Tallgrass Prairie Restoration Legacies, Summary

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```
packages_needed = c("tidyverse",
                    "png",
                    "knitr",
                    "conflicted")
packages_installed = packages_needed %in% rownames(installed.packages())
if (any(!packages_installed)) {
    install.packages(packages_needed[!packages_installed])
```

```
for (i in 1:length(packages_needed)) {
    library(packages_needed[i], character.only = T)
}
```

```
conflict_prefer("filter", "dplyr")
conflict_prefer("select", "dplyr")
```

Description

The goal is to present results and discuss whether a path forward exists. If so, we will determine the strategy that presents the best story, organization, and interpretation of these results.

Data

For this summary, I'll pull as many objects as possible from existing files to reduce the number of interspersed code chunks. A few quick new analyses will be necessary, though. Data are loaded here.

Methods

Sites and design

The survey followed an unbalanced complete block design. Corn, restored, and remnant fields are compared, with at least one of each field type in each block. I have called blocks "regions" so far. We collected samples and data from four regions, shown on the map below.

The design is unbalanced because there are more restored fields than corn or remnant. In all but one case, only single corn and remnant fields were available in each region. This means that we only have replication to separate field types when using the entire block design.

Table 1: Count of fields by type and region: BM = Blue Mounds, FG = Faville Grove, FL = Fermilab, LP = Lake Petite

	corn	restored	remnant
$\overline{\mathrm{BM}}$	1	7	1
FG	1	1	1
FL	2	6	1
$_{ m LP}$	1	2	1

corn restored remnant

The rules for establishing a chronosequence are strict. We cannot call fields from all regions a chronosequence. With seven restored fields in a small geographic area, the Blue Mounds fields are our best bet for this, but we will likely have to call this a "pseudochronosequence" and avoid some inferences. Mantel tests (not shown) failed to find correlations between soil variables and pairwise distance, giving us a little confidence that we're avoiding systemic bias.

Fungal communities

We collected soil cores from 10 haphazardly-selected locations in each field. Genomic DNA was extracted, and the lab and bioinformatics pipeline delivered community data from ITS or 18S regions clustered as OTUs or SVs. In preliminary work, inferences made with SVs were weaker but not qualitatively different. I continued with OTUs only.

A few samples had failed to amplify, leaving some fields represented with 9 samples instead of 10. This unbalance, which is normally not a problem, became unacceptable for a couple of reasons. First, I had planned to use permutation tests in ordinations, and application function how() from package permute requires balance at the plot (i.e., field) level. Second, I wanted to summarize the subsamples at the field level because fields are our replicates. Any comparisons with field metadata, where we have one data point per field, would be pseudoreplicated when regressed against subsample level data.

Finally, after rarefaction it was clear that several samples contained very few sequences compared with the others. I used an iterative process to remove subsamples with few sequences, choosing a rarefication cutoff near the plateau of OTU recovery at depth (see figures). This ended up being 8 subsamples for ITS and 7 for 18S datasets. These subsample sequence values were then rarefied and used as-is, or, when reporting at the field level, subsample sequence values were summed and rarefied.

This process didn't remove many OTUs and doesn't change any major interpretation.

Even after this process, it's clear that fungal communities were undersampled.

Plant data

Plant community data resulted from two independent surveys. In the Wisconsin regions, haphazard transects were established and 10 meter frames placed, with percent cover estimated for all species, resulting in a dataset with plant composition. In Fermi, relevé methods were used, resulting in presence/absence data only. Plant metadata were assembled from multiple sources and include plant traits and natural history details.

Soil data

In the field, soil was pooled from 10 haphazardly-selected locations, mixed, and sampled once for soil chemical analysis. Soil data includes abiotic macro and micronutrients, organic carbon, and properties like pH. Average precipitation was determined for each field using PRISM climate data and is included with the soil data.

Design/site data

- Field type: corn, restored, remnant
- Field age: years since restoration (NA for corn and remnant)
- Region: blocks



Figure 1: Labels show centroids of regions used for this work. BM = Blue Mounds, FG = Faville Grove, FL = Fermilab, LP = Lake Petite.

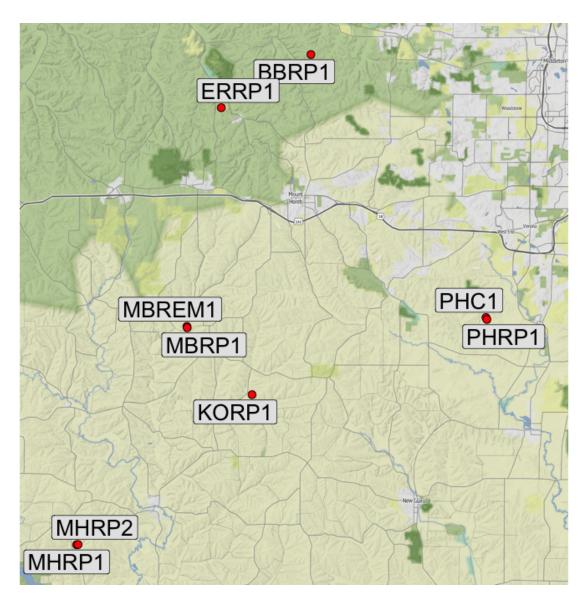
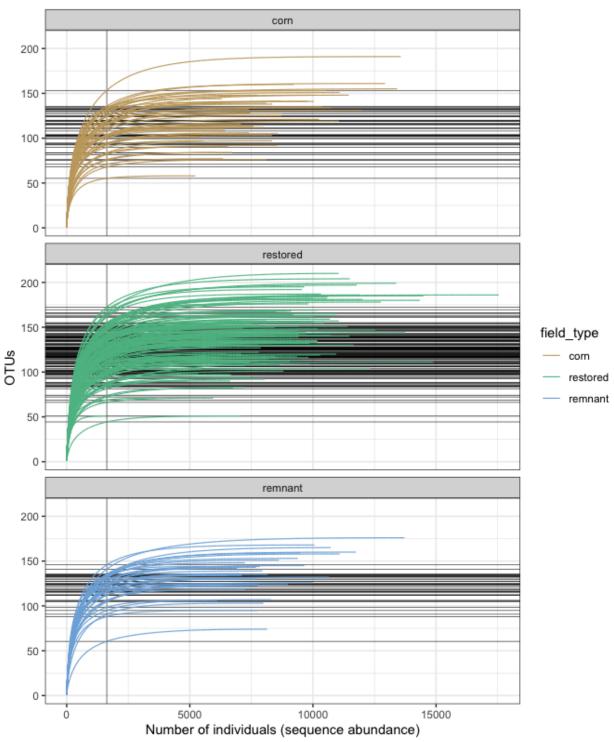


Figure 2: Labels show individual fields in the Blue Mounds region.

Rarefaction of ITS data



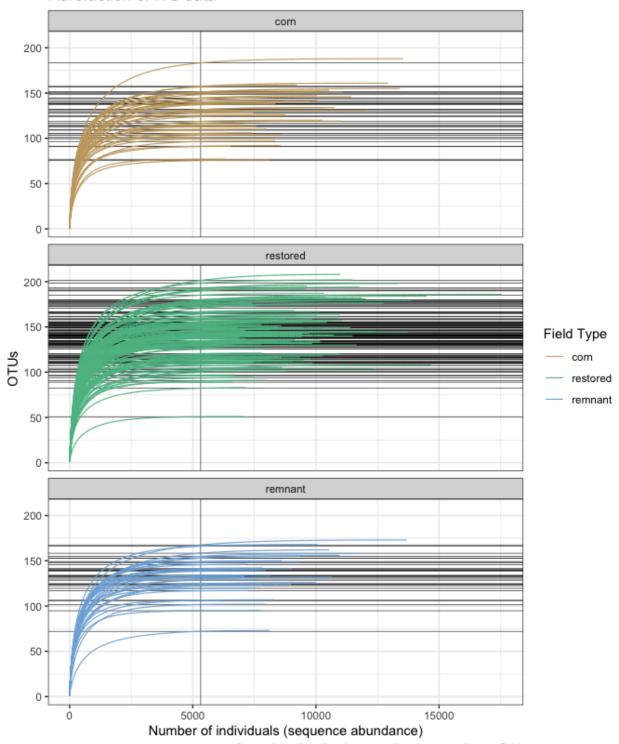
Curves based on the nine most abundant samples per field.

Vertical line shows the minimum sequence abundance for any field.

Horizontal lines show expected richness when rarefied to that abundance.

Figure 3: Pre

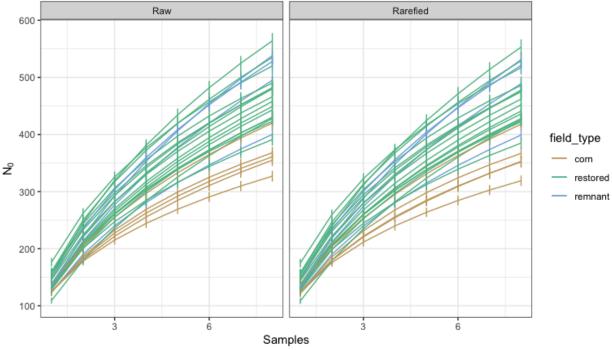
Rarefaction of ITS data



Curves based on the nine most abundant samples per field. Vertical line shows the minimum sequence abundance for any field. Horizontal lines show expected richness when rarefied to that abundance.

Figure 4: Post

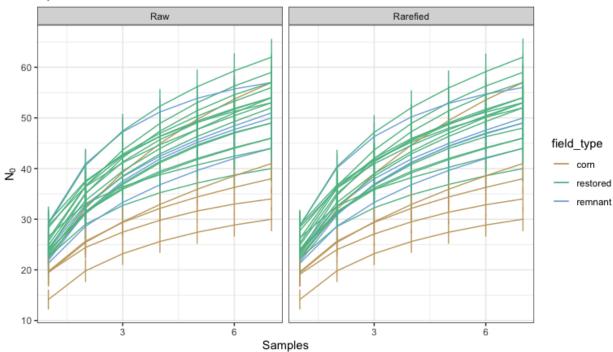
Species accumulation of ITS data



Species accumulation by the "exact" method; standard deviation (vertical lines) are conditioned by the empirical dataset.

Figure 5: ITS

Species accumulation of 18S data



Species accumulation by the "exact" method; standard deviation (vertical lines) conditioned by the empirical dataset.

Figure 6: 18S

Response data

Also taken from the pooled soil in each field, one sample was taken for analysis of Water Stable Aggregates (I don't know who did these), and one for quantification of microbial biomass. ## Data sources, summary

- Fungal genomic data, ITS and 18S, OTU clusters
- Plant community data, composition in Wisconsin and presence/absence in Fermi
- Plant traits and natural history
- Soil properties
- Fungal biomass
- Water stable aggregates
- Site metadata and design

Results

Fungal communities

Soil fungi, field types, and regions

Cornfields stand apart while remnants are mixed with restored fields. As the age of restored fields increases, their distance from corn and remnants grows. The high dimensionality here keeps the explanatory power of axes 1 & 2 low. Clusters based on field type were significant based on a permanova (with regions as blocks and subsamples held constant within fields, the replicate). However, a pairwise post-hoc permutation failed to separate remnant fields from any other type. This was also true when field centroids were used exclusively and the permutation test adjusted to this scheme.

```
## Permutation test for adonis under reduced model
## Terms added sequentially (first to last)
## Blocks: region
## Plots: field_key, plot permutation: free
## Permutation: none
## Number of permutations: 1999
##
##
  adonis2(formula = d ~ field_type, data = env_w, permutations = gl_perm_design)
##
               Df SumOfSqs
                                R2
                                         F Pr(>F)
## field_type
                2
                     5.790 0.08144 8.7335 5e-04 ***
## Residual
              197
                    65.303 0.91856
## Total
              199
                    71.094 1.00000
## ---
## Signif. codes: 0 '***' 0.001 '**' 0.05 '.' 0.1 ' ' 1
##
##
## group1
              group2
                            R2
                                  F_value
                                            df1
                                                  df2
                                                          p_value
                                                                     p_value_adj
## restored
              corn
                         0.066
                                   11.791
                                              1
                                                  166
                                                        0.0010000
                                                                           0.003
## restored
                         0.018
                                    2.835
                                              1
                                                  158
                                                        0.2110000
                                                                           0.211
              remnant
                                   10.946
                                                        0.1428571
                                                                           0.211
## corn
              remnant
                         0.135
                                              1
                                                   70
```

PCoA Ordination (ITS gene, 97% OTU) 250 -200 150 Seq. abund. 100 50 Field Type com soil sapr wood sapr plnt path 0.2 restored remnant Axis 2 (5%) Region ВМ FG -0.2 -0.2 0.0 0.2 -0.4 Axis 1 (8.2%) Blue Mounds Faville Grove 0.3 0.2 0.2 0.1 0.0 0.0 -0.1 -0.2 -0.2 -0.3 Axis 2 -0.4 -0.2 0.0 0.2 -0.4 -0.2 0.0 0.2 Fermilab Lake Petite 0.2 0.2 -0.0 0.0

ITS gene. Text indicates years since restoration.

0.2

0.4

0.0

Figure 7: ITS-based fungal communities, most abundant guilds in set. $10\,$

0.25

-0.2

-0.25

0.00

-0.2

Axis 1

-0.2

Soil fungal guilds, field types, and regions

By sequence abundance across the entire dataset, the top guilds are:

- 1. Unidentified (not shown on column charts)
- 2. plant pathogens
- 3. soil saprotrophs
- 4. wood saprotrophs
- 5. dung saprotrophs
- 6. litter saprotrophs

Compared with the sequence abundance in the NA group, plant pathogens and soil saprotrophs are abundant enough to feel somewhat confident about in terms of coverage. These are also the ones that varied most with age in restored fields as we'll see later, and they were most conspicuous in an indicator species analysis. Trends across field types within regions aren't very consistent...

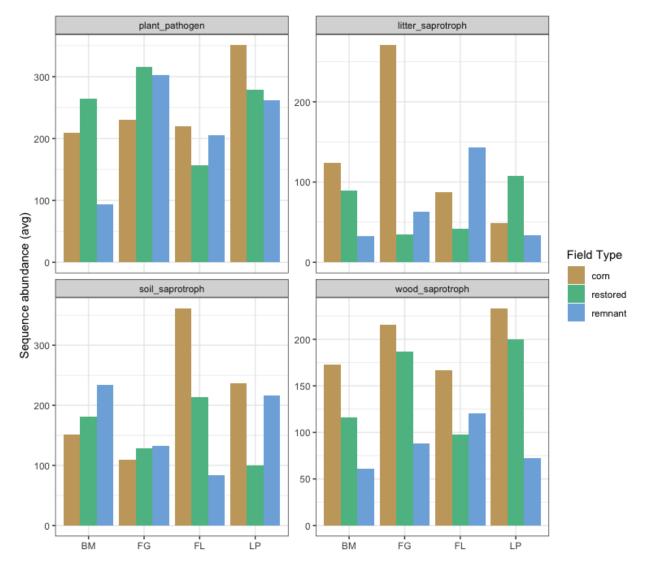


Figure 8: Abundant or dynamic guilds across field types in regions, ITS dataset.

heading

more text

heading again

PCoA Ordination (18S gene, 97% OTU) 0.2 Field Type corn restored 0.0 Axis 2 (5%) remnant Region ВМ FG Seq. abund. (avg) -0.4400 200 Cirdgi DvrsspGgsprcPrglmr -0.6 -0.25 -0.50 0.00 0.25 Axis 1 (7.1%) Blue Mounds Faville Grove 0.4 0.2 0.2 0.0 0.0 -0.2 -0.2 -0.4 Axis 2 -0.2 0.0 0.2 0.4 -0.4 -0.2 0.0 0.2 0.4 -0.4 Fermilab Lake Petite 0.4 0.2 0.25 0.0 0.00 -0.2 -0.25 -0.25 0.00 0.25 -0.25 0.00 0.25 0.50 Axis 1

Figure 9: 18S-based fungal communities, most important families in set. $13\,$

18S gene (AMF). Text indicates years since restoration.