Day 17: Nonparametric tests - Supplemental material **BSTA 511/611** Meike Niederhausen, PhD **OHSU-PSU School of Public Health** 11/29/2023

MoRitz's tip: write "nice" R code

Check out the tidyverse style guide: https://style.tidyverse.org/index.html

Especially, Chapter 4: Pipes and Chapter 5: ggplot2

```
```{r}
```

승 🔺 🕨 employ<-employ%>%mutate(disability=factor(disability),di sability=fct\_relevel(disability, "none"), disability=fct\_r ecode(disability.amputation="amputee")) summary(employ) . . .

VS.

```
```{r}
employ <- employ %>%
  mutate(
    disability = factor(disability),
# make "none" the first level
      disability = fct_relevel(disability,
                    "none"),
# change level name amputee to amputation
      disability = fct_recode(disability,
                 amputation = "amputee"))
summary(employ)
* * *
```

$^{} ^{} r}$

VS.

gqplot(employ,aes(x=disability,y=score,fill=disability,c olor=disability))+geom_dotplot(binaxis="y",alpha=.5)+geo m_hline(aes(yintercept=mean(score)),lty="dashed")+stat_s ummary(fun="mean",geom="point",size=3,color="grey33",alp ha=1)+theme(legend.position="none")

```
```{r}
 🎰 🔟 🕨
gqplot(employ,
 aes(x = disability, y=score,
 fill = disability,
 color = disability)) +
 geom_dotplot(binaxis = "y", alpha = .5) +
 geom_hline(aes(yintercept = mean(score)),
 lty = "dashed") +
 stat_summary(fun = "mean",
 geom = "point",
 size = 3.
 color = "grey33",
 alpha = 1) +
 theme(legend.position = "none")
```

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# Where are we?



# Goals for today (Supplemental material)

- Why us a nonparametric approach?
- What the following tests are & when to use them

### • Sign test

- for paired data or single samples
- (Wilcoxon) sign-rank test

• for paired data or single samples

accounts for sizes of differences

- Wilcoxon Rank-sum test
  - for two independent samples
  - a.k.a Mann-Whitney U test
- Kruskal-Wallis test
  - nonparametric ANOVA test
- How to use R for each test & interpret the results

### Additional resource

- Chapter 13: Nonparametric tests of Pagano's Principles of Biostatistics, 2022 edition
- Can download chapter from OHSU library eBook at https://ebookcentral.proquest.com/lib/ohsu/detail.action?docID=6950388&pqorigsite=primo

# Nonparametric tests

### Background: parametric vs nonparametric

- Prior inference of means methods had conditions assuming the underlying population(s) was (were) normal (or at least approximately normal).
- Normal distribution is completely described (parameterized) by two parameters:  $\mu$  and  $\sigma$ .
- The first was often the parameter of interest, while the latter was assumed known ( Z-test) or estimated ( t-tests).
- The above are therefore described as **parametric** procedures.
- Nonparametric procedures
  - Make fewer assumptions about the structure of the underlying population from which the samples were collected.
  - Work well when distributional assumptions are in doubt.

### The good and the bad about nonparametric tests

#### Good

- Fewer assumptions
- Tests are based on ranks
  - Therefore outliers have no greater influence than non-outliers.
  - Since tests are based on ranks we can apply these procedures to ordinal data
    - (where means and standard deviations are not meaningful).

### Drawbacks

- Less powerful than parametric tests (due to loss of information when data are converted to ranks)
- While the test is easy, it may require substantial (computer) work to develop a confidence interval [by "inverting" the test].
- Theory was developed for continuous data (where ties are not possible); if population or data contain many ties, then a correction to the procedures must be implemented.
- Some procedures have "large" and "small" sample versions; the small sample versions require special tables or a computer.



For paired data or single samples

# Example: Intraocular pressure of glaucoma patients

- Intraocular pressure of glaucoma patients is often reduced by treatment with adrenaline.
- A new synthetic drug is being considered, but it is more expensive than the current adrenaline alternative.
- 7 glaucoma patients were treated with both drugs:
  - one eye with adrenaline and
  - the other with the synthetic drug
- **Reduction in pressure** was recorded in each eye after following treatment (larger numbers indicate greater reduction)

Patient	Adren	Synth	d	Sign
1	3.5	3.2	-0.3	-
2	2.6	3.1	0.5	+
3	3.0	3.3	0.3	+
4	1.9	2.4	0.5	+
5	2.9	2.9	0.0	NA
6	2.4	2.8	0.4	+
7	2.0	2.6	0.6	+

- **d** is the difference in reduction of pressure: **Synth Adren**
- Sign is
  - + if the difference is positive and
  - - if the difference is negative

### Visualize the differences

Visualize the differences in reduction of pressure d : Synth - Adren



# Hypotheses & "statistic" (Sign test)

#### Hypotheses

 $H_0$  : The median difference in the population is 0  $H_a$  : The median difference in the population is NOT 0

#### "Statistic"

 $D^+$  = number of positive differences  $D^-$  = number of negative differences

What are  $D^+$  and  $D^-$  for our example?

Patient	Adren	Synth	d	Sign
1	3.5	3.2	-0.3	-
2	2.6	3.1	0.5	+
3	3.0	3.3	0.3	+
4	1.9	2.4	0.5	+
5	2.9	2.9	0.0	NA
6	2.4	2.8	0.4	+
7	2.0	2.6	0.6	+

## Exact p-value (Sign test)

- If the median difference is 0 (  $H_0$  is true) , then

half the population consists of positive differences
while half have negative differences.

• Let p = P(neg. diff.) = P(pos. diff.) = 0.5

- If the median difference is 0 (  $H_0$  is true),
  - $\circ$  then a sample of n differences
    - behaves like n trials in a binomial experiment
    - where "success" is analogous to seeing a positive difference.
  - $\circ\,$  By symmetry ( p=0.5 ), the same distribution applies to negative differences, i.e.,

$$D^+ ext{ and } D^- \sim \mathrm{Bin}(n,p=0.5)$$
 .

• Thus the (exact) p-value is calculated using the Binomial distribution

### Glaucoma example (exact) p-value

- 7 differences:
  - $\circ$  1 negative (  $D^-$  )
  - $\circ\,$  5 were positive (  $D^+$  )
  - 1 difference is 0 and is discarded
- Thus the effective sample size is n=6.

**One-sided p-value** = probability that we would see 1 or fewer negative signs among the n = 6 differences, if the median difference is really 0

**Two-sided p-value** = 2 × One-sided p-value

# 2-sided p-value: 2\*P(D^- <= 1)
2\*pbinom(1, size = 6, p = 0.5)</pre>

$$D^- \sim {
m Bin}(n=6,p=0.5)$$

 $p-value = P(D^- \le 1)$ =  $P(D^- = 0) + P(D^- = 1)$ =  $rac{6!}{0!6!}(0.5)^6 + rac{6!}{1!5!}(0.5)^6$ pprox 0.1094

p-value imes 2 pprox 0.2188

## [1] 0.21875

# Sign test in R: Glaucoma example

Below we create the dataset as a tibble (and add the signs):

Recall we're testing the population median. Here's the sample median:

median(IOP\$d)

## [1] 0.4

IOP %>% gt()

Patient	Adren	Synth	d	Sign
1	3.5	3.2	-0.3	-
2	2.6	3.1	0.5	+
3	3.0	3.3	0.3	+
4	1.9	2.4	0.5	+
5	2.9	2.9	0.0	NA
6	2.4	2.8	0.4	+
7	2.0	2.6	0.6	+

## Sign test in R: Glaucoma example (specifying both columns)

library(BSDA) # new package!! Make sure to first install it # Can't "tidy" the output SIGN.test(x = IOP\$Synth, y = IOP\$Adren, alternative = "two.sided", conf.level = 0.95)

```
##
 Dependent-samples Sign-Test
##
##
data: IOP$Synth and IOP$Adren
S = 5, p-value = 0.2187
alternative hypothesis: true median difference is not equal to 0
95 percent confidence interval:
-0.2057143 0.5685714
sample estimates:
median of x-y
##
 0.4
##
Achieved and Interpolated Confidence Intervals:
##
 Conf.Level L.E.pt U.E.pt
##
 Lower Achieved CI 0.8750 0.0000 0.5000
##
Interpolated CI 0.9500 -0.2057 0.5686
Upper Achieved CI 0.9844 -0.3000 0.6000
```

### Sign test in R: Glaucoma example (specifying differences)

```
Note output calls this a "One-sample Sign-Test"
SIGN.test(x = IOP$d, alternative = "two.sided", conf.level = 0.95)
```

```
##
##
 One-sample Sign-Test
##
data: IOP$d
s = 5, p-value = 0.2187
alternative hypothesis: true median is not equal to 0
95 percent confidence interval:
 -0.2057143 0.5685714
##
sample estimates:
median of x
##
 0.4
##
Achieved and Interpolated Confidence Intervals:
##
##
 Conf.Level L.E.pt U.E.pt
 Lower Achieved CI 0.8750 0.0000 0.5000
##
Interpolated CI 0.9500 -0.2057 0.5686
Upper Achieved CI 0.9844 -0.3000 0.6000
```

### Conclusion

Recall the hypotheses to the sign test:

 $H_0$  : The median population difference in reduction of intraocular pressure in treatment with adrenaline vs. new synthetic drug is 0.

 $H_a$ : The median population difference in reduction of intraocular pressure in treatment with adrenaline vs. new synthetic drug is NOT 0.

- Significance level:  $\alpha$  = 0.05
- p-value = 0.2188

### Conclusion:

The median difference in reduction of intraocular pressure between eyes being treated with the synthetic drug and adrenaline for seven glaucoma patients was 0.4 (95% CI: -0.2, 0.6). There is insufficient evidence the reduction in intraocular pressure differs when using the synthetic drug and adrenaline (2-sided sign test p-value = 0.219).

### Sign test with large samples: p-value normal approximation

- If the sample size is large, say greater than 20,
  - then binomial probabilities can be approximated using normal probabilities
- Normal approximation:

$$\mu = np = n(0.5) = n/2$$
 $\sigma = \sqrt{np(1-p)} = \sqrt{n(0.5)(0.5)} = \sqrt{n}/2$ 

• Thus we have the test statistic:

$$z=rac{D^--n/2}{\sqrt{n}/2}$$

• With access to a computer, it's better to use the exact binomial probabilities instead of the normal approximation.

### Sign test with one sample

- One can use the sign test when testing just one sample.
- Note that we did this when in R, when running the sign test using just the differences.
- For one sample, we can specify the null population median value:

 $H_0$  : The population median is m $H_a$  : The population median is NOT m

Example: Run sign test for paired data with null m = 0.7:

```
SIGN.test(x = IOP$d, md = 0.7, alternative = "two.sided", conf.level = 0.95)
```

```
##
One-sample Sign-Test
##
data: IOP$d
s = 0, p-value = 0.01563
alternative hypothesis: true median is not equal to 0.7
95 percent confidence interval:
-0.2057143 0.5685714
sample estimates:
median of x
```

# (Wilcoxon) Signed-rank test

For paired data or single samples; accounts for sizes of differences

# (Wilcoxon) Signed-rank test

- Like the sign test, the (Wilcoxon) signed-rank test is used for
  - paired samples (i.e., a single set of differences) or
    a one-sample comparison against a specified value
- However, this test *does* make use of information concerning the size of the differences.

#### Hypotheses

 $H_0$ : the population is symmetric around some value  $\tilde{\mu}_0$  $H_a$ : the population is not symmetric around some value  $\tilde{\mu}_0$ 

- Even if the population has a mean/median equal to  $\tilde{\mu}_0$ , the test may reject the null if the population isn't symmetric, thus only use if the data (differences) are symmetric.
- If the population is symmetric
  - then the mean and median coincide,
  - thus often the null hypothesis is phrased in terms of the median difference being 0

## Example: calculate signed ranks

- Rank the absolute values of the differences from smallest to largest
- For ties, take the average of the highest and lowest tied ranks
  - I.e. if ranks 3-7 are tied, then assign (3+7)/2 = 5 as the rank
- Then calculate the signed ranks as +/the rank depending on whether the sign is +/-

```
IOP_ranks <- IOP %>%
 mutate(abs_d = abs(d)) %>%
 arrange(abs_d) %>%
 mutate(
 Rank = c(NA, 1.5, 1.5, 3, 4.5, 4.5, 6),
 Signed_rank = case_when(
 d < 0 ~ -Rank,
 d > 0 ~ Rank))
```

IOP\_ranks %>% gt()

Patient	Adren	Synth	d	Sign	abs_d	Rank	Signed_rank
5	2.9	2.9	0.0	NA	0.0	NA	NA
1	3.5	3.2	-0.3	-	0.3	1.5	-1.5
3	3.0	3.3	0.3	+	0.3	1.5	1.5
6	2.4	2.8	0.4	+	0.4	3.0	3.0
2	2.6	3.1	0.5	+	0.5	4.5	4.5
4	1.9	2.4	0.5	+	0.5	4.5	4.5
7	2.0	2.6	0.6	+	0.6	6.0	6.0

## Test statistic (Wilcoxon) Signed-rank test

### If the null is true:

- The population is symmetric around some point (  $ilde{\mu}_0=0$  , typically), and
- The overall size of the positive ranks should be about the same as the overall size of negative ranks.

Note:

- The sum of the ranks  $1,2,\ldots,n$  is always n(n+1)/2,
- which can be broken down as the
  - $\circ\,$  sum of the positive ranks (  $T^+$  )
  - $\circ\,$  plus the sum of the negative ranks (  $T^-$  )

Thus, any of the following can be used as a test statistic and will lead to the same conclusion:

- $T^+$
- <u>T</u>-

•  $T_{min} = \min(T^+, T^-)$ 

### Example: calculate sums of signed ranks

IOP\_ranks %>% gt()

Patient	Adren	Synth	d	Sign	abs_d	Rank	Signed_rank
5	2.9	2.9	0.0	NA	0.0	NA	NA
1	3.5	3.2	-0.3	-	0.3	1.5	-1.5
3	3.0	3.3	0.3	+	0.3	1.5	1.5
6	2.4	2.8	0.4	+	0.4	3.0	3.0
2	2.6	3.1	0.5	+	0.5	4.5	4.5
4	1.9	2.4	0.5	+	0.5	4.5	4.5
7	2.0	2.6	0.6	+	0.6	6.0	6.0

• Sum of the positive ranks

 $\circ T^+ = 1.5 + 3 + 4.5 + 4.5 + 6 = 19.5$ 

• Sum of the negative ranks

∘ *T*<sup>−</sup> = -1.5

- The sum of the ranks  $1,2,\ldots,n$  is always n(n+1)/2:
  - $egin{array}{lll} \circ \ n(n+1)/2 = 6(7)/2 = 21 \ \circ \ T^+ + |T^-| = 19.5 + |-1.5| = 21 \end{array}$

# Exact p-value (Wilcoxon) Signed-rank test (fyi) (1/2)

- Exact p-value is preferable
  - This is the default method in R's wilcox.test()
    - if the samples contain less than 50 finite values
    - and there are no ties
      - *R* will automatically use normal approximation method if there are ties
- We will not be calculating the exact p-value "by hand." We will be using R for this.

$$p-value=2*P(\min(T^+,T^-)\leq t)$$

- t is the smaller of the calculated sums of the positive and negative ranks
- To calculate the exact p-value, we need the probability of each possible sum of ranks.

# Exact p-value (Wilcoxon) Signed-rank test (fyi) (2/2)

- To calculate the exact p-value, we need the probability of each possible sum of ranks:
  - list all possible combinations of positive and negative ranks for the sample size,
  - $\circ\,$  calculate the sum of the positive ranks (  $T^+$  ) for each possible combination (or  $T^-$  ), and
  - $\circ\,$  then figure out the probability of each possible  $T^+$  (assuming all combinations are equally likely)

Example when n = 3: (from https://online.stat.psu.edu/stat415/lesson/20/20.2)



See https://online.stat.psu.edu/stat415/lesson/20/20.2 for more details.

# Normal approx. p-value (Wilcoxon) Signed-rank test (fyi)

#### • Normal approximation method:

- If the number of non-zero differences is at least 16, then a normal approximation can be used.
- Have the option to apply a continuity correct (default) or not
- We will not be calculating the p-value "by hand." We will be using R for this.

Test statistic:

$$Z_{T_{min}} = rac{T_{min} - rac{n(n+1)}{4}}{\sqrt{rac{n(n+1)(2n+1)}{24}}}$$

- $T_{min}=\min(T^+,T^-)$
- *n* = sample size
- Test statistic  $Z_{T_{min}}$  follows a standard normal distribution N(0,1)
- Use normal distribution to calculate p-value

See https://online.stat.psu.edu/stat415/lesson/20/20.2 for more details.

### (Wilcoxon) Signed-rank test in R: Glaucoma example

"Attempt" with exact p-value & specifying columns for paired data

## Warning in wilcox.test.default(x = IOP\$Synth, y = IOP\$Adren, paired = TRUE, :
## cannot compute exact p-value with ties

## Warning in wilcox.test.default(x = IOP\$Synth, y = IOP\$Adren, paired = TRUE, :
## cannot compute exact p-value with zeroes

```
##
Wilcoxon signed rank test with continuity correction
##
data: IOP$Synth and IOP$Adren
V = 19.5, p-value = 0.07314
alternative hypothesis: true location shift is not equal to 0
```

### (Wilcoxon) Signed-rank test in R: Glaucoma example

"Attempt" with exact p-value & running one sample test with differences

## Warning in wilcox.test.default(x = IOP\$d, alternative = c("two.sided"), :
## cannot compute exact p-value with ties

## Warning in wilcox.test.default(x = IOP\$d, alternative = c("two.sided"), :
## cannot compute exact p-value with zeroes

```
##
Wilcoxon signed rank test with continuity correction
##
data: IOP$d
V = 19.5, p-value = 0.07314
alternative hypothesis: true location is not equal to 0
```

## (Wilcoxon) Signed-rank test in R: Glaucoma example

"Attempt" with approximate p-value & specifying columns for paired data

##
## Wilcoxon signed rank test with continuity correction
##
## data: IOP\$Synth and IOP\$Adren
## V = 19.5, p-value = 0.07314
## alternative hypothesis: true location shift is not equal to 0

No more warnings!! However,... should we be using the normal approximation here?

## Conclusion

Recall the hypotheses to the (Wilcoxon) Signed-rank test:

 $H_0$ : the population difference in reduction of intraocular pressure in treatment with adrenaline vs. new synthetic drug is **symmetric around**  $\tilde{\mu}_0 = 0$  $H_a$ : the population difference in reduction of intraocular pressure in treatment with adrenaline vs. new synthetic drug is **not symmetric around**  $\tilde{\mu}_0 = 0$ 

- Significance level:  $\alpha$  = 0.05
- p-value = 0.07314

### Conclusion:

There is insufficient evidence the differences in reduction in intraocular pressure differs between the synthetic drug and adrenaline are symmetric about 0 (2-sided Wilcoxon signed rank test p-value = 0.07314)

However,...

## Wilcoxon rank-sum test

For two independent samples a.k.a Mann-Whitney U test

### Wilcoxon rank-sum test

- The nonparametric alternative to the two-sample *t*-test
   used to analyze two samples selected from separate (independent) populations
- Also called the Mann-Whitney U test.
- Unlike the signed-rank test, there is no need to assume symmetry
- Necessary condition is that the two populations being compared
  - have the same shape (both right skewed, both left skewed, both symmetric, etc.),
    so any noted difference is due to a shift in the median
- Since they have the same shape, when summarizing the test, we can describe the results in terms of a difference in medians.

#### Hypotheses:

 $H_0:$  the two populations have the same median  $H_a:$  the two populations do NOT have the same median

## Example

gt()

Dr. Priya Chaudhary (OHSU) examined the evoked membrane current of dental sensory neurons (in rats) under control conditions and a mixture of capsaicin plus capsazepine (CPZ). J. Dental Research} 80:1518--23, 2001.

```
CPZdata <- tibble(
 control = c(3024, 2164, 864, 780, 125, 110),
 cap_CPZ = c(426, 232, 130, 94, 75, 55)
)
CPZdata %>%
 get_summary_stats(type = "median") %>%
```

variable n median

822

112

control 6

cap\_CPZ 6

CPZdata %>% gt()

control	cap_CPZ
3024	426
2164	232
864	130
780	94
125	75
110	55

## Visualize the data

Do the independent samples have the same distribution?

#### Control group



### Cap + CPZ group



## Calculating ranks and test statistic $\boldsymbol{W}$

- 1. Combine the two samples together (keep track of which observations came from each sample).
- 2. Rank the full set of  $N=n_1+n_2$  observations.
  - If ties exist, assign average ranks to the tied values (as with the signedrank test).
- 3. Sum the ranks corresponding to those observations from the smaller sample.
  - This is a time-saving step; you could also have used the larger sample.
  - $\circ\,$  Call this sum W.

$$W_{CPZ} = 1 + 2 + 3 + 6 + 7 + 8 = 27$$

 $W_{control} = 4 + 5 + 9 + 10 + 11 + 12 = 51$ 

Group	Current	Rank
cap_CPZ	55	1
cap_CPZ	75	2
cap_CPZ	94	3
control	110	4
control	125	5
cap_CPZ	130	6
cap_CPZ	232	7
cap_CPZ	426	8
control	780	9
control	864	10
control	2164	11
control	3024	12

In our example, both groups have equal n; choose either for computing W. 35/53

# Exact p-value approach (fyi)

- If  $n_1, n_2$  are both less than 10, then use an exact test,
  - otherwise use the large-sample normal approximation.
  - However, exact method can only be done if **no ties** are present
- p-value is the probability of getting a rank sum W as extreme or more extreme than the observed one.
  - Multiply the 1-tail probability by 2 for the 2-tailed probability

$$p-value = 2 \cdot P(W_{CPZ} \leq 27)$$

- To calculate  $P(W_{CPZ} \leq 27)$ ,
  - we need to enumerate all possible sets ranks for the sample size,
  - calculate the sum of ranks for each set of possible ranks,
  - and then the probability for each sum (assuming each set of ranks is equally likely).
- We will not be calculating the p-value "by hand." We will be using R for this.

# Normal approximation approach (fyi)

If the null hypothesis is true, then the mean of the sum of the ranks from the smallersized group will be

$$\mu_W = rac{n_s \cdot (n_s + n_l + 1)}{2},$$

with a standard deviation of

$$\sigma_W = \sqrt{rac{n_s \cdot n_l \cdot (n_s + n_l + 1)}{12}}.$$

Provided both groups are large (  $\geq 10$  ),

$$Z = rac{W-\mu_W}{\sigma_W} pprox Normal(0,1)$$

#### **Example:**

We have 
$$W=27$$
 and  $n_l=n_s=6$ :

$$\mu_W = \frac{6 \cdot (6 + 6 + 1)}{2} = 39$$

$$\sigma_W = \sqrt{\frac{6 \cdot 6 \cdot (6 + 6 + 1)}{12}} = \sqrt{39} \approx 6.2450$$

$$z \approx \frac{27 - 39}{6.2450} = -1.921538$$

The two-sided p-value is

 $p = 2 \cdot P(Z < -1.921538) = 0.05466394$ 

## R code for creating ranks on previous slide

CPZdata
<pre>## # A tibble: 6 × 2 ## control cap_CPZ ##</pre>
<pre>CPZdata_long &lt;- CPZdata %&gt;%   pivot_longer(cols = c(control,cap_CPZ),</pre>

CPZdata\_long %>% gt()

Group	Current	Rank
cap_CPZ	55	1
cap_CPZ	75	2
cap_CPZ	94	3
control	110	4
control	125	5
cap_CPZ	130	6
cap_CPZ	232	7
cap_CPZ	426	8
control	780	9
control	864	10
control	2164	11
control	3024	12

38/53

### Wilcoxon rank-sum test in R: with wide data

glimpse(CPZdata)

## Rows: 6
## Columns: 2
## \$ control <dbl> 3024, 2164, 864, 780, 125, 110
## \$ cap\_CPZ <dbl> 426, 232, 130, 94, 75, 55

#### Exact p-value

```
##
Wilcoxon rank sum exact test
##
data: CPZdata$cap_CPZ and CPZdata$control
W = 6, p-value = 0.06494
alternative hypothesis: true location shift is not equal to 0
```

### Wilcoxon rank-sum test in R: with wide data

Normal approximation p-value without CC

statistic	p.value	method	alternative
6	0.05466394	Wilcoxon rank sum test	two.sided

```
Normal approximation p-value with CC
```

statistic	p.value	method	alternative
6	0.06555216	Wilcoxon rank sum test with continuity correction	two.sided

### Wilcoxon rank-sum test in R: with long data

Make data long (if it's not already long):

head(CPZdata\_long)

##	#	A tibble	e: 6 × 2
##		Group	Current
##		<chr></chr>	<dbl></dbl>
##	1	control	3024
##	2	cap_CPZ	426
##	3	control	2164
##	4	cap_CPZ	232
##	5	control	864
##	6	cap_CPZ	130

Exact p-value

statistic	p.value	method	alternative
6	0.06493506	Wilcoxon rank sum exact test	two.sided

## Conclusion

Recall the hypotheses to the (Wilcoxon) Signed-rank test:

 $H_0:$  the control and treated populations have the same median  $H_a:$  the control and treated populations do NOT have the same median

- Significance level:  $\alpha$  = 0.05
- p-value = 0.06494

### **Conclusion**:

There is suggestive but inconclusive evidence that the evoked membrane current of dental sensory neurons (in rats) differs between the control group and the group exposed to a mixture of capsaicin plus capsazepine (2-sided Wilcoxon rank-sum test p-value = 0.06494).

# Kruskal-Wallis test

Nonparametric ANOVA test

# Kruskal-Wallis test: nonparametric ANOVA test

- Recall that an ANOVA tests means from more than 2 groups
- Conditions for ANOVA include
  - $\circ\,$  Sample sizes in each group group are large (each  $n\geq 30$ ),
    - OR the data are relatively normally distributed in each group
  - Variability is "similar" in all group groups
- If these conditions are in doubt, or if the response is ordinal, then the Kruskal-Wallis test is an alternative.

 $H_0: ext{pop median}_1 = ext{pop median}_2 = \ldots = ext{pop median}_k$ vs.  $H_A: ext{At least one pair pop median}_i \neq ext{pop median}_i ext{ for } i \neq j$ 

- K-W test is an extension of the (Wilcoxon) rank-sum test to more than 2 groups  $\,\circ\,$  With k=2 groups, the K-W test is the same as the rank-sum test

### K-W test statistic: *H* (fyi)

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

- k is the number of groups,
- $n_i$  is the number of observations in group i
- $N=n_1+\ldots+n_k$  is the total number of observations across all groups,
- $R_i$  is the sum of ranks for group i

The test statistic H has a Chi-squared distribution with k-1 degrees of freedom:

$$H\sim\chi^2_{k-1}$$

Ranks are calculated similarly to the (Wilcoxon) rank-sum test.

# Ranks for the K-W test

- 1. Combine the k samples together (keep track of which observations came from each sample).
- 2. Rank the full set of  $N=n_1+\ldots+n_k$  observations.
  - If ties exist, assign average ranks to the tied values (as with the signed-rank test).
- 3. Then sum the ranks within each of the k groups
  - $\circ$  Label the sums of the ranks for each group as  $R_1, \ldots + R_k$ .

If  $H_0$  is true, we expect the populations to have the same medians, and thus the ranks to be similar as well.

## Example: Ozone levels by month (1/2)

- airquality data included in base R no need to load it
- Daily air quality measurements in New York, May to September 1973.

## \$ Day <int> 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, 17, 18,...

• Question: do ozone levels differ by month?

glimpse(airquality)

# Example: Ozone levels by month (2/2)

```
Oz_mnth <- airquality %>%
 group_by(Month) %>%
 get_summary_stats(Ozone,
 show = c("n", "mean", "median", "sd"))
Oz_mnth %>% gt()
```

Month	variable	n	mean	median	sd
5	Ozone	26	23.615	18	22.224
6	Ozone	9	29.444	23	18.208
7	Ozone	26	59.115	60	31.636
8	Ozone	26	59.962	52	39.681
9	Ozone	29	31.448	23	24.142

max(Oz\_mnth\$sd) / min(Oz\_mnth\$sd)



## [1] 2.179317

## Example: calculate ranks (fyi) (1/2)

```
ranks_Oz_mnth <- airquality %>%
 select(Ozone, Month)
```

summary(ranks\_0z\_mnth)

##	Ozc	one	Month	
##	Min.	: 1.00	Min.	:5.000
##	1st Qu.	: 18.00	lst Qu.	:6.000
##	Median	: 31.50	Median	:7.000
##	Mean	: 42.13	Mean	:6.993
##	3rd Qu.	: 63.25	3rd Qu.	:8.000
##	Max.	:168.00	Max.	:9.000
##	NA's	:37		

<pre>ranks_0z_mnth &lt;-</pre>	ranks_Oz_mnth %>%
drop_na(Ozone)	%>%
arrange(Ozone)	%>%
mutate(Rank = 1	:nrow(.))

#### Ranks below do not take into account ties!!

#### ranks\_0z\_mnth

##		0zone	Month	Rank
##	1	1	5	1
##	2	4	5	2
##	3	6	5	3
##	4	7	5	4
##	5	7	7	5
##	6	7	9	6
##	7	8	5	7
##	8	9	8	8
##	9	9	8	9
##	10	9	9	10
##	11	10	7	11
##	12	11	5	12
##	13	11	5	13
##	14	11	5	14
##	15	12	5	15
##	16	12	6	16

## Example: calculate ranks (fyi) (2/2)

Ranks below do not take into account ties!!

#### ranks\_0z\_mnth

##		0zone	Month	Rank	
##	1	1	5	1	
##	2	4	5	2	
##	3	6	5	3	
##	4	7	5	4	
##	5	7	7	5	
##	6	7	9	6	
##	7	8	5	7	
##	8	9	8	8	
##	9	9	8	9	
##	10	9	9	10	
##	11	10	7	11	
##	12	11	5	12	
##	13	11	5	13	
##	14	11	5	14	
##	15	12	5	15	
##	16	12	6	16	

Sum of ranks for each group: (not taking into account ties!!)

ranks\_Oz\_mnth %>%
group\_by(Month) %>%
summarise(sumRank = sum(Rank))

##	#	A tib	ole:	5	×	2
##		Month	sum	Rar	۱k	
##		<int></int>	< -	int	_>	
##	1	5		93	39	
##	2	6		43	34	
##	3	7	-	202	23	
##	4	8	-	195	56	
##	5	9	-	143	34	

### K-W test in R

```
kruskal.test(Ozone ~ Month, data = airquality)
```

```
##
Kruskal-Wallis rank sum test
##
data: Ozone by Month
Kruskal-Wallis chi-squared = 29.267, df = 4, p-value = 6.901e-06
```

```
kruskal.test(Ozone ~ Month, data = airquality) %>% tidy() %>% gt()
```

statistic	p.value	parameter	method
29.26658	6.900714e-06	4	Kruskal-Wallis rank sum test

There is sufficient evidence that the median ozone levels are different in at least two months from May - September, 1973 in New York City (p < 0.001; Kruskal-Wallis test).

 (fyi) Since the K-W test is significant, follow-up with pairwise (Wilcoxon) rank-sum tests using a multiple comparison procedure to identify which months have different medians<sub>31/53</sub>

# Permutation tests & bootstrapping

another option to consider

### Permutation tests & bootstrapping

- In some cases we saw that the conditions failed or the sample size was too small for a normal approximation and there were ties in ranks preventing us from using an exact method.
- Another nonparametric option to consider is a permutation test or bootstrapping.
- If you're interested in learning more about this approach, check out the ModernDive Statistical Inference via Data Science book by Chester Ismay and Albert Kim.
  - Ch 7: Sampling
  - Ch 8: Bootstrapping and Confidence Intervals
  - Ch 9: Hypothesis Testing