

Session 3: Tidyverse

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06/29/2020

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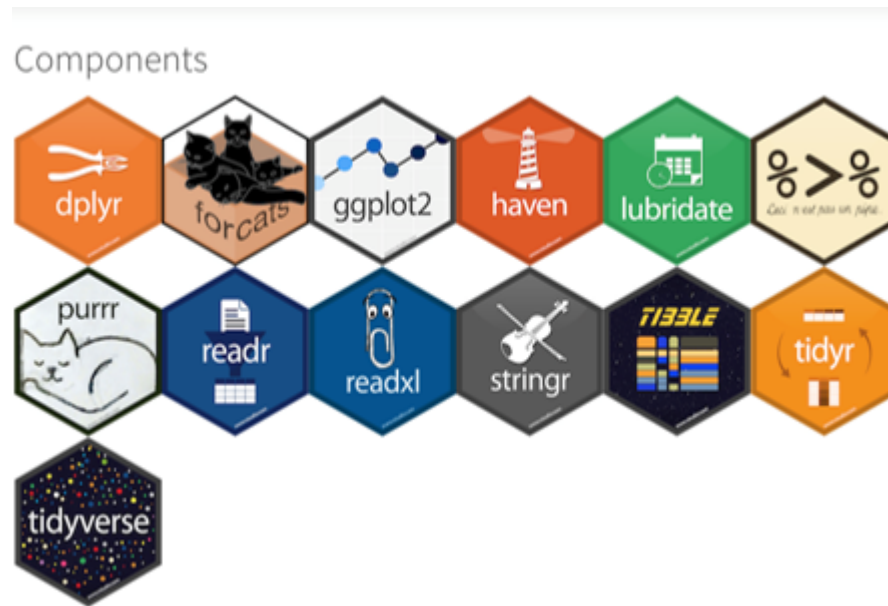
Objectives

- What is tidyverse and packages in tidyverse?
- readr
- dplyr

What is tidyverse?

The tidyverse is a collection of open source R packages introduced by Hadley Wickham and his team that “share an underlying design philosophy, grammar, and data structures” of **tidy data**. Tidy data is a standard method of displaying a multivariate set of data is in the form of a data matrix in which rows correspond to sample individuals and columns to variables, so that the entry in the i^{th} row and j^{th} column gives the value of the j^{th} variate as measured or observed on the i^{th} individual.

Packages in tidyverse



Installing tidyverse

```
install.packages("tidyverse")
```

```
# Call library  
library(tidyverse)
```

```
## -- Attaching packages ----- tidyverse_
```

```
## v ggplot2 3.2.1    v purrr   0.3.3  
## v tibble  2.1.3    v dplyr   0.8.3  
## v tidyr   1.0.0    v stringr 1.4.0  
## v readr   1.3.1    v forcats 0.4.0
```

```
## -- Conflicts ----- tidyverse_  
## x dplyr::filter() masks stats::filter()  
## x dplyr::lag()    masks stats::lag()
```

The readr package

Tidyverse packages providing rectangular data importing and saving functions. Cheatsheets can be found [here](#). `[readr]` is used for turning flat files into data frames.

Importing data

- `read_csv("file.csv")`: Comma Delimited Files
- `read_csv2("file2.csv")`: Semi-colon Delimited Files
- `read_delim("file.txt", delim = "|")`: Files with Any Delimiter
- `read_fwf("file.fwf", col_positions = c(1, 3, 5))`: Fixed Width Files
- `read_tsv("file.tsv")`: Tab Delimited Files

Let us import the Puromycin dataset

```
Puromycin <- read_csv("Puromycin.csv",  
  #col_names = FALSE, # skip header row  
  #col_names = c("x", "y", "z"), # provide header  
  #skip = n, # skip nth row  
  col_types = cols(conc = col_double(),  
                    rate = col_double(),  
                    state = col_character()))  
  
# Open a tab  
View(Puromycin)  
  
# First 5 rows  
head(Puromycin)  
  
# Last 5 rows  
tail(Puromycin)  
  
# A glimpse of the dataset  
glimpse(Puromycin)
```

Saving datafiles

- `write_csv(x, path)`: Comma delimited file
- `write_delim(x, path, delim = " ")`: File with arbitrary delimiter
- `write_excel_csv(x, path)`: CSV for excel
- `write_tsv(x, path)`: write_tsv

Let us save a file of our own. Let's the Theophylline data into a CSV file.

```
# Take a look at the data  
View(Theoph)  
  
write_csv(Theoph,  
  "Theophylline.csv",  
  # delim = " ", #for delimited files  
  # na = "-99", # String used for missing values. Defaults to NA  
  # append = T/F #If FALSE, will overwrite existing file.  
  #If TRUE, will append to existing file.  
  #In both cases, if file does not exist a new file is created.  
  # col_names = T/F #Write columns names at the top of the file?  
)
```

The dplyr package

dplyr is a grammar of data manipulation, providing a consistent set of verbs that help you solve the most common data manipulation challenges. But before we explore dplyr, we need to understand the pipe/%>% operator from the magrittr

The magrittr package

It offers a set of operators which make your code more readable. Pipes are a powerful tool for clearly expressing a sequence of multiple operations.

Some basic piping rules:

- `x %>% f` is equivalent to `f(x)`
- `x %>% f(y)` is equivalent to `f(x, y)`
- `x %>% f() %>% g() %>% h()` is equivalent to `h(g(f(x)))`
- `x %>% f(y, .)` is equivalent to `f(y, x)`
- `x %>% f(y, z = .)` is equivalent to `f(y, z = x)`

```
x <- rnorm(n=10,mean=20,sd=5)

# Let's say I want to round the vector to nearest integer and then take mean

# non-pipe formulation
mean(round(x,digits = 0))

#pipe formulation
x %>%
  round(digits = 0) %>% # same as round(.,digits=0)
  mean() # same as mean(.)
```

Back to dplyr

This package is massive and beautiful. We will be covering only the basics of this package, so please look at the [cheatsheets](#)

Let us use the Theophylline data to explore this package. First, let's start with summarizing the dataset

```
Theo_summ <-
  Theoph %>%
  #Summarise the data
  #Calculate number of subjects, weight and dose ditributions
  summarise(nID = n_distinct(Subject), # count the number of individuals
    `Weight (Mean,SD)` = paste(Wt %>% mean() %>% round(digits=1),
                                "(",
                                Wt %>% sd() %>% round(digits=1),
                                ")",
                                sep=""),
    `Dose (Median,Range)` = paste(Dose %>% median() %>% round(digits=1),
                                "(",
                                range(Dose)[1],
                                "-",
                                range(Dose)[2],
```

```

    ")",
    sep=""))

# Individual Cmax/Cmed

Cmax.med <-
  Theoph %>%
  group_by(Subject) %>%
  summarise(Cmax = max(conc),
            Cmed = median(conc))

```

Other useful summarize functions:

- `summarise_all()` - Apply funs to every column.
- `summarise_at()` - Apply funs to specific columns.
- `summarise_if()` - Apply funs to all cols of one type.

Now let's try to manipulate cases/vairables:

```

# Filter subjects with dose less than 4.5
# Add a column of weight descriptor
# Select only the Subject, Time and conc columns and weight descriptor
# Renmae Subject to ID and conc to DV

Theop2 <-
  Theoph %>%
  filter(Dose < 4.5) %>%
  mutate(WTLT60 = if_else(Wt <= 60,1,0)) %>%
  select(Subject, Time, conc, WTLT60) %>%
  rename(ID=Subject,DV=conc)

## Let's create a NONMEM dataset

# Let's start with the dosing rows

dosing <-
  Theoph %>%
  group_by(Subject) %>%
  filter(Time == 0) %>%
  mutate(conc = NA)

# Concentration rows

conc <-
  Theoph %>%
  group_by(Subject) %>%
  filter(!(Time == 0 & conc == 0)) %>%
  mutate(Time = if_else(Time == 0, 0.001, Time),
         Dose = NA,
         LNDV = log(conc) %>% round(2))

```

```
# join the two

Theop_NM <-
  dosing %>%
  bind_rows(conc) %>%
  arrange(Subject,.by_group=TRUE) %>%
  mutate(MDV = if_else(is.na(Dose),0,1),
         EVID = if_else(is.na(Dose),0,1),
         WTLT60 = if_else(Wt <= 60,1,0)) %>%
  rename(ID=Subject,DV=conc,AMT=Dose)
```

Other useful manipulation functions:

- `distinct(.data, ..., .keep_all = FALSE)` - Remove rows with duplicate values
- `sample_frac(tbl, size = 1, replace = FALSE)` - Randomly select fraction of rows
- `slice(.data, .)` - Select rows by position
- `transmute(.data, .)` - Compute new column(s), drop others
- `mutate_all(.tbl, .funs, .)` - Apply funs to every column
- `mutate_at(.tbl, .cols, .funs, .)` - Apply funs to specific columns
- `add_column(.data, ..., .before = NULL, .after = NULL)` - Add new column(s)

Now, let's look at combining tables

```
# Let's say you have a dataset with additional demographic data

demog <- data.frame(
  ID = unique(Theop_NM$ID),
  SEX = sample(c("M","F"),12,replace = TRUE),
  BSA = rnorm(12,1.73,0.2) %>% round(2),
  GENO = sample(c("PM","WT","FM"),12,replace = TRUE)
)

# Merge this with the NM dataset

Theop_NM2 <-
  Theop_NM %>%
  left_join(demog,by="ID")

# But we cannot have strings in NONMEM, so let us convert them to numeric

Theop_NM3 <-
  Theop_NM2 %>%
  mutate(SEX2 = if_else(SEX == "M",1,0),
         GENO2 = case_when(
           GENO == "WT" ~ 0,
           GENO == "PM" ~ 1,
           GENO == "FM" ~ 2)
  )
```

Other important join functions:

- `right_join(x, y, by = NULL,.)` - Join matching values from x to y

- `inner_join(x, y, by = NULL, .)` - Retain only rows with matches
- `full_join(x, y, by = NULL, .)` - Retain all values, all rows
- `semi_join(x, y, by = NULL, .)` - Return rows of x that have a match in y
- `anti_join(x, y, by = NULL, .)` - Return rows of x that do not have a match in y

Let's write out this file using the `write_csv()` function

```
write_csv(Theop_NM3, "Theop06292020.csv", na = ".")
```

Takehome exercises:

- The best exercise is to start working with your own dataset. The complexities and intricacies that exist in real world data cannot be replicated with dummy datasets
- Online exercises:
 - Basics
 - Intermediate
 - Advanced