Escherichia Coli Network

Example for GeneNet 1.2.7 (June 2013) or later

This note reproduces the "Escherichia coli" network example from J. Schäfer and K. Strimmer. 2005. A shrinkage approach to large-scale covariance estimation and implications for functional genomics. Statist. Appl. Genet. Mol. Biol. 4: 32 (http://dx.doi.org/10.2202/1544-6115.1175)

Load GeneNet package

```
library("GeneNet")

## Loading required package: corpcor

## Loading required package: longitudinal

## Loading required package: fdrtool

## Loading required package: igraph

E. Coli data set (9 time points for 102 genes):

data(ecoli)
dim(ecoli)

## [1] 9 102
```

Estimation of partial correlations

Estimate matrix of partial correlation using a shrinkage estimator:

```
pc = ggm.estimate.pcor(ecoli)

## Estimating optimal shrinkage intensity lambda (correlation matrix): 0.1804

dim(pc)

## [1] 102 102

Assign p-values, q-values and empirical posterior probabilities to all 5151 potential edges in the network:
ecoli.edges = network.test.edges(pc, direct=TRUE, fdr=TRUE)

## Estimate (local) false discovery rates (partial correlations):
## Step 1... determine cutoff point
## Step 2... estimate parameters of null distribution and eta0
## Step 3... compute p-values and estimate empirical PDF/CDF
## Step 4... compute q-values and local fdr
## Step 5... prepare for plotting
```

```
##
## Estimate (local) false discovery rates (log ratio of spvars):
## Step 1... determine cutoff point
## Step 2... estimate parameters of null distribution and eta0
## Step 3... compute p-values and estimate empirical PDF/CDF
## Step 4... compute q-values and local fdr
## Step 5... prepare for plotting
dim(ecoli.edges)
## [1] 5151
              10
```

[1] 125 11

The table lists all edges in the order strength of partial correlations:

```
ecoli.edges[1:5,]
```

```
##
          pcor node1 node2
                                    pval
                                                 qval
                                                           prob
                                                                   log.spvar
## 1 0.2318566
                  51
                        53 2.220446e-16 3.612205e-13 1.0000000 -0.043537019
                        53 2.220446e-16 3.612205e-13 1.0000000 -0.040249854
## 2
     0.2240555
                   52
## 3 0.2150782
                  51
                        52 2.220446e-16 3.612205e-13 1.0000000 -0.003287165
                  7
## 4 0.1732886
                        93 3.108624e-15 3.792816e-12 0.9999945 -0.025293430
## 5 -0.1341889
                   29
                         86 1.120813e-09 1.093998e-06 0.9999516 0.022305368
      pval.dir qval.dir prob.dir
## 1 0.3803869 0.7557272
## 2 0.4173922 0.7724561
                                0
## 3 0.9471949 0.8851073
                                0
## 4 0.6103234 0.8323249
                                0
## 5 0.6531371 0.8415749
                                0
```

Decide which edges to include in the network

To obtain a graph you need to select top ranking edges according to a suitable criterion. Here are some suggestions:

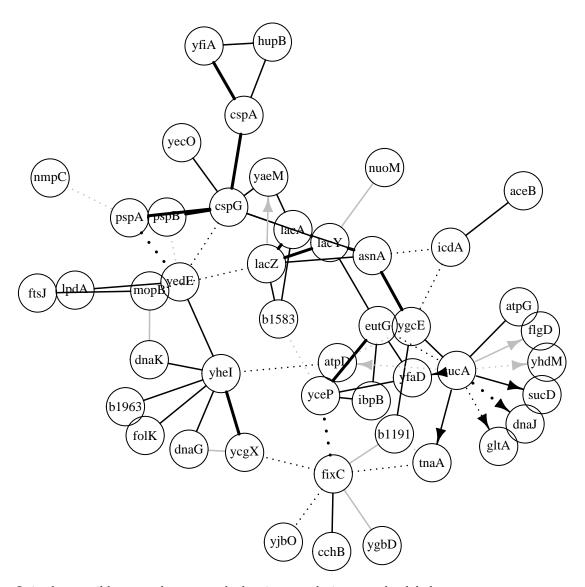
1. Use local fdr cutoff 0.2, i.e. include all edges with posterior probability of at least 0.8.

```
ecoli.net = extract.network(ecoli.edges)
##
## Significant edges: 125
##
      Corresponding to 2.43 % of possible edges
##
## Significant directions: 377
      Corresponding to 7.32 % of possible directions
## Significant directions in the network: 17
      Corresponding to 13.6 % of possible directions in the network
dim(ecoli.net)
```

2. Use local fdr cutoff 0.1, i.e. i.e. include all edges with posterior probability of at least 0.9.

```
ecoli.net = extract.network(ecoli.edges, cutoff.ggm=0.9, cutoff.dir=0.9)
##
## Significant edges: 65
       Corresponding to 1.26\ \% of possible edges
##
##
## Significant directions: 269
       Corresponding to 5.22 % of possible directions
##
## Significant directions in the network: 6
       Corresponding to 9.23 % of possible directions in the network
dim(ecoli.net)
## [1] 65 11
  3. Include a fixed number of edges, say the 70 strongest edges
ecoli.net = extract.network(ecoli.edges, method.ggm="number", cutoff.ggm=70)
##
## Significant edges: 70
##
       Corresponding to 1.36 % of possible edges
##
## Significant directions: 377
       Corresponding to 7.32 % of possible directions
## Significant directions in the network: 9
       Corresponding to 12.86 % of possible directions in the network
dim(ecoli.net)
## [1] 70 11
Plot network For plotting we use the igraph package (http://igraph.org). Note igraph is automatically
installed with GeneNet.
library("igraph") #
Create igraph object from the list of edges:
node.labels = colnames(ecoli)
igr1 = network.make.igraph(ecoli.net, node.labels)
igr1
## IGRAPH DN-- 46 70 --
## + attr: name (v/c), color (v/l), label.color (v/n), arrow.mode
     (e/n), width (e/n), color (e/c), lty (e/n)
Plot the network:
```

Ecoli Network



It is also possible to produce a graph showing correlations as edge labels:

```
igr2 = network.make.igraph(ecoli.net, node.labels, show.edge.labels=TRUE)
```

plot(igr2, main="Ecoli Network with Partial Correlations as Edge Labels",
vertex.label.cex=0.8, edge.arrow.size=0.8)

Ecoli Network with Partial Correlations as Edge Labels

