Project 8

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Problem 20: Classical NEMs

1. For each model, construct the transitive closure (by adding edges) and define the corresponding adjacency matrices Φ and Θ , which represent the signalling pathways and the E-gene attachments. Determine the corresponding expected effect patterns (F).

Construct phi

```
theta2["S3",] = c(0,0,0,0,0,0)
theta2["S4",] = c(0,0,1,0,0,0)
theta2["S5",] = c(0,0,0,0,1,0)
```

Construct thetha

```
F1 = phi1%*%theta1
F2 = phi2%*%theta2
print("F1")
Calculate F = \Phi\Theta
## [1] "F1"
F1
##
     E1 E2 E3 E4 E5 E6
## S1 1 1 1 0 1 0
## S2 0 0 0 1 1 1
## S3 1 1 1 0 1 0
## S4 0 0 1 0 1 0
## S5 0 0 0 0 1 0
print("F2")
## [1] "F2"
F2
     E1 E2 E3 E4 E5 E6
##
## S1 1 1 1 0 1 0
## S2 0 0 0 1 1 1
## S3 1 1 1 0 1 0
## S4 0 0 1 0 1 0
## S5 0 0 0 0 1 0
```

2. Assuming no noise, determine the discrete data D1 and D2 from both models. Given only the data, can you tell apart the two models?

```
D1 = array(dim = c(6, 5), dimnames = list(c("E1", "E2", "E3", "E4", "E5", "E6"), c("S1", "S2", "S3", "S4", "S5")))

D1["E1",] = c(1,0,1,0,0)

D1["E2",] = c(1,0,1,0,0)

D1["E3",] = c(1,0,1,1,0)

D1["E4",] = c(0,1,0,0,0)
```

```
D1["E5",] = c(1,1,1,1,0)
D1["E6",] = c(0,1,0,0,1)
D2 = array(dim = c(6, 5), dimnames = list(c("E1", "E2", "E3", "E4", "E5", "E6"),
                                         c("S1", "S2", "S3", "S4", "S5")))
D2["E1",] = c(1,0,1,0,0)
D2["E2",] = c(1,0,1,0,0)
D2["E3",] = c(1,0,1,1,0)
D2["E4",] = c(0,1,0,0,0)
D2["E5",] = c(1,1,1,1,0)
D2["E6",] = c(0,1,0,0,1)
D1
##
     S1 S2 S3 S4 S5
## E1 1 0 1 0 0
## E2 1 0 1 0 0
## E3 1 0 1 1 0
## E4 0 1 0 0 0
## E5
     1 1 1 1 0
## E6 0 1 0 0 1
D2
     S1 S2 S3 S4 S5
##
## E1 1 0 1 0 0
## E2 1 0 1 0 0
## E3 1 0 1 1 0
## E4 0 1 0 0 0
## E5
     1 1 1 1 0
## E6 0 1 0 0 1
Since the Data matrices D1 and D2 are identical, we cannot tell the two models apart.
library(mnem)
## Registered S3 methods overwritten by 'RcppEigen':
##
    method
                         from
##
    predict.fastLm
                         RcppArmadillo
##
    print.fastLm
                         RcppArmadillo
##
     summary.fastLm
                         RcppArmadillo
    print.summary.fastLm RcppArmadillo
nem1 = nem(D1, marginal = TRUE, fpfn = c(0.05, 0.01))
nem1$score
```

[1] 28.33379

```
nem2 = nem(D2,marginal = TRUE,fpfn = c(0.05,0.01))
nem2$score

## [1] 28.33379

###Not sure which one it is
#scoreAdj(D1,adj = phi1,method="disc",marginal=TRUE,fpfn=c(0.05,0.01))$score
#scoreAdj(D2,adj = phi2,method="disc",marginal=TRUE,fpfn=c(0.05,0.01))$score
```

Problem 21: Hidden Markov NEMs

1. Using the definitions for HM-NEMs from the lecture, compute the transition probabilities from $G_t = u$ to $G_{t+1} \in v1, v2$ for different smoothness parameter $\lambda \in 0.1, \dots 0.9$.

```
## not sure if the enumerate has to be used and how?
u = t(array(c(c(1,1,1,0)),
              c(0,1,1,1),
              c(0,0,1,1),
              c(0,0,0,1)),
              dim = c(4, 4), dimnames = list(c("S1", "S2", "S3", "S4"),
                                              c("S1", "S2", "S3", "S4"))))
v1 = t(array(c(c(1,1,1,0)),
              c(0,1,1,1),
              c(0,0,1,0),
              c(0,0,0,1)),
              dim = c(4, 4), dimnames = list(c("S1", "S2", "S3", "S4"),
                                             c("S1", "S2", "S3", "S4"))))
v2 = t(array(c(c(1,0,0,0)),
              c(1,1,1,0),
              c(1,0,1,0),
              c(1,0,0,1)),
              dim = c(4, 4), dimnames = list(c("S1", "S2", "S3", "S4"),
                                              c("S1", "S2", "S3", "S4"))))
lambda = seq(0.1, 0.9, by=0.1)
s_uv1 = sum(u!=v1)
s_uv2 = sum(u!=v2)
T = array(dim = c(9,2), dimnames = list(lambda, c("v1", "v2")))
C = array(dim = c(9,1), dimnames = list(lambda,c("C")))
for(i in lambda){
  C[as.character(i),] = ((1-i)^s_uv1)*i + ((1-i)^s_uv2)*i
  T[as.character(i),"v1"] = (1/C[as.character(i),])*((1-i)^s_uv1)*i
  T[as.character(i),"v2"] = (1/C[as.character(i),])*((1-i)^s_uv2)*i
}
Τ
```

```
## v1 v2
## 0.1 0.6529798 3.470202e-01
```

```
## 0.2 0.7923026 2.076974e-01

## 0.3 0.8947353 1.052647e-01

## 0.4 0.9554237 4.457625e-02

## 0.5 0.9846154 1.538462e-02

## 0.6 0.9959207 4.079291e-03

## 0.7 0.9992715 7.284689e-04

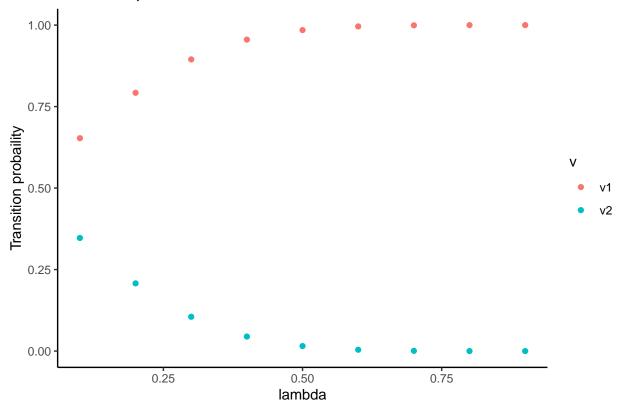
## 0.8 0.9999360 6.399590e-05

## 0.9 0.999990 9.999990e-07
```

2. Plot the transition probabilities for v_1 and v_2 as a function of λ . Describe the transition probabilities as a function of λ .

```
library(reshape2)
library(ggplot2)
library(RColorBrewer)
data = data.frame(melt(T))
colnames(data)<-c("lambda","v","T")
plot<-ggplot(data,aes(x=lambda,y=T,color=v))+
    geom_point()+
    theme_classic()+
    ylab("Transition probability")+
    labs(title="Transition probabilities as a function of lambda")
plot</pre>
```

Transition probabilities as a function of lambda



Problem 22: Mixture NEMs

1. Determine the the cellular perturbation map ρ , where $\rho_{ic} = 1$ if cell c is perturbed by a knock-down of S-gene i.

2. Assume that C_1, C_2 are generated from F_1 and C_3, C_4 are generated from F_2 , compute the noiseless log odds matrix R, where $R_{jc} > 0$ means that the perturbation on cell c has an effect on E-gene j:

```
phi_F1 = array(dim = c(2,2), dimnames = list(c("S1", "S2"),
                                           c("S1", "S2")))
phi_F1["S1",] = c(1,1)
phi_F1["S2",] = c(0,1)
phi_F2 = array(dim = c(2,2), dimnames = list(c("S1", "S2"),
                                           c("S1","S2")))
phi_F2["S1",] = c(1,0)
phi_F2["S2",] = c(1,1)
theta_F1 = array(\dim = c(2,2), \dim = list(c("S1","S2"),
                                              c("E1","E2")))
theta_F1["S1",] = c(1,0)
theta_F1["S2",] = c(0,1)
theta_F2 = array(\dim = c(2,2), \dim = list(c("S1","S2"),
                                              c("E1","E2")))
theta_F2["S1",] = c(0,1)
theta_F2["S2",] = c(1,0)
EEP_F1 = t(t(rho)%*\%phi_F1%*\%theta_F1)
EEP_F1[EEP_F1>1] = 1
EEP_F2 = t(t(rho)%*%phi_F1%*%theta_F2)
EEP_F2[EEP_F2>1] = 1
print("Expected effect pattern of F1")
```

- (a) For each component k, compute the expected effect pattern $(\rho^T \phi^k \theta^k)^T$. Replace all non-zeros by 1.
- ## [1] "Expected effect pattern of F1"

```
EEP_F1
      C1 C2 C3 C4
##
## E1 1 0 1 0
## E2 1 1 1
print("Expected effect pattern of F2")
## [1] "Expected effect pattern of F2"
EEP_F2
     C1 C2 C3 C4
##
## E1 1 1 1 1
## E2 1 0 1 0
R= cbind(EEP_F1[,1:2],EEP_F2[,3:4])
R[R==0] = -1
(b) Based on the component assignment for each cell, extract the corresponding column from
the expected effect patterns computed above and put it into R. Replace all zeros by -1.
```

3. Take R from the previous question. Given the vector of mixture weights $\pi = (0.44, 0.56)$, calculate the responsibilities Γ . Then, update the mixture weights.

```
L1 = t(EEP_F1)\%*\%R
L2 = t(EEP_F2)\%*\%R
pi = c(0.44, 0.56)
gamma = array(dim = c(2,4), dimnames = list(c("F1","F2"),
                                             c("C1", "C2", "C3", "C4")))
gamma["F1",] = pi[1]*diag(L1)/sum(pi[1]*diag(L1))
gamma["F2",] = pi[2]*diag(L2)/sum(pi[2]*diag(L2))
##Responsibilities should be in [0,1]??
print("Responsibilities")
```

[1] "Responsibilities"

gamma

```
##
      C1
            C2 C3
                      C4
## F1 0.5 0.25 0.5 -0.25
## F2 0.5 -0.25 0.5 0.25
```

```
pi[1] = sum(gamma["F1",])/(sum(gamma["F1",])+sum(gamma["F2",]))
pi[2] = sum(gamma["F2",])/(sum(gamma["F1",])+sum(gamma["F2",]))

print("Updated mixture weights")

## [1] "Updated mixture weights"

pi

## [1] 0.5 0.5
```