

1 ggtreeExtra: A universal package to visualize compact circular layers of phylogenetic tree

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1.1 Examples of mapping and visualizing associated data on circular layout tree.

It can help to find patterns and generate new hypotheses by integrating and visualizing external data to the phylogenetic tree. And no matter in which field, associated data type of phylogenetic tree can be roughly divided into continuous data and discrete data. For example, abundance of species, number of target genes and length of genome can be considered as continuous data. Endemic region information of virus, Taxonomy information of species and sampling location information can be considered as discrete data. They are also often multi-dimensional. To display them on the outer of phylogenetic tree more compactly, we developed *ggtreeExtra* to annotate multi-dimensional data to the outer ring of circular phylogenetic tree. It is a universal tool, based on the grammar of graphics(Wilkinson 2012). So user can easily map the associated dataset to aesthetic attributes (size, color, shape) of outer geometric objects (bar, point, boxplot) of circular phylogenetic tree using *ggtreeExtra*. Here, we present several examples to elucidate how the associated data can be mapped and displayed on the outer rings of circular phylogenetic trees using *ggtreeExtra*. More examples can be found on the *chapter10* of online book¹.

1.1.1 Displaying the continuous data to annotate circular phylogenetic trees.

It is efficient to annotate tree with circular layout, when there are multiple continuous datasets. The continuous dataset can be displayed using heat map, bar plot, box plot or dot plot etc. This example reproduce Fig.2 of (Morgan, Segata, and Huttenhower 2013). The data is provided by GraPhlAn (Asnicar et al. 2015), which contained the relative abundance of bacteria at different body sites. We use heat map to display the abundance of tip species, and the most abundance of specie at specific site also was visualized with bar plot. In addition, the commensal microbes and potential pathogens also were indicated with different shape of point. This example demonstrates the abilities of adding multiple layers (dot plot, heat map and bar plot) created with continuous data to a specific outer ring of circular phylogenetic tree (Fig.S1). The tree in *geom_fruit* can be fully annotated with multiple layers (high light, clade labels). The tree layer and other outer ring layers can be added step-by-step with *+*. More step-by-step instructions are also available at the vignette².

```
library(ggtreeExtra)
library(ggtree)
library(treeio)
library(tidytree)
library(ggstar)
library(ggplot2)
library(ggnewscale)
library(patchwork)

tree <- read.tree("../data/HMP_tree/hmptree.nwk")
# the abundance and types of microbes
dat1 <- read.csv("../data/HMP_tree/tippoint_attr.csv")
# the abundance of microbes at different body sites.
dat2 <- read.csv("../data/HMP_tree/ringheatmap_attr.csv")
# the abundance of microbes at the body sites of greatest prevalence.
dat3 <- read.csv("../data/HMP_tree/barplot_attr.csv")

# adjust the order
```

¹<http://yulab-smu.top/treedata-book>

²<http://bioconductor.org/packages/release/bioc/vignettes/ggtreeExtra/inst/doc/ggtreeExtra.html>

```

dat2$Sites <- factor(dat2$Sites, levels=c("Stool (prevalence)", "Cheek (prevalence)",
                                             "Plaque (prevalence)", "Tongue (prevalence)",
                                             "Nose (prevalence)", "Vagina (prevalence)",
                                             "Skin (prevalence)"))
dat3$Sites <- factor(dat3$Sites, levels=c("Stool (prevalence)", "Cheek (prevalence)",
                                             "Plaque (prevalence)", "Tongue (prevalence)",
                                             "Nose (prevalence)", "Vagina (prevalence)",
                                             "Skin (prevalence)"))

# extract the clade label information. Because some nodes of tree are annotated to genera,
# which can be displayed with high light using ggtree.
nodeids <- nodeid(tree, tree$node.label[nchar(tree$node.label)>4])
nodedf <- data.frame(node=nodeids)
nodelab <- gsub("[\\.\u0-9]", "", tree$node.label[nchar(tree$node.label)>4])
# The layers of clade and hightlight
poslist <- c(1.6, 1.4, 1.6, 0.8, 0.1, 0.25, 1.6, 1.6, 1.2, 0.4,
            1.2, 1.8, 0.3, 0.8, 0.4, 0.3, 0.4, 0.4, 0.4, 0.6,
            0.3, 0.4, 0.3)
labdf <- data.frame(node=nodeids, label=nodelab, pos=poslist)
# The circular layout tree.
p <- ggtree(
  tree,
  layout="fan",
  size=0.15,
  open.angle=5
) +
  geom_hilight(
    data=nodedf,
    mapping=aes(node=node),
    extendto=6.8,
    alpha=0.3,
    fill="grey",
    color="grey50",
    size=0.05
) +
  geom_cladelab(
    data=labdf,
    mapping=aes(node=node, label=label, offset.text=pos),
    barsize=NA,
    fontsize=0.7,
    angle="auto",
    hjust=0.5,
    horizontal=FALSE,
    fontface="italic"
) +
  ggtitle("A")

p1 <- p %<+% dat1 +
  geom_fruit(
    geom=geom_star,
    mapping=aes(fill=Phylum, starshape=Type, size=Size),
    position="identity",
    starstroke=0.05
) +
  scale_fill_manual(
    values=c("#FFC125", "#87CEFA", "#7B68EE", "#808080", "#800080",
             "#9ACD32", "#D15FEE", "#FFC0CB", "#EE6A50", "#8DEEEE",
             "#006400", "#800000", "#B0171F", "#191970"),
    guide=guide_legend(keywidth = 0.5, keyheight = 0.5, order=1,

```

```

    override.aes=list(starshape=15)),
    na.translate=FALSE
) +
scale_starshape_manual(
  values=c(15, 1),
  guide=guide_legend(keywidth = 0.5, keyheight = 0.5, order=2),
  na.translate=FALSE
) +
scale_size_continuous(
  range = c(0.5, 1.5),
  guide = guide_legend(keywidth = 0.5, keyheight = 0.5, order=3,
  override.aes=list(starshape=15))
) +
new_scale_fill() +
ggtitle("B") +
theme(legend.position="none")

p2 <- p1 +
geom_fruit(
  data=dat2,
  geom=geom_tile,
  mapping=aes(y=ID, x=Sites, alpha=Abundance, fill=Sites),
  color = "grey50",
  offset = 0.04,
  size = 0.02
) +
scale_alpha_continuous(
  range=c(0, 1),
  guide=guide_legend(keywidth = 0.3, keyheight = 0.3, order=5)
) +
scale_fill_manual(
  values=c("#0000FF", "#FFA500", "#FF0000", "#800000",
  "#006400", "#800080", "#696969"),
  guide=guide_legend(keywidth = 0.3, keyheight = 0.3, order=4)
) +
ggtitle("C") +
theme(legend.position="none")

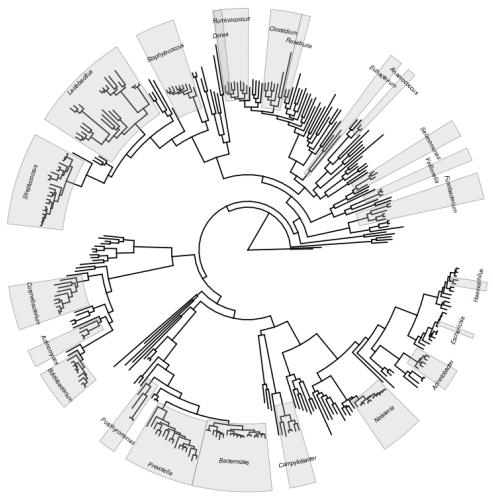
p3 <- p2 +
geom_fruit(
  data=dat3,
  geom=geom_bar,
  mapping=aes(y=ID, x=HigherAbundance, fill=Sites),
  pwidth=0.38,
  orientation="y",
  stat="identity"
) +
geom_treescale(fontsize=1.2, linesize=0.3, x=4.9, y=0.1) +
ggtitle("D") +
theme(legend.position="none")

p4 <- (p + p1 + plot_layout(width=c(3.4,4)))/ (p2 + p3 + plot_layout(width=c(3.4,4))) +
plot_layout(heights=c(3,4))

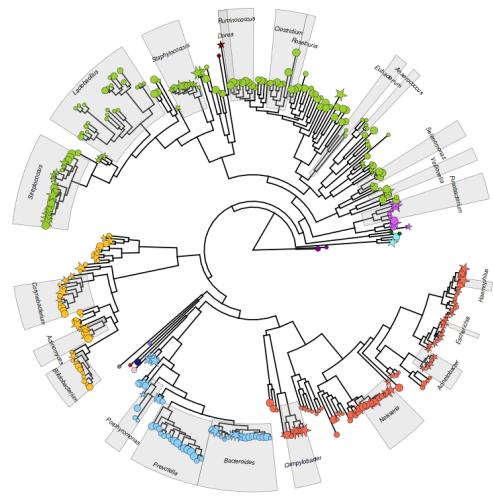
p4

```

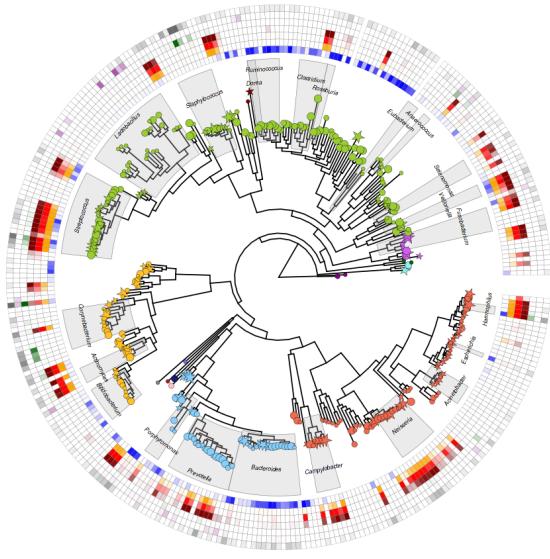
A



B



C



D

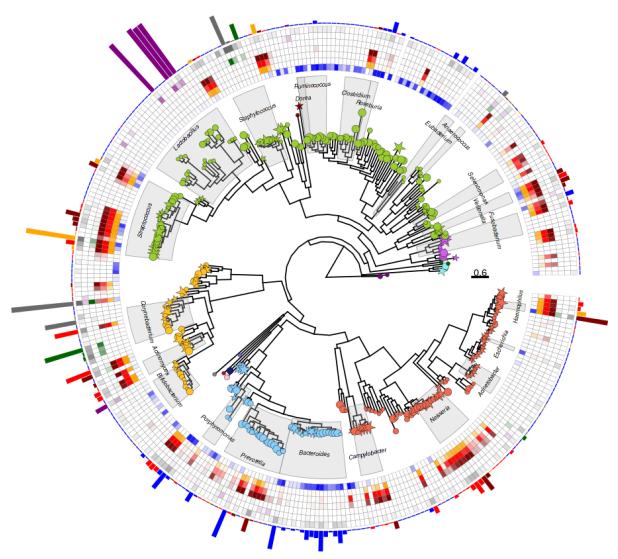


Fig. S1: The abundance of microbes at different sites of human. The shape of tip labels indicated the commensal microbes or potential pathogens. The transparency of heat map indicates the abundance of microbes, and colors of heat map indicate the different sites of human. The bar plot indicates the relative abundance at body site of the most abundance. Fig.S1.A contains the tree layer, high light layer and clade label layer; Fig.S1.B is a tip point layer added on the basis of Fig.S1.A; Fig.S1.C is a heatmap layer added on the basis of Fig.S1.B; Fig.S1.D is a bar chart layer added on the basis of Fig.S1.C. This example shows the abilities of annotate phylogenetic tree through layer overlay using *ggtree* and *ggtreeExtra*.

1.1.2 Visualizing discrete datasets to annotate circular phylogenetic trees.

The discrete datasets can be displayed using heat map. When the external datasets have a column of *node* or first column of tip labels. We can integrated the data to tree data by `%<+%` operator of *ggtree* (Yu et al. 2018). The *geom_fruit* can extract the tree data integrated automatically, and the attributes of datasets can mapped using related attributes of geometry layers. The *y* of *aes* can be ignored, it will be added automatically. We will reproduce Figure 3.3 of (Smith

and Wrighton 2019) to show how to map discrete datasets. The external ring heat maps represent the different types in corresponding categories (Fig. S2). And the source data is available at this repository³.

```

library(ggtreeExtra)
library(ggtree)
library(treeio)
library(ggplot2)
library(ggnewscale)
tree <- read.raxml("../data/Methanotroph/Methanotroph_rpS3_Modified_Alignment_RAxML")
# Root the tree to the archaea sequences
tree@phylo <- root(tree@phylo, node=1402, edgelabel=TRUE)
df <- read.csv("../data/Methanotroph/metadata.csv")
# reset the levels of columns to reproduce the order of original figure.
df$Specific.Ecosystem <- factor(df$Specific.Ecosystem,
                                 levels=c("Agriculture", "Alkaline/Hypersaline",
                                         "Contaminated/Wastewater", "Endosymbiont",
                                         "Freshwater", "Forest", "Geothermal",
                                         "Marine", "Natural Seep", "Peat",
                                         "Permafrost", "Wetland", "Unknown"))
df$MetaType <- factor(df$MetaType,
                       levels=c("Metatranscriptome", "Metagenome",
                               "Single-amplified genome", "Fosmid", NA))
df$Treatment <- factor(df$Treatment, levels=c("Native", "Enrichment", "Isolate", "Unknown"))
p <- ggtree(tree, layout="fan", open.angle=30)
p <- rotate_tree(p, 90)
p <- p + geom_treescaler(x=0.2, y=727*6/11, width=1, offset=20) +
  geom_nodepoint(aes(subset=as.numeric(bootstrap)>=70),
                 shape=21, fill="grey", size=1)

# we can use %<+% to integrate the external datasets to tree structure.
# and the y can not be specified.
p <- p %<+% df +
  geom_fruit(
    geom=geom_tile,
    mapping=aes(fill=Specific.Ecosystem),
    offset=0.13,
    width=0.35,
    axis.params=list(axis="x", text="Ecosystem",
                     text.angle=0, hjust=0,
                     text.size=3, family="Times",
                     fontface="bold"))
  ) +
  scale_fill_manual(
    values=c("green3", "turquoise", "maroon", "orchid",
            "deepskyblue", "forestgreen", "salmon", "cadetblue3",
            "slategray4", "yellowgreen", "gray90", "chocolate2",
            "yellow"),
    guide=guide_legend(title="Ecosystem", keywidth=0.5, keyheight=0.5, order=4),
    na.translate=FALSE
  ) +
  new_scale_fill() +
  geom_fruit(
    geom=geom_tile,
    mapping=aes(fill=MetaType),
    offset=0.13,
    width=0.35,
    axis.params=list(
      axis="x",

```

³https://github.com/TheWrightonLab/Methanotroph_rpS3Analyses_SmithWrighton2018

```

        text="Sequencing Type",
        text.angle=0,
        hjust=0,
        text.size=3,
        family="Times",
        fontface="bold"
    )
)
+
scale_fill_manual(
    values=c("red", "black", "dodgerblue", "gray50"),
    guide=guide_legend(title="Sequencing Type",
        keywidth=0.5, keyheight=0.5, order=3),
    na.translate=FALSE
)
+
new_scale_fill()+
geom_fruit(
    geom=geom_tile,
    mapping=aes(fill=Treatment),
    offset=0.13,
    width=0.35,
    axis.params=list(
        axis="x",
        text="Sample Treatment",
        text.angle=0,
        hjust=0,
        text.size=3,
        family="Times",
        fontface="bold"
    )
)
+
scale_fill_manual(
    values=c("red","gray50", "black", "yellow"),
    guide=guide_legend(title="Sample Treatment", keywidth=0.5,
        keyheight=0.5, order=2),
    na.translate=FALSE
)
+
theme(legend.background=element_rect(fill=NA), # the background of legend.
    legend.title=element_text(size=9, family="Times", face="bold"),
    legend.text=element_text(size=7, family="Times"), # the text size of legend.
    legend.spacing.y = unit(0.02, "cm"),
    legend.margin=margin(0.1, 0.9, 0.1,-0.9, unit="cm"), # t, r, b, l, cm
    legend.box.margin=margin(0.1, 0.9, 0.1, -0.9, unit="cm"),
    plot.margin = unit(c(-1.2, -1.2, -1.2, 0.1),"cm"))

# To build the external annotation of clades.
p <- p + geom_cladelabel(node=793, angle="auto", label="Gammaproteobacteria", horizontal=FALSE,
    offset=1.4, align=T, fontsize = 3, hjust=0.5, offset.text=0.12, barsize=1,
    family="Times", fontface="bold", color="gray30")+
geom_cladelabel(node=791, angle="auto", label="Alphaproteobacteria", horizontal=FALSE,
    offset=1.4, align=T, fontsize = 3, hjust=0.5, offset.text=0.12, barsize=1,
    family="Times", fontface="bold", color="black")+
geom_cladelabel(node=1384, label="Ca.Methylomirabilis", horizontal=TRUE,
    offset=1.4, align=T, fontsize = 3, angle="auto", barsize=1,
    family="Times", fontface="bold", color="gray30")+
geom_cladelabel(node=1394, label="Methylacidiphilae", angle="auto", horizontal=TRUE,
    offset=1.4, align=T, fontsize = 3, barsize=1,
    family="Times", fontface="bold", color="black")+
geom_cladelabel(node=1440, label="ANME-1", angle="auto", horizontal=TRUE,
    offset=1.4, align=T, fontsize = 3, barsize=1,

```

```

family="Times", fontface="bold", color="gray30")+
geom_cladelabel(node=1405, label="ANME-2", angle="auto", horizontal=TRUE,
offset=1.4, align=T, fontsize = 3, barsize=1,
family="Times", fontface="bold", color="black")

```

p

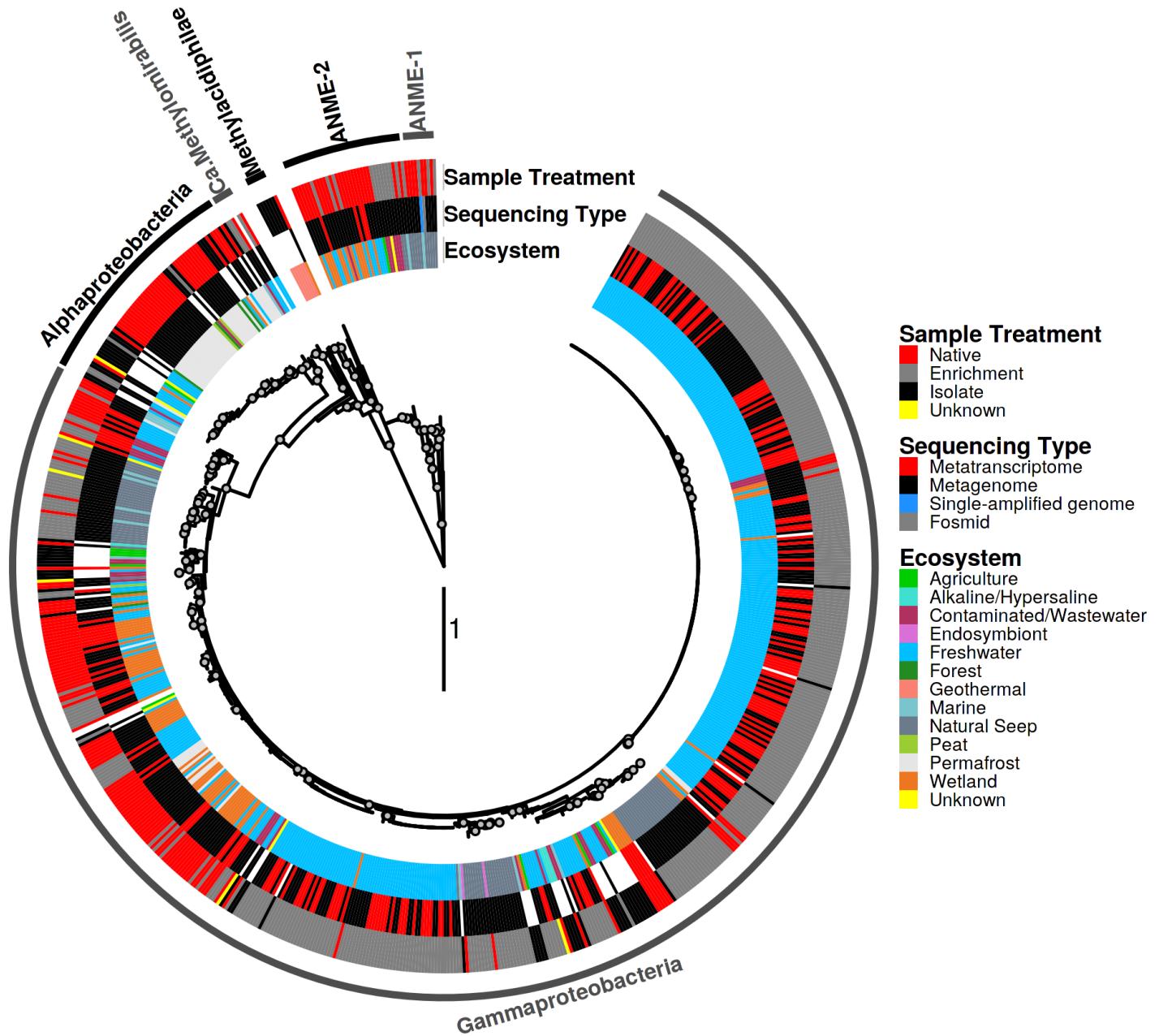


Fig. S2: Phylogeny of methanotroph ribosomal protein S39 (rpS3) genes from Figure 3.3 (Smith and Wrighton 2019). The external ring of circular tree is built with discrete datasets, which were added to tree structure in advance

1.1.3 Mapping the discrete data and continuous data to circular phylogenetic trees.

In this section, we show that feature of mapping the discrete datasets and continuous datasets to the circular tree. The discrete datasets also can be usually visualized using dot plot and heat map. We use two examples to show how to map the datasets. The Fig 1 of (Segata et al. 2013) and Fig 2 of (Asnicar et al. 2015) will be reproduced in the section.

1.1.3.1 The first example in this section This example will show that discrete dataset is visualized in external ring using dot plot, which is from geometric layer of *ggstar* (Xu 2020). The continuous data is also displayed with bar plot (Fig. S3).

```

library(ggtreeExtra)
library(ggtree)
library(treeio)
library(tidytree)
library(ggstar)
library(ggplot2)
library(ggnewscale)

tree <- read.tree("../data/PhyloPhlAn/ppal_tol.nwk")
# The attributes of tip point
da1 <- read.csv("../data/PhyloPhlAn/tipoint_attr.csv")
# The attributes of bar plot
da2 <- read.csv("../data/PhyloPhlAn/barplot_attr.csv")
# The attributes of triangle point
da3 <- read.csv("../data/PhyloPhlAn/ringpoint_attr.csv")

# to reproduce, reorder column Type
da3$Type <- factor(da3$Type, levels=c("Mislabelled", "Insertions",
                                         "Corrections", "Refinements"))

# extract the node label to annotate clade
nodelab <- tree$node.label[nchar(tree$node.label)>0]
nodeids <- nodeid(tree, nodelab)
# the position of high light extends to
extos <- c(rep(5.66, 12), 7.1, rep(5.66, 30))
nodedf <- data.frame(node=nodeids, extos=extos)

# the position of clade label
textex <- c(0.8, 0.8, 1.0, 0.6, 0, 0.8, 0.8, 0.2, -0.05, 0.5,
           0.1, 0.1, -0.1, 0.6, 0.4, 0, 1.2, 0.8, 1.0, 0.6,
           0.6, 1.0, 0.8, 0.2, -0.1, 1.2, 1, 1, 0.6, 0.6,
           0.4, 0.5, 1.2, 0.6, 1.4, 1.3, 0, -0.1, 0.8, 0.6,
           0.8, 1.4, 2.4)
cladelabels <- mapply(function(x, y, z){geom_cladelabel(node=x, label=y, barsize=NA, extend=0,
                                                       offset.text=z, fontsize=1.2, angle="auto",
                                                       hjust=0.5, horizontal=FALSE, fontface="italic")},
                       nodeids, nodelab, textex, SIMPLIFY=FALSE)

# To reproduce the color of original figures, we generated the same colors,
# user can custom their colors.
phylacolor <- c(Proteobacteria="#9ACD32", Firmicutes="#EE6A50",
                 Actinobacteria="#87CEFA", Bacteroidetes="#FFC125",
                 Euryarchaeota="#D15FEE", Tenericutes="#8DEEEE",
                 Cyanobacteria="#800000", Spirochaetes="#006400",
                 Chlamydiae="#800080", Crenarchaeota="#808080",
                 Fusobacteria="#FFC0CB", Thermi="#B0171F", Other="Black",
                 Chloroflexi="#191970", Thermotogae="#7B68EE",
                 Aquificae="#00CD00", Synergistetes="#8B4513",
                 Chlorobi="#BC8F8F", Verrucomicrobia="#303030",
                 Planctomycetes="#8E8E38", Acidobacteria="#CDCDC1")

tipcolors <- c("#9ACD32", "#EE6A50", "#87CEFA", "#FFC125", "#D15FEE",
               "#8DEEEE", "#800000", "#006400", "#800080", "#808080",
               "#FFC0CB", "#B0171F", "#191970", "#7B68EE", "#00CD00",
               "#8B4513", "#BC8F8F", "#303030", "#8E8E38", "#CDCDC1")

```

```

names(tipcolors) <- c("Proteobacteria", "Firmicutes", "Actinobacteria",
                      "Bacteroidetes", "Euryarchaeota", "Tenericutes",
                      "Cyanobacteria", "Spirochaetes", "Chlamydiae",
                      "Crenarchaeota", "Fusobacteria", "Thermi",
                      "Chloroflexi", "Thermotogae", "Aquificae",
                      "Synergistetes", "Chlorobi", "Verrucomicrobia",
                      "Planctomycetes", "Acidobacteria")

# We built a circular layout tree with open a small (6) angle.
p <- ggtree(
  tree,
  layout="fan",
  open.angle=6,
  size=0.1
) +
  geom_hilight(
    data=nodedef,
    mapping=aes(node=node, extendto=extos),
    alpha=0.3,
    fill="grey",
    color="grey50",
    size=0.05)

p <- p %<+% da1 +
  geom_tippoint(
    mapping=aes(fill=Phylum),
    size=1.2,
    shape=21,
    stroke=0.05,
    show.legend=FALSE
) +
  scale_fill_manual(values=tipcolors) +
  new_scale_fill() +
  geom_fruit(
    data=da2,
    geom=geom_bar,
    mapping=aes(x=Abundance, y=ID, fill=Phyla),
    offset=-0.2,
    pwidth=0.1,
    stat='identity',
    orientation="y"
) +
  scale_fill_manual(values=phylacolor,
                    guide=guide_legend(keywidth=0.6,
                                        keyheight=0.6,
                                        order=1, ncol=1)) +
  cladelabels +
  new_scale_fill() +
  geom_fruit(
    data=da3,
    geom=geom_star,
    mapping=aes(x=Pos, y=ID, fill=Type, size=Pos),
    starshape=26,
    alpha=0.8,
    starstroke=0,
    offset=0.024,
    pwidth=0.008
) +

```

```

scale_fill_manual(values=c("blue", "black", "green", "red"),
                  guide=guide_legend(keywidth=1, keyheight=1, order=2,
                                     override.aes=list(alpha=1, size=2)),
                  na.translate=FALSE)+
scale_size_continuous(range=c(1.4,2.2), guide="none")+
geom_treescale(fontsize=1.2, linesize=0.3) +
theme(legend.position=c(0.94, 0.5),
      legend.title=element_text(size=7),
      legend.background=element_rect(fill=NA),
      legend.text=element_text(size=6),
      legend.spacing.y = unit(0.02, "cm"))

```

p

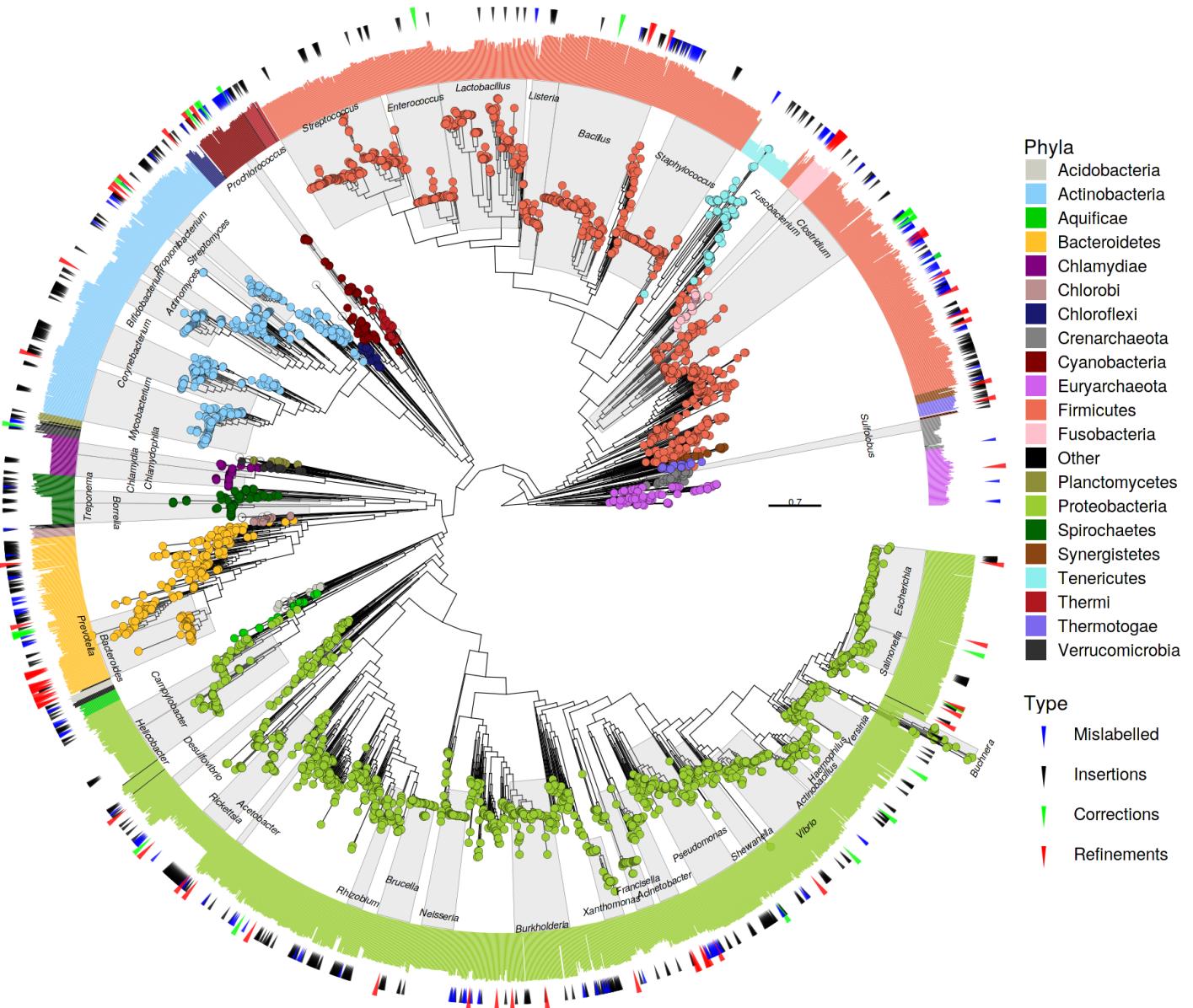


Fig. S3: The bacterial and archaeal (3737 genomes) phylogenetic tree built with 400 conserved proteins. The tip points are built using discrete data, the different colors represent the different phyla. The external bar is created using continuous data, the length represents the fraction of the 400 proteins contained in each genome. The external triangle is also created using discrete data, the different colors represent the different types of genomes (mislabelled and confidently replaced see Fig 1 of (Segata et al. 2013)).

1.1.3.2 The second example in this section This example reproduces Fig 2 of (Asnicar et al. 2015). The discrete data is displayed using heat map, and other continuous datasets are visualized with heat map and bar plot. These show that no restriction of data types or how the data should be plotted in *geom_fruit* (Fig. S4).

```

library(ggtreeExtra)
library(ggtree)
library(treeio)
library(tidytree)
library(ggstar)
library(ggplot2)
library(ggnewscale)

tree <- read.tree("../data/kegg/kegg.nwk")
# The attributes of tip point
dt1 <- read.csv("../data/kegg/tippoint_attr.csv")
# The attributes of first ring
dt2 <- read.csv("../data/kegg/firstring_attr.csv")
# The attributes of second ring
dt3 <- read.csv("../data/kegg/secondring_attr.csv")
# The attrihutes of bar plot
dt4 <- read.csv("../data/kegg/barplot_attr.csv")

#reorder the Phyla column
dt1$Phyla <- factor(dt1$Phyla, levels=c("Actinobacteria", "Aquificae", "Bacteroidetes",
                                             "Chlamydiae", "Chlorobi", "Chloroflexi", "Crenarchaeota",
                                             "Cyanobacteria", "Euryarchaeota", "Firmicutes", "Proteobacteria",
                                             "Spirochaetes", "Tenericutes", "Thermi", "Thermotogae", "Other"))

# reorder the Type2 column
dt3$Type2 <- factor(dt3$Type2, levels=c("FA synth init", "FA synth elong",
                                           "acyl-CoA synth", "beta-Oxidation",
                                           "Ketone biosynth"))

dt4$Phyla <- factor(dt4$Phyla, levels=c("Actinobacteria", "Aquificae", "Bacteroidetes",
                                             "Chlamydiae", "Chlorobi", "Chloroflexi", "Crenarchaeota",
                                             "Cyanobacteria", "Euryarchaeota", "Firmicutes", "Proteobacteria",
                                             "Spirochaetes", "Tenericutes", "Thermi", "Thermotogae", "Other"))

# extract node label for the clade layers
nodelab <- tree$node.label[nchar(tree$node.label)>0]
nodeids <- nodeid(tree, nodelab)

# the position of clade label
textex <- c(1.0, 0.4, 0.2, 1.4, 1.4, 0.4, 1.4, 1.4, 0.4, 0.4,
           0.8, 1, 0.6, 0.6, 0.4, 0.3, 0, 0.4, 0.1, 0.25,
           0.2, 0.3, 0.8, 0.8, 0.8, 0.6, 2.4)

# clade layers
cladelabels <- mapply(function(x, y, z){geom_cladelabel(node=x, label=y, barsize=NA, extend=0.3,
                                                          offset.text=z, fontsize=1.2, angle="auto",
                                                          hjust=0.5, horizontal=FALSE, fontface="italic")}, 
                       nodeids, nodelab, textex, SIMPLIFY=FALSE)

# high light layers
fills <- c("#808080", "#808080", "#808080", "#808080", "#808080",
           "#191970", "#87CEFA", "#FFC125", "#B0171F", "#B0171F",
           "#B0171F", "#B0171F", "#B0171F", "#B0171F", "#B0171F",
           "#B0171F", "#B0171F", "#B0171F", "#B0171F", "#B0171F",
           "#B0171F", "#B0171F", "#9ACD32", "#9ACD32", "#9ACD32",
           "#006400", "#800000")

highlights <- mapply(function(x, y){geom_hilight(node=x, extendto=5.8, alpha=0.3,

```

```

            fill=y, color=y, size=0.05)},
        nodeids, fills, SIMPLIFY=FALSE)

# to reproduce the original figures, we use the same colors.
# uses can custom set it.
colors <- c("#9ACD32", "#EE6A50", "#87CEFA", "#FFC125", "#D15FEE", "#8DEEEE", "#800000",
           "#006400", "#800080", "#808080", "#B0171F", "#B0171F", "#191970", "#7B68EE",
           "#00CD00", "Black")

p <- ggtree(tree, layout="circular", size=0.1) + highlights

p <- p +
  geom_fruit(
    data=dt1,
    geom=geom_point,
    mapping=aes(y=ID, fill=Phyla),
    shape=21,
    size=1.2,
    stroke=0.05,
    position="identity",
    show.legend=FALSE
  ) +
  scale_fill_manual(values=colors) +
  cladelabels +
  new_scale_fill() +
  geom_fruit(
    data=dt2,
    geom=geom_tile,
    mapping=aes(y=ID, x=ring, fill=Type1),
    offset=-0.02,
    pwidth=0.14,
    addbrink=TRUE
  ) +
  scale_fill_manual(
    name="ATP synthesis",
    values=c("#339933", "#dfac03"),
    guide=guide_legend(keywidth=0.5, keyheight=0.5, order=1)
  ) +
  new_scale_fill() +
  geom_fruit(
    data=dt3,
    geom=geom_tile,
    mapping=aes(y=ID, alpha=Abundance, x=Type2, fill=Type2),
    offset=0.001,
    pwidth=0.18
  ) +
  scale_fill_manual(
    name="Fatty Acid metabolism",
    values=c("#b22222", "#005500", "#0000be", "#9f1f9f", "#793a07"),
    guide=guide_legend(keywidth=0.5, keyheight=0.5, order=2)
  ) +
  scale_alpha_continuous(range=c(0, 0.4),
                        guide=guide_legend(keywidth=0.5, keyheight=0.5, order=3)) +
  new_scale_fill() +
  geom_fruit(data=dt4,
             geom=geom_bar,
             mapping=aes(y=ID, x=Length, fill=Phyla),
             stat="identity",

```

```
  orientation="y",
  pwidth=0.3,
  position=position_dodgex()) +
scale_fill_manual(values=colors,
                  guide=guide_legend(keywidth=0.5, keyheight=0.5, order=4)) +
geom_treescale(fontsize=1.2, linesize=0.3) +
theme(legend.position=c(0.95, 0.5),
      legend.background=element_rect(fill=NA),
      legend.title=element_text(size=7),
      legend.text=element_text(size=6),
      legend.spacing.y = unit(0.02, "cm"))
```

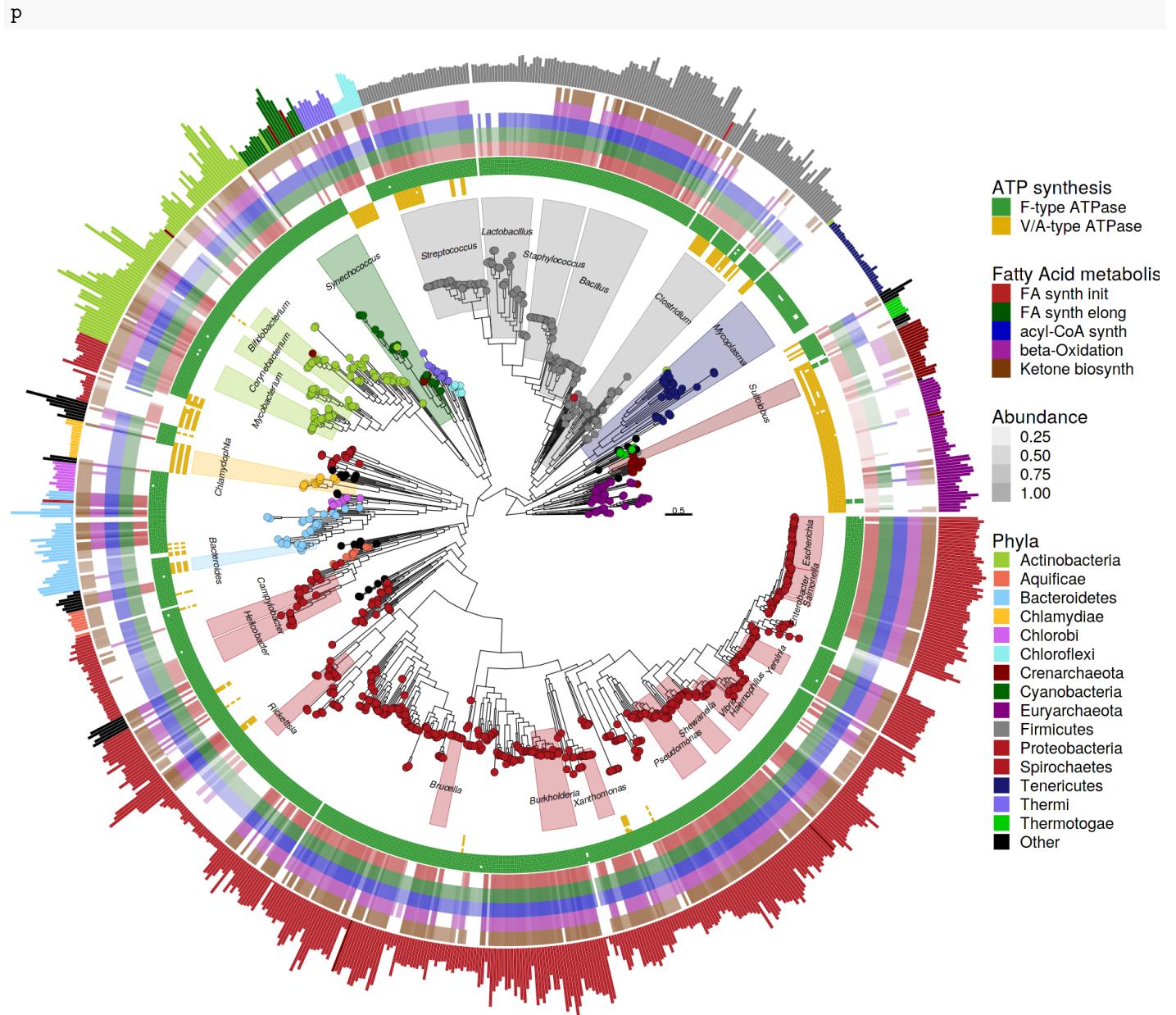


Fig. S4: The phylogenetic tree built using in (Segata et al. 2013) using 3737 microbial genomes. The first and second ring heat map were built with discrete data, it represents the presence or absence of each module, the other heat map rings were created with continuous data, the transparency of color represents the capability of fatty acid metabolism, the different colors represent the types of fatty acid metabolism. The length of bar represents the genome length of corresponding microbes.

1.1.4 Obtain more power with some packages

The *geom_fruit* not only work with some *geom* function defined in *ggplot2* (Wickham 2016), but also can work with *geom* function defined in ggplot2-based packages. This ability will make *ggtreeExtra* more power to present dataset. We use an example from Fig.1 of (Song et al. 2020) to show *geom_fruit* can work with *geom_phylopic* defined in *ggimage* (Yu 2020). The attributes (such as color, size) of subplot also can be mapped to variables of external dataset. This is often not available in the existing tools. The source data of the example is available at this repository⁴.

```

library(ggtree)
library(treeio)
library(ggplot2)
library(ggtreeExtra)
library(tidytree)
library(ggnewscale)
library(ggimage)

tr <- read.tree("../data/VertebrateGutMicrobiomes/annotated_host_tree.tre")
corda <- read.csv("../data/VertebrateGutMicrobiomes/mantel.jaccard.pearson.csv")
corda$r <- abs(corda$r)
barda <- read.csv("../data/VertebrateGutMicrobiomes/data_diet_bar.csv", check.names=F)
barda <- reshape2::melt(barda, id.vars="ID", variable.name="Diet", value.name="mete")
barda$Diet <- factor(barda$Diet, levels=c("Fruit", "Invertebrates",
                                             "Nectar", "Plants", "Scavenging",
                                             "Seeds", "Meat (Ectotherms)",
                                             "Meat (Endotherms)",
                                             "Meat (Fish)", "Meat (Unknown)"))

cladeda <- read.csv("../data/VertebrateGutMicrobiomes/data_clade_class.csv", check.names=F)
cladeda$id <- nodeid(tr, cladeda$id)
cladeda$class <- factor(cladeda$class, levels=c("Amphibia", "Chelonia", "Lepidosauria",
                                                 "Crocodylomorpha", "Aves", "Mammalia"))

flightda <- read.csv("../data/VertebrateGutMicrobiomes/data_flight_bar.csv")

phylopicda <- read.csv("../data/VertebrateGutMicrobiomes/data_phylopic_uid.csv")
phylopicda$class <- factor(phylopicda$class, levels=c("Amphibia", "Chelonia", "Lepidosauria",
                                                       "Aves", "Mammalia"))

p <- ggtree(tr, layout="fan", open.angle=15)

p <- p %<+% corda
p$data$width <- ifelse(is.na(p$data$r), 0.1, 0.6)
r <- NULL
p <- p +
  aes(color=r, size=I(width)) +
  scale_colour_viridis_c(
    name="Mantel Correlation",
    option="C",
    guide=guide_colorbar(
      barheight=0.6,
      order=4,
      title.position="top",
      label.position="bottom",
      direction="horizontal"
    )
  )
p1 <- p +

```

⁴https://github.com/tanaes/tetrapod_microbiome_analysis

```

geom_fruit(
  data=barda,
  geom=geom_bar,
  mapping=aes(x=mete, y=ID, fill=Diet),
  orientation="y",
  stat="identity",
  colour=NA,
  pwidth=0.25,
  offset=0.008
) +
scale_fill_manual(
  values=c("#a6cee3", "#cab2d6",
    "#1f78b4", "#33a02c",
    "#6a3d9a", "#b2df8a",
    "#fb9a99", "#e31a1c",
    "#ff7f00", "#fdbf6f"),
  guide=guide_legend(keywidth=0.5, keyheight=0.5, order=1)
)

p2 <- p1 +
  new_scale_colour() +
  geom_cladelab(
    data=cladeda,
    mapping=aes(node=id, label=class, colour=class),
    textcolour=NA,
    barsize=4,
    extend=0.2,
    offset=100) +
  scale_colour_manual(
    name="Host Class",
    values=c("#b2df8a", "#33a02c", "#fb9a99",
      "#e31a1c", "#EACB47", "#6a3d9a"),
    guide=guide_legend(
      keywidth=0.5,
      keyheight=0.5,
      order=2,
      override.aes=list(size=3, alpha=1)
    )
  )

p3 <- p2 +
  new_scale_fill() +
  geom_fruit(
    data=flightda,
    geom=geom_tile,
    mapping=aes(y=ID, fill=flight),
    size=0,
    width=14,
    offset=0.11,
    pwidth=0.4,
  ) +
  scale_fill_manual(
    name="Flight Status",
    values=c("black", "white"),
    guide=guide_legend(keywidth=0.5, keyheight=0.5, order=3,
      override.aes=list(color="black", size=0.3)))
p4 <- p3 +
  new_scale_colour() +

```

```

geom_fruit(
  data=phylopicda,
  geom=geom_phylopic,
  mapping=aes(y=taxa, image=uid, color=class),
  size=0.035,
  offset=0.16,
  alpha=0.8,
  position=position_identityx()
) +
scale_colour_manual(
  values=c("#b2df8a", "#33a02c", "#fb9a99",
           "#EACB47", "#6a3d9a"),
  guide="none"
) +
theme(
  legend.background=element_rect(fill=NA),
  legend.title=element_text(size=9),
  legend.text=element_text(size=6.6),
  legend.spacing.y = unit(0.02, "cm")
)

```

p4

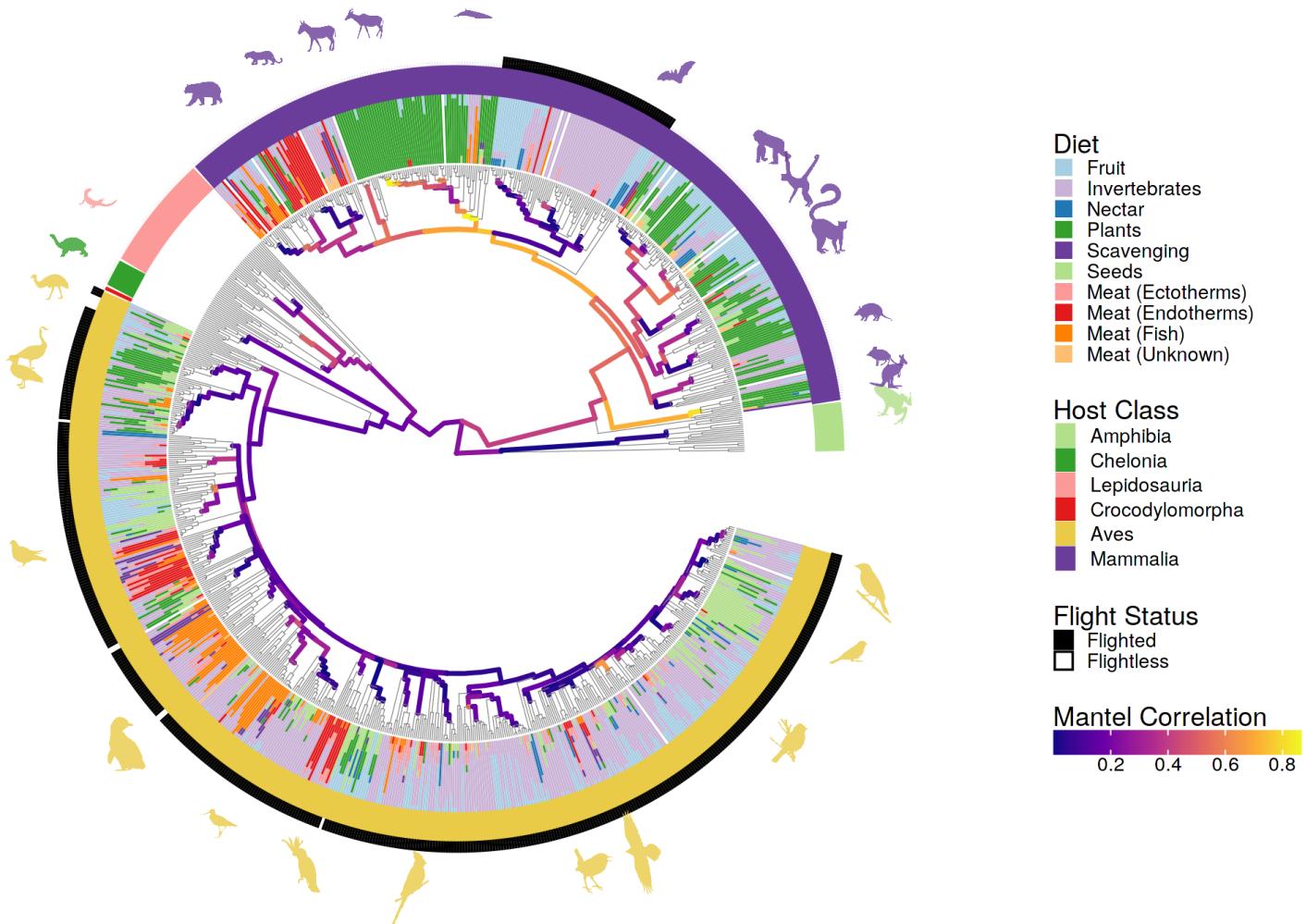


Fig. S5: Host tree was obtained from TimeTree (Kumar et al. 2017). The branch colour represents the Mantel Pearson correlation. The stack bar chart represents the diet composition of host. The inner ring represents the host taxonomic class, and some representative species are also displayed in the outermost circle. The middle ring represents flight status.

1.1.5 Obtain new patterns by annotating tree with multi-dimensional data

The `geom_fruit` is a general function to link `ggtree` (Yu et al. 2017) and `ggplot2` (Wickham 2016), so it can be applied in many fields. It can give user more convenience and reproducibility to visualize and annotate phylogenetic trees with associated data for better exploring phylogenetic patterns behind multi-dimensional data. For example, Fig.1b and Fig.2 of (Helfrich et al. 2018) show the directional interactions and the biosynthetic potential of isolates from *Arabidopsis* leaf microbiome in phylogenetic tree. However, because GraPhlAn (Asnicar et al. 2015) did not display the stack bar plot, some phylogenetic patterns behind multi-dimensional data were not be found easily. Here, we can use `ggtree` and `ggtreeExtra` to show the total information of Fig.1b and Fig.2 of (Helfrich et al. 2018). We found some strains from Firmicutes and from Grammaproteobacteria have more inhibitor interactions. However, many strains from Alphaproteobacteria and Betaproteobacteria prefer the interaction of sensitivity. In addition, These strains that prefer the interactions of sensitivity might be have more BGCs from ribosomally synthesized and post-translationally modified peptide (*RiPP*).

```

library(ggtree)
library(ggtreeExtra)
library(ggplot2)
library(MicrobiotaProcess)
library(ggstar)
library(ggnewscale)
library(grid)

alltax <- read.csv("../data/Arabidopsis_leaf_microbiome/all_stain_taxonomy.csv")
linktab <- read.csv("../data/Arabidopsis_leaf_microbiome/Interaction_link_tab.csv")
weighttab <- read.csv("../data/Arabidopsis_leaf_microbiome/Interaction_weight.csv")
tippoint <- read.csv("../data/Arabidopsis_leaf_microbiome/stain_tippoint.csv")
BGCsda <- read.csv("../data/Arabidopsis_leaf_microbiome/BGCs_heatmap.csv")

tippoint$Taxa <- factor(tippoint$Taxa,
                         levels=c("Actinobacteria",
                                 "Bacteroidetes",
                                 "Firmicutes",
                                 "Deinococcus-Thermus",
                                 "Alphaproteobacteria",
                                 "Betaproteobacteria",
                                 "Gammaproteobacteria"
                         )
)
tippoint$names <- gsub("s_Leaf","",tippoint$Isolation)

BGCsda$BGCs <- factor(BGCsda$BGCs,
                       levels=c("modular.PKS",
                               "modular.PKS.NRPS.hybrid",
                               "non_modular.PKS", "NRPS",
                               "RiPP",
                               "Quorum.sensing",
                               "terpene",
                               "other"
                       )
)
BGCsda$Count <- log10(BGCsda$Count+1)
BGCsda$Count <- ifelse(BGCsda$Count==0, NA, BGCsda$Count)

trda <- convert_to_treedata(alltax)
p <- ggtree(
  trda,
  layout="inward_circular",
  size=0.2,
  xlim=c(18,NA)
)

```

```

p <- p %<+% tippoint

p1 <- p +
  geom_tippoint(
    mapping=aes(
      color=Taxa,
      shape=Level
    ),
    size=1,
    alpha=0.8
  ) +
  scale_color_manual(values=c("#EF3B2C", "#1D91C0", "#FEB24C", "grey60",
                            "#7FBC41", "#4D9221", "#276419"),
                     guide=guide_legend(
                       keywidth=0.5,
                       keyheight=0.5,
                       order=2,
                       override.aes=list(shape=c("Actinobacteria"=20,
                                                 "Bacteroidetes" =20,
                                                 "Firmicutes" =20,
                                                 "Deinococcus-Thermus" =20,
                                                 "Alphaproteobacteria" =18,
                                                 "Betaproteobacteria" =18,
                                                 "Gammaproteobacteria" =18
                                               )),
                     size=2
  ),
  na.translate=TRUE
)
) +
scale_shape_manual(
  values=c("Phylum"=20, "Class"=18),
  guide="none"
)

p2 <- p1 +
  new_scale_color() +
  geom_taxalink(
    data=linktab,
    mapping=aes(
      taxa1=Inhibitor,
      taxa2=Sensitive,
      color=Interaction
    ),
    alpha=0.6,
    offset=0.1,
    size=0.15,
    ncp=10,
    hratio=1,
    arrow=grid::arrow(length = unit(0.005, "npc"))
  ) +
  scale_colour_manual(values=c("chocolate2", "#3690C0", "#009E73"),
                      guide=guide_legend(
                        keywidth=0.8,
                        keyheight=0.5,
                        order=1,
                        override.aes=list(alpha=1,
                                          size=0.5))

```

```

)
)

p3 <- p2 +
  geom_fruit(
    data=BGCsda,
    geom=geom_tile,
    mapping=aes(y=Strain, x=BGCs, alpha=Count, fill=BGCs),
    offset=-0.9,
    pwidth=1,
    size=0.02,
    color = "grey50"
) +
  scale_alpha_continuous(range=c(0.1, 1),
                        name=bquote(paste(Log[10], "(, .("Count+1"), ")")),
                        guide=guide_legend(
                          keywidth = 0.4,
                          keyheight = 0.4,
                          order=4
                        )
  ) +
  scale_fill_manual(
    values=c("#66C2A5", "#FC8D62", "#8DA0CB", "#E78AC3",
             "#A6D854", "#FFD92F", "#E5C494", "#B3B3B3"),
    guide=guide_legend(keywidth = 0.4, keyheight = 0.4, order=3)
  )

p4 <- p3 +
  geom_tiplab(
    mapping=aes(label=names),
    align=TRUE,
    size=1,
    linetype=NA,
    offset=7.8
  )

p5 <- p4 +
  new_scale_fill() +
  geom_fruit(
    data=weighttab,
    geom=geom_bar,
    mapping=aes(
      x=value,
      y=Strain,
      fill=Number
    ),
    stat="identity",
    orientation="y",
    offset=0.48,
    pwidth=2,
    axis.params=list(
      axis="x",
      text.angle=-45,
      hjust=0,
      vjust=0.5,
      nbreak=4,
    )
  ) +

```

```

scale_fill_manual(
  values=c("#E41A1C", "#377EB8", "#4DAF4A", "#984EA3"),
  guide=guide_legend(keywidth=0.5, keyheight=0.5, order=5)
) +
theme(
  legend.background=element_rect(fill=NA),
  legend.title=element_text(size=6.5),
  legend.text=element_text(size=5),
  legend.spacing.y = unit(0.02, "cm"),
  legend.margin=margin(0.1, 0.9, 0.1, -0.9, unit="cm"),
  legend.box.margin=margin(0.1, 0.9, 0.1, -0.9, unit="cm"),
  plot.margin = unit(c(-1.2, -1.2, -1.2, 0.1), "cm")
)

```

p5

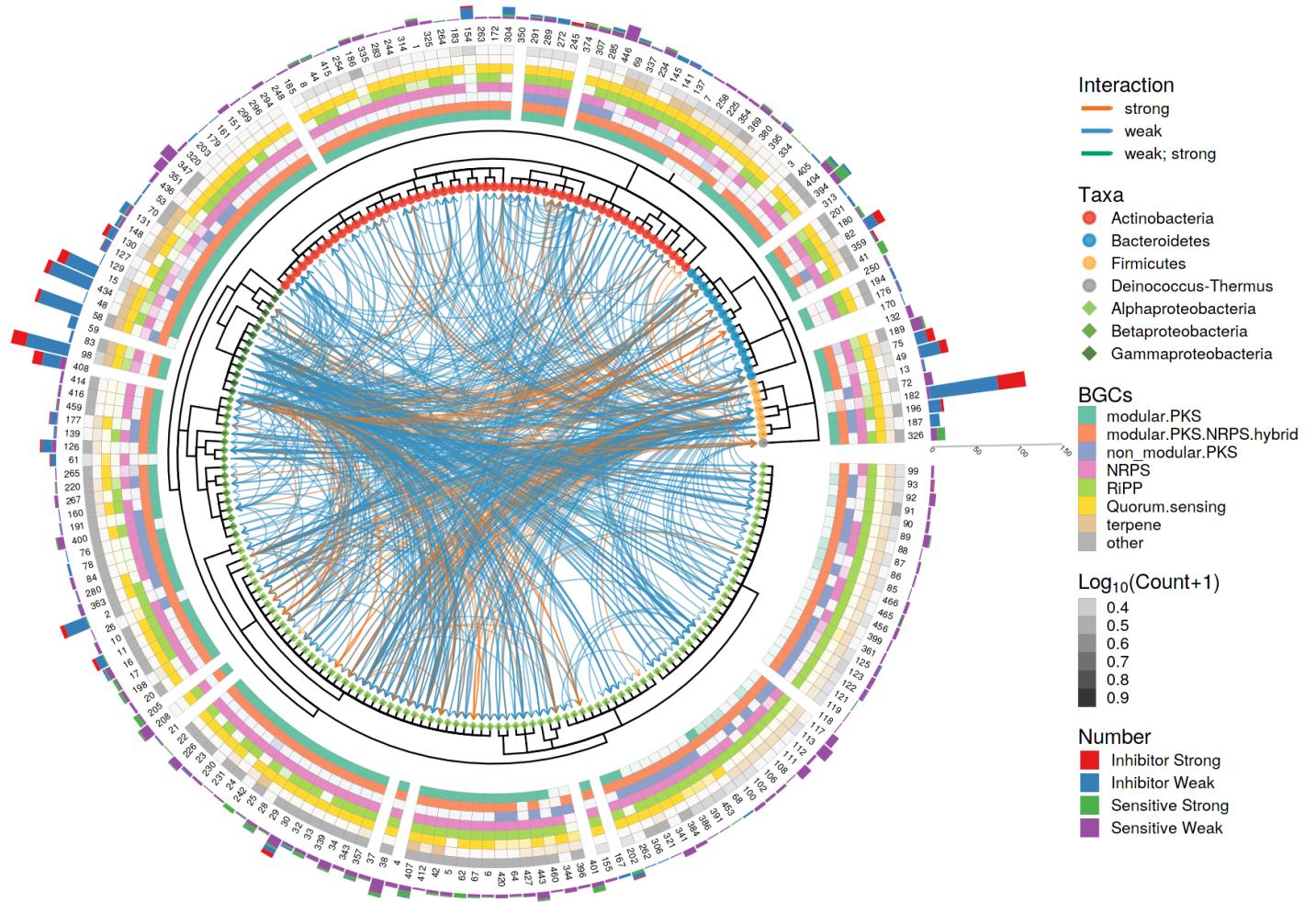


Fig. S6: The Phylogenetic tree of isolates from *Arabidopsis* leaf microbiome (Bai et al. 2015). The lines in inner represent the directional inhibitory interactions, the direction of arrow represents from inhibitor to sensitive, and the color of line represents the weak or strong interactions. The color of tip point represents the taxonomy annotation, circle represents the phylum, square represents the class from Proteobacteria. The heat map of external ring represents the number of detected BGCs (biosynthesis gene clusters). The ring from inner to outside represents the Modular polyketide synthase, Modular PKS non-ribosomal peptide synthetase (NRPS) hybrid, Non modular PKS, NRPS, ribosomally synthesized and post-translationally modified peptide (RiPP), Quorum sensing, Terpene, and Others. The external bar represent the number of interactions of inhibitions or sensitivities per strain.

1.2 Geometric layers that supported by *geom_fruit* of *ggtreeExtra*

As mentioned above, *geom_fruit* is an general function for linking multiple geometric graph to tree. It supports multiple layout type tree, such as circular layout, inward circular layout, rectangular layout etc. It can re-order the input data based on tree structure and displayed the data at the external of tree with the geometric function. It is designed to work with some *geom* layers defined in *ggplot2* (Wickham 2016) and other *ggplot2* extension packages. Here is the list of the geometric functions that work seamlessly with *geom_fruit* (Table S1). Since the *ggplot2* (Wickham 2016) community keeps expanding and more *geom* layers will be implemented in either *ggplot2* (Wickham 2016) or other extensions, *ggtreeExtra* also will support more *geom* layers and will gain more power to explore data in future.

Table S1: List of geometric layers supported by 'geom_fruit()'

Package	Geom Layer	Description
ggdist	geom_dotinterval	creates dots, intervals, and quantile dotplots
	geom_pointinterval	creates point and multiple uncertainty interval
	geom_slab	creates slab geom
	geom_slabinterval	creates slab, point and interval meta-geom
ggimage	geom_image	visualizes image files
	geom_phylopic	queries image files from phylopic database and visualizes them
ggpattern	geom_bar_pattern	draws bar charts with support for pattern fills
	geom_boxplot_pattern	draws box and whiskers plot with support for pattern fills
	geom_col_pattern	draws bar charts using 'stat_identity()' with support for pattern fills
	geom_tile_pattern	draws rectangle by using the center of the tile and its size with support for pattern fills
ggplot2	geom_bar	draws bar charts
	geom_boxplot	draws box and whiskers plot
	geom_col	draws bar charts using 'stat_identity()'
	geom_label	draws a rectangle behind the text
	geom_point	creates scatterplots
	geom_raster	a high performance special case for all the tiles are the same size
	geom_text	adds text to the plot
ggrepel	geom_text_repel	adds text to the plot. The text labels repel away from each other and away from the data points
	geom_label_repel	draws a rectangle underneath the text. The text labels repel away from each other and away from the data points
ggstance	geom_bbarh	horizontal version of 'geom_bar()'
	geom_boxploth	horizontal version of 'geom_boxplot()'
	geom_colh	horizontal version of 'geom_col()'
ggstar	geom_star	creates scatterplots
ggsymbol	geom_symbol	creates scatterplots
scatterpie	geom_scatterpie	creates scatter pie plot

1.3 Adding further annotation to specific layers

When the annotation information should be add to same panel, we developed *geom_fruit_list* to add multiple layers to the same external ring of circular tree (Fig S7).

```
library(ggtreeExtra)
library(ggtree)
library(treeio)
library(ggplot2)
set.seed(1024)
tree <- rtree(100)
df <- data.frame(id=tree$tip.label, value=sample(seq_len(200), 100, replace=TRUE),
                  group=c(rep("A",50),rep("B",50)))
p <- ggtree(tree, layout="fan", open.angle=180) + xlim(-2, NA)
p <- p + geom_fruit_list(geom_fruit(data=df,
                                       geom=geom_bar,
                                       mapping=aes(x=value, y=id, fill=group),
                                       orientation="y",
                                       
```

```

    stat="identity"
  ),
  geom_fruit(data=df,
    geom=geom_text,
    mapping=aes(x=value, y=id, label=value),
    size=2,
    offset=0.055,
    pwidth=0.26,
    position=position_identityx(),
  )) +
  scale_fill_manual(values=c("chocolate2", "#009E73"),
    guide=guide_legend(keywidth = 0.5, keyheight = 0.5))

```

p

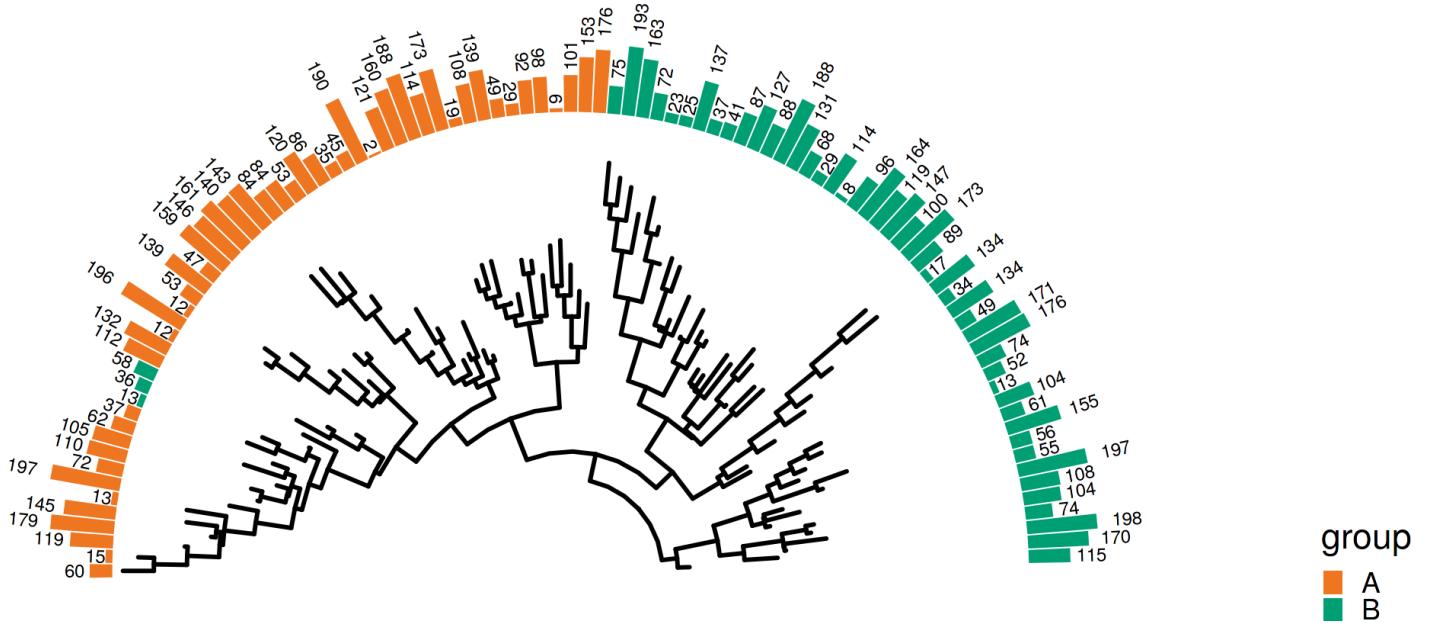


Fig. S7: visualizing multiple layers at them same external ring using *ggtreeExtra*

1.4 Summary

As far as we known, None of R packages support visualizing external data and align the plot to circular layout trees or other layout trees. There are some tools depending on python or other platform, they provide some features to annotate and visualize circular phylogenetic tree, but they are not general and still have some shortcomings. For example, *GraPhlAn* (Asnicar et al. 2015) and *ETE3* (Huerta-Cepas, Serra, and Bork 2016), both based on python, *GraPhlAn* (Asnicar et al. 2015) provided few graph layers to display associated data, while *ETE3* (Huerta-Cepas, Serra, and Bork 2016) did not support visualizing associated dataset in the outer ring of circular phylogenetic tree. There are some web tools support visualizing phylogenetic tree, such as *iTOL* (Letunic and Bork 2019), *Microreact* (Argimón et al. 2016) and *EvolView* (Subramanian et al. 2019). But the feature to link tree with external datasets is often missing, and they often need user to provide predefined data types, which make it tedious to use and debug. They are also not conducive to integration into the relevant data analysis pipeline.

ggtreeExtra is designed to work with *treeio*, *tidytree* and *gtree*. Since it depends R, an ideal environment for statistical analysis and visualization, the user can use it with other R package, such as *ggstar* (Xu 2020), *ggnewscale* (Campitelli 2020) and *ggplot2* (Wickham 2016). It inherits the merits of *ggplot2* (Wickham 2016), built based on grammar of graphics. So it is easy to use *ggtreeExtra*. In addition, there are no restriction of data types (Fig. S3 and S4). Combining multiple geometric layers to display associated data is supported (Fig. S4 and S1). The internal data of tree data or data added by %<+% operator also can be used in *geom_fruit* (Fig. S2). It is more efficient and convenient to visualize multi-dimensional data using multiple external ring layers (Fig. S4). In addition, the *radial*, *rectangular* and *slanted* layouts also are supported by *ggtreeExtra*. The tree in *geom_fruit* can be fully annotated with multiple layers (high light, clade

labels, group clade, etc functions of *ggtree*) (Fig. S1, S3 and S4).

Reference

- Argimón, S., K. Abudahab, R. J. E. Goater, A. Fedosejev, J. Bhai, C. Glasner, E. J. Feil, et al. 2016. “Microreact: Visualizing and Sharing Data for Genomic Epidemiology and Phylogeography.” Journal Article. *Microb Genom* 2 (11): e000093. <https://doi.org/10.1099/mgen.0.000093>.
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- Campitelli, Elio. 2020. *Ggnewscale: Multiple Fill and Colour Scales in 'Ggplot2'*. <https://CRAN.R-project.org/package=ggnewscale>.
- Helfrich, Eric J. N., Christine M. Vogel, Reiko Ueoka, Martin Schäfer, Florian Ryffel, Daniel B. Müller, Silke Probst, Markus Kreuzer, Jörn Piel, and Julia A. Vorholt. 2018. “Bipartite Interactions, Antibiotic Production and Biosynthetic Potential of the Arabidopsis Leaf Microbiome.” Journal Article. *Nature Microbiology* 3 (8): 909–19. <https://doi.org/10.1038/s41564-018-0200-0>.
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