```
\# xy = matrix(runif(10000), ncol = 2) xy = xy[xy[,1] < 0.1 | xy[,2] < 0.1,]
\# xy = xy \% \% \text{ matrix}(c(1, 0.4, 0.4, 1), ncol = 2) xy = xy[xy[,1] <= 1 &
\# xy[,2] <= 1,]
xyc = read.csv("synthetic_data.csv")
xy = xyc[, 1:2]
set.seed(1234)
subset = sample.int(nrow(xy), 300)
library(fastICA)
library(NMF)
## Loading required package: methods
## Loading required package: pkgmaker
## Loading required package: registry
## Loading required package: rngtools
## Loading required package: cluster
## NMF - BioConductor layer [OK] | Shared memory capabilities [OK] | Cores 7/8
fit.pca = prcomp(xy, center = TRUE, scale = FALSE)
temp = replicate(1000, fastICA(xy, 2, method = "C"), simplify = FALSE)
temp2 = sapply(temp, function(x) shapiro.test(x$S)$statistic)
fit.ica = temp[[which.max(temp2)]]
fit.nmf = nmf(t(xy[subset, ]), rank = 2, nrun = 20, method = "snmf/r")
library(NMF)
## Loading required package: methods
## Loading required package: pkgmaker
## Loading required package: registry
## Loading required package: rngtools
## Loading required package: cluster
## NMF - BioConductor layer [OK] | Shared memory capabilities [OK] | Cores 7/8
library(RColorBrewer)
pal = brewer.pal(3, "Set2")[c(2, 3, 1)]
pal = sapply(pal, function(col) do.call(rgb, c(as.list(col2rgb(col)/255), alpha = 0.5)))
syms = c(19, 4, 21)
```

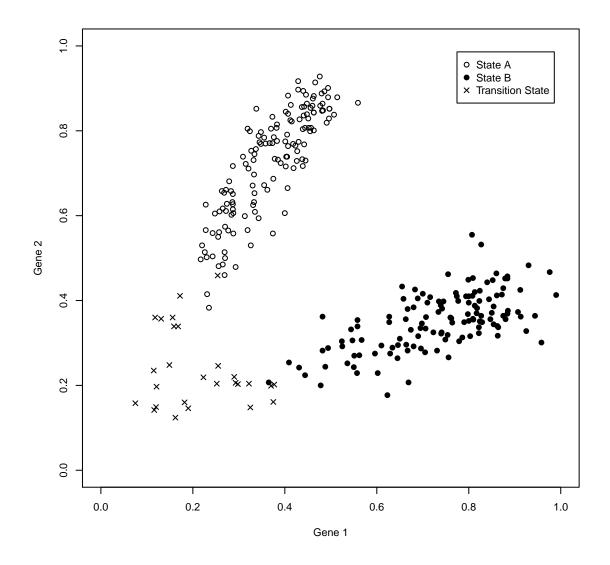
plot(xy[subset, 1], xy[subset, 2], col = "black", pch = pch[subset], xlab = "Gene 1",

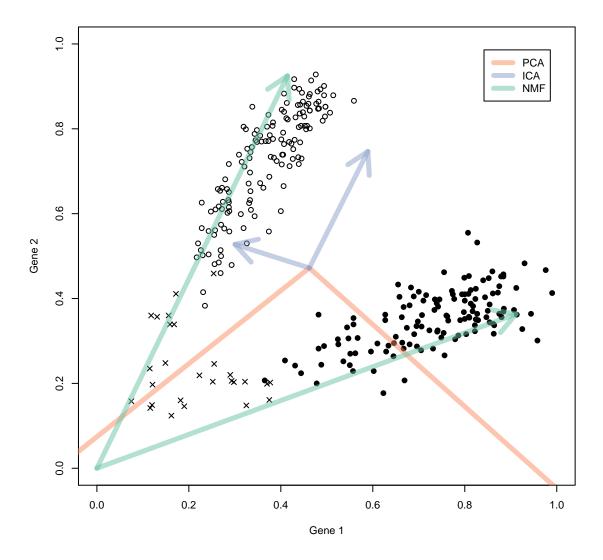
legend("topright", legend = c("State A", "State B", "Transition State"), pch = syms[c(3,

ylab = "Gene 2", xlim = c(0, 1), ylim = c(0, 1))

col = pal[xyc[, 3]]
pch = syms[xyc[, 3]]

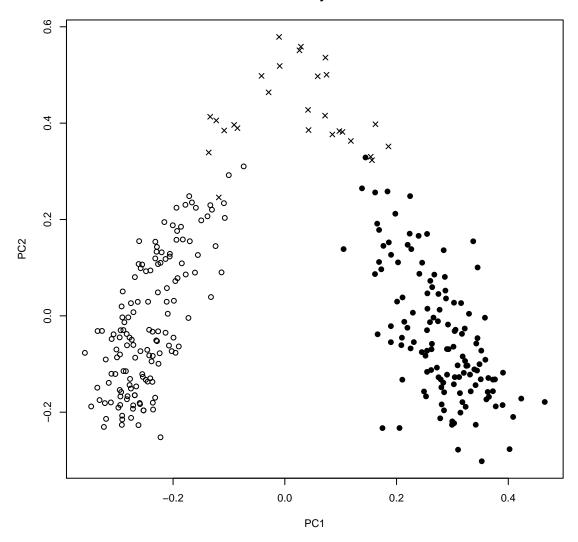
1, 2), inset = 0.05)





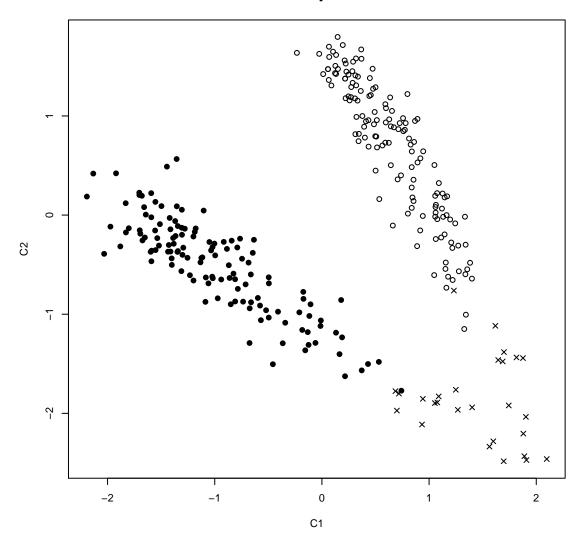
plot(fit.pca\$x[subset,], pch = pch[subset], xlab = "PC1", ylab = "PC2", main = "PCA Projection")

PCA Projection



plot(fit.ica\$S[subset,], pch = pch[subset], xlab = "C1", ylab = "C2", main = "ICA Projection")

ICA Projection



plot(t(coef(fit.nmf)), pch = pch[subset], xlab = "F1", ylab = "F2", main = "NMF Projection")

NMF Projection

