# A Report on Lake Ontario's Microbes

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## Prepare the R Environment

```r
# Load libraries/packages
library(tidyverse)
```

```
## -- Attaching core tidyverse packages ----------------------- tidyverse 2.0.0 --
## v dplyr     1.1.4      v readr     2.1.4
## v forcats   1.0.0      v stringr   1.5.1
## v ggplot2   3.4.4      v tibble    3.2.1
## v lubridate 1.9.3      v tidyr     1.3.0
## v purrr     1.0.2
## -- Conflicts ----------------------------------------- tidyverse_conflicts() --
## x dplyr::filter() masks stats::filter()
## x dplyr::lag()    masks stats::lag()
## i Use the conflicted package (<http://conflicted.r-lib.org/>) to force all conflicts to become errors
```

## Load in the Lake Ontario Data

```r
# Load in lake ontario microbial community data
sample_and_taxon <-
  read_csv("data/sample_and_taxon.csv")
```

```
## Rows: 71 Columns: 15
## -- Column specification -------------------------------------------------------
## Delimiter: ","
## chr  (2): sample_id, env_group
## dbl (13): depth, cells_per_ml, temperature, total_nitrogen, total_phosphorus...
##
## i Use `spec()` to retrieve the full column specification for this data.
## i Specify the column types or set `show_col_types = FALSE` to quiet this message.
```

```r
# Inspect the data
glimpse(sample_and_taxon)
```

```
## Rows: 71
## Columns: 15
## $ sample_id        <chr> "May_12_B", "May_12_E", "May_12_M", "May_17_E", "May~
## $ env_group        <chr> "Deep", "Shallow_May", "Shallow_May", "Shallow_May",~
## $ depth            <dbl> 102.8, 5.0, 15.0, 5.0, 27.0, 5.0, 19.0, 135.0, 5.0, ~
## $ cells_per_ml     <dbl> 2058864, 4696827, 4808339, 3738681, 2153086, 3124920~
## $ temperature      <dbl> 4.07380, 7.01270, 6.13500, 5.99160, 4.66955, 5.97390~
## $ total_nitrogen   <dbl> 465, 465, 474, 492, 525, 521, 539, 505, 473, 515, 47~
```

```
## $ total_phosphorus  <dbl> 3.78, 4.39, 5.37, 4.67, 4.44, 3.71, 4.23, 4.18, 6.64~
## $ diss_org_carbon    <dbl> 2.478, 2.380, 2.601, 2.435, 2.396, 2.283, 2.334, 2.3~
## $ chlorophyll        <dbl> 0.05, 2.53, 3.20, 0.55, 0.48, 0.79, 0.44, 0.22, 3.44~
## $ Proteobacteria     <dbl> 0.4120986, 0.3389293, 0.2762080, 0.4351188, 0.410063~
## $ Actinobacteriota   <dbl> 0.1288958, 0.1861232, 0.2866884, 0.1910769, 0.280123~
## $ Bacteroidota       <dbl> 0.08065717, 0.23470807, 0.21659843, 0.21576244, 0.11~
## $ Chloroflexi        <dbl> 0.19463564, 0.08086689, 0.07032061, 0.08498357, 0.13~
## $ Verrucomicrobiota  <dbl> 0.13249532, 0.10878214, 0.09991639, 0.05752092, 0.06~
## $ Cyanobacteria      <dbl> 2.482454e-04, 9.574640e-03, 1.262830e-02, 1.288730e-~
```

## Microbial Cell Abundance Versus Temperature

```r
# temp on the X
# cell abundance on the Y
# Colored by environmental Group
# make it look nice (e.g. labels, models, etc)

ggplot(data = sample_and_taxon,
       aes(x = temperature, y = cells_per_ml)) +
  geom_point(aes(color = env_group)) +
  labs(x = "Temp (Celsius)", y = "Cell Density (cells/mL)") +
  theme_classic() +
  scale_color_manual(values = c("cornflowerblue",
                                "goldenrod1",
                                "firebrick")) +
  geom_smooth(method = "lm", color = "black") +
  theme(legend.position = "bottom",
        legend.title = element_blank())
```

```
## `geom_smooth()` using formula = 'y ~ x'
```
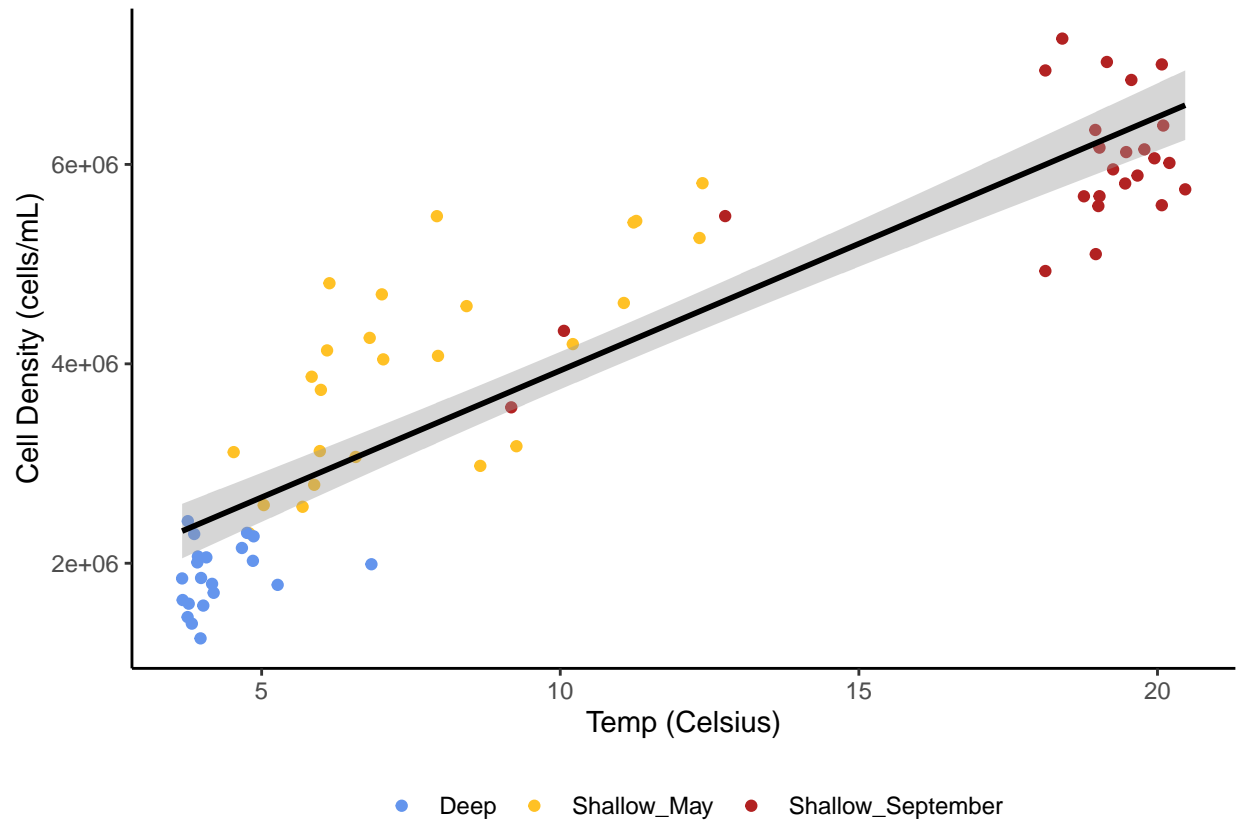
The above plot shows that:

- There's a positive relationship between temperature and cell abundances.
- For example, deep samples are the coldest and have the fewest cells.

The total number of samples is 71. For this set of samples, temperature ranged from a minimum of 3.7 celsius to a maximum of 20.5 celsius.