1. Done

```
2. mydata <- read.table("sle_b_cell.txt", header=T, row.names=1)
    > mydata <- read.table("sle_b_cell.txt", header=T, row.names=1)
    > length(mydata)
[1] 26
    > dim(mydata)
[1] 34853 26
```

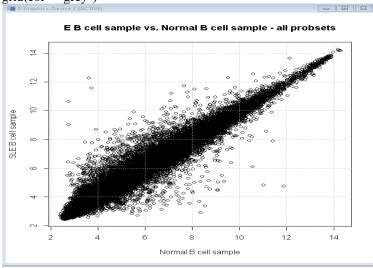
- 3. dim(mydata) produced 34853 and 26.
- 4. Mydata (prints the samples to the screen)

5. x <- c(mydata['control.1')]

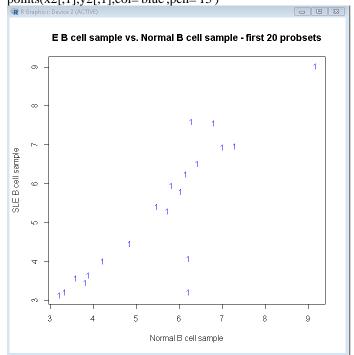
y <- (mydata['sle.2'])

plot(x[,1],y[,1],xlab='Normal B cell sample',ylab='SLE B cell sample',main='E B cell sample vs. Normal B cell sample - all probsets')

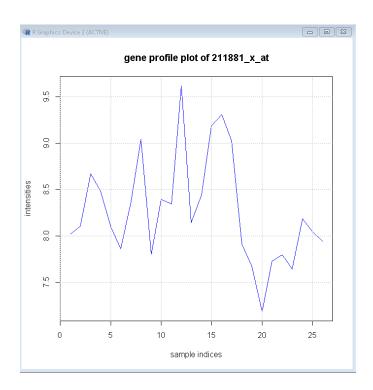




6. y2 <- data.frame(y = y[,1][1:20]) x2 <- data.frame(x = x[,1][1:20]) plot(x2[,1],y2[,1],type='n',xlab='Normal B cell sample',ylab='SLE B cell sample',main='E B cell sample vs. Normal B cell sample - first 20 probsets') points(x2[,1],y2[,1],col='blue',pch='15')



7. df <- data.frame(mydata) x.numeric = as.numeric(df['211881_x_at',]) plot(1:26,x.numeric,type ='n', xlab ='sample indices', ylab='intensities', main = 'gene profile plot of 211881_x_at') lines(1:26,x.numeric,col='blue') grid(col ='grey')



 $8. \quad f <- \ data.frame(x.numeric,c(rep("SLE",17),rep("Control",9))) \\ colnames(f) <- \ c('expression', 'sample') \\ box <- \ ggplot(f,aes(x=sample,y=expression,fill=sample)) + \\ ggtitle("gene \ profile \ across \ conditions") + \\ geom_boxplot()$

