STAT 516 Lec 05 One-way analysis of variance (review-ish)

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Rust inhibitors example

Data from Kutner et al. (2005).

Ten experimental units assigned to each of four brands of rust inhibitors.

link <- url("https://people.stat.sc.edu/gregorkb/data/KNNLrust.txt")
rust <- read.csv(link,col.names=c("score","brand","rep"),sep = "")
head(rust)</pre>



Do the brands differ in effectiveness? Is there a best brand?

Randomized experiments comparing treatments

- Start with N experimental units (EUs), e.g. subjects, mice, etc.
- Randomly assign each EU to one of a treatment groups.
- Measure on each EU after treatment a response Y.
- Compute the average of the responses in each treatment group...
- Questions we'd like to answer:
 - Is the response mean the same in all treatment groups?
 If not, then which pairs of means are different?

One-way ANOVA setup

reatment effects model.



- \triangleright Y_{ij} is the response for EU *j* in treatment group *i*.
- \blacktriangleright μ represents an overall or baseline mean.
- \triangleright τ_i is the treatment effect for treatment *i*.
- The ε_{ij} are independent Normal $(0, \sigma^2)$ error terms.

Of central interest are the hypotheses

$$H_0$$
: $\tau_i = 0$ for all i versus H_1 : At least one τ_i is nonzero.

If we reject H_0 , we may wish to sort/compare the treatments.

Alternative "cell means model" setup

An alternate version of the model is

$$Y_{ij} = \underbrace{\mu_i}_{\mu_i} + \varepsilon_{ij}, \quad j = 1, \dots, n_i, \quad i = 1, \dots, a,$$

where

In this version of the model the central hypotheses become

$$H_0: \mu_1 = \dots = \mu_a$$
 versus $H_1: \mu_i \neq \mu_j$ for some $i \neq j$.

Goals in one-way ANOVA

Under the one-way ANOVA setup

$$Y_{ij} = \mu + \tau_i + \varepsilon_{ij}, \quad j = 1, \dots, n_i, \quad i = 1, \dots, a,$$



7. Check whether the model assumptions are satisfied.

Rust inhibitors example (cont)

Visually compare the means of several treatment groups with boxplots.



brand

 $Y_{ij} = \mu + \tau_i + \varepsilon_{ij} ,$ Eij ~ Normal (o, or)













Treatment effect estimation in one-way ANOVA

Let
$$N = n_1 + \dots + n_a$$
 and define $\overline{Y} = \frac{1}{N} \sum_{i=1}^{a} \sum_{j=1}^{n_i} Y_{ij}$ as well as
 $\downarrow_{i+1} \downarrow_{i} \downarrow_{i}$
 $\downarrow_{i} \in U_s$
 $\overline{Y}_{i_0} = \frac{1}{n_i} \sum_{j=1}^{n_i} Y_{ij}$ for $i = 1, \dots, a$.
 $\downarrow_{frest mean}$
The two most common ways to estimate $\mu, \tau_1, \dots, \tau_a$:
1. Deviations from baseline parameterization:
 $\overline{Y}_{i_0} = \overline{T}_{i_1} \underbrace{\overline{\gamma}_{i_1} - \overline{\gamma}_{i_2}}_{i_1}$ for $i = 1, \dots, a - 1$ and set $\hat{\mu} = \overline{Y}_{a, \dots}$

2. Deviations from overall mean parameterization:

Set
$$\hat{\tau}_i = \bar{Y}_{i.} - \bar{Y}_{..}$$
 for $i = 1, \dots, a$ and $\hat{\mu} = \bar{Y}_{..}$

Under either parameterization, we have $\hat{\mu} + \hat{\tau}_i = \bar{Y}_{i.}$ for $i = 1, \dots, a$.

$$Y_{ij} = \mu + \tau_c + z_{ij}$$

$$E_{xample}: a = 3 \\ n_1 = n_2 = n_3 = 2 \\ (v_1 = v_2) = 0$$

	ا خر	f,	$ \begin{array}{c} \widehat{\nabla}_{1} = \overline{Y}_{1} \\ \widehat{\nabla}_{2} = \overline{Y}_{2} \\ \widehat{\nabla}_{2} = \overline{Y}_{2} \\ \widehat{\nabla}_{3} = \overline{Y}_{3} \end{array} $			
	ディア・ア・ア・ア	$ \begin{array}{c} & & & \\ & & \\$		$ \left(\begin{array}{c} \overline{Y}_{1}, \ - \overline{Y}_{2}, \\ \overline{Y}_{1}, \ - \overline{Y}_{2}, \\ \overline{Y}_{2}, \ - \overline{Y}_{2}, \\ \overline{Y}_{2}, \ - \overline{Y}_{2}, \\ \overline{Y}_{3}, \ - \overline{Y}_{2}, \\ \overline{Y}_{3}, \ - \overline{Y}_{2}, \\ \overline{Y}_{3}, \ - \overline{Y}_{3}, Y$) ~) ~ .) ~ .) ~ .) ~ .) ~	T. T	
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$$Y_{ij} = (\mu + c) + (2; -c) + \epsilon_{ij}$$

Problem is: I cannot uniquely identify the volues of g and $T_{1}, ..., T_{G}$.

= 1,..., nc.

Solution: Constrain the volves in som very (i) Set $T_a = 0$. (ii) $\prod_{i=1}^{a} T_i = 0$

Rust inhibitors example (cont)

R uses by default the deviations from baseline parameterization:



Estimation of the error term variance σ^2 $Y_{ij} = \rho + \tau_i + \Sigma_{ij}$ $\Sigma_{ij} \sim N_{orm} \wedge (o, \sigma^2)$ As in linear regression, define the $\hat{Y}_{ij} = \hat{\rho} + \hat{c}_i = \bar{Y}_i$. $\hat{fitted values} \hat{Y}_{ij}$ as $\hat{Y}_{ij} = \bar{Y}_i$ for $j = 1, ..., n_i$, and the $\hat{residuals} \hat{\varepsilon}_{ij}$ as $\hat{\varepsilon}_{ij} = Y_{ij} - \bar{Y}_i$. for $j = 1, ..., n_i$, i = 1, ..., a.

Then an unbiased estimator of σ^2 is given by

$$\hat{\sigma}^2 = \frac{1}{N-a} \sum_{i=1}^{a} \sum_{j=1}^{n_i} \hat{\varepsilon}_{ij}^2 = \frac{1}{N-a} \sum_{i=1}^{a} \sum_{j=1}^{n_i} (Y_{ij} - \bar{Y}_{i.})^2.$$
ned to estimate a proton that means to get neidenly

Divide by N-a since the N residuals depend on a estimated quantities...

Rust inhibitors example (cont)

tab <- cbind(rust\$brand,rust\$score,lm_out\$fitted.values,lm_out\$residuals)
colnames(tab) <- c("brand","score","Fitted value","Residual")
head(tab,n = 13)</pre>



The value of $\hat{\sigma}$ is printed in the summary() output:

summary(lm_out)

Call: lm(formula = score ~ as.factor(brand), data = rust) Residuals: Min 1Q Median 3Q Max -4.270 -1.597 0.395 1.275 4.730 Coefficients: Estimate Std. Error t value Pr(>|t|) (Intercept) 0.7836 55.056 <2e-16 *** 43.1400 as.factor(brand)2 46.3000 1.1081 41.782 <2e-16 *** as.factor(brand)3 24.8100 1.1081 22.389 <2e-16 *** 1.1081 -2.409 0.0212 * as.factor(brand)4 -2.6700 Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1 σ Residual standard error: 2.478 on 36 degrees of freedom Multiple R-squared: 0.9863, Adjusted R-squared: 0.9852 F-statistic: 866.1 on 3 and 36 DF, p-value: < 2.2e-16

Sums of squares in the one-way ANOVA model $Y_{ij} = p + \tau_i + \epsilon_{ij}$ $Y_{ij} = \int_{N} \frac{1}{\epsilon_{ij}} \frac{1}{\epsilon$

As in linear regression we decompose the variation in the Y_{ij} by defining:

Total sum of squares: SS_{Tot} =
$$\sum_{i=1}^{a} \sum_{j=1}^{n_i} (Y_{ij} - \bar{Y}_{..})^2$$
Treatment sum of squares: SS_{Trt} = $\sum_{i=1}^{a} n_i (\bar{Y}_{i.} - \bar{Y}_{..})^2$
Error sum of squares: SS_{Error} = $\sum_{i=1}^{a} \sum_{j=1}^{n_i} (Y_{ij} - \bar{Y}_{..})^2$
We have SS_{Tot} = SS_{Trt} + SS_{Error}.
Note that SS_{Trt} is computed just like SS_{Reg} in linear regression.
We again define $R^2 = \frac{SS_{Trt}}{SS_{Tot}}$.
Reg = $\sum_{i=1}^{a} \sum_{j=1}^{n_i} (\hat{Y}_{ij} - \bar{Y}_{..})^2 = \sum_{i=1}^{a} \sum_{j=1}^{n_i} (\bar{Y}_{i.} - \bar{Y}_{..})^2 = \sum_{i=1}^{a} \sum_{j=1}^{n_i} (\bar{Y}_{i.} - \bar{Y}_{..})^2$

Sampling distributions of our sums of squares

The SS, appropriately scaled, follow chi-square distributions:

$$SS_{Tot} / \sigma^2 \sim \chi^2_{N-1}(\phi_{Tot})$$

$$SS_{Trt} / \sigma^2 \sim \chi^2_{a-1}(\phi_{Trt})$$

$$SS_{Error} / \sigma^2 \sim \chi^2_{N-a},$$

where $\phi_{\rm Tot}$ and $\phi_{\rm Trt}$ are noncentrality parameters.

The mean squares in the one-way ANOVA model

Dividing $\mathrm{SS}_{\mathrm{Trt}}$ and $\mathrm{SS}_{\mathrm{Error}}$ by their dfs, we define:





$$N = n_1 + \dots + n_a$$

 $q = 4$
 $N = a = 40 - 4 = 36$
 $N = 40$

Rust inhibitors example (cont)

Obtain the ANOVA table with the anova() function on the lm() output.



Testing whether there is any difference in treatment means

Given
$$Y_{ij} = \mu + \tau_i + \varepsilon_{ij}$$
, $j = 1, ..., n_i$, $i = 1, ..., a$, we wish to test
 $H_0: \tau_i = 0$ for all i versus $H_1:$ At least one τ_i is nonzero.
none of the tradiment means differs from the baseline.
We use the overall F test of significance:
1. Compute $F_{\text{stat}} = \underbrace{MS_{\text{Trt}}}_{MS_{\text{Error}}}$ Under H_0
2. Reject H_0 at α if $F_{\text{stat}} > F_{a-1,N-a,\alpha}$.

3. Obtain p-value as $P(F>F_{\rm stat})$, where $F\sim F_{a-1,N-a}.$

The value of $F_{\rm stat}$ and the p-value are printed in the summary() output.



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Exercise: For which data set will the F-statistic be largest/smallest?





Exercise: Compute F_{stat} for the rust data using the summary info:





$$= \frac{1}{36} (10-1) \sum_{i=1}^{4} 5_{i}^{2} = \frac{9}{36} ((3.0)^{2} + (2.22)^{2} + (2.13)^{2} + (2.14)^{2})^{2}$$

$$= 6.15$$

Post-hoc comparisons of means

- Lafter F test rejects Ho: M= ... = M.
 - If we reject $H_0: \mu_1 = \dots = \mu_a$, then we may wish to compare means.
 - Call such comparisons post-hoc as they follow the F-test.
 - We may wish to compare several pairs of means, which is like testing several hypotheses at once.
 - When several hypotheses are tested at once, the <u>familywise Type I</u> error rate is the probability that any Type I error is committed.
 - We discuss two methods for post-hoc comparisons of means which control the familywise Type I error rate.

Comparing all pairs of means Minny Ma. Compone Mi and Mi for Ill ifj. Test Ho: Mi=Mi vs Hi: MifMi for ifj. We want to build a CI for $\mu_i - \mu_j$ for all pairs $i \neq j$. Suppose the design is balanced, i.e. $n_i = n$ for all i = 1, ..., a. $\frac{\bar{Y}_{i.} - \bar{Y}_{j.} \pm t_{a(n-1),\alpha/2} \hat{\sigma} \sqrt{2/n}}{\text{CF. for a Given a in difference in differe$ lf we build for all $i \neq j$ the ordinary $(1 - \alpha) \times 100\%$ CIs each one will cover its target with probability $1 - \alpha$. \triangleright We want simultaneous coverage with probability $1 - \alpha$. I.e., we want the familywise coverage of all the intervals to be $1 - \alpha$. q = 4 $\begin{pmatrix} 4 \\ 2 \end{pmatrix} = \frac{4!}{2!(4-2)!} = \frac{4\cdot 3\cdot 2\cdot 1}{2\cdot 1(2\cdot 1)} = 6.$ a groups $\begin{pmatrix} a \\ 2 \end{pmatrix} = \frac{a!}{2!(a-2)!}$

Multiple comparisons of means with Tukey's HSD HSD: Honest significant difference.

Suppose the design is <u>balanced</u>, i.e. n_i = n for all i = 1,..., a.
 Suppose we could find the value q_{a,a(n-1),α} such that

$$P\left(\max_{i\neq j}\left\{\frac{|(\bar{Y}_{i},-\bar{Y}_{j})-(\mu_{i}-\mu_{j})|}{\hat{\sigma}/\sqrt{n}}\right\} \leq q_{a,a(n-1),\alpha}\right) = 1-\alpha.$$
Then with probability $1-\alpha$ the Cls
$$\overline{Y}_{i},-\overline{Y}_{j},\pm q_{a,a(n-1),\alpha}\hat{\sigma}/\sqrt{n}$$
will simultaneously cover the targets $\mu_{i}-\mu_{j}$ for all $i\neq j$. Show!
Tukey made tables of the values $q_{a,a(n-1),\alpha}$.
Can use the simultaneous intervals to sort/compare the means.

t treatment groups

$$P\left(\begin{array}{c} m \cdot x \\ if_{j} \end{array}\right) \left(\begin{array}{c} (\overline{y}_{i}, -\overline{y}_{j},) - (m_{i} - m_{j}) \\ \hline \sigma / \sqrt{n} \end{array}\right) = g_{\alpha} = 1 - \alpha$$

$$P\left(\begin{array}{c} (1)\\ i\neq j\end{array}\right) \left\{ \begin{array}{c} (\overline{Y}_{i},-\overline{Y}_{j},)-(\mu_{i}-\mu_{j})\\ \hline \sigma/\sqrt{n}\end{array}\right\} = I-\alpha$$

"





Samittaneouly aver pi-pi, ifj, with prob 1-a.



Figure 1: Table A.6 from Mohr, Wilson, and Freund (2021)

Rust inhibitors example (cont)

For the rust data we have n = 10 and a = 4. At $\alpha = 0.05$ we have $q_{a,a(n-1),\alpha} = q_{4,36,0.05} \approx 3.85$ from table. Obtain exact value with qtukey(.95, 4, 36) = 3.8087984. Build the Tukey HSD CI for $\mu_2 - \mu_1$. A 2 0 MSE <- sum(lm_out\$residuals^2) / (40 - 4) <y1bar <- mean(rust\$score[rust\$brand == 1]) <- Y.. n=10 lo21 <- y2bar - y1bar - (qtukey(.95,4,36) * sqrt(MSE) / sqrt(10) up21 <- y2bar - y1bar + qtukey(.95,4,36) * sqrt(MSE) / sqrt(10) c(lo21,up21)



Rust inhibitors example (cont)

Use TukeyHSD() on aov() output to obtain the simultaneous Cls.

```
# must use the aov() function instead of the lm() function
       aov_out <- aov(score ~ as.factor(brand), data = rust)
       TukeyHSD(aov out)
         Tukey multiple comparisons of means
           95% family-wise confidence level
       Fit: aov(formula = score ~ as.factor(brand), data = rust)
                                                                       Dunnetté, for compension
       $`as.factor(brand)`
              diff
                                               p adj
                                       upr
       2-1 46.30 43.315536 49.2844635 0.0000000
M2- M1
       3-1 24.81 21.825536 27.7944635 0.0000000
                                                                    43.582516 49.017484
       4-1 -2.67 -5.654464 0.3144635 0.0933303
       3-2 -21.49 -24.474464 -18.5055365 0.000000
4-2 -48.97 -51.954464 -45.9855365 0.000000
       4-3 -27.48 -30.464464 -24.4955365 0.0000000
                          Con trust then composisons.
            Y: - Y;
```





Differences in mean levels of as.factor(brand)

Comparison of treatments with a baseline treatment

Tykey's is for comparing <u>all pairs</u> of means: (2) pairs. Let treatment a is baseline. Company Mi to Ma, i=1,..., a-1.

It may be that not all pairwise comparisons are of interest.

Then Tukey's method is too conservative (Cls wider than necessary).

Say we want to compare all treatments to a "baseline" treatment.

Build CIs for $\mu_j - \mu_a$, j = 1, ..., a - 1, a the baseline treatment.

This is a - 1 Cls instead of $\begin{pmatrix} a \\ 2 \end{pmatrix}$ Cls.

Can use Dunnett's method.

Dunnett's method for comparisons with a baseline

Assume n_i = n for all i (balanced case).
 Given a value d_{n,a(n-1),\alpha} such that

$$P\left(\max_{\substack{1 \leq i \leq a-1}} \left| \frac{(\bar{Y}_{i} - \bar{Y}_{a.}) - (\mu_{i} - \mu_{a})}{\hat{\sigma}\sqrt{2/n}} \right| \leq d_{n,a(n-1),\alpha} \right) = 1 - \alpha,$$

with probability $1-\alpha$ the CIs

$$\bar{Y}_{\rm L} - \bar{Y}_{a.} \pm d_{n,a(n-1),\alpha} \hat{\sigma} \sqrt{2/n}$$

will simultaneously cover the targets $\mu_i - \mu_a$ for all i = 1, ..., a - 1.

- Dunnett made tables of the values $d_{n,a(n-1),\alpha}$.
 - Cannot sort the means after Dunnett's.

- 4		Two-sided α	Wumber of Groups Counting Both Treatments and Control						
-	Error df		2	3	4	5	6	7	8
= 10	trol 5	0.05	2.57	3.03	3.29	3.48	3.62	3 73	3.82
the "	5	0.01	4.03	4.63	4.97	5.22	5.02	5.56	5.68
1 1 15 11	6	0.05	2.45	2.86	3.10	3.26	3 39	3.49	3.57
Brand	6	0.01	3.71	4.21	4 51	4 71	4.87	5.00	5.10
	7	0.05	2.36	2.75	2.97	3.12	3.24	3 33	3.41
	7	0.01	3.50	3.95	4.21	4.39	4 53	4.64	4.74
	8	0.05	2.31	2.67	2.88	3.02	3.13	3.22	3 29
	8	0.01	3.36	3.77	4.00	4 17	4 29	4 40	4.48
	9	0.05	2.26	2.61	2.81	2.95	3.05	3 1 4	3.20
2 - 191	9	0.01	3.25	3.63	3.85	4.01	4.12	4.22	4.30
	10	0.05	2.23	2.57	2.76	2.89	2.99	3.07	3.14
$\chi = \mu_1$	10	0.01	3.17	3.53	3.74	3.88	3.99	4.08	4.16
• /	11	0.05	2.20	2.53	2.72	2.84	2.94	3.02	3.08
	11	0.01	3.11	3.45	3.65	3.79	3.89	3.98	4.05
$\mu - \mu_1$	12	0.05	2.18	2.50	2.68	2.81	2.90	2.98	3.04
• /	12	0.01	3.05	3.39	3.58	3.71	3.81	3.89	3.96
	13	0.05	2.16	2.48	2.65	2.78	2.87	2.94	3.00
	13	0.01	3.01	3.33	3.52	3.65	3.74	3.82	3.89
	14	0.05	2.14	2.46	2.63	2.75	2.84	2.91	2.97
= 0.05	14	0.01	2.98	3.29	3.47	3.59	3.69	3.76	3.83
~ 2	15	0.05	2.13	2.44	2.61	2.73	2.82	2.89	2.95
- + (4)0 -	15	0.01	2.95	3.25	3.43	3.55	3.64	3.71	3.78
4 1 (2.77) VN	16	0.05	2.12	2.42	2.59	2.71	2.80	2.87	2.92
	16	0.01	2.92	3.22	3.39	3.51	3.60	3.67	3.73
	17	0.05	2.11	2.41	2.58	2.69	2.78	2.85	2.90
	17	0.01	2.90	3.19	3.36	3.47	3.56	3.63	3.69
	18	0.05	2.10	2.40	2.56	2.68	2.76	2.83	2.89
	18	0.01	2.88	3.17	3.33	3.44	3.53	3.60	3.66
da a(n-1)	19	0.05	2.09	2.39	2.55	2.66	2.75	2.81	2.87
	19	0.01	2.86	3.15	3.31	3.42	3.50	3.57	3.63
	20	0.05	2.09	2.38	2.54	2.65	2.73	2.80	2.86
	20	0.01	2.85	3.13	3.29	3.40	3.48	3.55	3.60
du 30,000 \	20	0.01	2.06	2.34	2.50	2.61	2.69	2.75	2.81
	25	0.05	2 79	3.06	3.21	3.31	3.39	3.45	3.51
	30	0.05	2.04	2.32	2.47	2.58	2.66	2.72	2.77
1	30	0.03	2.75	3.01	3.15	3.25	3.33	3.39	3.44
T	6 1 1	0.01	2.02	2.29	2.44	2.54	2.62	2.68	2.73
	40	0.03	2.70	2.95	3.09	3.19	3.26	3.52	3.37
	- 0 40 60	0.01	2.00	2.27	2.41	2.51	2.58	2.64	2.69
1.1.6	00	0.05	2.66	2.90	3.03	3.12	5.19	5.25	5.29

Figure 2: Table A.5 from Mohr, Wilson, and Freund (2021)

Rust inhibitor data (cont)

For the rust data we have n = 10 and a = 4.

At
$$\alpha = 0.05$$
 we have $d_{a,a(n-1),\alpha} = d_{4,36,0.05} \approx 2.47$
Use value 2.4 in the table (should be close).

[1] 43.59615 49.00385

```
Rust inhibitor data (cont)
     Use DunnettTest() from R package DescTools.
     library(DescTools) # first time run install.packages("DescTools")
     Dunnett_out <- DunnettTest(score ~ as.factor(brand), data = rust, control = "1")</pre>
     Dunnett_out
                                                                        specify control/
bealine treatment
       Dunnett's test for comparing several treatments with a control :
         95% family-wise confidence level
     $`1`
          diff
                  lwr ci
                          <u>upr.ci</u> pval
         46.30 43.582516 49.017484 20 16 ***
     3-1 24.81 22.092516 27.527484 <2e-16 ***
     4-1 -2.67 -5.387484 0.047484 0.0549 .
     Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```



Dunnett's vs Tukey's

- Tukey's is for comparisons between all pairs of means.
- Dunnett's is for comparison of means with a baseline.
- So Tukey's must make greater adjustments to control the familywise Type I error.
- Tukey intervals will be wider than Dunnett intervals.
- Tukey's allows you to sort the means, while Dunnett's does not.

If building *B* CIs you can ALWAYS use the Bonferroni correction:

Build each CI ordinarily, but use α/B instead of α.
 Ensures simultaneous coverage of all CIs with probability ≥ 1 - α.
 True probability of simultaneous coverage may be greater.
 Bonferroni-corrected CIs will be wider than Dunnett's and wider than Tukey's if used for making those same comparisons.

Use when we do not know how to adjust for multiple comparisons.

$$d = 0.05$$

$$35\%$$

$$3 = 0.005$$

$$3 = 0.005$$

$$3 = 0.005$$

$$3 = 0.005$$

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$$3 = 0.005$$

$$3 = 0.005$$

$$3 = 0.005$$

Rust inhibitor data (cont) $\overline{Y}_{3.} - \overline{Y}_{9.} \neq \pm 36, \begin{pmatrix} 0.05\\2\\2\\ \end{array}$ 40-4 = 36 Compare Brand 3 to 4 and Brand 1 to 3, using the Bonferroni correction to control the familywise error rate. y1bar <- mean(rust\$score[rust\$brand == 1]) %.</pre> y3bar <- mean(rust\$score[rust\$brand == 3]) $\overline{Y_3}$. y4bar <- mean(rust\$score[rust\$brand == 4]) Yy. alpha <- 0.05 (t36, (0:05)/2 5 B <- 2 = # (.I.S $me \leftarrow qt(1 - (alpha/B)/2, a*(n-1)) * sqrt(MSE) * sqrt(2/n)$ tab <- rbind(c(y3bar - y4bar - me,y3bar - y4bar + me),</pre> c(y1bar - y3bar - me,y1bar - y3bar + me)) rownames(tab) <- c("3-4","1-3") colnames(tab) <- c("lower","upper")</pre> tab lower upper 3-4 24.888 30.072 1-3 -27.402 -22.218

Checking model assumptions

$$Y_{ij} = \mu + \tau_i + \epsilon_{ij}, \quad \epsilon_{ij} \sim N(0, \sigma^2)$$

treatment men

Validity of the foregoing analyses depends on these assumptions:

The responses are normally distributed around the treatment means (Check QQ plot of residuals).

The response has the same variance in all treatment groups (Check residuals vs fitted values plot).

3. The response values are independent of each other (No way to check; must trust experimental design).

Rust inhibitors example (cont)

plot(lm_out,which = 2)



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Rust inhibitors example (cont)

plot(lm_out,which = 1)



Residuals vs Fitted

Perception of slope example WILL NOT TEST ABOUT LEVENES TEST.

Do axis re-scalings affect how we perceive an x-y relationship?

For a single data set with data pairs (X_i, Y_i) , with $X_i \sim \text{Normal}(0, 1)$ and $Y_i = \text{Normal}(X_i, 1)$ for $i = 1, \dots, 50$, three scatterplot treatments were constructed:

- 1. "Control" used x and y plotting limits given by the range of the data.
- 2. "X" extended the x-limits by 1.5 in each direction.
- 3. "Y" extended the y-limits by 1.5 in each direction.

Each student in a class was randomly assigned a scatterplot and told to draw with a ruler the best-fitting line through the data. The slope of each student-drawn line was measured and recorded as the response.

Is the response mean the same in the three treatment groups?

An artifact from each treatment group:



Figure 3: "Control"





boxplot(slope ~ trt)



```
lm_slope <- lm(slope ~ as.factor(trt))
summary(lm_slope)</pre>
```

Call: lm(formula = slope ~ as.factor(trt)) Residuals: Min 1Q Median 3Q Max -0.9222 -0.2847 -0.1293 0.2628 1.3478 Coefficients:

Estimate Std. Error t value Pr(>|t|) (Intercept) 1.36857 0.18161 7.536 2.12e-07 *** as.factor(trt)X 0.05143 0.24868 0.207 0.838 as.factor(trt)Y 0.17365 0.24215 0.717 0.481 ---Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Residual standard error: 0.4805 on 21 degrees of freedom Multiple R-squared: 0.02614, Adjusted R-squared: -0.06661 F-statistic: 0.2818 on 2 and 21 DF, p-value: 0.7572



Q–Q Residuals

lm(slope ~ as.factor(trt))



lm(slope ~ as.factor(trt))

Levene's test for equality of variances

$$H_0: G_1^2 = G_2^2 = \sigma_3^2$$

Checks if the mean magnitude of the residuals is equal across groups:

1. Obtain the residuals $\hat{\varepsilon}_{ij}$ from the one-way ANOVA model.

- 2. Treat the absolute values $|\hat{\varepsilon}_{ij}|$ of the residuals as *new* responses.
- 3. Test for equal means of the new responses with the F test.

So, do the ordinary F-test with the $|\hat{\varepsilon}_{ij}|$ as the responses.

Perception of slope example (cont)



Can also use the leveneTest() function in the R package car.

```
library(car)
leveneTest(slope~as.factor(trt),center = mean)
```

```
Levene's Test for Homogeneity of Variance (center = mean)

Df F value Pr(>F)

group 2 4.5652 0.02258 *

21

----

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

We conclude that the variances are *not* equal across treatment groups.

References

Kutner, Michael H, Christopher J Nachtsheim, John Neter, and William Li. 2005. *Applied Linear Statistical Models*. McGraw-hill.
Mohr, Donna L, William J Wilson, and Rudolf J Freund. 2021. *Statistical Methods*. Academic Press.