Lecture 10: Bootstrap

STAT 324

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Spring 2020

Confidence Interval: The Short Story

To find confidence interval, we use the distribution of $\frac{\bar{X}-E(\bar{X})}{\mathrm{SD}(\bar{X})}$.

If
$$ar{X} \sim N$$
, and $\mathrm{SD}(ar{X})$ is known, then $rac{ar{X} - E(ar{X})}{\mathrm{SD}(ar{X})} \sim N$.

If
$$ar{X}\sim N$$
, and $\mathrm{SD}(ar{X})$ is unknown, then $rac{ar{X}-E(ar{X})}{\widehat{\mathrm{SD}}(ar{X})}\sim t_{n-1}.$

So, when is $ar{X} \sim N$?

- 1. If the data are normal, i.e. $X_1, \ldots, X_n \sim N$.
 - o check using histogram and/or QQ-plot
- 2. If $n \geq 30$, then CLT tells us $ar{X} \sim N$ (in most real life scenarios...)

What if the data are not normal, and n < 30?!?!?!?!



What is the "gold standard" for finding the distribution of anything?

Sample from the population many, many times, and create a histogram.

That's all fun and games in theory, but in practice we cannot really do that.

Remember, all of statistics is build on one fundamental assumption: the sample looks like the population.

So what if we just... resample from the sample...?

This approach is called *bootstraping*. How it works:

- 1. Grab your bootstraps
- 2. Pull yourself up!



This approach is called *bootstraping*. How it works:

- 1. Given a sample, calculate \bar{x} .
- 2. Generate a new sample of size n from the original sample by sampling with replacement (!)
 - \circ we call the first new sample $x_{11}, x_{12}, \ldots, x_{1n}$, the second new sample $x_{11}, x_{12}, \ldots, x_{1n}$, ..., the B'th new sample $x_{B1}, x_{B2}, \ldots, x_{Bn}$
 - these new samples are called *bootstrap samples*
- 3. For each bootstrap sample, calculate $t_j=rac{ar{x}_{\cdot j}-ar{x}}{s_j/\sqrt{n}}$.
 - here, $\bar{x}_{\cdot j}$ is the average of the j'th bootstrap sample, while \bar{x} is the average of the original sample.
- 4. Estimate the distribution of $rac{ar{X}-E(ar{X})}{\widehat{ ext{SD}}(ar{X})}$ by the distribution of t_1,t_2,\ldots,t_B

To find a confidence interval, we need to find x_1, x_2 such that

$$P\left(x_1 \leq rac{ar{X} - E(ar{X})}{\widehat{ ext{SD}}(ar{X})} \leq x_2
ight) = 1 - lpha.$$

We can use the bootstrap samples to estimate the distribution of $\frac{X-E(X)}{\widehat{\mathrm{SD}}(\bar{X})}$, and find the cut-offs such that there's $\alpha/2$ to the left of x_1 and $\alpha/2$ to the right of x_2 .

We will call $x_1=\hat{t}_{\,1-\alpha/2}$, and $x_2=\hat{t}_{\,\alpha/2}$ – the $1-\alpha/2$ and $\alpha/2$ critical values of the distribution of the $\hat{t}_{\,j}$'s.

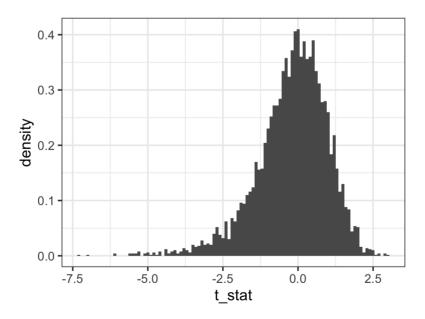
Note: in this case, it is most likely that $\hat{t}_{1-\alpha/2} \neq -\hat{t}_{\alpha/2}$!!! There is no guarentee that the distribution is symmetrical, so no reason to think one of the values will be the negative of the other.

This code takes an origina sample (orig_sample), creates 5000 bootstrap samples, and calculates $\hat{t}_1, \ldots, \hat{t}_{5000}$.

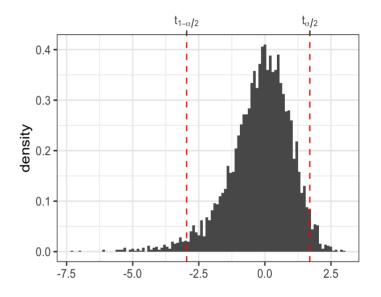
All of this will be in the data set bootstrap_samples.

We can then create a histogram of the \hat{t}_j 's:

```
ggplot(bootstrap_samples,
    aes(x = t_stat)) +
geom_histogram(binwidth = 0.1,
    aes(y = ..density..))
```



 $\hat{t}_{\,1-lpha/2}$ and $\hat{t}_{\,lpha/2}$ are by definition the numbers that cut-off 1-lpha/2 and lpha/2 of the area to the right, respectively.



In this case, the numbers are:

A tibble: 1 x 2

So,

$$\begin{split} 1 - \alpha &= P\left(\hat{t}_{1-\alpha/2} \leq \frac{\bar{X} - E(\bar{X})}{\widehat{\mathrm{SD}}(\bar{X})} \leq \hat{t}_{\alpha/2}\right) \\ &= P\left(\hat{t}_{1-\alpha/2}\widehat{\mathrm{SD}}(\bar{X}) \leq \bar{X} - \mu \leq \hat{t}_{\alpha/2}\widehat{\mathrm{SD}}(\bar{X})\right) \\ &= P\left(-\bar{X} + \hat{t}_{1-\alpha/2}\widehat{\mathrm{SD}}(\bar{X}) \leq -\mu \leq -\bar{X} + \hat{t}_{\alpha/2}\widehat{\mathrm{SD}}(\bar{X})\right) \\ &= P\left(\bar{X} - \hat{t}_{1-\alpha/2}\widehat{\mathrm{SD}}(\bar{X}) \geq \mu \geq \bar{X} - \hat{t}_{\alpha/2}\widehat{\mathrm{SD}}(\bar{X})\right) \\ &= P\left(\bar{X} - \hat{t}_{\alpha/2}\widehat{\mathrm{SD}}(\bar{X}) \leq \mu \leq \bar{X} - \hat{t}_{1-\alpha/2}\widehat{\mathrm{SD}}(\bar{X})\right) \end{split}$$

A $(1-\alpha)\cdot 100\%$ Confidence Interval for the true mean μ is $[\bar{X}-\hat{t}_{\alpha/2}\widehat{\mathrm{SD}}(\bar{X}), \bar{X}-\hat{t}_{1-\alpha/2}\widehat{\mathrm{SD}}(\bar{X})].$

We are $(1-\alpha)\cdot 100\%$ confident that the true value is in this interval.

The ChickWeight data have data regarding the effect of diet on early growth of chicks.

```
ChickWeight
```

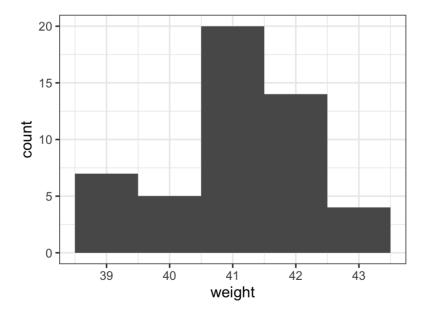
```
## # A tibble: 578 x 4
    weight Time Chick Diet
##
  <dbl> <dbl> <ord> <fct>
##
##
       42
  1
                   1
## 2 51
##
  3 59 4 1
## 4 64 61
## 5 76 8 1
## 6 93 10 1
  7
      106 12 1
##
## 8
      125 14 1
      149 16 1
##
## 10
      171
            18 1
## # ... with 568 more rows
```

We are interested in the mean birth weight of the chicks. This would not be affected by the diet, so treat as one big sample.

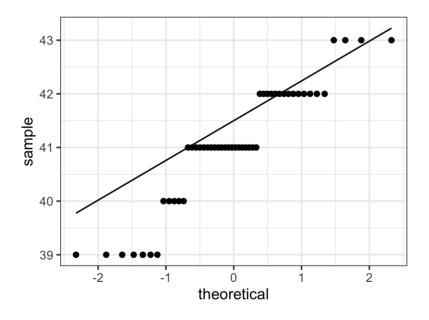
Want to find a confidence interval for μ = true mean birth weight.

```
birth_weights <- ChickWeight %>% filter(Time == 0)

ggplot(birth_weights,
        aes(x = weight)) +
   geom_histogram(binwidth = 1)
```



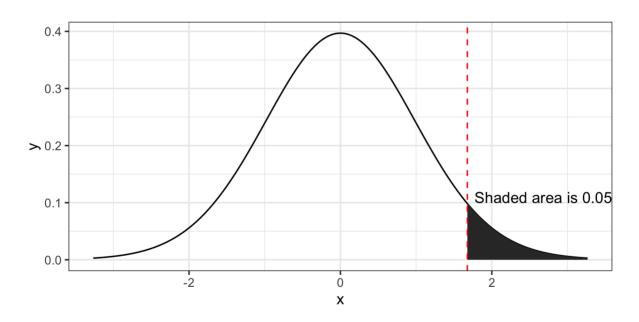
```
ggplot(birth_weights,
    aes(sample = weight)) +
geom_qq() +
geom_qq_line()
```



Definitely not normal.

BUT... n=50! So, by CLT $\bar{X}\sim N$. Therefore, can construct a CI. Since we do not know true σ , find a 90% CI as $\bar{x}\pm t_{n-1,0.05}\frac{s}{\sqrt{n}}$.

What is $t_{n-1,0.05}$? The value on x-axis such that we cut-off 0.05 to the right.



In R: remember that quanile finds the cut-off that cuts off to the left. To cut off 0.05 to the right, we cut off 0.95 to the left:

```
T 49 <- StudentsT(df = 49) # n-1
 (t_crit \leftarrow quantile(T_49, 0.95))
## [1] 1.676551
So, 90\% CI is
birth_weights %>%
   summarize(mean = mean(weight),
             sd = sd(weight),
             LL = mean - t_crit * sd/sqrt(50),
             UL = mean + t_crit * sd/sqrt(50))
## # A tibble: 1 x 4
## mean sd LL
                        UI
## <dbl> <dbl> <dbl> <dbl>
```

1 41.1 1.13 40.8 41.3

Measuring water quality over time. Done by measuring biochemical oxygen demand.

Data:

```
BOD
```

```
## Time demand

## 1 1 8.3

## 2 2 10.3

## 3 3 19.0

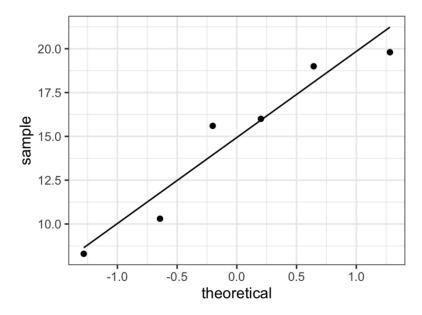
## 4 4 16.0

## 5 5 15.6

## 6 7 19.8
```

n small, so cannot use CLT to conclude that $ar{X} \sim N$. However, if the data is normal, we can still get to that same conclusion!

```
ggplot(BOD, aes(sample = demand)) +
  geom_qq() +
  geom_qq_line()
```



Is it straight enough? Not sure. Compare to other samples of same size that are *actually* from a normal with same mean and SD as our sample. Then ask: does our sample seem that much different?

```
X <- Normal(mu = mean(BOD$demand), sigma = sd(BOD$demand))
normal_samples <- tibble(i = 1:9) %>%
  mutate(data = map(i, ~random(X, n = nrow(BOD)))) %>%
  unnest_longer(data)
```

I would probably say no.

So, we assume $X_1,\dots,X_6\sim N.$ We do not know σ , so find 99% CI as $ar x\pm t_{n-1,0.005}rac{s}{\sqrt n}.$

```
T_5 <- StudentsT(df = 5) # n-1
(t_crit <- quantile(T_5, 0.995))</pre>
```

[1] 4.032143

So, 99% CI is

```
BOD %>%
summarize(mean = mean(demand),
sd = sd(demand),
LL = mean - t_crit * sd/sqrt(50),
UL = mean + t_crit * sd/sqrt(50))
```

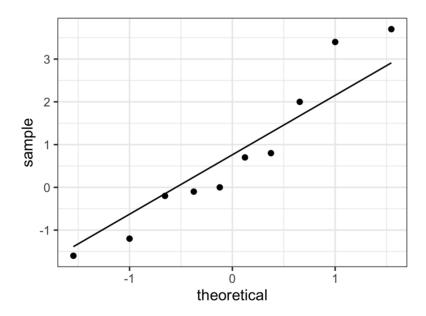
Scientists are interested in the effect of soporific drugs on amount of sleep. Data actually has data for 10 patients in 2 groups, but we will only consider one of the groups.

```
sleep1 <- sleep %>% filter(group == 1)
sleep1
```

```
## 1 0.7 1 1
## 2 -1.6 1 2
## 3 -0.2 1 3
## 4 -1.2 1 4
## 5 -0.1 1 5
## 6 3.4 1 6
## 7 3.7 1 7
## 8 0.8 1 8
## 9 0.0 1 9
## 10 2.0 1 10
```

Small *n*, so no CLT. Is it normal?

```
ggplot(sleep1,
    aes(sample = extra)) +
  geom_qq() +
  geom_qq_line()
```



Is it straight enough? Not sure. Compare to other samples of same size that are *actually* from a normal with same mean and SD as our sample. Then ask: does our sample seem that much different?

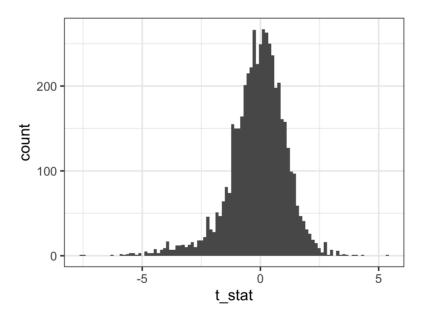
```
X <- Normal(mu = mean(sleep1$extra), sigma = sd(sleep1$extra))
normal_samples <- tibble(i = 1:9) %>%
  mutate(data = map(i, ~random(X, n = nrow(sleep1)))) %>%
  unnest_longer(data)
```

Not sure. After consulting with the scientists in charge of the study, it is decided that we do **NOT** want to assume normality.

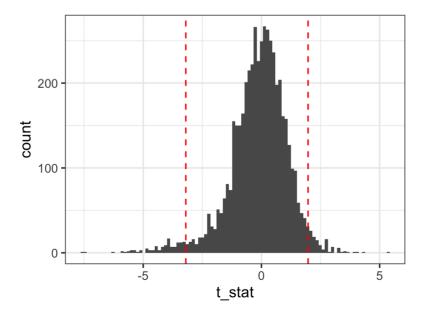
So, non-normal data, and n too small for CLT. So, we opt for a bootstrap approach:

Distribution from bootstrap:

```
ggplot(data = bootstrap_samples,
    aes(x = t_stat)) +
    geom_histogram(bins = 100)
```



Want to create a 95% CI for the true mean. Find $\hat{t}_{0.025}$ and $\hat{t}_{0.975}$:



We find these, estimated mean ($ar{x}$), and standard deviation (s) in R

```
bootstrap_samples %>%
  summarize(t_left = quantile(t_stat, 0.025),
            t_right = quantile(t_stat, 0.975))
## # A tibble: 1 x 2
## t_left t_right
## <dbl> <dbl>
## 1 -3.20 1.97
sleep1 %>%
  summarize(mean = mean(extra),
            sd = sd(extra),
            n = n()
##
              sd n
    mean
## 1 0.75 1.78901 10
```

So, we find the lower limit of 95% CI as

$$ar{x} - \hat{t}_{0.975} \frac{s}{\sqrt{n}} = 0.75 - 1.973 \frac{1.789}{\sqrt{10}}$$

$$= -0.37$$

and the upper limit as

So, we find the lower limit of 95% CI as

$$egin{aligned} ar{x} - \hat{t}_{\,0.025} rac{s}{\sqrt{n}} &= 0.75 - (-3.198) rac{1.789}{\sqrt{10}} \ &= 2.56 \end{aligned}$$