Project 5: Statistical phylogenetics

Team C - Minghang Li, Xiaocheng Yang, Xinyi Chen

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Problem 12: Transition matrix, rate matrix, and stationary distribution

1. Show that $\frac{dP(t)}{dt} = R \cdot P(t)$

Proof.

$$\begin{split} \frac{dP(t)}{dt} &= \frac{P(dt+t) - P(t)}{dt} & \text{(Definition of derivative)} \\ &= \frac{P(dt)P(t) - P(t)}{dt} & \text{(Chapman-Kolmogorov's equation)} \\ &= \frac{(I+Rdt)P(t) - P(t)}{dt} & \text{(Definition of } P(dt)) \\ &= \frac{P(t) + RP(t)dt - P(t)}{dt} \\ &= \frac{RP(t)dt}{dt} \\ &= RP(t) \end{split}$$

2. Assume that the given Markov chain is ergodic with (unique) stationary distribution π , show that $R\pi=0$

Proof. The stationary distribution π satisfies:

$$P(t)\pi = \pi$$

after a sufficiently long time t and any time point that follows. Hence, the following equation also holds:

$$P(t+dt)\pi = \pi$$

We can further re-write the left-hand side of the equation:

$$\begin{split} P(dt+t)\pi &= P(dt)P(t)\pi & \text{(Chapman-Kologomorov's equation)} \\ &= (I+Rdt)P(t)\pi & \text{(Definition of } P(dt)) \\ &= P(t)\pi + R\cdot P(t)dt\pi \\ &= \pi + Rdt\pi & \text{(Plug in } P(t)\pi = \pi) \end{split}$$

which leads us to the following equality:

$$\pi + Rdt\pi = \pi$$

$$Rdt\pi = 0$$

$$R\pi = 0$$

Problem 13: Phylogenetic trees as Bayesian networks

1. What is the joint probability P(X, Z|T) of the tree?

$$P(X,Z|T) = \pi(Z_4)P(X_5|Z_4)P(Z_3|Z_4)P(Z_2|Z_3)P(Z_1|Z_3)P(X_4|Z_2)P(X_3|Z_2)P(X_2|Z_1)P(X_1|Z_1)$$

2. How many summation steps would be required for the naive calculation of P(X|T) via brute-force marginalization over the hidden nodes Z?

Since there are 4 possibilities (A, C, G, T) for each hidden node:

$$4 \times 4 \times 4 \times 4 = 256$$

3. Rearrange the expression P(X|T) such that the number of operations is minimized. How many summation steps are required now for the calculation of P(X|T)?

$$\begin{split} P(X|T) &= \sum_{Z_4} \sum_{Z_3} \sum_{Z_2} \sum_{Z_1} \pi(Z_4) P(X_5|Z_4) P(Z_3|Z_4) P(Z_2|Z_3) P(Z_1|Z_3) P(X_4|Z_2) P(X_3|Z_2) P(X_2|Z_1) P(X_1|Z_1) \\ &= \sum_{Z_4} \pi(Z_4) P(X_5|Z_4) \sum_{Z_3} P(Z_3|Z_4) \sum_{Z_2} P(Z_2|Z_3) P(X_4|Z_2) P(X_3|Z_2) \sum_{Z_1} P(Z_1|Z_3) P(X_2|Z_1) P(X_1|Z_1) \\ &= \underbrace{\sum_{Z_4} \pi(Z_4) P(X_5|Z_4)}_{4 \times 4} \underbrace{\sum_{Z_3} P(Z_3|Z_4)}_{4 \times 4} \underbrace{\sum_{Z_2} P(Z_2|Z_3) P(X_4|Z_2) P(X_3|Z_2)}_{4 \times 4(Z_2 \to X_4 \text{ branch}) + 4 \times 4(Z_2 \to X_3 \text{ branch})} \underbrace{\sum_{Z_1} P(Z_1|Z_3) P(X_2|Z_1) P(X_1|Z_1)}_{4 \times 4(Z_1 \to X_2 \text{ branch}) + 4 \times 4(Z_1 \to X_3 \text{ branch})} \underbrace{\sum_{Z_1} P(Z_1|Z_3) P(X_2|Z_1) P(X_1|Z_1)}_{4 \times 4(Z_1 \to X_4 \text{ branch}) + 4 \times 4(Z_2 \to X_3 \text{ branch})} \underbrace{\sum_{Z_1} P(Z_1|Z_3) P(X_2|Z_1) P(X_1|Z_1)}_{4 \times 4(Z_1 \to X_4 \text{ branch}) + 4 \times 4(Z_2 \to X_3 \text{ branch})} \underbrace{\sum_{Z_1} P(Z_1|Z_2) P(X_2|Z_1) P(X_1|Z_1)}_{4 \times 4(Z_1 \to X_2 \text{ branch}) + 4 \times 4(Z_1 \to X_3 \text{ branch})} \underbrace{\sum_{Z_1} P(Z_1|Z_2) P(X_2|Z_1) P(X_1|Z_1)}_{4 \times 4(Z_1 \to X_4 \text{ branch}) + 4 \times 4(Z_2 \to X_3 \text{ branch})} \underbrace{\sum_{Z_1} P(Z_1|Z_2) P(X_2|Z_1) P(X_2|Z_1) P(X_2|Z_1)}_{4 \times 4(Z_1 \to X_4 \text{ branch}) + 4 \times 4(Z_2 \to X_3 \text{ branch})} \underbrace{\sum_{Z_1} P(Z_1|Z_2) P(X_2|Z_1) P(X_2|Z_1) P(X_2|Z_1)}_{4 \times 4(Z_1 \to X_4 \text{ branch}) + 4 \times 4(Z_2 \to X_3 \text{ branch})} \underbrace{\sum_{Z_1} P(Z_1|Z_2) P(X_2|Z_1) P(X_2|Z_1)}_{4 \times 4(Z_1 \to X_4 \text{ branch}) + 4 \times 4(Z_1 \to X_4 \text{ branch})} \underbrace{\sum_{Z_1} P(Z_1|Z_2) P(X_2|Z_2) P(X_2|Z_2)}_{4 \times 4(Z_1 \to X_4 \text{ branch}) + 4 \times 4(Z_1 \to X_4 \text{ branch})} \underbrace{\sum_{Z_1} P(Z_1|Z_2) P(X_2|Z_2) P(X_2|Z_2)}_{4 \times 4(Z_1 \to X_4 \text{ branch})} \underbrace{\sum_{Z_1} P(Z_1|Z_2) P(X_2|Z_2) P(X_2|Z_2)}_{4 \times 4(Z_1 \to X_4 \text{ branch})} \underbrace{\sum_{Z_1} P(Z_1|Z_2) P(Z_1|Z_2) P(Z_1|Z_2)}_{4 \times 4(Z_1 \to X_4 \text{ branch})} \underbrace{\sum_{Z_1} P(Z_1|Z_2) P(Z_1|Z_2)}_{4 \times 4(Z_1 \to X_4 \text{ branch})} \underbrace{\sum_{Z_1} P(Z_1|Z_2) P(Z_1|Z_2)}_{4 \times 4(Z_1 \to X_4 \text{ branch})} \underbrace{\sum_{Z_1} P(Z_1|Z_2)}_{4 \times 4(Z_$$

In total we have $(16+16) \times 2 + 16 + 4 = 52$ summations.

Problem 14: Learning phylogenetic trees from sequence alignment data

Load the dataset ParisRT.txt.

```
data <- read.dna("ParisRT.txt", format="sequential")
data

## 17 DNA sequences in binary format stored in a matrix.

##
## All sequences of same length: 618

##
## Labels:
## B_OYI
## B_HXB2
## D_eli
## Mr_D
## Mme_L
## A_ibng
## ...
##</pre>
```

```
## Base composition:
## a c g t
## 0.404 0.162 0.195 0.239
## (Total: 10.51 kb)
```

Create initial tree topology for the alignment using neighbor joining and K80 model.

```
set.seed(42)
tree.ini <- NJ(dist.dna(data, model = "K80"))</pre>
```

Plot the initial tree.

```
plot.phylo(tree.ini)
```

```
A ibng
                            -A u455
                           Mr D
                                                                                      O MVP5180
                                                                                      O ant
               D Z2Z6
             D ndk
                -D eli
       B manc
       -Mme S
      -Mme L
     -B JRFL
    -B yu2
  <sub>I</sub>B lai
  LB HXB2
    -B OYI
    −B weau
tree.ML <- pml(tree.ini, phyDat(data), model = "K80")</pre>
tree.ML
## model: K80
## loglikelihood: -3003.487
## unconstrained loglikelihood: -2098.897
##
## Rate matrix:
##
   acgt
## a 0 1 1 1
## c 1 0 1 1
## g 1 1 0 1
## t 1 1 1 0
##
## Base frequencies:
          С
                g
```

Find the optimal parameters of K80 model for rate matrix.

0.25 0.25 0.25 0.25

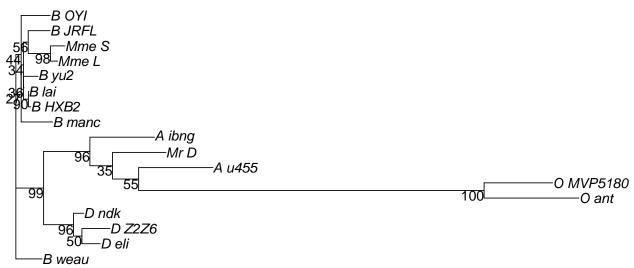
```
tree.optim.Q <- optim.pml(
    tree.ML,
    optQ = TRUE,
    optNni = FALSE,
    optBf = FALSE,
    optInv = FALSE,
    optGamma = FALSE,
    optEdge = FALSE,</pre>
```

```
optRate = FALSE,
  optRooted = FALSE,
  model = "K80"
## optimize rate matrix: -3003.487 --> -2884.408
## optimize rate matrix: -2884.408 --> -2884.408
tree.optim.Q
## model: K80
## loglikelihood: -2884.408
## unconstrained loglikelihood: -2098.897
##
## Rate matrix:
##
            a
                      С
                               g
## a 0.000000 1.000000 4.976955 1.000000
## c 1.000000 0.000000 1.000000 4.976955
## g 4.976955 1.000000 0.000000 1.000000
## t 1.000000 4.976955 1.000000 0.000000
##
## Base frequencies:
##
   a
         c g
## 0.25 0.25 0.25 0.25
Optimize for branch lengths, nucelotide substitution rates and tree topology simultaneously.
tree.optim <- optim.pml(</pre>
 tree.ML,
  optQ = TRUE,
                   # rate matrix
 optNni = TRUE,  # tree topology
optBf = FALSE,  # base frequencies
optInv = FALSE,  # proportion of var size
  optGamma = FALSE, # gamma rate param
 optEdge = TRUE, # edge lengths
  optRate = FALSE, # overall rate
  optRooted = FALSE, # edge lengths of a rooted tree
  model = "K80"
)
## optimize edge weights: -3003.487 --> -2992.981
## optimize rate matrix: -2992.981 --> -2873.703
## optimize edge weights: -2873.703 --> -2872.892
## optimize topology: -2872.892 --> -2859.775 NNI moves: 5
## optimize rate matrix: -2859.775 --> -2859.682
## optimize edge weights: -2859.682 --> -2859.681
## optimize topology: -2859.681 --> -2859.681 NNI moves: 0
## optimize rate matrix: -2859.681 --> -2859.681
## optimize edge weights: -2859.681 --> -2859.681
tree.optim
## model: K80
## loglikelihood: -2859.681
## unconstrained loglikelihood: -2098.897
##
## Rate matrix:
```

```
## a c g t
## a 0.000000 1.000000 5.262144 1.000000
## c 1.000000 0.000000 1.000000 5.262144
## g 5.262144 1.000000 0.000000 1.000000
## t 1.000000 5.262144 1.000000 0.000000
##
## Base frequencies:
## a c g t
## 0.25 0.25 0.25 0.25
```

Bootstrap on optimised model. The program resampled the nucleotides in the alignment (columns in the input matrix).

```
plotBS(tree.optim$tree, bootstrap, type = "phylogram")
```



 Mme_S is more likely to affect patient Mme_L . The bootstrap support is 98% (in my run with seed 42), suggesting Mme_S and Mme_L share a more recent ancestor (closer relationship) than Mme_D