

## Exercise 9

### Basic graphs

#### Exercise 9a – scatter plot

1. Read in file “/users/bi/public-docs/sbonnin/Rcourse/gene\_counts.txt” in object **genes**.

*Note: this file contains a header.*

2. Create a scatter plot showing **sample1** (x-axis) vs **sample2** (y-axis) of **genes**.  
**plot()**

3. Change the point type and colors.

*Note: see options **pch** and **col**.*

*example of **pch**:*

<b>0</b> □	<b>1</b> ○	<b>2</b> △	<b>3</b> +	<b>4</b> ×	
<b>5</b> ◇	<b>6</b> ▽	<b>7</b> ⊠	<b>8</b> ✱	<b>9</b> ⊞	
<b>10</b> ⊕	<b>11</b> ⊗	<b>12</b> ⊞	<b>13</b> ⊗	<b>14</b> ⊞	
<b>15</b> ■	<b>16</b> ●	<b>17</b> ▲	<b>18</b> ◆	<b>19</b> ●	
<b>20</b> ●	<b>21</b> ●	<b>22</b> ■	<b>23</b> ◆	<b>24</b> ▲	<b>25</b> ▼

*For **col**, you can use:*

- integers (1, 2, 3 etc.) but limited to 8 different colors!

- or pick up the name of a color using: **colors()**; for color picked randomly, you can use **sample(colors(), 1)**.

4. Change axis labels to “**Sample 1**” and “**Sample 2**”.

*Note: see options **xlab** and **ylab**.*

5. Add a title to the plot.

*Note: see option **main**.*

6. Add a **red** vertical line at the **median** expression value of sample 1.

a. calculate the median expression of genes in sample 1.

**median()**

b. plot a vertical line using **abline()**.

*Note: **plot(...)** must be called before **abline()** is called. **abline** is not an argument but a function!*

*See **abline** help page.*

#### Exercise 9b – bar plot + pie chart

1. Read in file “/nfs/users/bi/public-docs/sbonnin/Rcourse/genes\_counts\_significance.txt” in object **de**.

*Note: this file contains a header.*

2. The column **updown** specifies whether a gene is up- or down- regulated, or neither of them (none).

Produce a **barplot** displaying this information: how many genes are up- or down-regulated, or neither of them.

**barplot(table())**

3. Color the bars of the boxplot, each in a different color: up in red, down in blue, none in grey.

4. Produce the same plot, but this time set argument **space** to 0.

What is **space** used for?

5. Now use option **names.arg** in **barplot()** to rename the bars.

6. The **las** argument allows you to rotate labels for a better visibility.

Try it!

*Note: try values 0, 1, 2, 3. Observe how labels on both axis are oriented.*

4. Create a pie chart of the same information (up, down, none)

**pie(table())**

*Note: Try arguments **color**, **main** and **labels**.*

### **Exercise 9c – histogram**

1. Using **genes** object from exercise 9a, create a **histogram** of the gene expression distribution of sample 1.

**hist()**

2. Repeat the histogram but change argument **breaks** to 50.

What is the difference?

3. Color this histogram in light blue.

*Note: there is color called “lightblue”*

4. Overlap a second histogram to that first one, corresponding to sample 2 expression values.

*Note: you should call **hist()** once for sample 1, once for sample 2. The second histogram called should have option **add** set to **TRUE**. Read help page of **hist**.*

5. “Zoom” in the histogram: show only the distribution of expression values from 7 to 12 (x-axis).

*Note: use **xlim** option. Adjust also **ylim** if necessary for a better visibility.*

6. Save plot in a pdf file.

a. Try with RStudio Plots window (Export)

b. Try in the console:

```
pdf(...)  
hist(...)  
dev.off()
```

## **Exercise 10.**

### **Introduction to ggplot2.**

#### **Exercise 10a – scatter plot**

1. Load **ggplot2** package.

2. Using **de** object from Exercise 9, create a simple scatter plot for plotting gene expression of sample 1 and sample 2.

*Note: remember the structure:*

```
ggplot(dataframe, aes(x=, y=)) + geom_point()
```

3. Color points according to the **updown** column. Save in object **p**.

*Note: remember the structure:*

```
p <- ggplot(dataframe, aes(x=, y=, color=)) + geom_point()
```

4. Change colors of the points to blue, grey and red. Save to p2.

```
p2 <- p + scale_color_manual()
```

5. Save p2 into a jpeg file.

a. Try with RStudio Plots window (Export)

b. Try in the console:

```
jpeg(...)  
plot(...)  
dev.off()
```

#### **Exercise 10b – box plot**

1. Convert **de** from a wide format to a long format, save in **de\_long**.

*Note: remember **melt** function from **reshape2** package.*

```
de_long <- melt(de)
```

2. Produce a boxplot of the expression of sample 1 and sample 2 (each sample should be represented by a box)

```
ggplot(dataframe, aes(x, y)) + geom_boxplot()
```

3. Modify the previous boxplot so as to obtain 3 “sub”-boxplots per sample, each representing the expression of either UP, DOWN or NONE genes.

```
ggplot(dataframe, aes(x, y, col)) + geom_boxplot()
```

#### **Exercise 10c – bar plot**

1. Produce a bar plot of how many UP/DOWN/NONE genes are in **de**.

```
ggplot(de, aes(x=)) + geom_bar()
```

2. Add an horizontal line at counts **1000**.  
**geom\_hline()**

3. Swap x and y axis.  
**coord\_flip()**

4. Add a title to the graph.  
**ggtitle()**

### **Exercise 10d - histogram**

1. Create a simple histogram using **de\_long**.  
*ggplot(de\_long, aes(x=)) + geom\_histogram()*

2. Notice that you get the following warning message “*stat\_bin() using `bins = 30`. Pick better value with `binwidth`.*”  
Change **bins** parameter in **geom\_histogram()** to **50**.

3. This histogram plots expression values for both sample1 and sample2.  
Change the plot so as to obtain 2 histograms on the same plot: one corresponding to sample1, one corresponding to sample2.  
*ggplot(...fill=) + geom\_histogram()*

4. By default, geom\_histogram produces a stacked histogram.  
Change the position of the bars to **dodge**.  
*ggplot(...fill=) + geom\_histogram(position=...)*

5. Zoom in the plot: reduce the x-axis to values from 7 to 12.  
**xlim()** layer.

6. Finally, change the colors of the bars to colors of your choice.  
**scale\_fill\_manual()**  
*Note: Try the rainbow() function for coloring!*