

MEDICAL MICROBIOLOGY AND INFECTIOUS DISEASES CODING WORKSHOP

Presents

Data visualization using ggtree

INSTRUCTED BY

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INFORMATION FOR PARTICIPANTS

All workshops are being recorded and posted to the MMID Coding Workshop - YouTube

Question and Answer period will not be recorded.

LEARNING OBJECTIVES

- 1. Install and load packages into RStudio
- 2. Load a newick tree and compliment files
- 3. Learn how to create a basic tree
- 4. Changing tree layout
- 5. Adding and customizing tip labels
- 6. How to merge your tree with a heatmap
- 7. How to export the final tree

Data for workshop

- Data set was modified from Arteaga et al. 2020.
 Microbial Genomics. https://doi.org/10.1099/mgen.0.000340
- Based on 26 samples (reduced from 70)

MICROBIAL GENOMICS

SHORT COMMUNICATION

Arteaga et al., Microbial Genomics DOI 10.1099/mgen.0.000340





Genomic characterization of the non-01/non-0139 *Vibrio* cholerae strain that caused a gastroenteritis outbreak in Santiago, Chile, 2018

Mónica Arteaga¹‡, Juliana Velasco¹‡, Shelly Rodriguez¹, Maricel Vidal², Carolina Arellano³, Francisco Silva⁴, Leandro J. Carreño⁵.⁶. Roberto Vidal³.⁶.* and David A. Montero³.⁵.*

Data for workshop

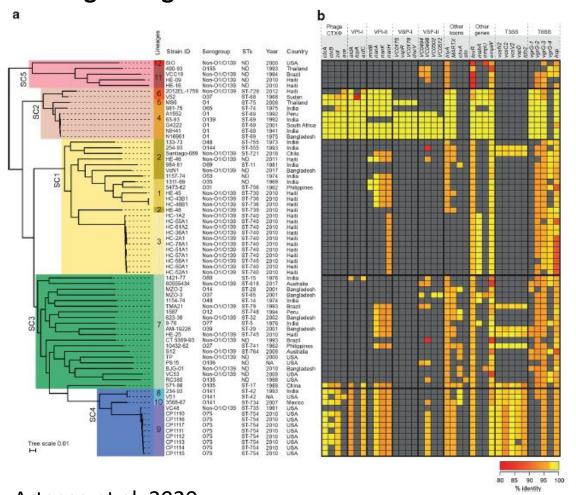
Data files we'll be using for today's workshop:

- 1. Tree file (newick)
 - Created by downloading genome accessions listed in manuscript and running SNVPhyl pipeline in Galaxy
- 2. Metadata (xlsx)
 - Information compiled from manuscript (strain ID, serogroup, year etc.)
- 3. Simulated BlastN (xlsx) 7 genes (reduced from 37)
 - Percent identity was produced using random number generator so will not reflect the results in the paper

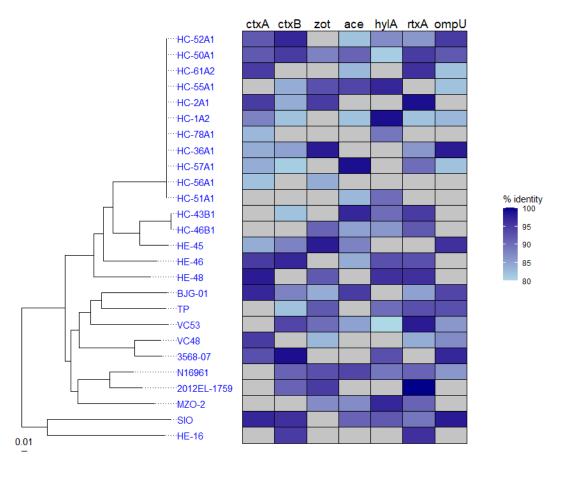
Please **download** and **save** materials in your working directory https://github.com/MMID-coding-workshop

Preview

Original Figure



What we'll be creating!



Arteaga et al. 2020

ggtree

ggtree

- Package for R programming language
- Under Bioconductor project
- Guangchuang Yu: https://yulab-smu.top/treedata-book/
- Data integration, manipulation and visualization of phylogenetic trees
- Customized annotation of tree

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Packages for today's workshop

```
# for reading xlsx files
Readxl
BiocManager # for installing ggtree
Treeio
              # for read.newick function
Phytools
             # for midpoint.root (also has read.newick)
Tidyverse
             # tidying data
              # tree visualization & annotation
ggtree
             # for additional plotting support
ggplot2
```

Install packages

Install the following packages using install.packages function: readxl, BiocManager, treeio, and tidyverse

> install.packages("package name")

Then install ggtree using BiocManager:

> BiocManager::install("ggtree")

Load packages

Load the packages using:

> library("package name")

Note: package installation only has to be done once, but we must load our libraries each time we want to use them

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Load in files

```
# tree file
tree <- read.newick("data/sample tree.newick")</pre>
# metadata file
metadata <- read xlsx("data/metadata.xlsx")</pre>
# blast results file
blast raw <- read xlsx("data/blast results.xlsx")</pre>
```

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Creating a basic tree

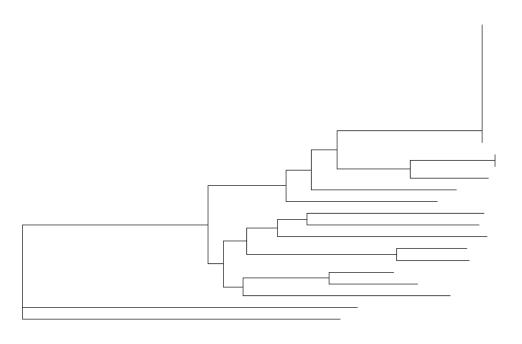
To create a tree, use the ggtree() function:

```
> ggtree(tr)
# or
> tr %>% ggtree()
```

- tr is the phylo object, so for us this would be:
- > ggtree(tree)

Note: for a list of other arguments check out the ggtree help page

> ?ggtree



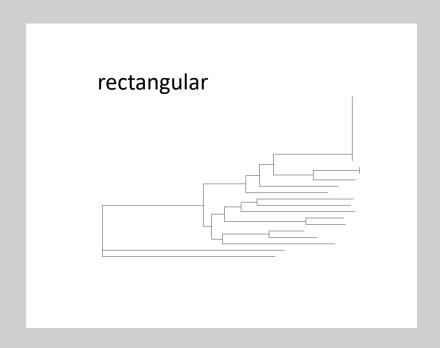
LEARNING OBJECTIVES

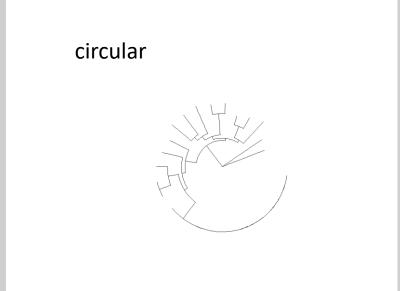
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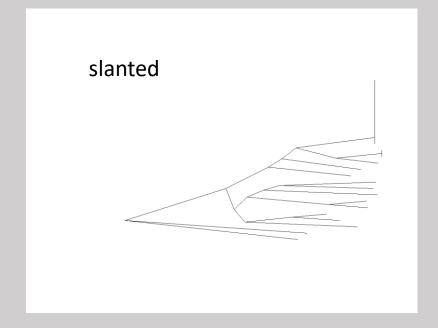
Changing tree layout

To change the layout in ggtree use the **layout** = "shape" argument:

> ggtree(tree, layout = "rectangular")







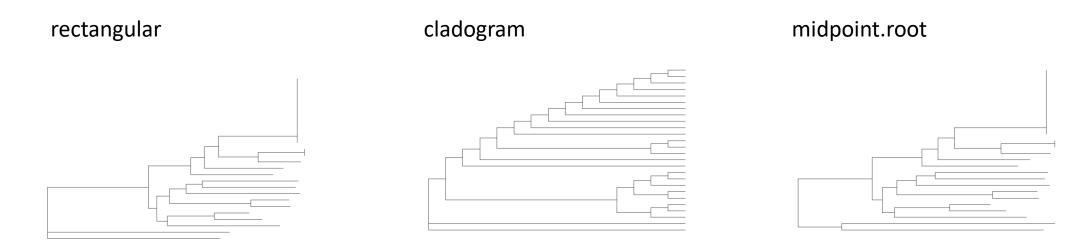
Changing tree layout

A cladogram will show topology without branch length information

> ggtree(tree, branch.length = "none")

Midpoint root: roots the tree at the midpoint of the longest point between two tips.

> ggtree(midpoint.root(tree))

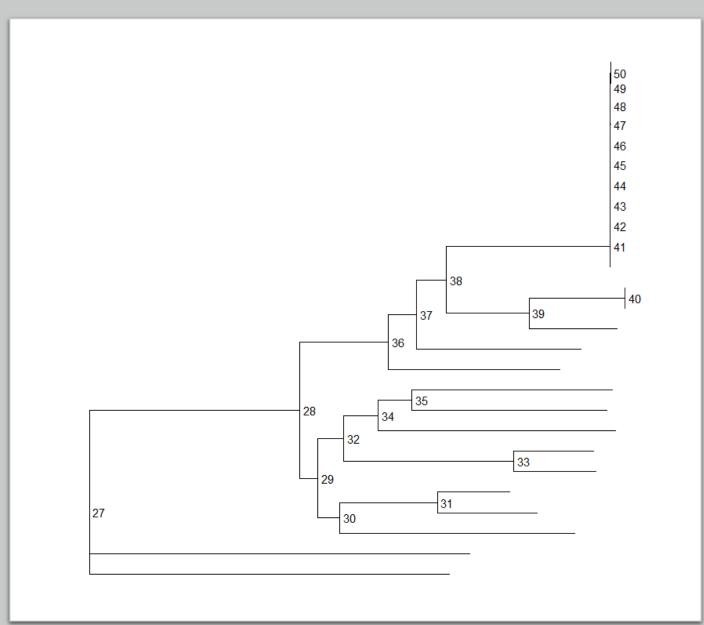


Try changing position of the root node using **root.position** argument

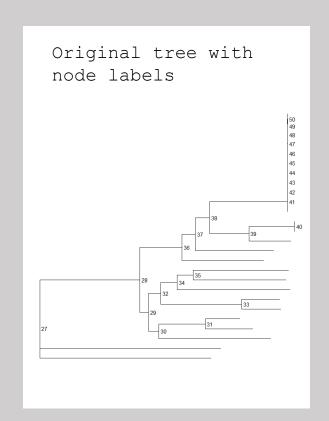
^{**}Caution: rooting can drastically change tree topology and you must use an appropriate method

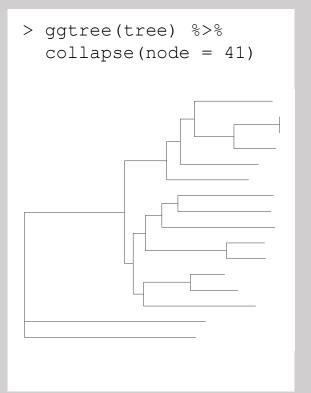
Identify nodes

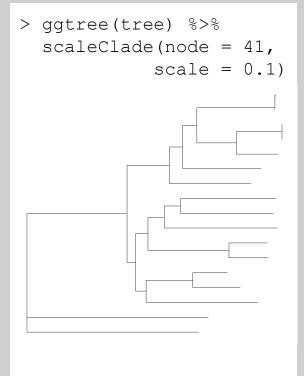
First, we need to identify the nodes
We will do this using geom_text2

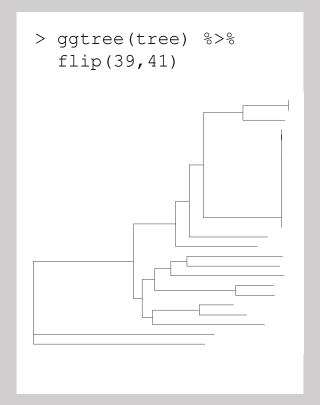


Tree manipulation



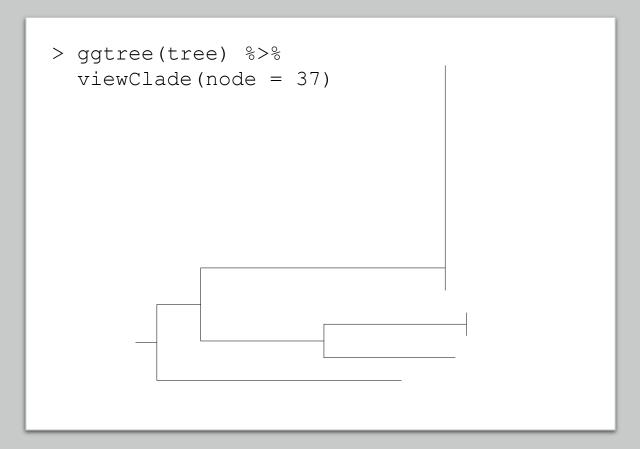






Tree manipulation

- To view a particular clade, we can use viewClade()
- Notice the difference between operators
 - + Plus
 - %>% Pipe





LEARNING OBJECTIVES

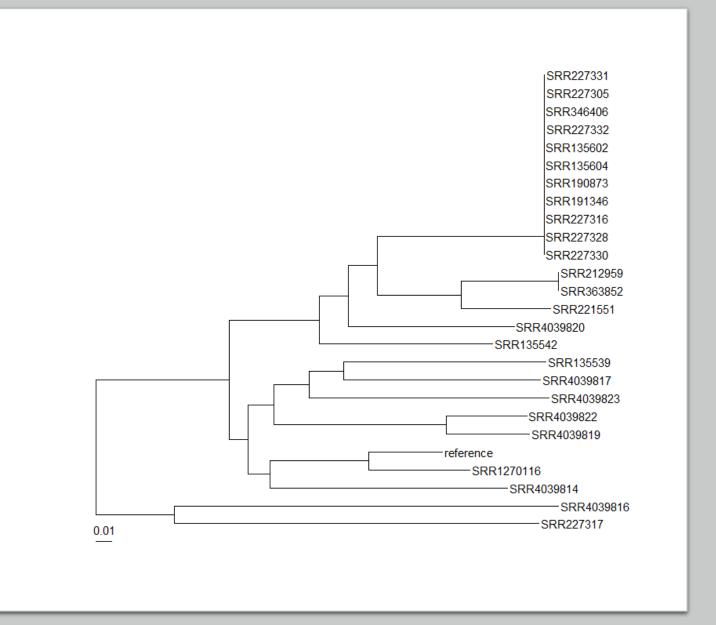
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Adding tip labels

Check tip labels:

```
> head(tree$tip.label)
[1] "SRR227317" "SRR4039816" "SRR4039814" "SRR1270116"
 "reference" "SRR4039819"
> ggtree(midpoint.root(tree)) +
 geom treescale (x = 0, y = 0, # x and y position of the tree scale
                 width = 0.01) + # width of scale
 geom tiplab(size = 4) + # displaying tip labels (size four)
 coord cartesian(clip = 'off') + # turning off the plot limits
 theme (plot.margin = margin (1, 2, 1, 1, "cm")) # add margin
```

Adding tip labels

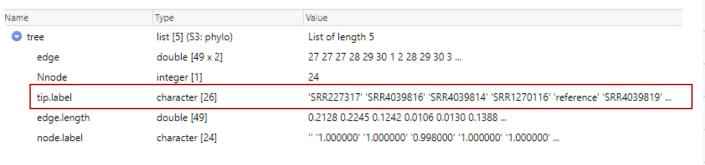


Since we will be linking the metadata file to the tree, we can use a vector to check if there are any file_name observations that are not in the tree

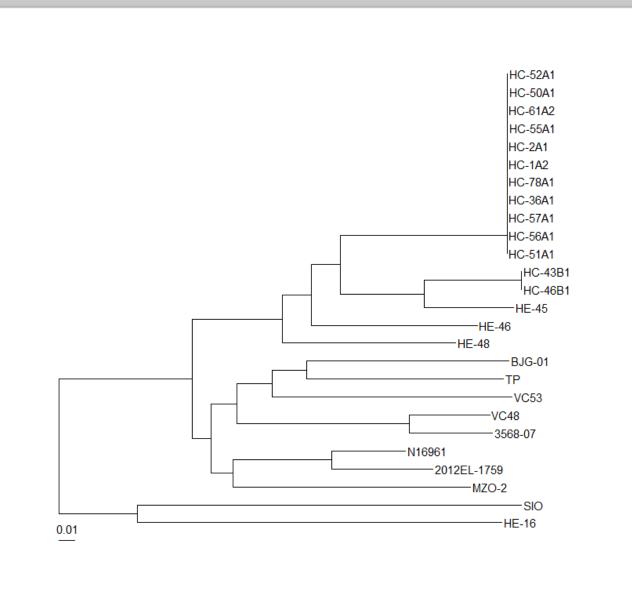
```
> metadata$file_name[!tree$tip.label %in% metadata$file_name]
```

[1] character(0) # all 26 observations match b/w tree and metadata file

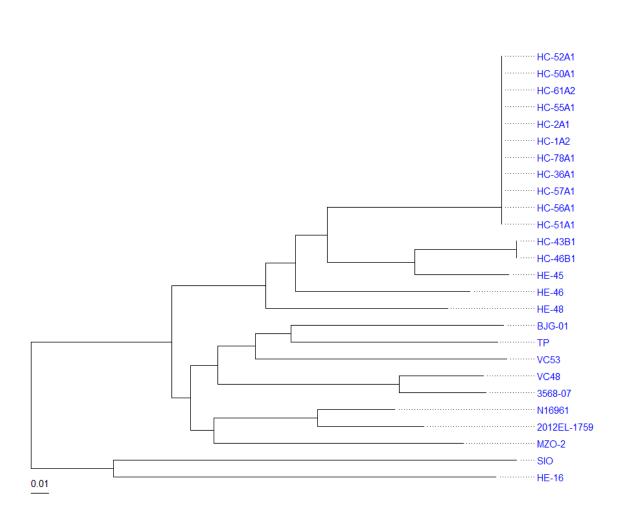
We can also view the data files from our environment:



*	file_name [‡]	strain_ID [‡]	serogroup [‡]
1	reference	N16961	01
2	SRR1270116	2012EL-1759	Non-O1/O139
3	SRR135539	BJG-01	Non-O1/O139
4	SRR135542	HE-48	Non-O1/O139
5	SRR135602	HC-2A1	Non-O1/O139



```
> ggtree(midpoint.root(tree)) %<+%
 metadata +
 geom treescale (x = 0, y = 0,
                 width = 0.01) +
 coord cartesian(clip = 'off') +
 theme(plot.margin = margin(1,2,1,1, "cm")) +
 geom tiplab(aes(label = strain ID),
            color = "blue", # changing font color
            size = 4, # changing font size
            offset = 0.01, # horizontal adjustment of tip labels
            align = TRUE) # this creates a dotted leader line
```



Adding another layer of tip labels

You can keep adding layer by layer by adding a new geom_tiplab, but will need to keep adjusting the offset

Adding another layer of tip labels

```
> gg_simple +
  geom_tiplab(aes(label =
    serogroup), # add in serogroup
  information

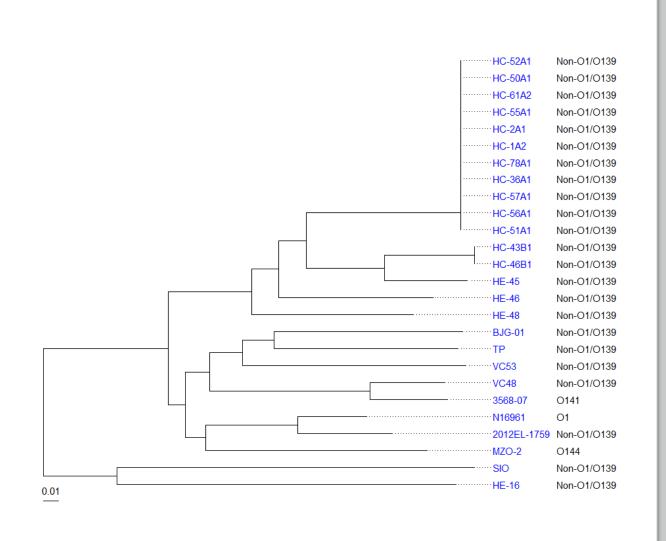
  color = "black",

  offset = .05,

  size = 4,

  align = TRUE,

  linetype = NA)
```



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Merging tree with a heatmap

gheatmap(): a function in ggTree that joins a heatmap matrix and phylogenetic tree

Merging tree with heatmap

We need to prepare the blast file so it will align with tree

- 1. Tidy the blast file
- 2. Set row names to file_name
- 3. Select columns of interest

Merging tree with heatmap

1. Tidy blast file

```
blast df <- blast raw %>% filter(percent identical >= 80) %>%
  pivot wider (names from = gene name,
                  values_from = percent identical)%>%
   relocate(ctxA, .before = ctxB) %>% as.data.frame()
# A tibble: 182 x 3
  file_name gene_name percent_identical
  <chr>>
            <chr>
                               \langle db 1 \rangle
1 reference ctxA
                                                       file_name ctxA ctxB zot ace hylA rtxA ompU
                                 73
 2 reference ctxB
                                                     1 reference
                                                                            93
                                                                                 94
                                                                                      89
                                                                                            91
                                                                                                 86
3 reference zot
                                                     2 SRR1270116
                                                                                          100
                                                                                                 NΑ
4 reference ace
                                                       SRR135539
                                                                                            85
                                                                                                 94
5 reference hyla
                                                       SRR135542
                                                                         NA 92 NA
                                                                                                 NΑ
6 reference rtxA
                                 91
                                                       SRR135602
                                                                                                 NΑ
7 reference ompU
                                 86
8 SRR1270116 ctxA
                                 79
                                                       SRR135604
                                                                                                 83
9 SRR1270116 ctxB
10 SRR1270116 zot
# ... with 172 more rows
```

Merging tree with heatmap

2. Set row names to file name

```
file_name ctxA ctxB zot ace hylA rtxA ompU
                                           reference
                                                                           86
                                                                      100
                                                                           NΑ
  row names before
                                                                           94
                                                     98 NA 92 NA
  rownames (blast df)
                                                                           NΑ
                                                     95 84 95 NA
                                           SRR135602
                                                                           NΑ
               "3"
                                           SRR135604
  set row names
  rownames(blast df) <- blast df$file name
                                                 file_name ctxA ctxB zot ace hylA rtxA ompU
  row names after
> rownames(blast df)
                                                 SRR135602
   "reference" "SRR1270116" "SRR135539"
```

3. Select columns of interest

We can now get rid of the redundant file_name column

We are only interested in columns 2:8 with the genes

Double check before removing, using view() or:

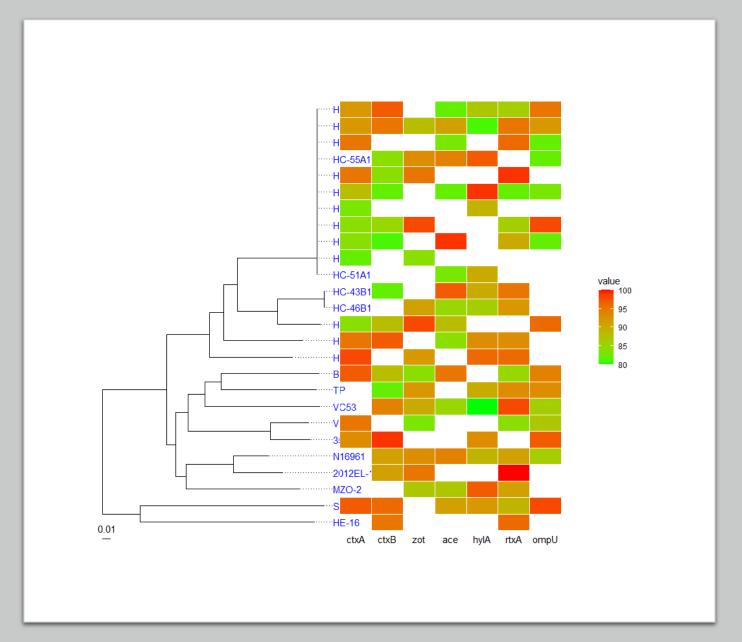
```
> blast_df %>% head(5) file_name ctxA ctxB zot ace hyla rtxA ompU
reference reference NA 91 93 94 89 91 86
SRR1270116 SRR1270116 NA 91 95 NA NA 100 NA
SRR135539 SRR135539 97 88 84 95 NA 85 94
SRR135542 SRR135542 98 NA 92 NA 96 96 NA
SRR135602 SRR135602 95 84 95 NA NA 99 NA
```

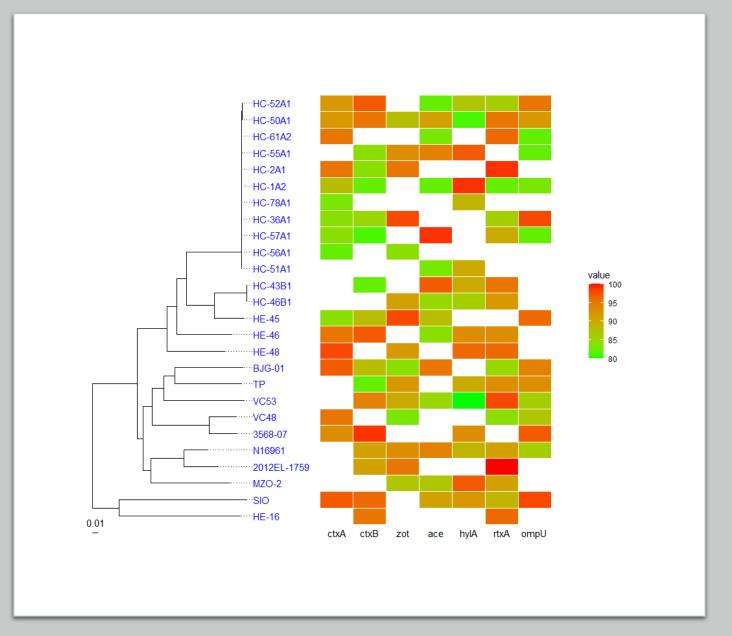
Select columns of interest (! means not)

```
> blast df <- select(blast df, !file name)</pre>
                                           ctxA ctxB zot ace hylA rtxA ompU
> blast df %>% head(5)
                                 reference
                                                                    86
                                                                91
                                                               100
                                                                    NΑ
                                                            NA
                                                                85
                                  SRR135539
                                            98
                                                NA 92 NA
                                                            96
                                                                96
                                                                    NΑ
                                  SRR135602
                                                                    NΑ
```

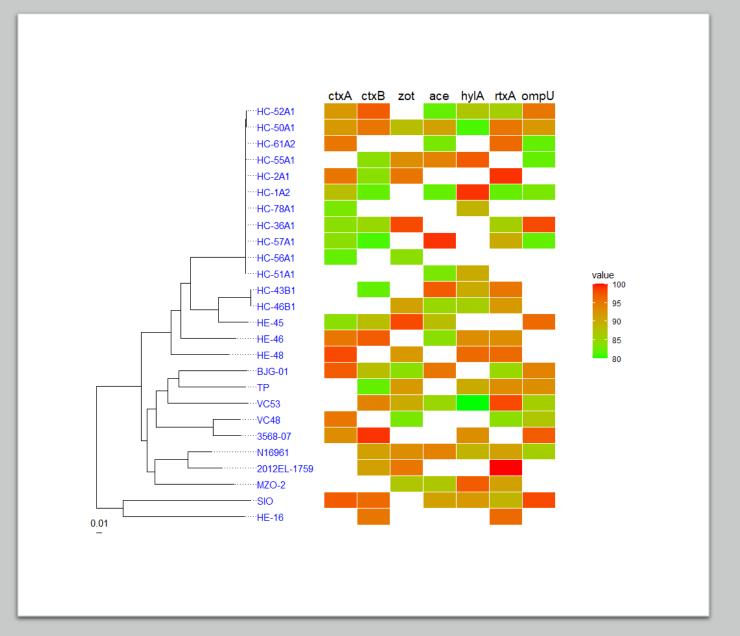
```
> gheatmap(gg_simple, # tree

blast_df) # heatmap
```





```
> gheatmap(gg_simple,
    blast_df,
    offset = 0.1,
    width = 1.5,
    font.size = 5,
    colnames_position = "top")
```



Changing heatmap colors: Option 1

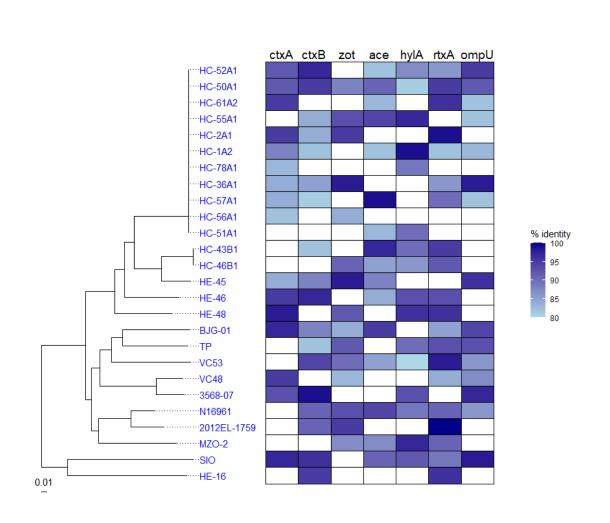
You can change the heatmap colors directly in gheatmap using arguments color, low and high

```
> gheatmap(gg simple,
        blast df,
         offset = 0.1,
         width = 1.5,
         font.size = 5,
         colnames position = "top",
         color = "black", # color of cell border
         legend title = "% identity", # title of legend
         low = "lightblue", # color of lowest value
         high = "darkblue") # color of highest value
```

The NA values will remain white

Option 1

```
> gheatmap(gg_simple,
    blast_df,
    offset = 0.1,
    width = 1.5,
    font.size = 5,
    colnames_position = "top",
    color = "black",
    legend_title = "% identity",
    low = "lightblue",
    high = "darkblue")
```



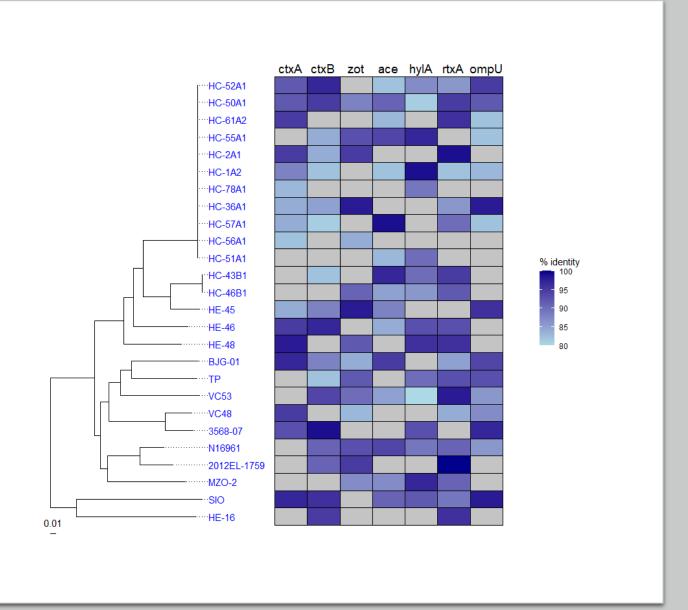
Changing heatmap colors: Option 2

You can change the heatmap colors using ggplot2:

A warning message will appear because we are overriding the previous color scale from gheatmap with the new scale from ggplot2

Option 2

```
> gheatmap(gg_simple,
  blast df,
   offset = 0.1,
   width = 1.5,
   font.size = 5,
   colnames position = "top",
   color = "black") +
scale_fill_gradient(name = "%
  identity",
   low = "lightblue",
   high = "darkblue",
   na.value = "grey77")
```



LEARNING OBJECTIVES

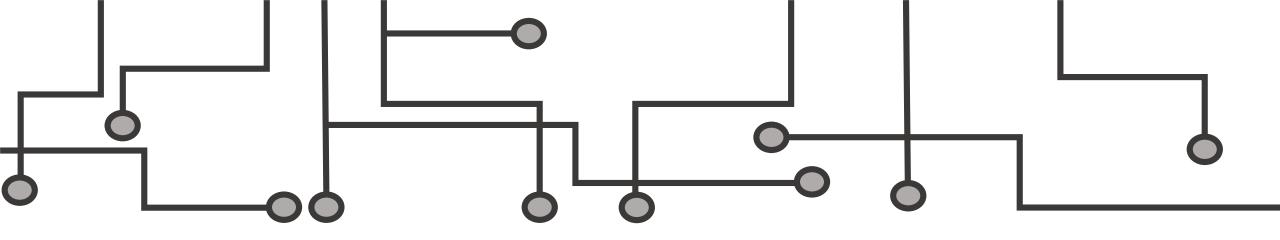
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Exporting the final tree

```
> tree heat <- gheatmap(gg simple,</pre>
           blast df,
           offset = 0.1,
           width = 1.5,
           font.size = 5,
           colnames position = "top",
           color = "black") +
      scale fill gradient(name = "% identity", # title of legend
                       low = "lightblue", high = "darkblue",
                      na.value = "grey77")
ggsave("output/tree heat.jpeg", tree heat, dpi = 300)
```

HELPFUL RESOURCES

- 1. Data integration, manipulation and visualization of phylogenetic
 - trees: https://yulab-smu.top/treedata-book/index.html
- 2. ggtree github: https://github.com/YuLab-SMU/ggtree
- 3. Enhanced annotation practice: http://www.randigriffin.com/2017/05/11/primate-phylogeny-ggtree.html
- 4. Colors and scales: https://ggplot2-book.org/scale-colour.html



THANK YOU FOR ATTENDING! The Q&A Session will now begin.

Please make sure to fill out the Exit Survey
We value your feedback!

More questions? Please email us at mmid.coding.workshop@gmail.com or post them to the workshop slack channel

