

# Business Analytics Using Computational Statistics

Week 6  
Permutation Tests

Week 7  
Multigroup Comparisons

Week 8  
No Class!

## Visualizing Normality

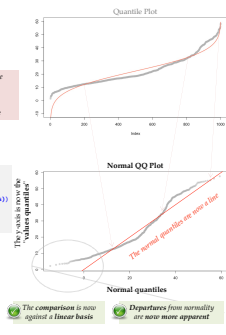
### Normal Quantile-Quantile Plot

For each normal quantile, what is quantile of our values?

Differences in the plots are difficult to distinguish  
Both curves change at the same time

```
quantile_quantile_plot <- function(values) {  
  probs1000 <- seq(0, 1, 0.001)  
  q_vals <- quantile(values, probs1000)  
  q_norm <- qnorm(probs1000, mean=mean(values), sd=sd(values))  
  plot(q_norm, q_vals, lty=2)  
  abline(a=0, b=1, col="red", lwd=2)  
}
```

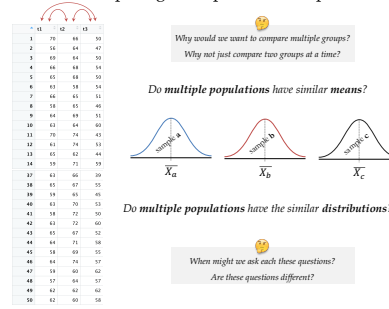
Compare to R's built-in Normal QQ plot functions:  
qqnorm(d123) # Shows normal qq plot  
qqline(d123) # Shows diagonal normal line



The comparison is now against a linear basis  
Departures from normality are now more apparent

## Multigroup Tests

### Comparing Multiple (3+) Groups



## Posthoc Tests

### Multigroup Post-Hoc Tests

	1	2	3
1	55	58	58
2	56	64	47
3	69	64	59
4	66	68	54

It is tempting to use multiple two-sample Tests Post-hoc

ANOVA → 3 t-tests

```
t.test(adverts$t1, adverts$t2)  
t.test(adverts$t1, adverts$t3)  
t.test(adverts$t2, adverts$t3)
```

Kruskal-Wallis → 3 Wilcoxon Tests

```
wilcox.test(adverts$t1, adverts$t2)  
wilcox.test(adverts$t1, adverts$t3)  
wilcox.test(adverts$t2, adverts$t3)
```

But do not use two-sample methods to test two-way differences between multiple groups!

# Picking Packages

## TidyR

Well maintained and updated

Simpler Syntax

Used in many books/tutorials



Part of larger “**tidy**” universe  
*(might be better for large analytics project)*



*There's a **lot of decisions** that should go into  
adopting a package for your work.*

*Consider the factors that are **important for you***

***There are no right or wrong choices**  
(you may learn more by switching between packages)*

```
library(tidyR)  
verizon <- read.csv("verizon_wide.csv")
```

```
verizon_long <- gather(verizon_wide, na.rm = TRUE, key = "provider", value = "time")  
providers <- split(x = verizon_long$time, f = verizon_long$provider)
```

## Reshape2



Seems to be “superseded” by TidyR  
*(will it stop being maintained one day?)*



Works on more than data.frames

Still found in online tutorials

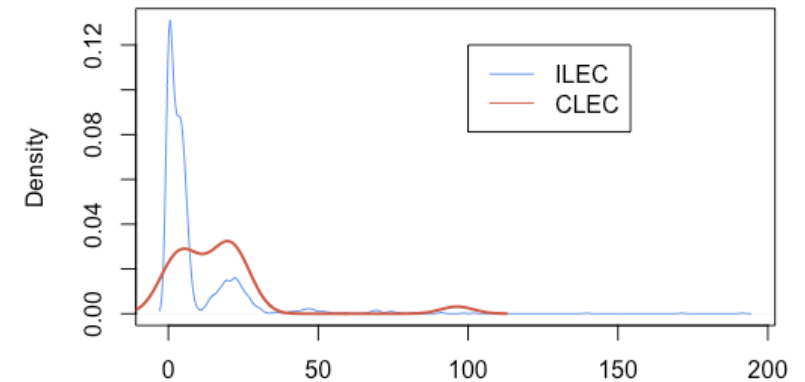


Custom built for reshaping  
*(might be better for using in your own packages)*

# t-Test of Sample Means

## Assumptions of Two-Sample t-Tests:

1. Ratio/Interval scale (continuous numbers)
2. Normal distribution of data
3. Homogeneity of variance (constant variance)



## Student's t-Test assuming equal variances

```
t.test(providers$CLEC, providers$ILEC, alt="greater", var.equal=TRUE, conf.level = 0.99)
```

```
# t = 2.6125, df = 1685, p-value = 0.004534
```

*Reject  $H_{null}$*



*Assumptions about variances are important:  
we arrive at contrary conclusions at  $\alpha = 0.01$*

## Welch's t-Test assuming unequal variances

```
t.test(providers$CLEC, providers$ILEC, alt="greater", var.equal=FALSE, conf.level = 0.99)
```

```
# t = 1.9834, df = 22.346, p-value = 0.02987
```

*Cannot Reject  $H_{null}$*



*We must also confirm normality assumption*

# Assessing Normality

*How can we visually compare distributions?*

## Density Plots

We can plot the distributions of our data against perfect normal

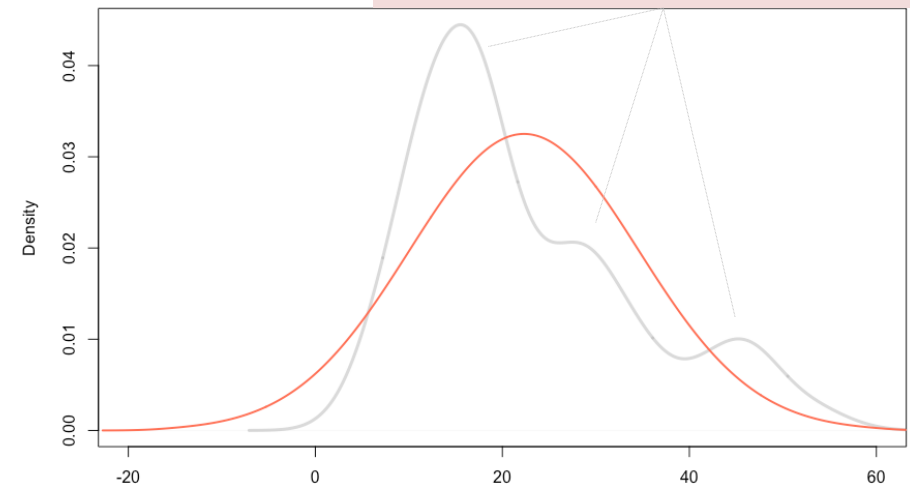
*Let's take 1000 quantiles from a normal distribution with same central tendency and dispersion as our original data*

```
probs1000 <- seq(0, 1, 0.001)
qnorm1000 <- qnorm(probs1000, mean=mean(d123), sd=sd(d123))
```

```
plot(density(d123), main=NA, xlab=NA)
lines(density(qnorm1000))
```



*Our data has many distracting features that make comparisons hard*



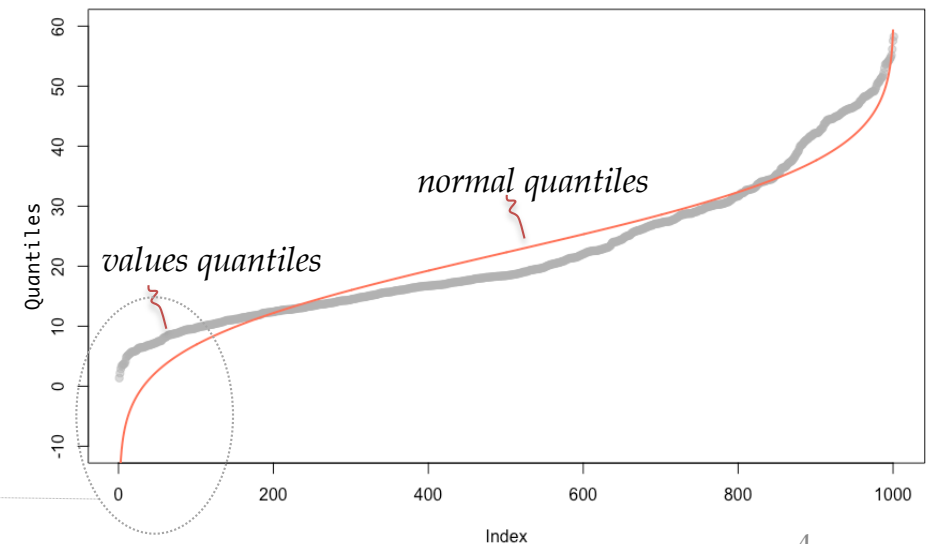
## Quantile Plots

We can plot the quantiles of our data against perfect normal

```
plot(quantile(d123, probs=probs1000))
lines(qnorm1000)
```



*Removes many of the features that make comparisons difficult*



# Normal Quantile-Quantile Plot

For each normal quantile,  
what is quantile of our values?



*Differences in the plots are  
difficult to distinguish*

*Both curves  
change at the same time*

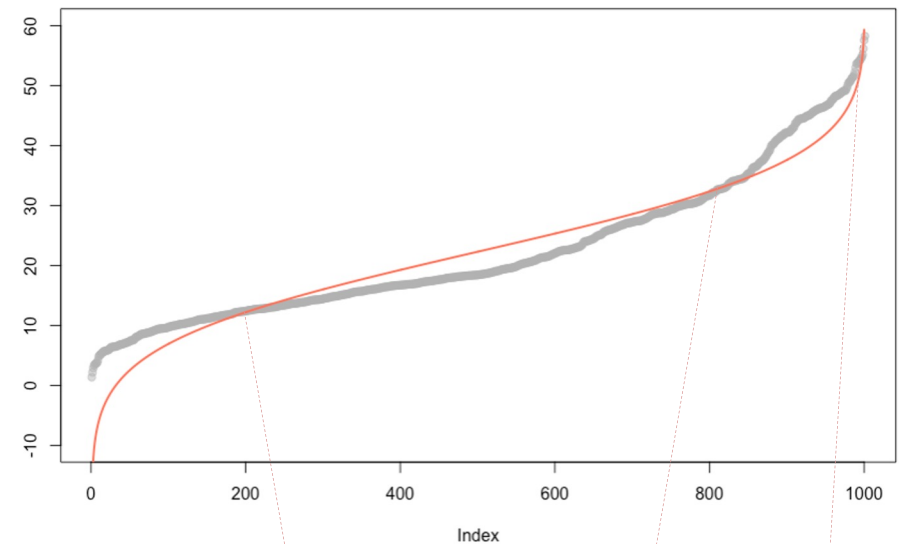
```
quantile_quantile_plot <- function(values) {  
  probs1000 <- seq(0, 1, 0.001)  
  q_vals <- quantile(values, probs1000)  
  q_norm <- qnorm(probs1000, mean=mean(values), sd=sd(values))  
  plot(q_norm, q_vals, ...)  
  abline(a=0, b=1, col="red", lwd=2)  
}
```

`quantile_quantile_plot(d123)`

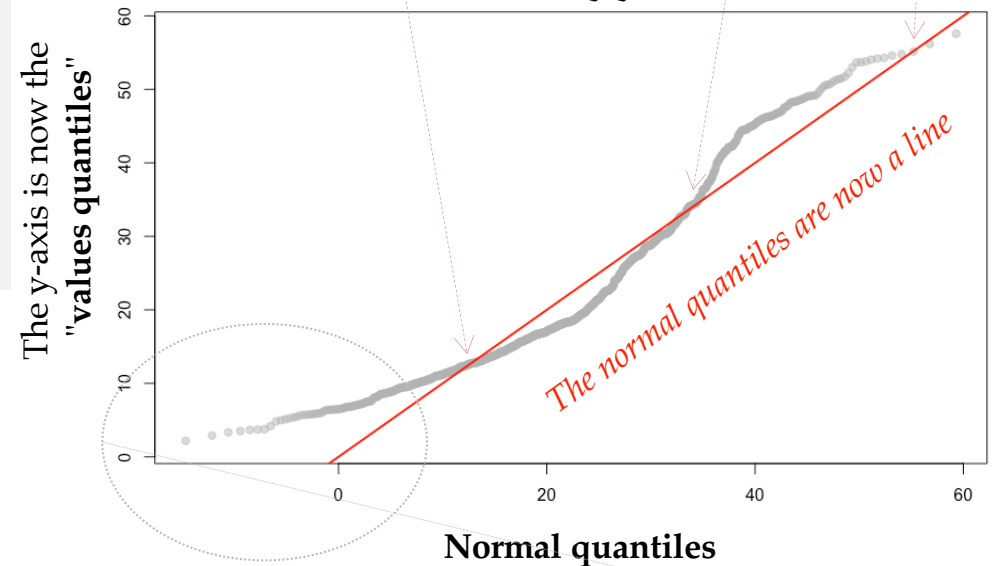
Compare to R's built-in Normal QQ plot functions:

```
qqnorm(d123) # Draws normal qq plot  
qqline(d123) # Draws diagonal normal line
```

Quantile Plot



Normal QQ Plot



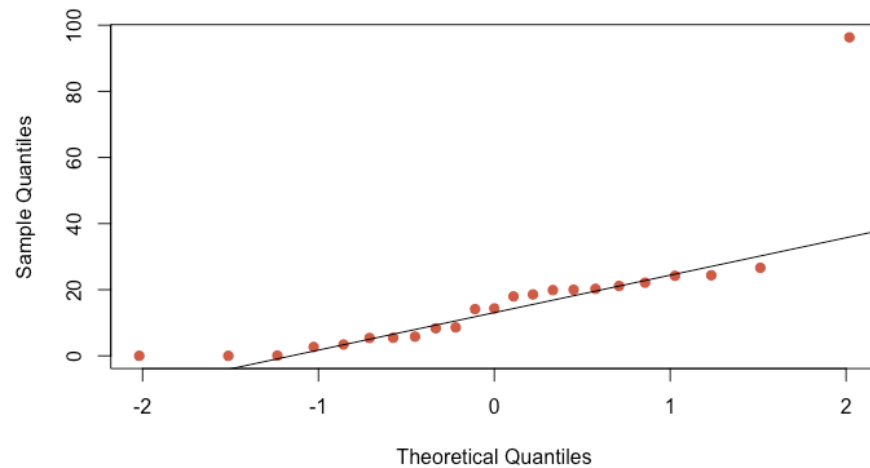
The **comparison** is now  
against a **linear basis**



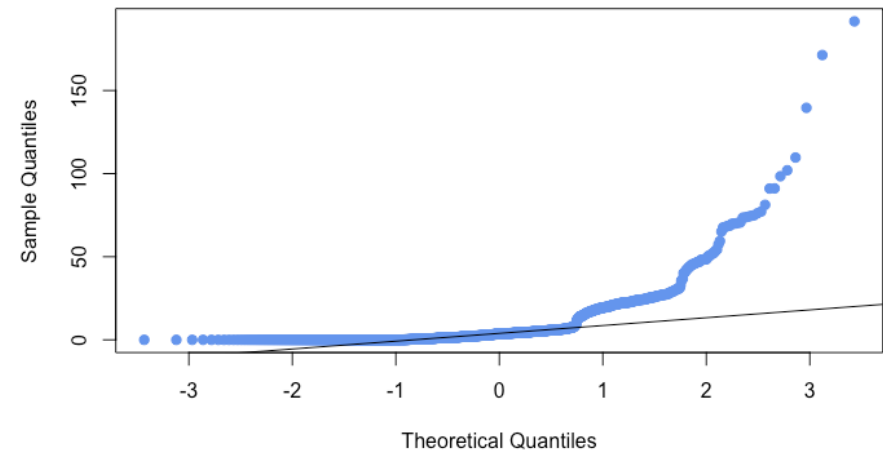
**Departures from normality**  
are now **more apparent**

## Does our data match assumptions of normality?

```
qqnorm(providers$CLEC, col="coral3", pch=19)  
qqline(providers$CLEC)
```



```
qqnorm(providers$ILEC, col="cornflowerblue", pch=19)  
qqline(providers$ILEC)
```



*We should likely prefer our **non-parametric tests***

*You now know exactly how Normal-QQ plots work,  
**But use the built-in functions qqnorm(), qqline() in your work***

# Permutation Tests of Sample Means

```
observed_diff <- mean(providers$CLEC) - mean(providers$ILEC)
```

```
# Simulate Permuted differences
```

```
permute_diff <- function(values, groups) {  
  permuted <- sample(values, replace = FALSE)  
  grouped <- split(permuted, groups)  
  mean(grouped$CLEC) - mean(grouped$ILEC)  
}
```

```
nperms <- 10000
```

```
set.seed(895702709)
```

```
permuted_diffs <- replicate(nperms, permute_diff(verizon_long$time, verizon_long$provider))
```

```
# Visualize permuted differences
```

```
hist(permuted_diffs, breaks = "fd", probability = TRUE, ...)
```

```
lines(density(permuted_diffs), lwd=2)
```

```
abline(v=observed_diff, col="coral3", lwd=3)
```

```
# Significance of permuted differences
```

```
p_1tailed <- sum(permuted_diffs > observed_diff) / nperm
```

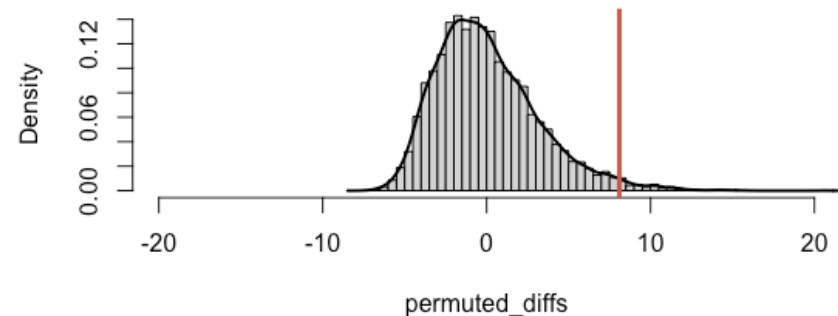
```
# [1] 0.0186
```

*Cannot Reject  $H_{null}$*



*Assumption free conclusion: we cannot reject  $H_{null}$*

Null Distribution of Differences



*Would two-tailed p-value be different?*

# Wilcoxon Test of Samples

```
gt_eq <- function(a, b) {  
  ifelse(a > b, 1, 0) + ifelse(a == b, 0.5, 0)  
}
```

```
W <- sum(outer(providers$CLEC, providers$ILEC, FUN = gt_eq))  
# [1] 26820
```

```
n1 = length(providers$CLEC) # 23  
n2 = length(providers$ILEC) # 1664
```

```
wilcox_p_1tail <- 1 - pwilcox(W, n1, n2) # 0.0003688341  
wilcox_p_2tail <- 2 * wilcox_p_1tail    # 0.0007376683
```

```
wilcox.test(providers$CLEC, providers$ILEC, alternative = "greater", paired = FALSE)  
W = 26820, p-value = 0.0004565
```

*Reject  $H_{null}$*



***Assumption free conclusion:***  
*Values in the two samples are distinct  
(values seem shifted)*

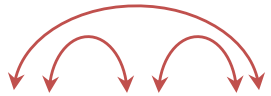
***Mean of samples are not distinguishable***

***BUT***

***CLEC-vs-ILEC customers seem to be treated differently***



# Comparing Multiple (3+) Groups



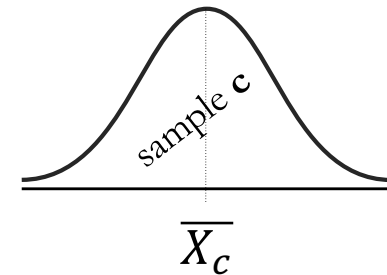
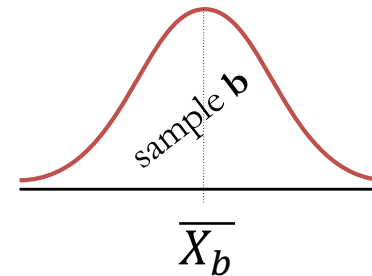
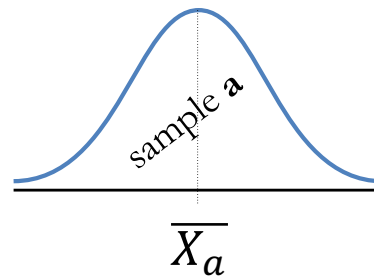
	t1	t2	t3
1	70	66	50
2	56	64	47
3	69	64	50
4	66	68	54
5	65	68	50
6	63	58	54
7	66	65	51
8	58	65	46
9	64	69	51
10	63	64	60
11	70	74	43
12	61	74	53
13	65	62	44
14	59	71	59
37	63	66	39
38	65	67	55
39	59	65	45
40	63	70	53
41	58	72	50
42	63	72	60
43	65	67	52
44	64	71	58
45	58	69	55
46	64	74	57
47	59	60	62
48	57	64	57
49	62	62	62
50	62	60	58



*Why would we want to compare multiple groups?*

*Why not just compare two groups at a time?*

*Do multiple populations have similar means?*



*Do multiple populations have the similar distributions?*



*When might we ask each these questions?*

*Are these questions different?*

# Advertising Example

A marketing manager is trying to develop a new nation-wide advertising campaign. She is not sure whether to advertise the low price, the high quality, or the great features of the product. To find out which strategy works best, she conducts an experiment using three different advertisements in three different cities that have nearly identical sales and demographics. She monitors the sales in each city for 50 days.

*Assume: the 3 strategies might not have run at the same time; the 3 cities are very similar; people did not share their reviews between cities*

She treats the three advertisements differently:

- the *low product price* is advertised in the first city (t1)
- the *high product quality* is advertised in the second city (t2)
- the *number of product features* is advertised in the third city (t3)



1. Did the three strategies produce different sales, on average?
2. Did the three strategies produce difference sales numbers?

```
adverts <- read.table("marketing_strategies.txt", header=TRUE)
```

Descriptive Statistics:

```
sapply(adverts, mean)
```

t1	t2	t3
62.36	66.74	51.96

```
sapply(adverts, sd)
```

t1	t2	t3
4.279805	4.173825	6.630049

$\bar{x}_1 : 62.36$

$s_1 : 4.28$

$\bar{x}_2 : 66.74$

$s_2 : 4.17$

$\bar{x}_3 : 51.96$

$s_3 : 6.63$

Overall sample mean (grand mean):

```
mean(sapply(adverts, mean))
```

```
[1] 60.35333
```

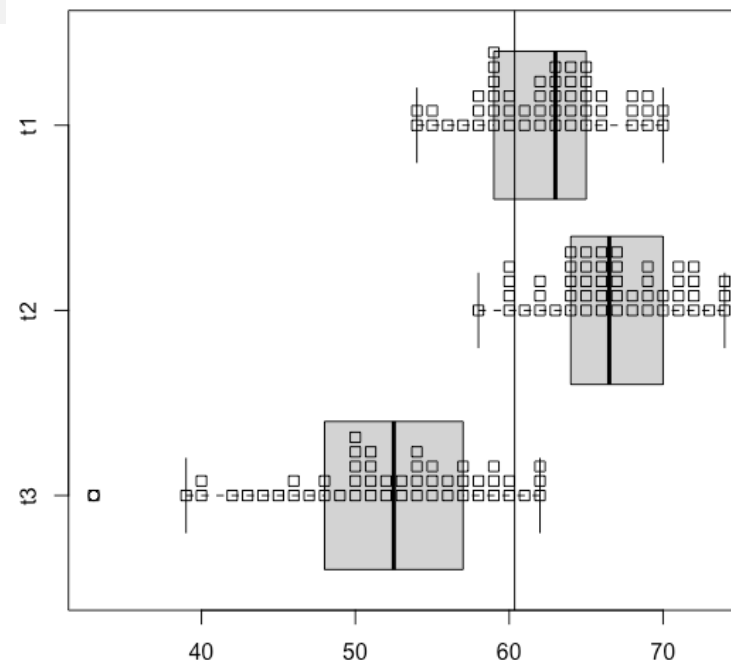
$\bar{x} : 60.35$

rev() reverses order

```
boxplot(rev(adverts), horizontal=TRUE)
```

```
stripchart(rev(adverts), method="stack", add=TRUE)
```

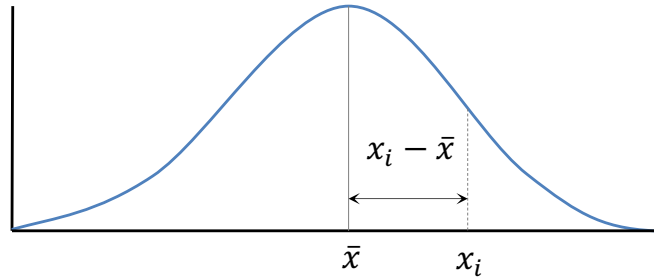
```
abline(v=mean(sapply(adverts, mean)))
```



wide dataframe

	t1	t2	t3
1	70	66	50
2	56	64	47
3	69	64	50
4	66	68	54
5	65	68	50
6	63	58	54
7	66	65	51
8	58	65	46
9	64	69	51
•	•	•	•
49	62	62	62
50	62	60	58

# Parametric Approach: Analyze Variances



**Absolute Deviation**

$$|x_i - \bar{x}|$$

**Variability**

$$\sum (x_i - \bar{x})^2 \quad \text{“Sum of Squares”}$$

**Mean Absolute Deviation**

$$\frac{\sum |x_i - \bar{x}|}{N}$$

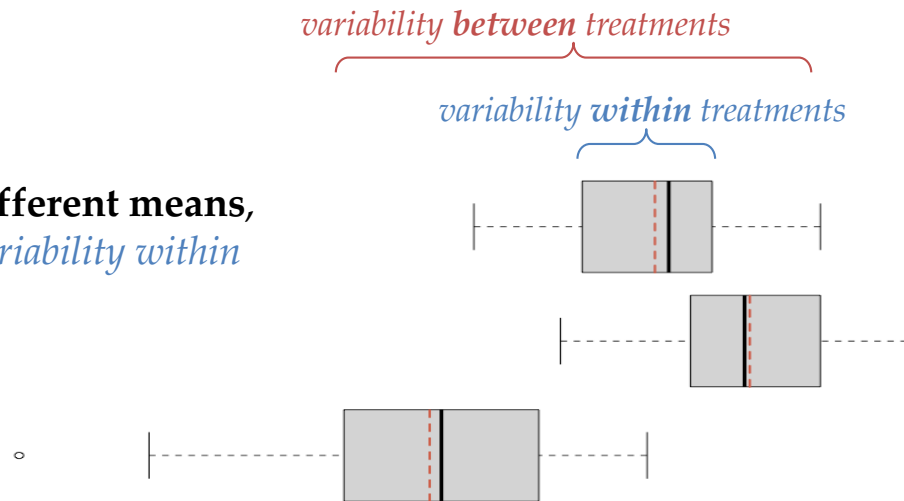
**Variance**

$$\frac{\sum (x_i - \bar{x})^2}{n - 1} \quad \begin{array}{l} \text{“Mean Squares”} \\ \text{(average variability?)} \end{array}$$

# Sources of Variance: Between/Within Treatments

**Treatment:** what was changed to create different groups (e.g., advertising strategy)

If "treatments" have **different means**,  
*variability between* > *variability within*



**Treatment variability**  
(variability between treatments)

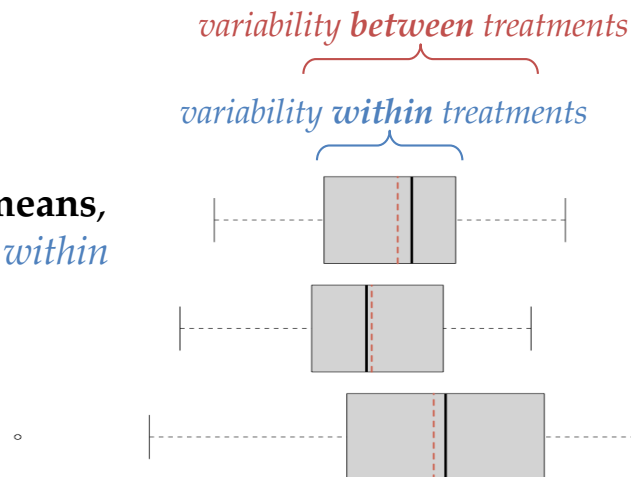
+

**Error variability**  
(variability within treatments)

---

**TOTAL Variability**

If "treatments" have **similar means**,  
*variability between*  $\approx$  *variability within*



We can *analyze variabilities* to see  
if treatments were effective!

# One-way Analysis of Variance (ANOVA)

**H<sub>null</sub>** : The means of the three treatment populations are the same

$$\mu_1 = \mu_2 = \mu_3$$

**H<sub>alt</sub>** : The means of the three treatments populations are not the same

*at least one mean is different*

**SSTR**: sum of squares due to treatments (**treatment variability**)

**MSTR**: mean square due to treatments (**treatment variance**)

**SSE**: sum of squares due to error (**error variability**)

**MSE**: mean square due to error (**error variance**)

*k treatments of 1 factor*

	t1	t2	t3
<i>n<sub>j</sub> observations of each treatment</i>			
1	70	66	50
2	56	64	47
3	69	64	50
4	66	68	54

$n_j \times k = n_T$  total observaitons

**between-treatments variance:**  
(variance of treatment means  
from grand mean)

$$MSTR = \frac{SSTR}{df_{MSTR}} = \frac{\sum_{j=1}^k n_j (\bar{x}_j - \bar{\bar{x}})^2}{k - 1}$$

**within-treatments variance:**  
(average variance within treatments)

$$MSE = \frac{SSE}{df_{MSE}} = \frac{\sum_{j=1}^k (n_j - 1) s_j^2}{n_T - k}$$

$$F = \frac{MSTR}{MSE}$$

*Ratio of variances  
follows F distrbution*

k: number of treatments

n<sub>j</sub>: sample size of treatment j

s<sub>j</sub>: standard deviation of treatment j

n<sub>T</sub>: size of all treatments combined

$\bar{x}_j$ : mean of sample j

$\bar{\bar{x}}$ : grand mean (mean of all sample means)

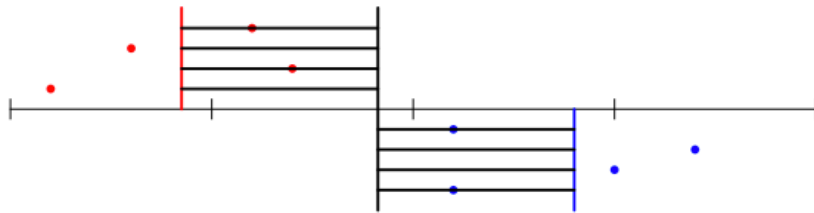
df: degrees of freedom

**between-treatments estimate of variance**  
 “variance between treatment means”

$$s^2 = (\text{treatment variance}) + (\text{error variance})$$

Will be larger than MSE if  $H_{\text{null}}$  is false

$$MSTR = \frac{SSTR}{df_{MSTR}} = \frac{\sum_{j=1}^k n_j (\bar{x}_j - \bar{\bar{x}})^2}{k-1}$$

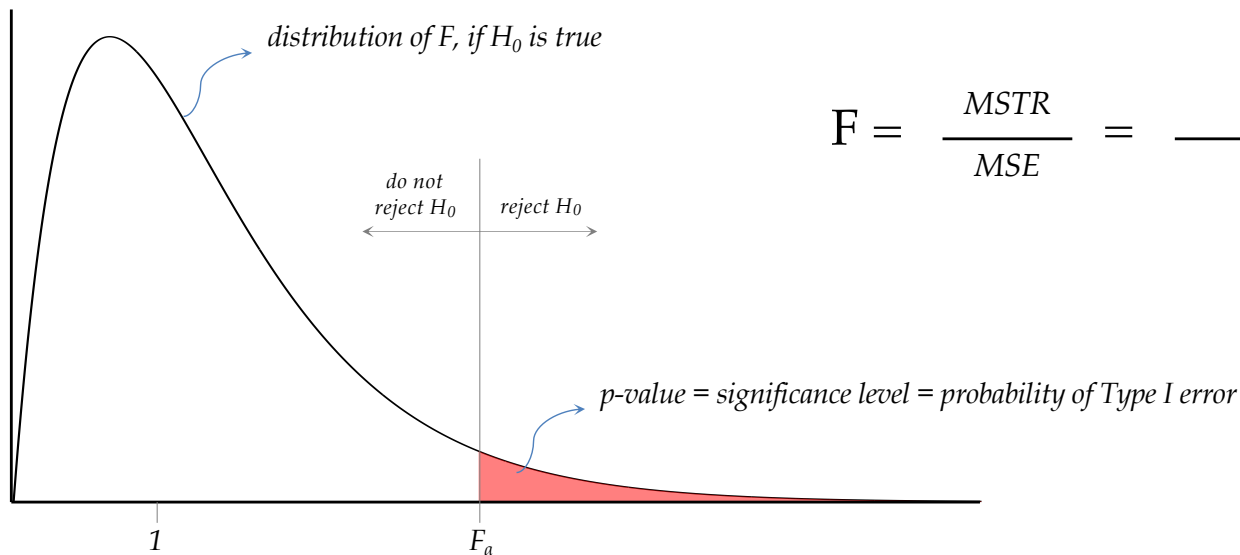
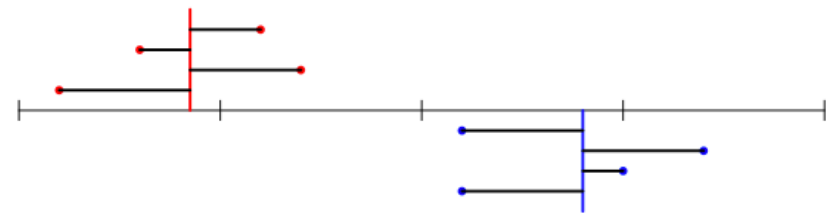


**within-treatments estimate of variance**  
 “average variance of all treatments”

$$s^2 = (\text{error variance})$$

$$MSE = \frac{SSE}{df_{MSE}} = \frac{\sum_{j=1}^k s_j^2}{k} \quad \text{if treatment sizes are the same}$$

$$MSE = \frac{SSE}{df_{MSE}} = \frac{\sum_{j=1}^k (n_j - 1) s_j^2}{n_T - k} \quad \text{if treatment sizes are different}$$



$$F = \frac{MSTR}{MSE} = \frac{(\text{treatment variance}) + (\text{error variance})}{(\text{error variance})}$$

# One-way ANOVA the long way in R

*functional iteration -- we don't need to explicitly loop through all the treatments!*

$$MSTR = \frac{SSTR}{df_{MSTR}} = \frac{\sum_{j=1}^k n_j (\bar{x}_j - \bar{\bar{x}})^2}{k-1} = \frac{5763.21}{3-1} = 2881.61$$

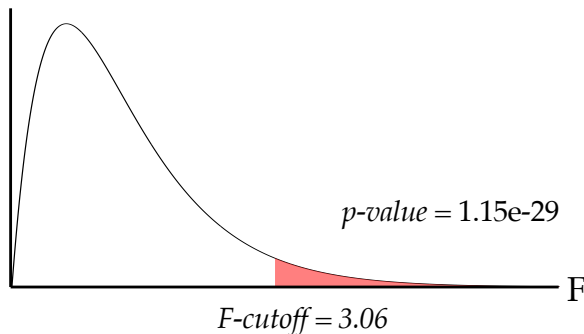
```
sstr <- 50*sum((sapply(adverts, mean) - mean(sapply(adverts, mean)))^2)
df_mstr <- 3-1
mstr <- sstr/df_mstr
```

$$MSE = \frac{SSE}{df_{MSE}} = \frac{\sum_{j=1}^k (n_j - 1) s_j^2}{n_T - k} = \frac{3905.06}{150-3} = 26.56$$

```
sse <- sum((50-1)*sapply(adverts, var))
df_mse <- 50*3 - 3
mse <- sse/df_mse
```

$$F = \frac{MSTR}{MSE} = \frac{2881.61}{26.56} = 108.47$$

```
f_value <- mstr/mse
```



```
qf(p=0.95, df1=df_mstr, df2=df_mse)
```

```
p_value <- pf(f_value, df_mstr, df_mse, lower.tail=FALSE)
```

## One-way ANOVA the easy way in R

	t1	t2	t3
1	70	66	50
2	56	64	47
3	69	64	50
4	66	68	54



	strategy	sales
1	t1	70
2	t1	56
3	t1	69
4	t1	66
...	...	...
51	t2	66
52	t2	64
53	t2	64
54	t2	68
...	...	...
101	t3	50
102	t3	47
103	t3	50
104	t3	54
...	...	...

*Change shape data of data to row-wise format*

```
library(reshape2)
ads <- melt(adverts, id.vars = NULL,
            variable.name = "strategy",
            value.name = "sales")
```

*Run **oneway.test()** function for one-way ANOVA*

*The tilde ~ separates dependent and independent variables*

***factor()** tells R that this is a **categorical** variable*

```
oneway.test(ads$sales ~ factor(ads$strategy), var.equal=TRUE)
```

One-way analysis of means

data: ads\$sales and ads\$strategy

F = 108.4737, num df = 2, denom df = 147, p-value < 2.2e-16

*Alternatively, run the more general **aov()** function for ANOVA*

```
anova_model <- aov( ads$sales ~ factor(ads$strategy))
summary(anova_model)
```

***summary()** often gives a more in-depth report on an estimated model*

```
              Df Sum Sq Mean Sq F value Pr(>F)
factor(ads$strategy)  2    5763   2881.6   108.5 <2e-16 ***
Residuals          147    3905    26.6
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```



# Requirements for ANOVA

ANOVA requires some assumptions to be met:

1. Each treatment/population's **response variable is normally distributed**
2. The **variance ( $s^2$ ) of the response variables is the same** for all treatments/populations
3. The **observations are independent**: the response variables are not related between groups



But it is fairly robust to minor violations



*ANOVA is robust to  
minor violations of  
normality and variances*



*ANOVA is more robust with  
more observations and  
fairly equal group sizes*



*But what should we do if we must greatly  
violate these assumptions?*

# Kruskal Wallis Test of Multiple Samples

*An extension of Mann-Whitney/Wilcoxon for multiple groups!*

$H_{\text{null}}$  : All groups would give you similar a value if randomly drawn from them

$H_{\text{alt}}$  : At least one group would give you a larger value than another if randomly drawn

	strategy	sales	ranks
1	t1	70	136.5
2	t1	56	40.0
3	t1	69	131.0
4	t1	66	113.5
5	t1	65	104.5
. . .			
51	t2	66	113.5
52	t2	64	94.5
53	t2	64	94.5
54	t2	68	125.0
55	t2	68	125.0
. . .			
101	t3	50	17.0
102	t3	47	11.0
103	t3	50	17.0
104	t3	54	30.5
105	t3	50	17.0
. . .			

*These rank sums ( $R_i$ ) give us a sense of how big the values are in each group*

## 1. Rank all the combined values across groups ( $i=3$ )

*Use the rank() function in R*

```
sales_ranks <- rank(. . .)
```

```
[1] 136.5 40.0 131.0 113.5 104.5 86.5 113.5 48.5 94.5 86.5 136.5 71.5 104.5 56.0
[15] 131.0 56.0 94.5 65.0 56.0 131.0 30.5 30.5 113.5 104.5 94.5 71.5 36.0 125.0
[29] 36.0 56.0 78.5 125.0 65.0 78.5 125.0 65.0 86.5 104.5 56.0 86.5 48.5 86.5
[43] 104.5 94.5 48.5 94.5 56.0 43.5 78.5 78.5 113.5 94.5 94.5 125.0 125.0 48.5
[57] 104.5 104.5 131.0 94.5 149.0 149.0 78.5 140.5 131.0 113.5 104.5 94.5 140.5 144.5
[71] 86.5 113.5 104.5 147.0 120.0 65.0 78.5 131.0 136.5 120.0 140.5 113.5 120.0 65.0
[85] 71.5 144.5 113.5 120.0 104.5 136.5 144.5 144.5 120.0 140.5 131.0 149.0 65.0 94.5
[99] 78.5 65.0 17.0 11.0 17.0 30.5 17.0 30.5 21.5 9.5 21.5 65.0 6.0 26.5
[113] 7.0 56.0 9.5 36.0 5.0 3.5 21.5 56.0 21.5 78.5 56.0 30.5 43.5 1.0
[127] 40.0 40.0 12.5 12.5 71.5 14.0 3.5 30.5 17.0 24.5 2.0 36.0 8.0 26.5
[141] 17.0 65.0 24.5 48.5 36.0 43.5 78.5 43.5 78.5 48.5
```

## 2. Group the ranks into original groups

```
group_ranks <- split(. . .)
```

```
$t1
[1] 136.5 40.0 131.0 113.5 104.5 86.5 113.5 48.5 94.5 86.5 136.5 71.5 104.5 56.0
[15] 131.0 56.0 94.5 65.0 56.0 131.0 30.5 30.5 113.5 104.5 94.5 71.5 36.0 125.0
[29] 36.0 56.0 78.5 125.0 65.0 78.5 125.0 65.0 86.5 104.5 56.0 86.5 48.5 86.5
[43] 104.5 94.5 48.5 94.5 56.0 43.5 78.5 78.5

$t2
[1] 113.5 94.5 94.5 125.0 125.0 48.5 104.5 104.5 131.0 94.5 149.0 149.0 78.5 140.5
[15] 131.0 113.5 104.5 94.5 140.5 144.5 86.5 113.5 104.5 147.0 120.0 65.0 78.5 131.0
[29] 136.5 120.0 140.5 113.5 120.0 65.0 71.5 144.5 113.5 120.0 104.5 136.5 144.5 144.5
[43] 120.0 140.5 131.0 149.0 65.0 94.5 78.5 65.0

$t3
[1] 17.0 11.0 17.0 30.5 17.0 30.5 21.5 9.5 21.5 65.0 6.0 26.5 7.0 56.0 9.5 36.0 5.0
[18] 3.5 21.5 56.0 21.5 78.5 56.0 30.5 43.5 1.0 40.0 40.0 12.5 12.5 71.5 14.0 3.5 30.5
[35] 17.0 24.5 2.0 36.0 8.0 26.5 17.0 65.0 24.5 48.5 36.0 43.5 78.5 43.5 78.5 48.5
```

## 3. Sum the ranks for each group

```
sapply(group_ranks, sum)
```

```
      t1      t2      t3
4159.0 5645.5 1520.5
```

*But these rank sums also depend on the size of each group ( $n_i$ )*

There are multiple ways of computing  $H$   
We will look at the rank sum approach...

4. Apply the **Kruskal Wallis formula** to sum the squared ranks

$$\text{Kruskal Wallis } H = \frac{12}{N(N+1)} \sum_{i=1}^k \frac{R_i^2}{n_i} - 3(N+1)$$

$R_i$  – sum of ranks of group  $i$

$n_i$  – number of observations in group  $i$

We adjust the rank sums by their group size, and compute a value that follows a chi-square distribution

```
H # [1] 92.49268
```

5. Find **p-value** of  $H$  from  $\chi^2$  distribution

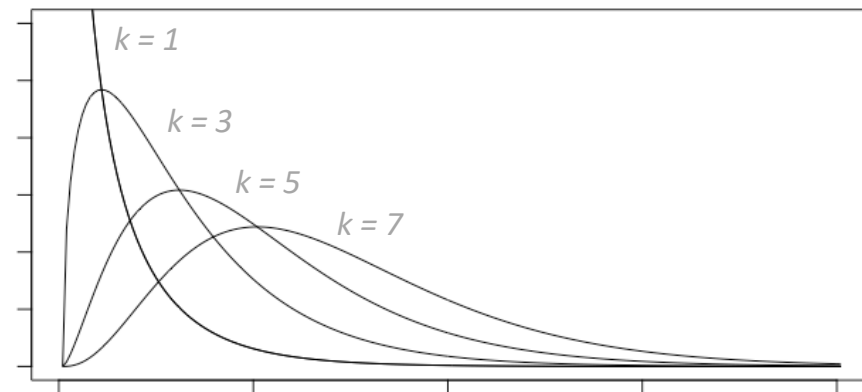
*p-value* tells you significance of test

```
kw_p <- 1 - pchisq(H, df=k-1)
```

$k$  – number of groups

```
kw_p  
[1] 0 Reject  $H_{null}$ 
```

$\chi^2$  “Chi-square” distribution



**R has a built-in Kruskal-Wallis Test!**


```
kruskal.test(sales ~ strategy, data = ads)
```

```
Kruskal-Wallis chi-squared = 92.679, df = 2, p-value < 2.2e-16
```


**Note:** There will be some difference in the calculation of  $H$  because `kruskal.test()` accounts for ties in ranks...

# Multigroup Post hoc Tests

*After data or results are seen*



	<u>t1</u>	<u>t2</u>	<u>t3</u>
1	70	66	50
2	56	64	47
3	69	64	50
4	66	68	54


  
*There are three possible group-wise differences:  
t1-vs-t2, t1-vs-t3, t2-vs-t3*

*So which groups are different from which others?*

*It is tempting to use multiple two-Sample Tests Post-hoc*

ANOVA  3 t-tests

```
t.test(adverts$t1, adverts$t2)
t.test(adverts$t1, adverts$t3)
t.test(adverts$t2, adverts$t3)
```

  
*But do not use two-sample methods to test  
two-way differences between multiple groups!*

Kruskal-Wallis  3 Wilcoxon Tests

```
wilcox.test(adverts$t1, adverts$t2)
wilcox.test(adverts$t1, adverts$t3)
wilcox.test(adverts$t2, adverts$t3)
```

# Family-Wise Error



*single die*

Running a test at 5% significance ( $\alpha=0.05$ ) is like rolling a 20-sided dice

Imagine rolling **one** 20-sided dice; probability of getting a “1” each time is 5%

```
rbinom(n=1, size=1, prob=0.05) 0, 0, 0, 0, 0, 0, 0, 0, 0, 0
```

This is the **single** risk of Type I error at 5% significance

```
sum(replicate(n, sum(rbinom(n=1, size=1, prob=0.05) >= 1))) / n  
[1] 0.05055
```



*family of dice*

Imagine rolling **five** 20-sided dice; probability of getting a “1” each time is **22%**

```
rbinom(n=1, size=5, prob=0.05) 0, 0, 1, 0, 0, 0, 0, 0, 1, 0
```

This is the **family-wise** risk of type I error at 5% significance

```
sum(replicate(n, sum(rbinom(n=1, size=5, prob=0.05) >= 1))) / n  
[1] 0.22659
```



*When evaluating many hypotheses,  
you are quite likely to commit type I error*



*So we cannot assess a family of tests using  
the same method to assess a single test*

# Tukey Test: Post-Hoc Test for ANOVA

## Confidence interval of difference of means

$$\underbrace{\bar{y}_{i.} - \bar{y}_{j.}}_{\text{difference of group means}} \pm \frac{1}{\sqrt{2}} q_{\alpha; r, N-r} \hat{\sigma}_{\epsilon} \sqrt{\frac{2}{n}}$$

$\hat{\sigma}_{\epsilon}$  – standard error of ANOVA ( $\sqrt{MSE}$ )

$q$  – adjustment factor based on number of groups

$n$  – size of a single group

```
TukeyHSD(anova_model, conf.level = 0.01)
```

	diff	lwr	upr	p adj
t2-t1	4.38	4.240815	4.519185	<b>0.000112</b>
t3-t1	-10.40	-10.539185	-10.260815	<b>0.000000</b>
t3-t2	-14.78	-14.919185	-14.640815	<b>0.000000</b>



Considers number of groups  
Considers unexplained variance (MSE)



## Assumptions of Tukey Test

1. The **variance** ( $s^2$ ) of the response variables is the same for all treatments/populations
2. The **observations are independent**: the response variables are not related between groups

## Dunn Test: Posthoc Tests for Kruskal Wallis

$$Z_{kw} = \frac{\overbrace{\bar{R}_i - \bar{R}_j}^{\text{difference of mean group sum-ranks}}}{\sqrt{\underbrace{\frac{N(N+1)}{12}}_{N - \text{total number of data observations}} \left( \underbrace{\frac{1}{n_i} + \frac{1}{n_j}}_{n_i, n_j - \text{size of groups } i \text{ and } j} \right)}}$$

$\bar{R}_i, \bar{R}_j$  – mean of sum of ranks for groups  $i$  and  $j$

```
# install.packages("FSA")
library(FSA)
dunnTest(sales ~ strategy, data = ads, method = "bonferroni")
```

	Comparison	Z	P.unadj	P.adj
1	t1 - t2	-3.42499	6.148226e-04	<b>1.844468e-03</b>
2	t1 - t3	6.07927	1.207306e-09	<b>3.621918e-09</b>
3	t2 - t3	9.50426	2.014750e-21	<b>6.044251e-21</b>



**No assumptions of variance homogeneity**  
**No assumptions of equal group sizes**

# Parametric vs. Non-Parametric Methods

	Parametric Methods	Non-parametric Methods
<b>Data Distribution Assumptions</b>	<i>Distribution parameters</i> of original data	--
<b>Approach</b>	<i>Statistical</i> (simple, elegant formulas)	<i>Computational</i> (resampling/ranking)
<b>Advantage</b>	More statistical power if assumptions are met	Better when assumptions violated; Useful for new / different statistics
<b>Disadvantage</b>	Often not applicable to noisy real-world data; Often not applicable to new / different statistics	Procedures get complicated quickly; Many variations to choose from



## Parametric Tests

## Non-parametric Tests

**Assumptions**  
that need to be met

*Distribution parameters* of original data

--

**Information**  
that is tested

*Mean, Variance*

*Median, Quantiles, Ranks, Deviation, ...*

Example of  
**One-sample test**

t-Tests

Bootstrap

Example of  
**Two-sample test**

t-Tests

Bootstrapped t-Tests  
Parametric Test  
Wilcoxon Test  
Signed-Rank Test

Example of  
**Multi-group test**

ANOVA

Bootstrapped ANOVA  
Kruskal Wallis Test

**Posthoc**  
multigroup  
comparisons

Tukey Test

Dunn Test

**Major**  
**Disadvantage**

Assumptions often cannot be met  
in real world data

Gets very complicated quickly;  
Many variations and alternatives