

# Pre-processing and plotting data

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## Workflow

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If you haven't already, load the necessary packages.

```
library(gcplyr)

library(dplyr)
library(ggplot2)
library(lubridate)
```

```
#This code was explained in sections 2 and 3
#Here we're re-running it so it's available for us to work with
example_tidydata <- trans_wide_to_tidy(example_widedata,
                                     id_cols = "Time")

example_design <- make_design(
  pattern_split = ",", nrows = 8, ncols = 12,
  "Bacteria_strain" = make_designpattern(
```

```

values = paste("Strain", 1:48),
rows = 1:8, cols = 1:6,
pattern = 1:48,
byrow = TRUE),
"Bacteria_strain" = make_designpattern(
  values = paste("Strain", 1:48),
  rows = 1:8, cols = 7:12,
  pattern = 1:48,
  byrow = TRUE),
"Phage" = make_designpattern(
  values = c("No Phage"),
  rows = 1:8, cols = 1:6,
  pattern = "1"),
"Phage" = make_designpattern(
  values = c("Phage Added"),
  rows = 1:8, cols = 7:12,
  pattern = "1"))
ex_dat_mrg <- merge_dfs(example_tidydata, example_design)
#> Joining, by = "Well"

```

## Pre-processing

Now that we have our data and designs merged, we're almost ready to start processing and analyzing them. However, first we need to carry out any necessary pre-processing steps, like excluding wells that were contaminated or empty, and converting time formats to numeric.

### Pre-processing: excluding data

In some cases, we want to remove some of the wells from our growth curves data before we carry on with downstream analyses. For instance, they may have been left empty, contained negative controls, or were contaminated. We can use `dplyr`'s `filter` function to remove those wells that meet criteria we want to exclude.

For instance, let's imagine that we realized that we put the wrong media into Well B1, and so we should remove it from our analyses. In that case, we can simply:

```

example_data_and_designs_filtered <- filter(ex_dat_mrg, Well != "B1")
head(example_data_and_designs_filtered)
#>   Time Well Measurements Bacteria_strain   Phage
#> 1    0   A1         0.003      Strain 1 No Phage
#> 2    0   C1         0.002      Strain 13 No Phage
#> 3    0   D1         0.002      Strain 19 No Phage
#> 4    0   E1         0.002      Strain 25 No Phage
#> 5    0   F1         0.001      Strain 31 No Phage
#> 6    0   G1         0.002      Strain 37 No Phage

```

Now we can see that all rows from Well B1 have been excluded. We could do something similar if we realized that a Bacterial strain was contaminated. For instance, if strain 13 was contaminated, we could exclude it (and Well B1) as follows:

```
example_data_and_designs_filtered <-
  filter(ex_dat_mrg,
    Well != "B1", Bacteria_strain != "Strain 13")
head(example_data_and_designs_filtered)
#>   Time Well Measurements Bacteria_strain   Phage
#> 1 0 A1 0.003 Strain 1 No Phage
#> 2 0 D1 0.002 Strain 19 No Phage
#> 3 0 E1 0.002 Strain 25 No Phage
#> 4 0 F1 0.001 Strain 31 No Phage
#> 5 0 G1 0.002 Strain 37 No Phage
#> 6 0 H1 0.002 Strain 43 No Phage
```

## Pre-processing: converting dates & times into numeric

Growth curve data produced by a plate reader often encodes the timestamp information as a string (e.g. “2:45:11” for 2 hours, 45 minutes, and 11 seconds), while downstream analyses need timestamp information as a numeric (e.g. number of seconds elapsed). Luckily, others have written great packages that make it easy to convert from common date-time text formats into plain numeric formats. Here, we’ll see how to use `lubridate` to do so:

First we have to create a data frame with time saved as it might be by a plate reader. As usual, **don’t worry how this block of code works**, since it’s just creating an example file in the same format as that output by a plate reader.

```
ex_dat_mrg$Time <-
  paste(ex_dat_mrg$Time %/% 3600,
    formatC((ex_dat_mrg$Time %/% 3600) %/% 60,
      width = 2, flag = 0),
    formatC((ex_dat_mrg$Time %/% 3600) %/% 60,
      width = 2, flag = 0),
    sep = ":")
```

Let’s take a look at this data.frame. This shows the `Time` column as it might be written by a plate reader.

```
head(ex_dat_mrg)
#>   Time Well Measurements Bacteria_strain   Phage
#> 1 0:00:00 A1 0.003 Strain 1 No Phage
#> 2 0:00:00 B1 0.001 Strain 7 No Phage
#> 3 0:00:00 C1 0.002 Strain 13 No Phage
#> 4 0:00:00 D1 0.002 Strain 19 No Phage
#> 5 0:00:00 E1 0.002 Strain 25 No Phage
#> 6 0:00:00 F1 0.001 Strain 31 No Phage
```

We can see that our `Time` aren’t written in an easy numeric. Instead, they’re in a format that’s easy for a human to understand (but unfortunately not very usable for analysis).

Let’s use `lubridate` to convert this text into a usable format. `lubridate` has a whole family of functions that can parse text with hour, minute, and/or second components. You can use `hms` if your text contains hour, minute, and second information, `hm` if it only contains hour and minute information, and `ms` if it only contains minute and second information.

Since the example has all three, we’ll use `hms`. Once `hms` has parsed the text, we’ll use another function to convert the output of `hms` into a pure numeric value: `time_length`. By default, `time_length` returns in units

of seconds, but you can change that by changing the `unit` argument to `time_length`. See `?time_length` for details.

```
#We have previously loaded lubridate, but if you haven't already then  
#make sure to add the line:  
# library(lubridate)
```

```
ex_dat_mrg$Time <- time_length(hms(ex_dat_mrg$Time))
```

```
head(ex_dat_mrg)
```

```
#>   Time Well Measurements Bacteria_strain    Phage  
#> 1    0   A1         0.003      Strain 1 No Phage  
#> 2    0   B1         0.001      Strain 7 No Phage  
#> 3    0   C1         0.002      Strain 13 No Phage  
#> 4    0   D1         0.002      Strain 19 No Phage  
#> 5    0   E1         0.002      Strain 25 No Phage  
#> 6    0   F1         0.001      Strain 31 No Phage
```

And now we can see that we've gotten nice numeric `Time` values! So we can proceed with the next steps of the analysis.

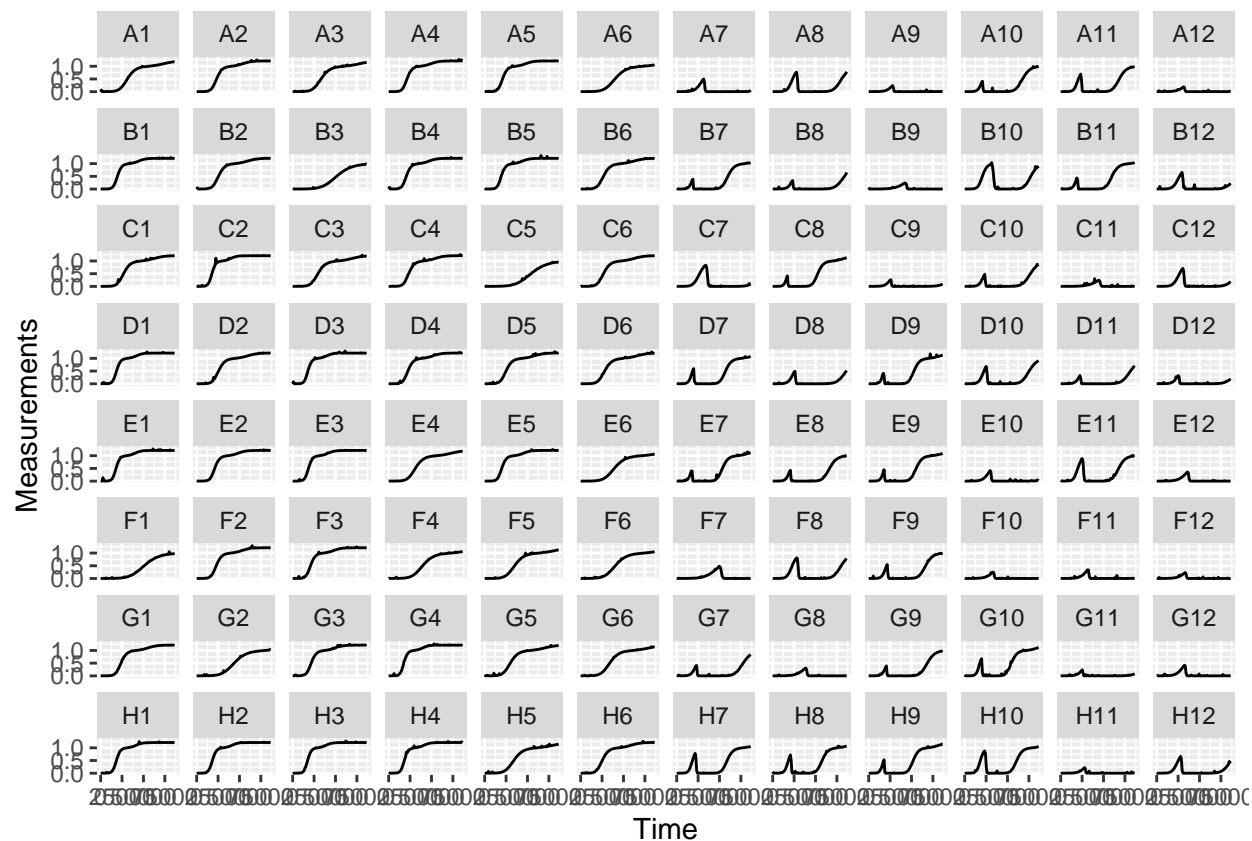
## Plotting your data

Once your data has been merged and times have been converted to numeric, we can easily plot our data using the `ggplot2` package. That's because `ggplot2` was specifically built on the assumption that data would be tidy-shaped, which ours is! We won't go into depth on how to use `ggplot` here, but there are three main commands to the plot below:

- `ggplot` - the `ggplot` function is where you specify the `data.frame` you would like to use and the *aesthetics* of the plot (the x and y axes you would like)
- `geom_line` - tells `ggplot` how we would like to plot the data, in this case with a line (another common `geom` for time-series data is `geom_point`)
- `facet_wrap` - tells `ggplot` to plot each `Well` in a separate facet

We'll be using this format to plot our data throughout the remainder of this vignette

```
#We have previously loaded ggplot2, but if you haven't already then  
#make sure to add the line:  
# library(ggplot2)  
  
#First, we'll reorder the Well levels so they plot in the correct order  
ex_dat_mrg$Well <-  
  factor(ex_dat_mrg$Well,  
         levels = paste(rep(LETTERS[1:8], each = 12), 1:12, sep = ""))  
  
ggplot(data = ex_dat_mrg, aes(x = Time, y = Measurements)) +  
  geom_line() +  
  facet_wrap(~Well, nrow = 8, ncol = 12)
```



Generally speaking, **from here on you should plot your data frequently**, and in every way you can think of! **After every processing and analysis step, visualize both the input data and output data** to understand what the processing and analysis steps are doing and whether they are the right choices for your particular data (this vignette will be doing that too!)

## What's next?

Now that you've pre-processed and visualized your data, it's time to process (in most cases) and analyze (pretty much always) it!

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