

## Class\_16

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11/19/2021

```
library(DESeq2)
```

```
## Loading required package: S4Vectors
```

```
## Loading required package: stats4
```

```
## Loading required package: BiocGenerics
```

```
##
```

```
## Attaching package: 'BiocGenerics'
```

```
## The following objects are masked from 'package:stats':
```

```
##
```

```
##      IQR, mad, sd, var, xtabs
```

```
## The following objects are masked from 'package:base':
```

```
##
```

```
##      anyDuplicated, append, as.data.frame, basename, cbind, colnames,  
##      dirname, do.call, duplicated, eval, evalq, Filter, Find, get, grep,  
##      grepl, intersect, is.unsorted, lapply, Map, mapply, match, mget,  
##      order, paste, pmax, pmax.int, pmin, pmin.int, Position, rank,  
##      rbind, Reduce, rownames, sapply, setdiff, sort, table, tapply,  
##      union, unique, unsplit, which.max, which.min
```

```
##
```

```
## Attaching package: 'S4Vectors'
```

```
## The following objects are masked from 'package:base':
```

```
##
```

```
##      expand.grid, I, unname
```

```
## Loading required package: IRanges
```

```
## Loading required package: GenomicRanges
```

```
## Loading required package: GenomeInfoDb
```

```
## Loading required package: SummarizedExperiment
```

```

## Loading required package: MatrixGenerics

## Loading required package: matrixStats

##
## Attaching package: 'MatrixGenerics'

## The following objects are masked from 'package:matrixStats':
##
##   colAlls, colAnyNAs, colAnys, colAvgsPerRowSet, colCollapse,
##   colCounts, colCummaxs, colCummins, colCumprods, colCumsums,
##   colDiffs, colIQRDiffs, colIQRs, colLogSumExps, colMadDiffs,
##   colMads, colMaxs, colMeans2, colMedians, colMins, colOrderStats,
##   colProds, colQuantiles, colRanges, colRanks, colSdDiffs, colSds,
##   colSums2, colTabulates, colVarDiffs, colVars, colWeightedMads,
##   colWeightedMeans, colWeightedMedians, colWeightedSds,
##   colWeightedVars, rowAlls, rowAnyNAs, rowAnys, rowAvgsPerColSet,
##   rowCollapse, rowCounts, rowCummaxs, rowCummins, rowCumprods,
##   rowCumsums, rowDiffs, rowIQRDiffs, rowIQRs, rowLogSumExps,
##   rowMadDiffs, rowMads, rowMaxs, rowMeans2, rowMedians, rowMins,
##   rowOrderStats, rowProds, rowQuantiles, rowRanges, rowRanks,
##   rowSdDiffs, rowSds, rowSums2, rowTabulates, rowVarDiffs, rowVars,
##   rowWeightedMads, rowWeightedMeans, rowWeightedMedians,
##   rowWeightedSds, rowWeightedVars

## Loading required package: Biobase

## Welcome to Bioconductor
##
##   Vignettes contain introductory material; view with
##   'browseVignettes()'. To cite Bioconductor, see
##   'citation("Biobase)"', and for packages 'citation("pkgname)"'.

##
## Attaching package: 'Biobase'

## The following object is masked from 'package:MatrixGenerics':
##
##   rowMedians

## The following objects are masked from 'package:matrixStats':
##
##   anyMissing, rowMedians

metaFile <- "GSE37704_metadata.csv"
countFile <- "GSE37704_featurecounts.csv"

# Import metadata and take a peak
colData = read.csv(metaFile, row.names=1)
head(colData)

```

```
##                condition
## SRR493366 control_sirna
## SRR493367 control_sirna
## SRR493368 control_sirna
## SRR493369      hoxa1_kd
## SRR493370      hoxa1_kd
## SRR493371      hoxa1_kd
```

```
countData = read.csv(countFile, row.names=1)
head(countData)
```

```
##                length SRR493366 SRR493367 SRR493368 SRR493369 SRR493370
## ENSG00000186092      918         0         0         0         0         0
## ENSG00000279928      718         0         0         0         0         0
## ENSG00000279457     1982        23        28        29        29        28
## ENSG00000278566      939         0         0         0         0         0
## ENSG00000273547      939         0         0         0         0         0
## ENSG00000187634     3214        124        123        205        207        212
##                SRR493371
## ENSG00000186092         0
## ENSG00000279928         0
## ENSG00000279457        46
## ENSG00000278566         0
## ENSG00000273547         0
## ENSG00000187634        258
```

```
#Remove the first $length col
```

```
countData1 <- as.matrix(countData[, -1])
head(countData1)
```

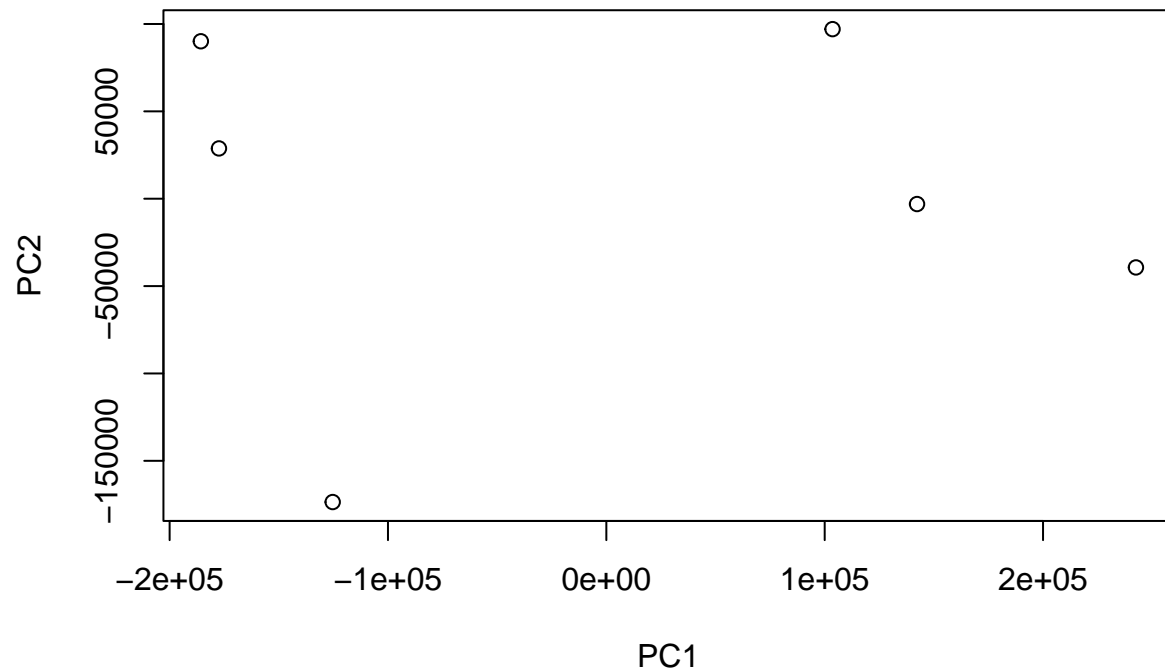
```
##                SRR493366 SRR493367 SRR493368 SRR493369 SRR493370 SRR493371
## ENSG00000186092         0         0         0         0         0         0
## ENSG00000279928         0         0         0         0         0         0
## ENSG00000279457        23        28        29        29        28        46
## ENSG00000278566         0         0         0         0         0         0
## ENSG00000273547         0         0         0         0         0         0
## ENSG00000187634        124        123        205        207        212        258
```

```
countData1 = countData1[-which(rowSums(countData1) == 0), ]
head(countData1)
```

```
##                SRR493366 SRR493367 SRR493368 SRR493369 SRR493370 SRR493371
## ENSG00000279457        23        28        29        29        28        46
## ENSG00000187634        124        123        205        207        212        258
## ENSG00000188976       1637       1831       2383       1226       1326       1504
## ENSG00000187961        120        153        180        236        255        357
## ENSG00000187583         24         48         65         44         48         64
## ENSG00000187642          4          9         16         14         16         16
```

```
#Run PCR To see that the data looks ok
```

```
pca <- prcomp(t(countData1))
plot(pca$x[,1:2])
```



#setup the DESeqDataSet object required for the DESeq() function and then run the DESeq pipeline

```
dds = DESeqDataSetFromMatrix(countData=countData1,
                              colData=colData,
                              design=~condition)
```

```
## Warning in DESeqDataSet(se, design = design, ignoreRank): some variables in
## design formula are characters, converting to factors
```

```
dds = DESeq(dds)
```

```
## estimating size factors
```

```
## estimating dispersions
```

```
## gene-wise dispersion estimates
```

```
## mean-dispersion relationship
```

```
## final dispersion estimates
```

```
## fitting model and testing
```

```
dds
```

```
## class: DESeqDataSet
## dim: 15975 6
## metadata(1): version
## assays(4): counts mu H cooks
## rownames(15975): ENSG00000279457 ENSG00000187634 ... ENSG00000276345
## ENSG00000271254
## rowData names(22): baseMean baseVar ... deviance maxCooks
## colnames(6): SRR493366 SRR493367 ... SRR493370 SRR493371
## colData names(2): condition sizeFactor
```

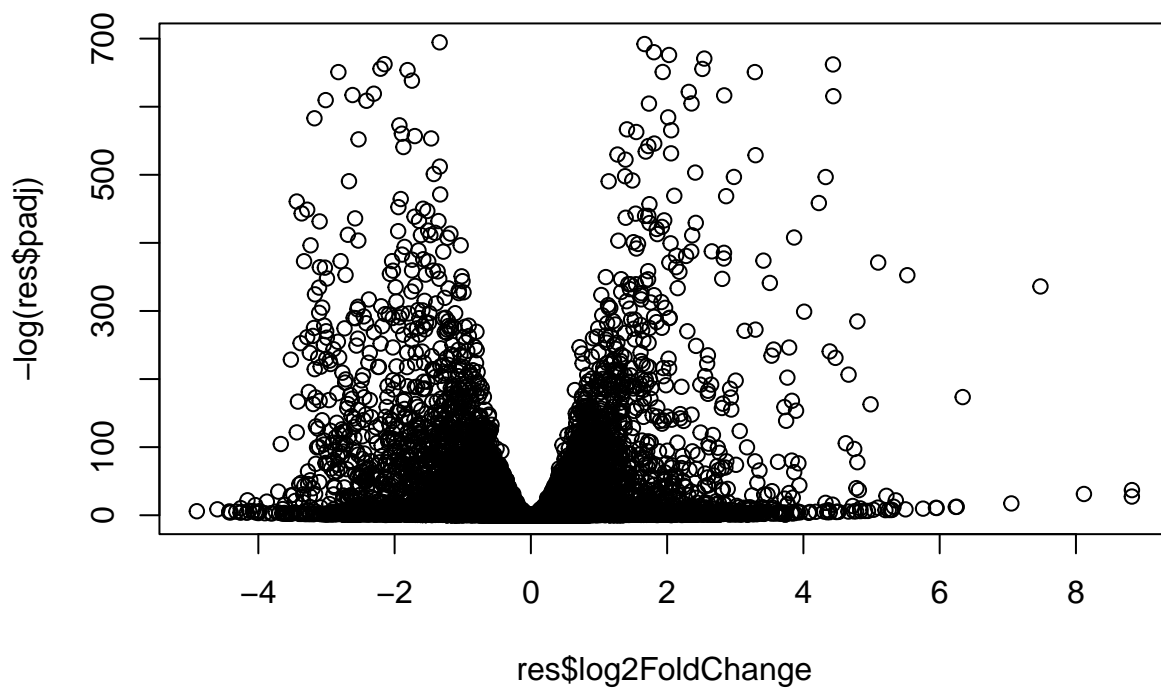
#get results for the HoxA1 knockdown versus control siRNA (remember that these were labeled as “hoxa1\_kd” and “control\_siRNA” in our original colData metaFile input to DESeq, you can check this above and by running resultsNames(dds) command)

```
res = results(dds)
```

```
summary(res)
```

```
##
## out of 15975 with nonzero total read count
## adjusted p-value < 0.1
## LFC > 0 (up)      : 4349, 27%
## LFC < 0 (down)    : 4396, 28%
## outliers [1]      : 0, 0%
## low counts [2]    : 1237, 7.7%
## (mean count < 0)
## [1] see 'cooksCutoff' argument of ?results
## [2] see 'independentFiltering' argument of ?results
```

```
plot( res$log2FoldChange, -log(res$padj) )
```

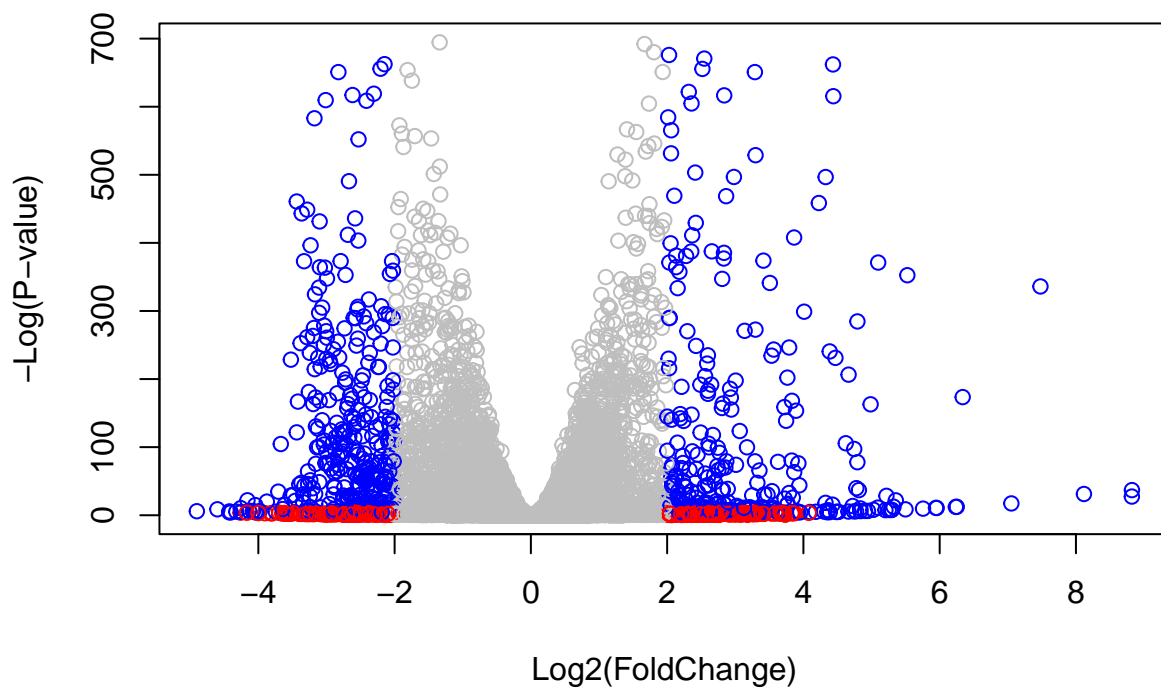


```
# Make a color vector for all genes
mycols <- rep("gray", nrow(res) )

# Color red the genes with absolute fold change above 2
mycols[ abs(res$log2FoldChange) > 2 ] <- "red"

# Color blue those with adjusted p-value less than 0.01
# and absolute fold change more than 2
inds <- (res$pvalue < 0.01) & (abs(res$log2FoldChange) > 2 )
mycols[ inds ] <- "blue"

plot( res$log2FoldChange, -log(res$padj), col=mycols, xlab="Log2(FoldChange)", ylab="-Log(P-value)" )
```



```
library("AnnotationDbi")
```

```
## Warning: package 'AnnotationDbi' was built under R version 4.1.2
```

```
library("org.Hs.eg.db")
```

```
##
```

```
columns(org.Hs.eg.db)
```

```
## [1] "ACCNUM"      "ALIAS"       "ENSEMBL"     "ENSEMBLPROT" "ENSEMBLTRANS"
## [6] "ENTREZID"    "ENZYME"      "EVIDENCE"    "EVIDENCEALL"  "GENENAME"
## [11] "GENETYPE"    "GO"          "GOALL"       "IPI"          "MAP"
## [16] "OMIM"        "ONTOLOGY"    "ONTOLOGYALL" "PATH"         "PFAM"
## [21] "PMID"        "PROSITE"     "REFSEQ"      "SYMBOL"       "UCSCKG"
## [26] "UNIPROT"
```

```
res$symbol = mapIds(org.Hs.eg.db,
  keys=row.names(res),
  keytype="ENSEMBL",
  column="SYMBOL",
  multiVals="first")
```

```
## 'select()' returned 1:many mapping between keys and columns
```

```
res$entrez = mapIds(org.Hs.eg.db,
                    keys=row.names(res),
                    keytype="ENSEMBL",
                    column="ENTREZID",
                    multiVals="first")
```

```
## 'select()' returned 1:many mapping between keys and columns
```

```
res$name = mapIds(org.Hs.eg.db,
                  keys=row.names(res),
                  keytype="ENSEMBL",
                  column="GENENAME",
                  multiVals="first")
```

```
## 'select()' returned 1:many mapping between keys and columns
```

```
head(res, 10)
```

```
## log2 fold change (MLE): condition hoxa1 kd vs control sirna
```

```
## Wald test p-value: condition hoxa1 kd vs control sirna
```

```
## DataFrame with 10 rows and 9 columns
```

##		baseMean	log2FoldChange	lfcSE	stat	pvalue
##		<numeric>	<numeric>	<numeric>	<numeric>	<numeric>
##	ENSG00000279457	29.913579	0.1792571	0.3248216	0.551863	5.81042e-01
##	ENSG00000187634	183.229650	0.4264571	0.1402658	3.040350	2.36304e-03
##	ENSG00000188976	1651.188076	-0.6927205	0.0548465	-12.630158	1.43990e-36
##	ENSG00000187961	209.637938	0.7297556	0.1318599	5.534326	3.12428e-08
##	ENSG00000187583	47.255123	0.0405765	0.2718928	0.149237	8.81366e-01
##	ENSG00000187642	11.979750	0.5428105	0.5215598	1.040744	2.97994e-01
##	ENSG00000188290	108.922128	2.0570638	0.1969053	10.446970	1.51282e-25
##	ENSG00000187608	350.716868	0.2573837	0.1027266	2.505522	1.22271e-02
##	ENSG00000188157	9128.439422	0.3899088	0.0467163	8.346304	7.04321e-17
##	ENSG00000237330	0.158192	0.7859552	4.0804729	0.192614	8.47261e-01
##		padj	symbol	entrez	name	
##		<numeric>	<character>	<character>	<character>	
##	ENSG00000279457	6.86555e-01	WASH9P	102723897	WAS protein family h..	
##	ENSG00000187634	5.15718e-03	SAMD11	148398	sterile alpha motif ..	
##	ENSG00000188976	1.76549e-35	NOC2L	26155	NOC2 like nucleolar ..	
##	ENSG00000187961	1.13413e-07	KLHL17	339451	kelch like family me..	
##	ENSG00000187583	9.19031e-01	PLEKHN1	84069	pleckstrin homology ..	
##	ENSG00000187642	4.03379e-01	PERM1	84808	PPARGC1 and ESRR ind..	
##	ENSG00000188290	1.30538e-24	HES4	57801	hes family bHLH tran..	
##	ENSG00000187608	2.37452e-02	ISG15	9636	ISG15 ubiquitin like..	
##	ENSG00000188157	4.21963e-16	AGRN	375790	agrin	
##	ENSG00000237330	NA	RNF223	401934	ring finger protein ..	

```
#Reorder the results by adjusted p-value and save them to a CSV file in your current project directory.
```

```
res = res[order(res$pvalue),]
write.csv(res, file="deseq_results.csv")
```



```
library(pathview)
```

```
## #####  
## Pathview is an open source software package distributed under GNU General  
## Public License version 3 (GPLv3). Details of GPLv3 is available at  
## http://www.gnu.org/licenses/gpl-3.0.html. Particullary, users are required to  
## formally cite the original Pathview paper (not just mention it) in publications  
## or products. For details, do citation("pathview") within R.  
##  
## The pathview downloads and uses KEGG data. Non-academic uses may require a KEGG  
## license agreement (details at http://www.kegg.jp/kegg/legal.html).  
## #####
```

```
library(gage)
```

```
##
```

```
library(gageData)
```

```
data(kegg.sets.hs)
```

```
data(sigmet.idx.hs)
```

```
# Focus on signaling and metabolic pathways only
```

```
kegg.sets.hs = kegg.sets.hs[sigmet.idx.hs]
```

```
# Examine the first 3 pathways
```

```
head(kegg.sets.hs, 3)
```

```
## $'hsa00232 Caffeine metabolism'
```

```
## [1] "10" "1544" "1548" "1549" "1553" "7498" "9"
```

```
##
```

```
## $'hsa00983 Drug metabolism - other enzymes'
```

```
## [1] "10" "1066" "10720" "10941" "151531" "1548" "1549" "1551"
```

```
## [9] "1553" "1576" "1577" "1806" "1807" "1890" "221223" "2990"
```

```
## [17] "3251" "3614" "3615" "3704" "51733" "54490" "54575" "54576"
```

```
## [25] "54577" "54578" "54579" "54600" "54657" "54658" "54659" "54963"
```

```
## [33] "574537" "64816" "7083" "7084" "7172" "7363" "7364" "7365"
```

```
## [41] "7366" "7367" "7371" "7372" "7378" "7498" "79799" "83549"
```

```
## [49] "8824" "8833" "9" "978"
```

```
##
```

```
## $'hsa00230 Purine metabolism'
```

```
## [1] "100" "10201" "10606" "10621" "10622" "10623" "107" "10714"
```

```
## [9] "108" "10846" "109" "111" "11128" "11164" "112" "113"
```

```
## [17] "114" "115" "122481" "122622" "124583" "132" "158" "159"
```

```
## [25] "1633" "171568" "1716" "196883" "203" "204" "205" "221823"
```

```
## [33] "2272" "22978" "23649" "246721" "25885" "2618" "26289" "270"
```

```
## [41] "271" "27115" "272" "2766" "2977" "2982" "2983" "2984"
```

```
## [49] "2986" "2987" "29922" "3000" "30833" "30834" "318" "3251"
```

```
## [57] "353" "3614" "3615" "3704" "377841" "471" "4830" "4831"
```

```
## [65] "4832" "4833" "4860" "4881" "4882" "4907" "50484" "50940"
```

```
## [73] "51082" "51251" "51292" "5136" "5137" "5138" "5139" "5140"
```

```
## [81] "5141" "5142" "5143" "5144" "5145" "5146" "5147" "5148"
## [89] "5149" "5150" "5151" "5152" "5153" "5158" "5167" "5169"
## [97] "51728" "5198" "5236" "5313" "5315" "53343" "54107" "5422"
## [105] "5424" "5425" "5426" "5427" "5430" "5431" "5432" "5433"
## [113] "5434" "5435" "5436" "5437" "5438" "5439" "5440" "5441"
## [121] "5471" "548644" "55276" "5557" "5558" "55703" "55811" "55821"
## [129] "5631" "5634" "56655" "56953" "56985" "57804" "58497" "6240"
## [137] "6241" "64425" "646625" "654364" "661" "7498" "8382" "84172"
## [145] "84265" "84284" "84618" "8622" "8654" "87178" "8833" "9060"
## [153] "9061" "93034" "953" "9533" "954" "955" "956" "957"
## [161] "9583" "9615"
```

```
foldchanges = res$log2FoldChange
names(foldchanges) = res$entrez
head(foldchanges)
```

```
##      1266      54855      1465      51232      2034      2317
## -2.422719  3.201955 -2.313738 -2.059631 -1.888019 -1.649792
```

#Now, let's run the gage pathway analysis.

```
keggres = gage(foldchanges, gsets=kegg.sets.hs)
attributes(keggres)
```

```
## $names
## [1] "greater" "less" "stats"
```

```
head(keggres$less)
```

```
##                p.geomean stat.mean      p.val
## hsa04110 Cell cycle      8.995727e-06 -4.378644 8.995727e-06
## hsa03030 DNA replication  9.424076e-05 -3.951803 9.424076e-05
## hsa03013 RNA transport    1.375901e-03 -3.028500 1.375901e-03
## hsa03440 Homologous recombination 3.066756e-03 -2.852899 3.066756e-03
## hsa04114 Oocyte meiosis    3.784520e-03 -2.698128 3.784520e-03
## hsa00010 Glycolysis / Gluconeogenesis 8.961413e-03 -2.405398 8.961413e-03
##                q.val set.size      exp1
## hsa04110 Cell cycle      0.001448312      121 8.995727e-06
## hsa03030 DNA replication  0.007586381       36 9.424076e-05
## hsa03013 RNA transport    0.073840037      144 1.375901e-03
## hsa03440 Homologous recombination 0.121861535       28 3.066756e-03
## hsa04114 Oocyte meiosis    0.121861535      102 3.784520e-03
## hsa00010 Glycolysis / Gluconeogenesis 0.212222694       53 8.961413e-03
```

```
pathview(gene.data=foldchanges, pathway.id="hsa04110")
```

```
## 'select()' returned 1:1 mapping between keys and columns
```

```
## Info: Working in directory /Users/elizabethwest/Documents/Classes/Bioinformatics/R Files/bggn213/Class
```

```
## Info: Writing image file hsa04110.pathview.png
```



```
## Info: Writing image file hsa04630.pathview.png

## 'select()' returned 1:1 mapping between keys and columns

## Info: Working in directory /Users/elizabethwest/Documents/Classes/Bioinformatics/R Files/bggn213/Clas

## Info: Writing image file hsa00140.pathview.png

## 'select()' returned 1:1 mapping between keys and columns

## Info: Working in directory /Users/elizabethwest/Documents/Classes/Bioinformatics/R Files/bggn213/Clas

## Info: Writing image file hsa04142.pathview.png

## Info: some node width is different from others, and hence adjusted!

## 'select()' returned 1:1 mapping between keys and columns

## Info: Working in directory /Users/elizabethwest/Documents/Classes/Bioinformatics/R Files/bggn213/Clas

## Info: Writing image file hsa04330.pathview.png
```

```
data(go.sets.hs)
data(go.subs.hs)

# Focus on Biological Process subset of GO
gobpsets = go.sets.hs[go.subs.hs$BP]

gobpres = gage(foldchanges, gsets=gobpsets, same.dir=TRUE)

lapply(gobpres, head)
```

```
## $greater
##
##           p.geomean stat.mean      p.val
## GO:0007156 homophilic cell adhesion 8.519724e-05 3.824205 8.519724e-05
## GO:0002009 morphogenesis of an epithelium 1.396681e-04 3.653886 1.396681e-04
## GO:0048729 tissue morphogenesis 1.432451e-04 3.643242 1.432451e-04
## GO:0007610 behavior 2.195494e-04 3.530241 2.195494e-04
## GO:0060562 epithelial tube morphogenesis 5.932837e-04 3.261376 5.932837e-04
## GO:0035295 tube development 5.953254e-04 3.253665 5.953254e-04
##
##           q.val set.size      exp1
## GO:0007156 homophilic cell adhesion 0.1951953      113 8.519724e-05
## GO:0002009 morphogenesis of an epithelium 0.1951953      339 1.396681e-04
## GO:0048729 tissue morphogenesis 0.1951953      424 1.432451e-04
## GO:0007610 behavior 0.2243795      427 2.195494e-04
## GO:0060562 epithelial tube morphogenesis 0.3711390      257 5.932837e-04
## GO:0035295 tube development 0.3711390      391 5.953254e-04
##
## $less
##
##           p.geomean stat.mean      p.val
## GO:0048285 organelle fission 1.536227e-15 -8.063910 1.536227e-15
```

```
## G0:0000280 nuclear division 4.286961e-15 -7.939217 4.286961e-15
## G0:0007067 mitosis 4.286961e-15 -7.939217 4.286961e-15
## G0:0000087 M phase of mitotic cell cycle 1.169934e-14 -7.797496 1.169934e-14
## G0:0007059 chromosome segregation 2.028624e-11 -6.878340 2.028624e-11
## G0:0000236 mitotic prometaphase 1.729553e-10 -6.695966 1.729553e-10
## q.val set.size exp1
## G0:0048285 organelle fission 5.841698e-12 376 1.536227e-15
## G0:0000280 nuclear division 5.841698e-12 352 4.286961e-15
## G0:0007067 mitosis 5.841698e-12 352 4.286961e-15
## G0:0000087 M phase of mitotic cell cycle 1.195672e-11 362 1.169934e-14
## G0:0007059 chromosome segregation 1.658603e-08 142 2.028624e-11
## G0:0000236 mitotic prometaphase 1.178402e-07 84 1.729553e-10
##
## $stats
## stat.mean exp1
## G0:0007156 homophilic cell adhesion 3.824205 3.824205
## G0:0002009 morphogenesis of an epithelium 3.653886 3.653886
## G0:0048729 tissue morphogenesis 3.643242 3.643242
## G0:0007610 behavior 3.530241 3.530241
## G0:0060562 epithelial tube morphogenesis 3.261376 3.261376
## G0:0035295 tube development 3.253665 3.253665
```

```
sig_genes <- res[res$padj <= 0.05 & !is.na(res$padj), "symbol"]
print(paste("Total number of significant genes:", length(sig_genes)))
```

```
## [1] "Total number of significant genes: 8147"
```

```
write.table(sig_genes, file="significant_genes.txt", row.names=FALSE, col.names=FALSE, quote=FALSE)
```





