# Pre-processing and plotting data

### Mike Blazanin

## Contents

where are we so far!	1
Pre-processing	2
Pre-processing: excluding data	2
Pre-processing: converting dates & times into numeric	2
Plotting your data	3
What's next?	4

### Where are we so far?

- 1. Introduction: vignette("gc01\_gcplyr")
- 2. Importing and transforming data: vignette("gc02 import reshape")
- 3. Incorporating experimental designs: vignette("gc03\_incorporate\_designs")
- 4. Pre-processing and plotting your data: vignette("gc04\_preprocess\_plot")
- 5. Processing your data: vignette("gc05\_process")
- 6. Analyzing your data: vignette("gc06\_analyze")
- 7. Dealing with noise: vignette("gc07 noise")
- 8. Statistics, merging other data, and other resources: vignette("gc08\_conclusion")
- 9. Working with multiple plates: vignette("gc09\_multiple\_plates")

So far, we've imported and transformed our measures, then combined them with our design information. Now we're going to do some final pre-processing steps and show how to easily plot our data with ggplot.

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

```
library(gcplyr)

library(ggplot2)
library(lubridate)
#>
#> Attaching package: 'lubridate'
#> The following objects are masked from 'package:base':
#>
#> date, intersect, setdiff, union
```

## **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 that strain 13 was contaminated. To exclude them from our analyses, we can simply:

```
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
       O A1
                    0.002
                                Strain 1 No Phage
#> 2
          D1
                    0.002
                                Strain 19 No Phage
       0
#> 3
       0 E1
                    0.002
                                Strain 25 No Phage
     0 F1
#> 4
                    0.002
                                Strain 31 No Phage
#> 5
     0 G1
                    0.002
                                Strain 37 No Phage
#> 6
                                Strain 43 No Phage
           H1
                     0.002
```

### 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.

```
#> 2 0:00:00
               B1
                          0.002
                                       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
                                      Strain 25 No Phage
                          0.002
#> 6 0:00:00
               F1
                          0.002
                                      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.

Once hms has parsed the text, we'll use time\_length to convert the output of hms into a pure numeric value. By default, time\_length returns in units of seconds, but you can change that by changing the unit argument to time\_length.

```
# 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), unit = "hour")</pre>
head(ex_dat_mrg)
     Time Well Measurements Bacteria_strain
                                               Phage
#> 1
        O A1
                      0.002
                                  Strain 1 No Phage
#> 2
                      0.002
                                  Strain 7 No Phage
        0
          B1
#> 3
          C1
                      0.002
                                  Strain 13 No Phage
        0
#> 4
        0
           D1
                      0.002
                                  Strain 19 No Phage
#> 5
        0
           E1
                                  Strain 25 No Phage
                      0.002
#> 6
            F1
                      0.002
                                  Strain 31 No Phage
```

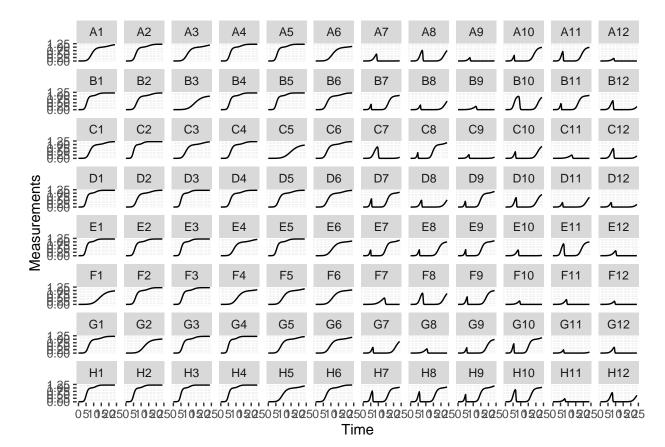
And now we can see that we've gotten nice numeric Time values! So we can proceed with the next steps of the analysis.

# 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



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!

1. Introduction: vignette("gc01\_gcplyr")

- 2. Importing and transforming data: vignette("gc02\_import\_reshape")
- 3. Incorporating experimental designs: vignette("gc03\_incorporate\_designs")
- 4. Pre-processing and plotting your data: vignette("gc04\_preprocess\_plot")
- 5. Processing your data: vignette("gc05\_process")
- 6. Analyzing your data: vignette("gc06\_analyze")
- 7. Dealing with noise: vignette("gc07\_noise")
- 8. Statistics, merging other data, and other resources: vignette("gc08\_conclusion")
- 9. Working with multiple plates: vignette("gc09\_multiple\_plates")