

Graphics and Data Visualization in R

Overview

Thomas Girke

December 13, 2013

Overview

Graphics Environments

Base Graphics

Grid Graphics

lattice

ggplot2

Specialty Graphics

Genome Graphics

ggbio

Additional Genome Graphics

Clustering

Background

Hierarchical Clustering Example

Non-Hierarchical Clustering Examples

Outline

Overview

Graphics Environments

- Base Graphics
- Grid Graphics
- lattice
- ggplot2

Specialty Graphics

Genome Graphics

- ggbio
- Additional Genome Graphics

Clustering

- Background
- Hierarchical Clustering Example
- Non-Hierarchical Clustering Examples

Graphics in R

- Powerful environment for visualizing scientific data
- Integrated graphics and statistics infrastructure
- Publication quality graphics
- Fully programmable
- Highly reproducible
- Full \LaTeX [Link](#) & Sweave [Link](#) support
- Vast number of R packages with graphics utilities

Documentation on Graphics in R

General

- Graphics Task Page [Link](#)
- R Graph Gallery [Link](#)
- R Graphical Manual [Link](#)
- Paul Murrell's book R (Grid) Graphics [Link](#)

Interactive graphics

- rggobi (GGobi) [Link](#)
- ipplots [Link](#)
- Open GL (rgl) [Link](#)

Graphics Environments

Viewing and saving graphics in R

- On-screen graphics
- postscript, pdf, svg
- jpeg/png/wmf/tiff/...

Four major graphic environments

- Low-level infrastructure
 - R Base Graphics (low- and high-level)
 - *grid*: Manual [Link](#), Book [Link](#)
- High-level infrastructure
 - *lattice*: Manual [Link](#), Intro [Link](#), Book [Link](#)
 - *ggplot2*: Manual [Link](#), Intro [Link](#), Book [Link](#)

Outline

Overview

Graphics Environments

- Base Graphics
- Grid Graphics
- lattice*
- ggplot2*

Specialty Graphics

Genome Graphics

- ggbio*
- Additional Genome Graphics

Clustering

- Background
- Hierarchical Clustering Example
- Non-Hierarchical Clustering Examples

Outline

Overview

Graphics Environments

Base Graphics

Grid Graphics

lattice

ggplot2

Specialty Graphics

Genome Graphics

ggbio

Additional Genome Graphics

Clustering

Background

Hierarchical Clustering Example

Non-Hierarchical Clustering Examples

Base Graphics: Overview

Important high-level plotting functions

- `plot`: generic x-y plotting
- `barplot`: bar plots
- `boxplot`: box-and-whisker plot
- `hist`: histograms
- `pie`: pie charts
- `dotchart`: cleveland dot plots
- `image`, `heatmap`, `contour`, `persp`: functions to generate image-like plots
- `qqnorm`, `qqline`, `qqplot`: distribution comparison plots
- `pairs`, `coplot`: display of multivariate data

Help on these functions

- `?myfct`
- `?plot`
- `?par`

Base Graphics: Preferred Input Data Objects

- Matrices and data frames
- Vectors
- Named vectors

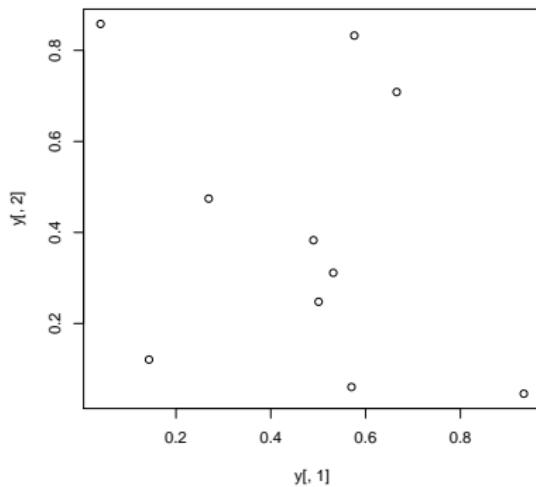
Scatter Plot: very basic

Sample data set for subsequent plots

```
> set.seed(1410)
> y <- matrix(runif(30), ncol=3, dimnames=list(letters[1:10], LETTERS[1:3]))
```

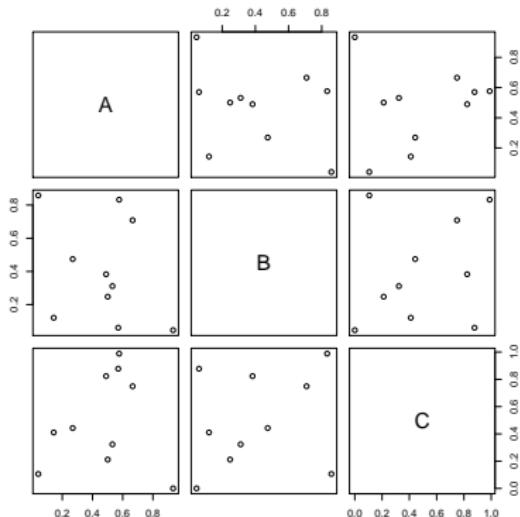


```
> plot(y[,1], y[,2])
```



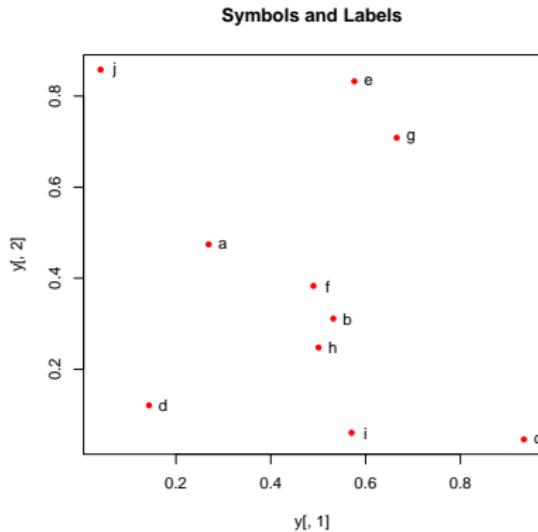
Scatter Plot: all pairs

```
> pairs(y)
```



Scatter Plot: with labels

```
> plot(y[,1], y[,2], pch=20, col="red", main="Symbols and Labels")
> text(y[,1]+0.03, y[,2], rownames(y))
```



Scatter Plots: more examples

Print instead of symbols the row names

```
> plot(y[,1], y[,2], type="n", main="Plot of Labels")
> text(y[,1], y[,2], rownames(y))
```

Usage of important plotting parameters

```
> grid(5, 5, lwd = 2)
> op <- par(mar=c(8,8,8,8), bg="lightblue")
> plot(y[,1], y[,2], type="p", col="red", cex.lab=1.2, cex.axis=1.2,
+       cex.main=1.2, cex.sub=1, lwd=4, pch=20, xlab="x label",
+       ylab="y label", main="My Main", sub="My Sub")
> par(op)
```

Important arguments

- `mar`: specifies the margin sizes around the plotting area in order: `c(bottom, left, top, right)`
- `col`: color of symbols
- `pch`: type of symbols, samples: `example(points)`
- `lwd`: size of symbols
- `cex.*`: control font sizes
- For details see `?par`

Scatter Plots: more examples

Add a regression line to a plot

```
> plot(y[,1], y[,2])
> myline <- lm(y[,2]~y[,1]); abline(myline, lwd=2)
> summary(myline)
```

Same plot as above, but on log scale

```
> plot(y[,1], y[,2], log="xy")
```

Add a mathematical expression to a plot

```
> plot(y[,1], y[,2]); text(y[1,1], y[1,2],
>     expression(sum(frac(1,sqrt(x^2*pi)))), cex=1.3)
```

Exercise 1: Scatter Plots

- Task 1** Generate scatter plot for first two columns in `iris` data frame and color dots by its `Species` column.
- Task 2** Use the `xlim`/`ylim` arguments to set limits on the x- and y-axes so that all data points are restricted to the left bottom quadrant of the plot.

Structure of `iris` data set:

```
> class(iris)
[1] "data.frame"

> iris[1:4,]

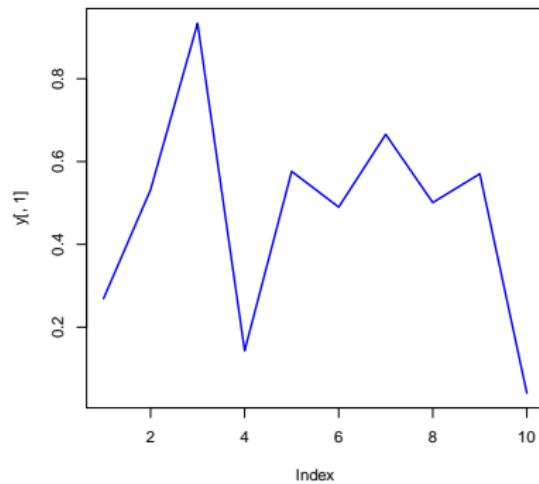
  Sepal.Length Sepal.Width Petal.Length Petal.Width Species
1          5.1       3.5        1.4       0.2   setosa
2          4.9       3.0        1.4       0.2   setosa
3          4.7       3.2        1.3       0.2   setosa
4          4.6       3.1        1.5       0.2   setosa

> table(iris$Species)

setosa versicolor virginica
      50         50         50
```

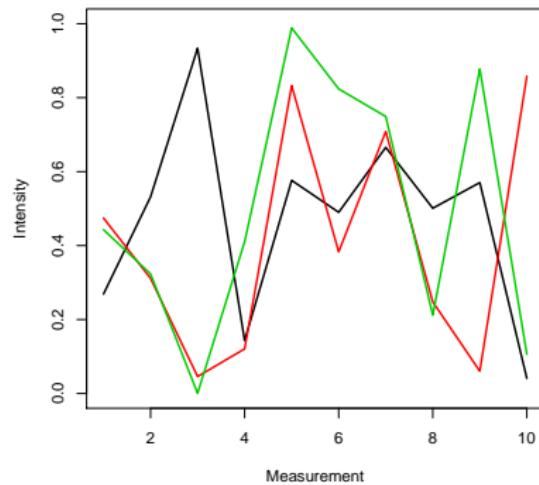
Line Plot: Single Data Set

```
> plot(y[,1], type="l", lwd=2, col="blue")
```



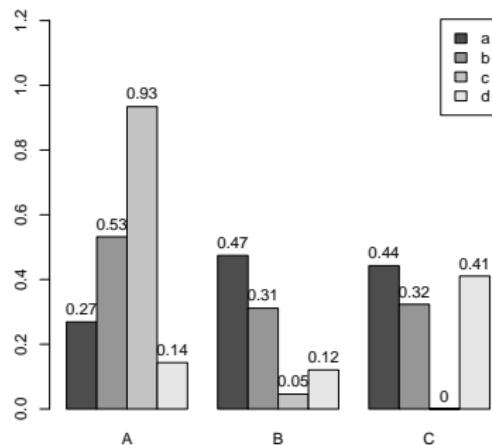
Line Plots: Many Data Sets

```
> split.screen(c(1,1));
[1] 1
> plot(y[,1], ylim=c(0,1), xlab="Measurement", ylab="Intensity", type="l", lwd=2, col=1)
> for(i in 2:length(y[,])) {
+   screen(1, new=FALSE)
+   plot(y[,i], ylim=c(0,1), type="l", lwd=2, col=i, xaxt="n", yaxt="n", ylab="",
+       xlab="", main="", bty="n")
+ }
> close.screen(all=TRUE)
```



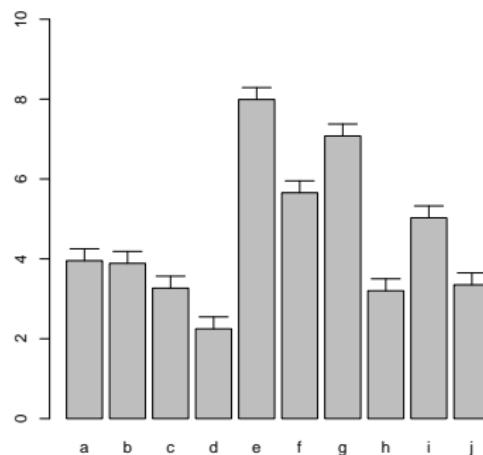
Bar Plot Basics

```
> barplot(y[1:4,], ylim=c(0, max(y[1:4,])+0.3), beside=TRUE,  
+           legend=letters[1:4])  
> text(labels=round(as.vector(as.matrix(y[1:4,])),2), x=seq(1.5, 13, by=1)  
+       +sort(rep(c(0,1,2), 4)), y=as.vector(as.matrix(y[1:4,]))+0.04)
```



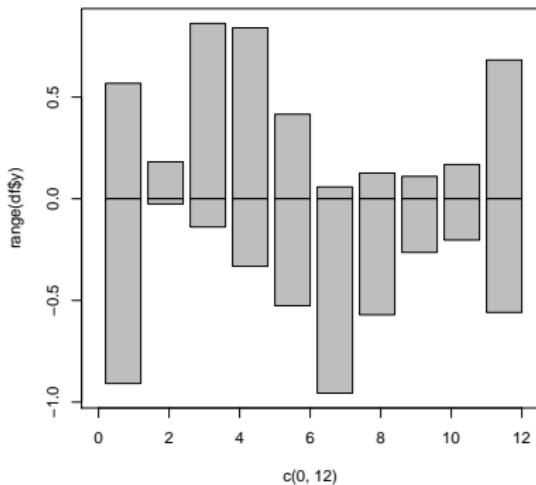
Bar Plots with Error Bars

```
> bar <- barplot(m <- rowMeans(y) * 10, ylim=c(0, 10))
> stdev <- sd(t(y))
> arrows(bar, m, bar, m + stdev, length=0.15, angle = 90)
```



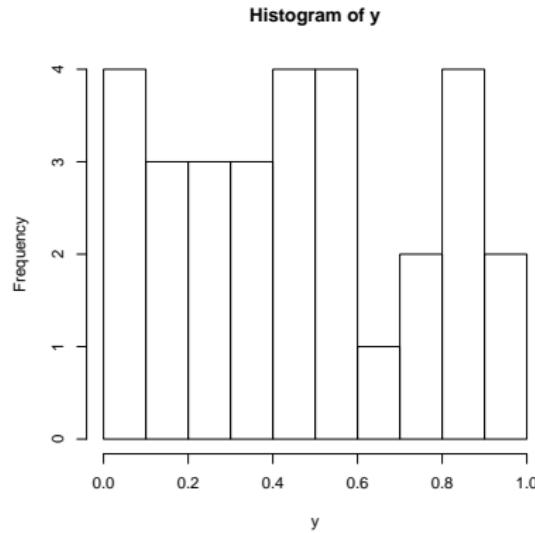
Mirrored Bar Plots

```
> df <- data.frame(group = rep(c("Above", "Below"), each=10), x = rep(1:10, 2),  
> plot(c(0,12),range(df$y),type = "n")  
> barplot(height = df$y[df$group == 'Above'], add = TRUE,axes = FALSE)  
> barplot(height = df$y[df$group == 'Below'], add = TRUE,axes = FALSE)
```



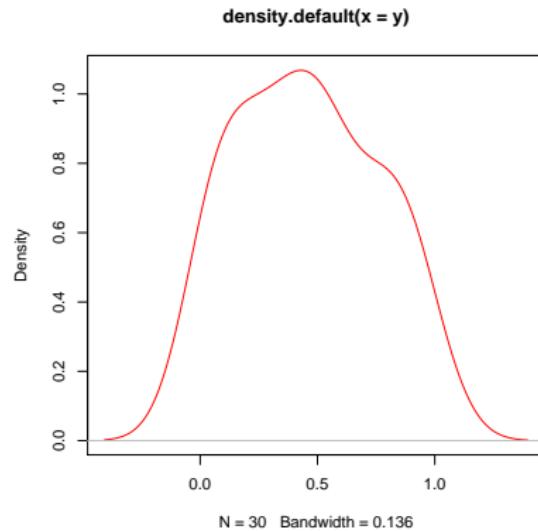
Histograms

```
> hist(y, freq=TRUE, breaks=10)
```



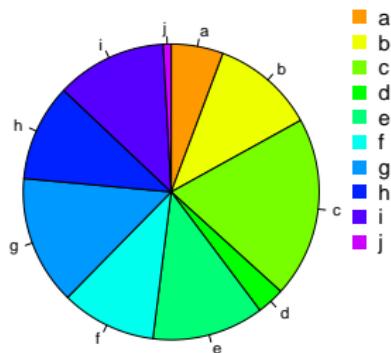
Density Plots

```
> plot(density(y), col="red")
```



Pie Charts

```
> pie(y[,1], col=rainbow(length(y[,1])), start=0.1, end=0.8, clockwise=TRUE)
> legend("topright", legend=row.names(y), cex=1.3, bty="n", pch=15, pt.cex=1.8,
+ col=rainbow(length(y[,1])), start=0.1, end=0.8), ncol=1)
```



Color Selection Utilities

Default color palette and how to change it

```
> palette()  
[1] "black"    "red"      "green3"    "blue"     "cyan"     "magenta"  "yellow"   "gray"  
  
> palette(rainbow(5, start=0.1, end=0.2))  
> palette()  
[1] "#FF9900"  "#FFBF00"  "#FFE600"  "#F2FF00"  "#CCFF00"  
  
> palette("default")
```

The gray function allows to select any type of gray shades by providing values from 0 to 1

```
> gray(seq(0.1, 1, by= 0.2))  
[1] "#1A1A1A"  "#4D4D4D"  "#808080"  "#B3B3B3"  "#E6E6E6"
```

Color gradients with `colorpanel` function from `gplots` library

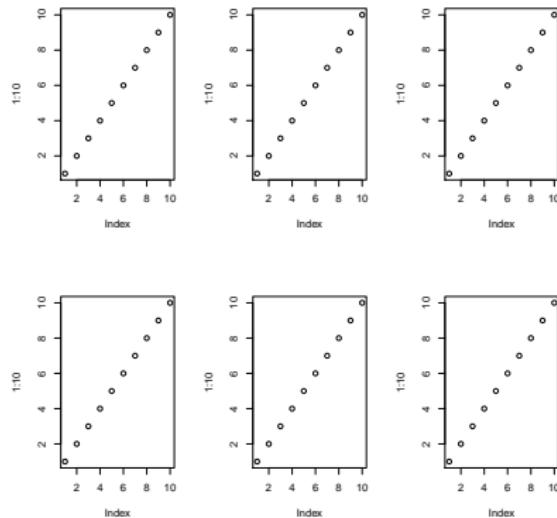
```
> library(gplots)  
> colorpanel(5, "darkblue", "yellow", "white")
```

Much more on colors in R see Earl Glynn's color chart [Link](#)

Arranging Several Plots on Single Page

With `par(mfrow=c(nrow,ncol))` one can define how several plots are arranged next to each other.

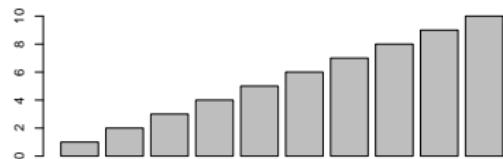
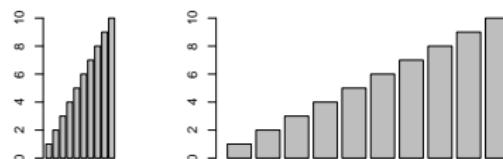
```
> par(mfrow=c(2,3)); for(i in 1:6) { plot(1:10) }
```



Arranging Plots with Variable Width

The `layout` function allows to divide the plotting device into variable numbers of rows and columns with the column-widths and the row-heights specified in the respective arguments.

```
> nf <- layout(matrix(c(1,2,3,3), 2, 2, byrow=TRUE), c(3,7), c(5,5),  
+                 respect=TRUE)  
> # layout.show(nf)  
> for(i in 1:3) { barplot(1:10) }
```



Saving Graphics to Files

After the `pdf()` command all graphs are redirected to file `test.pdf`. Works for all common formats similarly: jpeg, png, ps, tiff, ...

```
> pdf("test.pdf"); plot(1:10, 1:10); dev.off()
```

Generates Scalable Vector Graphics (SVG) files that can be edited in vector graphics programs, such as Inkscape.

```
> svg("test.svg"); plot(1:10, 1:10); dev.off()
```

Exercise 2: Bar Plots

- Task 1** Calculate the mean values for the Species components of the first four columns in the iris data set. Organize the results in a matrix where the row names are the unique values from the iris Species column and the column names are the same as in the first four iris columns.
- Task 2** Generate two bar plots: one with stacked bars and one with horizontally arranged bars.

Structure of iris data set:

```
> class(iris)
[1] "data.frame"
> iris[1:4,]
  Sepal.Length Sepal.Width Petal.Length Petal.Width Species
1          5.1       3.5        1.4       0.2   setosa
2          4.9       3.0        1.4       0.2   setosa
3          4.7       3.2        1.3       0.2   setosa
4          4.6       3.1        1.5       0.2   setosa
> table(iris$Species)
setosa versicolor virginica
      50           50          50
```

Outline

Overview

Graphics Environments

Base Graphics

Grid Graphics

`lattice`

`ggplot2`

Specialty Graphics

Genome Graphics

`ggbio`

Additional Genome Graphics

Clustering

Background

Hierarchical Clustering Example

Non-Hierarchical Clustering Examples

grid Graphics Environment

- What is *grid*?
 - Low-level graphics system
 - Highly flexible and controllable system
 - Does not provide high-level functions
 - Intended as development environment for custom plotting functions
 - Pre-installed on new R distributions
- Documentation and Help
 - Manual [Link](#)
 - Book [Link](#)

Outline

Overview

Graphics Environments

Base Graphics

Grid Graphics

`lattice`

`ggplot2`

Specialty Graphics

Genome Graphics

`ggbio`

Additional Genome Graphics

Clustering

Background

Hierarchical Clustering Example

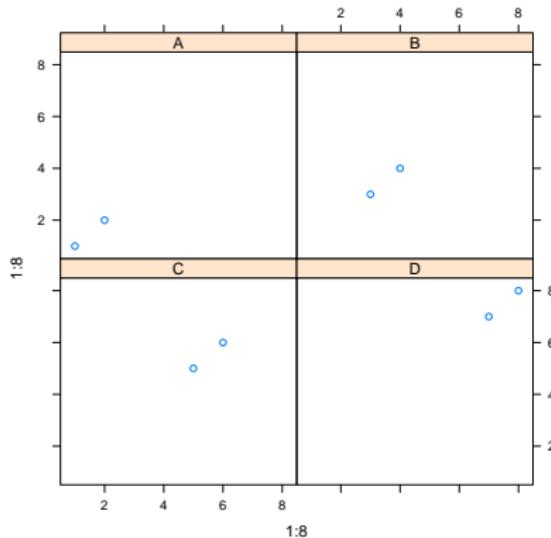
Non-Hierarchical Clustering Examples

lattice Environment

- What is *lattice*?
 - High-level graphics system
 - Developed by Deepayan Sarkar
 - Implements Trellis graphics system from S-Plus
 - Simplifies high-level plotting tasks: arranging complex graphical features
 - Syntax similar to R's base graphics
- Documentation and Help
 - Manual [Link](#)
 - Intro [Link](#)
 - Book [Link](#)
 - `library(help=lattice)` opens a list of all functions available in the lattice package
 - Accessing and changing global parameters:
`?lattice.options` and `?trellis.device`

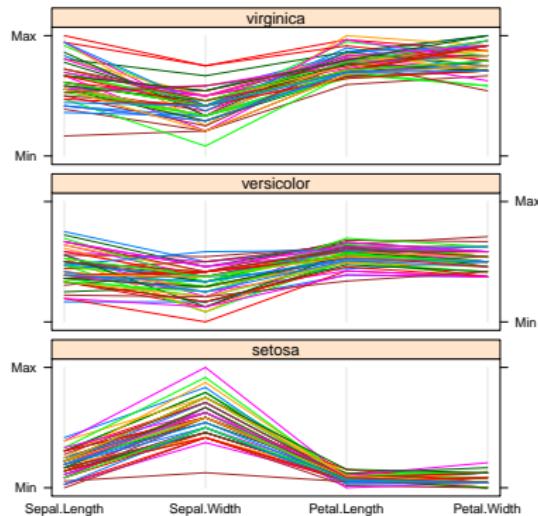
Scatter Plot Sample

```
> library(lattice)
> p1 <- xyplot(1:8 ~ 1:8 | rep(LETTERS[1:4], each=2), as.table=TRUE)
> plot(p1)
```



Line Plot Sample

```
> library(lattice)
> p2 <- parallelplot(~iris[1:4] | Species, iris, horizontal.axis = FALSE,
+                      layout = c(1, 3, 1))
> plot(p2)
```



Outline

Overview

Graphics Environments

Base Graphics

Grid Graphics

lattice

ggplot2

Specialty Graphics

Genome Graphics

ggbio

Additional Genome Graphics

Clustering

Background

Hierarchical Clustering Example

Non-Hierarchical Clustering Examples

ggplot2 Environment

- What is *ggplot2*?
 - High-level graphics system
 - Implements grammar of graphics from Leland Wilkinson [Link](#)
 - Streamlines many graphics workflows for complex plots
 - Syntax centered around main `ggplot` function
 - Simpler `qplot` function provides many shortcuts
- Documentation and Help
 - Manual [Link](#)
 - Intro [Link](#)
 - Book [Link](#)
 - Cookbook for R [Link](#)

ggplot2 Usage

- ggplot function accepts two arguments
 - Data set to be plotted
 - Aesthetic mappings provided by aes function
- Additional parameters such as geometric objects (e.g. points, lines, bars) are passed on by appending them with + as separator.
- List of available geom_* functions: [Link](#)
- Settings of plotting theme can be accessed with the command theme_get() and its settings can be changed with theme().
- Preferred input data object
 - qgplot: data.frame (support for vector, matrix, ...)
 - ggplot: data.frame
- Packages with convenience utilities to create expected inputs
 - *plyr*
 - *reshape*

qplot Function

- qplot syntax is similar to R's basic plot function
- Arguments:
 - x: x-coordinates (e.g. col1)
 - y: y-coordinates (e.g. col2)
 - data: data frame with corresponding column names
 - xlim, ylim: e.g. xlim=c(0,10)
 - log: e.g. log="x" or log="xy"
 - main: main title; see ?plotmath for mathematical formula
 - xlab, ylab: labels for the x- and y-axes
 - color, shape, size
 - ...: many arguments accepted by plot function

qplot: Scatter Plots

Create sample data

```
> library(ggplot2)
> x <- sample(1:10, 10); y <- sample(1:10, 10); cat <- rep(c("A", "B"), 5)
```

Simple scatter plot

```
> qplot(x, y, geom="point")
```

Prints dots with different sizes and colors

```
> qplot(x, y, geom="point", size=x, color=cat,
+         main="Dot Size and Color Relative to Some Values")
```

Drops legend

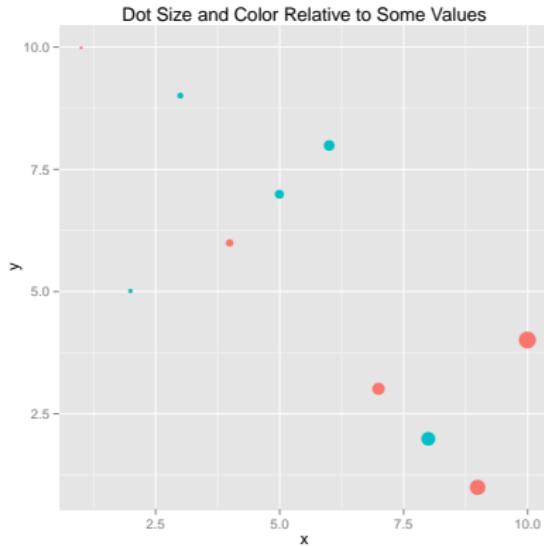
```
> qplot(x, y, geom="point", size=x, color=cat) +
+     theme(legend.position = "none")
```

Plot different shapes

```
> qplot(x, y, geom="point", size=5, shape=cat)
```

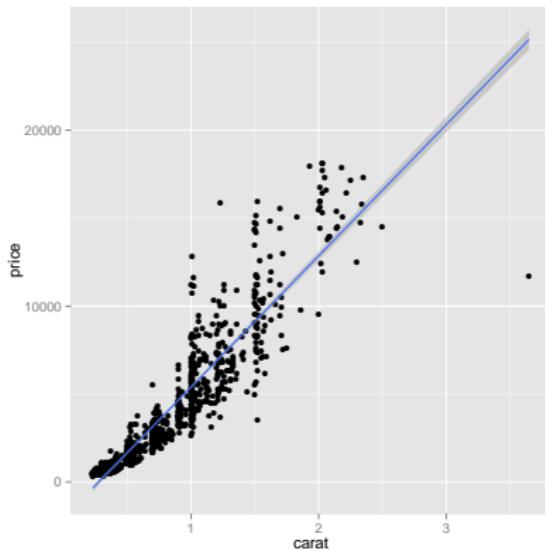
qplot: Scatter Plot with qplot

```
> p <- qplot(x, y, geom="point", size=x, color=cat,
+             main="Dot Size and Color Relative to Some Values") +
+     theme(legend.position = "none")
> print(p)
```



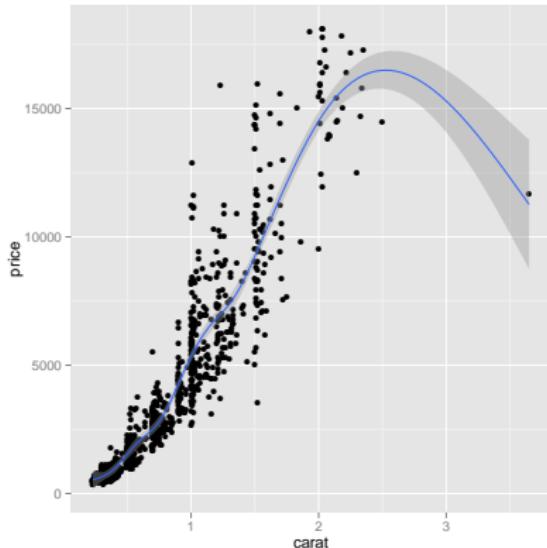
qplot: Scatter Plot with Regression Line

```
> set.seed(1410)
> dsmall <- diamonds[sample(nrow(diamonds), 1000), ]
> p <- qplot(carat, price, data = dsmall, geom = c("point", "smooth"),
+             method = "lm")
> print(p)
```



qplot: Scatter Plot with Local Regression Curve (loess)

```
> p <- qplot(carat, price, data=dsmall, geom=c("point", "smooth"), span=0.4)
> print(p) # Setting 'se=FALSE' removes error shade
```



ggplot Function

- More important than qplot to access full functionality of *ggplot2*
- Main arguments
 - data set, usually a `data.frame`
 - aesthetic mappings provided by `aes` function
- General ggplot syntax
 - `ggplot(data, aes(...)) + geom_*(...) + ... + stat_*(...) + ...`
- Layer specifications
 - `geom_*(mapping, data, ..., geom, position)`
 - `stat_*(mapping, data, ..., stat, position)`
- Additional components
 - `scales`
 - `coordinates`
 - `facet`
- `aes()` mappings can be passed on to all components (`ggplot`, `geom_*`, etc.).
Effects are global when passed on to `ggplot()` and local for other components.
 - `x`, `y`
 - `color`: grouping vector (`factor`)
 - `group`: grouping vector (`factor`)

Changing Plotting Themes with ggplot

- Theme settings can be accessed with `theme_get()`
- Their settings can be changed with `theme()`
- Some examples
 - Change background color to white

```
... + theme(panel.background=element_rect(fill = "white", colour = "black"))
```

Storing ggplot Specifications

Plots and layers can be stored in variables

```
> p <- ggplot(dsmall, aes(carat, price)) + geom_point()  
> p # or print(p)
```

Returns information about data and aesthetic mappings followed by each layer

```
> summary(p)
```

Prints dots with different sizes and colors

```
> bestfit <- geom_smooth(methodw = "lm", se = F, color = alpha("steelblue", 0.5)  
> p + bestfit # Plot with custom regression line
```

Syntax to pass on other data sets

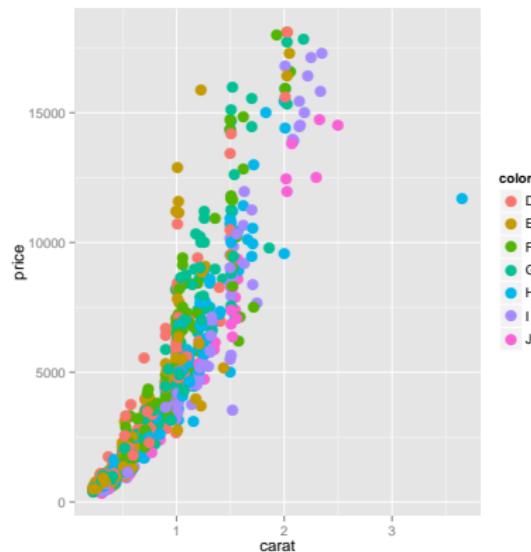
```
> p %>% diamonds[sample(nrow(diamonds), 100),]
```

Saves plot stored in variable p to file

```
> ggsave(p, file="myplot.pdf")
```

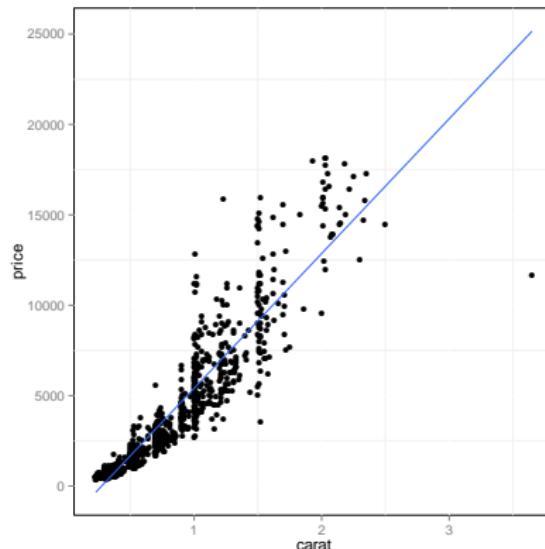
ggplot: Scatter Plot

```
> p <- ggplot(dsmall, aes(carat, price, color=color)) +  
+         geom_point(size=4)  
> print(p)
```



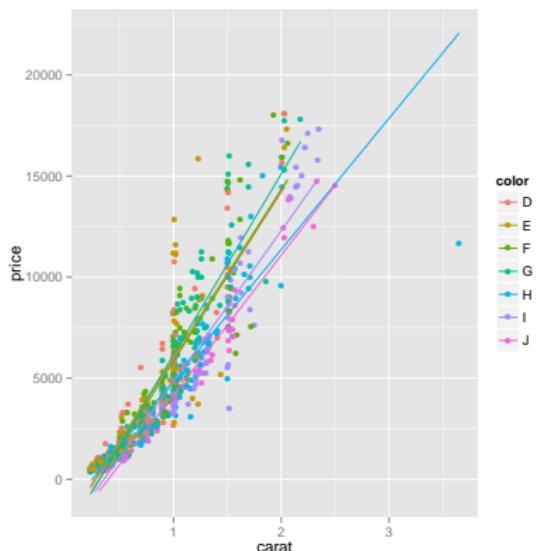
ggplot: Scatter Plot with Regression Line

```
> p <- ggplot(dsmall, aes(carat, price)) + geom_point() +  
+         geom_smooth(method="lm", se=FALSE) +  
+         theme(panel.background=element_rect(fill = "white", colour = "black"))  
> print(p)
```



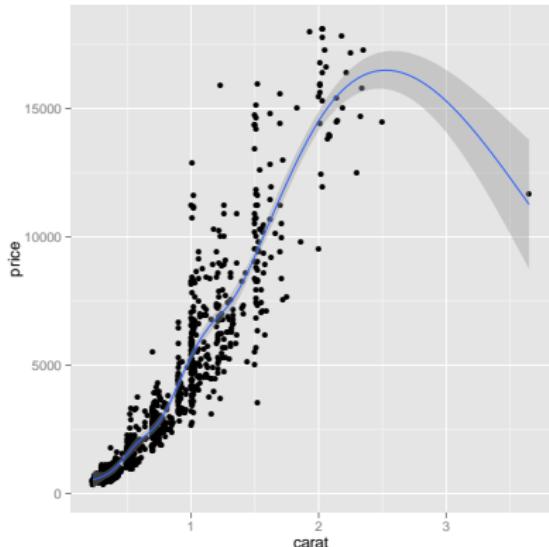
ggplot: Scatter Plot with Several Regression Lines

```
> p <- ggplot(dsmall, aes(carat, price, group=color)) +  
+     geom_point(aes(color=color), size=2) +  
+     geom_smooth(aes(color=color), method = "lm", se=FALSE)  
> print(p)
```



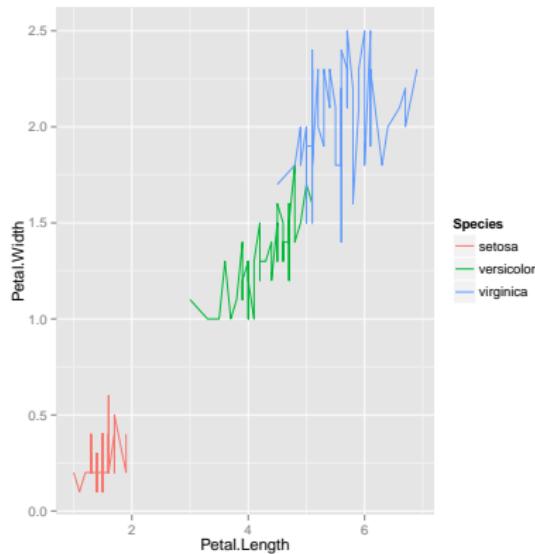
ggplot: Scatter Plot with Local Regression Curve (loess)

```
> p <- ggplot(dsmall, aes(carat, price)) + geom_point() + geom_smooth()  
> print(p) # Setting 'se=FALSE' removes error shade
```



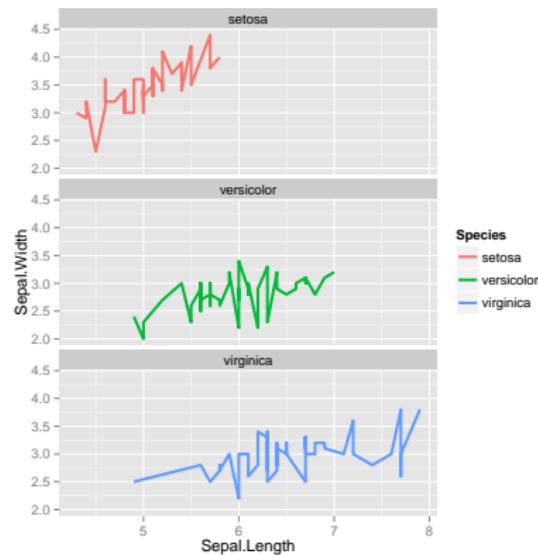
ggplot: Line Plot

```
> p <- ggplot(iris, aes(Petal.Length, Petal.Width, group=Species,  
+ color=Species)) + geom_line()  
> print(p)
```



ggplot: Faceting

```
> p <- ggplot(iris, aes(Sepal.Length, Sepal.Width)) +  
+     geom_line(aes(color=Species), size=1) +  
+     facet_wrap(~Species, ncol=1)  
> print(p)
```



Exercise 3: Scatter Plots

- Task 1** Generate scatter plot for first two columns in `iris` data frame and color dots by its `Species` column.
- Task 2** Use the `xlim`, `ylim` functions to set limits on the x- and y-axes so that all data points are restricted to the left bottom quadrant of the plot.
- Task 3** Generate corresponding line plot with faceting show individual data sets in separate plots.

Structure of `iris` data set:

```
> class(iris)
[1] "data.frame"
> iris[1:4,]
  Sepal.Length Sepal.Width Petal.Length Petal.Width Species
1          5.1       3.5        1.4       0.2   setosa
2          4.9       3.0        1.4       0.2   setosa
3          4.7       3.2        1.3       0.2   setosa
4          4.6       3.1        1.5       0.2   setosa
> table(iris$Species)
setosa versicolor virginica
      50         50         50
```

ggplot: Bar Plots

Sample Set: the following transforms the iris data set into a ggplot2-friendly format.

Calculate mean values for aggregates given by Species column in iris data set

```
> iris_mean <- aggregate(iris[,1:4], by=list(Species=iris$Species), FUN=mean)
```

Calculate standard deviations for aggregates given by Species column in iris data set

```
> iris_sd <- aggregate(iris[,1:4], by=list(Species=iris$Species), FUN=sd)
```

Convert iris_mean with melt

```
> library(reshape2) # Defines melt function
```

```
> df_mean <- melt(iris_mean, id.vars=c("Species"), variable.name = "Samples", value.name = "Mean")
```

Convert iris_sd with melt

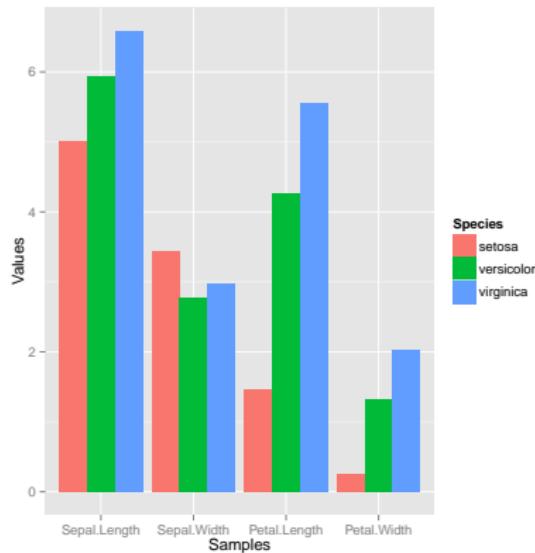
```
> df_sd <- melt(iris_sd, id.vars=c("Species"), variable.name = "Samples", value.name = "SD")
```

Define standard deviation limits

```
> limits <- aes(ymax = df_mean[, "Values"] + df_sd[, "Values"], ymin=df_mean[, "Values"] - df_sd[, "Values"])
```

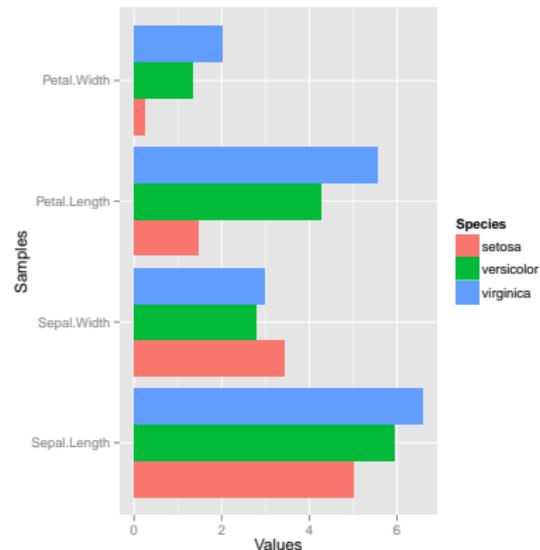
ggplot: Bar Plot

```
> p <- ggplot(df_mean, aes(Samples, Values, fill = Species)) +  
+         geom_bar(position="dodge", stat="identity")  
> print(p)
```



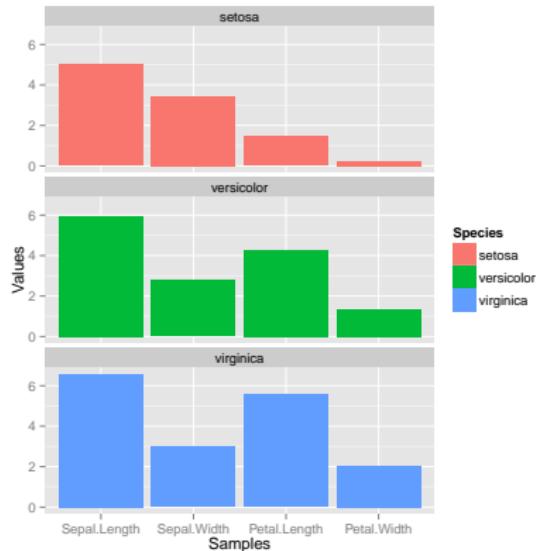
ggplot: Bar Plot Sideways

```
> p <- ggplot(df_mean, aes(Samples, Values, fill = Species)) +  
+     geom_bar(position="dodge", stat="identity") + coord_flip() +  
+     theme(axis.text.y=theme_text(angle=0, hjust=1))  
> print(p)
```



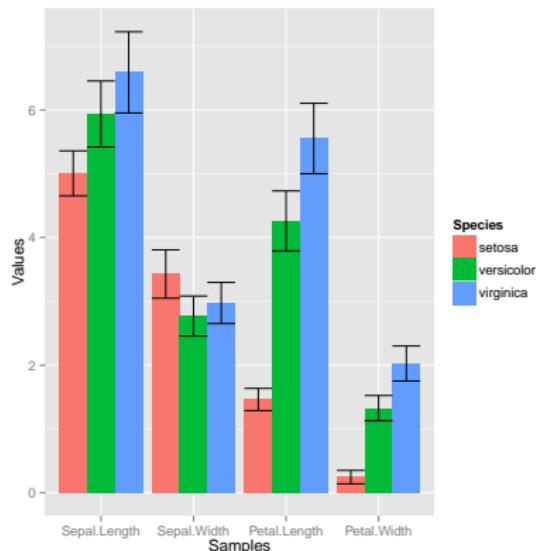
ggplot: Bar Plot with Faceting

```
> p <- ggplot(df_mean, aes(Samples, Values)) + geom_bar(aes(fill = Species), stat="sum")  
+ facet_wrap(~Species, ncol=1)  
> print(p)
```



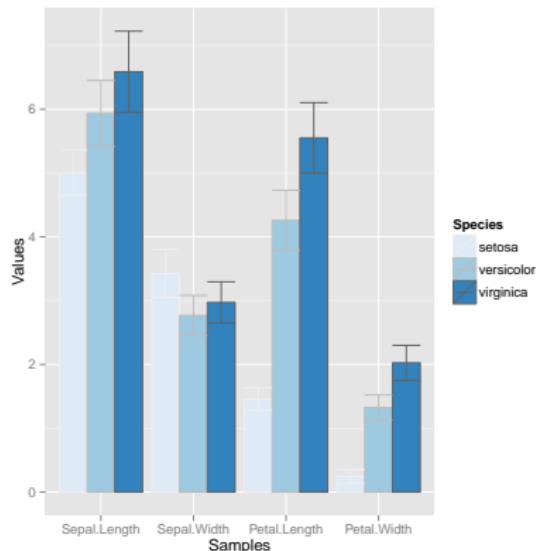
ggplot: Bar Plot with Error Bars

```
> p <- ggplot(df_mean, aes(Samples, Values, fill = Species)) +  
+         geom_bar(position="dodge", stat="identity") + geom_errorbar(limits  
> print(p)
```



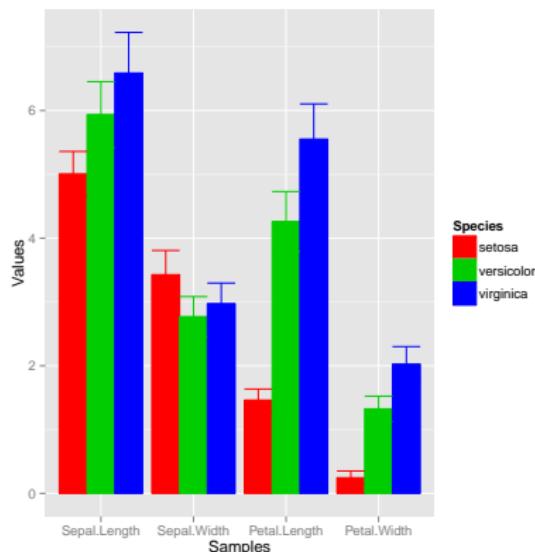
ggplot: Changing Color Settings

```
> library(RColorBrewer)
> # display.brewer.all()
> p <- ggplot(df_mean, aes(Samples, Values, fill=Species, color=Species)) +
+     geom_bar(position="dodge", stat="identity") + geom_errorbar(limits, position="dodge") +
+     scale_fill_brewer(palette="Blues") + scale_color_brewer(palette = "Greys")
> print(p)
```



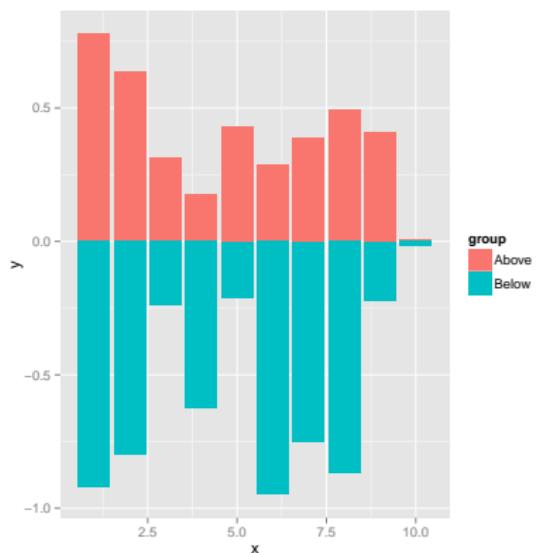
ggplot: Using Standard Colors

```
> p <- ggplot(df_mean, aes(Samples, Values, fill=Species, color=Species)) +  
+     geom_bar(position="dodge", stat="identity") + geom_errorbar(limits, position="dodge") +  
+     scale_fill_manual(values=c("red", "green3", "blue")) +  
+     scale_color_manual(values=c("red", "green3", "blue"))  
> print(p)
```



ggplot: Mirrored Bar Plots

```
> df <- data.frame(group = rep(c("Above", "Below"), each=10), x = rep(1:10, 2), y = c(runif(10, 0, 1), runif(10, -1, 0)))
> p <- ggplot(df, aes(x=x, y=y, fill=group)) +
+     geom_bar(stat="identity", position="identity")
> print(p)
```



Exercise 4: Bar Plots

- Task 1** Calculate the mean values for the Species components of the first four columns in the iris data set. Use the `melt` function from the `reshape2` package to bring the results into the expected format for `ggplot`.
- Task 2** Generate two bar plots: one with stacked bars and one with horizontally arranged bars.

Structure of iris data set:

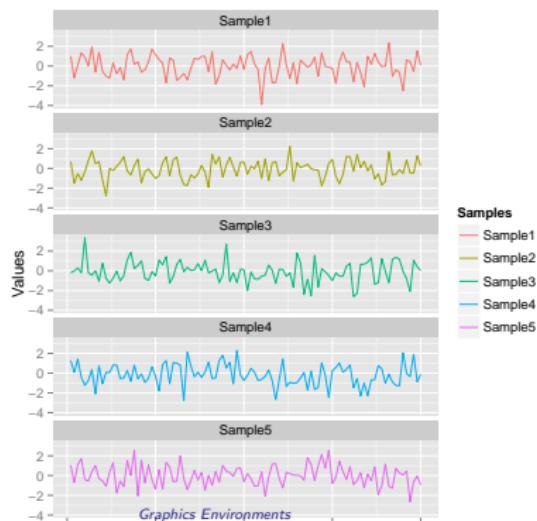
```
> class(iris)
[1] "data.frame"

> iris[1:4,]
   Sepal.Length Sepal.Width Petal.Length Petal.Width Species
1          5.1       3.5        1.4       0.2    setosa
2          4.9       3.0        1.4       0.2    setosa
3          4.7       3.2        1.3       0.2    setosa
4          4.6       3.1        1.5       0.2    setosa

> table(iris$Species)
setosa versicolor virginica
      50         50         50
```

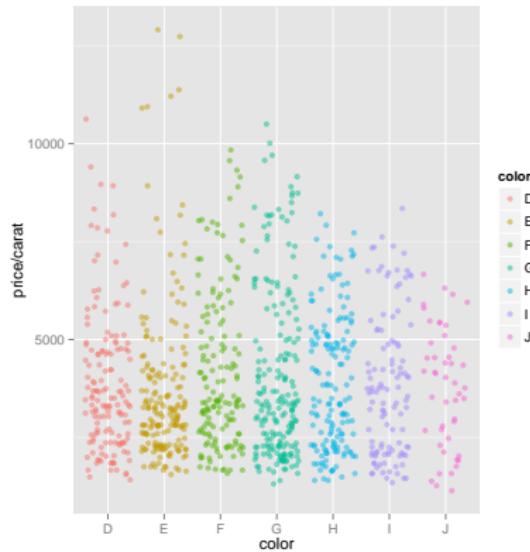
ggplot: Data Reformatting Example for Line Plot

```
> y <- matrix(rnorm(500), 100, 5, dimnames=list(paste("g", 1:100, sep=""), paste("Sample", 1:5, sep="")))
> y <- data.frame(Position=1:length(y[,1]), y)
> y[1:4, ] # First rows of input format expected by melt()
>
  Position     Sample1     Sample2     Sample3     Sample4     Sample5
g1          1  1.0002088  0.6850199 -0.21324932  1.27195056  1.0479301
g2          2 -1.2024596 -1.5004962 -0.01111579  0.07584497 -0.7100662
g3          3  0.1023678 -0.5153367  0.28564390  1.41522878  1.1084695
g4          4  1.3294248 -1.2084007 -0.19581898 -0.42361768  1.7139697
>
> df <- melt(y, id.vars=c("Position"), variable.name = "Samples", value.name="Values")
> p <- ggplot(df, aes(Position, Values)) + geom_line(aes(color=Samples)) + facet_wrap(~Samples, ncol=1)
> print(p)
> ## Represent same data in box plot
> ## ggplot(df, aes(Samples, Values, fill=Samples)) + geom_boxplot()
```



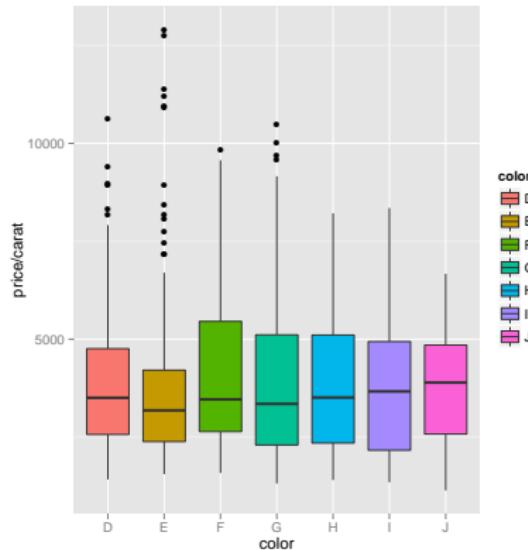
ggplot: Jitter Plots

```
> p <- ggplot(dsmall, aes(color, price/carat)) +  
+     geom_jitter(alpha = I(1 / 2), aes(color=color))  
> print(p)
```



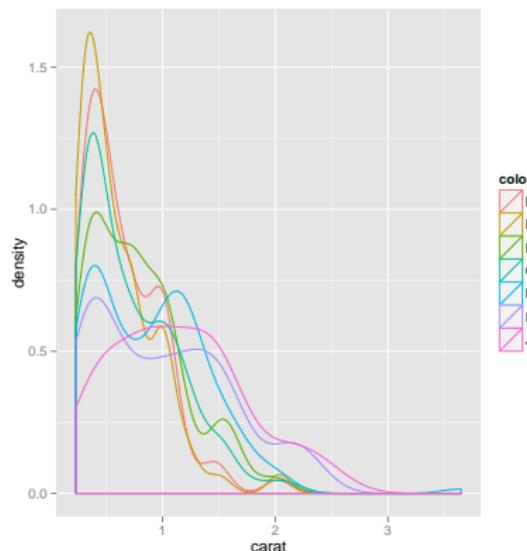
ggplot: Box Plots

```
> p <- ggplot(dsmall, aes(color, price/carat, fill=color)) + geom_boxplot()  
> print(p)
```



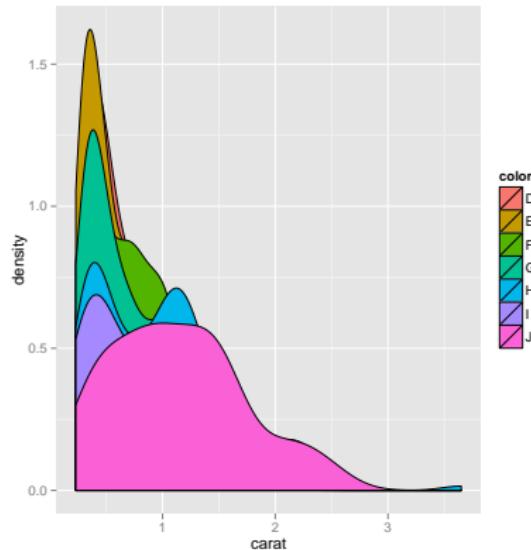
ggplot: Density Plot with Line Coloring

```
> p <- ggplot(dsmall, aes(carat)) + geom_density(aes(color = color))
> print(p)
```



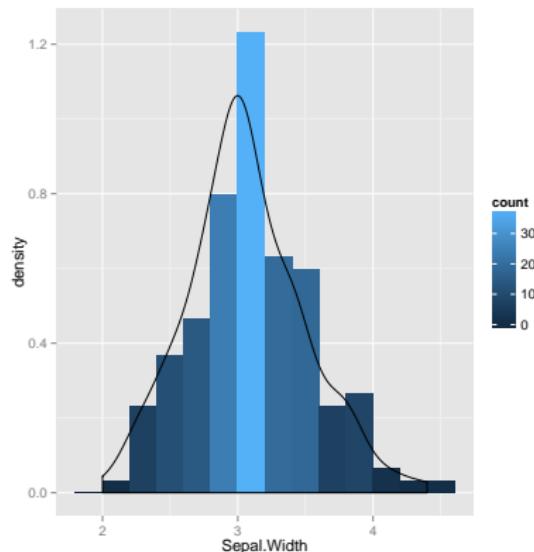
ggplot: Density Plot with Area Coloring

```
> p <- ggplot(dsmall, aes(carat)) + geom_density(aes(fill = color))
> print(p)
```



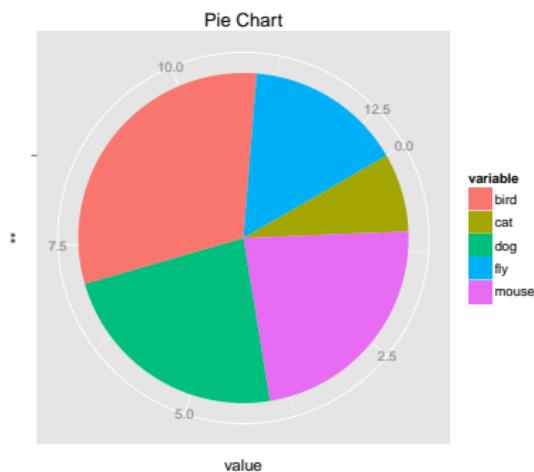
ggplot: Histograms

```
> p <- ggplot(iris, aes(x=Sepal.Width)) + geom_histogram(aes(y = ..density..,  
+                                         fill = ..count..), binwidth=0.2) + geom_density()  
> print(p)
```



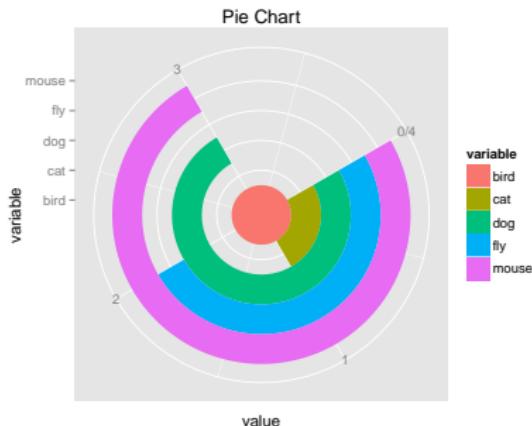
ggplot: Pie Chart

```
> df <- data.frame(variable=rep(c("cat", "mouse", "dog", "bird", "fly")),  
+                     value=c(1,3,3,4,2))  
> p <- ggplot(df, aes(x = "", y = value, fill = variable)) +  
+         geom_bar(width = 1, stat="identity") +  
+         coord_polar("y", start=pi / 3) + ggtitle("Pie Chart")  
> print(p)
```



ggplot: Wind Rose Pie Chart

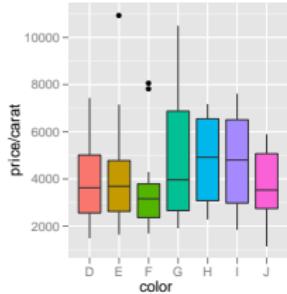
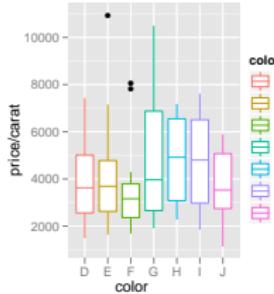
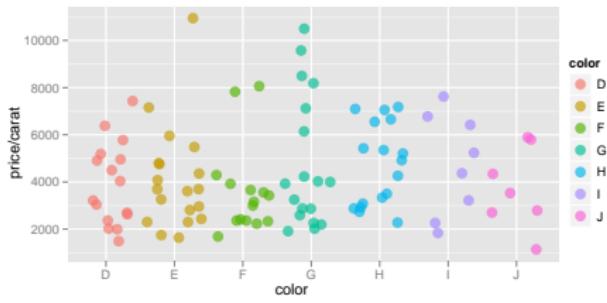
```
> p <- ggplot(df, aes(x = variable, y = value, fill = variable)) +  
+     geom_bar(width = 1, stat="identity") + coord_polar("y", start=pi / 3) +  
+     ggtitle("Pie Chart")  
> print(p)
```



ggplot: Arranging Graphics on One Page

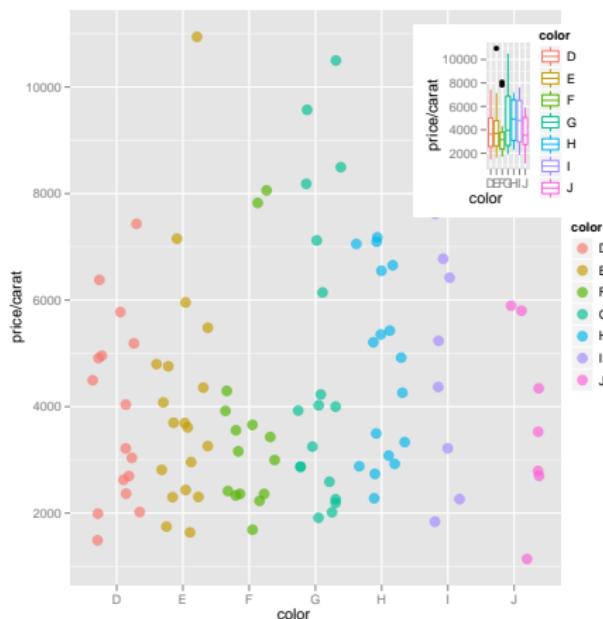
```
> library(grid)
> a <- ggplot(dsmall, aes(color, price/carat)) + geom_jitter(size=4, alpha = I(1 / 1.5), aes(color=color))
> b <- ggplot(dsmall, aes(color, price/carat, color=color)) + geom_boxplot()
> c <- ggplot(dsmall, aes(color, price/carat, fill=color)) + geom_boxplot() + theme(legend.position = "none")
> grid.newpage() # Open a new page on grid device
> pushViewport(viewport(layout = grid.layout(2, 2))) # Assign to device viewport with 2 by 2 grid layout
> print(a, vp = viewport(layout.pos.row = 1, layout.pos.col = 1:2))
> print(b, vp = viewport(layout.pos.row = 2, layout.pos.col = 1))
> print(c, vp = viewport(layout.pos.row = 2, layout.pos.col = 2, width=0.3, height=0.3, x=0.8, y=0.8))
```

ggplot: Arranging Graphics on One Page



ggplot: Inserting Graphics into Plots

```
> # pdf("insert.pdf")
> print(a)
> print(b, vp=viewport(width=0.3, height=0.3, x=0.8, y=0.8))
> # dev.off()
```



Outline

Overview

Graphics Environments

Base Graphics

Grid Graphics

lattice

ggplot2

Specialty Graphics

Genome Graphics

ggbio

Additional Genome Graphics

Clustering

Background

Hierarchical Clustering Example

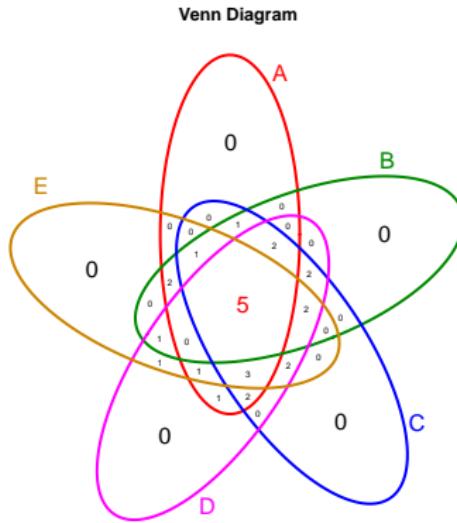
Non-Hierarchical Clustering Examples

Venn Diagrams (Code)

```
> source("http://faculty.ucr.edu/~tgirke/Documents/R_BioCond/My_R_Scripts/overLapper.R")

> setlist5 <- list(A=sample(letters, 18), B=sample(letters, 16), C=sample(letters, 20), D=sample(letters, 22))
> OLList5 <- overLapper(setlist=setlist5, sep="_", type="vennsets")
> counts <- sapply(OLList5$Venn_List, length)
> # pdf("venn.pdf")
> vennPlot(counts=counts, ccol=c(rep(1,30),2), lceex=1.5, ccex=c(rep(1.5,5), rep(0.6,25),1.5))
> # dev.off()
```

Venn Diagram (Plot)



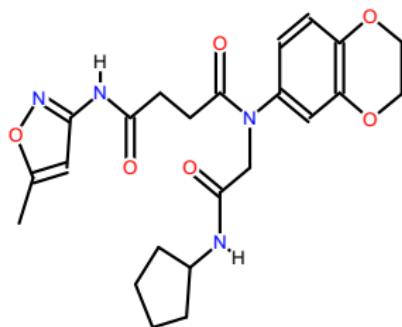
Unique objects: All = 26; S1 = 18; S2 = 16; S3 = 20; S4 = 22; S5 = 18

Figure: Venn Diagram

Compound Depictions with ChemmineR

```
> library(ChemmineR)
> data(sdfsamp)
> plot(sdfsamp[1], print=FALSE)
```

CMP1



Outline

Overview

Graphics Environments

Base Graphics

Grid Graphics

lattice

ggplot2

Specialty Graphics

Genome Graphics

ggbio

Additional Genome Graphics

Clustering

Background

Hierarchical Clustering Example

Non-Hierarchical Clustering Examples

Outline

Overview

Graphics Environments

Base Graphics

Grid Graphics

lattice

ggplot2

Specialty Graphics

Genome Graphics

ggbio

Additional Genome Graphics

Clustering

Background

Hierarchical Clustering Example

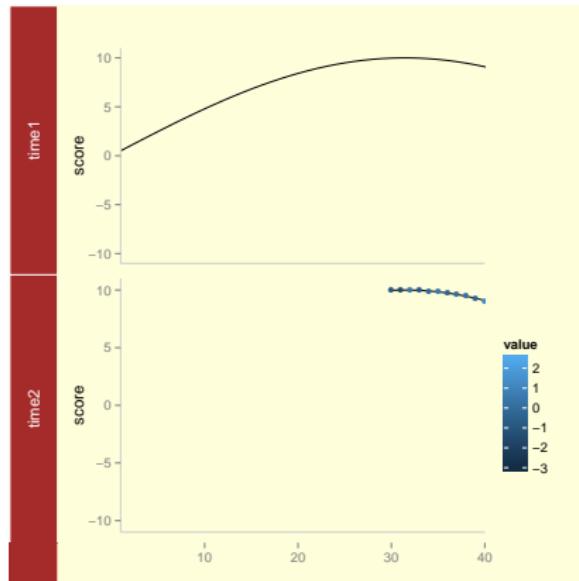
Non-Hierarchical Clustering Examples

ggbio: A Programmable Genome Browser

- A genome browser is a visualization tool for plotting different types of genomic data in separate tracks along chromosomes.
- The *ggbio* package (Yin et al., 2012) facilitates plotting of complex genome data objects, such as read alignments (SAM/BAM), genomic context/annotation information (gff/txdb), variant calls (VCF/BCF), and more. To easily compare these data sets, it extends the faceting facility of *ggplot2* to genome browser-like tracks.
- Most of the core object types for handling genomic data with R/Bioconductor are supported: GRanges, GAlignments, VCF, etc. For more details, see Table 1.1 of the *ggbio* vignette [Link](#).
- *ggbio*'s convenience plotting function is `autoplot`. For more customizable plots, one can use the generic `ggplot` function.
- Apart from the standard *ggplot2* plotting components, *ggbio* defines several new components useful for genomic data visualization. A detailed list is given in Table 1.2 of the vignette [Link](#).
- Useful web sites:
 - *ggbio* manual [Link](#)
 - *ggbio* functions [Link](#)
 - *autoplot* demo [Link](#)

Tracks: Aligning Plots Along Chromosomes

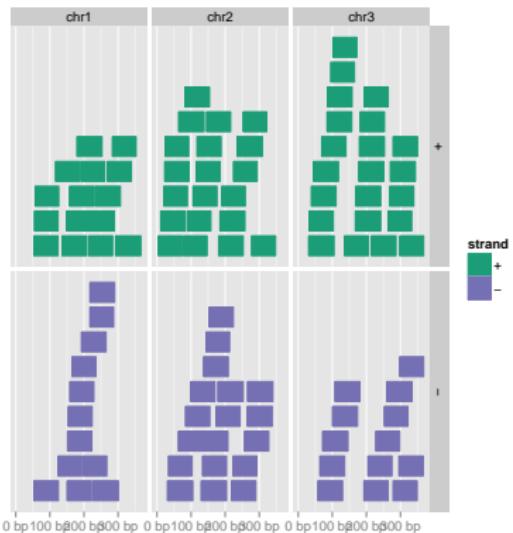
```
> library(ggbio)
> df1 <- data.frame(time = 1:100, score = sin((1:100)/20)*10)
> p1 <- qplot(data = df1, x = time, y = score, geom = "line")
> df2 <- data.frame(time = 30:120, score = sin((30:120)/20)*10, value = rnorm(120-30 +1))
> p2 <- ggplot(data = df2, aes(x = time, y = score)) + geom_line() + geom_point(size = 2, aes(color = value))
> tracks(time1 = p1, time2 = p2) + xlim(1, 40) + theme_tracks_sunset()
```



Plotting Genomic Ranges

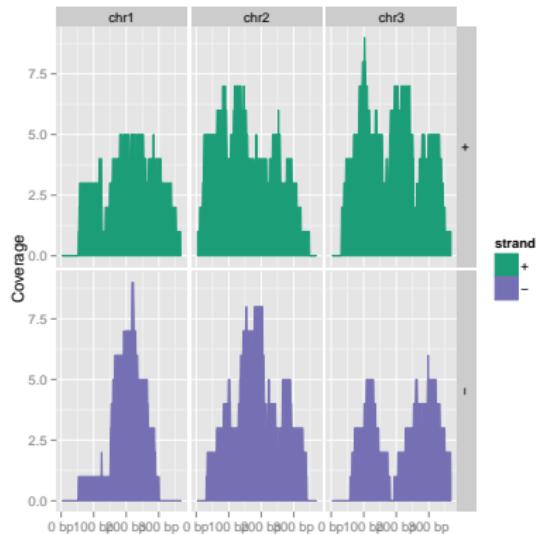
GRanges objects are essential for storing alignment or annotation ranges in R/Bioconductor. The following creates a sample GRanges object and plots its content.

```
> library(GenomicRanges)
> set.seed(1); N <- 100; gr <- GRanges(seqnames = sample(c("chr1", "chr2", "chr3"), size = N, replace = TRUE),
> autplot(gr, aes(color = strand, fill = strand), facets = strand ~ seqnames)
```



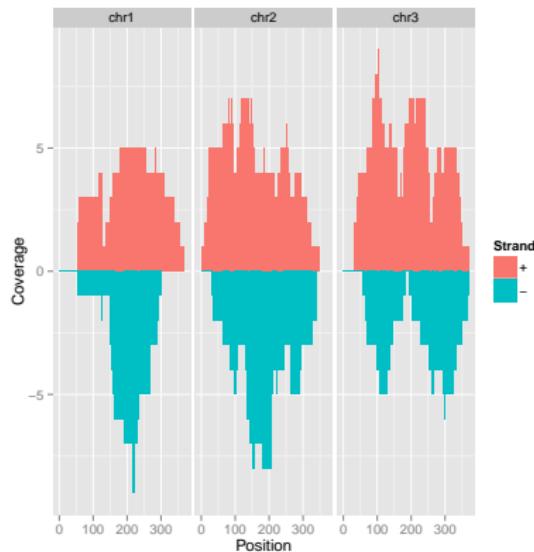
Plotting Coverage Instead of Ranges

```
> autoplot(gr, aes(color = strand, fill = strand), facets = strand ~ seqnames, stat = "coverage")
```



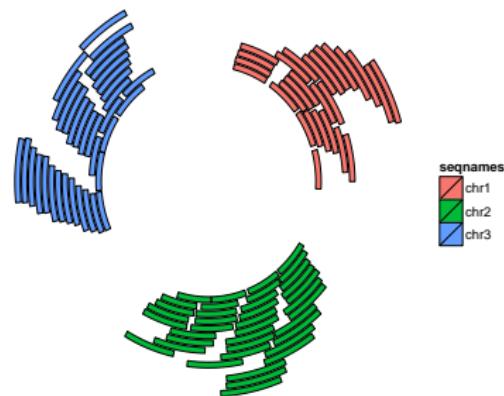
Mirrored Coverage Plot

```
> pos <- sapply(coverage(gr$strand(gr)=="+"), as.numeric)
> pos <- data.frame(Chr=rep(names(pos), sapply(pos, length)), Strand=rep("+", length(unlist(pos))), Position=unlist(pos))
> neg <- sapply(coverage(gr$strand(gr)=="-"), as.numeric)
> neg <- data.frame(Chr=rep(names(neg), sapply(neg, length)), Strand=rep("-", length(unlist(neg))), Position=unlist(neg))
> covdf <- rbind(pos, neg)
> p <- ggplot(covdf, aes(Position, Coverage, fill=Strand)) +
+     geom_bar(stat="identity", position="identity") + facet_wrap(~Chr)
> p
```



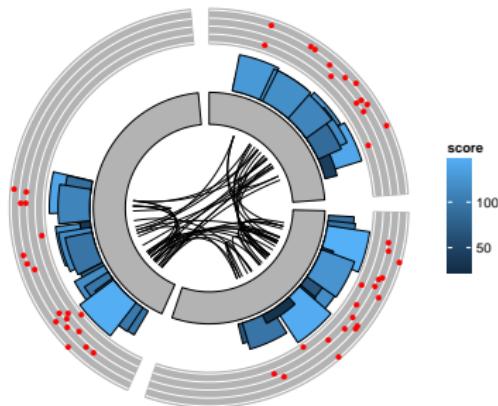
Circular Layout

```
> autoplot(gr, layout = "circle", aes(fill = seqnames))
```



More Complex Circular Example

```
> seqlengths(gr) <- c(400, 500, 700)
> values(gr)$to.gr <- gr[sample(1:length(gr), size = length(gr))]
> idx <- sample(1:length(gr), size = 50)
> gr <- gr[idx]
> ggplot() + layout_circle(gr, geom = "ideo", fill = "gray70", radius = 7, trackWidth = 3) +
+   layout_circle(gr, geom = "bar", radius = 10, trackWidth = 4,
+                 aes(fill = score, y = score)) +
+   layout_circle(gr, geom = "point", color = "red", radius = 14,
+                 trackWidth = 3, grid = TRUE, aes(y = score)) +
+   layout_circle(gr, geom = "link", linked.to = "to.gr", radius = 6, trackWidth = 1)
```

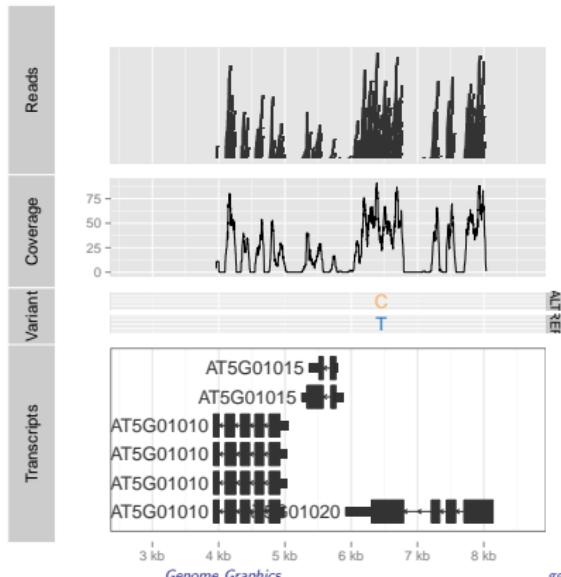


Viewing Alignments and Variants

To make the following example work, please download and unpack this data archive

[Link](#) containing GFF, BAM and VCF sample files.

```
> library(rtracklayer); library(GenomicFeatures); library(Rsamtools); library(VariantAnnotation)
> ga <- readGAlignmentsFromBam("./data/SRR064167.fastq.bam", use.names=TRUE, param=ScanBamParam(which=GRanges))
> p1 <- autoplot(ga, geom = "rect")
> p2 <- autoplot(ga, geom = "line", stat = "coverage")
> vcf <- readVcf(file="data/varianttools_gnsap.vcf", genome="ATH1")
> p3 <- autoplot(vcf[seqnames(vcf)=="Chr5"], type = "fixed") + xlim(4000, 8000) + theme(legend.position =
> txdb <- makeTranscriptDbFromGFF(file="./data/TAIR10_GFF3_trunc.gff", format="gff3")
> p4 <- autoplot(txdb, which=GRanges("Chr5", IRanges(4000, 8000)), names.expr = "gene_id")
> tracks(Reads=p1, Coverage=p2, Variant=p3, Transcripts=p4, heights = c(0.3, 0.2, 0.1, 0.35)) + ylab("")
```



Additional Sample Plots

- autoplot demo [Link](#)

Outline

Overview

Graphics Environments

Base Graphics

Grid Graphics

lattice

ggplot2

Specialty Graphics

Genome Graphics

ggbio

Additional Genome Graphics

Clustering

Background

Hierarchical Clustering Example

Non-Hierarchical Clustering Examples

Additional Packages for Visualizing Genome Data

- Gviz [Link](#)
- RCircos (Zhang et al., 2013) [Link](#)
- Genome Graphs [Link](#)
- genoPlotR [Link](#)

Outline

Overview

Graphics Environments

Base Graphics

Grid Graphics

lattice

ggplot2

Specialty Graphics

Genome Graphics

ggbio

Additional Genome Graphics

Clustering

Background

Hierarchical Clustering Example

Non-Hierarchical Clustering Examples

Outline

Overview

Graphics Environments

Base Graphics

Grid Graphics

lattice

ggplot2

Specialty Graphics

Genome Graphics

ggbio

Additional Genome Graphics

Clustering

Background

Hierarchical Clustering Example

Non-Hierarchical Clustering Examples

What is Clustering?

- Clustering is the classification/partitioning of data objects into similarity groups (clusters) according to a defined distance measure.
- It is used in many fields, such as machine learning, data mining, pattern recognition, image analysis, genomics, systems biology, etc.
- Machine learning typically regards data clustering as a form of unsupervised learning.

Why Clustering and Data Mining in R?

- Efficient data structures and functions for clustering.
- Efficient environment for algorithm prototyping and benchmarking.
- Comprehensive set of clustering and machine learning libraries.
- Standard for data analysis in many areas.
- Overview: [Clustering Task View on CRAN](#)

- Center & standardize
 - ① Center: subtract vector mean from each value
 - ② Standardize: devide by standard deviation

⇒ $\text{Mean} = 0$ and $\text{STDEV} = 1$
- Center & scale with the `scale()` fuction
 - ① Center: subtract vector mean from each value
 - ② Scale: divide centered vector by their root mean square (rms)

$$x_{rms} = \sqrt{\frac{1}{n-1} \sum_{i=1}^n x_i^2}$$

⇒ $\text{Mean} = 0$ and $\text{STDEV} = 1$

- Log transformation
- Rank transformation: replace measured values by ranks
- No transformation

- Euclidean distance for two profiles X and Y

$$d(X, Y) = \sqrt{\sum_{i=1}^n (x_i - y_i)^2}$$

Disadvantages: not scale invariant, not for negative correlations

- Maximum, Manhattan, Canberra, binary, Minowski, ...
- Correlation-based distance: $1 - r$
 - Pearson correlation coefficient (PCC)

$$r = \frac{n \sum_{i=1}^n x_i y_i - \sum_{i=1}^n x_i \sum_{i=1}^n y_i}{\sqrt{(\sum_{i=1}^n x_i^2 - (\sum_{i=1}^n x_i)^2)(\sum_{i=1}^n y_i^2 - (\sum_{i=1}^n y_i)^2)}}$$

Disadvantage: outlier sensitive

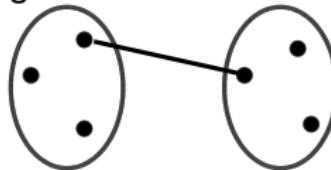
- Spearman correlation coefficient (SCC)
Same calculation as PCC but with ranked values!

There Are Many more Distance Measures

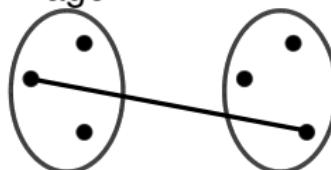
- If the distances among items are quantifiable, then clustering is possible.
- Choose the most accurate and meaningful distance measure for a given field of application.
- If uncertain then choose several distance measures and compare the results.

Cluster Linkage

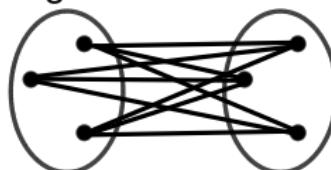
Single Linkage



Complete Linkage



Average Linkage



Outline

Overview

Graphics Environments

Base Graphics

Grid Graphics

lattice

ggplot2

Specialty Graphics

Genome Graphics

ggbio

Additional Genome Graphics

Clustering

Background

Hierarchical Clustering Example

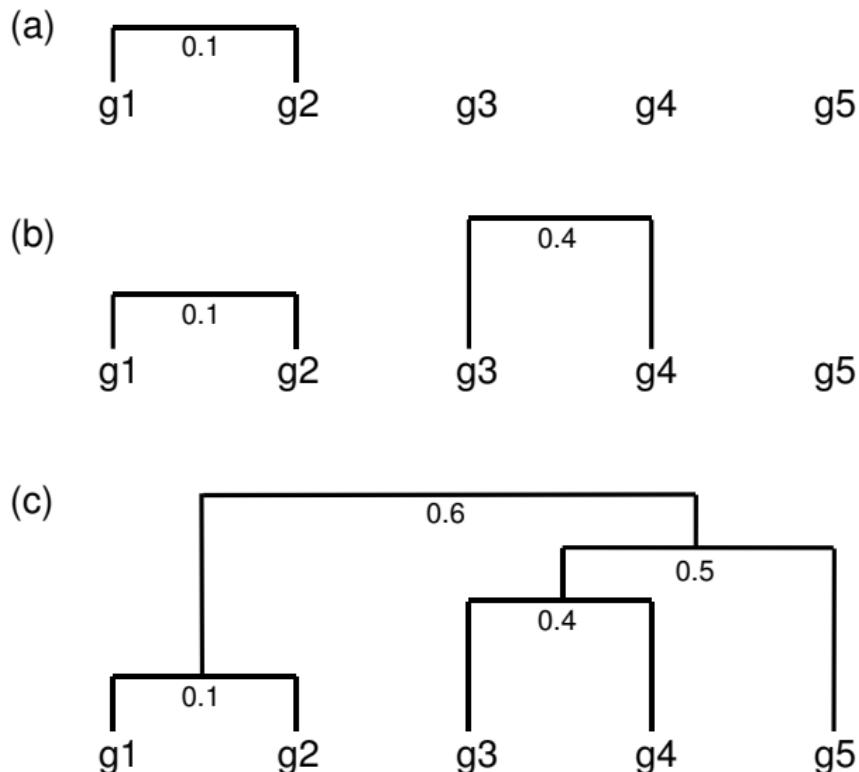
Non-Hierarchical Clustering Examples

Hierarchical Clustering Steps

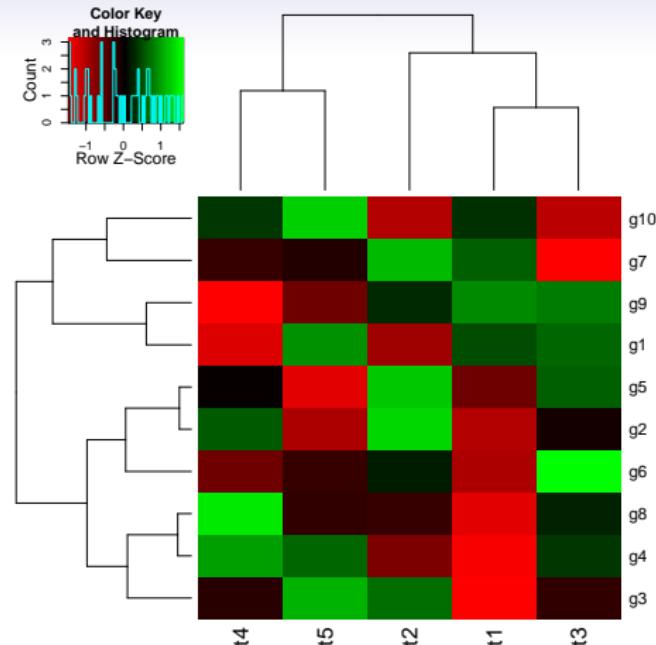
- ① Identify clusters (items) with closest distance
- ② Join them to new clusters
- ③ Compute distance between clusters (items)
- ④ Return to step 1

Hierarchical Clustering

Agglomerative Approach



Hierarchical Clustering Result with Heatmap



- A heatmap is a color coded table. To visually identify patterns, the rows and columns of a heatmap are often sorted by hierarchical clustering trees.
- In case of gene expression data, the row tree usually represents the genes, the column tree the treatments and the colors in the heat table represent the intensities or ratios of the underlying gene expression data set.

Hierarchical Clustering Approaches in R

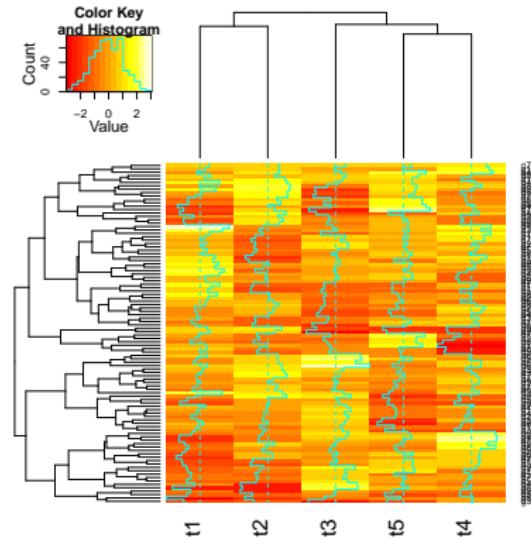
- ① Agglomerative approach (bottom-up)
`hclust()` and `agnes()`
- ② Divisive approach (top-down)
`diana()`

Tree Cutting to Obtain Discrete Clusters

- ① Node height in tree
- ② Number of clusters
- ③ Search tree nodes by distance cutoff

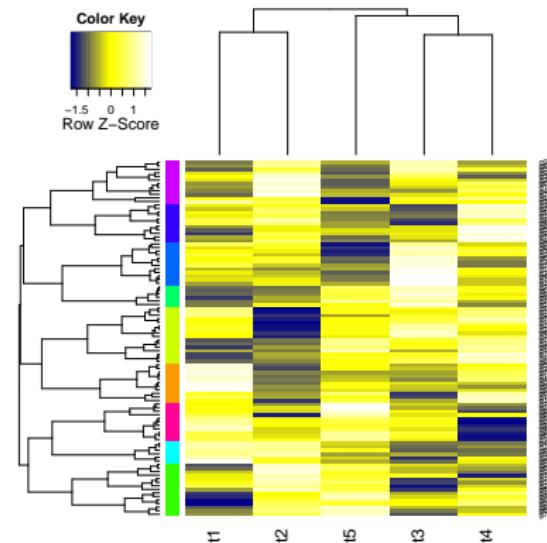
Example: hclust and heatmap.2

```
> library(gplots)
> y <- matrix(rnorm(500), 100, 5, dimnames=list(paste("g", 1:100, sep=""), paste("t", 1:5, sep="")))
> heatmap.2(y) # Shortcut to final result
```



Example: Stepwise Approach with Tree Cutting

```
> ## Row- and column-wise clustering  
> hr <- hclust(as.dist(1-cor(t(y), method="pearson")), method="complete")  
> hc <- hclust(as.dist(1-cor(y, method="spearman")), method="complete")  
> ## Tree cutting  
> mycl <- cutree(hr, h=max(hr$height)/1.5); mycolhc <- rainbow(length(unique(mycl)), start=0.1, end=0.9); mycolhc  
> ## Plot heatmap  
> mycol <- colorpanel(40, "darkblue", "yellow", "white") # or try redgreen(75)  
> heatmap.2(y, Rowv=as.dendrogram(hr), Colv=as.dendrogram(hc), col=mycol, scale="row", density.info="none", trace=0)
```



Outline

Overview

Graphics Environments

Base Graphics

Grid Graphics

lattice

ggplot2

Specialty Graphics

Genome Graphics

ggbio

Additional Genome Graphics

Clustering

Background

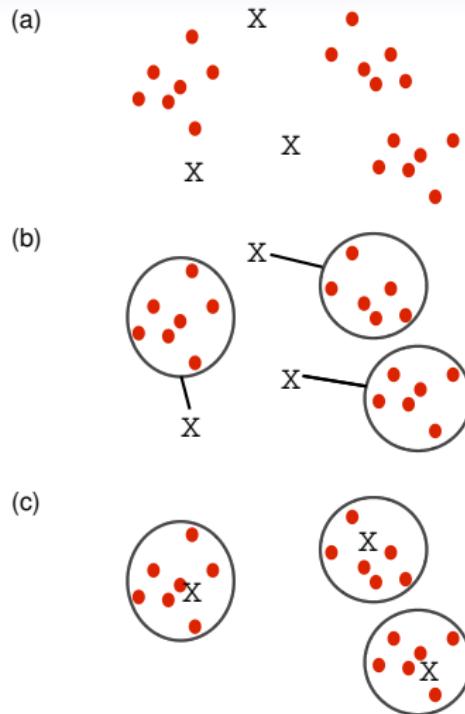
Hierarchical Clustering Example

Non-Hierarchical Clustering Examples

K-Means Clustering

- ① Choose the number of k clusters
- ② Randomly assign items to the k clusters
- ③ Calculate new centroid for each of the k clusters
- ④ Calculate the distance of all items to the k centroids
- ⑤ Assign items to closest centroid
- ⑥ Repeat until clusters assignments are stable

K-Means



Example: Clustering with kmeans Function

```
> km <- kmeans(t(scale(t(y))), 3)
```

```
> km$cluster
```

g1	g2	g3	g4	g5	g6	g7	g8	g9	g10	g11	g12	g13	g14	g15	g16	g17
2	1	2	3	1	3	1	2	2	1	3	1	3	2	3	3	3
g45	g46	g47	g48	g49	g50	g51	g52	g53	g54	g55	g56	g57	g58	g59	g60	g61
3	3	3	2	2	2	1	2	1	3	2	3	1	1	2	3	3
g89	g90	g91	g92	g93	g94	g95	g96	g97	g98	g99	g100					
3	1	1	3	2	1	2	1	1	3	3	2					

Fuzzy C-Means Clustering

- ① In contrast to strict (hard) clustering approaches, fuzzy (soft) clustering methods allow multiple cluster memberships of the clustered items.
- ② This is commonly achieved by assigning to each item a weight of belonging to each cluster.
- ③ Thus, items on the edge of a cluster, may be in the cluster to a lesser degree than items in the center of a cluster.
- ④ Typically, each item has as many coefficients (weights) as there are clusters that sum up for each item to one.

Example: Fuzzy Clustering with fanny

```
> library(cluster) # Loads the cluster library.  
> fannyy <- fanny(y, k=4, metric = "euclidean", memb.exp = 1.2)  
> round(fannyy$membership, 2)[1:4,]  
  
 [,1] [,2] [,3] [,4]  
g1 0.82 0.04 0.10 0.05  
g2 0.82 0.05 0.12 0.01  
g3 0.98 0.01 0.01 0.01  
g4 0.03 0.82 0.03 0.12  
  
> fannyy$clustering  
  
 g1   g2   g3   g4   g5   g6   g7   g8   g9   g10  g11  g12  g13  g14  g15  g16  g17  
  1    1    1    2    3    4    1    1    1    1    1    3    4    4    4    2    4  
g45  g46  g47  g48  g49  g50  g51  g52  g53  g54  g55  g56  g57  g58  g59  g60  g61  
  2    4    2    4    1    1    3    1    3    4    1    2    3    3    3    3    3    2  
g89  g90  g91  g92  g93  g94  g95  g96  g97  g98  g99  g100  
  2    3    3    2    4    3    1    2    1    4    4    4
```

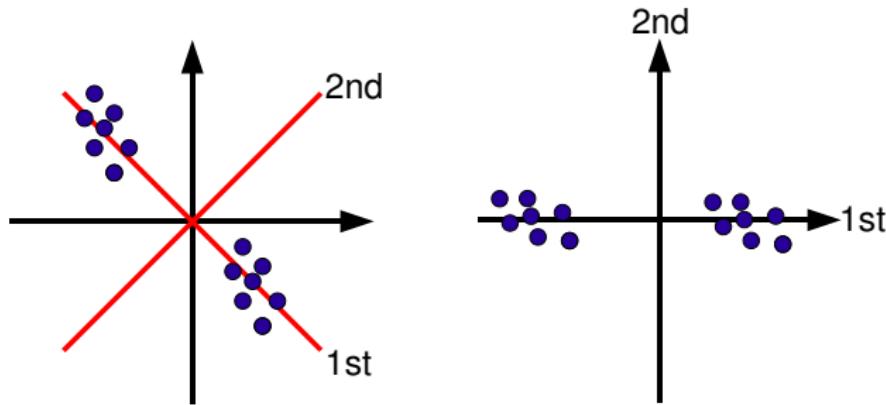
Principal Component Analysis (PCA)

Principal components analysis (PCA) is a data reduction technique that allows to simplify multidimensional data sets to 2 or 3 dimensions for plotting purposes and visual variance analysis.

Basic PCA Steps

- Center (and standardize) data
- First principal component axis
 - Across centroid of data cloud
 - Distance of each point to that line is minimized, so that it crosses the maximum variation of the data cloud
- Second principal component axis
 - Orthogonal to first principal component
 - Along maximum variation in the data
- 1st PCA axis becomes x-axis and 2nd PCA axis y-axis
- Continue process until the necessary number of principal components is obtained

PCA on Two-Dimensional Data Set



Identifies the Amount of Variability between Components

Example

Principal Component	1st	2nd	3rd	Other
Proportion of Variance	62%	34%	3%	rest

1st and 2nd principal components explain 96% of variance.

Example: PCA

```
> pca <- prcomp(y, scale=T)
> summary(pca) # Prints variance summary for all principal components
```

Importance of components:

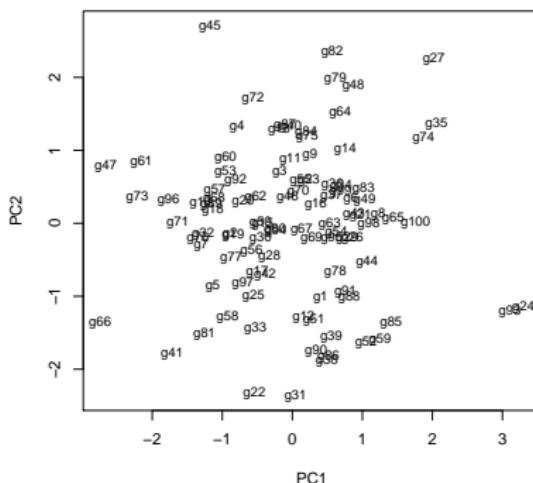
	PC1	PC2	PC3	PC4	PC5
--	-----	-----	-----	-----	-----

Standard deviation 1.0996 1.0505 1.0247 0.9219 0.8873

Proportion of Variance 0.2418 0.2207 0.2100 0.1700 0.1575

Cumulative Proportion 0.2418 0.4626 0.6725 0.8425 1.0000

```
> plot(pca$x, pch=20, col="blue", type="n") # To plot dots, drop type="n"
> text(pca$x, rownames(pca$x), cex=0.8)
```



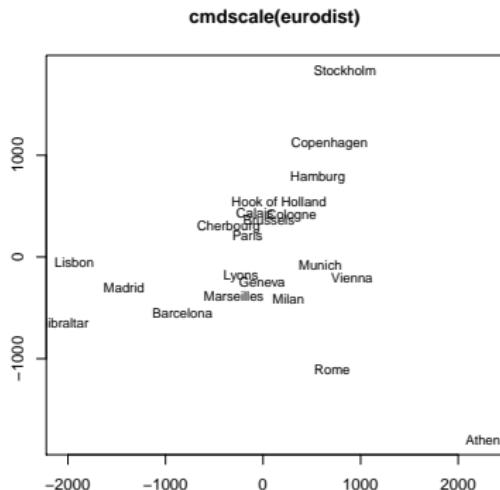
Multidimensional Scaling (MDS)

- Alternative dimensionality reduction approach
- Represents distances in 2D or 3D space
- Starts from distance matrix (PCA uses data points)

Example: MDS with cmdscale

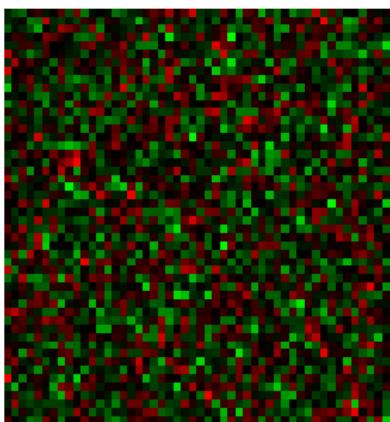
The following example performs MDS analysis on the geographic distances among European cities.

```
> loc <- cmdscale(eurodist)
> plot(loc[,1], -loc[,2], type="n", xlab="", ylab="", main="cmdscale(eurodist)")
> text(loc[,1], -loc[,2], rownames(loc), cex=0.8)
```

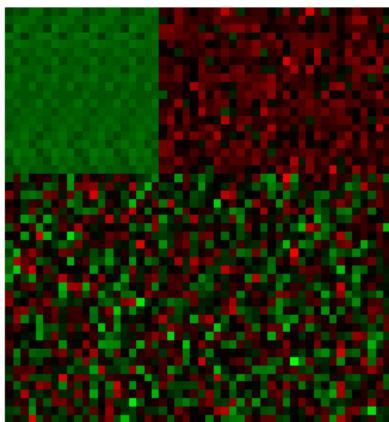


Biclustering

Finds in matrix subgroups of rows and columns which are as similar as possible to each other and as different as possible to the remaining data points.



Unclustered



Clustered

Bibliography I

Yin, T., Cook, D., Lawrence, M., Aug 2012. ggbio: an R package for extending the grammar of graphics for genomic data. *Genome Biol* 13 (8).

URL <http://www.hubmed.org/display.cgi?uids=22937822>

Zhang, H., Meltzer, P., Davis, S., 2013. RCircos: an R package for Circos 2D track plots. *BMC Bioinformatics* 14, 244–244.

URL <http://www.hubmed.org/display.cgi?uids=23937229>