

GENOTYPIC VARIATION INFLUENCES LICHEN CO-OCCURRENCE NETWORK STRUCTURE

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1. THINGS TO DO

- (1) Do a whole network analysis using PerMANOVA
- (2) Can we do pathway de-proliferation?
- (3) Use eigenvector centrality. What is the dominant eigenvalue for each matrix?
- (4) Use Fath and Patten's 1998 network mutualism in utility analysis
- (5) Fix the gplot inputs for genotype vertex attributes (it is currently using the tree level data)
- (6)
- (7)

2. GENERATE LICHEN NETWORK MODELS FOR ALL GENOTYPES IN EACH GARDEN

2.1. Load packages.

```
> require(RLRsim)
> require(lme4)
> require(sna)
  Tools for Social Network Analysis
Version      2.2-0 created on      2010-11-21.
copyright (c) 2005, Carter T. Butts, University of California-Irvine
Type help(package="sna") to get started.
> require(vegan)
> require(xtable)
> require(gplots)
gdata: read.xls support for 'XLS' (Excel 97-2004) files ENABLED.

gdata: read.xls support for 'XLSX' (Excel 2007+) files ENABLED.
> source("source/CorNets.R")
> source("source/pairs.R")
> se = function(x) {
+   sd(x)/sqrt(length(x))
+ }
> binary = function(x, bin = 0) {
+   x[x > bin] = 1
```

```

+   return(x)
+ }
> richness = function(x) {
+   x = apply(x, 2, sum)
+   x[x != 0] = 1
+   sum(x)
+ }
> bplot = function(x, y, ylab = "") {
+   mu. = tapply(y, x, mean)
+   se. = tapply(y, x, se)
+   barplot2(mu., plot.ci = TRUE, ci.u = mu. + se., ci.l = mu. -
+     se., las = 2, ylab = ylab)
+ }
> which.garden = function(x) {
+   x = unlist(strsplit(as.character(x), split = ".", fixed = TRUE))[1]
+   x = unlist(strsplit(x, split = ""))
+   if (length(x) < 3) {
+     x = "ONC"
+   }
+   else {
+     x = "PIT"
+   }
+   return(x)
+ }
> lco2quad = function(x) {
+   com = x[, -1:-6]
+   com = apply(com, 1, sum)
+   x = x[, 5:6]
+   out = array(0, dim = c(max(x), max(x)))
+   for (i in (1:nrow(x))) {
+     out[x[i, 1], x[i, 2]] = com[i]
+   }
+   return(out)
+ }
> exp.m <- function(mat, n) {
+   if (n == 1)
+     return(mat)
+   result <- diag(1, ncol(mat))
+   while (n > 0) {
+     if (n%%2 != 0) {
+       result <- result %*% mat
+       n <- n - 1
+     }

```

```
+      mat <- mat %*% mat
+      n <- n/2
+    }
+    return(result)
+  }
> proliferate = function(g = "graph", lim = 10000) {
+   n = numeric()
+   for (i in 1:lim) {
+     n[i] = sum(exp.m(g, i))
+     if (n[i] == Inf) {
+       break
+     }
+   }
+   n = n[-i]
+   return(n)
+ }
```

2.2. **Data Summary.** NOTE: there was a data entry error for tree N4.30, changed 11 to 1 in LCO_data_ONC_PIT.csv.

```
> lco = read.csv("data/LCO_data_ONC_PIT.csv")
> xtable(summary(lco[, 1:6]), table.placement = "tbp")
```

	Tree	Geno	Year	Quadrat	Row	Col
1	N1.10 : 100	996 :1000	Min. :2010	n45.55:4850	Min. : 1.0	Min. : 1.0
2	N1.11 : 100	1008 : 900	1st Qu.:2011	n80.90:3050	1st Qu.: 3.0	1st Qu.: 3.0
3	N1.24 : 100	1017 : 750	Median :2011		Median : 5.5	Median : 5.5
4	N1.27 : 100	11 : 700	Mean :2011		Mean : 5.5	Mean : 5.5
5	N1.28 : 100	10 : 600	3rd Qu.:2011		3rd Qu.: 8.0	3rd Qu.: 8.0
6	N1.31 : 100	1023 : 600	Max. :2011		Max. :10.0	Max. :10.0
7	(Other):7300	(Other):3350				

```
> xtable(summary(lco[, 7:10]), table.placement = "tbp")
```

	Xgal	Csub	Lsp	Chol
1	Min. :0.0000	Min. :0.00000	Min. :0.00000	Min. :0.000000
2	1st Qu.:0.0000	1st Qu.:0.00000	1st Qu.:0.00000	1st Qu.:0.000000
3	Median :1.0000	Median :0.00000	Median :0.00000	Median :0.000000
4	Mean :0.5084	Mean :0.09165	Mean :0.02076	Mean :0.009494
5	3rd Qu.:1.0000	3rd Qu.:0.00000	3rd Qu.:0.00000	3rd Qu.:0.000000
6	Max. :1.0000	Max. :1.00000	Max. :1.00000	Max. :1.000000

```
> xtable(summary(lco[, 13:ncol(lco)]), table.placement = "tbp")
```

	Pads	Pund	Rsp
1	Min. :0.000000	Min. :0.000000	Min. :0.000000
2	1st Qu.:0.000000	1st Qu.:0.000000	1st Qu.:0.000000
3	Median :0.000000	Median :0.000000	Median :0.000000
4	Mean :0.002532	Mean :0.002025	Mean :0.002532
5	3rd Qu.:0.000000	3rd Qu.:0.000000	3rd Qu.:0.000000
6	Max. :1.000000	Max. :1.000000	Max. :1.000000

2.3. Separate Community Matrixes by Tree.

```
> lco = lco[lco$Quadrat == "n45.55", ]
> lco.l = list()
> tree.names = as.character(unique(lco$Tree))
> for (i in (1:length(tree.names))) {
+   lco.l[[i]] = lco[lco$Tree == tree.names[i], ]
+ }
> names(lco.l) = tree.names
> any(table(tree.names) != 1)

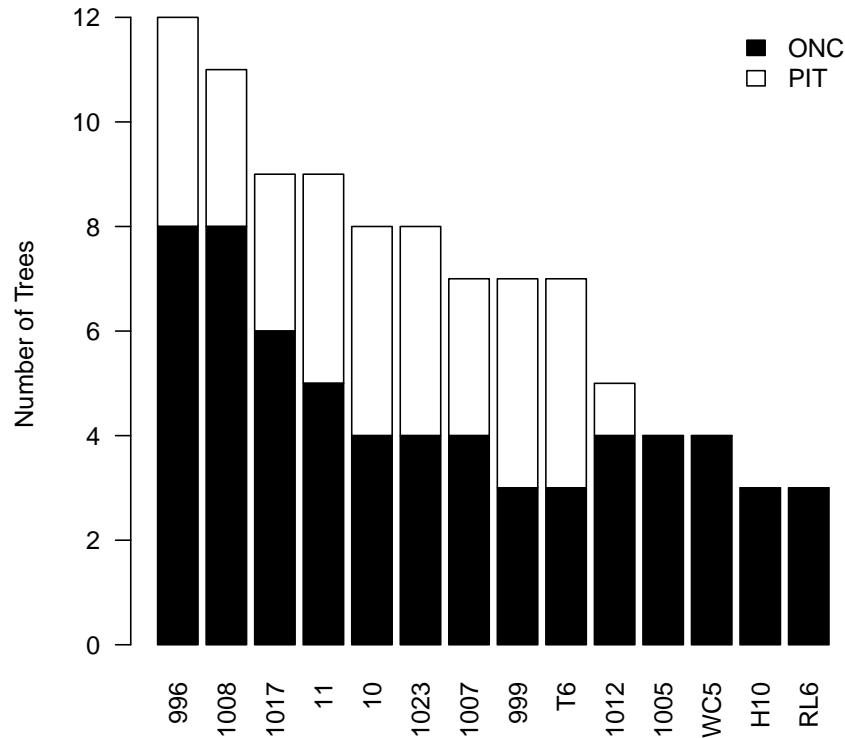
[1] FALSE
```

2.4. Separate garden, genotype and community data.

```
> garden = sapply(tree.names, which.garden)
> geno = unlist(lapply(lco.l, function(x) as.character(x$Geno[1])))
> com.l = lapply(lco.l, function(x) x[, (1:ncol(x))[colnames(x) ==
+   "Xgal"]:ncol(x)])
> geno.table = tapply(factor(geno), garden, table)
> geno.table = rbind(geno.table$ONC, geno.table$PIT)
> rownames(geno.table) = c("ONC", "PIT")
> geno.table = geno.table[, order(apply(geno.table, 2, sum), decreasing = TRUE)]
> xtable(geno.table)
```

	996	1008	1017	11	10	1023	1007	999	T6	1012	1005	WC5	H10	RL6
ONC	8	8	6	5	4	4	4	3	3	4	4	4	3	3
PIT	4	3	3	4	4	4	3	4	4	1	0	0	0	0

```
> barplot(geno.table, beside = FALSE, las = 2, ylab = "Number of Trees",
+   col = c(1, 0))
> legend("topright", c("ONC", "PIT"), fill = c(1, 0), bty = "n")
```



2.5. Convert to Quadrat View.

```
> quad.l = lapply(lco.l, lco2quad)
```

Using the `kendall.pairs` function at an α level of 0.05 and adjusting for multiple tests, we can model each community network using the pairwise correlations of all species pairs.

```
> zero.na = function(x) {
+   x[is.na(x)] = 0
+   return(x)
+ }
> cor.l = lapply(com.l, kendall.pairs, alpha = 0.05, p.adj = TRUE)
> cor.l = lapply(cor.l, zero.na)
```

3. NETWORK STATISTICS

```
> A = lapply(com.l, function(x) apply(x, 2, sum))
> A. = lapply(A, binary)
> R = unlist(lapply(com.l, richness))
```

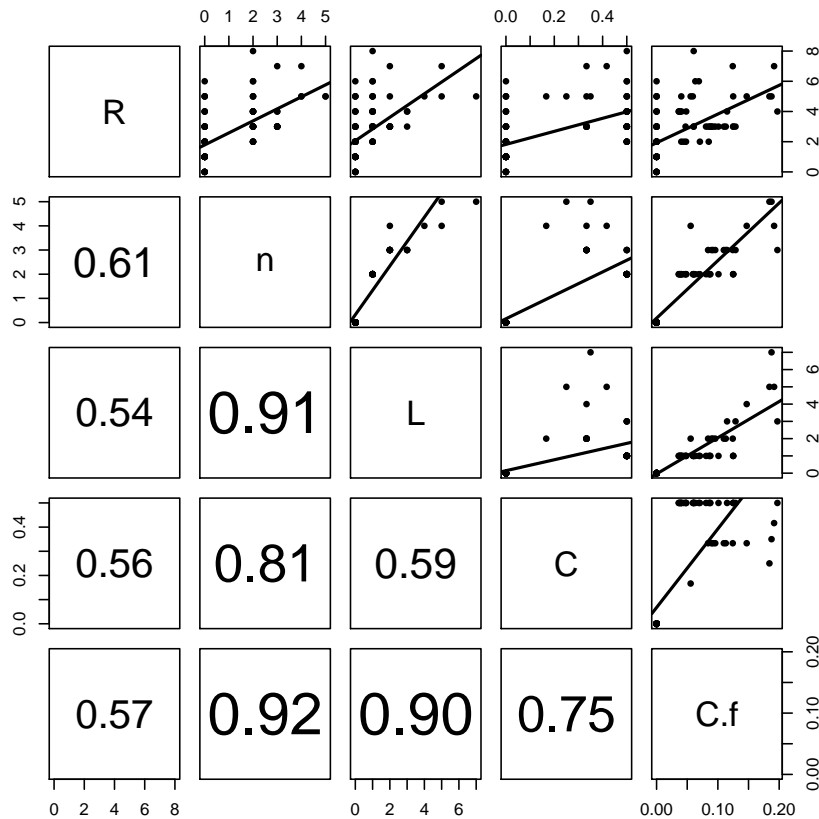
```

> n = unlist(lapply(cor.l, function(x) length(apply(x, 1, sum)[abs(apply(x,
+ 1, sum)) > 0])))
> L = unlist(lapply(cor.l, function(x) length(x[x != 0])/2))
> C = L/(n * (n - 1))
> C[is.na(C)] = 0
> abs.cor.l = lapply(cor.l, abs)
> C.f = unlist(lapply(abs.cor.l, function(x) centralization(x,
+ degree, mode = "graph"))))

> input.pairs = cbind(R, n, L, C, C.f)
> pairs.onc = input.pairs[garden == "ONC", ]
> pairs.pit = input.pairs[garden == "PIT", ]

> pairs(input.pairs, upper.panel = panel.lm, lower.panel = panel.cor)

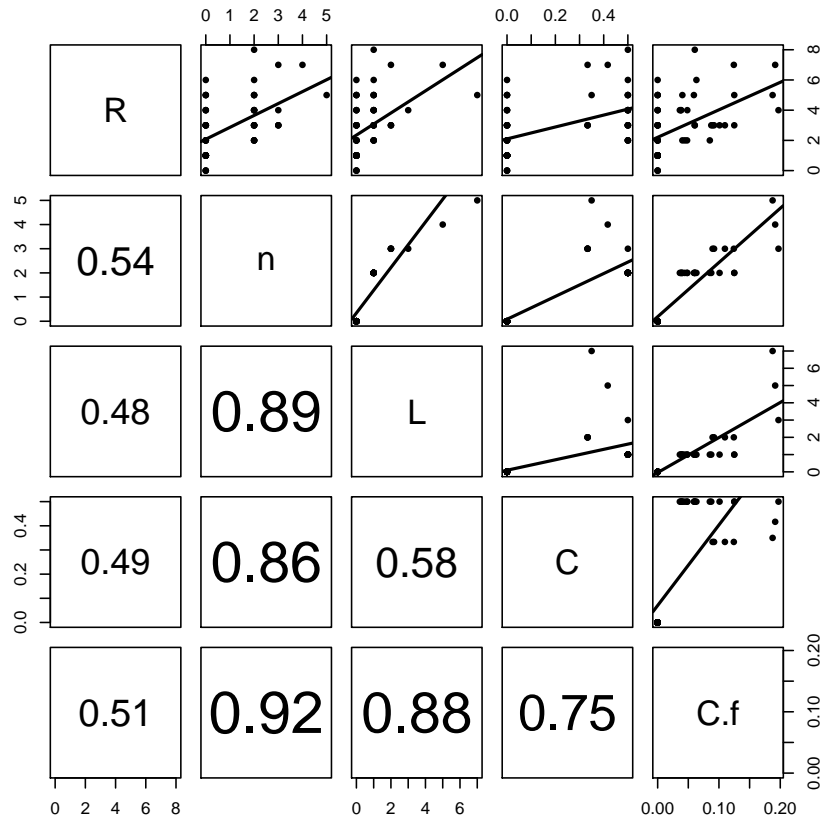
```



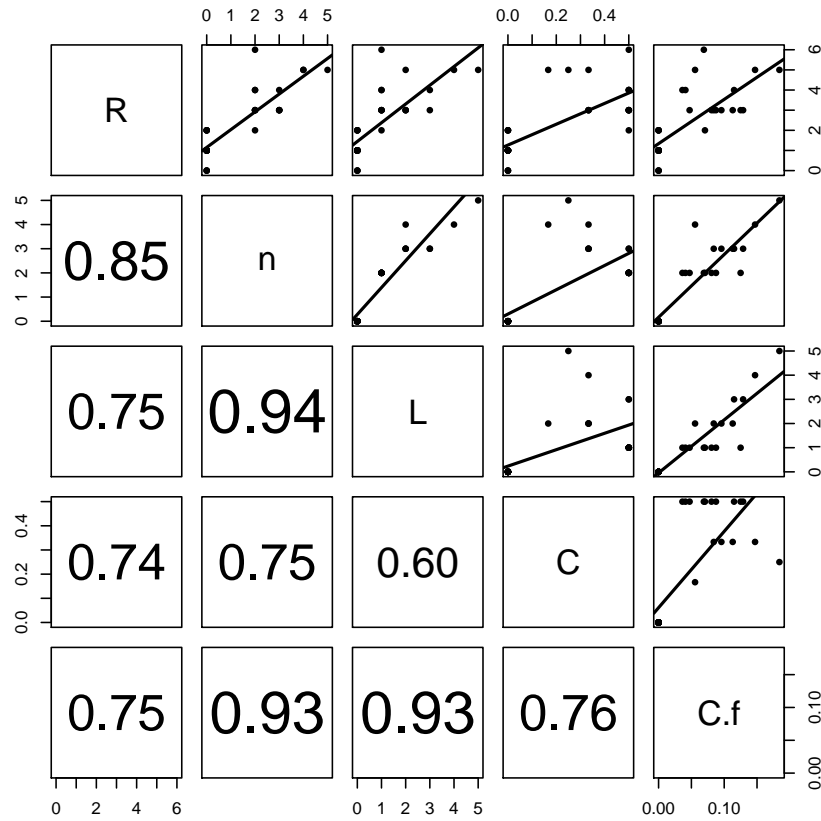
```

> pairs(pairs.onc, upper.panel = panel.lm, lower.panel = panel.cor)

```



```
> pairs(pairs.pit, upper.panel = panel.lm, lower.panel = panel.cor)
```



3.1. REML.

Separate the analysis of ONC and PIT due to unequal representation of genotypes at both gardens.

```
> geno.onc = geno[garden == "ONC"]
> n.onc = n[garden == "ONC"]
> log.n.onc = log(n.onc + 0.5)
> L.onc = L[garden == "ONC"]
> log.L.onc = log(L.onc + 0.5)
> C.onc = C[garden == "ONC"]
> Cf.onc = C.f[garden == "ONC"]
> asin.sqrt.Cf.onc = asin(sqrt(Cf.onc))
> exactRLRT(lmer(n.onc ~ 1 | geno.onc))
```

simulated finite sample distribution of RLRT. (p-value based on 10000 simulated values)

data: RLRT = 3.151, p-value = 0.0325

```
> exactRLRT(lmer(log.n.onc ~ 1 | geno.onc))
```



```

simulated finite sample distribution of RLRT. (p-value based on 10000 simulated values)
  data: RLRT = 4.9866, p-value = 0.0099
> exactRLRT(lmer(L.unc ~ 1 | geno.unc))
simulated finite sample distribution of RLRT. (p-value based on 10000 simulated values)
  data: RLRT = 0, p-value = 0.4589
> exactRLRT(lmer(log.L.unc ~ 1 | geno.unc))
simulated finite sample distribution of RLRT. (p-value based on 10000 simulated values)
  data: RLRT = 3.3797, p-value = 0.0287
> exactRLRT(lmer(C.unc ~ 1 | geno.unc))
simulated finite sample distribution of RLRT. (p-value based on 10000 simulated values)
  data: RLRT = 7.0159, p-value = 0.0038
> exactRLRT(lmer(Cf.unc ~ 1 | geno.unc))
simulated finite sample distribution of RLRT. (p-value based on 10000 simulated values)
  data: RLRT = 0.8348, p-value = 0.16
> exactRLRT(lmer(asin.sqrt.CF.unc ~ 1 | geno.unc))
simulated finite sample distribution of RLRT. (p-value based on 10000 simulated values)
  data: RLRT = 2.8031, p-value = 0.0423
> geno.pit = geno[garden == "PIT"]
> n.pit = n[garden == "PIT"]
> log.n.pit = log(n.pit + 0.5)
> L.pit = L[garden == "PIT"]
> log.L.pit = log(L.pit + 0.5)
> C.pit = C[garden == "PIT"]
> Cf.pit = Cf[garden == "PIT"]
> asin.sqrt.CF.pit = asin(sqrt(Cf.pit))
> exactRLRT(lmer(n.pit ~ 1 | geno.pit))
simulated finite sample distribution of RLRT. (p-value based on 10000 simulated values)
  data: RLRT = 0, p-value = 1
> exactRLRT(lmer(log.n.pit ~ 1 | geno.pit))
simulated finite sample distribution of RLRT. (p-value based on 10000 simulated values)
  data: RLRT = 0, p-value = 1
> exactRLRT(lmer(L.pit ~ 1 | geno.pit))
simulated finite sample distribution of RLRT. (p-value based on 10000 simulated values)
  data: RLRT = 0, p-value = 1
> exactRLRT(lmer(log.L.pit ~ 1 | geno.pit))
simulated finite sample distribution of RLRT. (p-value based on 10000 simulated values)
  data: RLRT = 0, p-value = 1
> exactRLRT(lmer(C.pit ~ 1 | geno.pit))

```

simulated finite sample distribution of RLRT. (p-value based on 10000 simulated values)

data: RLRT = 0, p-value = 1

```
> exactRLRT(lmer(Cf.pit ~ 1 | geno.pit))
```

simulated finite sample distribution of RLRT. (p-value based on 10000 simulated values)

data: RLRT = 0, p-value = 1

```
> exactRLRT(lmer(asin.sqrt.CF.pit ~ 1 | geno.pit))
```

simulated finite sample distribution of RLRT. (p-value based on 10000 simulated values)

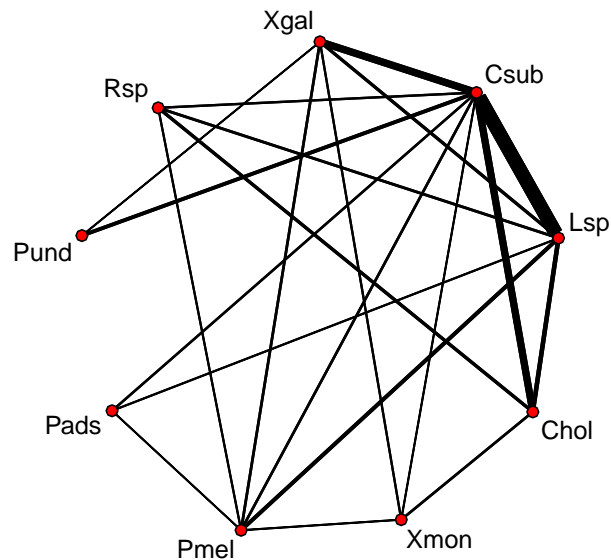
data: RLRT = 0, p-value = 1

4. PLOTS

4.1. Network Graphs.

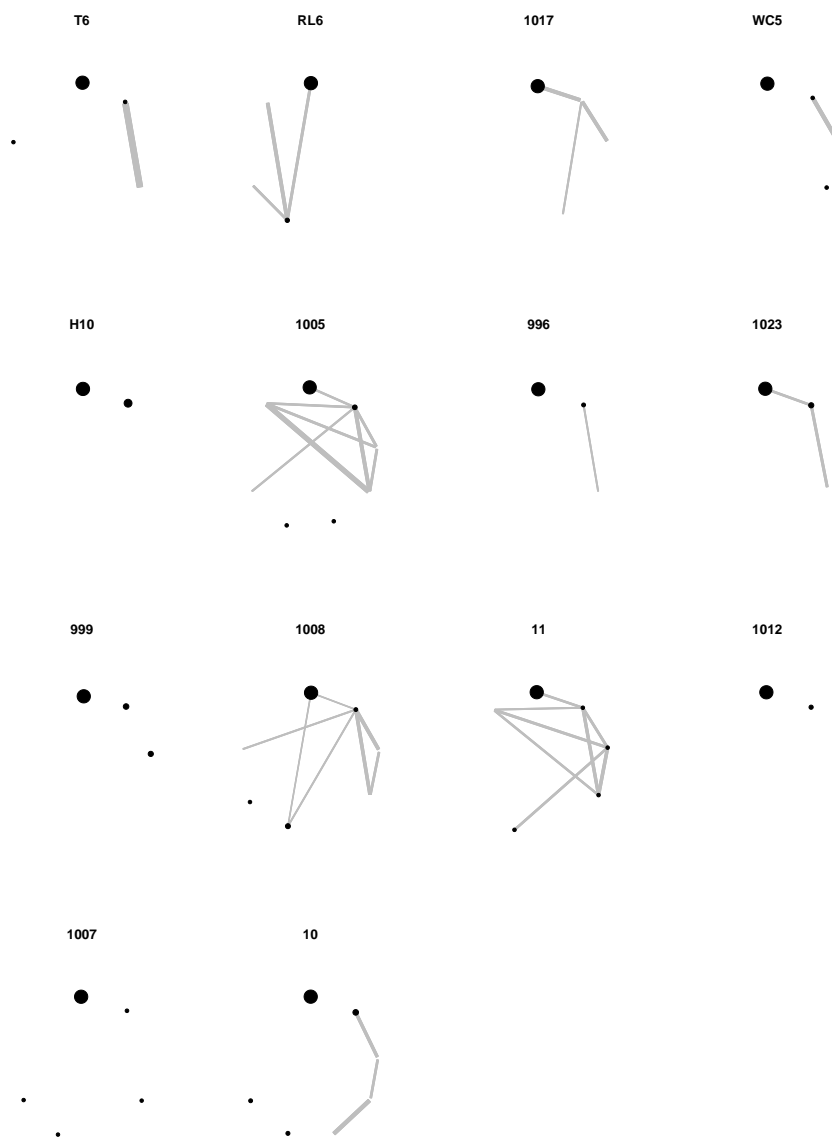
Graph of the mean connections across all trees.

```
> gplot(G.graph, gmode = "graph", displaylabels = TRUE, mode = "circle",
+       edge.lwd = G.graph * 200)
```



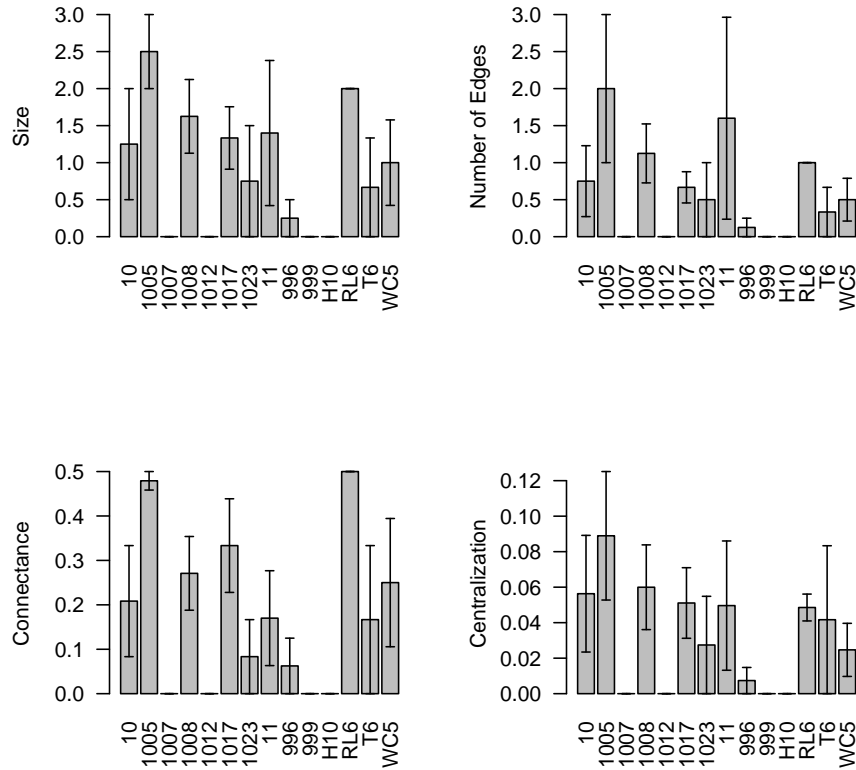
Mean graphs for each genotype.

```
> cor.l. = cor.l[garden == "ONC"]
> geno.A = A[garden == "ONC"]
> geno.A. = A.[garden == "ONC"]
> geno.net = list()
> for (i in (1:length(unique(geno.onc)))) {
+   x = cor.l.[geno.onc == unique(geno.onc)[i]]
+   geno.net[[i]] = x[[1]]
+   for (j in (2:length(x))) {
+     geno.net[[i]] = geno.net[[i]] + x[[j]]
+   }
+   geno.net[[i]] = geno.net[[i]]/length(x)
+ }
> names(geno.net) = unique(geno)
```



ONC Network Statistics

```
> par(mfrow = c(2, 2))
> bplot(geno.onc, n.onc, "Size")
> bplot(geno.onc, L.onc, "Number of Edges")
> bplot(geno.onc, C.onc, "Connectance")
> bplot(geno.onc, Cf.onc, "Centralization")
```



PIT Network Statistics

```
> par(mfrow = c(2, 2))
> bplot(geno.pit, n.pit, "Size")
> bplot(geno.pit, L.pit, "Number of Edges")
> bplot(geno.pit, C.pit, "Connectance")
> bplot(geno.pit, Cf.pit, "Centralization")
```

