# Advanced Tabular Data Manipulation

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- compute cumulative, offset, and sliding-window transformations
- simultaneously transform or summarize multiple columns
- transform between long and wide data formats
- combine multiple data frames using joins on one or more columns

## Window transformations

#### Offsets

• lead() and lag() return either forward- or backward-shifted versions of their input vectors

```
lead(c(1,2,3))
[1] 2 3 NA
lag(c(1,2,3))
[1] NA 1 2
```

• This is most useful to compute offsets, which we often do when looking at time series.

#### Offsets Example

• Let's look at some time series data about when GTEx samples were collected, and calculate the change over time

```
gtex_samples_time_link = "https://raw.githubusercontent.com/alejandroschuler/r4ds-
courses/af6056f9a8d999a80fd787f89aa3483157d43681/data/gtex_metadata/gtex_samples_time.csv"

gtex_samples_by_month = read_csv(file = gtex_samples_time_link, col_types = cols())

gtex_samples_by_month %>%
  head(2L)

# A tibble: 2 x 3
  month year num_samples
  <dbl> <dbl> <dbl> <dbl> 1 5 2011 20
  2 6 2011 44
```

```
gtex_samples_by_month %>%
  mutate(increase_in_samples = total_num_samples - lag(total_num_samples)) %>%
  head(3L)
Error: Problem with `mutate()` input `increase_in_samples`.
x object 'total_num_samples' not found
i Input `increase_in_samples` is `total_num_samples - lag(total_num_samples)`.
```

#### Exercise: multiple offsets

```
gtex_samples_by_month %>%
 filter (month %in% c(3, 6, 9, 12)) %>%
 head(n = 6L)
# A tibble: 6 x 3
 month year num_samples
 <dbl> <dbl>
                <db1>
     6 2011
                   44
   9 2011
           100
   12 2011 111
    3 2012
               141
   6 2012
                63
    9 2012
                  115
```

Let's say you want to compute the change in number of samples within months from one year to the next. We've shortened the data to only include four months each year (March, June, September, and December) to simplify this. (i.e. comparing March 2015 to March 2016 and June 2015 to June 2016)

- 1. Figure out a way to do this using <code>lead()</code> or <code>lag()</code> in a single <code>mutate()</code> statement (hint: check the documentation).
- 2. Figure out a different way to do this with group\_by () instead. Which seems more natural or robust to you? Why?
- 3. In both solutions you end up with some NAs since the March 2011 counts are unknown. If we wanted to assume that the March 2011 would be the same as the June 2011, how might we modify our code to reflect that in order to make sure we don't get NAs in the result?

#### Rolling functions

 slide\_vec applies a function using a sliding window across a vector (sometimes called a "rolling" function)

```
library("slider")
numbers = c(9, 6, 8, 4, 7, 3, 8, 4, 2,
1, 3, 2)
slide_vec(numbers, sum, .after = 3,
.step = 2L)
[1] 27 NA 22 NA 22 NA 15 NA 8 NA 5
NA
```

- the .after argument specifies how many elements after the "index" element are included in the rolling window
- .step specifies how to move from one index element to the next
- .before is the backward-looking equivalent of .after

```
gtex_samples_by_month %>%
   mutate(avg_samples_2_month =
slide_vec(total_num_samples, mean,
.before = 1L)) %>%
   select(-year) %>%
   head(2L)
Error: Problem with `mutate()` input
`avg_samples_2_month`.
x object 'total_num_samples' not found
i Input `avg_samples_2_month` is
`slide_vec(total_num_samples, mean,
.before = 1L)`.
```

#### Cumulative functions

- A cumulative function is like a rolling window function except that the window expands with each iteration instead of shifting over
- For example, cumsum takes the cumulative sum of a vector. See ?cumsum for similar functions

```
cumsum(c(1, 2, 3))
[1] 1 3 6

gtex_samples_by_month %>%
  mutate(num_samples_to_date = cumsum(total_num_samples))
Error: Problem with `mutate()` input `num_samples_to_date`.
  x object 'total_num_samples' not found
i Input `num_samples_to_date` is `cumsum(total_num_samples)`.
```

#### Turning any function into a cumulative function

- you can use slider::slide\_vec() to turn any function that accepts a vector and returns a number into a cumulative function
- Use .before=Inf to achieve this

```
library(slider) # imports slide_vec() function

gtex_samples_by_month %>%
  mutate(samples_to_date = slide_vec(total_num_samples, sum, .before = Inf))
Error: Problem with `mutate()` input `samples_to_date`.
x object 'total_num_samples' not found
i Input `samples_to_date` is `slide_vec(total_num_samples, sum, .before = Inf)`.
```

• it is usually better (computationally faster) to use a built-in cumulative function (e.g. cumsum ()), but if none exists this is a great solution

#### Turning any function into a cumulative function

• If the function you want to transform takes additional arguments, you can give those to slide\_vec and it will pass them through for you

```
gtex_samples_by_month %>%
   mutate(
     avg_samples_by_month = slide_vec(total_num_samples, mean, .before = Inf, na.rm
= TRUE)
   )
Error: Problem with `mutate()` input `avg_samples_by_month`.
x object 'total_num_samples' not found
i Input `avg_samples_by_month` is `slide_vec(total_num_samples, mean, .before = Inf, na.rm = TRUE)`.
```

#### Exercise: total number of samples in the last twelve months

```
library(lubridate) # this helps us with dates
```

```
gtex_samples_by_month
# A tibble: 66 x 3
  month year num_samples
  <dbl> <dbl>
                  <dbl>
      5 2011
                     20
    6 2011
                   44
   7 2011
   8 2011
                   132
   9 2011
                   100
  10 2011
                   110
    11 2011
                   203
   12 2011
                  111
   1 2012
                    208
10
      2 2012
                    95
# ... with 56 more rows
```

Starting with this gtex\_samples\_by\_month dataframe you, add a column that has the total number of samples in the last twelve months relative to the row we are on.

# Column-wise operations

#### Repeating operations on columns

```
df = tibble(
    a = rnorm(10),
    b = rnorm(10),
    c = rnorm(10),
    g = rbinom(10, 1, 0.5)
)
```

- Let's say we have these data and we want to take the mean of each column a, b, and c within the groups g.
- One way to do it is with a normal summarize:

```
df %>%
  group_by(g) %>%
  summarize(
    mean_a = mean(a),
    mean_b = mean(b),
    mean_b = mean(c)
)

# A tibble: 2 x 3
    g mean_a mean_b
  <int> <dbl> <dbl>
1 0 -0.249 -0.0148
2 1 1.06 -0.569
```

- Copy-pasting code like this frequently creates errors and bugs that are hard to see
- It's even worse if you want to do multiple summaries

```
df %>%
 group_by(g) %>%
 summarize(
   mean a = mean(a),
   mean b = mean(b),
   mean b = mean(c),
   median a = median(a),
   median b = median(b),
   median_c = median(c)
# A tibble: 2 x 6
     g mean_a mean_b median_a
median_b median c
 <int> <dbl> <dbl> <dbl>
<dbl> <dbl>
1 0 -0.249 -0.0148
                        -0.686
-0.218 \quad -0.0994
2 1 1.06 -0.569
                         1.15
0.570 - 0.824
```

#### Columnwise operations

• The solution is to use across () in your summarize:

- The first argument to across () is a selection of columns. You can use anything that would work in a select () here
- We've explicitly included the argument names for .cols and .fns here, but in R code out-in-the-wild they're usually omitted.

 The second argument is the function you'd like to apply to each column. You can provide multiple functions by wrapping them in a "list()". Lists are like vectors but their elements can be of different types and each element has a name (more on that later)

```
fns = list(
  avg = mean,
  max = max
)

df %>%
  group_by(g) %>%
  summarize(across(c(a,b), fns))
# A tibble: 2 x 5
        g a_avg a_max b_avg b_max
  <int> <dbl> <dbl> <dbl> <dbl> <dbl> <dbl> 1
        0 -0.249 2.05 -0.236 1.79
        2 1 1.06 1.93 0.788 2.61
```

• see ?across() to find out how to control how these columns get named in the output

#### Columnwise operations with where()

```
df = tibble(
    a = rnorm(10),
    b = rnorm(10),
    c = as.character(rnorm(10)),
    g = rbinom(10, 1, 0.5)
)
```

- Sometimes its nice to apply a transformation to all columns of a given type or all columns that match some condition
- where () is a handy function for that

#### Columnwise mutate

```
df = tibble(
    a = rnorm(10),
    b = rnorm(10),
    c = as.character(rnorm(10)),
    g = rbinom(10, 1, 0.5)
)
```

• across() works with any dplyr "verb", including mutate():

#### Columnwise mutate

• Most often you will need to write your own mini function to do what you want. To do that you put  $\sim$  before your expression and use . where you would put the name of the column

```
df %>%
 mutate(across(
                  # columns to mutate
   a:b,
   ~ . - lag(.), # function to mutate them with
   .names = '{col}_offset' # how to name the outputs
# A tibble: 10 x 6
            bс
                                 q a_offset b_offset
   <dbl> <dbl> <chr>
                   <int>
                                     <dbl>
                                            <dbl>
1 -2.22 1.43 1.48582091419658
                                 0 NA
                                          NA
2 1.52 0.757 1.5111112894678 0 3.74 -0.677
3 0.935 0.972 0.605762865798173
                                 1 -0.586 0.215
  0.710 1.34 0.204771748793352
                                1 -0.225 0.369
                                1 -2.17 -1.01
5 -1.46 0.327 -1.74629427588112
  0.385 0.764 0.634975230582194
                                0 1.85 0.437
7 -0.444 -1.38 2.46701997718975 1 -0.829 -2.14
  1.82 0.609 0.0912452808035654
                                 1 2.27 1.98
                             1 -1.59 -1.28
  0.228 -0.675 -0.214257824130902
10 -0.139 0.978 0.382695567242115
                                1 -0.367
                                          1.65
```

• Note that I've also used the . names argument to control how the output columns get named

### Exercise: Filtering out or replacing NAs

Let's go back to the GTEx expression data we've been looking at:

- 1. Replacing NAs with some other value is a very common operation so it gets its own function:  $replace_na()$ . Use this function to replace all NAs present in any numeric column with 0s
- 2. Instead of replacing these values, we may want to filter them all out instead. Starting with the original data, use filter() and across() to remove all rows from the data that have any NAs in any column. Recall that is.na() checks which elements in a vector are NA.

Tidy data: rearranging a data frame

#### Messy data

• Sometimes data are organized in a way that makes it difficult to compute in a vector-oriented way. For example, look at this dataset:

```
gtex time link =
 "https://raw.githubusercontent.com/alejandroschuler/r4ds-courses/advance-
2021/data/gtex_metadata/gtex_time_tissue.csv"
gtex time tissue data = read csv(file = gtex time link, col types = cols())
head(gtex_time_tissue_data, 3L)
# A tibble: 3 x 8
               `2011` `2012` `2013` `2014` `2015` `2016` `2017`
 tissue
 <chr> <dbl> <dbl> <dbl>
                                   <dbl> <dbl> <dbl> <dbl> <
                               243
1 Adipose Tissue
                   56 107
                                     206
                                                  134
2 Adrenal Gland
                   28 41 84
                                   65
                                            20 31
                                                          0
                      18
3 Bladder
                              0
                                            0
                                                          0
```

- the values in the table represent how many samples of that tissue were collected during that year.
- How could I use ggplot to make this plot? It's hard!

#### Messy data

```
head(gtex_time_tissue_data, 3L)
# A tibble: 3 x 8
 tissue
                `2011` `2012` `2013` `2014` `2015` `2016` `2017`
 <chr>
                 <dbl> <dbl> <dbl>
                                     <dbl> <dbl> <dbl>
                                                         <dbl>
1 Adipose Tissue
                    56
                         107
                                243
                                       206
                                                     134
2 Adrenal Gland
                    28
                                 84
                                        65
                                                      31
                           41
                                                              0
3 Bladder
                           18
                                   0
                                                              0
```

- One of the problems with the way these data are formatted is that the year collected, which is a property of the samples, is stuck into the names of the columns.
- Because of this, it's also not obvious what the numbers in the table mean (although we know they are counts)

#### Tidy data

• Here's a better way to organize the data:

```
# A tibble: 6 x 3
 tissue
                year count
  <chr>
                 <chr> <dbl>
1 Adipose Tissue 2011
                          56
2 Adipose Tissue 2012
                         107
3 Adipose Tissue 2013
                         243
4 Adipose Tissue 2014
                         206
5 Adipose Tissue 2015
                        84
6 Adipose Tissue 2016
                         134
```

This data is tidy. Tidy data follows three precepts:

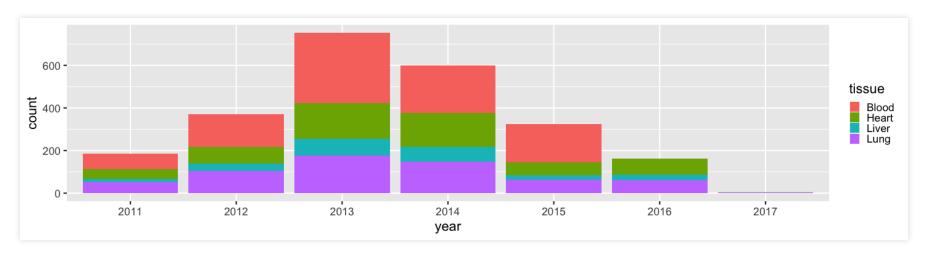
- 1. each "variable" has its own dedicated column
- 2. each "observation" has its own row
- 3. each type of observational unit has its own data frame

In our example, each of the **observations** are different **groups of samples**, each of which has an associated *tissue*, *year*, and *count*. These are the *variables* that are associated with the groups of samples.

## Tidy data

Tidy data is easy to work with.

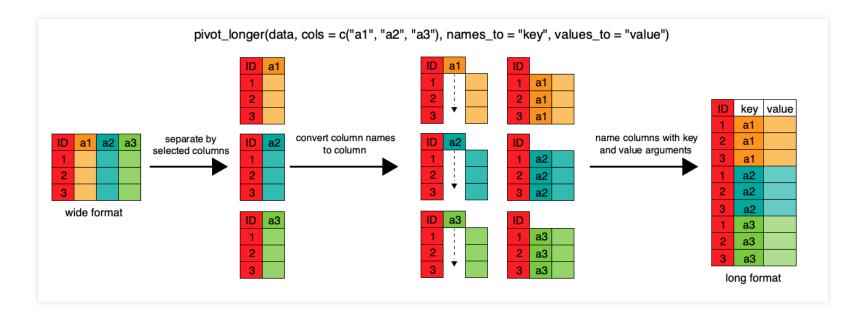
```
tidy %>%
  filter(tissue %in% c("Blood", "Heart", "Liver", "Lung")) %>%
  ggplot() +
  geom_bar(aes(x = year, y = count, fill = tissue), stat = 'identity')
```



### Tidying data with pivot\_longer()

• tidyr::pivot\_longer() is the function you will most often want to use to tidy your data

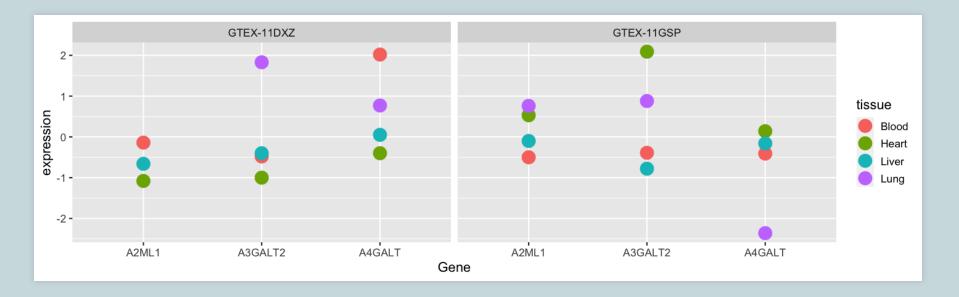
• the three important arguments are: a) a selection of columns, b) the name of the new key column, and c) the name of the new value column



#### Exercise: cleaning GTEX

```
head(gtex_data, 3L)
# A tibble: 3 x 7
Gene Ind Blood Heart Lung Liver NTissues
<chr> <chr> <chr> <chr> <dbl> 3
A2ML1 GTEX-11DXZ -0.14 -1.08 NA -0.66 3
A2 A2ML1 GTEX-11GSP -0.5 0.53 0.76 -0.1 4
A2ML1 GTEX-11NUK -0.08 -0.4 -0.26 -0.13 4
```

#### Use the GTEX data to reproduce the following plot:



The individuals and genes of interest are c ('GTEX-11GSP', 'GTEX-11DXZ') and c ('A2ML1', 'A3GALT2', 'A4GALT'), respectively.

#### "Messy" data is relative and not always bad

#### Pivoting wider

- As we saw with the mouse example, sometimes our data is actually easier to work with in the "wide" format.
- wide data is also often nice to make tables for presentations, or is (unfortunately) sometimes required as input for other software packages
- To go from long to wide, we use pivot\_wider():

```
long_mice
# A tibble: 8 x 3
 mouse time
            weight
 <dbl> <chr>
             <dbl>
     1 before 11.2
     1 after
             13.8
     2 before 10.1
     2 after 12.9
    3 before 10.4
     3 after
               9.72
     4 before 7.96
               8.39
     4 after
```

#### Names prefix

• you can use names\_prefix to make variables names that are more clear in the result

• this can also be used to *remove* a prefix when going from wide to long:

```
wide_mice %>%
  pivot_longer(
    -mouse
    names_to = "time",
    values_to = "weight",
    names_prefix = "weight_"
)
```

### Exercise: creating a table

Use the GTEX data to make the following table:

The numbers in the table are the number of tissues in each individual for which the gene in question was missing.

### Multi-pivoting

Have a look at the following data. How do you think we might want to make it look?

```
gtex time chunk link =
 "https://raw.githubusercontent.com/alejandroschuler/r4ds-courses/advance-
2021/data/gtex metadata/gtex samples tiss time chunk.csv"
gtex_samples_time_chunk =
 read_csv(file = gtex_time_chunk_link, col_types = cols())
head(gtex_samples_time_chunk)
# A tibble: 6 x 9
 tissue `Sept-2015` `Sept-2016` `Oct-2015` `Oct-2016` `Nov-2015` `Nov-2016`
 <chr>
                 <dbl> <dbl>
                                     <dbl>
                                               <dbl>
                                                         <dbl>
                                                                   <dbl>
1 Adipose T...
                              36
                                                            15
                                                                      16
2 Adrenal G...
3 Blood
                                                            33
4 Blood Ves... 9
                              24
                                                  26
                                                                      17
5 Brain
                                                            17
                                                                      24
6 Breast 9
                              19
                                                  20
# ... with 2 more variables: Dec-2016 <dbl>, Dec-2015 <dbl>
```

The problem here is that the column names contain two pieces of data:

- 1. the year
- 2. the month it came from

Our use of pivot\_longer has so far been to extract a single piece of information from the column name

### Multi-pivoting

• Turns out this problem can be tackled too:

```
gtex_samples_time_chunk %>%
 pivot_longer(
   cols=contains("-201"), # selects columns that contain this
   names_pattern = "(\D+)-(\d+)", # a "regular expression" - we'll learn about
these later
   names_to = c(".value", "year")
# A tibble: 54 x 6
  tissue year Sept Oct
                                    Nov
                                          Dec
  <chr> <chr> <dbl> <dbl> <dbl> <dbl> <dbl>
1 Adipose Tissue 2015 5 4 15 0
2 Adipose Tissue 2016 36 20 16 15
3 Adrenal Gland 2015 4 1 2
4 Adrenal Gland 2016 5 5 6
                     2 6 33 17
0 0 0 0
9 7 9 0
24 26 17 12
5 Blood 2015
6 Blood 2016
7 Blood Vessel 2015
8 Blood Vessel 2016
9 Brain 2015
                      12 3 17
10 Brain
                                     2.4
                 2016
# ... with 44 more rows
```

- We won't dig into this, but you should know that almost any kind of data-tidying problem can be solved with some combination of the functions in the tidyr package.
- See the online docs and vignettes for more info

## Combining multiple tables with joins

#### Relational data

- Relational data are interconnected data that is spread across multiple tables, each of which usually has a different unit of observation
- When we get an expression dataset, the data is usually divided into an expression matrix with the expression values of each sample, and table(s) with metadata about the samples themselves.
- For the GTEx dataset, we have information about the samples, subjects, and experiment batches in additional data frames in addition to the expression matrix we've been working with.

```
gtex_metadata_link =
  "https://raw.githubusercontent.com/alejandroschuler/r4ds-courses/advance-
2021/data/gtex_metadata/gtex_sample_metadata.csv"
gtex sample data = read csv(file = gtex metadata link, col types = cols())
head(gtex_sample_data, 2L)
# A tibble: 2 x 6
 subject_id sample_id batch_id center_id tissue rin_score
 <chr>
       <chr>
                          <chr>
                                   <chr>
                                             <chr>
                                                        <dbl>
1 GTEX-11DXZ 0003-SM-5807X BP-39216 B1
                                             Blood
                                                        NA
2 GTEX-11DXZ 0126-SM-5EGGY BP-44460 B1
                                                         7.9
                                             Liver
```

• The sample data has information about the tissue and the subject who contributed the sample, the batch it was processed in, the center the sample was processed at, and the RNA integrity number (RIN score) for the sample.

#### Relational data

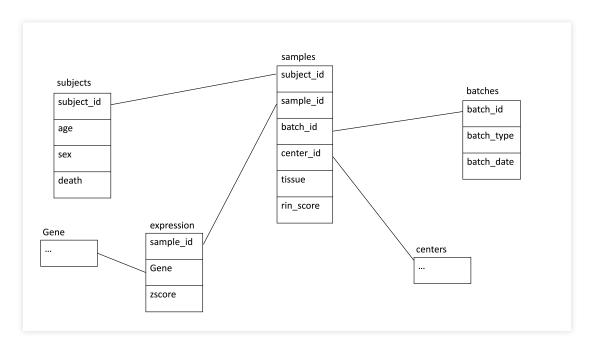
The subject data table contains some subject demographic information. Death refers to circumstances surrounding death.

The batch data containts the batch type and the dates the batches were run (we were been working a bit with this date data aggregated into counts of samples earlier).

```
gtex_batch_link = "https://raw.githubusercontent.com/alejandroschuler/r4ds-
courses/9e4fb21ccf93a83e2b6004b9aa467426806f8589/data/gtex_metadata/gtex_batch_metadata.csv"
gtex_batch_data = read_csv(file = gtex_batch_link, col_types = cols())
head(gtex_batch_data, 2L)
# A tibble: 2 x 3
```

#### Relational data

• These data are not independent of each other. Subjects described in the subject data are referenced in the sample data, and the batches referenced in the sample data are in the batch data. The sample ids from the sample data are used for accessing expression data.



- subject connects to sample via a single variable, subject\_id.
- sample connects to batch through the batch\_id variable.

#### Relational + tidy data

For the expression data, we have been using the gtex\_data expression data frame:

The expression data on the previous slide is formatted slightly differently:

#### An example join

- Imagine we want to add subject information to the sample data
- We can accomplish that with a **join**:

```
qtex sample data %>%
  inner_join(gtex_subject_data, by = "subject_id")
# A tibble: 312 x 9
   subject_id sample_id batch_id center_id tissue rin_score sex
                                                                            death
                                                                      age
   <chr>
              <chr>
                          <chr>
                                   <chr>
                                              <chr>
                                                         <dbl> <chr> <chr> <chr>
1 GTEX-11DXZ 0003-SM-5... BP-39216 B1
                                              Blood
                                                                male 50-59 ventil...
                                                          NA
2 GTEX-11DXZ 0126-SM-5... BP-44460 B1
                                              Liver
                                                          7.9 male 50-59 ventil...
                                                           8.3 male 50-59 ventil...
 3 GTEX-11DXZ 0326-SM-5... BP-44460 B1
                                              Heart
 4 GTEX-11DXZ 0726-SM-5... BP-43956 B1
                                                           7.8 male 50-59 ventil...
                                              Lung
 5 GTEX-11GSP 0004-SM-5... BP-39412 B1
                                              Blood
                                                                fema... 60-69 sudden...
                                                           6.2 fema... 60-69 sudden...
 6 GTEX-11GSP 0626-SM-5... BP-44902 B1
                                              Liver
7 GTEX-11GSP 0726-SM-5... BP-44902 B1
                                              Lung
                                                           6.9 fema... 60-69 sudden...
8 GTEX-11GSP 1226-SM-5... BP-44902 B1
                                              Heart
                                                           7.9 fema... 60-69 sudden...
 9 GTEX-11NUK 0004-SM-5... BP-39723 B1
                                              Blood
                                                                male 50-59 sudden...
10 GTEX-11NUK 0826-SM-5... BP-43730 B1
                                                           7.4 male 50-59 sudden...
                                              Lung
# ... with 302 more rows
```

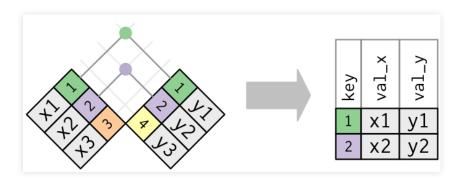
#### Joins

```
x = tibble(
  key = c(1, 2, 3),
  val_x = c("x1", "x2", "x3")
)

y = tibble(
  key = c(1, 2, 4),
  val_y = c("y1", "y2", "y3")
)
```

X		у		
1	x1		1	у1
2	x2		2	y2
3	x3		4	y3

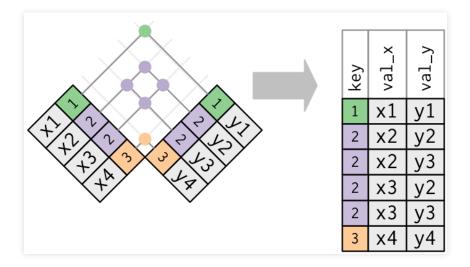
- An inner join matches pairs of observations when their "keys" are equal
- the column that is joined on is specified as a "key" with the argument by="column"



### Duplicate keys

```
x = tibble(
  key = c(1, 2, 2, 3),
  val_x = c("x1", "x2", "x3", "x4")
)

y = tibble(
  key = c(1, 2, 2, 4),
  val_y = c("y1", "y2", "y3", "y4")
)
```



When keys are duplicated, multiple rows can match multiple rows, so each possible combination is produced

## Specifying the keys

```
gtex_sample_data %>%
  inner_join(gtex_subject_data, by = "center_id")
Error: Join columns must be present in data.
x Problem with `center_id`.
```

• Why does this fail?

#### Specifying the keys

• When keys have different names in different dataframes, the syntax to join is:

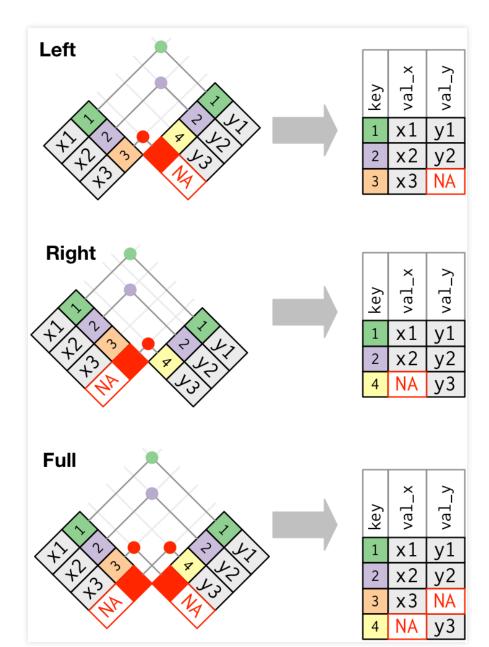
```
head (gtex data, 2)
# A tibble: 2 \times 7
 Gene Ind Blood Heart Lung Liver NTissues
 <chr> <chr> <dbl> <dbl> <dbl> <dbl> <dbl>
                                            <dbl>
1 A2ML1 GTEX-11DXZ -0.14 -1.08 NA -0.66
2 A2ML1 GTEX-11GSP -0.5 0.53 0.76 -0.1
head(gtex_subject_data, 2)
# A tibble: 2 x 4
  subject id sex age death
 <chr> <chr> <chr> <chr>
1 GTEX-11DXZ male 50-59 ventilator
2 GTEX-11GSP female 60-69 sudden but natural causes
gtex data %>%
  inner join (gtex subject data, by = c("Ind" = "subject id")) %>%
 head(5L)
# A tibble: 5 x 10
 Gene Ind Blood Heart Lung Liver NTissues sex
                                                      age
                                                            death
 <chr> <chr> <dbl> <dbl> <dbl> <dbl> <dbl> <chr> <chr> <<rp> <dbl> <chr> <chr> 
1 A2ML1 GTEX-11D... -0.14 -1.08 NA
                                  -0.66
                                              3 male 50-59 ventilator
2 A2ML1 GTEX-11G... -0.5 0.53 0.76 -0.1
                                              4 fema... 60-69 sudden but natur...
3 A2ML1 GTEX-11N... -0.08 -0.4 -0.26 -0.13
                                              4 male 50-59 sudden but natur...
4 A2ML1 GTEX-11N... -0.37 0.11 -0.42 -0.61
                                              4 male 60-69 sudden but natur...
5 A2ML1 GTEX-11T... 0.3 -1.11 0.59 -0.12
                                              4 male 20-29 ventilator
```

### Exercise: finding expression of specific samples

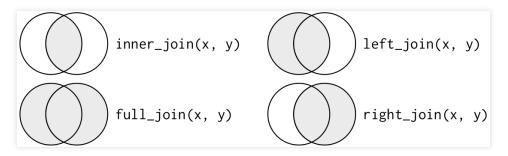
Use joins to find the samples collected in 2015 with high blood expression (Z>3) of "KRT19" in males. Start with the batch\_data\_year; this data has an extra extracted column with the year (we'll go over how this worked in the next lecture).

Note that you'll have to join to other data frames the sample data frame to put this together.

## Other joins



- A left join keeps all observations in x.
- A right join keeps all observations in y.
- A full join keeps all observations in x and y.



- Left join should be your default
  - it looks up additional information in other tables
  - preserves all rows in the table you're most interested in

#### Joining on multiple columns

• It is often desirable to find matches along more than one column, such as month and year in this example. Here we're joining tissue sample counts with total sample counts.

```
gtex_tissue_month_link = "https://raw.githubusercontent.com/alejandroschuler/r4ds-
courses/9e4fb21ccf93a83e2b6004b9aa467426806f8589/data/gtex metadata/gtex tissue month year.csv"
gtex_tissue_month =
  read_csv(file = gtex_tissue_month_link, col_types = cols()) %>%
 filter(tissue %in% c("Blood", "Heart", "Liver", "Lung"))
head(gtex_tissue_month, 2L)
# A tibble: 2 \times 4
 tissue month year tiss_samples
 <chr> <dbl> <dbl> <dbl>
1 Blood 1 2012 25
2 Blood 1 2013 16
gtex_samples_by_month =
  read_csv(file = gtex_samples_time_link, col_types = cols()) %>%
 rename (total num samples = num samples)
head(gtex_samples_by_month, 2L)
# A tibble: 2 \times 3
 month year total_num_samples
 <dbl> <dbl> <dbl>
     5 2011
                           20
     6 2011
                           44
```

### Joining on multiple columns

This is also possible if the columns have different names:

```
gtex_data_long = gtex_data %>%
 pivot longer(cols = c("Blood", "Heart", "Lung", "Liver"), names to = "tissue",
   values to = "zscore")
head (gtex data long, n = 2L)
# A tibble: 2 x 5
 Gene Ind NTissues tissue zscore
 <chr> <chr> <dbl> <chr>
1 A2ML1 GTEX-11DXZ 3 Blood -0.14
2 A2ML1 GTEX-11DXZ 3 Heart -1.08
head (gtex sample data, n = 2L)
# A tibble: 2 x 6
 subject_id sample_id batch_id center_id tissue rin_score
 <chr> <chr> <chr> <chr>
                                                   <dbl>
1 GTEX-11DXZ 0003-SM-5807X BP-39216 B1 Blood NA
2 GTEX-11DXZ 0126-SM-5EGGY BP-44460 B1 Liver 7.9
gtex data long %>%
 inner join(gtex sample data, by = c("tissue", "Ind" = "subject id")) %>%
 head(n = 4L)
# A tibble: 4 x 9
 Gene Ind NTissues tissue zscore sample id batch id center id rin score
 <chr> <chr> <dbl> <chr> <dbl> <chr> <dbl> <chr>
                                                                   <dbl>
1 A2ML1 GTEX-11... 3 Blood -0.14 0003-SM-58... BP-39216 B1 NA
2 A2ML1 GTEX-11... 3 Heart -1.08 0326-SM-5E... BP-44460 B1 8.3
3 A2ML1 GTEX-11... 3 Lung NA 0726-SM-5N... BP-43956 B1 4 A2ML1 GTEX-11... 3 Liver -0.66 0126-SM-5E... BP-44460 B1
                                                                    7.8
                                                                     7.9
```

#### Join problems

- Joins can be a source of subtle errors in your code
- check for NAs in variables you are going to join on
- make sure rows aren't being dropped if you don't intend to drop rows
  - checking the number of rows before and after the join is not sufficient. If you have an inner join with duplicate
    keys in both tables, you might get unlucky as the number of dropped rows might exactly equal the number of
    duplicated rows
- anti\_join() and semi\_join() are useful tools (filtering joins) to diagnose problems
  - anti\_join() keeps only the rows in x that don't have a match in y
  - semi\_join() keeps only the rows in x that do have a matchin y

## Exercise: Looking for variables related to data missingness

It is important to make sure that the missingness in the expression data is not related to variables present in the data. Use the tables <code>batch\_data\_year</code>, <code>sample\_data</code>, <code>subject\_data</code>, <code>and</code> the <code>gtex\_data</code> to look at the relationship between missing gene values and other variables in the data.