

lab13_deseq2

AUTHOR

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Section 1: Differential Expression Analysis

```
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

```
countFile <- "https://bioboot.github.io/bimm143_W18/class-material/counts/
metaFile <- "https://bioboot.github.io/bimm143_W18/class-material/metadata/
```

```
#Import metadata
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
```

```
# Import countdata
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 column from countData
```

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

```
head(countData)
```

```

                                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

```

```
# Filter count data where you have 0 read count across all samples
```

```
countData = countData[rowSums(countData[])>0,]
head(countData)
```

	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			

```
# Set up the DESeqDataSet object and run the DESeq pipeline
```

```
dds= DESeqDataSetFromMatrix(countData=countData, 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
```

```
resultsNames(dds)
```

```
[1] "Intercept"
"condition_hoxa1_kd_vs_control_sirna"
```

```
res=results(dds, contrast=c("condition", "hoxa1_kd", "control
_sirna"))
```

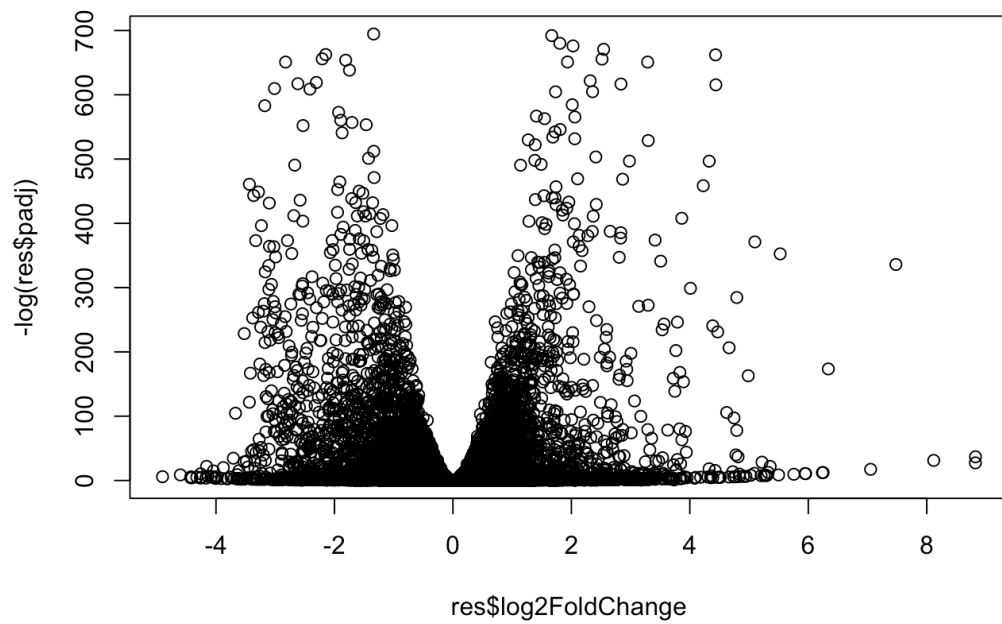
Call the summary() function to get a sense of how many genes

```
summary(res)
```

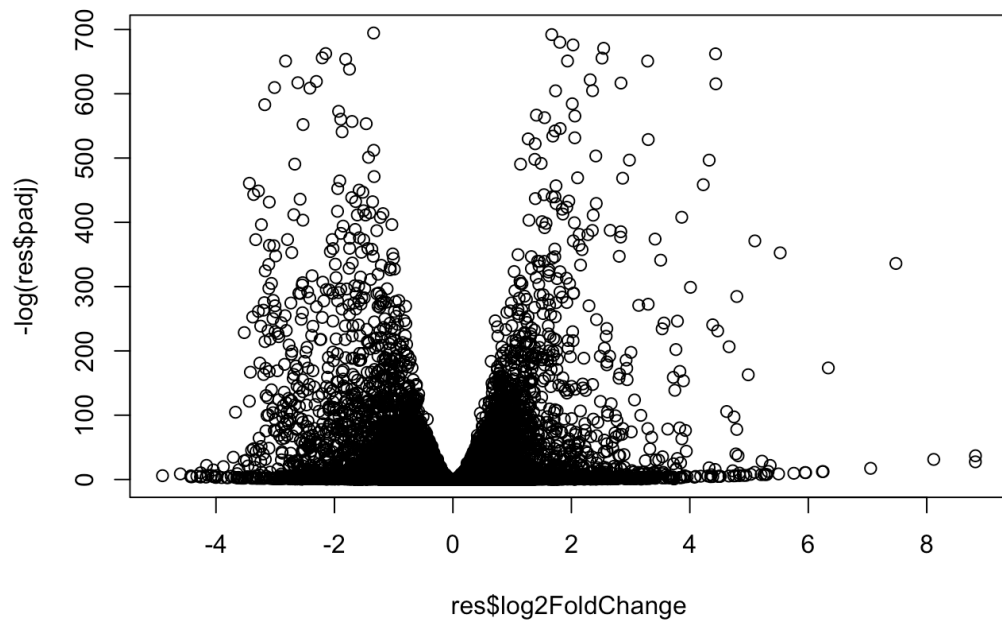
```
out of 15975 with nonzero total read count
adjusted p-value < 0.1
LFC > 0 (up)      : 4349, 27%
LFC < 0 (down)    : 4396, 28%
outliers [1]      : 0, 0%
low counts [2]    : 1237, 7.7%
(mean count < 0)
[1] see 'cooksCutoff' argument of ?results
[2] see 'independentFiltering' argument of ?results
```

Volcano plot

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



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



```
# Make a color vector for all genes
```

```

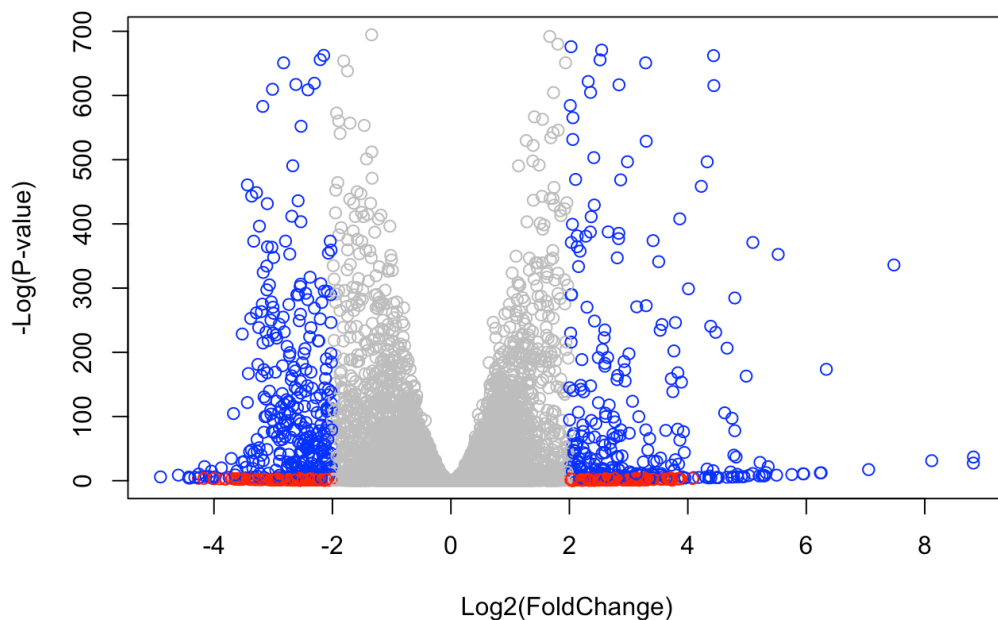
mycols <- rep("gray", nrow(res) )

# 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 <- (abs(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)" )

```



```
?mapIds()
```

```
# Use the mapIds() function to add SYMBOL, ENTREZID, and GENENAME
```

```

library("AnnotationDbi")
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	NA	NA	
NA				
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 ..

```

Order these results by adjusted p-value and save them to a CSV

```

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

```

Section 2: Pathway Analysis

```
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

```

```

# Get the results
keggres = gage(foldchanges, gsets=kegg.sets.hs)

```

```
?gage()
```

```

# Look at the first few down (less) pathways
head(keggres$less)

```

```

p.val      p.geomean stat.mean
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/cathy/BIMM 143/lab13

Info: Writing image file hsa04110.pathview.png

```
# A different PDF based output of the same data
pathview(gene.data=foldchanges, pathway.id="hsa04110", kegg.name="hsa04110")
```

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

Info: Working in directory /Users/cathy/BIMM 143/lab13

Info: Writing image file hsa04110.pathview.pdf

```
## Focus on top 5 upregulated pathways here for demo purposes
```

```
keggrespathways <- rownames(keggres$greater)[1:5]

# Extract the 8 character long IDs part of each string
keggresids = substr(keggrespathways, start=1, stop=8)
keggresids
```

```
[1] "hsa04640" "hsa04630" "hsa00140" "hsa04142" "hsa04330"
```

```
# Plots for all 5 pathways
pathview(gene.data=foldchanges, pathway.id=keggresids, species=
  "")
```

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

Info: Working in directory /Users/cathy/BIMM 143/lab13

Info: Writing image file hsa04640.pathview.png

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

Info: Working in directory /Users/cathy/BIMM 143/lab13

Info: Writing image file hsa04630.pathview.png

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

Info: Working in directory /Users/cathy/BIMM 143/lab13

Info: Writing image file hsa00140.pathview.png

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

Info: Working in directory /Users/cathy/BIMM 143/lab13

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/cathy/BIMM 143/lab13

Info: Writing image file hsa04330.pathview.png

Q. Can you do the same procedure as above to plot the pathway

```
## Focus on top 5 downregulated pathways here
keggrespathways <- rownames(keggres$less)[1:5]
```

```
# Extract the 8 character long IDs part of each string
keggresids = substr(keggrespathways, start=1, stop=8)
keggresids
```

```
[1] "hsa04110" "hsa03030" "hsa03013" "hsa03440" "hsa04114"
```

```
# Plots for all 5 pathways
pathview(gene.data=foldchanges, pathway.id=keggresids, species=
  "")
```

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

Info: Working in directory /Users/cathy/BIMM 143/lab13

Info: Writing image file hsa04110.pathview.png

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

Info: Working in directory /Users/cathy/BIMM 143/lab13

Info: Writing image file hsa03030.pathview.png

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

Info: Working in directory /Users/cathy/BIMM 143/lab13

Info: Writing image file hsa03013.pathview.png

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

Info: Working in directory /Users/cathy/BIMM 143/lab13

Info: Writing image file hsa03440.pathview.png

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

Info: Working in directory /Users/cathy/BIMM 143/lab13

Info: Writing image file hsa04114.pathview.png

Section 3. Gene Ontology

```
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

stat.mean	p.val	p.geomean
G0:0007156 homophilic cell adhesion	8.519724e-05	
3.824205 8.519724e-05		
G0:0002009 morphogenesis of an epithelium	1.396681e-04	
3.653886 1.396681e-04		
G0:0048729 tissue morphogenesis	1.432451e-04	
3.643242 1.432451e-04		
G0:0007610 behavior	2.195494e-04	
3.530241 2.195494e-04		
G0:0060562 epithelial tube morphogenesis	5.932837e-04	
3.261376 5.932837e-04		
G0:0035295 tube development	5.953254e-04	
3.253665 5.953254e-04		

exp1	q.val	set.size
G0:0007156 homophilic cell adhesion	0.1951953	113
8.519724e-05		
G0:0002009 morphogenesis of an epithelium	0.1951953	339
1.396681e-04		
G0:0048729 tissue morphogenesis	0.1951953	424
1.432451e-04		
G0:0007610 behavior	0.2243795	427
2.195494e-04		
G0:0060562 epithelial tube morphogenesis	0.3711390	257
5.932837e-04		
G0:0035295 tube development	0.3711390	391
5.953254e-04		

\$less

stat.mean	p.val	p.geomean
G0:0048285 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		

exp1	q.val	set.size
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

Section 4. Reactome Analysis

```
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=
```

```
sessionInfo()
```

R version 4.2.1 (2022-06-23)

Platform: x86_64-apple-darwin17.0 (64-bit)

Running under: macOS Catalina 10.15.7

Matrix products: default

BLAS:

/Library/Frameworks/R.framework/Versions/4.2/Resources/lib/lib
Rblas.0.dylib

LAPACK:

/Library/Frameworks/R.framework/Versions/4.2/Resources/lib/lib
Rlapack.dylib

locale:

[1] en_US.UTF-8/en_US.UTF-8/en_US.UTF-8/C/en_US.UTF-
8/en_US.UTF-8

attached base packages:

[1] stats4 stats graphics grDevices utils datasets
methods
[8] base

other attached packages:

[1] gageData_2.34.0 gage_2.46.1
[3] pathview_1.36.1 org.Hs.eg.db_3.15.0
[5] AnnotationDbi_1.58.0 DESeq2_1.36.0
[7] SummarizedExperiment_1.26.1 Biobase_2.56.0
[9] MatrixGenerics_1.8.1 matrixStats_0.63.0
[11] GenomicRanges_1.48.0 GenomeInfoDb_1.32.4
[13] IRanges_2.30.1 S4Vectors_0.34.0
[15] BiocGenerics_0.42.0

loaded via a namespace (and not attached):

[1] http_1.4.5 bit64_4.0.5
jsonlite_1.8.4
[4] splines_4.2.1 blob_1.2.3
GenomeInfoDbData_1.2.8

[7] yaml_2.3.7	pillar_1.8.1	
RSQLite_2.3.0		
[10] lattice_0.20-45	glue_1.6.2	
digest_0.6.31		
[13] RColorBrewer_1.1-3	XVector_0.36.0	
colorspace_2.1-0		
[16] htmltools_0.5.4	Matrix_1.5-3	XML_3.99-
0.13		
[19] pkgconfig_2.0.3	genefilter_1.78.0	
zlibbioc_1.42.0		
[22] GO.db_3.15.0	xtable_1.8-4	
scales_1.2.1		
[25] BiocParallel_1.30.4	tibble_3.1.8	
annotate_1.74.0		
[28] KEGGREST_1.36.3	generics_0.1.3	
ggplot2_3.4.1		
[31] cachem_1.0.7	cli_3.6.0	
survival_3.5-3		
[34] magrittr_2.0.3	crayon_1.5.2	
KEGGgraph_1.56.0		
[37] memoise_2.0.1	evaluate_0.20	fansi_1.0.4
[40] graph_1.74.0	tools_4.2.1	
lifecycle_1.0.3		
[43] munsell_0.5.0	locfit_1.5-9.7	
DelayedArray_0.22.0		
[46] Biostrings_2.64.1	compiler_4.2.1	rlang_1.0.6
[49] grid_4.2.1	RCurl_1.98-1.10	
rstudioapi_0.14		
[52] htmlwidgets_1.6.1	bitops_1.0-7	
rmarkdown_2.20		
[55] gtable_0.3.1	codetools_0.2-19	DBI_1.1.3
[58] R6_2.5.1	knitr_1.42	dplyr_1.1.0
[61] fastmap_1.1.1	bit_4.0.5	utf8_1.2.3
[64] Rgraphviz_2.40.0	parallel_4.2.1	Rcpp_1.0.10
[67] vctrs_0.5.2	geneplotter_1.74.0	png_0.1-8
[70] tidyselect_1.2.0	xfun_0.37	