

Lesson 8: Chip Melting Times

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Background

Industrial-scale baking requires a precise understanding of how recipe ingredients behave when baked. For example, bakers want to know and understand how different types of baking chips respond to different oven temperatures. Researchers selected a sample of 31 participants to compare the melting times of chocolate chips and butterscotch chips. The response variable will be the amount of time, from placing a chip on a participant's tongue and holding it steadily to the roof of their mouth, until the chip is completely melted.

```
library(tidyverse)
library(pwr) # see https://www.statmethods.net/stats/power.html for good tutorial.

## Load data
chipmelt <- read.table('http://www.isi-stats.com/isi2/data/MeltingTimesUnstacked.txt', header = TRUE, stringsAsFactors = TRUE)

chipmelt2 <- read.table('MeltingTimes.txt', header = TRUE, stringsAsFactors = TRUE)
```

Step 1: Ask a research question: “Is there an association between chip type and melting time?”

Step 2: Design a study and collect data:

1. Create a sources of variation table.

Observed variation in:	Sources of explained variation:	Sources of unexplained variation:
melting time	type of chip (chocolate/butterscotch)	body temperature
		saliva amount (by person)
		chip size
Inclusion criteria: ?? (class? people?)		chip temperature
Design: same technique for “melting technique” for each person		Unknown

2. Develop a study protocol to investigate this question. What is your inclusion criteria? Will you use randomization?

Have a lot of participants and randomly assign them to a chocolate or butterscotch chip. Use a stopwatch to measure how long the chip is completely dissolved. Inclusion criteria can be any group within reason.

3. Think about the sources of variation diagram. Do any of your answers have anything in common? Can you suggest a way to improve the model?

Many of the unexplained sources of variation come from the (same) participant. We could improve the model by having each participant melt both types of chips.

4. Is a repeated measures type of matched pairs design feasible for this study? Describe how it would work and the benefits. Would randomization still need to apply here? How?

Yes a repeated measures, matched pairs design is feasible. Each participant can receive both treatments (both chips). This would keep person-specific variables more constant (and reduce variability). Yes, randomization will still need to apply. If we don't randomize order of taking the chip we will have the order confound the treatment.

5. Update the sources of variation table when using a repeated measures, matched pairs design.

Observed variation in:	Sources of explained variation:	Sources of unexplained variation:
melting time	type of chip (chocolate/butterscotch)	chip size
	body temperature	chip temperature
	saliva amount (by person)	Unknown
Inclusion criteria: ?? (class? people?)		
Design: same technique for "melting technique" for each person		

6. Would a two-sample t-test (or separate means model) as in Chapter 1 be an appropriate analysis to compare the two melting time means for our matched pairs study design? Explain.

No. The two-sample t-test requires independent observations. Since we have people take both treatments we cannot assume independence.

Step 3: Explore the Data.

7. The data set you are currently working with is in what we consider "wide format" where each participant appears in only one row. Often times in our analysis it helps to have our data in "long format" where participants appear multiple times based on the levels of another variable. Use the `pivot_longer()` to pivot the data set and rename it `chips_long`. Keep in mind that you must tell R the columns you want to pivot (`cols =`), the new variable that you want to contain the current variable names of the columns you are pivoting (`names_to =`) and the new variable that you want to contain the current values of the columns you are pivoting (`values_to =`). Hint: use the help file in R for `pivot_longer`.

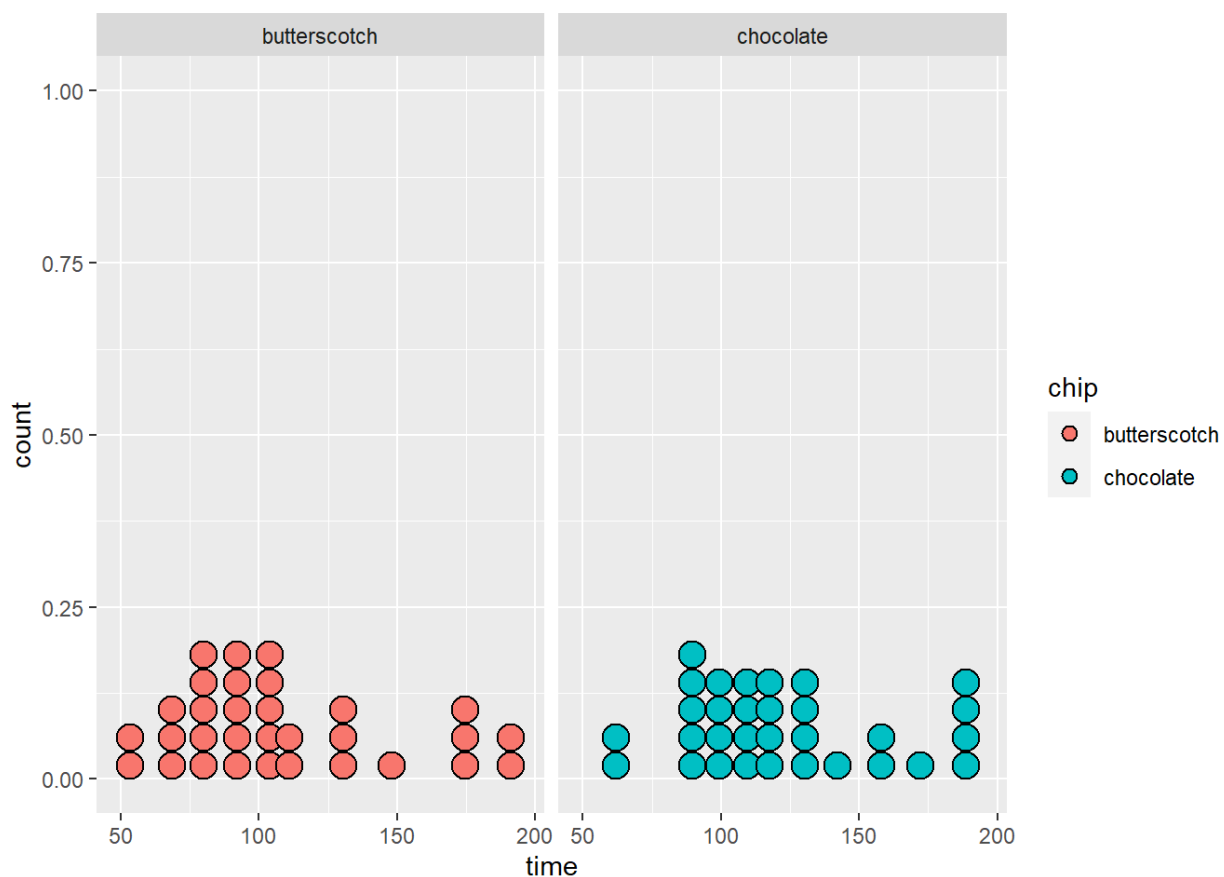
```
chips_long <- chipmelt %>% pivot_longer(cols=-student, names_to = "chip", values_to = "time")
```

Update the “chip” column to be a factor variable.

```
chips_long <- chips_long %>% mutate(chip = as.factor(chip))
```

8. Create a dotplot (using `ggplot` and `geom_dotplot`), stratified by chip type. Does it appear that one chip has a longer melting time than the other?

```
chips_long %>% ggplot(aes(x=time, fill = chip)) + geom_dotplot(binwidth = 10) + facet_wrap(~chip)
```



Yes, it appears that the chocolate chips take slightly longer to melt.

9. Calculate the effect of each type of chip.

```
mn <- mean(chips_long$time)
chips_long %>% group_by(chip) %>% summarise(mean(time) - mn)
```

chip <fct>	mean(time) - mn <dbl>
butterscotch	-7.048387
chocolate	7.048387
2 rows	

10. Conduct a separate means model using the type of chip as our explanatory variable. Remember: we can't use the R-squared value here.

```
sep.model <- lm(time~0+chip, data=chips_long)
summary(sep.model) # do not use R-squared!
```

```
##
## Call:
## lm(formula = time ~ 0 + chip, data = chips_long)
##
## Residuals:
##      Min       1Q   Median       3Q      Max
## -62.484 -27.460  -7.935  20.339  86.613
##
## Coefficients:
##              Estimate Std. Error t value Pr(>|t|)
## chipbutterscotch  108.387      6.795   15.95  <2e-16 ***
## chipchocolate    122.484      6.795   18.02  <2e-16 ***
## ---
## Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
##
## Residual standard error: 37.83 on 60 degrees of freedom
## Multiple R-squared:  0.9061, Adjusted R-squared:  0.903
## F-statistic: 289.7 on 2 and 60 DF,  p-value: < 2.2e-16
```

Step 4: Draw inferences beyond the data.

11. Conduct a separate means model using the type of chip as explanatory variable, but use effect encoding (use `contrasts()` and `contr.sum`). This will allow us to use R to find the correct R-squared value (and other things)). Also use `anova()` and `aov()` to calculate SST, SSE, SSM, F statistic, p-value.

```
contrasts(chips_long$chip) <- contr.sum  
contrasts(chips_long$chip)
```

```
##           [,1]  
## butterscotch    1  
## chocolate      -1
```

```
eff.model <- lm(time ~ chip, data = chips_long)  
summary(eff.model)
```

```
##  
## Call:  
## lm(formula = time ~ chip, data = chips_long)  
##  
## Residuals:  
##      Min       1Q   Median       3Q      Max   
## -62.484 -27.460  -7.935   20.339   86.613   
##  
## Coefficients:  
##              Estimate Std. Error t value Pr(>|t|)      
## (Intercept)  115.435      4.805   24.024  <2e-16 ***  
## chip1        -7.048      4.805   -1.467    0.148   
## ---  
## Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1  
##  
## Residual standard error: 37.83 on 60 degrees of freedom  
## Multiple R-squared:  0.03462,    Adjusted R-squared:  0.01853   
## F-statistic: 2.152 on 1 and 60 DF,  p-value: 0.1476
```

```
aov(eff.model)
```

```
## Call:
##   aov(formula = eff.model)
##
## Terms:
##             chip Residuals
## Sum of Squares   3080.15 85887.10
## Deg. of Freedom      1      60
##
## Residual standard error: 37.83453
## Estimated effects are balanced
```

```
anova(eff.model)
```

	Df <int>	Sum Sq <dbl>	Mean Sq <dbl>	F value <dbl>	Pr(>F) <dbl>
chip	1	3080.145	3080.145	2.151763	0.1476284
Residuals	60	85887.097	1431.452	NA	NA
2 rows					

$R^2 = 0.03462$, $SST = 88,967.24$, $SSE = 85,894.42$, $SSM (SS_{Chip}) = 3,072.82$, $F\text{-stat} = 2.1518$, $p\text{-value} = 0.1476$.

12. How do we take into account the 'person effect'? Two approaches: 1) take the differences for each individual. 2) Use a separate means model. Which method is preferred and why?

For the differences model we can conduct a regular t-test. However, this will only tell us that there is a significant association between chip type and melting time. The separate means model will tell us how much of the variation in mean melting time is explained by the chip type used.

13. Conduct a separate means model using the type of chip as explanatory variable and account for the person to person variation. Use effect encoding (use `contrasts()`) as we did in the previous model. Provide the R^2 , SST , SSE , SSM , F statistic(s), p -value(s).

```
contrasts(chips_long$student) <- contr.sum
contrasts(chips_long$student)
```

##	[,1]	[,2]	[,3]	[,4]	[,5]	[,6]	[,7]	[,8]	[,9]	[,10]	[,11]	[,12]	[,13]	[,14]
## BB	1	0	0	0	0	0	0	0	0	0	0	0	0	0
## CL	0	1	0	0	0	0	0	0	0	0	0	0	0	0
## DC	0	0	1	0	0	0	0	0	0	0	0	0	0	0
## EH	0	0	0	1	0	0	0	0	0	0	0	0	0	0
## EP	0	0	0	0	1	0	0	0	0	0	0	0	0	0
## GA	0	0	0	0	0	1	0	0	0	0	0	0	0	0
## HH	0	0	0	0	0	0	1	0	0	0	0	0	0	0
## IQ	0	0	0	0	0	0	0	1	0	0	0	0	0	0
## JO	0	0	0	0	0	0	0	0	1	0	0	0	0	0
## JR	0	0	0	0	0	0	0	0	0	1	0	0	0	0
## JS	0	0	0	0	0	0	0	0	0	0	1	0	0	0
## JvK	0	0	0	0	0	0	0	0	0	0	0	1	0	0
## KH	0	0	0	0	0	0	0	0	0	0	0	0	1	0
## KH2	0	0	0	0	0	0	0	0	0	0	0	0	0	1
## KH3	0	0	0	0	0	0	0	0	0	0	0	0	0	0
## KJ	0	0	0	0	0	0	0	0	0	0	0	0	0	0
## LC	0	0	0	0	0	0	0	0	0	0	0	0	0	0
## LO	0	0	0	0	0	0	0	0	0	0	0	0	0	0
## LR	0	0	0	0	0	0	0	0	0	0	0	0	0	0
## MB	0	0	0	0	0	0	0	0	0	0	0	0	0	0
## MG	0	0	0	0	0	0	0	0	0	0	0	0	0	0
## ML	0	0	0	0	0	0	0	0	0	0	0	0	0	0
## NH	0	0	0	0	0	0	0	0	0	0	0	0	0	0
## NL	0	0	0	0	0	0	0	0	0	0	0	0	0	0
## NN	0	0	0	0	0	0	0	0	0	0	0	0	0	0
## PS	0	0	0	0	0	0	0	0	0	0	0	0	0	0
## RUR	0	0	0	0	0	0	0	0	0	0	0	0	0	0
## SG	0	0	0	0	0	0	0	0	0	0	0	0	0	0
## SS	0	0	0	0	0	0	0	0	0	0	0	0	0	0
## TA	0	0	0	0	0	0	0	0	0	0	0	0	0	0
## TS	-1	-1	-1	-1	-1	-1	-1	-1	-1	-1	-1	-1	-1	-1
##	[,15]	[,16]	[,17]	[,18]	[,19]	[,20]	[,21]	[,22]	[,23]	[,24]	[,25]	[,26]		
## BB	0	0	0	0	0	0	0	0	0	0	0	0		
## CL	0	0	0	0	0	0	0	0	0	0	0	0		
## DC	0	0	0	0	0	0	0	0	0	0	0	0		
## EH	0	0	0	0	0	0	0	0	0	0	0	0		
## EP	0	0	0	0	0	0	0	0	0	0	0	0		
## GA	0	0	0	0	0	0	0	0	0	0	0	0		
## HH	0	0	0	0	0	0	0	0	0	0	0	0		
## IQ	0	0	0	0	0	0	0	0	0	0	0	0		
## JO	0	0	0	0	0	0	0	0	0	0	0	0		

## JR	0	0	0	0	0	0	0	0	0	0	0	0
## JS	0	0	0	0	0	0	0	0	0	0	0	0
## JvK	0	0	0	0	0	0	0	0	0	0	0	0
## KH	0	0	0	0	0	0	0	0	0	0	0	0
## KH2	0	0	0	0	0	0	0	0	0	0	0	0
## KH3	1	0	0	0	0	0	0	0	0	0	0	0
## KJ	0	1	0	0	0	0	0	0	0	0	0	0
## LC	0	0	1	0	0	0	0	0	0	0	0	0
## LO	0	0	0	1	0	0	0	0	0	0	0	0
## LR	0	0	0	0	1	0	0	0	0	0	0	0
## MB	0	0	0	0	0	1	0	0	0	0	0	0
## MG	0	0	0	0	0	0	1	0	0	0	0	0
## ML	0	0	0	0	0	0	0	1	0	0	0	0
## NH	0	0	0	0	0	0	0	0	1	0	0	0
## NL	0	0	0	0	0	0	0	0	0	1	0	0
## NN	0	0	0	0	0	0	0	0	0	0	1	0
## PS	0	0	0	0	0	0	0	0	0	0	0	1
## RUR	0	0	0	0	0	0	0	0	0	0	0	0
## SG	0	0	0	0	0	0	0	0	0	0	0	0
## SS	0	0	0	0	0	0	0	0	0	0	0	0
## TA	0	0	0	0	0	0	0	0	0	0	0	0
## TS	-1	-1	-1	-1	-1	-1	-1	-1	-1	-1	-1	-1
##	[,27]	[,28]	[,29]	[,30]								
## BB	0	0	0	0								
## CL	0	0	0	0								
## DC	0	0	0	0								
## EH	0	0	0	0								
## EP	0	0	0	0								
## GA	0	0	0	0								
## HH	0	0	0	0								
## IQ	0	0	0	0								
## JO	0	0	0	0								
## JR	0	0	0	0								
## JS	0	0	0	0								
## JvK	0	0	0	0								
## KH	0	0	0	0								
## KH2	0	0	0	0								
## KH3	0	0	0	0								
## KJ	0	0	0	0								
## LC	0	0	0	0								
## LO	0	0	0	0								
## LR	0	0	0	0								

## MB	0	0	0	0
## MG	0	0	0	0
## ML	0	0	0	0
## NH	0	0	0	0
## NL	0	0	0	0
## NN	0	0	0	0
## PS	0	0	0	0
## RUR	1	0	0	0
## SG	0	1	0	0
## SS	0	0	1	0
## TA	0	0	0	1
## TS	-1	-1	-1	-1

```
eff.model2 <- lm(time ~ chip + student, data = chips_long)
summary(eff.model2)
```

```
##
## Call:
## lm(formula = time ~ chip + student, data = chips_long)
##
## Residuals:
```

	Min	1Q	Median	3Q	Max
	-23.452	-4.202	0.000	4.202	23.452

```
##
## Coefficients:
```

	Estimate	Std. Error	t value	Pr(> t)
(Intercept)	115.43548	1.86584	61.868	< 2e-16 ***
chip1	-7.04839	1.86584	-3.778	0.000701 ***
student1	0.06452	10.21965	0.006	0.995005
student2	68.06452	10.21965	6.660	2.24e-07 ***
student3	-35.93548	10.21965	-3.516	0.001414 **
student4	-50.93548	10.21965	-4.984	2.44e-05 ***
student5	-4.93548	10.21965	-0.483	0.632644
student6	-16.93548	10.21965	-1.657	0.107912
student7	0.56452	10.21965	0.055	0.956315
student8	40.56452	10.21965	3.969	0.000415 ***
student9	-33.43548	10.21965	-3.272	0.002690 **
student10	18.06452	10.21965	1.768	0.087295 .
student11	16.06452	10.21965	1.572	0.126457
student12	-2.43548	10.21965	-0.238	0.813258
student13	-1.93548	10.21965	-0.189	0.851064
student14	-16.93548	10.21965	-1.657	0.107912
student15	-41.93548	10.21965	-4.103	0.000287 ***
student16	-17.93548	10.21965	-1.755	0.089468 .
student17	-17.43548	10.21965	-1.706	0.098328 .
student18	26.06452	10.21965	2.550	0.016103 *
student19	-7.93548	10.21965	-0.776	0.443537
student20	-27.43548	10.21965	-2.685	0.011711 *
student21	-11.93548	10.21965	-1.168	0.252047
student22	68.06452	10.21965	6.660	2.24e-07 ***
student23	-14.43548	10.21965	-1.413	0.168087
student24	5.06452	10.21965	0.496	0.623811
student25	69.56452	10.21965	6.807	1.50e-07 ***
student26	-26.43548	10.21965	-2.587	0.014784 *
student27	70.56452	10.21965	6.905	1.15e-07 ***
student28	-18.93548	10.21965	-1.853	0.073764 .
student29	54.56452	10.21965	5.339	8.94e-06 ***
student30	-28.93548	10.21965	-2.831	0.008198 **

```
## ---
## Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
##
## Residual standard error: 14.69 on 30 degrees of freedom
## Multiple R-squared:  0.9272, Adjusted R-squared:  0.852
## F-statistic: 12.33 on 31 and 30 DF,  p-value: 3.375e-10
```

```
anova(eff.model2)
```

	Df <int>	Sum Sq <dbl>	Mean Sq <dbl>	F value <dbl>	Pr(>F) <dbl>
chip	1	3080.145	3080.1452	14.27016	7.009264e-04
student	30	79411.742	2647.0581	12.26369	4.026186e-10
Residuals	30	6475.355	215.8452	NA	NA
3 rows					

Step 5: Formulate conclusions.

14. After accounting for student to student variation is there a significant difference between the melting time of the two chip types?

Yes, we can see that we account for variation by student, we can see a significant difference between the melting times of the two chips.