# Graphics and Data Visualization in R

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### Overview

### Graphics in R

- Powerful environment for visualizing scientific data
- Integrated graphics and statistics infrastructure
- Publication quality graphics
- Fully programmable
- Highly reproducible
- Full LaTeX, Sweave, knitr and R Markdown support.
- Vast number of R packages with graphics utilities

### Documentation on Graphics in R

- General
  - Graphics Task Page
  - R Graph Gallery
  - R Graphical Manual
  - Paul Murrell's book R (Grid) Graphics
- Interactive graphics
  - rggobi (GGobi)
  - iplots
  - Open GL (rgl)

#### **Graphics Environments**

- Viewing and savings graphics in R
  - On-screen graphics
  - postscript, pdf, svg
  - jpeg/png/wmf/tiff/...
- Four major graphics environments
  - Low-level infrastructure
    - \* R Base Graphics (low- and high-level)
    - \* grid: Manual, Book
  - High-level infrastructure
    - \* lattice: Manual, Intro, Book
    - \* ggplot2: Manual, Intro, Book

# **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 multivariant data
- Help on these functions
  - ?myfct
  - ?plot
  - ?par

### Preferred Input Data Objects

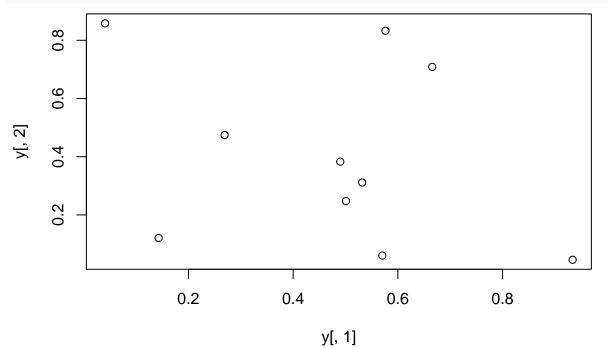
- Matrices and data frames
- Vectors
- Named vectors

### **Scatter Plots**

### Basic scatter plots

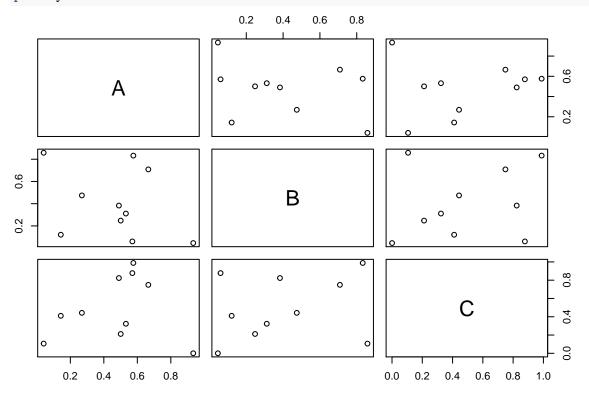
Sample data set for subsequent plots

```
set.seed(1410)
y <- matrix(runif(30), ncol=3, dimnames=list(letters[1:10], LETTERS[1:3]))
plot(y[,1], y[,2])</pre>
```



# All pairs

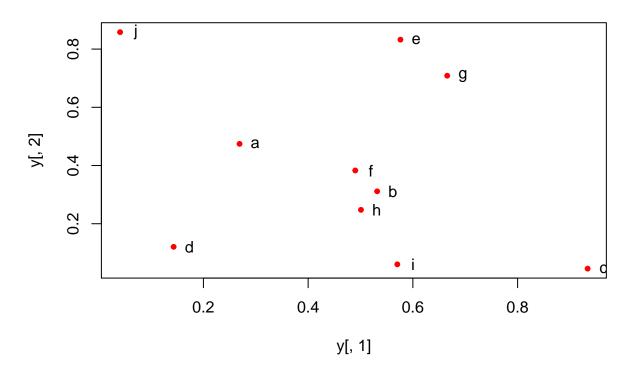
# pairs(y)



### Plot labels

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

# **Symbols and Labels**



### 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))
```

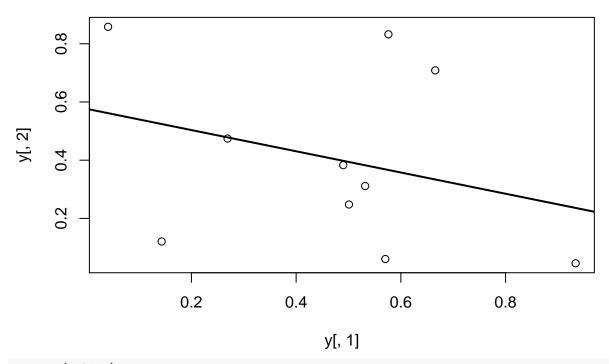
Usage of important plotting parameters

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

Add a regression line to a plot

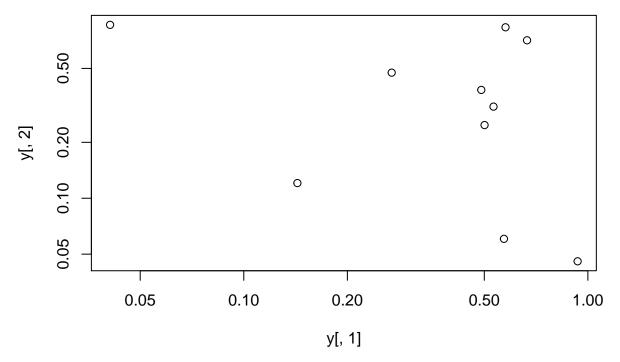
```
plot(y[,1], y[,2])

myline \leftarrow lm(y[,2]~y[,1]); abline(myline, lwd=2)
```

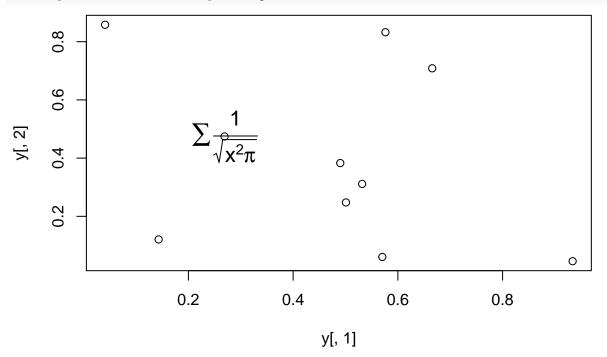


### summary(myline)

```
##
## Call:
## lm(formula = y[, 2] ~ y[, 1])
##
## Residuals:
##
       Min
                  1Q
                      Median
                                    3Q
                                            Max
## -0.40357 -0.17912 -0.04299 0.22147 0.46623
##
## Coefficients:
##
              Estimate Std. Error t value Pr(>|t|)
                0.5764
                            0.2110
                                     2.732
                                            0.0258 *
## (Intercept)
                -0.3647
                            0.3959 -0.921
                                            0.3839
## y[, 1]
## ---
## Signif. codes: 0 '***' 0.001 '**' 0.05 '.' 0.1 ' ' 1
## Residual standard error: 0.3095 on 8 degrees of freedom
## Multiple R-squared: 0.09589, Adjusted R-squared: -0.01712
## F-statistic: 0.8485 on 1 and 8 DF, p-value: 0.3839
Same plot as above, but on log scale
plot(y[,1], y[,2], log="xy")
```



Add a mathematical expression to a plot



### Exercise 1

• Task 1: Generate scatter plot for first two columns in iris data frame and color dots by its Species column.

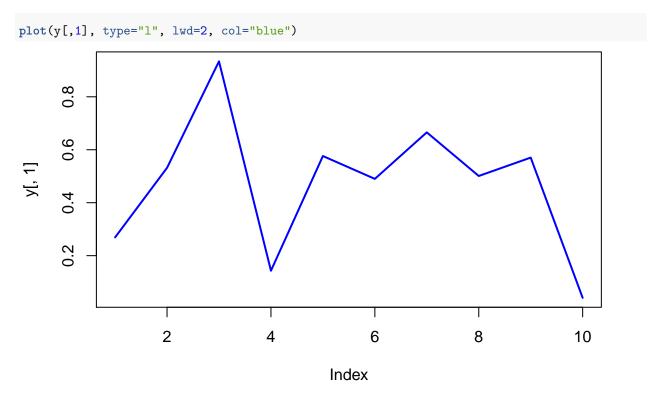
• Task 2: Use the xlim/ylim arguments to set limits on the x- and y-axes so that all data points are restricted to the left bottom quadrant of the plot.

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
                                                      0.2
                                         1.3
                                                           setosa
## 4
              4.6
                           3.1
                                         1.5
                                                      0.2
                                                           setosa
table(iris$Species)
##
##
       setosa versicolor
                           virginica
##
           50
                       50
```

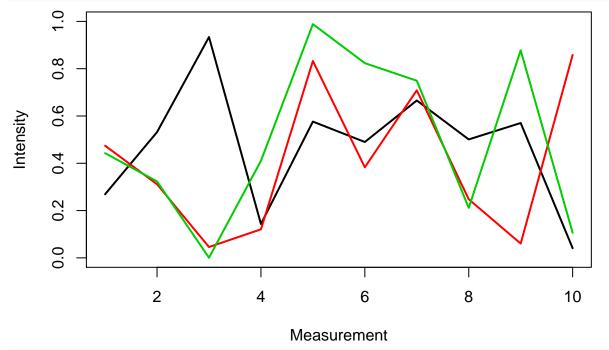
### Line Plots

### Single Data Set



### Many Data Sets

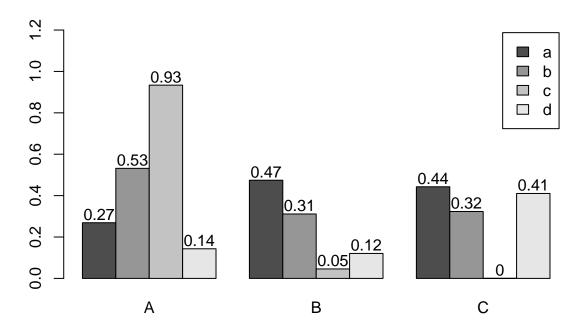
Plots line graph for all columns in data frame y. The split.screen function is used in this example in a for loop to overlay several line graphs in the same plot.



close.screen(all=TRUE)

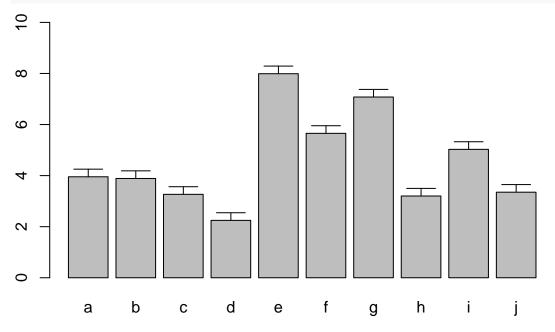
### **Bar Plots**

#### Basics



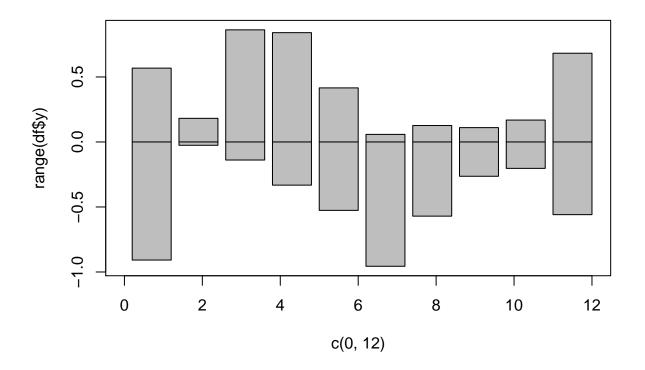
#### Error bars

```
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)</pre>
```



### Mirrored bar plot

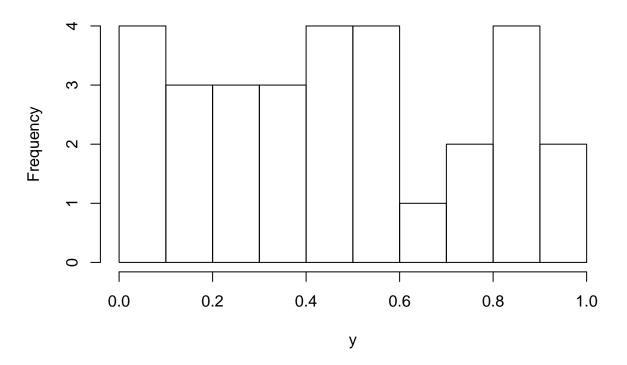
```
df <- data.frame(group = rep(c("Above", "Below"), each=10), x = rep(1:10, 2), y = c(runif(10, 0, 1), runplot(c(0,12),range(df$y),type = "n")
barplot(height = df$y[df$group == "Above"], add = TRUE,axes = FALSE)
barplot(height = df$y[df$group == "Below"], add = TRUE,axes = FALSE)</pre>
```



# Histograms

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

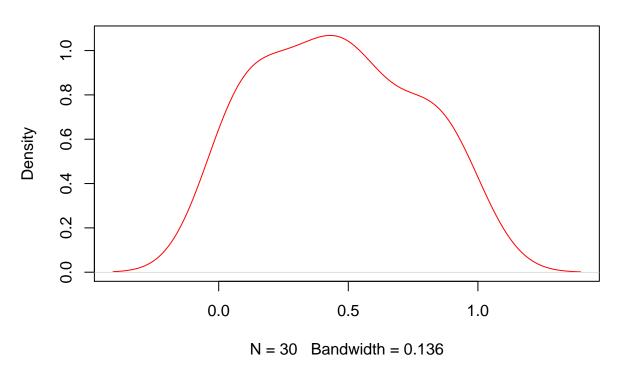
# Histogram of y



# Density Plots}

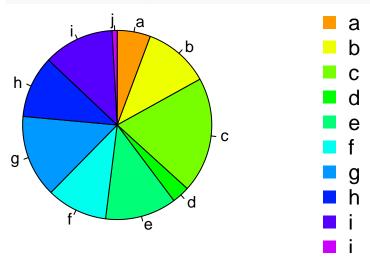
plot(density(y), col="red")

# density.default(x = y)



# 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

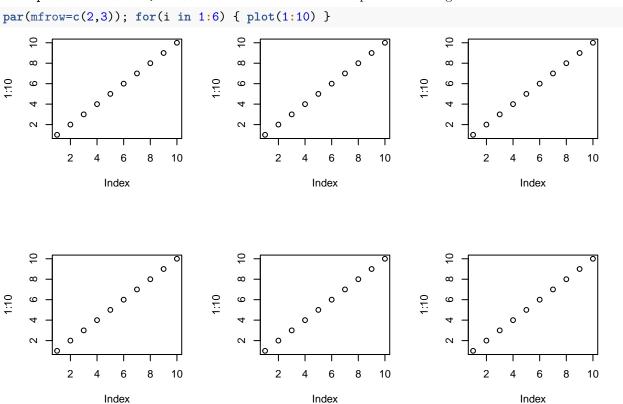
Default color palette and how to change it

```
palette()
## [1] "black"
                  "red"
                             "green3"
                                       "blue"
                                                  "cyan"
                                                             "magenta" "yellow"
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

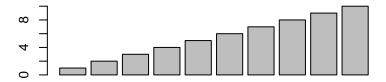
### Arranging Several Plots on Single Page

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



### 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.



### 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 3.5 5.1 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
```

# **Grid Graphics**

- 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
  - Book

# lattice Graphics

- 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
  - Intro
  - Book

Open a list of all functions available in the lattice package

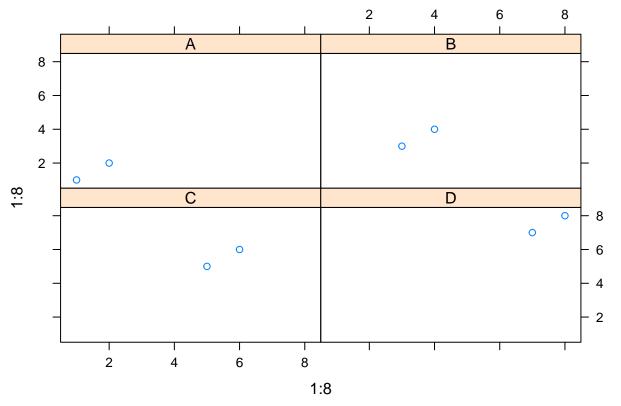
```
library(help=lattice)
```

Accessing and changing global parameters:

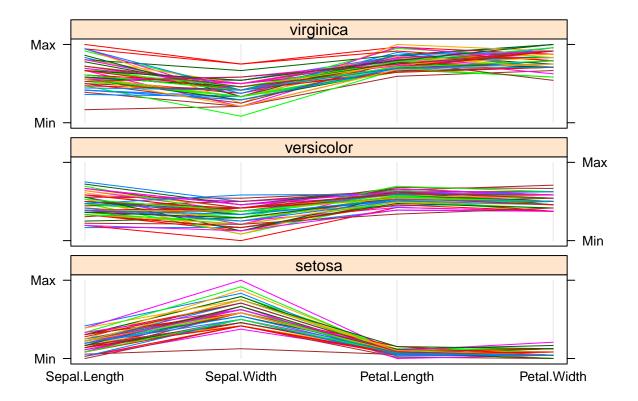
```
?lattice.options
?trellis.device
```

### Scatter Plot Sample

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



# Line Plot Sample



# ggplot2 Graphics

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

### 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 see here
- Settings of plotting theme can be accessed with the command theme\_get() and its settings can be changed with theme().
- Preferred input data object
  - qplot: data.frame (support for vector, matrix, ...)
  - ggplot: data.frame

- Packages with convenience utilities to create expected inputs
  - plyr
  - reshape

### qplot Function

The syntax of qplot is similar as R's basic plot function

```
• Arguments
```

```
x: x-coordinates (e.g. col1)
y: y-coordinates (e.g. col2)
data: data.frame or tibble 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
```

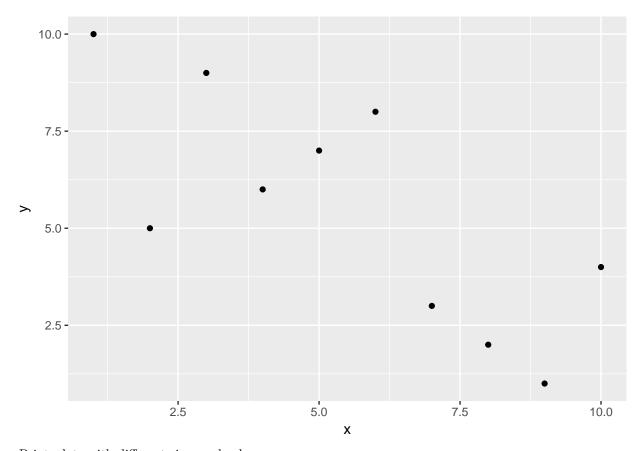
- ...: many arguments accepted by plot function

### qplot: scatter plot basics

Create sample data

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

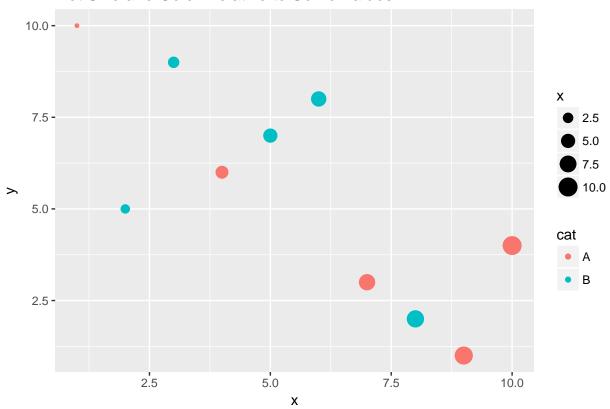
```
qplot(x, y, geom="point")
```



Prints dots with different sizes and colors

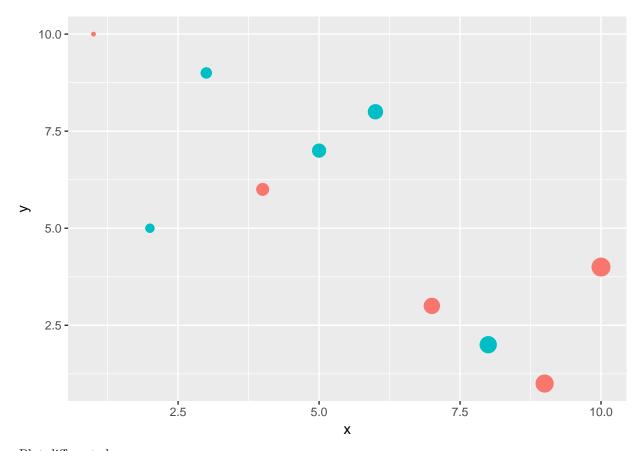
```
qplot(x, y, geom="point", size=x, color=cat,
    main="Dot Size and Color Relative to Some Values")
```

# Dot Size and Color Relative to Some Values



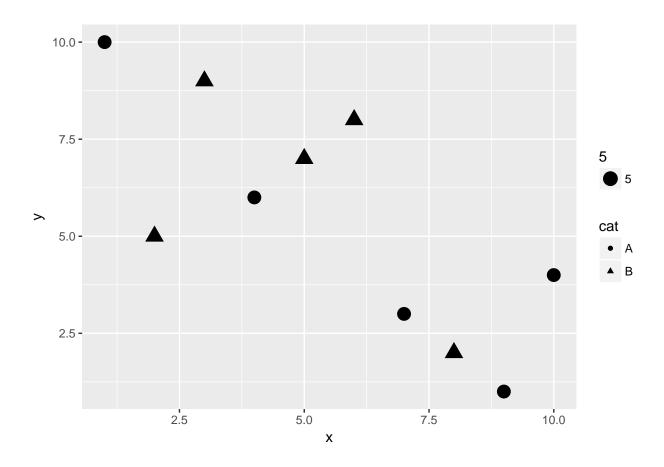
# Drops legend

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



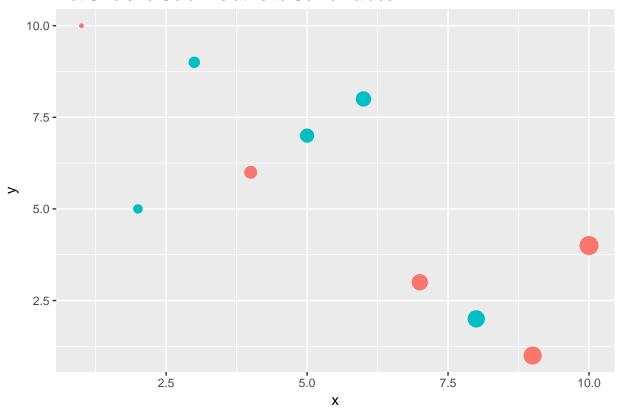
Plot different shapes

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

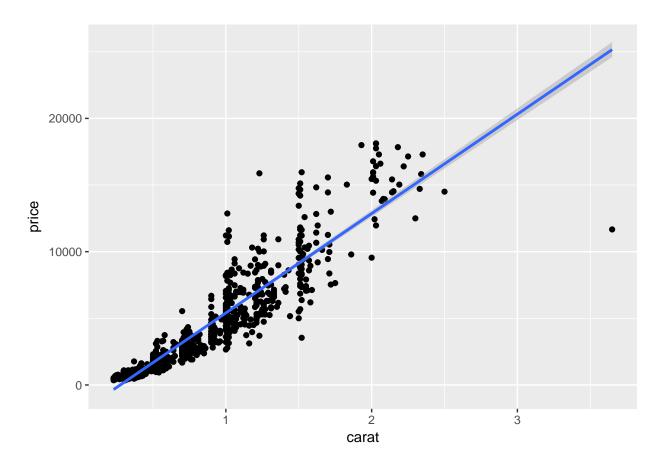


# Colored groups

# Dot Size and Color Relative to Some Values



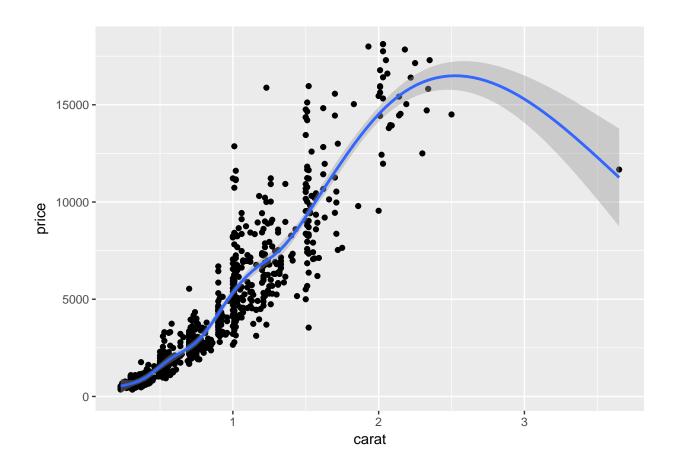
### Regression line



### Local regression curve (loess)

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

## `geom\_smooth()` using method = 'gam'



### ggplot Function

- More important than qplot to access full functionality of ggplot2
- Main arguments
  - data set, usually a data.frame or tibble
  - 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 in ggplot

- Theme settings can be accessed with theme\_get()
- Their settings can be changed with theme()

Example how to 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)</pre>
```

Returns information about data and aesthetic mappings followed by each layer

```
summary(p)
```

Print dots with different sizes and colors

```
bestfit <- geom_smooth(method = "lm", se = F, color = alpha("steelblue", 0.5), size = 2)
p + bestfit # Plot with custom regression line</pre>
```

Syntax to pass on other data sets

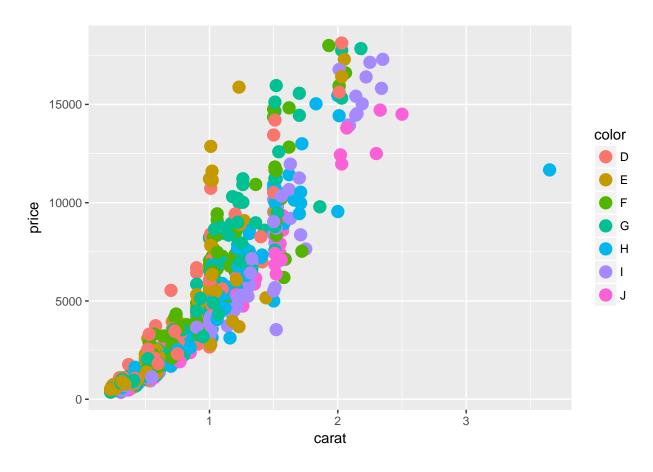
```
p %+% diamonds[sample(nrow(diamonds), 100),]
```

Saves plot stored in variable p to file

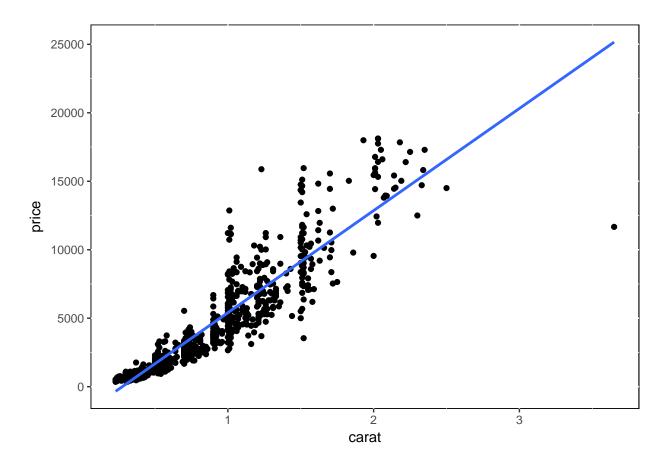
```
ggsave(p, file="myplot.pdf")
```

### ggplot: scatter plots

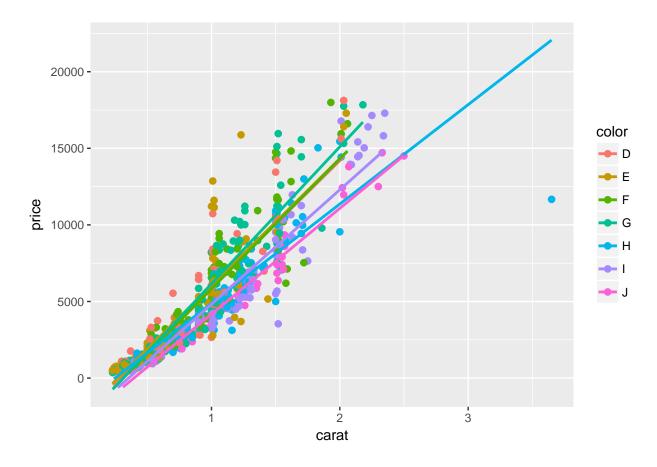
### Basic example



### Regression line



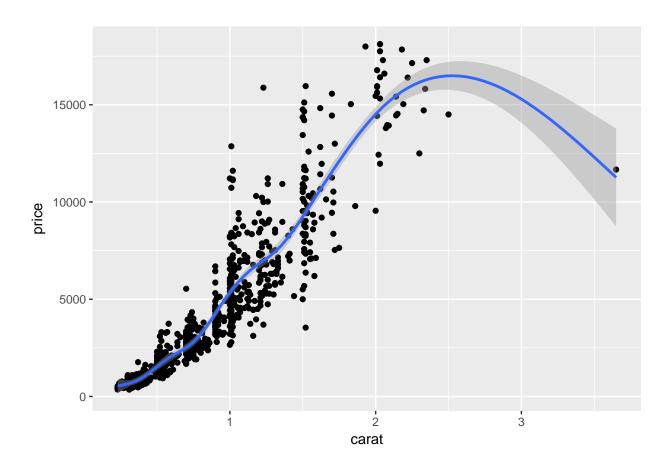
### Several regression lines



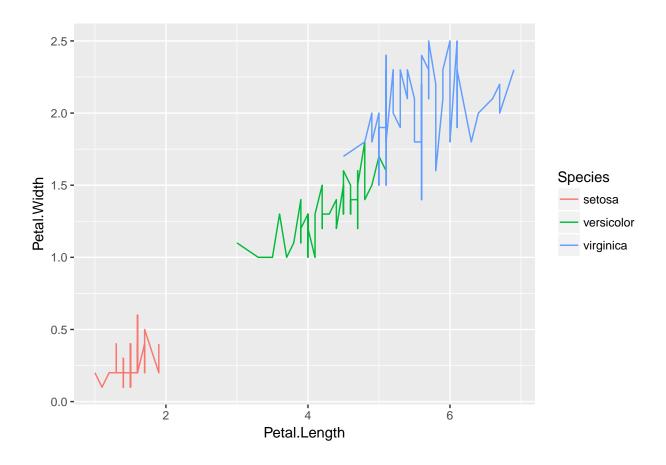
### Local regression curve (loess)

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

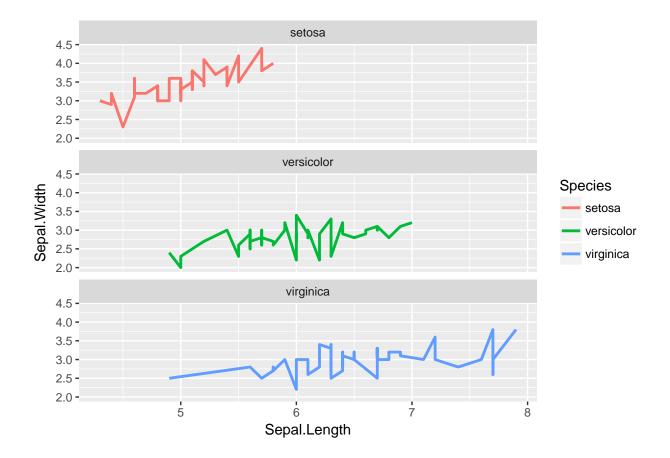
## `geom\_smooth()` using method = 'gam'



# ggplot: line plot



# Faceting



#### Exercise 3

Scatter plots with ggplot2

- 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 and 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.
- Task 3: Generate corresponding line plot with faceting show individual data sets in saparate 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
                           3.2
              4.7
                                         1.3
                                                      0.2
                                                           setosa
## 4
              4.6
                                                      0.2
                           3.1
                                         1.5
                                                           setosa
table(iris$Species)
##
##
       setosa versicolor
                           virginica
##
           50
                       50
                                   50
```

### **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)

Reformat iris_mean with melt

library(reshape2) # Defines melt function df_mean <- melt(iris_mean, id.vars=c("Species"), variable.name = "Samples", value.name="Values")

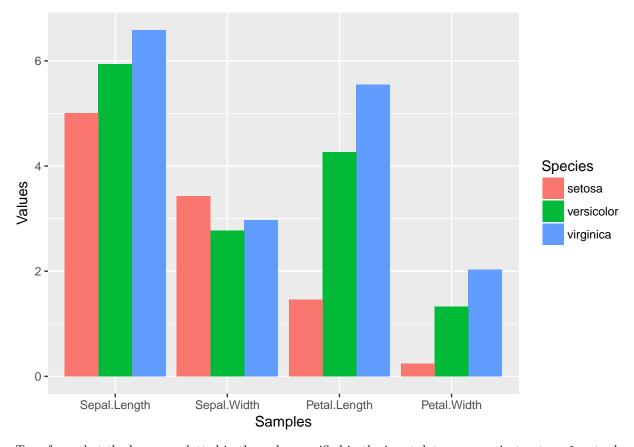
Reformat iris_sd with melt
```

Define standard deviation limits

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

### Verical orientation

df\_sd <- melt(iris\_sd, id.vars=c("Species"), variable.name = "Samples", value.name="Values")</pre>

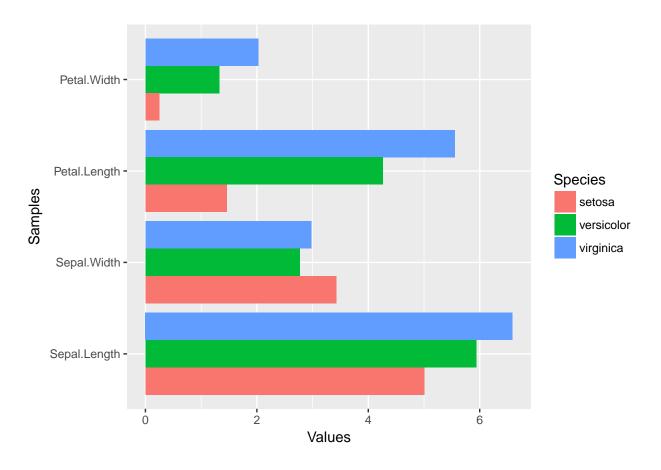


To enforce that the bars are plotted in the order specified in the input data, one can instruct ggplot to do so by turning the corresponding column (here Species) into an ordered factor as follows.

```
df_mean$Species <- factor(df_mean$Species, levels=unique(df_mean$Species), ordered=TRUE)</pre>
```

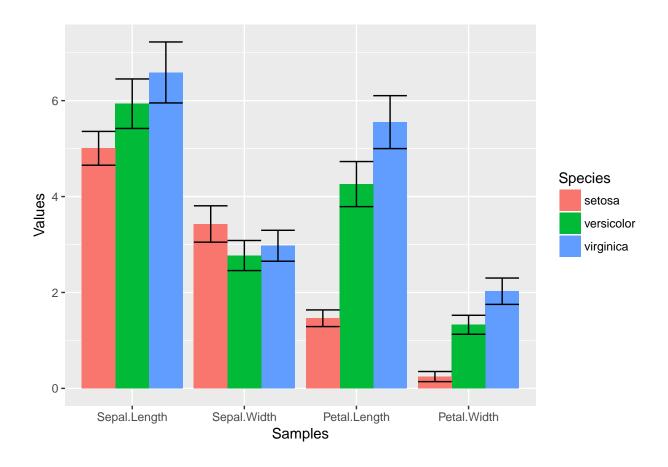
In the above example this is not necessary since ggplot uses this order already.

#### Horizontal orientation

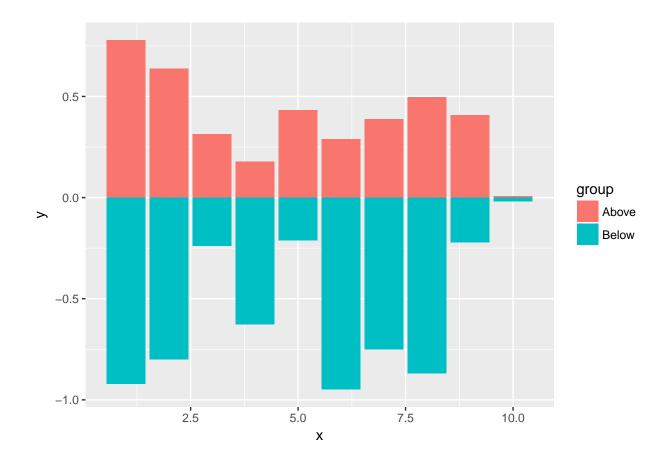


### Faceting

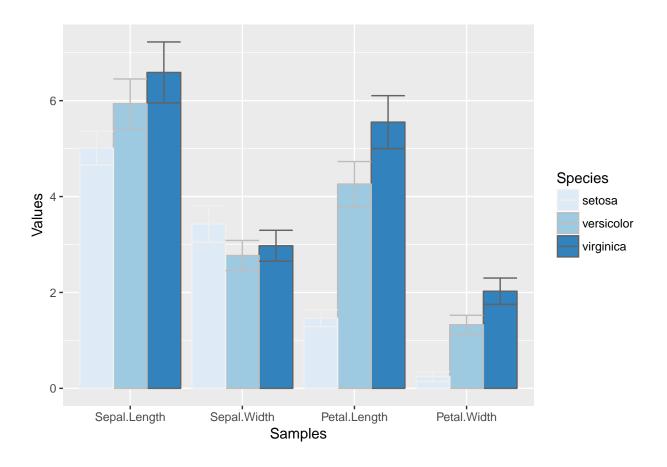
### Error bars



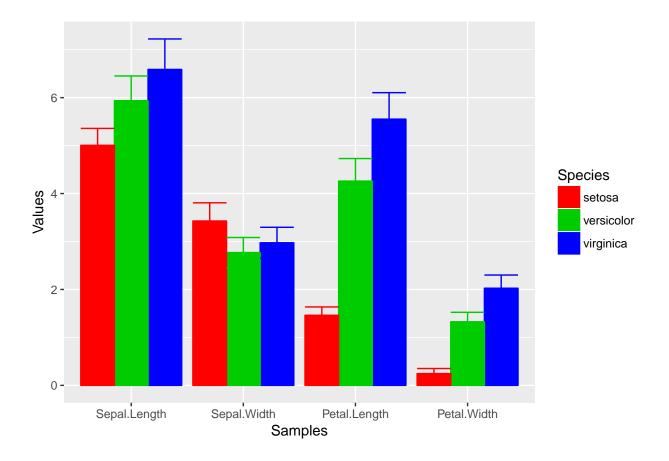
### Mirrored



# **Changing Color Settings**



## Using standard colors



#### 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 data into the expected format
- Task 2: Generate two bar plots: one with stacked bars and one with horizontally arranged bars.

Structure of iris data set

setosa versicolor

50

50

virginica

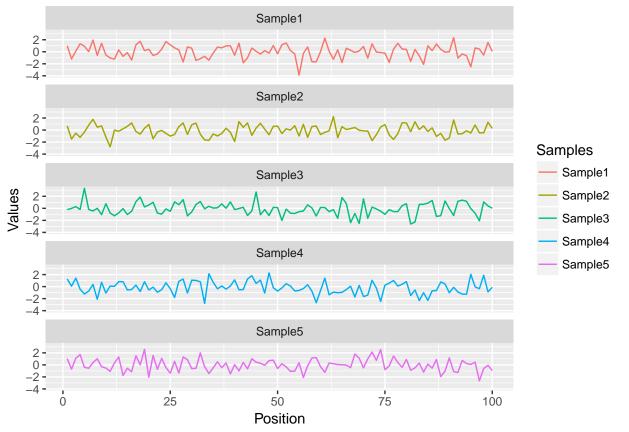
50

```
class(iris)
## [1] "data.frame"
iris[1:4,]
     Sepal.Length Sepal.Width Petal.Length Petal.Width Species
##
## 1
                           3.5
                                         1.4
              5.1
                                                          setosa
## 2
                           3.0
              4.9
                                         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)
##
```

### Data reformatting example

```
Here for line plot
```

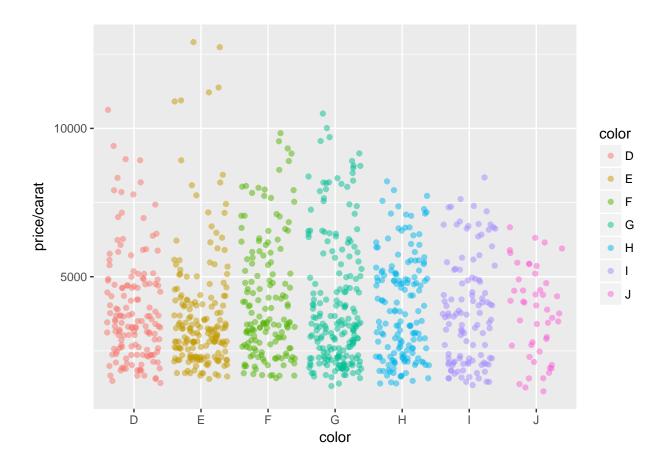
```
y <- matrix(rnorm(500), 100, 5, dimnames=list(paste("g", 1:100, sep=""), paste("Sample", 1:5, sep="")))
y <- data.frame(Position=1:length(y[,1]), y)</pre>
y[1:4, ] # First rows of input format expected by melt()
##
      Position
                  Sample1
                             Sample2
                                          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
             4 1.3294248 -1.2084007 -0.19581898 -0.42361768
## g4
df <- melt(y, id.vars=c("Position"), variable.name = "Samples", value.name="Values")</pre>
p <- ggplot(df, aes(Position, Values)) + geom_line(aes(color=Samples)) + facet_wrap(~Samples, ncol=1)</pre>
print(p)
```



Same data can be represented in box plot as follows

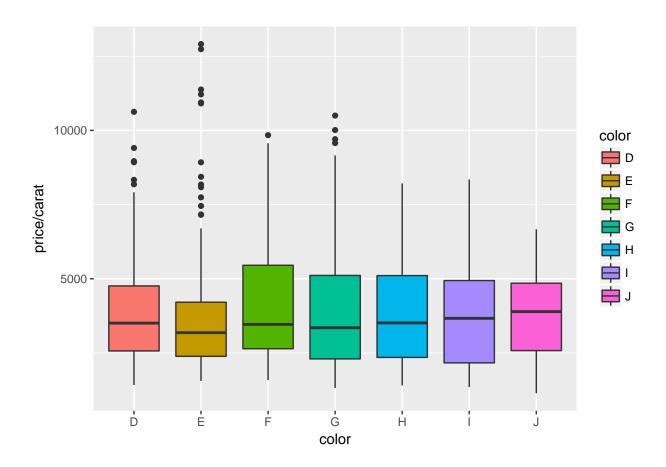
```
ggplot(df, aes(Samples, Values, fill=Samples)) + geom_boxplot()
```

#### Jitter Plots



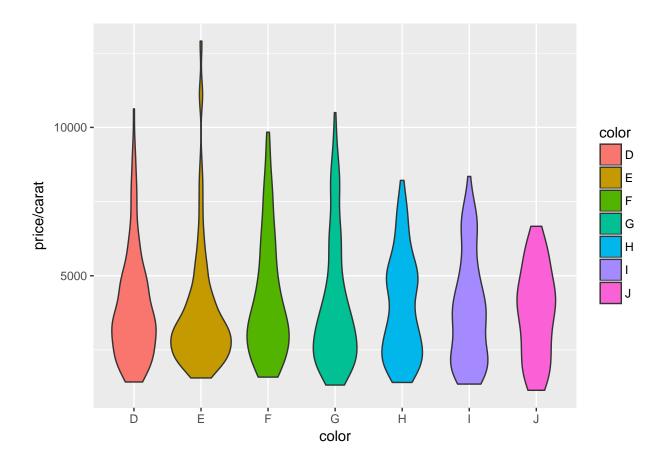
# Box plots

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



# Violin plots

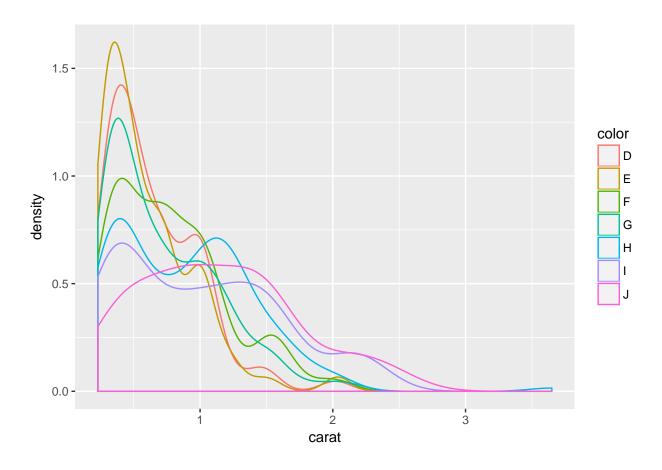
```
p <- ggplot(dsmall, aes(color, price/carat, fill=color)) + geom_violin()
print(p)</pre>
```



# Density plots

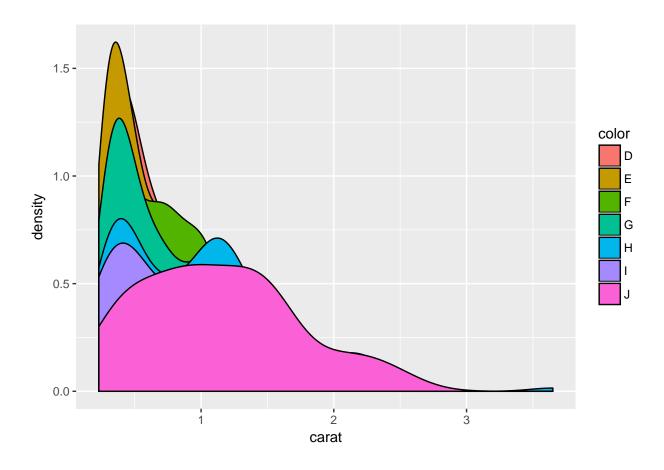
# Line coloring

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

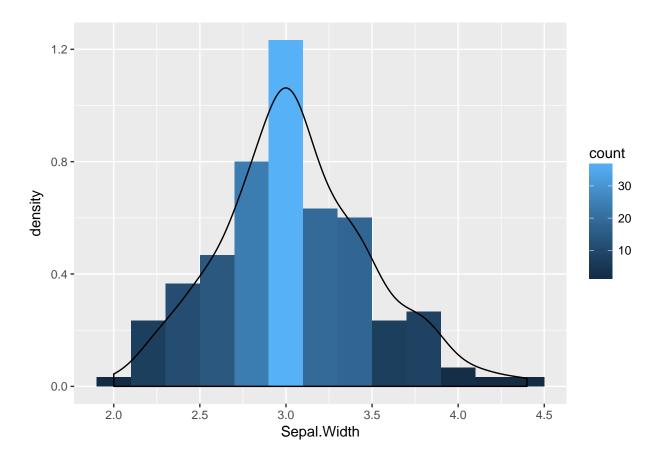


# Area coloring

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



# Histograms



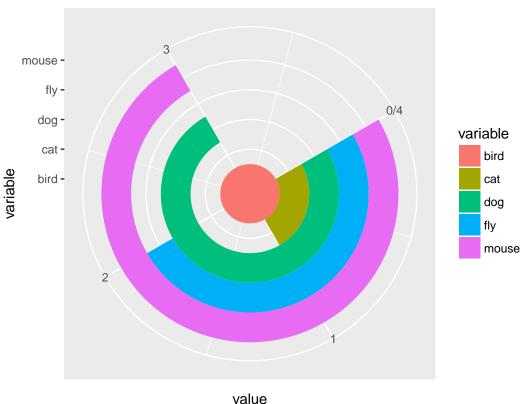
# Pie Chart

# Pie Chart



## Wind Rose Pie Chart

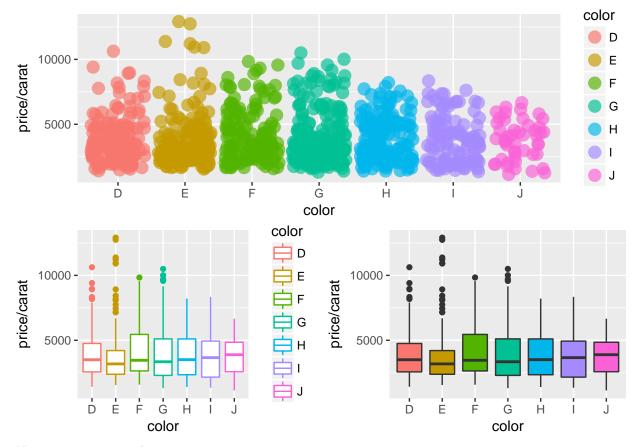
## Pie Chart



## Arranging Graphics on Page

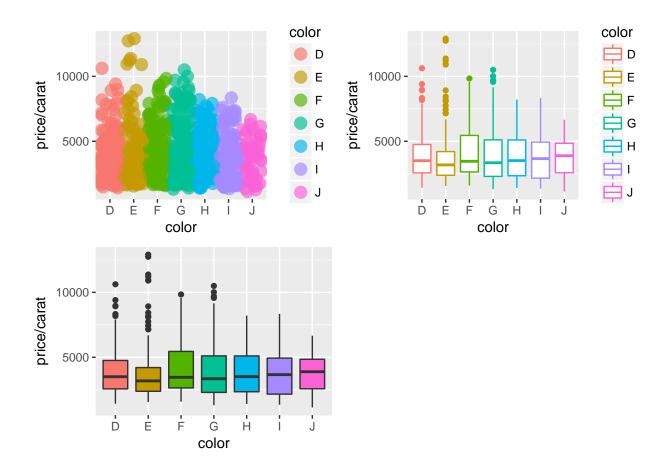
Using grid package

```
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 = "not 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))</pre>
```



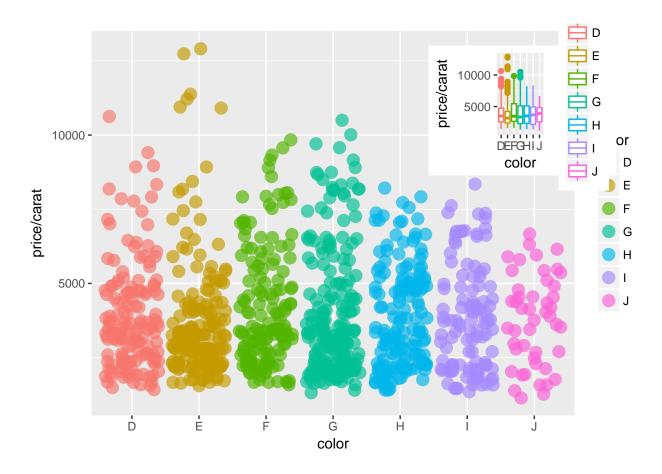
Using gridExtra package

library(gridExtra)
grid.arrange(a, b, c, nrow = 2, ncol=2)



# **Inserting Graphics into Plots**

```
library(grid)
print(a)
print(b, vp=viewport(width=0.3, height=0.3, x=0.8, y=0.8))
```



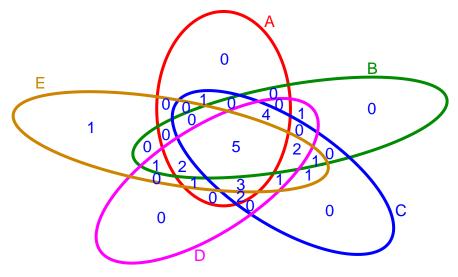
# **Specialty Graphics**

# Venn Diagrams

```
library(systemPipeR)
```

```
## Loading required package: ShortRead
## Loading required package: BiocParallel
##
## Attaching package: 'ShortRead'
## The following object is masked from 'package:ggbio':
##
##
       zoom
## The following object is masked from 'package:ape':
##
##
       zoom
## The following object is masked from 'package:ChemmineR':
##
##
       view
##
```

```
##
## Attaching package: 'systemPipeR'
## The following object is masked from 'package:VariantAnnotation':
##
## reference
setlist5 <- list(A=sample(letters, 18), B=sample(letters, 16), C=sample(letters, 20), D=sample(letters, 0Llist5 <- overLapper(setlist=setlist5, sep="_", type="vennsets")
vennPlot(OLlist5, mymain="", mysub="", colmode=2, ccol=c("blue", "red"))</pre>
```



## Compound Structures

Plots depictions of small molecules with  ${\tt ChemmineR}$  package

```
library(ChemmineR)

## Loading required package: methods
data(sdfsample)
plot(sdfsample[1], print=FALSE)
```

### CMP1

### **ROC Plots**

A variety of libraries are available for plotting receiver operating characteristic (ROC) curves in R:

- ROCR
- ROC
- pROC
- ggplot2

#### Example

Most commonly, in an ROC we plot the true positive rate (y-axis) against the false positive rate (x-axis) at decreasing thresholds. An illustrative example is provided in the ROCR package where one wants to inspect the content of the ROCR.simple object defining the structure of the input data in two vectors.

```
# install.packages("ROCR") # Install if necessary on your laptop
library(ROCR)
data(ROCR.simple)
ROCR.simple
```

```
## $predictions
##
              [1] 0.612547843 0.364270971 0.432136142 0.140291078 0.384895941 0.244415489 0.970641299
              [8] 0.890172812 0.781781371 0.868751832 0.716680598 0.360168796 0.547983407 0.385240464
##
##
           [15] \quad 0.423739359 \quad 0.101699993 \quad 0.628095575 \quad 0.744769966 \quad 0.657732644 \quad 0.490119891 \quad 0.072369921 \quad 0.07236
##
           [22] 0.172741714 0.105722115 0.890078186 0.945548941 0.984667270 0.360180429 0.448687336
##
            [29] \ \ 0.014823599 \ \ 0.543533783 \ \ 0.292368449 \ \ 0.701561487 \ \ 0.715459280 \ \ 0.714985914 \ \ 0.120604738 
##
           ##
##
            [50] \ \ 0.876086217 \ \ 0.353281048 \ \ 0.212014560 \ \ 0.703293499 \ \ 0.689075677 \ \ 0.627012496 \ \ 0.240911145 
           [57] 0.402801992 0.134794140 0.120473353 0.665444679 0.536339509 0.623494622 0.885179651
##
           [64] 0.353777439 0.408939895 0.265686095 0.932159806 0.248500489 0.858876675 0.491735594
##
            [71] \quad 0.151350957 \quad 0.694457482 \quad 0.496513160 \quad 0.123504905 \quad 0.499788081 \quad 0.310718619 \quad 0.907651100 
           ##
           ##
```

```
[99] 0.212404891 0.930846938 0.083048377 0.468610247 0.393378108 0.663367560 0.349540913
## [106] 0.194398425 0.844415442 0.959417835 0.211378771 0.943432189 0.598162949 0.834803976
## [113] 0.576836208 0.380396459 0.161874325 0.912325837 0.642933593 0.392173971 0.122284044
## [120] 0.586857799 0.180631658 0.085993218 0.700501359 0.060413627 0.531464015 0.084254795
## [127] 0.448484671 0.938583020 0.531006532 0.785213140 0.905121019 0.748438143 0.605235403
## [134] 0.842974300 0.835981859 0.364288579 0.492596896 0.488179708 0.259278968 0.991096434
## [141] 0.757364019 0.288258273 0.773336236 0.040906997 0.110241034 0.760726142 0.984599159
## [148] 0.253271061 0.697235328 0.620501132 0.814586047 0.300973098 0.378092079 0.016694412
## [155] 0.698826511 0.658692553 0.470206008 0.501489336 0.239143340 0.050999138 0.088450984
  [162] 0.107031842 0.746588080 0.480100183 0.336592126 0.579511087 0.118555284 0.233160827
## [169] 0.461150807 0.370549294 0.770178504 0.537336015 0.463227453 0.790240205 0.883431431
## [176] 0.745110673 0.007746305 0.012653524 0.868331219 0.439399995 0.540221346 0.567043171
## [183] 0.035815400 0.806543942 0.248707470 0.696702150 0.081439129 0.336315317 0.126480399
## [190] 0.636728451 0.030235062 0.268138293 0.983494405 0.728536415 0.739554341 0.522384507
## [197] 0.858970526 0.383807972 0.606960209 0.138387070
##
## $labels
##
    [1] 1 1 0 0 0 1 1 1 1 1 0 1 0 1 0 1 0 0 0 1 1 1 1 0 0 0 0 1 0 1 0 1 1 0 1 1 1 0 0 1 1 1 0 1 0 1 0 1 0 1 0
   ## [189] 0 0 0 1 0 1 1 0 1 0 1 0
pred <- prediction(ROCR.simple$predictions, ROCR.simple$labels)</pre>
perf <- performance( pred, "tpr", "fpr" )</pre>
plot(perf)
     \infty
     o.
True positive rate
     9
     o.
     0.4
     \sim
     Ö
     0
                      0.2
          0.0
                                  0.4
                                              0.6
                                                          8.0
                                                                      1.0
                                 False positive rate
```

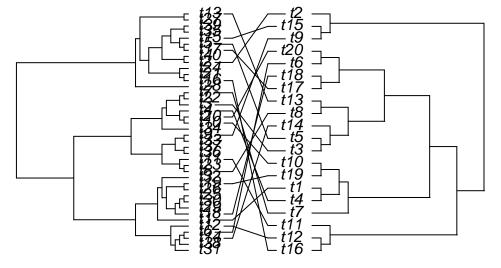
```
Obtain area under the curve (AUC)
```

```
auc <- performance( pred, "tpr", "fpr", measure = "auc")
auc@y.values[[1]]</pre>
```

## [1] 0.8341875

#### Trees

The ape package provides many useful utilities for phylogenetic analysis and tree plotting. Another useful package for plotting trees is ggtree. The following example plots two trees face to face with links to identical leaf labels.



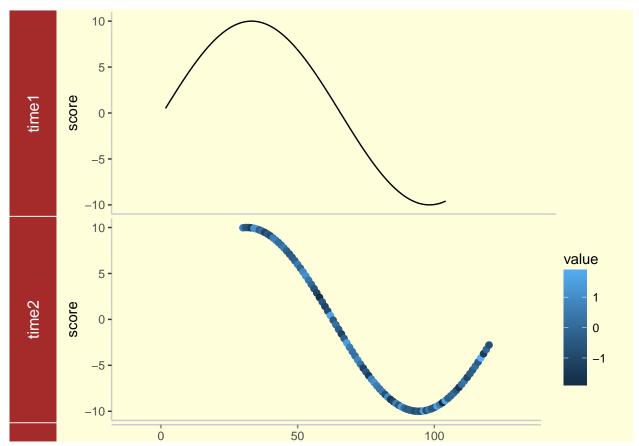
# Genome Graphics

#### ggbio

- What is ggbio?
  - A programmable genome browser environment
- Genome broswer concepts
  - A genome browser is a visualization tool for plotting different types of genomic data in separate tracks along chromosomes.
  - The ggbio package (Yin, Cook, and Lawrence 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 here.
  - 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 serval new components useful for genomic data visualization. A detailed list is given in Table 1.2 of the vignette here.
  - Useful web sites: ggbio manual
    - \* ggbio functions
    - \* autoplot demo

#### 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()</pre>
```



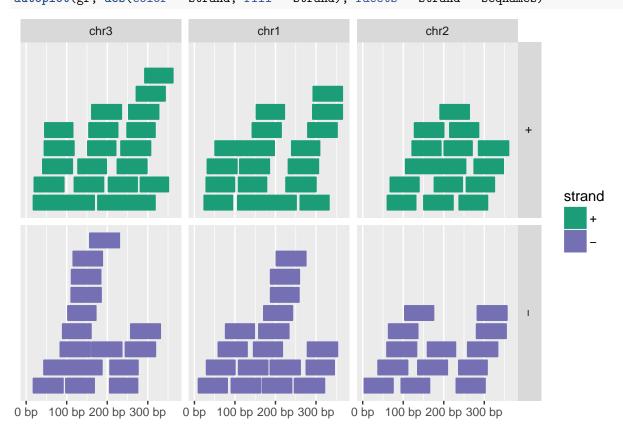
#### Plotting genomic ranges

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

```
library(GenomicRanges)
```

```
## Loading required package: stats4
## Loading required package: S4Vectors
##
## Attaching package: 'S4Vectors'
## The following object is masked from 'package:ChemmineR':
##
## fold
```

```
##
## expand.grid
## Loading required package: IRanges
## Loading required package: GenomeInfoDb
set.seed(1); N <- 100; gr <- GRanges(seqnames = sample(c("chr1", "chr2", "chr3"), size = N, replace = T
autoplot(gr, aes(color = strand, fill = strand), facets = strand ~ seqnames)</pre>
```

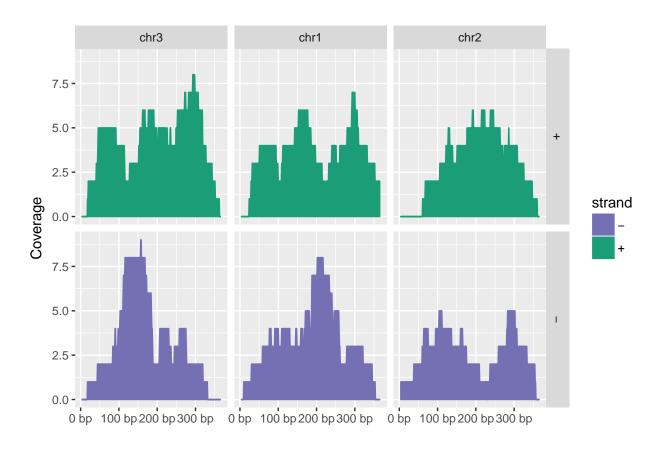


## The following object is masked from 'package:base':

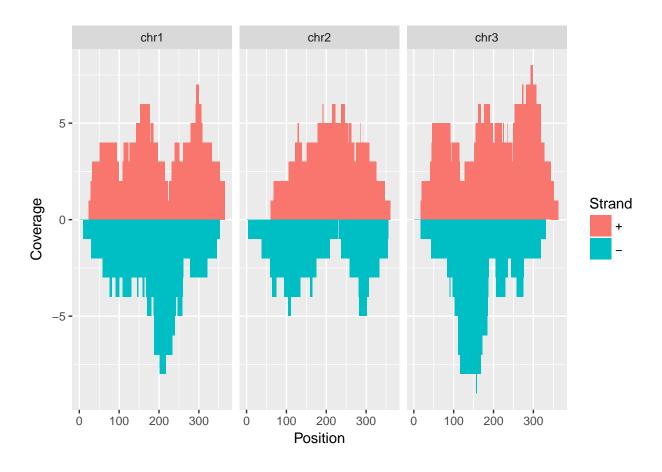
### Plotting coverage

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

## Scale for 'x' is already present. Adding another scale for 'x', which will replace the existing ## scale.

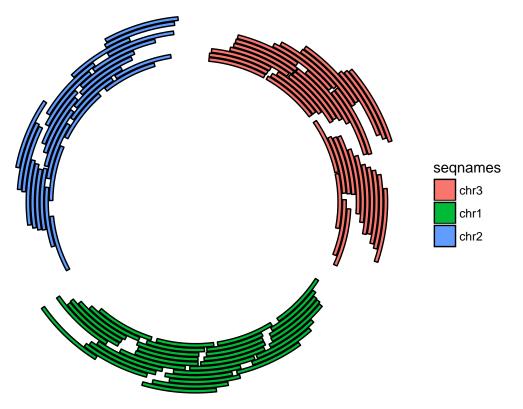


### Mirrored coverage



# Circular genome plots

```
ggplot(gr) + layout_circle(aes(fill = seqnames), geom = "rect")
```

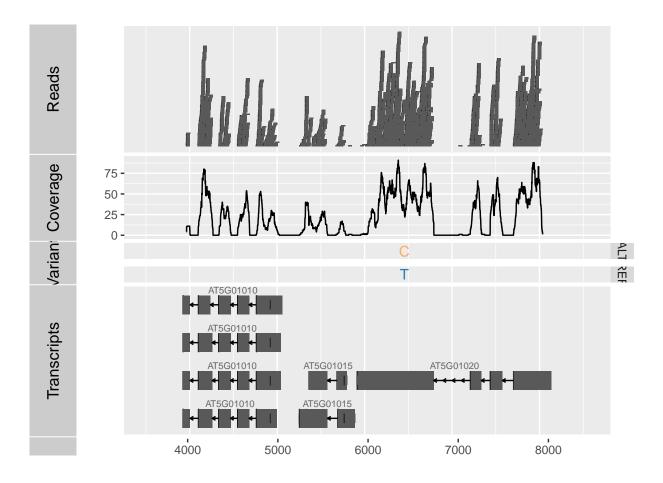


More complex circular example

### Alignments and variants

To make the following example work, please download and unpack this data archive containing GFF, BAM and VCF sample files.

```
library(rtracklayer); library(GenomicFeatures); library(Rsamtools); library(GenomicAlignments); library
ga <- readGAlignments("./data/SRR064167.fastq.bam", use.names=TRUE, param=ScanBamParam(which=GRanges("County parametry))
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 <- makeTxDbFromGFF(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("")</pre>
```



## Additional examples

See autoplot demo here

### Additional genome graphics

- Gviz
- RCircos (Zhang, Meltzer, and Davis 2013)
- Genome Graphs
- genoPlotR

### Genome Browser: IGV

View genome data in IGV

- Download and open IGV
- Select in menu in top left corner A. thaliana (TAIR10)
- Upload the following indexed/sorted Bam files with File -> Load from URL...

http://faculty.ucr.edu/~tgirke/HTML\_Presentations/Manuals/Workshop\_Dec\_6\_10\_2012/Rrnaseq/results/SRR064http://faculty.ucr.edu/~tgirke/HTML\_Presentations/Manuals/Workshop\_Dec\_6\_10\_2012/Rrnaseq/results/SRR064http://faculty.ucr.edu/~tgirke/HTML\_Presentations/Manuals/Workshop\_Dec\_6\_10\_2012/Rrnaseq/results/SRR064http://faculty.ucr.edu/~tgirke/HTML\_Presentations/Manuals/Workshop\_Dec\_6\_10\_2012/Rrnaseq/results/SRR064http://faculty.ucr.edu/~tgirke/HTML\_Presentations/Manuals/Workshop\_Dec\_6\_10\_2012/Rrnaseq/results/SRR064

• To view area of interest, enter its coordinates Chr1:49,457-51,457 in position menu on top.

#### Create symbolic links

For viewing BAM files in IGV as part of systemPipeR workflows.

• systemPipeR: utilities for building NGS analysis pipelines

#### Controlling IGV from R

Open IGV before running the following routine. Alternatively, open IGV from within R with startIGV("lm") . Note this may not work on all systems.

## References

Yin, T, D Cook, and M Lawrence. 2012. "Ggbio: An R Package for Extending the Grammar of Graphics for Genomic Data." *Genome Biol.* 13 (8). doi:10.1186/gb-2012-13-8-r77.

Zhang, H, P Meltzer, and S Davis. 2013. "RCircos: An R Package for Circos 2d Track Plots." BMC Bioinformatics 14: 244–44. doi:10.1186/1471-2105-14-244.