UCLA STAT 13

Introduction to Statistical Methods for the Life and Health Sciences

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University of California, Los Angeles, Fall 2005 http://www.stat.ucla.edu/~dinov/courses_students.html

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Lecture Set 8

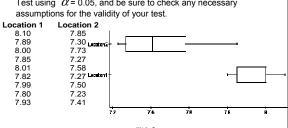
The T Test

Wilcoxon-Mann-Whitney Test

Application

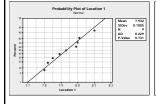
Example: Nine observations of surface soil pH were made two different locations. Does the data suggest that the true mean soil pH values differ for the two locations?

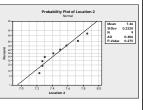
Test using $\alpha = 0.05$, and be sure to check any necessary



Application

To meet the assumption of <u>normality</u> (necessary for the t-test with such a small sample size in each group), we will calculate a normal probability plot for each group.





Application

#1 Formulate hypotheses

 H_o : $\mu_1 - \mu_2 = 0$

(there is no difference between the true mean soil pH of location1 and location2)

 H_a : $\mu_1 - \mu_2$!= 0

(there is a difference between the true mean soil pH of location1 and location2)

Application

• #2 Calculate the test statistic

Descriptive Statistics: Location 1, Location 2

Variable N N* Mean SE Mean StDev Minimum Q1 Median Q3 Location 1 9 0 7.9322 0.0335 0.1005 7.8000 7.8350 7.9300 8.0550 Location 2 9 0 7.4600 0.0740 0.2220 7.2300 7.2700 7.4100 7.6550

Variable Maximum Location 1 8.1000 Location 2 7.8500 $SE_{\overline{y_1}-\overline{y_2}} = \sqrt{\frac{s_1^2}{n_1} + \frac{s_2^2}{n_2}} = \sqrt{\frac{0.1005^2}{9} + \frac{0.222^2}{9}} = 0.081$

$$t_s = \frac{\overline{y}_1 - \overline{y}_2 - 0}{SE_{\overline{y}_1 - \overline{y}_2}} = \frac{7.9322 - 7.460 - 0}{0.081} = 5.827$$

Application

• #3 Calculate the p-value

$$df = \frac{\left(SE_1^2 + SE_2^2\right)^2}{SE_1^4 / n_1 - 1} = \frac{\left(0.0335^2 + 0.074^2\right)^2}{0.0335^4 / 9 - 1} = 11.03 \approx 11 df$$

$$p < 2(0.0005) = 0.001 \text{ (SOCR)}$$

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Application

• #4 Conclusion

Because p < 0.001 < 0.05, we will reject H_o .

CONCLUSION: These data show that there <u>is a statistically significant true mean difference</u> in the <u>pH</u> of <u>Location 1 and Location 2</u> (P < 0.001).

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Application

- ullet Confidence interval for $\mu_1 \mu_2$
 - Suppose we calculated a 95% confidence interval to be:

$$\begin{split} &(\overline{y}_1 - \overline{y}_2) \pm t(df)_{0.025} \left(SE_{\overline{y}_1 - \overline{y}_2}\right) = (7.932 - 7.460) \pm t(11)_{0.025} (0.081) \\ &= (0.472) \pm 2.201 (0.081) \\ &= (0.294, 0.650) \end{split}$$

■ Does this interval surprise you?

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Application

• Corresponding computer output:

Two-Sample T-Test and CI: Location 1, Location 2

Two-sample T for Location 1 vs Location 2 $\,$

N Mean StDev SE Mean Location 1 9 7.932 0.100 0.033 Location 2 9 7.460 0.222 0.074

Difference = mu (Location 1) - mu (Location 2)
Estimate for difference: 0.472222
95% CI for difference: (0.293459, 0.650985)
T-Test of difference = 0 (vs not =): T-Value = 5.81
P-Value = 0.000 DF = 11

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CI and Hypothesis-Testing relationship

- \bullet Consider a 95% confidence interval for $\mu_1-\mu_2$ and it's relationship to the t test at $~\alpha$ = 0.05
 - Both use $\overline{y}_1 \overline{y}_2$ and $SE_{\overline{y}_1 \overline{y}_2}$ in their calculations

CI:
$$(\bar{y}_1 - \bar{y}_2) \pm t(df)_{\alpha/2} (SE_{\bar{y}_1 - \bar{y}_2})$$

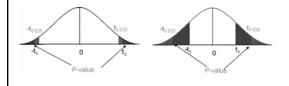
$$\mathsf{T}_{\mathsf{s}} \text{:} \qquad t_{s} = \frac{\left(\overline{y}_{1} - \overline{y}_{2}\right) - 0}{SE_{\overline{y}_{1} - \overline{y}_{2}}}$$

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CI and Hypothesis-Testing relationship

 \bullet With a t test we reject H $_{\!_0}$ if the p-value is less than $~\alpha$ then we reject H $_{\!_0},$ and fail to reject otherwise

 \Box this is the same thing as saying we reject if t_s is beyond $\pm t_{0.025}$, and fail to reject otherwise



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CI and Hypothesis-Testing relationship

• Focusing on the upper half of the distribution and remembering the symmetry: we fail to reject when

$$|T_s| = \frac{|\overline{y}_1 - \overline{y}_2|}{SE_{\overline{y}_1 - \overline{y}_2}} < t_{0.025}$$

■ Further manipulation gives us:

$$\begin{split} & \left| \vec{y}_1 - \vec{y}_2 \right| < t_{0.025}(SE_{\vec{y}_1 - \vec{y}_2}) \\ &= -t_{0.025}(SE_{\vec{y}_1 - \vec{y}_2}) < \vec{y}_1 - \vec{y}_2 < t_{0.025}(SE_{\vec{y}_1 - \vec{y}_2}) \\ &= -(\vec{y}_1 - \vec{y}_2) - t_{0.025}(SE_{\vec{y}_1 - \vec{y}_2}) < 0 < -(\vec{y}_1 - \vec{y}_2) + t_{0.025}(SE_{\vec{y}_1 - \vec{y}_2}) \\ &= (\vec{y}_1 - \vec{y}_2) + t_{0.025}(SE_{\vec{y}_1 - \vec{y}_2}) > 0 > (\vec{y}_1 - \vec{y}_2) - t_{0.025}(SE_{\vec{y}_1 - \vec{y}_2}) \end{split}$$

•Therefore, we fail to reject H_0 : $\mu_1 - \mu_2 = 0$ (for the not equal to alternative), if the confidence interval contains 0.

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CI and Hypothesis-Testing relationship

- If a two-tailed t test and a confidence interval give us the same result, why learn both?
 - There are advantages to each one

□ Confidence interval:

shows magnitude of difference between $\mu_{\rm 1}$ and $\mu_{\rm 2}$

□T test:

has p-value which describes the strength of evidence that $\mu_{\rm 1}$ and $\mu_{\rm 2}$ are really different.

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More on the significance level lpha

· Choose a significance level BEFORE analyzing the data

Example: Say df = 15 and a = 0.05

- If t_s is in either tail we will reject H_o . The chance of this happening is 0.05 -- P(reject H_o) = 0.05, if H_o is true.
- \bullet Because we are assuming that Ho is true, all $t_{\rm s}$ values on the t curve would only deviate from 0 because of sampling error.
- This means:
 - 95% would fail to reject H_o
 - 2.5% would reject H_o (-t_s)
 - 2.5% would reject H_o (t_s)

In other words, a total of 5% would reject Ho when Ho is actually true. This is an incorrect conclusion just because of sampling error!

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More on the significance level lpha

- When we are analyzing one data set in real life at the 0.05 level and our conclusion is to reject Ho there are two possible scenarios:
 - 1. H_o is in fact false
 - 2. H_o is true, but we were unlucky (5%)

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Type I and Type II Errors

- There are two possible mistakes that can be made when conducting a hypothesis test:
 - \blacksquare A type I error is when we reject $\mathrm{H_o}$ and $\mathrm{H_o}$ was true
 - □ P(type I error) = α
 - \Box When we choose lpha before we conduct our test, we are actually protecting ourselves against a type I error
 - ☐ This choice will depend on your experiment
 - A type II error is when we fail to reject H_o and H_o is false
 - \square P(type II error) = β
 - \square β can also be specified before we collect our data
 - $\hfill \square$ will have more to do with the number of observations in our sample

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Type I and Type II Errors

• Table (below) is the best way to summarize

Fail To Reject Ho

rtounty	
H₀ True	H₀ False
Correct	Type II
TN	FN
Type I	Correct
FP	TP
	Correct TN

- You cannot make both errors at the same time
 - Once you have reached a conclusion (reject or fail to reject) based on the data from your experiment you've either made a correct decision or you've made an error (type I for a reject conclusion and type II for a failing-to-reject conclusion)

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Type I and Type II Errors

- Analogy: Think of a car with a car alarm being broken
 - If the alarm goes off for no reason (reject H₀ when H₀ is true) type I error
 - If the car gets broken into and the alarm doesn't go off (fail to reject Ho when Ho is false) - type II error
 - Also consider the sensitivity of the alarm
 - REMEMBER: Fail To Reject Ho means "nothing is going on" or the data do not show otherwise
- Consequences of Type I / II errors are quite different

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Type I and Type II Errors

Example: Measuring pollution in a lake. Say the EPA institutes a rule that companies near bodies of water must test their pollution output. If the company doesn't find any statistical significance in their results, they may continue their current practices.

H_o: No significant pollution

Ha: Significant pollution

In this case a type II error would be much worse (probability of failing to reject H_o when H_o is false – saying no significant pollution when there really is)

An "ethical" company would want to make sure they tested enough samples to guarantee that $\,eta\,$ is small

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Type I and Type II Errors

Example: Drug Treatments. Say a doctor would like to study a new highly toxic drug treatment for cancer patients. There are many risks and side effects of the new drug, but would be of benefit if the proportion of patients responding is greater

 H_o : No significant response (Proportion responding to TX is \leq 0.5) H_a : Significant response (Proportion responding to TX is > 0.5) In this case a type I error is much worse (probability of rejecting H_o when H_o is true – like saying that the TX does something when it really doesn't)

An ethical researcher would want to make sure they keep $\,arphi\,$ small before collecting and analyzing the data

Type I and Type II Errors

- ullet Because lpha is chosen beforehand, we are protected against type I errors. However, type II errors depend on many things, such as sample size (section 7.8)
- β = P(fail to reject H_o) when H_o is false.
 - The chance of rejecting H_o when it is actually false is called the power of
 - \square Power = 1 β = P(reject H_o) when H_o is false
 - measures the ability of the test to detect a difference when a difference really
 - ☐ Power depends on sample size. A larger sample gives more information and
 - ☐ When you plan an analysis you always need to take power into account (ie
 - ☐ decide desired SE and calculate n ☐ analysis of power (7.8)

One Tailed t Tests

- The previous hypothesis test was called a two-tailed (or non-directional) test because Ho was rejected if ts fell in either tail
- In some analyses it is reasonable that there will be a certain direction of a deviation from H_o
 - This means that we are looking to see if one group mean is smaller/larger than the other
- The hypotheses for a one-tailed (or directional) test are:

 H_0 : $\mu_1 - \mu_2 = 0$ H_a : μ_1 - μ_2 > 0 OR

Ha: $\mu_1 - \mu_2 < 0$

■ Note: the null hypothesis doesn't change

One Tailed t Tests

- One-Tailed Test Procedure:
 - Step 1: Check direction to see if data deviate from Ho in the direction specified by Ha (If $\mu_1 < \mu_2$ then we expect to be negative, if $\mu_1 >$ μ_2 then we expect t_s to be positive.)
 - a. If no, then p-value > 0.5
 - b. If yes, then proceed to step 2
 - Step 2: The p-value of the data is the one-tailed area beyond t

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One Tailed t Tests

Example: Cholesterol (cont')

RECALL: Group 1 = Medication, Group 2 = Placebo

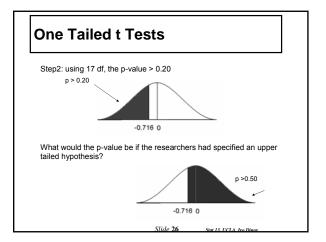
Suppose it is reasonable to assume that $\mu_1 < \mu_2$, in other words the researcher is hoping to show that this new medication lowers cholesterol The appropriate hypotheses would be

 H_0 : $\mu_1 - \mu_2 = 0$ H_a : $\mu_1 - \mu_2 < 0$

Calculate the p-value for this test

Step 1: $t_{\rm s}$ was calculated as -0.716. Check that the data deviate in the direction of H...

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One Tailed t Tests

Example: Soil pH (cont')

Suppose researchers had reason to believe that the soil pH for Location 1 was greater than Location 2.

Two-Sample T-Test and CI: Location 1, Location 2

Two-sample T for Location 1 vs Location 2

N Mean StDev SE Mean

Location 1 9 7.932 0.100 0.033

Location 2 9 7.460 0.222 0.074

Difference = $\mu_1 - \mu_2$

Estimate for difference: 0.472222

95% lower bound for difference: $\,0.326361\,$

T-Test of difference = 0 (vs >): T-Value = 5.81

P-Value = 0.000 DF = 11

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One Tailed t Tests

- P-values for a directional alternative are 1/2 of a nondirectional
 - assuming the direction matched H_a
- It is easier to reject H_o with a one-tailed alternative
 - However it is important that we decide on the direction of H_a before the data is collected
- \bullet If the data do not match the direction of ${\rm H}_a$ we conclude that the data do not indicate that the true means differ
 - However t_s may be statistically significant in the other tail
 □ In this case we would want to look for methodological errors in the experiment

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The Wilcoxon-Mann-Whitney

- Also known as the <u>rank sum test</u>
- This hypothesis test is also used to compare two independent samples
 - This procedure is different from the independent t test because it is valid even if the population distributions are not normal
 - In other words, we can use this test as a fair substitute when we cannot not meet the required normality assumption of the t test
- WMW is called a **distribution-free** type of test or a non-parametric test
 - This test doesn't focus on a parameter like the mean, instead it examines the distributions of the two groups

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The Wilcoxon-Mann-Whitney

- Keep in mind that this is another hypothesis test, so there are still four major parts to consider
- #1 The hypotheses:
 - \blacksquare $\mathrm{H_{o}}\mathrm{:}\$ The population distributions of $\mathrm{Y_{1}}$ and $\mathrm{Y_{2}}$ are the same
 - H_a: The population distributions of Y₁ and Y₂ are the different
 □ This could also be directional: distribution of Y₁ is less than Y₂; OR distribution of Y₁ is greater than Y₂
- #2 The test statistic:
 - denoted by U_s
 - measures the degree of separation between the two samples
 □ a large value of U_s indicates that the two samples are well separated with little overlap
 - $\hfill \square$ a small value of $\hfill U_s$ indicates that the two samples are not well separated with much overlap

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The Wilcoxon-Mann-Whitney

- #3 The p-value:
 - New table!
 - ■http://socr.stat.ucla.edu/Applets.dir/WilcoxonRankSumTable.html
 - Critical Values are in table 6 on p.680
 - Method very similar to using the t table

 $\hfill \Box$ find the appropriate row and then search for a number closest to

- don't need to worry about doubling the p-value for a two-tailed test (assuming we go to the right row header)
- #4 Conclusion:
 - Similar to the conclusion of an independent t test, but not linked to any parameter (for example the difference in means)

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The Wilcoxon-Mann-Whitney

- The Method:
 - Step 1: Arrange the data in increasing order
 - Step 2: Determine K₁ and K₂

 \square K_1 : for each observation in group 1, count the number of observations in the second group that are smaller. Use 1/2 for tied observations. \square K_2 : for each observation in group 2, count the number of observations in the first group that are smaller. Use 1/2 for tied observations.

□ CHECK: if you have done the procedure correctly K₁ + K₂ = n₁n₂

■ Step 3: If the test is non-directional then Us is the larger of K1 and K2. If the test is directional then Us is the K that jives with the direction of Ha (if Ha is Y1>Y2 then Us = K1, if Ha is Y1<Y2 then Us =

■Step 4: Determine the critical value

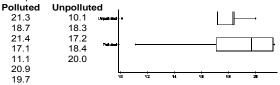
n = larger of n, and n,

 \square n' = smaller of n₁ and n₂

■ Step 5: Bracket the p-value

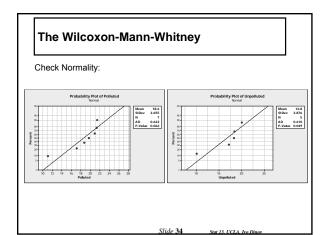
The Wilcoxon-Mann-Whitney

Example: The urinary fluoride concentration (ppm) was measured both for a sample of livestock grazing in an area previously exposed to fluoride pollution and also for a similar sample of livestock grazing in an unpolluted area.



Does the data suggest that the fluoride concentration for livestock grazing in the polluted region is larger that for the unpolluted region? Test using $\alpha = 0.01$.

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The Wilcoxon-Mann-Whitney

#1 The hypotheses:

H_o: urinary fluoride values do not differ between the polluted and unpolluted regions.

H_a: the polluted region has a higher livestock urinary fluoride than the unpolluted region.

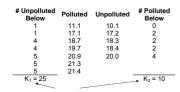
#2 The test statistic:

For this we need to deploy the WMW method shown a few slides earlier.

The Wilcoxon-Mann-Whitney

Let Polluted be group 1, and Unpolluted be group 2

Step 1: arrange the data in increasing order



Step 2: Determine K₄ and K₅

CHECK: 25 + 10 = 35 = (7)(5)

The Wilcoxon-Mann-Whitney

Step 3: H_a : Polluted (Y_1) > Unpolluted (Y_2) so Us is K_1

$$K_2 = 10$$

Step 4:

n = 7

n' = 5

 $\alpha = 0.01$

#3 The p-value:

p > 0.1

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The Wilcoxon-Mann-Whitney

#4 CONCLUSION: These data show that the <u>urinary fluoride concentration</u> ppm for $\underline{\text{livestock grazing}}$ in $\underline{\text{polluted region}}$ is not greater than in the unpolluted region (P>0.1)

NOTE: No mention of the population means!

Corresponding Minitab output:

is significant at 0.1278 \leftarrow

Mann-Whitney Test and CI: Polluted, Unpolluted

N Median Polluted 7 19.700 Unpolluted 5 18.300

Test statistic is calculated using a different formula than our text, but W is the test statistic from

Point estimate for ETA1-ETA2 is 1.400
96.5 Percent of for ETA1-ETA2 is (-2.897,8.602)
W = 53.0 Test of ETA1 = ETA2 vs ETA1 > ETA2

The p-value is calculated using the computer, but is not labeled well.

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The Wilcoxon-Mann-Whitney

Say n = 7, n' = 5 and $U_s = 32$

Two-tailed p-value: 0.01 < p < 0.02 One-tailed p-value: 0.005 < p < 0.01

Say n = 7, n' = 5 and U_s = 36

Impossible, for these sample sizes U_s cannot be larger than 35!

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The Wilcoxon-Mann-Whitney

• Why does this procedure make sense?

Suppose $n_1 = 3$ and $n_2 = 2$

 $K_1 + K_2 = (3)(2) = 6$

we know that $K_1 + K_2$ should sum to 6

The relative magnitudes of K₁ and K₂ indicate the overlap in Y₁



The Wilcoxon-Mann-Whitney

- Conditions for the WMW:
 - Data are from random samples
 - Observations are independent
 - Samples are independent
- Remember: normality will not matter for this test

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Wilcoxon-Mann-Whitney vs. Independent Test

- Both try to answer the same question, but treat data
 - W-M-W uses rank ordering
 - ☐ Positive: doesn't depend on normality or population parameters
 - ☐ Negative: distribution free lacks power because it doesn't use all
 - T-test uses actual Y values
 - $\hfill \square$ Positive : Incorporates all of the data into calculations
 - ☐ Negative : Must meet normality assumption
 - neither is superior
- So...
 - If your data are normally distributed use the t-test
 - If your data are not normal use the WMW test