## 11\_30\_22\_lab

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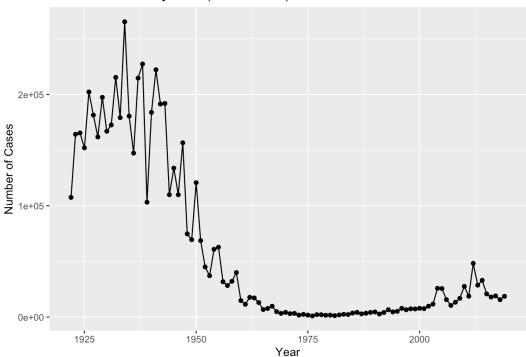
Q1. With the help of the R "addin" package datapasta assign the CDC pertussis case number data to a data frame called cdc and use ggplot to make a plot of cases numbers over time.

```
pertussis_cases_by_year <- data.frame(</pre>
                             Year = c(1922L,
                                        1923L, 1924L, 1925L, 1926L, 1927
                                        1929L, 1930L, 1931L, 1932L, 1933
                                        1936L, 1937L, 1938L, 1939L, 1940
                                        1942L, 1943L, 1944L, 1945L, 1946
                                        1949L, 1950L, 1951L, 1952L, 1953
                                        1955L, 1956L, 1957L, 1958L, 1959
                                        1961L, 1962L, 1963L, 1964L, 1965
                                        1968L, 1969L, 1970L, 1971L, 1972
                                        1974L, 1975L, 1976L, 1977L, 1978
                                        1981L, 1982L, 1983L, 1984L, 1985
                                        1987L, 1988L, 1989L, 1990L, 1991
                                        1994L, 1995L, 1996L, 1997L, 1998
                                        2000L, 2001L, 2002L, 2003L, 2004
                                        2006L, 2007L, 2008L, 2009L, 2016
                                        2013L, 2014L, 2015L, 2016L, 2017
                                        2019L),
  No. Reported Pertussis Cases = c(107473)
                                        164191, 165418, 152003, 202210,
                                        161799, 197371, 166914, 172559,
                                        265269, 180518, 147237, 214652,
                                        183866, 222202, 191383, 191890,
                                        133792, 109860, 156517, 74715, 6
                                        68687,45030,37129,60886,6278
                                        32148,40005,14809,11468,1774
                                        13005,6799,7717,9718,4810,32
                                        3036,3287,1759,2402,1738,101
                                        1623, 1730, 1248, 1895, 2463, 227
                                        4195, 2823, 3450, 4157, 4570, 271
                                        4617,5137,7796,6564,7405,729
                                        7580,9771,11647,25827,25616,
                                        13278, 16858, 27550, 18719, 4827
                                        32971, 20762, 17972, 18975, 156(
```

)

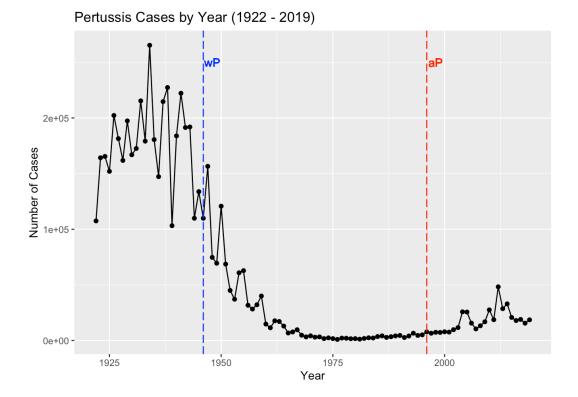
```
library(ggplot2)
ggplot(pertussis_cases_by_year) +
  aes(Year, No..Reported.Pertussis.Cases) +
  geom_point() +
  geom_line() +
  labs(x = "Year", y="Number of Cases", title = "Pertussis Cases")
```

## Pertussis Cases by Year (1922 - 2019)



Q2. Using the ggplot geom\_vline() function add lines to your previous plot for the 1946 introduction of the wP vaccine and the 1996 switch to aP vaccine (see example in the hint below). What do you notice?

```
ggplot(pertussis_cases_by_year) +
  aes(Year, No..Reported.Pertussis.Cases) +
  geom_point() +
  geom_line() +
  labs(x = "Year", y="Number of Cases", title = "Pertussis Case
  geom_vline(xintercept = c(1946, 1996), colour=c("blue", "rec
  geom_text(aes(x = 1946+2, y = 250000, label = "wP"), colour =
  geom_text(aes(x = 1996+2, y = 250000, label = "aP"), colour =
```



Q3. Describe what happened after the introduction of the aP vaccine? Do you have a possible explanation for the observed trend?

After the introduction of the aP vaccine there was a slight uptick in cases of Pertussis. It did take a while though, this gives credence to maybe it was due to other factors such as testing becoming more sensitive and reporting more cases. Another possibility is the vaccine is slightly weaker but gives fewer side effects in a cost benefit analysis concluding that it is better. Another explanation is if the vaccine isn't updating often enough, the bacterial evolution could outpace vaccine development, creating new strains that could get around vaccination. A final explanation is that with time, people begin fearing the whooping cough less as they have never seen anyone get it. Because of this, they don't vaccinate their children for it as it doesn't seem like a problem. This could lead to a rise in cases. Between the fall in cases to almost 0 to the uptick, there are about 30 years, enough for one generation to grow up without fear or respect of the disease.

```
subject_id infancy_vac biological_sex
                                                       ethnicity
race
1
           1
                       wP
                                  Female Not Hispanic or Latino
White
           2
                                  Female Not Hispanic or Latino
2
                       wP
White
           3
                                  Female
                                                         Unknown
                       wP
White
  year_of_birth date_of_boost
                                    dataset
                    2016-09-12 2020_dataset
1
     1986-01-01
2
     1968-01-01
                    2019-01-28 2020 dataset
3
     1983-01-01
                    2016-10-10 2020_dataset
```

Q4. How many aP and wP infancy vaccinated subjects are in the dataset?

```
table(subject$infancy_vac)
```

aP wP 47 49

47 aP infancy vaccinated subjects and 49 wP infancy vaccinated subjects

Q5. How many Male and Female subjects/patients are in the dataset?

```
table(subject$biological_sex)
```

```
Female Male 66 30
```

66 females and 30 males are in the dataset

Q6. What is the breakdown of race and biological sex (e.g. number of Asian females, White males etc...)?

```
table(subject$race, subject$biological_sex)
```

			Female	Male
American	Indian/Alaska	Native	0	1
Asian			18	9
Black or	African Americ	can	2	0

```
More Than One Race 8 2
Native Hawaiian or Other Pacific Islander 1 1
Unknown or Not Reported 10 4
White 27 13
```

Here is the breakdown

```
library(lubridate)
```

Loading required package: timechange

Attaching package: 'lubridate'

The following objects are masked from 'package:base':

date, intersect, setdiff, union

- Q7. Using this approach determine (i) the average age of wP individuals,
- (ii) the average age of aP individuals; and (iii) are they significantly different?

```
subject$age <- today() - ymd(subject$year_of_birth)</pre>
```

```
library(dplyr)
```

Attaching package: 'dplyr'

The following objects are masked from 'package:stats':

filter, lag

The following objects are masked from 'package:base':

intersect, setdiff, setequal, union

```
aP <- subject %>% filter(infancy_vac == "aP")
round(summary(time_length(aP$age, "years")))
```

```
Min. 1st Qu. Median Mean 3rd Qu. Max. 23 25 26 25 26 27
```

```
wP <- subject %>% filter(infancy_vac == "wP")
round(summary(time_length(wP$age, "years")))
```

```
Min. 1st Qu. Median Mean 3rd Qu. Max. 28 32 35 36 40 55
```

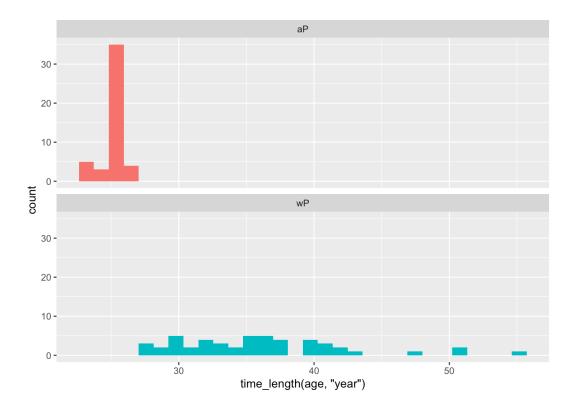
Q8. Determine the age of all individuals at time of boost?

```
time_diff_recieved_boost <- ymd(subject$date_of_boost) - ymd(st
age_at_boost <- time_length(time_diff_recieved_boost, "year")
head(age_at_boost)</pre>
```

- [1] 30.69678 51.07461 33.77413 28.65982 25.65914 28.77481
- Q9. With the help of a faceted boxplot or histogram (see below), do you think these two groups are significantly different?

```
ggplot(subject) +
  aes(time_length(age, "year"),
  fill=as.factor(infancy_vac)) +
  geom_histogram(show.legend=FALSE) +
  facet_wrap(vars(infancy_vac), nrow=2)
```

<sup>`</sup>stat\_bin()` using `bins = 30`. Pick better value with
`binwidth`.



Yes they look significantly different. In the past people seem to get the vaccine whenever they could, but not at a young age. This leads to a distribution ranging from 25ish to almost 60. With the new vaccine everyone gets it before 30.

```
# Complete the API URLs...
specimen <- read_json("https://www.cmi-pb.org/api/specimen", si
titer <- read_json("https://www.cmi-pb.org/api/ab_titer", simpl</pre>
```

Q9. Complete the code to join specimen and subject tables to make a new merged data frame containing all specimen records along with their associated subject details:

```
meta <- full_join(specimen, subject)

Joining, by = "subject_id"

dim(meta)

[1] 729 14</pre>
```

head(meta)

specimen_id s	subject_id	actual	_day_relative_t	o_boost						
1 1	1			-3						
2 2	1			736						
3 3	1			1						
4 4	1			3						
5 5	1			7						
6 6	1			11						
<pre>planned_day_relative_to_boost specimen_type visit</pre>										
infancy_vac biological_sex										
1	.1.	0	Blood	1						
wP Fema	ale	726	D1	10						
2	-1-	736	Blood	10						
wP Fema	ate	1	Blood	2						
wP Fema	ale	1	D (000	2						
4		3	Blood	3						
wP Fema	ale									
5	1	7	Blood	4						
wP Fema	ale	14	Blood	5						
wP Fema	ale	14	Dicod	3						
Wi i cinc		race	year_of_birth d	late_of_boost						
dataset	-									
1 Not Hispanic	or Latino	White	1986-01-01	2016-09-12						
2020_dataset										
2 Not Hispanic	or Latino	White	1986-01-01	2016-09-12						
2020_dataset										
3 Not Hispanic	or Latino	White	1986-01-01	2016-09-12						
2020_dataset 4 Not Hispanic	or Latino	White	1986-01-01	2016-09-12						
2020 dataset	or Lacino	WIIICC	1300 01 01	2010 05 12						
5 Not Hispanic	or Latino	White	1986-01-01	2016-09-12						
2020_dataset										
6 Not Hispanic	or Latino	White	1986-01-01	2016-09-12						
2020_dataset										
age										
1 13483 days										
2 13483 days										
3 13483 days										
4 13483 days										
5 13483 days										
6 13483 days										

Q10. Now using the same procedure join meta with titer data so we can

further analyze this data in terms of time of visit aP/wP, male/female etc.

```
abdata <- inner_join(titer, meta)</pre>
```

Joining, by = "specimen\_id"

```
dim(abdata)
```

[1] 32675 21

Q11. How many specimens (i.e. entries in abdata) do we have for each isotype?

```
table(abdata$isotype)
```

```
IgE IgG IgG1 IgG2 IgG3 IgG4
6698 1413 6141 6141 6141 6141
```

Q12. What do you notice about the number of visit 8 specimens compared to other visits?

```
table(abdata$visit)
```

```
1 2 3 4 5 6 7 8
5795 4640 4640 4640 4640 4320 3920 80
```

There are very few visit 8 specimens compared to all the other visits.

```
ig1 <- abdata %>% filter(isotype == "IgG1", visit!=8)
head(ig1)
```

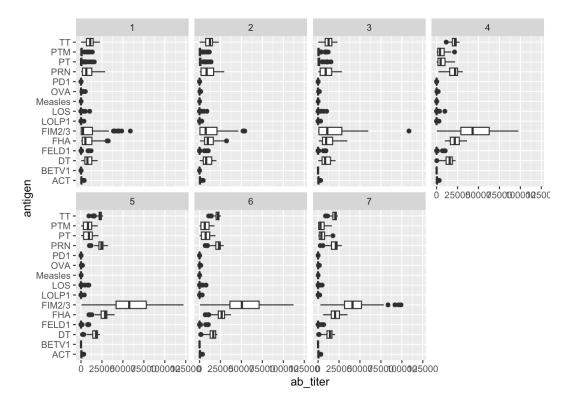
```
specimen_id isotype is_antigen_specific antigen
                                                           MFI
MFI_normalised
                                      TRUE
            1
                 IgG1
                                                ACT 274.355068
0.6928058
                 IgG1
                                      TRUE
                                                L0S
                                                     10.974026
            1
2.1645083
            1
                 IgG1
                                      TRUE
                                              FELD1
                                                      1.448796
0.8080941
            1
                 IgG1
                                      TRUE
                                              BETV1
                                                      0.100000
```

1.0000000							
5	1	IgG1		TRUE I	L0LP1	0.100000	
1.0000000 6	1	IgG1		TDIIF Mo	aclac	36.277417	
1.6638332	1	1961		TRUE ME	as (es	30.2//41/	
<pre>unit lower_limit_of_detection subject_id</pre>							
actual_day_	_relat	ive_to_	boost				
1 IU/ML -3			3.848750	1			
2 IU/ML -3			4.357917	1			
3 IU/ML			2.699944	1			
-3 4 IU/ML			1.734784	1			
-3				_			
5 IU/ML -3			2.550606	1			
6 IU/ML			4.438966	1			
-3	4a., r	alativa :	to boost (	specimen type	o vici	L	
infancy_vac				specimen_type	S ATZT	L	
1	010	rogica c_	0	Blood	d :	1	
wP	Fema 7	Le	-				
2			0	Blood	d :	1	
wP	Fema 7	Le					
3			0	Blood	d :	1	
wP	Femal	Le					
4			0	Blood	d :	1	
wP	Fema 7	Le					
5			0	Blood	d :	1	
wP	Fema 7	Le	•				
6	F 1		0	Blood	d :	1	
wP	Fema 7			aar of birth	data d	of boost	
dataset	E	ernnicir	y race ye	ear_of_birth	date_d	01_00051	
1 Not Hispa	nic c	vr Latin	o White	1986-01-01	201	16-09-12	
2020_datase		) Latin	O MILLICE	1900-01-01	20.	10-09-12	
2 Not Hispa		or Latin	o White	1986-01-01	201	16-09-12	
2020_datase		) Lacin	o wiiicc	1300 01 01	20.	10 05 12	
3 Not Hispa		r Latin	o White	1986-01-01	203	16-09-12	
2020_datase	et						
4 Not Hispa		r Latin	o White	1986-01-01	203	16-09-12	
2020_datase				1000 01 01		16 06 15	
5 Not Hispa 2020_datase		or Latin	o White	1986-01-01	201	16-09-12	
ZwZw_uataSt	- L						

```
6 Not Hispanic or Latino White 1986-01-01 2016-09-12 2020_dataset age 1 13483 days 2 13483 days 3 13483 days 4 13483 days 5 13483 days 6 13483 days
```

Q13. Complete the following code to make a summary boxplot of Ab titer levels for all antigens:

```
ggplot(ig1) +
  aes(MFI, antigen) +
  geom_boxplot() +
  facet_wrap(vars(visit), nrow=2)+
  labs(x = "ab_titer")
```



Q14. What antigens show differences in the level of IgG1 antibody titers recognizing them over time? Why these and not others?

FIM2/3 seems the most unique in terms of the level of IgG1 antibody titers. I didn't find anything on the website so I'm not entirely sure what that

means. I am somewhat confused.