
| AN75 | A. saccharum | Remnant | Busse | 42.03561 | -88.00299 | 10.5 | good |
| AN76 | A. saccharum | Remnant | Busse | 42.03536 | -88.00322 | 12.7 | good |
| AN76 | A. saccharum | Remnant | Busse | 42.03452 | -87.99978 | 11.5 | good |
| AU01 | A. saccharum | Planted | Northeast | 41.93965 | -87.67651 | 21.2 | fair |
| AU02 | A. saccharum | Planted | Northwest | 41.99043 | -87.73347 | 14.1 | good |
| AU03 | A. saccharum | Planted | Northwest | 41.99566 | -87.73827 | 34.7 | fair |
| AU04 | A. saccharum | Planted | Northwest | 41.99578 | -87.73823 | 36.2 | poor |
| AU05 | A. saccharum | Planted | Northwest | 41.99573 | -87.73499 | 44.5 | poor |
| AU06 | A. saccharum | Planted | Northwest | 41.99253 | -87.73508 | 8.3 | good |
| AU07 | A. saccharum | Planted | Northwest | 41.99218 | -87.73487 | 13.7 | good |
| AU08 | A. saccharum | Planted | Northwest | 41.99085 | -87.73377 | 22.2 | good |
| AU09 | A. saccharum | Planted | Northwest | 41.99312 | -87.73381 | 67.2 | good |
| AU10 | A. saccharum | Planted | Northwest | 41.99325 | -87.73388 | 38.8 | good |
| AU11 | A. saccharum | Planted | Northwest | 41.99333 | -87.73254 | 34.9 | good |
| AU12 | A. saccharum | Planted | Northwest | 41.99245 | -87.73256 | 48.2 | good |
| AU13 | A. saccharum | Planted | Northwest | 41.99294 | -87.73256 | 34.6 | poor |
| AU14 | A. saccharum | Planted | Northwest | 41.99221 | -87.73355 | 43.6 | fair |
| AU15 | A. saccharum | Planted | Northwest | 41.97511 | -87.70739 | 21.8 | fair |


| AU16 | A. saccharum | Planted | Northeast | 41.91325 | -87.62966 | 21.0 | good |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| AU17 | A. saccharum | Planted | Northeast | 41.91322 | -87.62961 | 22.0 | good |
| AU18 | A. saccharum | Planted | Northeast | 41.9132 | -87.62949 | 22.4 | good |
| AU19 | A. saccharum | Planted | Northeast | 41.91381 | -87.63058 | 16.2 | good |
| AU20 | A. saccharum | Planted | Northeast | 41.91846 | -87.64054 | 27.2 | poor |
| AU21 | A. saccharum | Planted | Northeast | 41.91057 | -87.64092 | 21.2 | poor |
| AU22 | A. saccharum | Planted | Northeast | 41.89834 | -87.67469 | 11.7 | good |
| AU23 | A. saccharum | Planted | Northeast | 41.89816 | -87.67961 | 18.0 | good |
| AU24 | A. saccharum | Planted | Northeast | 41.96984 | -87.64729 | 32.0 | poor |
| AU25 | A. saccharum | Planted | Northeast | 41.96254 | -87.64323 | 59.6 | fair |
| AU26 | A. saccharum | Planted | Northeast | 41.96871 | -87.64686 | 6.2 | good |
| AU27 | A. saccharum | Planted | Northeast | 41.93783 | -87.67568 | 25.1 | good |
| AU28 | A. saccharum | Planted | Northeast | 41.93507 | -87.67801 | 44.5 | fair |
| AU29 | A. saccharum | Planted | Northeast | 41.93881 | -87.67309 | 21.1 | good |
| AU30 | A. saccharum | Planted | Northeast | 41.9369 | -87.66959 | 39.6 | fair |
| AU31 | A. saccharum | Planted | Northeast | 41.93686 | -87.6698 | 43.5 | poor |
| AU32 | A. saccharum | Planted | Northeast | 41.93581 | -87.67426 | 14.3 | good |
| AU33 | A. saccharum | Planted | Northwest | 41.92717 | -87.71218 | 7.5 | good |
| AU34 | A. saccharum | Planted | Northwest | 41.92641 | -87.71337 | 52.6 | good |
| AU35 | A. saccharum | Planted | Northwest | 41.92842 | -87.71089 | 7.1 | poor |
| AU36 | A. saccharum | Planted | Northwest | 41.9254 | -87.70702 | 18.5 | good |
| AU37 | A. saccharum | Planted | Northwest | 41.92499 | -87.70697 | 11.2 | poor |
| AU38 | A. saccharum | Planted | Northwest | 41.92273 | -87.70747 | 19.5 | good |
| AU39 | A. saccharum | Planted | Northwest | 41.92417 | -87.70747 | 15.4 | good |
| AU40 | A. saccharum | Planted | South Central | 41.85385 | -87.66144 | 17.2 | fair |
| AU41 | A. saccharum | Planted | South Central | 41.85389 | -87.66142 | 12.2 | fair |
| AU42 | A. saccharum | Planted | South Central | 41.85245 | -87.66506 | 14.9 | fair |
| AU43 | A. saccharum | Planted | South Central | 41.85244 | -87.66526 | 10.6 | poor |
| AU44 | A. saccharum | Planted | South Central | 41.85246 | -87.66552 | 20.3 | good |
| AU45 | A. saccharum | Planted | South Central | 41.82513 | -87.67024 | 12.2 | good |
| AU46 | A. saccharum | Planted | South Central | 41.82501 | -87.66661 | 21.4 | good |
| AU47 | A. saccharum | Planted | South Central | 41.82468 | -87.68023 | 27.5 | good |
| AU48 | A. saccharum | Planted | South Central | 41.82461 | -87.68025 | 26.7 | good |
| AU49 | A. saccharum | Planted | South Central | 41.82455 | -87.68025 | 22.2 | good |
| AU50 | A. saccharum | Planted | South Central | 41.82258 | -87.6831 | 30.9 | poor |
| AU51 | A. saccharum | Planted | South Central | 41.82319 | -87.68333 | 19.8 | good |
| AU52 | A. saccharum | Planted | South Central | 41.82315 | -87.68356 | 15.9 | good |
| AU53 | A. saccharum | Planted | South Central | 41.82325 | -87.68353 | 10.1 | good |
| AU54 | A. saccharum | Planted | South Central | 41.82325 | -87.68365 | 17.3 | good |
| AU55 | A. saccharum | Planted | South Central | 41.82318 | -87.68363 | 16.5 | good |
| AU56 | A. saccharum | Planted | South Central | 41.82645 | -87.68331 | 16.5 | good |
| AU57 | A. saccharum | Planted | South Central | 41.82618 | -87.68295 | 9.1 | good |
| AU58 | A. saccharum | Planted | South Central | 41.82622 | -87.68264 | 7.9 | good |
| AU59 | A. saccharum | Planted | South Central | 41.82623 | -87.68268 | 8.4 | good |
| AU59 | A. saccharum | Planted | South | 41.78436 | -87.71088 | 23.3 | poor |
| AU60 | A. saccharum | Planted | South | 41.78432 | -87.71102 | 24.5 | fair |
| AU61 | A. saccharum | Planted | South | 41.78419 | -87.71444 | 34.2 | good |
| AU62 | A. saccharum | Planted | South | 41.78065 | -87.71627 | 35.3 | good |
| AU63 | A. saccharum | Planted | South | 41.77782 | -87.72003 | 9.0 | good |


| AU64 | A. saccharum Planted | South | 41.77693 | -87.71515 | 34.2 | good |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| AU65 | A. saccharum Planted | South | 41.77772 | -87.71529 | 34.5 | fair |
| AU66 | A. saccharum Planted | South | 41.77302 | -87.65757 | 23.7 | good |
| AU67 | A. saccharum Planted | South | 41.77312 | -87.6568 | 9.4 | good |
| AU68 | A. saccharum Planted | South | 41.77312 | -87.65654 | 9.1 | good |
| AU69 | A. saccharum Planted | South | 41.77328 | -87.65611 | 14.5 | fair |
| AU70 | A. saccharum Planted | South | 41.77325 | -87.65696 | 15.2 | fair |
| AU71 | A. saccharum Planted | South | 41.77634 | -87.65594 | 25.5 | good |
| AU72 | A. saccharum Planted | South | 41.76239 | -87.59135 | 16.3 | good |
| AU73 | A. saccharum Planted | South | 41.76929 | -87.60109 | 53.5 | good |
| AU74 | A. saccharum Planted | South | 41.79342 | -87.59644 | 11.2 | good |
| AU75 | A. saccharum Planted | South | 41.79338 | -87.59643 | 10.1 | good |
| FM01 | F. pennslyvanica Spontaneous | Montrose sanctuary | 41.96231 | -87.63399 | 16.6 | fair |
| FM02 | F. pennslyvanica Spontaneous | Montrose sanctuary | 41.96247 | -87.6346 | 5.0 | good |
| FM03 | F. pennslyvanica Spontaneous | Montrose sanctuary | 41.96249 | -87.63489 | 2.5 | good |
| FM04 | F. pennslyvanica Spontaneous | Montrose sanctuary | 41.96241 | -87.63414 | 12.2 | poor |
| FM05 | F. pennslyvanica Spontaneous | Montrose sanctuary | 41.96241 | -87.63414 | 8.4 | fair |
| FM06 | F. pennslyvanica Spontaneous | Montrose sanctuary | 41.96235 | -87.63417 | 3.7 | poor |
| FM07 | F. pennslyvanica Spontaneous | Montrose sanctuary | 41.96243 | -87.63379 | 5.3 | fair |
| FM08 | F. pennslyvanica Spontaneous | Montrose sanctuary | 41.96257 | -87.63375 | 24.3 | fair |
| FM09 | F. pennslyvanica Spontaneous | Montrose sanctuary | 41.96262 | -87.63304 | 6.8 | fair |
| FM10 | F. pennslyvanica Spontaneous | Labagh woods | 41.97637 | -87.74253 | 27.0 | fair |
| FM101 | F. pennslyvanica Spontaneous | Catherine Chevalier | 41.97659 | -87.84737 | 54.8 | fair |
| FM102 | F. pennslyvanica Spontaneous | Catherine Chevalier | 41.9767 | -87.84982 | 24.4 | poor |
| FM103 | F. pennslyvanica Spontaneous | Catherine Chevalier | 41.97645 | -87.84974 | 24.6 | poor |
| FM104 | F. pennslyvanica Spontaneous | Catherine Chevalier | 41.97672 | -87.85013 | 23.6 | fair |
| FM105 | F. pennslyvanica Spontaneous | Catherine Chevalier | 41.97608 | -87.84987 | 18.4 | fair |
| FM106 | F. pennslyvanica Spontaneous | Catherine Chevalier | 41.97602 | -87.8489 | 42.3 | fair |
| FM107 | F. pennslyvanica Spontaneous | Catherine Chevalier | 41.97615 | -87.84883 | 65.2 | poor |
| FM108 | F. pennslyvanica Spontaneous | Catherine Chevalier | 41.9758 | -87.84985 | 11.1 | poor |
| FM109 | F. pennslyvanica Spontaneous | Catherine Chevalier | 41.97593 | -87.85001 | 18.2 | poor |
| FM11 | F. pennslyvanica Spontaneous | Labagh woods | 41.97704 | -87.74193 | 33.7 | fair |
| FM110 | F. pennslyvanica Spontaneous | Catherine Chevalier | 41.9675 | -87.85288 | 19.2 | fair |
| FM111 | F. pennslyvanica Spontaneous | Catherine Chevalier | 41.96773 | -87.853 | 15.1 | poor |
| FM112 | F. pennslyvanica Spontaneous | Catherine Chevalier | 41.96764 | -87.85268 | 14.3 | fair |
| FM113 | F. pennslyvanica Spontaneous | Catherine Chevalier | 41.977 | -87.85164 | 9.7 | good |
| FM114 | F. pennslyvanica Spontaneous | Catherine Chevalier | 41.97715 | -87.85171 | 13.7 | fair |
| FM115 | F. pennslyvanica Spontaneous | Catherine Chevalier | 41.97702 | -87.85203 | 27.6 | poor |
| FM116 | F. pennslyvanica Spontaneous | Catherine Chevalier | 41.97706 | -87.85217 | 18.8 | poor |
| FM117 | F. pennslyvanica Spontaneous | Catherine Chevalier | 41.97716 | -87.8524 | 16.2 | fair |
| FM12 | F. pennslyvanica Spontaneous | Labagh woods | 41.97752 | -87.7417 | 32.5 | fair |
| FM13 | F. pennslyvanica Spontaneous | Labagh woods | 41.97839 | -87.74155 | 71.1 | good |
| FM14 | F. pennslyvanica Spontaneous | Labagh woods | 41.97856 | -87.74411 | 41.4 | fair |
| FM15 | F. pennslyvanica Spontaneous | Labagh woods | 41.97955 | -87.74467 | 9.8 | fair |
| FM16 | F. pennslyvanica Spontaneous | Labagh woods | 41.98027 | -87.74363 | 36.2 | good |
| FM19 | F. pennslyvanica Spontaneous | Labagh woods | 41.98057 | -87.74355 | 46.1 | poor |
| FM20 | F. pennslyvanica Spontaneous | Labagh woods | 41.98169 | -87.74302 | 57.3 | fair |
| FM21 | F. pennslyvanica Spontaneous | Labagh woods | 41.98151 | -87.74323 | 4.5 | good |
| FM22 | F. pennslyvanica Spontaneous | Labagh woods | 41.98188 | -87.74391 | 38.2 | fair |


| FM23 | F. pennslyvanica Spontaneous | Labagh woods | 41.98209 | -87.74557 | 9.7 | poor |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| FM24 | F. pennslyvanica Spontaneous | Labagh woods | 41.98218 | -87.74724 | 16.9 | r |
| FM25 | F. pennslyvanica Spontaneous | Labagh woods | 41.98206 | -87.74624 | 17.5 | good |
| FM26 | F. pennslyvanica Spontaneous | Labagh woods | 41.98231 | -87.7419 | 81.9 | or |
| FM27 | F. pennslyvanica Spontaneous | Labagh woods | 41.98182 | -87.7412 | 30.1 | or |
| FM28 | F. pennslyvanica Spontaneous | Labagh woods | 41.98161 | -87.74053 | 74.1 | oor |
| FM51 | F. pennslyvanica Spontaneous | Dan Ryan Preserve | 41.73938 | -87.68222 | 21.4 | or |
| FM52 | F. pennslyvanica Spontaneous | Dan Ryan Preserve | 41.73863 | -87.68223 | 11.8 | fair |
| FM53 | F. pennslyvanica Spontaneous | Dan Ryan Preserve | 41.73817 | -87.68194 | 26.0 | fair |
| FM54 | F. pennslyvanica Spontaneous | Dan Ryan Preserve | 41.73807 | -87.68176 | 6.5 | air |
| FM55 | F. pennslyvanica Spontaneous | Dan Ryan Preserve | 41.73805 | -87.68203 | 52.3 | good |
| FM56 | F. pennslyvanica Spontaneous | Dan Ryan Preserve | 41.73812 | -87.68225 | 35.0 | fair |
| FM57 | F. pennslyvanica Spontaneous | Dan Ryan Preserve | 41.73813 | -87.68228 | 24.7 | poor |
| FM58 | F. pennslyvanica Spontaneous | Dan Ryan Preserve | 41.73804 | -87.68208 | 47.6 | fair |
| FM58 | F. pennslyvanica Spontaneous | Dan Ryan Preserve | 41.73811 | -87.68232 | 55.2 | air |
| FM59 | F. pennslyvanica Spontaneous | Dan Ryan Preserve | 41.73808 | -87.68211 | 10.2 | poor |
| FM60 | F. pennslyvanica Spontaneous | Dan Ryan Preserve | 41.73907 | -87.6769 | 9.0 | poor |
| FM61 | F. pennslyvanica Spontaneous | Dan Ryan Preserve | 41.73864 | -87.67648 | 63.5 | poor |
| FM62 | F. pennslyvanica Spontaneous | Dan Ryan Preserve | 41.73844 | -87.67612 | 11.3 | fair |
| FM63 | F. pennslyvanica Spontaneous | Dan Ryan Preserve | 41.73839 | -87.67625 | 17.8 | poor |
| FM64 | F. pennslyvanica Spontaneous | Dan Ryan Preserve | 41.73853 | -87.67644 | 54.5 | poor |
| FM65 | F. pennslyvanica Spontaneous | Dan Ryan Preserve | 41.73854 | -87.67645 | 72.5 | fair |
| FM66 | F. pennslyvanica Spontaneous | Dan Ryan Preserve | 41.73854 | -87.67664 | 37.7 | poor |
| FM67 | F. pennslyvanica Spontaneous | Dan Ryan Preserve | 41.73844 | -87.67645 | 17.7 | fair |
| FM69 | F. pennslyvanica Spontaneous | Dan Ryan Preserve | 41.74047 | -87.67703 | 4.2 | good |
| FM70 | F. pennslyvanica Spontaneous | Dan Ryan Preserve | 41.74485 | -87.67961 | 17.0 | fair |
| FM71 | F. pennslyvanica Spontaneous | Dan Ryan Preserve | 41.74435 | -87.68119 | 13.1 | poor |
| FM72 | F. pennslyvanica Spontaneous | Dan Ryan Preserve | 41.74434 | -87.6811 | 13.0 | poor |
| FM73 | F. pennslyvanica Spontaneous | Dan Ryan Preserve | 41.73674 | -87.67466 | 39.9 | poor |
| FM74 | F. pennslyvanica Spontaneous | Dan Ryan Preserve | 41.73716 | -87.67507 | 18.8 | fair |
| FM75 | F. pennslyvanica Spontaneous | Dan Ryan Preserve | 41.73714 | -87.67515 | 11.2 | fair |
| FM76 | F. pennslyvanica Spontaneous | Dan Ryan Preserve | 41.73673 | -87.67482 | 34.5 | fair |
| FM77 | F. pennslyvanica Spontaneous | Dan Ryan Preserve | 41.73619 | -87.67468 | 18.7 | fair |
| FN01 | F. pennslyvanica Remnant | Somme | 42.1392 | -87.82268 | 28.4 | poor |
| FN02 | F. pennslyvanica Remnant | Somme | 42.13835 | -87.81044 | 9.2 | poor |
| FN03 | F. pennslyvanica Remnant | Somme | 42.13883 | -87.81091 | 6.2 | poor |
| FN04 | F. pennslyvanica Remnant | Somme | 42.13909 | -87.81105 | 31.5 | poor |
| FN10 | F. pennslyvanica Remnant | Somme | 42.13893 | -87.82443 | 15.4 | poor |
| FN11 | F. pennslyvanica Remnant | Somme | 42.14061 | -87.82683 | 13.5 | poor |
| FN12 | F. pennslyvanica Remnant | Somme | 42.14062 | -87.82592 | 9.7 | poor |
| FN13 | F. pennslyvanica Remnant | Somme | 42.14044 | -87.82566 | 23.0 | fair |
| FN14 | F. pennslyvanica Remnant | Somme | 42.14036 | -87.82553 | 25.6 | fair |
| FN15 | F. pennslyvanica Remnant | Somme | 42.14214 | -87.82514 | 7.4 | good |
| FN16 | F. pennslyvanica Remnant | Somme | 42.14202 | -87.82512 | 14.0 | good |
| FN17 | F. pennslyvanica Remnant | Somme | 42.14265 | -87.82414 | 14.6 | fair |
| FN18 | F. pennslyvanica Remnant | Somme | 42.14262 | -87.82414 | 8.0 | fair |
| FN19 | F. pennslyvanica Remnant | Somme | 42.13976 | -87.81985 | 17.0 | fair |
| FN20 | F. pennslyvanica Remnant | Somme | 42.13988 | -87.8193 | 31.1 | good |
| FN21 | F. pennslyvanica Remnant | Somme | 42.13988 | -87.81981 | 33.7 | good |


| FN22 | F. pennslyvanica Remnant | Somme | 42.14385 | -87.81929 | 13.7 | poor |
| :--- | :--- | :--- | :--- | :--- | :--- | :--- |
| FN23 | F. pennslyvanica Remnant | Somme | 42.14253 | -87.82223 | 24.3 | poor |
| FN24 | F. pennslyvanica Remnant | Somme | 42.142 | -87.82254 | 17.9 | poor |
| FN25 | F. pennslyvanica Remnant | Somme | 42.14193 | -87.82272 | 17.1 | poor |
| FN26 | F. pennslyvanica Remnant | Somme | 42.14182 | -87.82292 | 21.7 | poor |
| FN27 | F. pennslyvanica Remnant | Somme | 42.14098 | -87.82143 | 12.7 | good |
| FN28 | F. pennslyvanica Remnant | Somme | 42.14101 | -87.82142 | 28.0 | fair |
| FN50 | F. pennslyvanica Remnant | Busse | 42.04023 | -88.00277 | 21.6 | poor |
| FN51 | F. pennslyvanica Remnant | Busse | 42.04021 | -88.00289 | 37.6 | poor |
| FN52 | F. pennslyvanica Remnant | Busse | 42.03964 | -88.00433 | 37.0 | fair |
| FN53 | F. pennslyvanica Remnant | Busse | 42.03951 | -88.00486 | 7.1 | poor |
| FN54 | F. pennslyvanica Remnant | Busse | 42.03798 | -88.00586 | 54.3 | poor |
| FN55 | F. pennslyvanica Remnant | Busse | 42.03805 | -88.00604 | 38.3 | fair |
| FN56 | F. pennslyvanica Remnant | Busse | 42.03764 | -88.00659 | 15.0 | poor |
| FN57 | F. pennslyvanica Remnant | Busse | 42.03648 | -88.00875 | 43.7 | poor |
| FN58 | F. pennslyvanica Remnant | Busse | 42.03626 | -88.00705 | 15.7 | poor |
| FN59 | F. pennslyvanica Remnant | Busse | 42.0368 | -88.0064 | 5.5 | poor |
| FN60 | F. pennslyvanica Remnant | Busse | 42.03751 | -88.00338 | 9.2 | poor |
| FN61 | F. pennslyvanica Remnant | Busse | 42.03748 | -88.00333 | 5.3 | fair |
| FN62 | F. pennslyvanica Remnant | Busse | 42.03752 | -88.00319 | 14.9 | fair |
| FN63 | F. pennslyvanica Remnant | Busse | 42.03738 | -88.00312 | 18.6 | fair |
| FN64 | F. pennslyvanica Remnant | Busse | 41.89831 | -87.67464 | 23.8 | poor |
| FN65 | F. pennslyvanica Remnant | Busse | 41.99684 | -87.73375 | 77.5 | poor |
| FN66 | F. pennslyvanica Remnant | Busse | 41.97879 | -87.73045 | 89.1 | poor |
| FN67 | F. pennslyvanica Remnant | Busse | 41.97807 | -87.73304 | 65.4 | poor |
| FN68 | F. pennslyvanica Remnant | Busse | -87.73369 | 64.2 | fair |  |
| FN69 | F. pennslyvanica Remnant | Busse |  | 42.03234 | -88.00655 | 6.9 |


| FU18 | F. pennslyvanica Planted | Northeast | 41.89841 | -87.67967 | 51.0 | fair |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| FU19 | F. pennslyvanica Planted | Northeast | 41.90008 | -87.67972 | 55.5 | fair |
| FU20 | F. pennslyvanica Planted | Northeast | 41.90227 | -87.68121 | 28.3 | good |
| FU23 | F. pennslyvanica Planted | Northeast | 41.9698 | -87.64732 | 52.6 | fair |
| FU24 | F. pennslyvanica Planted | Northeast | 41.96988 | -87.64768 | 37.8 | fair |
| FU25 | F. pennslyvanica Planted | Northeast | 41.93715 | -87.63344 | 26.0 | poor |
| FU26 | F. pennslyvanica Planted | Northeast | 41.94585 | -87.64029 | 57.5 | fair |
| FU27 | F. pennslyvanica Planted | Northeast | 41.9375 | -87.63225 | 36.4 | fair |
| FU28 | F. pennslyvanica Planted | Northeast | 41.93787 | -87.6707 | 49.4 | good |
| FU29 | F. pennslyvanica Planted | Northwest | 41.926 | -87.71212 | 30.1 | good |
| FU30 | F. pennslyvanica Planted | Northeast | 41.93691 | -87.67008 | 51.7 | fair |
| FU31 | F. pennslyvanica Planted | Northwest | 41.92712 | -87.71345 | 27.3 | good |
| FU32 | F. pennslyvanica Planted | Northwest | 41.92767 | -87.71334 | 59.3 | good |
| FU33 | F. pennslyvanica Planted | Northwest | 41.92651 | -87.70717 | 25.0 | poor |
| FU34 | F. pennslyvanica Planted | Northwest | 41.92522 | -87.70723 | 30.5 | fair |
| FU35 | F. pennslyvanica Planted | Northwest | 41.92312 | -87.70741 | 37.3 | poor |
| FU36 | F. pennslyvanica Planted | Northwest | 41.92411 | -87.70748 | 66.9 | poor |
| FU37 | F. pennslyvanica Planted | Northwest | 41.92423 | -87.70755 | 30.3 | poor |
| FU38 | F. pennslyvanica Planted | Northwest | 41.92683 | -87.70765 | 50.4 | good |
| FU39 | F. pennslyvanica Planted | Northwest | 41.9269 | -87.70929 | 31.4 | fair |
| FU40 | F. pennslyvanica Planted | Southwest | 41.84787 | -87.68123 | 26.1 | fair |
| FU41 | F. pennslyvanica Planted | Southwest | 41.84746 | -87.68117 | 33.0 | good |
| FU42 | F. pennslyvanica Planted | Southwest | 41.83239 | -87.67658 | 39.7 | fair |
| FU43 | F. pennslyvanica Planted | Southwest | 41.8337 | -87.67654 | 24.3 | fair |
| FU44 | F. pennslyvanica Planted | Southwest | 41.83272 | -87.67043 | 35.6 | good |
| FU45 | F. pennslyvanica Planted | Southwest | 41.82452 | -87.67035 | 40.3 | fair |
| FU46 | F. pennslyvanica Planted | Southwest | 41.82503 | -87.66666 | 32.9 | poor |
| FU47 | F. pennslyvanica Planted | Southwest | 41.82458 | -87.67995 | 38.1 | fair |
| FU48 | F. pennslyvanica Planted | Southwest | 41.82457 | -87.68032 | 21.0 | fair |
| FU49 | F. pennslyvanica Planted | Southwest | 41.82313 | -87.68841 | 34.2 | good |
| FU50 | F. pennslyvanica Planted | Southwest | 41.82594 | -87.68404 | 30.3 | good |
| FU50B | F. pennslyvanica Planted | Southwest | 41.78416 | -87.7134 | 64.8 | good |
| FU51 | F. pennslyvanica Planted | Southwest | 41.78434 | -87.71545 | 28.7 | good |
| FU52 | F. pennslyvanica Planted | Southwest | 41.78458 | -87.71566 | 23.0 | good |
| FU53 | F. pennslyvanica Planted | Southwest | 41.78142 | -87.71602 | 80.8 | fair |
| FU54 | F. pennslyvanica Planted | Southwest | 41.78064 | -87.71641 | 44.4 | fair |
| FU55 | F. pennslyvanica Planted | Southeast | 41.77292 | -87.65758 | 35.5 | good |
| FU56 | F. pennslyvanica Planted | Southeast | 41.77305 | -87.6562 | 39.1 | fair |
| FU57 | F. pennslyvanica Planted | Southeast | 41.77347 | -87.65584 | 50.5 | poor |
| FU58 | F. pennslyvanica Planted | Southeast | 41.77346 | -87.65595 | 79.4 | fair |
| FU59 | F. pennslyvanica Planted | Southeast | 41.77628 | -87.65605 | 45.0 | good |
| FU60 | F. pennslyvanica Planted | Southeast | 41.77625 | -87.65604 | 42.5 | good |
| FU60B | F. pennslyvanica Planted | Southeast | 41.76333 | -87.59317 | 52.5 | poor |
| FU61 | F. pennslyvanica Planted | Southeast | 41.77624 | -87.65602 | 42.8 | fair |
| FU62 | F. pennslyvanica Planted | Southeast | 41.76413 | -87.58836 | 22.1 | good |
| FU63 | F. pennslyvanica Planted | Southeast | 41.7689 | -87.5965 | 107.0 | good |
| FU64 | F. pennslyvanica Planted | Southeast | 41.76777 | -87.60218 | 51.7 | good |
| FU65 | F. pennslyvanica Planted | Southeast | 41.78648 | -87.60856 | 16.0 | good |
| FU66 | F. pennslyvanica Planted | Southeast | 41.78645 | -87.60851 | 13.9 | good |


| FU67 | F. pennslyvanica | Planted | Southeast | 41.78651 | -87.60841 | 24.5 | good |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| FU68 | F. pennslyvanica | Planted | Southeast | 41.78662 | -87.60849 | 31.8 | good |
| FU69 | F. pennslyvanica | Planted | Southeast | 41.78975 | -87.60821 | 74.6 | good |
| FU70 | F. pennslyvanica | Planted | Southeast | 41.79833 | -87.61502 | 39.2 | good |
| FU75 | F. pennslyvanica | Planted | Southeast | 41.74516 | -87.67883 | 31.1 | poor |
| QM01 | Q. rubra | Spontaneous | Labagh woods | 41.97851 | -87.74407 | 97.6 | fair |
| QM02 | Q. rubra | Spontaneous | Labagh woods | 41.97968 | -87.74446 | 16.8 | good |
| QM03 | Q. rubra | Spontaneous | Labagh woods | 41.97981 | -87.74416 | 6.6 | good |
| QM04 | Q. rubra | Spontaneous | Labagh woods | 41.9805 | -87.74307 | 62.4 | fair |
| QM05 | Q. rubra | Spontaneous | Labagh woods | 41.98071 | -87.74304 | 62.7 | fair |
| QM06 | Q. rubra | Spontaneous | Labagh woods | 41.98157 | -87.74422 | 11.3 | good |
| QM07 | Q. rubra | Spontaneous | Labagh woods | 41.98159 | -87.74396 | 4.7 | fair |
| QM08 | Q. rubra | Spontaneous | Labagh woods | 41.98206 | -87.74557 | 2.5 | fair |
| QM09 | Q. rubra | Spontaneous | Labagh woods | 41.98185 | -87.74633 | 11.3 | fair |
| QM10 | Q. rubra | Spontaneous | Labagh woods | 41.98403 | -87.74789 | 23.8 | fair |
| QM101 | Q. rubra | Spontaneous | Catherine Chevalier | 41.9759 | -87.84716 | 30.0 | good |
| QM102 | Q. rubra | Spontaneous | Catherine Chevalier | 41.97612 | -87.8474 | 8.4 | fair |
| QM103 | Q. rubra | Spontaneous | Catherine Chevalier | 41.97626 | -87.84754 | 21.4 | fair |
| QM104 | Q. rubra | Spontaneous | Catherine Chevalier | 41.97689 | -87.84794 | 19.5 | good |
| QM105 | Q. rubra | Spontaneous | Catherine Chevalier | 41.97725 | -87.84775 | 4.2 | poor |
| QM106 | Q. rubra | Spontaneous | Catherine Chevalier | 41.97692 | -87.84942 | 39.8 | good |
| QM107 | Q. rubra | Spontaneous | Catherine Chevalier | 41.97677 | -87.84939 | 74.1 | fair |
| QM108 | Q. rubra | Spontaneous | Catherine Chevalier | 41.97637 | -87.8496 | 81.2 | good |
| QM109 | Q. rubra | Spontaneous | Catherine Chevalier | 41.97563 | -87.84878 | 66.3 | good |
| QM11 | Q. rubra | Spontaneous | Labagh woods | 41.98204 | -87.74202 | 50.0 | fair |
| QM110 | Q. rubra | Spontaneous | Catherine Chevalier | 41.97555 | -87.84936 | 62.8 | fair |
| QM111 | Q. rubra | Spontaneous | Catherine Chevalier | 41.97567 | -87.84951 | 27.0 | good |
| QM112 | Q. rubra | Spontaneous | Catherine Chevalier | 41.97586 | -87.84993 | 46.5 | poor |
| QM113 | Q. rubra | Spontaneous | Catherine Chevalier | 41.97532 | -87.85135 | 17.8 | good |
| QM114 | Q. rubra | Spontaneous | Catherine Chevalier | 41.97515 | -87.85194 | 57.5 | fair |
| QM115 | Q. rubra | Spontaneous | Catherine Chevalier | 41.97278 | -87.85252 | 7.8 | poor |
| QM116 | Q. rubra | Spontaneous | Catherine Chevalier | 41.97109 | -87.85149 | 125.7 | fair |
| QM117 | Q. rubra | Spontaneous | Catherine Chevalier | 41.96981 | -87.85159 | 21.0 | good |
| QM118 | Q. rubra | Spontaneous | Catherine Chevalier | 41.96914 | -87.85175 | 80.3 | fair |
| QM119 | Q. rubra | Spontaneous | Catherine Chevalier | 41.96786 | -87.8522 | 10.8 | fair |
| QM12 | Q. rubra | Spontaneous | Labagh woods | 41.98182 | -87.78124 | 32.9 | fair |
| QM120 | Q. rubra | Spontaneous | Catherine Chevalier | 41.96767 | -87.85231 | 4.5 | poor |
| QM121 | Q. rubra | Spontaneous | Catherine Chevalier | 41.96773 | -87.85336 | 10.9 | fair |
| QM122 | Q. rubra | Spontaneous | Catherine Chevalier | 41.96924 | -87.85308 | 87.3 | good |
| QM123 | Q. rubra | Spontaneous | Catherine Chevalier | 41.97583 | -87.85154 | 7.6 | good |
| QM124 | Q. rubra | Spontaneous | Catherine Chevalier | 41.97626 | -87.85181 | 13.9 | good |
| QM125 | Q. rubra | Spontaneous | Catherine Chevalier | 41.97802 | -87.85205 | 8.4 | fair |
| QM13 | Q. rubra | Spontaneous | Labagh woods | 41.98177 | -87.74109 | 3.6 | fair |
| QM14 | Q. rubra | Spontaneous | Labagh woods | 41.98197 | -87.74092 | 20.5 | good |
| QM15 | Q. rubra | Spontaneous | Labagh woods | 41.98205 | -87.7408 | 60.3 | good |
| QM17 | Q. rubra | Spontaneous | Labagh woods | 41.98212 | -87.74085 | 11.7 | good |
| QM18 | Q. rubra | Spontaneous | Labagh woods | 41.98217 | -87.74073 | 5.4 | fair |
| QM19 | Q. rubra | Spontaneous | Labagh woods | 41.98168 | -87.74063 | 87.6 | fair |
| QM51 | Q. rubra | Spontaneous | Dan Ryan Preserve | 41.7378 | -87.68179 | 35.3 | fair |


| QM52 | Q. rubra | Spontaneous | Dan Ryan Preserve | 41.73834 | -87.6818 | 54.0 | poor |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| QM53 | Q. rubra | Spontaneous | Dan Ryan Preserve | 41.73825 | -87.68173 | 65.0 | good |
| QM54 | Q. rubra | Spontaneous | Dan Ryan Preserve | 41.73794 | -87.68173 | 19.6 | poor |
| QM55 | Q. rubra | Spontaneous | Dan Ryan Preserve | 41.73805 | -87.68176 | 19.0 | poor |
| QM56 | Q. rubra | Spontaneous | Dan Ryan Preserve | 41.73802 | -87.68168 | 4.0 | good |
| QM57 | Q. rubra | Spontaneous | Dan Ryan Preserve | 41.73818 | -87.68237 | 14.3 | good |
| QM59 | Q. rubra | Spontaneous | Dan Ryan Preserve | 41.73839 | -87.67612 | 29.2 | good |
| QM60 | Q. rubra | Spontaneous | Dan Ryan Preserve | 41.73913 | -87.67648 | 63.9 | good |
| QM61 | Q. rubra | Spontaneous | Dan Ryan Preserve | 41.7406 | -87.67721 | 28.7 | fair |
| QM62 | Q. rubra | Spontaneous | Dan Ryan Preserve | 41.74061 | -87.67741 | 29.8 | poor |
| QM63 | Q. rubra | Spontaneous | Dan Ryan Preserve | 41.74056 | -87.67724 | 9.4 | fair |
| QN01 | Q. rubra | Remnant | Somme | 42.1391 | -87.82277 | 23.2 | poor |
| QN02 | Q. rubra | Remnant | Somme | 42.13838 | -87.82136 | 31.5 | fair |
| QN03 | Q. rubra | Remnant | Somme | 42.13882 | -87.82135 | 34.0 | fair |
| QN04 | Q. rubra | Remnant | Somme | 42.13945 | -87.81111 | 43.3 | fair |
| QN05 | Q. rubra | Remnant | Somme | 42.13986 | -87.81223 | 40.4 | fair |
| QN06 | Q. rubra | Remnant | Somme | 42.14082 | -87.81312 | 10.7 | fair |
| QN07 | Q. rubra | Remnant | Somme | 42.14056 | -87.8156 | 43.5 | good |
| QN08 | Q. rubra | Remnant | Somme | 42.14069 | -87.81609 | 54.5 | poor |
| QN09 | Q. rubra | Remnant | Somme | 42.13972 | -87.81818 | 21.3 | fair |
| QN10 | Q. rubra | Remnant | Somme | 42.13929 | -87.81911 | 65.4 | poor |
| QN101 | Q. rubra | Remnant | Morton | 41.81482 | -88.0408 | 10.3 | good |
| QN102 | Q. rubra | Remnant | Morton | 41.8139 | -88.04067 | 19.5 | good |
| QN103 | Q. rubra | Remnant | Morton | 41.81311 | -88.04001 | 33.6 | fair |
| QN104 | Q. rubra | Remnant | Morton | 41.81305 | -88.0398 | 48.1 | fair |
| QN105 | Q. rubra | Remnant | Morton | 41.813 | -88.04023 | 43.5 | good |
| QN106 | Q. rubra | Remnant | Morton | 41.81292 | -88.04013 | 28.2 | good |
| QN107 | Q. rubra | Remnant | Morton | 41.81346 | -88.04254 | 27.0 | fair |
| QN108 | Q. rubra | Remnant | Morton | 41.81544 | -88.04699 | 47.2 | good |
| QN11 | Q. rubra | Remnant | Somme | 42.13952 | -87.81981 | 59.2 | poor |
| QN110 | Q. rubra | Remnant | Morton | 41.81768 | -88.04153 | 71.2 | fair |
| QN111 | Q. rubra | Remnant | Morton | 41.81696 | -88.04173 | 2.5 | good |
| QN112 | Q. rubra | Remnant | Morton | 41.81222 | -88.04227 | 2.5 | good |
| QN113 | Q. rubra | Remnant | Morton | 41.81737 | -88.04401 | 3.7 | fair |
| QN12 | Q. rubra | Remnant | Somme | 42.14022 | -87.82058 | 70.0 | fair |
| QN12b | Q. rubra | Remnant | Somme | 42.13897 | -87.82424 | 26.4 | poor |
| QN13 | Q. rubra | Remnant | Somme | 42.13896 | -87.82429 | 27.8 | fair |
| QN14 | Q. rubra | Remnant | Somme | 42.14046 | -87.82561 | 10.4 | fair |
| QN15 | Q. rubra | Remnant | Somme | 42.14246 | -87.82544 | 17.7 | good |
| QN16 | Q. rubra | Remnant | Somme | 42.14264 | -87.82309 | 6.8 | fair |
| QN17 | Q. rubra | Remnant | Somme | 42.14273 | -87.82302 | 8.0 | fair |
| QN18 | Q. rubra | Remnant | Somme | 42.14274 | -87.8229 | 11.3 | good |
| QN19 | Q. rubra | Remnant | Somme | 42.14284 | -87.82293 | 12.4 | good |
| QN20 | Q. rubra | Remnant | Somme | 42.1403 | -87.81776 | 51.0 | good |
| QN20B | Q. rubra | Remnant | Somme | 42.14266 | -87.81704 | 12.5 | fair |
| QN21 | Q. rubra | Remnant | Somme | 42.14262 | -87.81719 | 7.0 | fair |
| QN22 | Q. rubra | Remnant | Somme | 42.14357 | -87.81894 | 5.1 | fair |
| QN23 | Q. rubra | Remnant | Somme | 42.14413 | -87.81823 | 29.2 | fair |
| QN24 | Q. rubra | Remnant | Somme | 42.14438 | -87.81925 | 36.3 | fair |


| QN25 | Q. rubra | Remnant | Somme | 42.14432 | -87.81944 | 25.3 | good |
| :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- |
| QN26 | Q. rubra | Remnant | Somme | 42.14248 | -87.82097 | 25.9 | good |
| QN27 | Q. rubra | Remnant | Somme | 42.14243 | -87.82126 | 5.2 | fair |
| QN28 | Q. rubra | Remnant | Somme | 42.14255 | -87.8221 | 15.7 | fair |
| QN29 | Q. rubra | Remnant | Somme | 42.14258 | -87.82215 | 26.6 | fair |
| QN30 | Q. rubra | Remnant | Somme | 42.14255 | -87.82225 | 6.5 | good |
| QN31 | Q. rubra | Remnant | Somme | 42.14244 | -87.82237 | 20.8 | good |
| QN50 | Q. rubra | Remnant | Busse | 42.0385 | -88.00249 | 37.7 | fair |
| QN51 | Q. rubra | Remnant | Busse | 42.03629 | -88.00863 | 15.0 | good |
| QN52 | Q. rubra | Remnant | Busse | Remnant | Busse | Russe | Remnant |


| QU16 | Q. rubra | Planted | Northeast | 41.94955 | -87.64103 | 13.6 | fair |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| QU17 | Q. rubra | Planted | Northeast | 41.94966 | -87.64056 | 17.1 | fair |
| QU18 | Q. rubra | Planted | Northeast | 41.95 | -87.64074 | 63.3 | good |
| QU19 | Q. rubra | Planted | Northeast | 41.96094 | -87.64502 | 16.9 | fair |
| QU20 | Q. rubra | Planted | Northeast | 41.93576 | -87.67248 | 16.6 | good |
| QU21 | Q. rubra | Planted | Northeast | 41.93592 | -87.67237 | 15.1 | fair |
| QU22 | Q. rubra | Planted | Northwest | 41.92824 | -87.70762 | 21.4 | fair |
| QU23 | Q. rubra | Planted | Northwest | 41.92742 | -87.70707 | 10.0 | good |
| QU24 | Q. rubra | Planted | Northwest | 41.92666 | -87.70711 | 12.8 | fair |
| QU25 | Q. rubra | Planted | Northwest | 41.92519 | -87.70708 | 12.2 | fair |
| QU26 | Q. rubra | Planted | Northwest | 41.92263 | -87.7073 | 21.4 | good |
| QU27 | Q. rubra | Planted | Northwest | 41.92655 | -87.70779 | 20.7 | fair |
| QU28 | Q. rubra | Planted | Northwest | 41.9267 | -87.7077 | 15.9 | fair |
| QU40 | Q. rubra | Planted | Southwest | 41.84722 | -87.6816 | 13.1 | poor |
| QU41 | Q. rubra | Planted | Southwest | 41.84726 | -87.68169 | 11.3 | poor |
| QU42 | Q. rubra | Planted | Southwest | 41.84753 | -87.68099 | 14.0 | poor |
| QU43 | Q. rubra | Planted | Southwest | 41.84758 | -87.68086 | 13.8 | poor |
| QU44 | Q. rubra | Planted | Southwest | 41.83233 | -87.67659 | 32.8 | fair |
| QU45 | Q. rubra | Planted | Southwest | 41.78148 | -87.71867 | 70.4 | good |
| QU46 | Q. rubra | Planted | Southwest | 41.77771 | -87.72155 | 52.0 | fair |
| QU47 | Q. rubra | Planted | Southwest | 41.77321 | -87.65676 | 10.2 | good |
| QU48 | Q. rubra | Planted | Southwest | 41.76503 | -87.58888 | 39.0 | fair |
| QU49 | Q. rubra | Planted | Southeast | 41.82608 | -87.68309 | 5.2 | good |
| QU50 | Q. rubra | Planted | Southeast | 41.82609 | -87.68323 | 5.0 | fair |
| QU51 | Q. rubra | Planted | Southeast | 41.82609 | -87.68309 | 5.5 | fair |
| QU53 | Q. rubra | Planted | Southeast | 41.82612 | -87.68291 | 5.2 | good |
| QU54 | Q. rubra | Planted | Southeast | 41.76696 | -87.60112 | 74.7 | fair |
| QU55 | Q. rubra | Planted | Southeast | 41.77104 | -87.60126 | 57.1 | poor |
| QU56 | Q. rubra | Planted | Southeast | 41.77111 | -87.60132 | 45.7 | good |
| QU57 | Q. rubra | Planted | Southeast | 41.76956 | -87.60134 | 85.2 | fair |
| QU58 | Q. rubra | Planted | Southeast | 41.76981 | -87.60198 | 65.5 | fair |
| QU59 | Q. rubra | Planted | Southeast | 41.77013 | -87.60142 | 81.3 | fair |
| QU60 | Q. rubra | Planted | Southeast | 41.78641 | -87.60856 | 49.2 | fair |
| QU61 | Q. rubra | Planted | Southeast | 41.78642 | -87.60833 | 36.3 | good |
| QU62 | Q. rubra | Planted | Southeast | 41.78882 | -87.60816 | 57.5 | fair |
| QU63 | Q. rubra | Planted | Southeast | 41.78929 | -87.60822 | 60.8 | good |
| QU64 | Q. rubra | Planted | Southeast | 41.78953 | -87.60803 | 56.4 | fair |
| QU65 | Q. rubra | Planted | Southeast | 41.79167 | -87.60861 | 79.0 | fair |
| QU66 | Q. rubra | Planted | Southeast | 41.79189 | -87.60881 | 62.9 | fair |
| QU67 | Q. rubra | Planted | Southeast | 41.79193 | -87.60886 | 53.0 | fair |
| QU68 | Q. rubra | Planted | Southeast | 41.79571 | -87.61503 | 49.1 | good |
| QU69 | Q. rubra | Planted | Southeast | 41.79631 | -87.61509 | 51.8 | good |
| QU70 | Q. rubra | Planted | Southeast | 41.79731 | -87.61535 | 41.0 | fair |
| QU71 | Q. rubra | Planted | Southeast | 41.79773 | -87.61553 | 59.4 | fair |
| QU72 | Q. rubra | Planted | Southeast | 41.79842 | -87.61437 | 70.5 | good |
| QU75 | Q. rubra | Planted | Southwest | 41.73958 | -87.68102 | 73.4 | fair |

