Class 5: Data Viisualization with ggplot

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Our first plot

Our first plot

R has base graphic

```
#ggplot(cars)
```

How would I plot this with 'ggplot'? NO! We need to install package first We use "install_package()"

```
# install.packages("ggplot2")
```

Before I can use this package I need to load it with a "library()" call

```
library(ggplot2)
```

Every ggplot needs at least three layers"

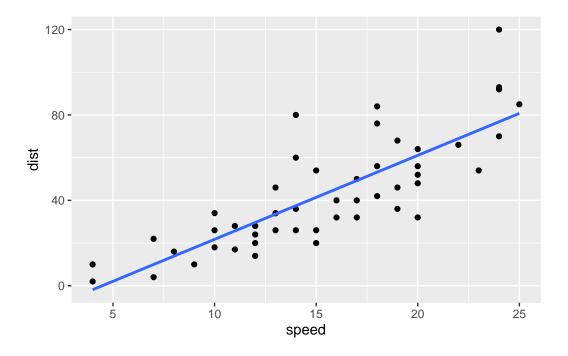
- **Data** (i.e. the data.frame we have)
- Aes (the aesthetic mapping of our data to what we want to plot)
- Geoms (How we want to plot this stuff!)

add a straight line without SE

```
ggplot(data=cars)+
  aes(x=speed, y=dist)+
  geom_point()+
```

```
geom_smooth(se=FALSE, method="lm")
```

`geom_smooth()` using formula 'y ~ x'

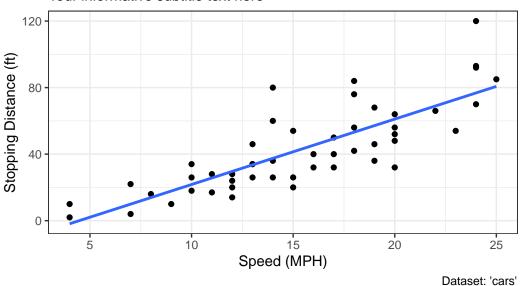


add lable and change background

[`]geom_smooth()` using formula 'y ~ x'

Speed and Stopping Distances of Cars

Your informative subtitle text here



url <- "https://bioboot.github.io/bimm143_S20/class-material/up_down_expression.txt"
genes <- read.delim(url)
head(genes)</pre>

```
Gene Condition1 Condition2 State
1 A4GNT -3.6808610 -3.4401355 unchanging
2 AAAS 4.5479580 4.3864126 unchanging
3 AASDH 3.7190695 3.4787276 unchanging
4 AATF 5.0784720 5.0151916 unchanging
5 AATK 0.4711421 0.5598642 unchanging
6 AB015752.4 -3.6808610 -3.5921390 unchanging
```

Q. Use the nrow() function to find out how many genes are in this dataset. What is your nrow(genes)

[1] 5196

Q. Use the colnames() function and the ncol() function on the genes data frame to find colnames(genes)

```
[1] "Gene" "Condition1" "Condition2" "State"
```

#Q. Use the table() function on the State column of this data.frame to find out how many 'ncol(genes)

[1] 4

```
table(genes$State)
```

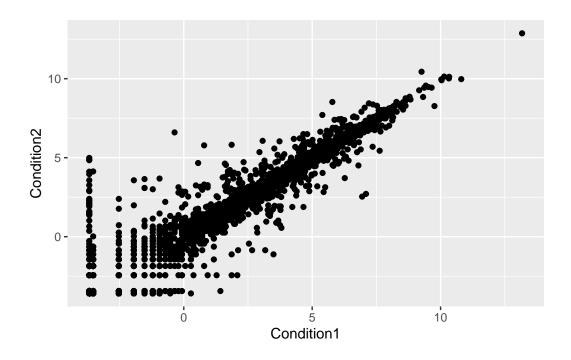
```
down unchanging up
72 4997 127
```

Using your values above and 2 significant figures. What fraction of total genes is up-reround(table(genes\$State)/nrow(genes)*100,2)

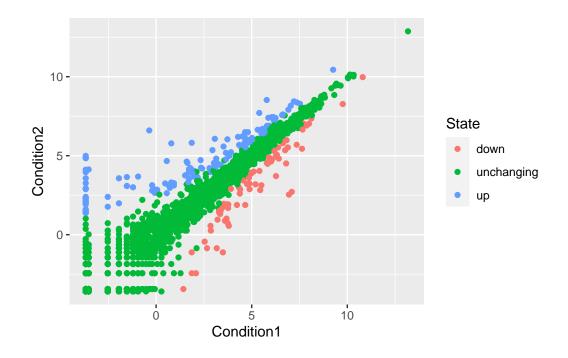
```
down unchanging up 1.39 96.17 2.44
```

There are 'r nrow(genes)' genes in this data set

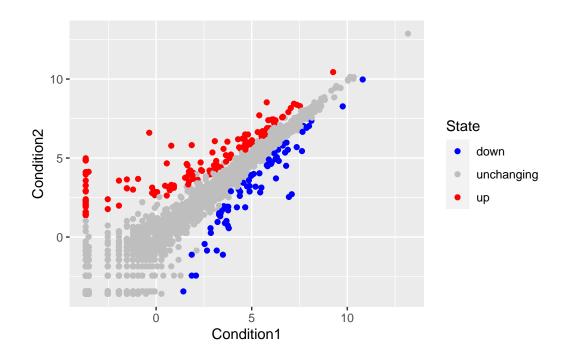
```
ggplot(genes) +
   aes(x=Condition1, y=Condition2) +
   geom_point()
```



```
p <- ggplot(genes) +
    aes(x=Condition1, y=Condition2, col=State) +
    geom_point()
p</pre>
```

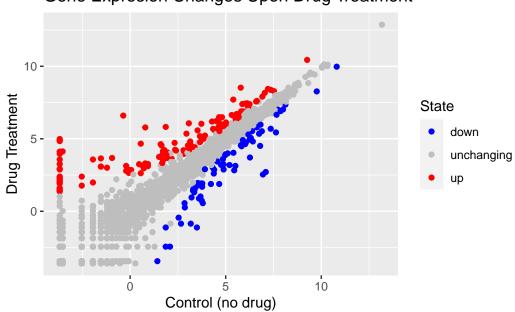


p + scale_colour_manual(values=c("blue", "gray", "red"))



```
#Q. Nice, now add some plot annotations to the p object with the labs() function so your p
p + scale_colour_manual( values=c("blue","gray","red") )+
labs(title="Gene Expression Changes Upon Drug Treatment", x="Control (no drug)", y="Drug")
```

Gene Expresion Changes Upon Drug Treatment



```
# File location online
url <- "https://raw.githubusercontent.com/jennybc/gapminder/master/inst/extdata/gapminder.
gapminder <- read.delim(url)

# install.packages("dplyr") ## un-comment to install if needed</pre>
```

Attaching package: 'dplyr'

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

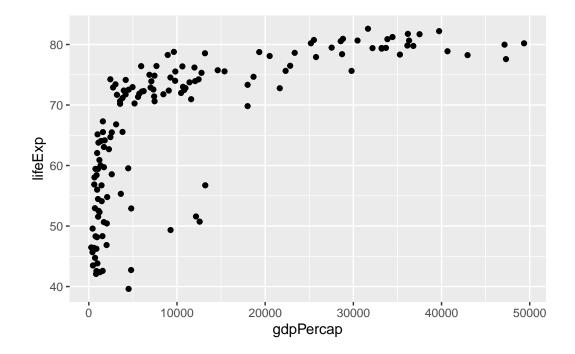
filter, lag

library(dplyr)

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

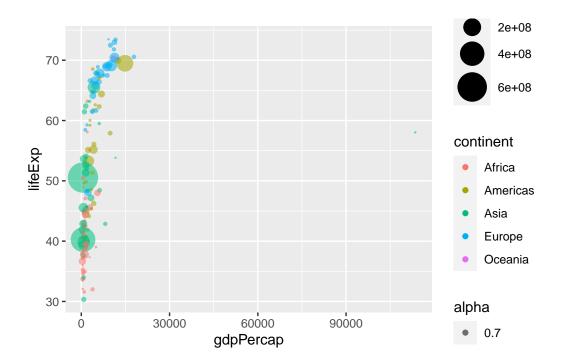
intersect, setdiff, setequal, union

```
gapminder_2007 <- gapminder %>%
  filter(year==2007)
ggplot(gapminder_2007) +
  aes(x=gdpPercap, y=lifeExp) +
  geom_point()
```



Q. Can you adapt the code you have learned thus far to reproduce our gapminder scatter plot for the year 1957? What do you notice about this plot is it easy to compare with the one for 2007?

```
gapminder_1957 <- gapminder %>%
  filter(year==1957)
ggplot(gapminder_1957) +
  aes(x = gdpPercap, y = lifeExp, color=continent, size = pop, alpha=0.7) +
  geom_point() +
  scale_size_area(max_size = 10)
```



Q. Do the same steps above but include 1957 and 2007 in your input dataset for ggplot(). You should now include the layer facet_wrap(~year) to produce the following plot:

```
gapminder_1957 <- gapminder %>%
  filter(year==1957|year==2007)
ggplot(gapminder_1957) +
  aes(x = gdpPercap, y = lifeExp, color=continent, size = pop, alpha=0.7) +
  geom_point() +
  scale_size_area(max_size = 10) +
  facet_wrap(~year)
```

