

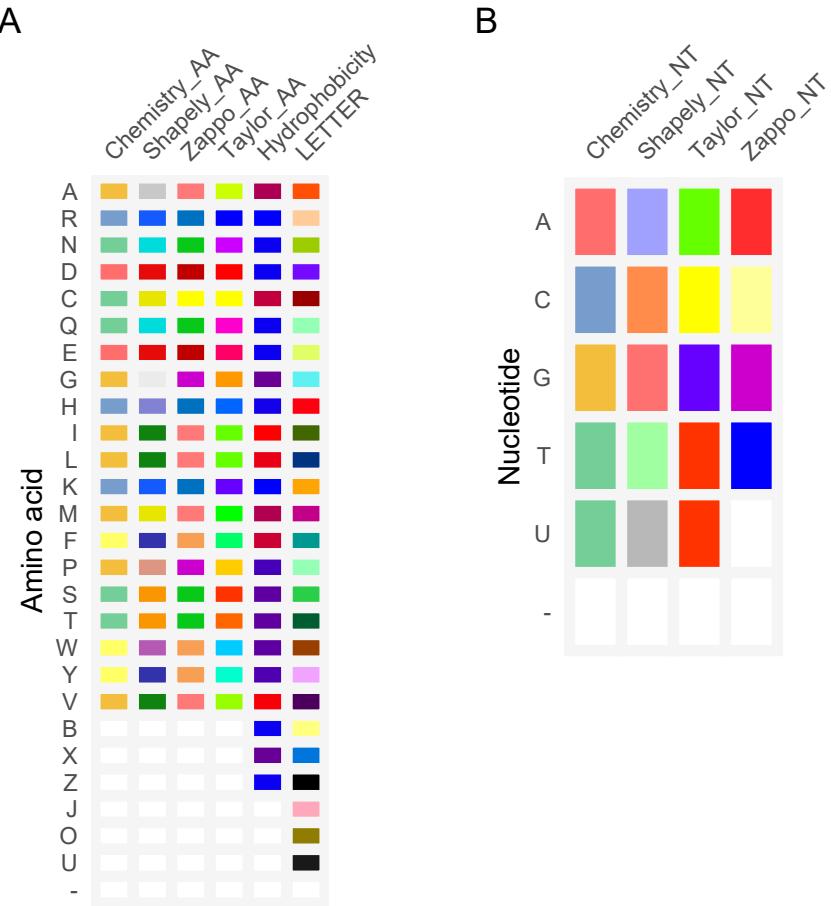
Supplemental File of

**ggmsa: a visual exploration tool for multiple sequence alignment and associated data**

```
library(ggmsa)
library(ggplot2)
library(ggtree)
library(gggenes)
library(ape)
library(Biostrings)
library(ggnewscale)
library(dplyr)
library(ggtreeExtra)
library(phangorn)
library(RColorBrewer)
library(patchwork)
library(ggplotify)
library(aplot)
library(magick)
library(treeio)

protein_sequences <- system.file("extdata", "sample.fasta", package = "ggmsa")
nt_sequence <- system.file("extdata", "LeaderRepeat_All.fa", package = "ggmsa")
miRNA_sequences <- system.file("extdata", "seedSample.fa", package = "ggmsa")
tp53_sequences <- system.file("extdata", "tp53.fa", package = "ggmsa")
tp53_genes <- system.file("extdata", "TP53_genes.xlsx", package = "ggmsa")
```

**Fig. S1: Amino acid and nucleotide color schemes in ggmsa**



**Fig. S1: Color schemes in ggmsa.** (A) Amino acid color schemes. Schemes are either quantitative, reflecting empirical or statistical properties of amino acids; or qualitative, reflecting physicochemical attributes. Chemistry is colored according to side-chain chemistry and is also used in DNASTAR applications (Burland 2000); Shapely matches the RasMol amino acid color schemes, which are, in turn, based on Robert Fletterick's Shapely models. Zappo is a qualitative scheme developed by M. Clamp. The residues are colored according to their physicochemical properties; Taylor (Taylor and W. 1997) is taken from Taylor and is also used in JalView (Waterhouse et al. 2009); Hydrophobicity colors the residues in the alignment based on the hydrophobicity table (Kyte and Doolittle 1982). B, X, Z, J, O, and U are amino acid ambiguity codes: B is aspartate or asparagine; Z is glutamate or glutamine; X, J, O, U is an unknown (or 'other'); “-” indicate a gap. (B) Nucleotide color schemes used by ggmsa.

## Fig. S2: MSA view with break down layout

`facet_msa()` module allows for showing more alignment data in a restricted canvas. The long sequence is broken down and displayed in several lines.

```
# 4 fields
ggmsa(protein_sequences, start = 0, end = 400, font = NULL, color = "Chemistry_AA") +
  facet_msa(field = 100)
```

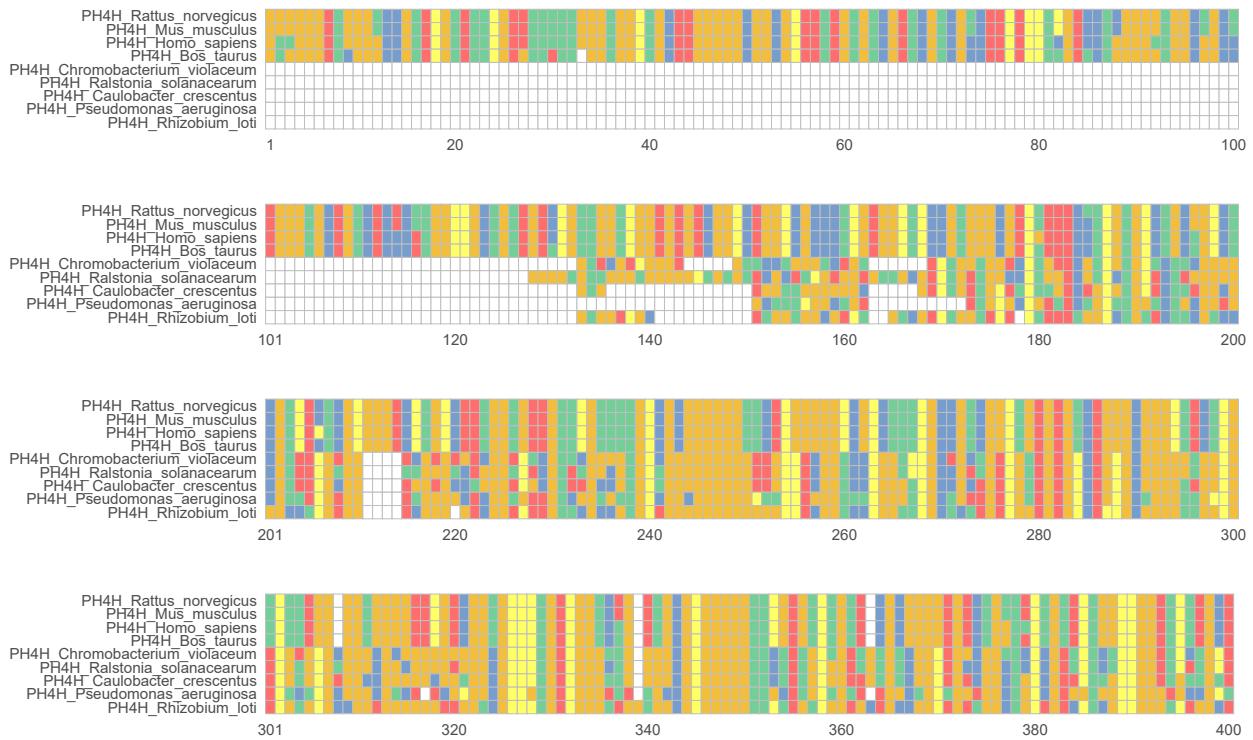


Fig. S2: MSA view with break down layout. The long sequence is displayed in several lines.

## Fig. S3: MSA view with circular layout tree

With `ggtreeExtra` (Xu et al. 2021), displaying alignment in a circular layout can also be supported. The `geom_fruit()` will automatically align the MSA graphs to the tree with a circular layout.

```
library(ggtree)
library(ggtreeExtra)
sequences <- system.file("extdata", "sequence-link-tree.fasta", package = "ggmsa")

x <- readAAStringSet(sequences)
d <- as.dist(stringDist(x, method = "hamming"))/width(x)[1])
tree <- bionj(d)
data <- tidy_msa(x, 120, 200)

p1 <- ggtree(tree, layout = 'circular') +
```

```
geom_tiplab(align = TRUE, offset = 0.545, size = 2) +  
  xlim(NA, 1.3)  
p1 + geom_fruit(data = data, geom = geom_msa, offset = 0,  
  pwidht = 1.2, font = NULL, border = NA)
```

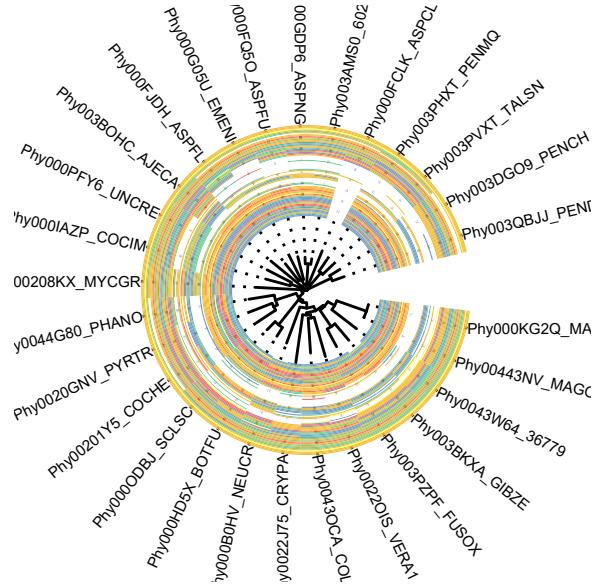


Fig. S3: Example of MSA view with circular layout phylogenetic tree.

Fig. S4: Visualizing MSA with a phylogenetic tree

Similar to `geom_fruit()`, the `geom_facet()` provided in `ggtree` (Yu et al. 2017) also works with `ggmsa` to display MSA with a phylogenetic tree.

```
x <- readAAStringSet(protein_sequences)
d <- as.dist(stringDist(x, method = "hamming")/width(x)[1])
library(ape)
tree <- bionj(d)
library(ggtree)
p <- ggtree(tree) + geom_tiplab()

data = tidy_msa(x)
p + geom_facet(geom = geom_msa, data = data, panel = 'msa',
                font = NULL, border = NA, color = "Chemistry_AA") +
    xlim_tree(1)
```

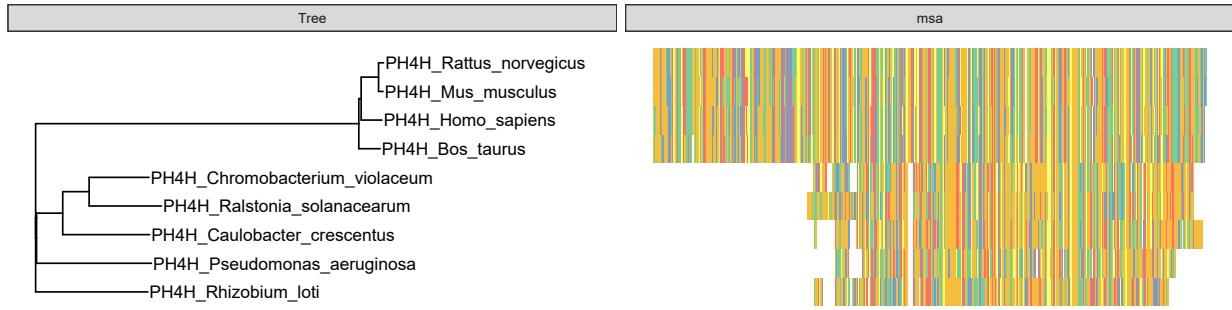


Fig. S4: A phylogenetic tree with MSA that it was built from.

### Fig. S5: Reassign residue order in sequence bundle

```

negative <- system.file("extdata", "Gram-negative_AKL.fasta",
                       package = "ggmsa")
positive <- system.file("extdata", "Gram-positive_AKL.fasta",
                       package = "ggmsa")

ggSeqBundle(list(negative, positive),
            alpha = 0.1,
            bundle_color = c("#FC8D62", "#8DA0CB"),
            lev_molecule = c("-", "W", "Y", "R", "F", "H", "M", "E",
                            "K", "Q", "D", "N", "L", "I", "C", "T",
                            "V", "P", "S", "A", "G"))

```

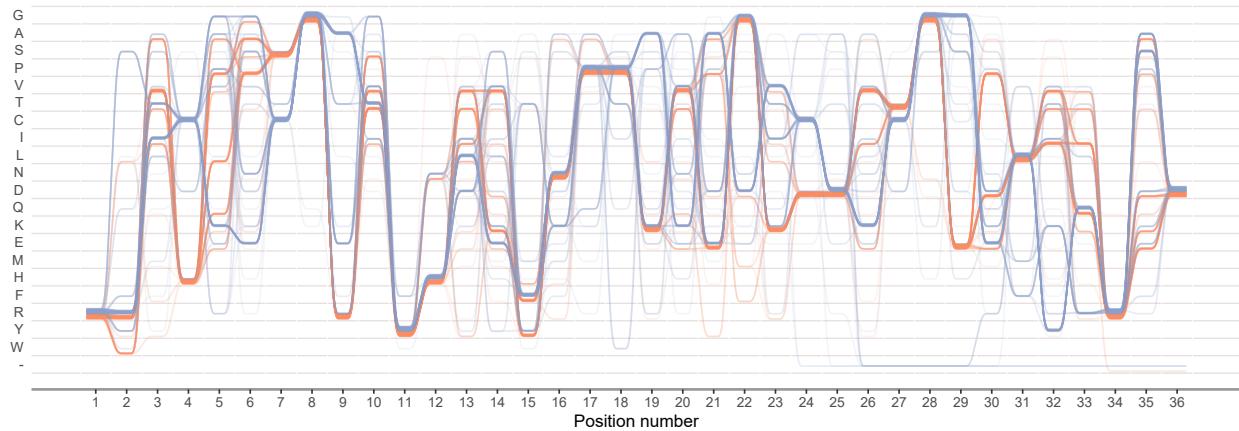
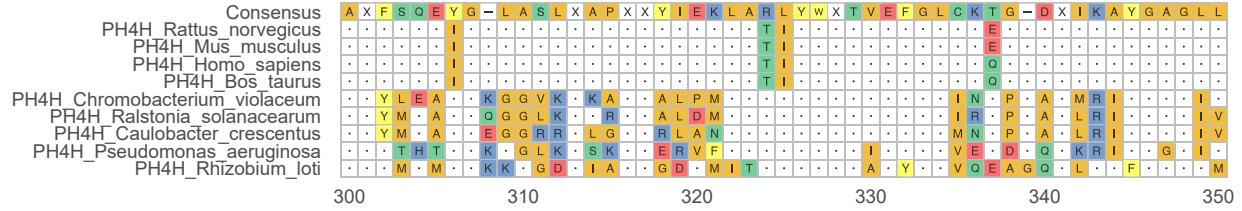


Fig. S5: Reassign residue order in sequence bundle. The preference of residues' physicochemical properties can be reflected by adjusting the order of the letters on the y-axis. This figure reassigns residue order with amino acid molecule weight from light (top) to heavy (bottom).

**Fig. S6: MSA with consensus sequence**

```
ggmsa(protein_sequences, 300, 350, char_width = 0.5,
       seq_name = TRUE, consensus_views = TRUE, use_dot = TRUE)
```

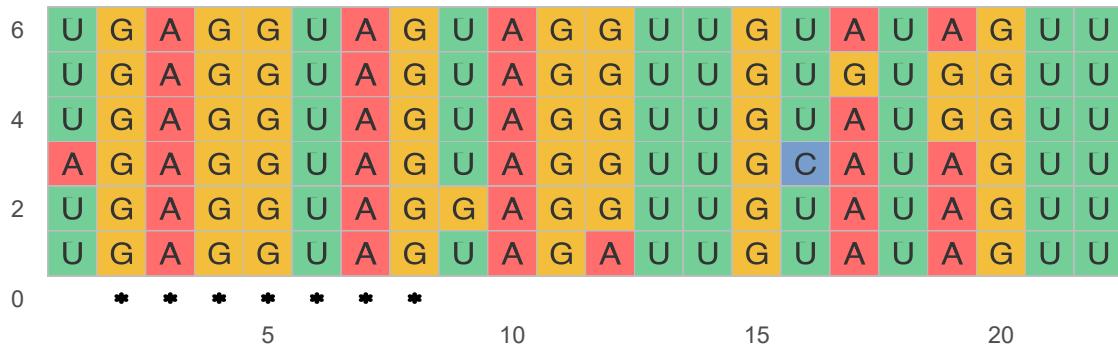


**Fig. S6: Example of displaying consensus sequence for an MSA.** The consensus sequence is displayed above the alignment and shows which residues are conserved and which residues are variable. A consensus is constructed from the most frequent residue at each site.

**Fig. S7: Seed region annotation (asterisks) for miRNA sequences**

The `geom_seed()` helps to identify microRNA seed region by adding asterisks or a shaded block. The seed region is a conserved heptameric sequence that is mostly situated at positions 2-7 from the miRNA 5'-end.

```
miRNA_sequences <- system.file("extdata", "seedSample.fa", package = "ggmsa")
ggmsa(miRNA_sequences, char_width = 0.5, color="Chemistry_NT") +
  geom_seed(seed = "GAGGUAG", star = TRUE) + coord_cartesian()
```



**Fig. S7: Example of highlighting miRNA seed region.** Asterisks are used for marking the seed region.

Fig. S8: Seed region annotation (shaded block) for miRNA sequences

```
ggmsa(miRNA_sequences, char_width = 0.5, seq_name = TRUE, none_bg = TRUE) +
  geom_seed(seed = "GAGGUAG") + coord_cartesian()
```

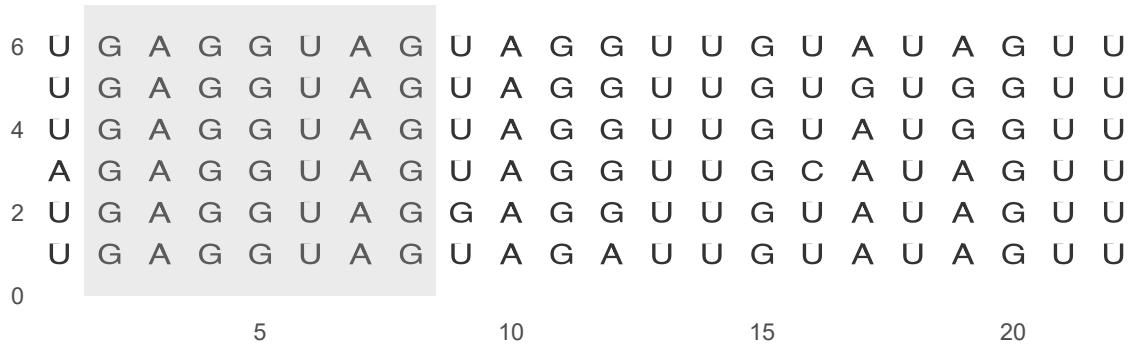


Fig. S8: Example of highlighting miRNA seed region. Annotation of sequence seed region with a shaded block

Fig. S9: Displaying ancestral sequences in Tree-MSA plot

```
x <- read.phyloxml("data/msa_phyloxml.xml")
p <- ggtree(x, size = 2) + xlim_tree(0.12)
tidymsa <- extract_seq(p)

p1 <- treeMSA_plot(p,
                     tidymsa,
                     ancestral_node = 11,
                     sub = FALSE,
                     color = "Chemistry_NT",
                     seq_colname = "mol_seq")

p2 <- treeMSA_plot(p,
                     tidymsa,
                     ancestral_node = 11,
                     sub = TRUE,
                     color = "Chemistry_NT",
                     seq_colname = "mol_seq")

p3 <- treeMSA_plot(p,
                     tidymsa,
                     ancestral_node = c(11,14,16,17),
                     sub = FALSE,
                     color = "Chemistry_NT",
                     seq_colname = "mol_seq")

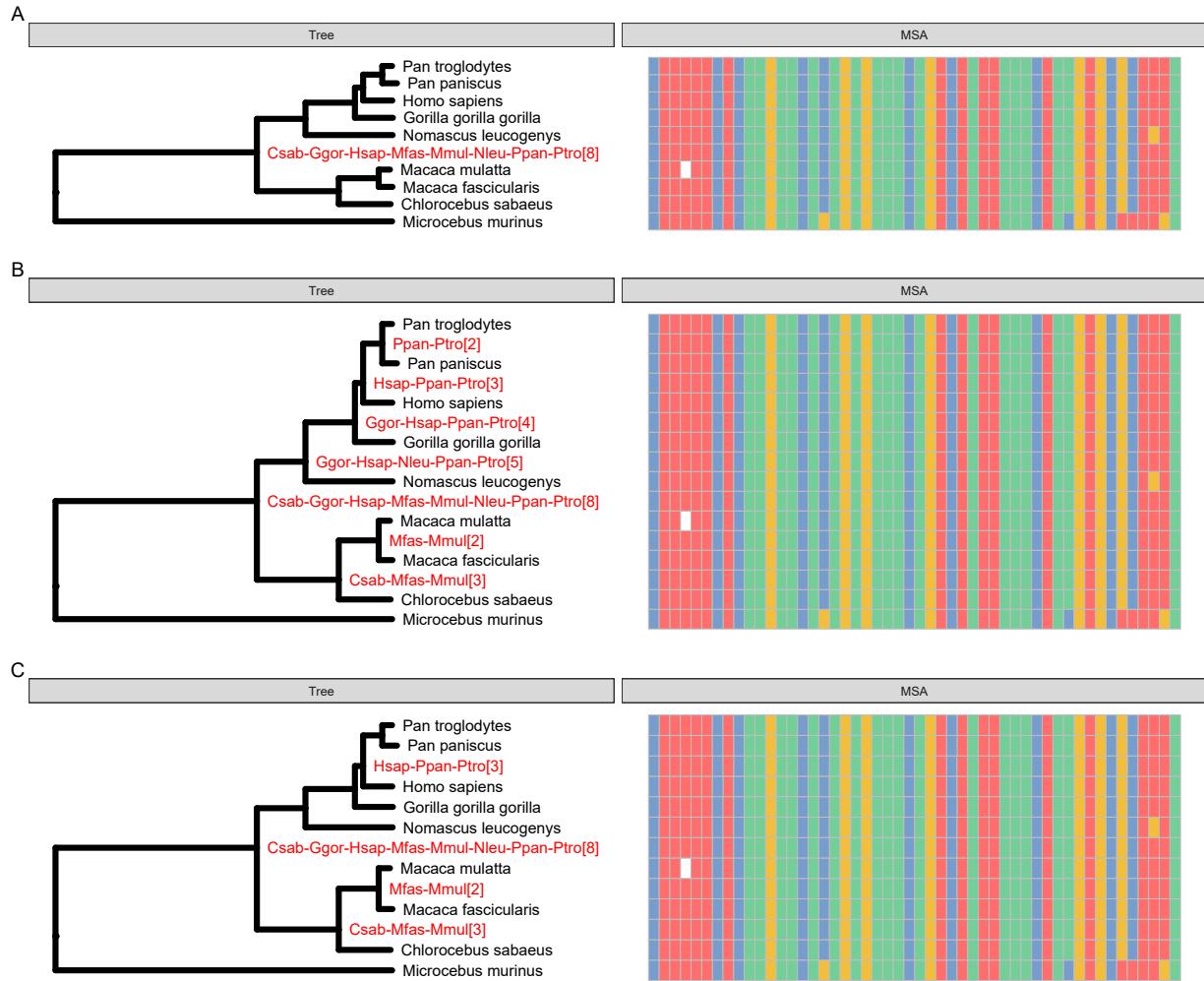
pp <- plot_list(gglist = list(p1,p2,p3),
```

```

    ncol = 1,
    tag_levels = "A",
    heights = c(0.4,0.7,0.6))

```

pp



**Fig. S9: Displaying ancestral sequences in tree-MSA plot.** Different display modes of `treeMSA_plot()`: (A) Only displaying the sequence corresponding to the selected ancestral node; (B) Displaying all ancestral sequences of the subtree. (C) Displaying corresponding ancestor sequences of multiple selected internal nodes.

**Fig. S10: MSA of SARS-CoV-2 Spike protein RBD coloring by RBD-hACE2 binding affinity**

We applied the Deep Mutational Scanning (DMS) data of ACE2-binding affinity to MSA to functionally annotate the mutated sites in the sequence. In addition, the application scope of the arc diagram is extended to the interaction process of ACE2-RBD (Fig. S11). Through MSA that used ACE2-binding affinity annotation and ACE2-RBD interaction arc diagram, we can observe how the mutation of Omicron affects ACE2-RBD interaction. Although DMS data only reveal the binding affinity of a single amino acid mutation (Fig. S10), we believe that an MSA plot annotated with DMS data can reflect changes in ACE2-RBD interactions to a certain extent. Arc plot representing ACE2-RBD interactions prove this (Fig. S11, ??). In the ACE2-RBD arc diagram, Omicron mutation sites K417N, G446S, and G496S show loss of hydrogen bond interaction, and these sites all show decreased binding ability in the binding affinity annotation diagram. The N501Y mutation in the binding affinity diagram shows an increased binding ability, which may form a new binding site. This has not been shown in the Arc plot (possible cause: the interaction site information contained in the 7WPB is not completely accurate) but has been reported in other literature (Yin et al. 2022).

```

colRD <- colorRampPalette(rev(brewer.pal(n = 9, name = "OrRd")))
colBU <- colorRampPalette(colors = rev(c("#185089", "#FFF7EC")))

data <- "data/s_RBD.fasta"
dms <- read.csv("data/DMS.csv")
del <- c("expr_lib1", "expr_lib2",
       "expr_avg", "bind_lib1",
       "bind_lib2")
dms <- dms[,!colnames(dms) %in% del]

tidymsa <- tidy_msa(data)
tidymsa <- assign_dms(tidymsa, dms)
#Mapping the position to the protein-protein interaction plot
tidymsa$position <- tidymsa$position + 330

p <- ggplot() +
  geom_msa(data = tidymsa,
            char_width = 0.5,
            dms = TRUE,
            seq_name = TRUE,
            show.legend = TRUE) +
  theme_msa() +
  scale_fill_gradientn(name = "ACE2 binding",
                        colors = c(colRD(75), colBU(25))) +
  facet_msa(50)
p

```

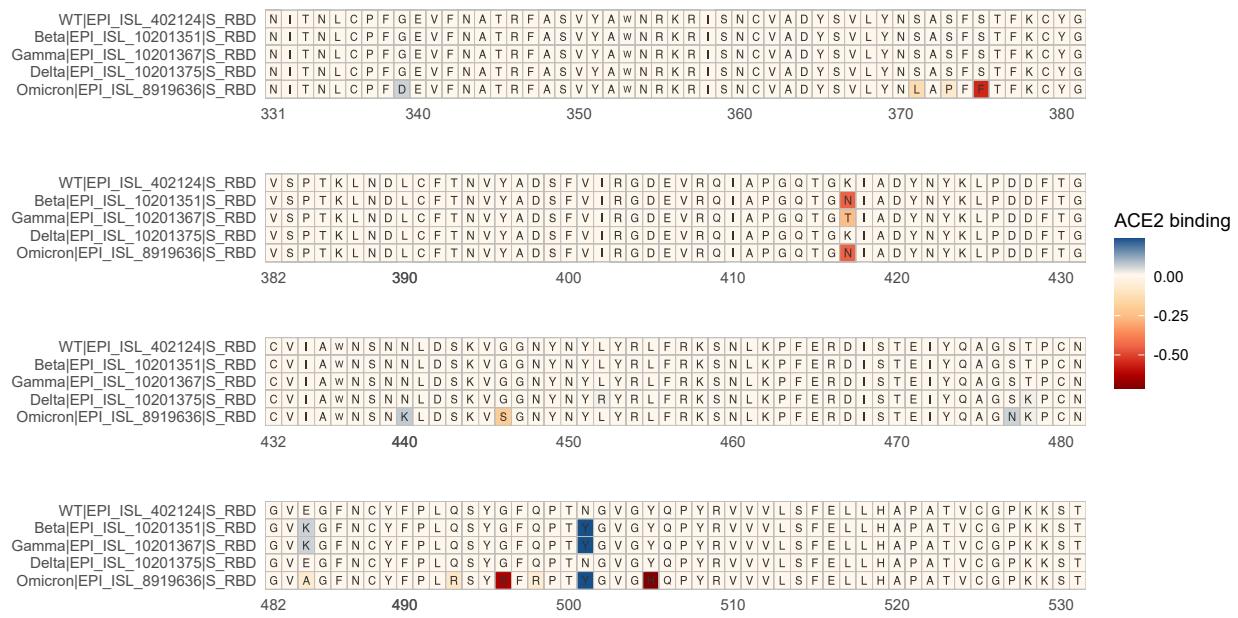


Fig. S10: **MSA of SARS-CoV-2 Spike protein RBD.** Mutations in Spike protein Receptor Binding Domain (RBD) are compared with wild-type SARS-CoV-2 and four other strains (Beta, Gamma, Delta, and Omicron, all from GISAID). The colors in MSA show RBD-hACE2 (human Angiotensin-converting enzyme 2) binding affinity scores. ACE2-binding affinity is shown in shades of blue or red representing higher or lower ACE2 affinity, respectively. The ACE2 binding scores are from the Deep Mutational Scanning (DMS) of SARS-CoV-2 RBD. Scores represent binding constants ( $\Delta\log_{10} \text{KD}$ ) relative to the wild-type reference amino acid.

Fig. S11: simplified and global arc diagram of ACE2-RBD interaction

```

#read sequences
x <- readAAStringSet("data/ACE2.fasta")
y <- readAAStringSet("data/Spike_RBD.fasta")

#read protein-protein position
inter1 <- read.csv("data/6m0j_inter.csv")
inter2 <- read.csv("data/7wpb_inter.csv")

#tidy data
t1 <- tidy_msa(x, start = 19)
t2 <- tidy_msa(y, start = 331)
t_merge <- merge_seq(t1, gap = 100, t2)

h1 <- tidy_hdata(100, inter1, t1, t2)
h2 <- tidy_hdata(100, inter2, t1, t2)

#protein-protein interactive plot
p_interactive <- ggplot() +
  geom_msa(data = t_merge, border = NA, font = NULL, seq_name = FALSE) +
  theme_msa() + theme(axis.text = element_blank()) +
  geom_helix(helix_data = list(known = h1, predicted = h2))

#simplified p-p interactive plot
ACE2 <- t_merge[t_merge$position %in% c(h1$i,h2$i),]
spike <- t_merge[t_merge$position %in% c(h1$j,h2$j),]
t1_reset <- reset_pos(ACE2)
t2_reset <- reset_pos(spike)
simplify_merge <- merge_seq(previous_seq = t1_reset,
                               subsequent_seq = t2_reset,
                               gap = 5,
                               adjust_name = FALSE)

sim_h1 <- simplify_hdata(hdata = h1, sim_msa = simplify_merge)
sim_h2 <- simplify_hdata(hdata = h2, sim_msa = simplify_merge)

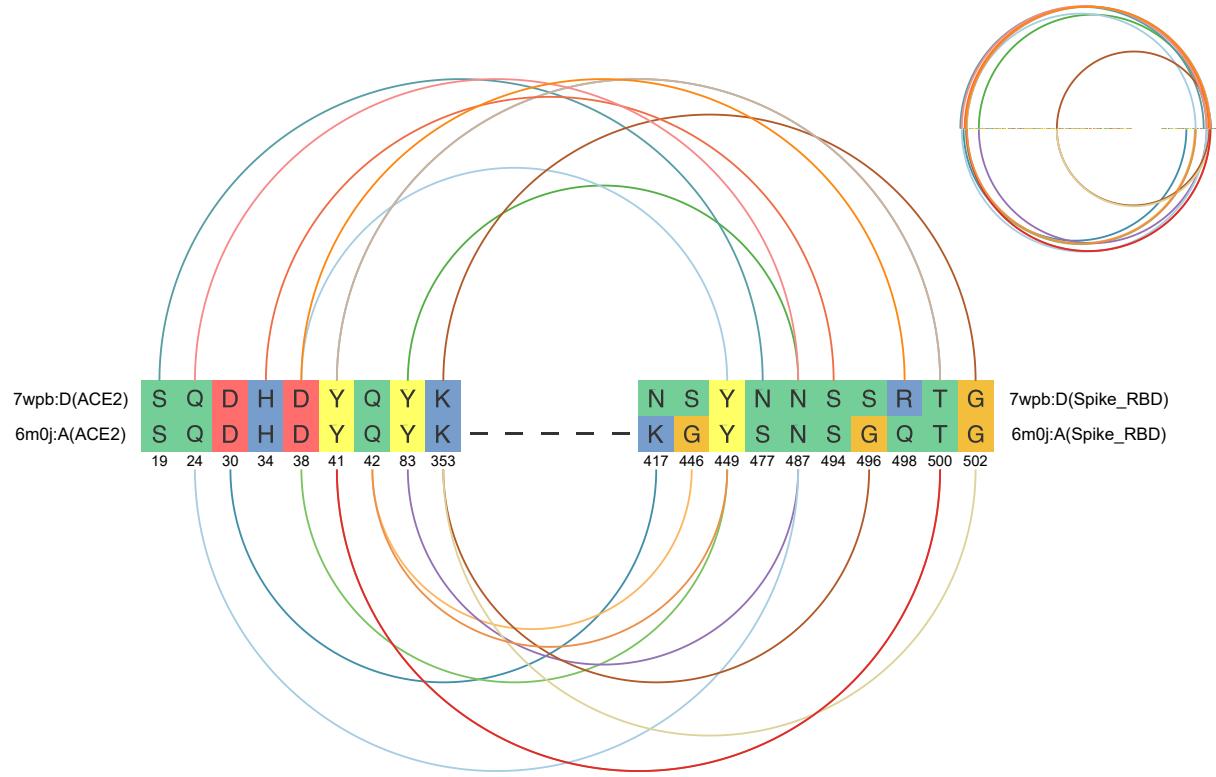
##break and label
b <- simplify_merge[simplify_merge$character != "-", "position"] %>% unique()
l <- c(inter1$Res.no.1,inter1$Res.no..2,
       inter2$Res.no.1,inter2$Res.no..2) %>% unique

p_sim <- ggplot() +
  geom_msa(data = simplify_merge, border = NA, char_width = 0.5, seq_name = FALSE) +
  ggmsa:::theme_msa() +
  geom_helix(helix_data = list(known = sim_h1,
                               predicted = sim_h2),
             overlap = FALSE) +
  geom_text(mapping = aes(x = b,
                         y = 0.25,
                         label = l[order(l)]),
            size = 3) +
  theme(axis.text.x = element_blank(),
        axis.ticks.x = element_blank())

```

```
axis.text.y = element_blank() +
scale_y_discrete(labels = c("6m0j:A(ACE2)", "7wpb:D(ACE2)")) +
geom_text(aes(x = c(27.2,27.2),
y = c(1,2),
label = c("6m0j:A(Spike_RBD)",
"7wpb:D(Spike_RBD)")),
size = 3.5) +
geom_text(aes(x = c(-1.5,-1.5),
y = c(1,2),
label = c("6m0j:A(ACE2)",
"7wpb:D(ACE2)")),
size = 3.5) + xlim(-2,29)

require(patchwork)
p_sim + inset_element(p_interactive, 0.7, 0.7, 1.1, 1.1)
```



**Fig. S11: Arc diagram of ACE2-RBD interaction.** The two sequences on the left panel are ACE2, and the two sequences on the right panel are RBD sequences of S protein of wild-type and Omicron, respectively. ACE2 and RBD are placed in the same horizontal position, and the residual sites that generated hydrogen bonds were connected by arcs. The upper arc represents the ACE2-Omicron\_RBD and the lower arc represents the ACE2-WT\_RBD. The main arc plot only shows the residues at the interaction sites, and the global arc diagram is shown in the upper right corner. Data of interacting sites was obtained from the PDBsum database, and the corresponding PDB ID were 7WPB and 6M0J.

**Table S1: Comparison of ggmsa with popular free MSA visualization tools**

We compared MSA visualization methods among ggmsa and other tools (**msa** (Bodenhofer et al. 2015), **MSAViewer** (Yachdav et al. 2016), **AliView** (Larsson 2014), **Jalview** (Waterhouse et al. 2009), **ALVIS** (Schwarz et al. 2016))

Table S1: Comparison of ggmsa with popular free MSA visualization tools

Tools	Platform	Sequence Logos	Sequence bundles	Stacked MSA <sup>1</sup>	Layouts for stacked MSA	Integrating external data into stacked MSA <sup>2</sup>		Exploring sequence recombination <sup>3</sup>	User interface
						YES	YES		
ggmsa	R package	YES	YES	YES	rectangular fragmen- tary circular	YES	YES	programming and command line	
msa	R package	YES	NO	YES	rectangular	NO	NO	programming and command line	
MSAViewer	Web service	YES	NO	YES	rectangular	NO	NO	interactive	
AliView	Desktop application	NO	NO	YES	rectangular	NO	NO	interactive	
Jalview	Desktop application and web service	YES	NO	YES	rectangular	NO	NO	interactive	
ALVIS	Desktop application	YES	YES	YES	rectangular	NO	NO	interactive	

<sup>1</sup> Stacked MSA: it represents all sequences as rows and homologous residue positions as columns

<sup>2</sup> Extended MSA: adding associated into stacked MSA plots

<sup>3</sup> A visualization method that designed for detecting sequence recombination signals

**Fig. 2A:** The combination of sequence logos and sequence bundles (R code)

```

p2a <- seqlogo("data/Gram-NP-merge.fa",
                color = "Chemistry_AA",
                font = "DroidSansMono") +
coord_cartesian()

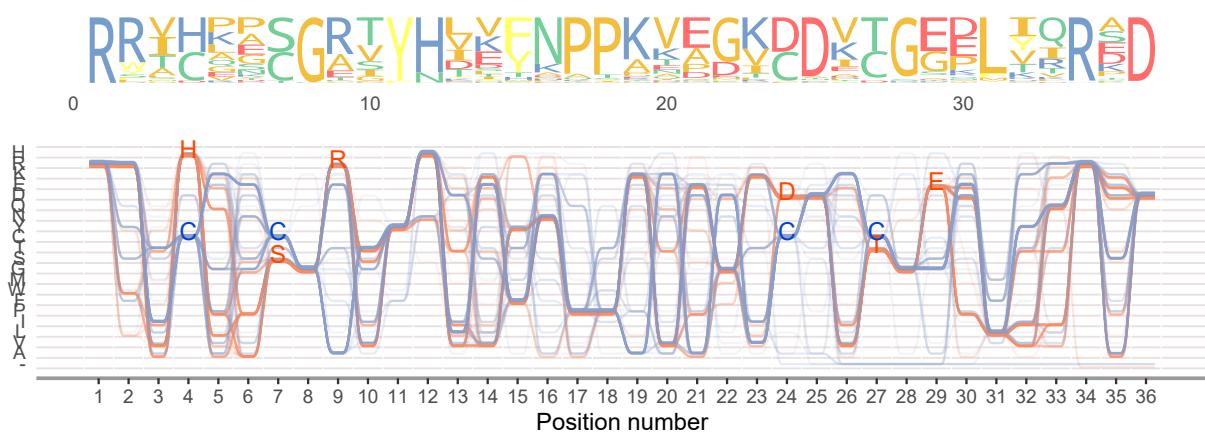
negative <- system.file("extdata", "Gram-negative_AKL.fasta",
                        package = "ggmsa")
positive <- system.file("extdata", "Gram-positive_AKL.fasta",
                        package = "ggmsa")

pos <- data.frame(x= c(4, 7, 9, 24, 27, 29,
                      4, 7, 24, 27),
                   y = c(c(21, 11, 20, 17, 12, 18) + .3,
                         c(13, 13, 13, 13) + .5
                     ),
                   label = c("H", "S", "R", "D", "T", "E",
                             "C", "C", "C", "C"),
                   color = c(rep("#ff4700",6),
                             rep("#0443d0",4)))

p2b <- ggSeqBundle(list(negative, positive),
                    alpha = 0.1,
                    bundle_color = c("#FC8D62","#8DA0CB"))+ #RColorBrewer: Set2:2-3
geom_text(data = pos,
          mapping = aes(x, y,
                        label = label,
                        color = I(color)),
          inherit.aes = FALSE,
          size = 4)

plot_list(gglist = list(p2a, p2b), ncol = 1, heights = c(0.3,1))

```

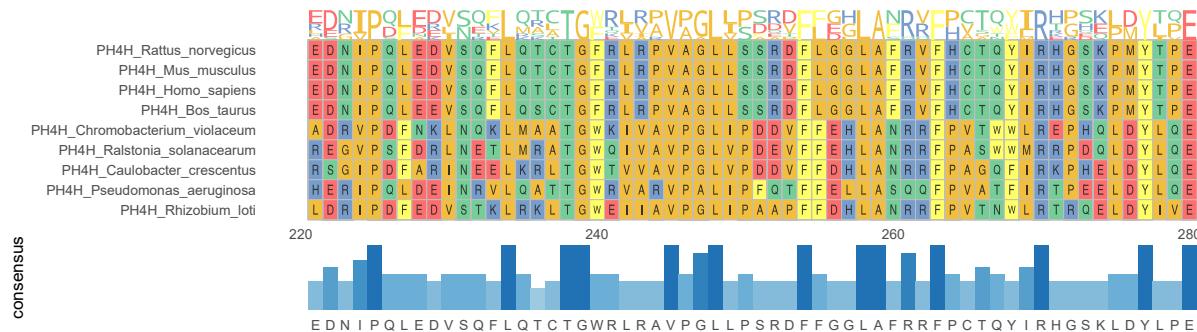


**Fig. 2A| The combination of sequence logos and sequence bundles.** The data contain the adenylate kinase lid (AKL) domain both of Gram-negative and Gram-negative bacteria. 100 sequences for each group. The sequence logo (top panel) represents the AKL sequence pattern and the sequence

bundle (bottom panel) represents the different residues relationship between Gram-negative (orange) and Gram-negative (purple) bacteria. The site at 4, 7, 9, 24, 27, and 29 has an exclusive pattern (His4, Ser7, Arg9, Asp24, Thr27, Glu29) in the Gram-negative sequences. And the site at 4, 7, 24, and 27 both contain the Cysteines in the Gram-positive sequences. These residues relationship in agreement with the structural stability of the AKL domain. Gram-negatives form a hydrogen-bonding network by the exclusive pattern and Gram-positive bound metal ion to form coordinated tetrahedrally by Cys. (Date from BioVis2013 and repeated example from Science Practice)

**Fig. 2B: An example of MSA visualization and MSA annotations (R code)**

```
ggmsa(protein_sequences, 221, 280, seq_name = TRUE, char_width = 0.5) +
  geom_seqlogo(color = "Chemistry_AA") +
  geom_msabar()
```



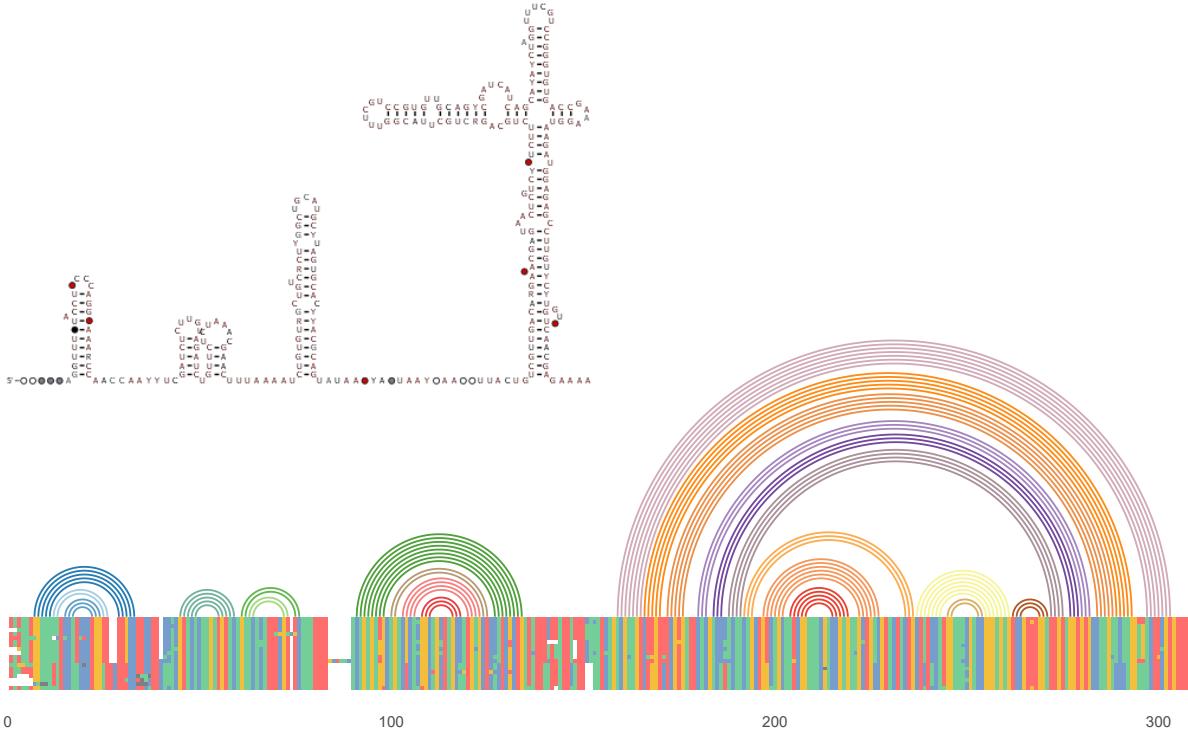
**Fig. 2B| A local visualization of the sequence alignment of the phenylalanine hydroxylase protein (PH4H) within nine species.** The center panel is the main MSA plot with residues in the alignment colored according to the Chemistry color scheme (amino acids are colored according to their side-chain chemistry). The top and bottom panels are corresponding annotations with MSAs, showing the conservation patterns at each position by sequence logos and the distribution of the high-frequency residue by a bar chart, respectively.

Fig. 3A: Visual methods of surveying RNA co-variation (R code)

```
p_RF03120_SS <- image_read_svg("data/RF03120_SS.svg")
q_RF03120_SS <- as.ggplot(p_RF03120_SS)

RF03120_msa<- system.file("extdata", "Rfam", "RF03120.fasta", package = "ggmsa")
RF03120_ss <- system.file("extdata", "Rfam", "RF03120_SS.txt", package = "ggmsa")
RF_arc <- readSSfile(RF03120_ss, type = "Vienna" )
p3a <- ggmsa(RF03120_msa,
              font = NULL,
              color = "Chemistry_NT",
              seq_name = F,
              show.legend = F,
              border = NA) +
  geom_helix(helix_data = RF_arc) +
  theme(axis.text.y = element_blank())
require(patchwork)
layout <- c(
  area(t = 0, b = 4, l = 1, r = 4),
  area(t = 0, b = 2, l = 1, r = 2))

p3A <- p3a + q_RF03120_SS + plot_layout(design = layout)
p3A
```



**Fig. 3A| Visual methods of surveying RNA co-variation.** The data from the Rfam database [family RF03120] include 19 seed alignments of Sarbecovirus 5'UTR (including 6 SARS-CoV-2 isolates sequences) and the corresponding consensus RNA secondary structure. Compensatory mutations in MSA can be detected by checking alignment columns in positions corresponding to arcs.

Fig. 3B: Visual RNA structural changes (R code)

```
color_589 = c(rep("#67a9cf",5),
             rep("#fed976",2),
             rep("#67a9cf",3),
             rep("#008837",1),
             rep("#fed976",2),
             rep("#67a9cf",3),
             rep("#fc8d59",10),
             rep("#67a9cf",3),
             rep("#fed976",2),
             rep("#fc8d59",3),
             rep("#67a9cf",1),
             rep("#fed976",1),
             rep("#67a9cf",1),
             rep("#008837",2),
             rep("#67a9cf",4),
             rep("#fed976",1),
             rep("#67a9cf",1),
             rep("#a6dba0",2),
             rep("#67a9cf",5),
             rep("#008837",2),
             rep("#67a9cf",4),
             rep("#fc8d59",6),
             rep("#67a9cf",4),
             rep("#008837",1),
             rep("#67a9cf",1),
             rep("#a6dba0",1),
             rep("#67a9cf",4),
             rep("#fed976",2),
             rep("#67a9cf",5),
             rep("#969696",1))

color_590 = c(rep("#67a9cf",5),
              rep("#fed976",2),
              rep("#67a9cf",3),
              rep("#008837",1),
              rep("#fed976",2),
              rep("#67a9cf",2),
              rep("#fc8d59",12),
              rep("#67a9cf",2),
              rep("#fed976",2),
              rep("#fc8d59",3),
              rep("#67a9cf",1),
              rep("#fed976",1),
              rep("#67a9cf",1),
              rep("#008837",2),
              rep("#67a9cf",4),
              rep("#fed976",1),
              rep("#67a9cf",1),
              rep("#a6dba0",2),
              rep("#67a9cf",4),
              rep("#008837",6),
```

```

rep("#67a9cf",1),
rep("#fc8d59",7),
rep("#67a9cf",1),
rep("#008837",5),
rep("#67a9cf",4),
rep("#fed976",2),
rep("#67a9cf",5),
rep("#969696",1))

col <- data.frame(name = c(rep("TPP_Riboswitch_590", time = 83),
                           rep("TPP_Riboswitch_589", time = 83)),
                  position = rep(1:83, time=2),
                  pos_color = c(color_590,color_589))

tpp_seq <- "data/tpp_riboswitch.fasta"
arc_4NYG <- "data/riboswitch_thiamine.txt"
arc_4NYD <- "data/riboswitch_hypoxanthine.txt"
thiamine <- readSSfile(arc_4NYG, type = "Vienna" )
hypoxanthine <- readSSfile(arc_4NYD, type = "Vienna")

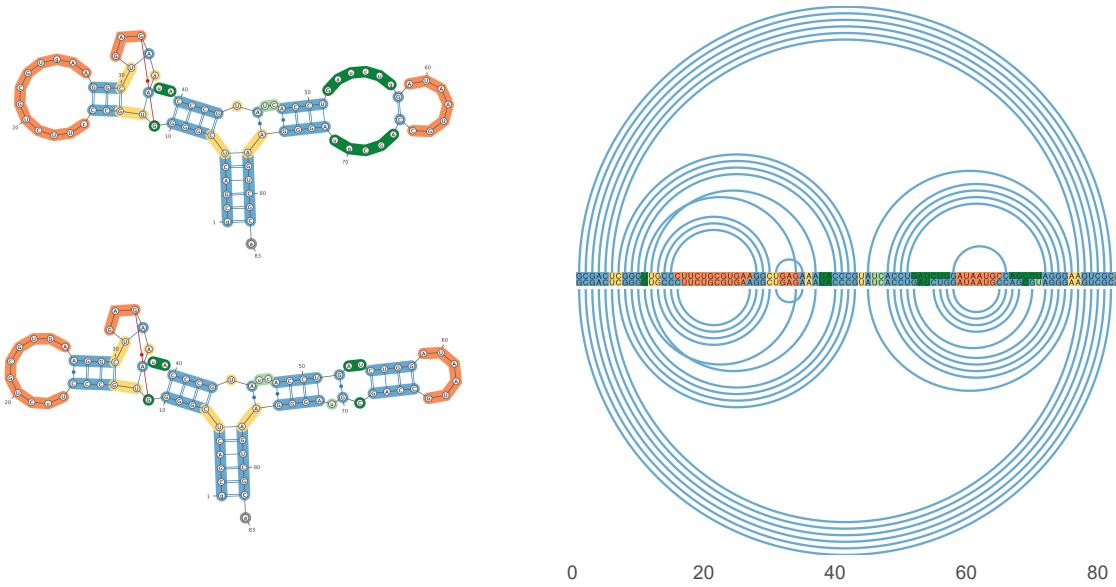
tidy_tpp <- tidy_msa(tpp_seq)
tidy_tpp <- merge(tidy_tpp, col, by = c("name","position"))

p_double_arc <- ggplot() +
  geom_msa(data = tidy_tpp,
            position_color = TRUE,
            seq_name = TRUE,
            border = NA,
            show.legend = F) +
  theme_msa() +
  geom_helix(helix_data = list(known = hypoxanthine,
                               predicted = thiamine),
             color_by = "#67a9cf") +
  theme(axis.text.y = element_blank())

p_loop1 <- image_read_pdf(path = "data/bpRNA_PDB_590_ColorCodedStructures_4NYG.pdf",
                           density = 300)
p_loop2 <- image_read_pdf(path = "data/bpRNA_PDB_589_ColorCodedStructures_4NYD.pdf",
                           density = 300)
q1 <- as.ggplot(p_loop1)
q2 <- as.ggplot(p_loop2)

p3B <- (q1/q2) | p_double_arc
p3B

```



**Fig. 3B| Exploring RNA structural changes by geom\_helix annotation.** Stem-loop diagram and arc diagram of secondary structure changes of TPP-riboswitches after binding to Hypoxanthine and Thiamine respectively. The sequence of the arc plot is the TPP-riboswitch RNA. The arc above the sequence represents the structure of the TPP-riboswitch bound to Thiamine and the bottom arc depicts TPP-riboswitch RNA structural changes binding to Hypoxanthine. The two stem-loop diagrams on the right correspond to the upper and lower arcs respectively. Stem-loop annotation of TPP-riboswitch from bpRNA-1m database (bpRNA ID: bpRNA\_PDB\_589 and bpRNA\_PDB\_590), and also corresponding PDB ID: 4NYD, 4NYG.

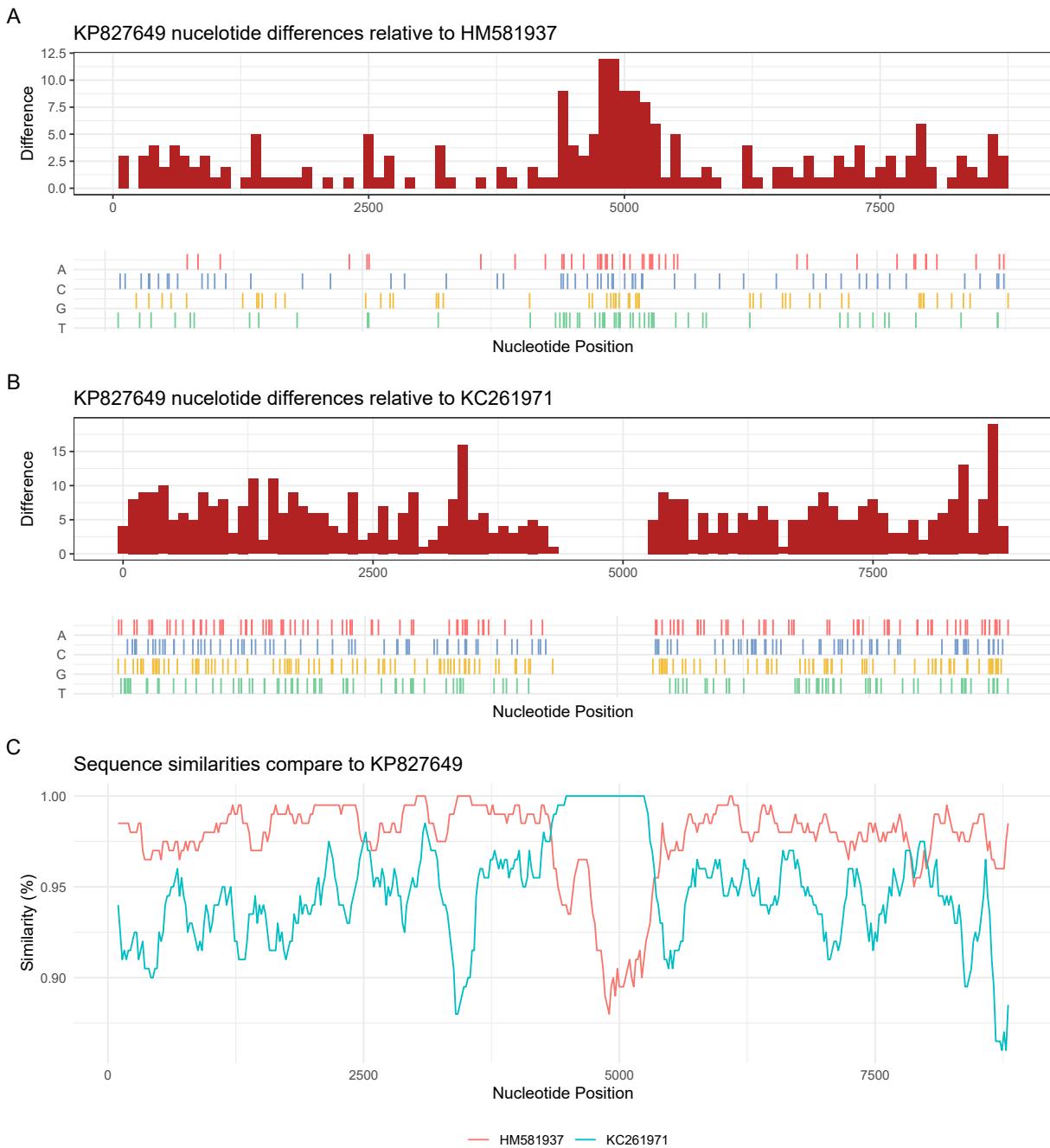
**Fig. 4: Visual exploration for sequence recombination signal (R code)**

```

fas <- c("data/HM_KP.fa", "data/CK_KP.fa")
xx <- lapply(fas, seqdiff)
plts <- lapply(xx, plot, width = 100)
plts[[3]] <- simplot("data/CK_HM_KP.fa", 'KP827649', smooth = FALSE) +
  theme(legend.position = "bottom")

plot_list(gglist=plts, ncol=1, tag_levels = list(c("A", ' ', "B", ' ', "C")))

```

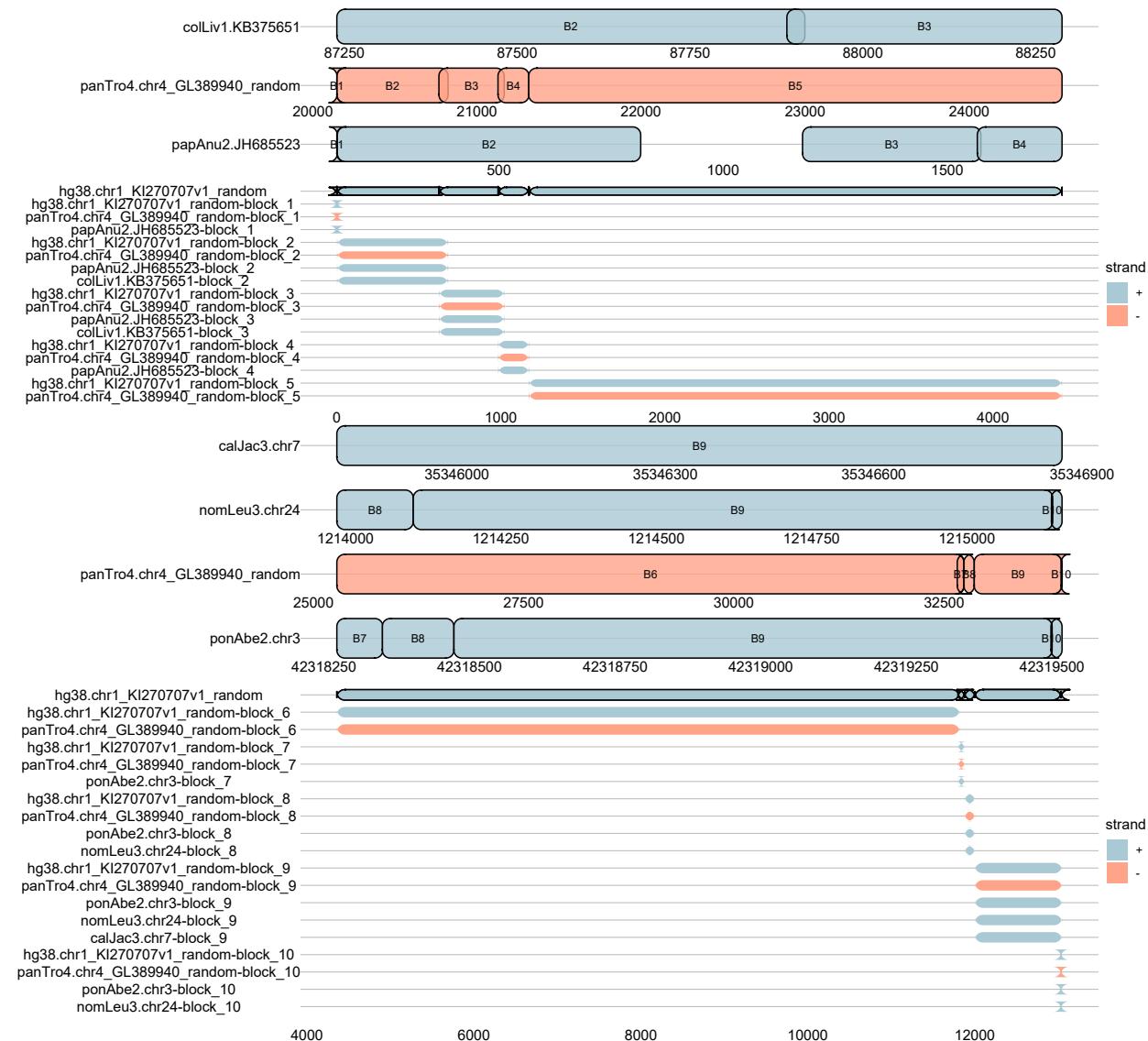


**Fig. 4| Visual exploration for sequence recombination signal.** The nucleotide different plots were generated using the comparison of the KP827649 sequence (Tomato Spotted Wilt Virus large RNA genomic segment) with that of (A) HM581979 sequence (Major parent) and (B) KC261971 sequence (Minor parent). (C) The similarity curves investigate the sequence similarity between the KP827649 sequence and potential parents. Two recombination signals (The intersection of two curves in a similarity plot, the start breakpoint is 4534 and the end breakpoint is 5536) were detected in the TSWV LRNA genomic segment.

**Fig. 5: Visualization of genome alignment in MAF format (R code)**

```
maf <- "data/chr1_KI270707v1_random.txt.maf"
ref = "hg38.chr1_KI270707v1_random"
seq_df <- read_maf(maf)
tidy_df <- tidy_maf_df(seq_df, ref = ref)

ggmaf(data = tidy_df,
      ref = ref,
      block_start = 1,
      block_end = 10,
      facet_field = 5,
      facet_heights = c(0.4,0.6))
```



shows the original genome coordinates and the lower panel shows genome alignments, with colors indicating positive and negative strands. In the genome alignments panel, the first genome with black border represents the reference genome hg38.chr1 and subsequences in each blocks does not show border color. In each block, the first sequence is the reference genome fragment, followed by other species fragments. Fragments in each block can correspond to labels B1-B10 on the original genome coordinates panel.

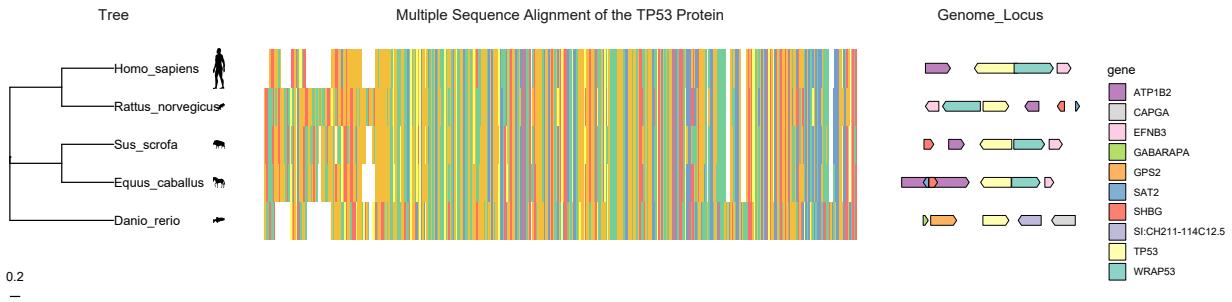
**Fig. 6A: An example of graphics combination (R code)**

```
##Fig 6A tree + msa + genes locus
dat <- read_aa(tp53_sequences, format = "fasta") %>% phyDat(type = "AA", levels = NULL)
tree <- dist.ml(dat, model = "JTT") %>% bionj()
dd <- ggimage::phylopic_uid(tree$tip.label)

p_tp53 <- ggtree(tree, branch.length = 'none') %<+% dd +
  geom_tiplab(aes(image=uid), geom = "phylopic", offset =1.9) +
  geom_tiplab(aes(label=label)) +
  geom_treescale(x = 0,y = -1)
#msa
data_53 <- readAAMultipleAlignment(tp53_sequences) %>% tidy_msa()
#gene maps
TP53_arrow <- readxl::read_xlsx(tp53_genes)
TP53_arrow$direction <- 1
TP53_arrow[TP53_arrow$strand == "reverse","direction"] <- -1

#color
mapping = aes(xmin = start, xmax = end, fill = gene, forward = direction)
my_pal <- colorRampPalette(rev(brewer.pal(n = 10, name = "Set3")))

#tree + gene maps + msa
p6a <- p_tp53 + xlim_tree(4) +
  geom_facet(geom = geom_msa, data = data_53,
             panel = 'Multiple Sequence Alignment of the TP53 Protein', font = NULL,
             border = NA) +
  new_scale_fill() +
  scale_fill_manual(values = my_pal(10)) +
  geom_facet(geom = geom_motif,
             mapping = mapping, data = TP53_arrow,
             panel = 'Genome_Locus', on = 'TP53',
             arrowhead_height = unit(3, "mm"),
             arrowhead_width = unit(1, "mm")) +
  theme(strip.background=element_blank(),
        strip.text = element_text(size = 13))
p6A <- facet_widths(p6a, c(Tree = 0.35, Genome_Locus = 0.3))
p6A
```



**Fig. 6A| An example of graphics combination.** The MSA-tree panel in conjunction with the external genome locus data set. Comparative genome locus structure (genome\_Locus panel), sequence alignment of TP53 protein (the middle panel), and the corresponding phylogenetic tree (Tree panel) among six species. The local genome map shows the 30000 sites around the TP53 gene, and the phylogenetic tree that represents evolutionary relationships inferred from TP53 protein sequences using the Neighbor-Joining method based on the evolutionary distances of the JTT matrix-based method.

**Fig. 6B: Another example of graphics combination (R code)**

```

##Fig 6B tree + msa + 2boxplot
seq <- readDNAStringSet("data//btuR.fa")
aln <- tidy_msa(seq)
btuR_tree <- read.tree("data/btuR.nwk")
meta_dat <- read.csv("data/meta_data_47.csv")

#Pathotype_fill_colors
Pathotype_cols <- RColorBrewer::brewer.pal(7, "Set3")
names(Pathotype_cols) <- meta_dat$Pathotypes %>% factor %>% levels

####tree OTU
Phylo_group <- list(A= meta_dat$Lineage[meta_dat$Phylogroup == "A"]%>% unique,
                      B1=meta_dat$Lineage[meta_dat$Phylogroup == "B1"]%>% unique,
                      B2=meta_dat$Lineage[meta_dat$Phylogroup == "B2"]%>% unique,
                      C=meta_dat$Lineage[meta_dat$Phylogroup == "C"]%>% unique,
                      D =meta_dat$Lineage[meta_dat$Phylogroup == "D"]%>% unique,
                      E =meta_dat$Lineage[meta_dat$Phylogroup == "E"]%>% unique,
                      `F`=meta_dat$Lineage[meta_dat$Phylogroup == "F"]%>% unique,
                      Shigella=meta_dat$Lineage[meta_dat$Phylogroup == "Shigella"]%>% unique)

Phylo_cols <- RColorBrewer::brewer.pal(8, "Dark2")
names(Phylo_cols) <- names(Phylo_group)

## plot tree
p_btuR_tree <- ggtree(btuR_tree) + geom_tiplab(align = TRUE)
p_btuR_tree <- groupOTU(p_btuR_tree ,Phylo_group)+aes(color=group) +
  scale_color_manual(values = c(Phylo_cols, "black"), na.value = "black", name = "Lineage",
                     breaks = c("A", "B1", "B2", "C", "D", "E", "F", "Shigella"), guide="none")

p_btuR_tree <- p_btuR_tree +
  geom_strip('L29', 'L20', barsize=2, color=Phylo_cols[["B2"]],
             label="B2", offset = .01, offset.text = 0.0015) +
  geom_strip('L28','L29', barsize=2, color=Phylo_cols[["A"]],
```

```

        label="A", offset = .01, offset.text = 0.0015) +
geom_strip('L15','L28', barsize=2, color=Phylo_cols[["B1"]],
           label="B1", offset = .01, offset.text = 0.0015) +
geom_strip('L45','L15', barsize=2, color=Phylo_cols[["Shigella"]],
           label="S.", offset = .01, offset.text = 0.0015) +
geom_strip('L36','L45', barsize=2, color=Phylo_cols[["B1"]],
           label="B1", offset = .01, offset.text = 0.0015) +
geom_strip('L30','L36', barsize=2, color=Phylo_cols[["Shigella"]],
           label="S.", offset = .01, offset.text = 0.0015) +
geom_strip('L39','L30', barsize=2, color=Phylo_cols[["B1"]],
           label="B1", offset = .01, offset.text = 0.0015) +
geom_strip('L40','L39', barsize=2, color=Phylo_cols[["C"]],
           label="C", offset = .01, offset.text = 0.0015) +
geom_strip('L48','L40', barsize=2, color=Phylo_cols[["B1"]],
           label="B1", offset = .01, offset.text = 0.0015) +
geom_strip('L10','L48', barsize=2, color=Phylo_cols[["E"]],
           label="E", offset = .01, offset.text = 0.0015) +
geom_strip('L37','L10', barsize=2, color=Phylo_cols[["D"]],
           label="D", offset = .01, offset.text = 0.0015) +
geom_strip('L33','L37', barsize=2, color=Phylo_cols[["F"]],
           label="F", offset = .01, offset.text = 0.0015) +
geom_strip('L1','L33', barsize=2, color=Phylo_cols[["E"]],
           label="E", offset = .01, offset.text = 0.0015)

##tree + meta data boxplots
p6B <- p_btuR_tree +
  geom_treescale(x = 0,y = -1) +
  geom_fruit(data = aln,
             geom = geom_msa,
             end = 200,
             font = NULL,
             color = "Chemistry_NT",
             border = NA,
             consensus_views = TRUE,
             ref = "L38",
             pwidth = 3.5,
             offset = 0.3,
             axis.params = list(title = "Multiple Sequence Alignment of the btuR Gene",
                                title.height = 0.05,
                                title.size = 4.5,
                                axis = "x",
                                vjust = 1.1,
                                text.size = 3,
                                line.size = 1,
                                line.color = "black")) +
  new_scale_fill() +
  geom_fruit(mapping = aes(x = AMR_genes, y = Lineage, fill = MDR),
             data = meta_dat,
             geom = geom_boxplot,
             outlier.size = 0.5,
             pwidth=1,
             offset = 0.1,
             axis.params = list(title = "Antimicrobial Classes",

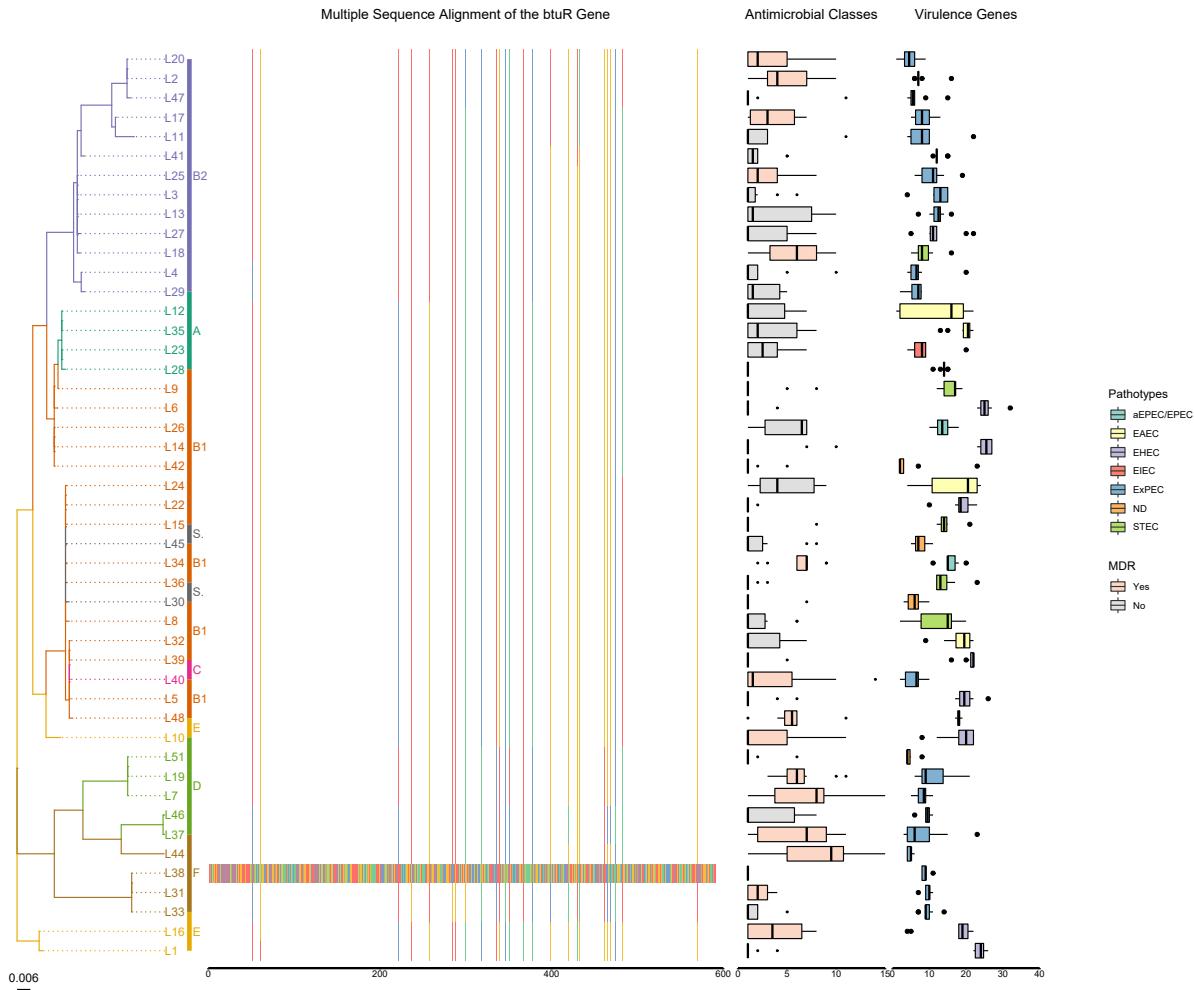
```

```

        title.height = 0.05,
        title.size = 4.5,
        axis = "x",
        vjust = 1.1,
        text.size = 3,
        line.size = 1,
        line.color = "black")) +
scale_fill_manual(values=c("Yes" = "#fcd7c5", "No" = "#dfdfdf")) +
new_scale_fill() +
geom_fruit(mapping = aes(x = virulence_genes, y = Lineage, fill = Pathotypes),
           data = meta_dat,
           geom = geom_boxplot,
           pwidth=1,
           offset = 0.05,
           axis.params = list(title = "Virulence Genes",
                               title.height = 0.05,
                               title.size = 4.5,
                               axis = "x",
                               vjust = 1.1,
                               text.size = 3,
                               line.size = 1,
                               line.color = "black"))+
scale_fill_manual(values = Pathotype_cols)

```

p6B



**Fig. 6B| Another example of graphics combination.** The summarized visualization for different *E.coli* lineages. The alignment of the *btuR* gene was selected to generate a phylogenetic tree which was stained according to the corresponding lineages. The boxplots represent antimicrobial resistance (AMR) and virulence profiles of the lineages. A number of antimicrobial classes and virulence genes per isolate are colored by multidrug-resistant (MDR) classes and the most prevalent predicted pathotype in the lineage.

Here is the output of sessionInfo() on the system on which this document was compiled:

```
## R version 4.1.1 (2021-08-10)
## Platform: x86_64-w64-mingw32/x64 (64-bit)
## Running under: Windows 10 x64 (build 19044)
##
## Matrix products: default
##
## locale:
## [1] LC_COLLATE=Chinese (Simplified)_China.936
## [2] LC_CTYPE=Chinese (Simplified)_China.936
## [3] LC_MONETARY=Chinese (Simplified)_China.936
## [4] LC_NUMERIC=C
## [5] LC_TIME=Chinese (Simplified)_China.936
##
## attached base packages:
## [1] stats4      stats       graphics    grDevices   utils       datasets    methods
```

```

## [8] base
##
## other attached packages:
## [1] kableExtra_1.3.4      treeio_1.18.0       magick_2.7.3
## [4] aplot_0.1.1          ggplotify_0.1.0    patchwork_1.1.1
## [7] RColorBrewer_1.1-2   phangorn_2.7.1     ggtreeExtra_1.4.0
## [10] dplyr_1.0.7          ggnewscale_0.4.5  Biostrings_2.62.0
## [13] GenomeInfoDb_1.30.0  XVector_0.34.0    IRanges_2.28.0
## [16] S4Vectors_0.32.2    BiocGenerics_0.40.0 ape_5.5
## [19] ggggenes_0.4.1      ggtree_3.3.1.901  ggplot2_3.3.5
## [22] ggmsa_1.1.7        knitr_1.36

##
## loaded via a namespace (and not attached):
## [1] nlme_3.1-153           bitops_1.0-7       webshot_0.5.2
## [4] ash_1.0-15             httr_1.4.2         tools_4.1.1
## [7] utf8_1.2.2            R6_2.5.1          KernSmooth_2.23-20
## [10] lazyeval_0.2.2        colorspace_2.0-2  withr_2.4.2
## [13] tidyselect_1.1.1      ggalt_0.4.0       curl_4.3.2
## [16] compiler_4.1.1        extrafontdb_1.0   rvest_1.0.2
## [19] statebins_1.4.0       xml2_1.3.2       labeling_0.4.2
## [22] scales_1.1.1         proj4_1.0-10.1   quadprog_1.5-8
## [25] askpass_1.1           systemfonts_1.0.3 stringr_1.4.0
## [28] digest_0.6.28         yulab.utils_0.0.4 R4RNA_1.22.0
## [31] rmarkdown_2.11         svglite_2.0.0     pkgconfig_2.0.3
## [34] htmltools_0.5.2       extrafont_0.17   fastmap_1.1.0
## [37] highr_0.9             maps_3.4.0        readxl_1.3.1
## [40] rlang_0.4.11          rstudioapi_0.13  gridGraphics_0.5-1
## [43] farver_2.1.0          generics_0.1.1   jsonlite_1.7.2
## [46] RCurl_1.98-1.5        magrittr_2.0.1   GenomeInfoDbData_1.2.7
## [49] Matrix_1.3-4          Rcpp_1.0.8.3     munsell_0.5.0
## [52] fansi_0.5.0           ggfittext_0.9.1  lifecycle_1.0.1
## [55] stringi_1.7.5         yaml_2.2.1       seqmagick_0.1.5
## [58] MASS_7.3-54            zlibbioc_1.40.0  grid_4.1.1
## [61] parallel_4.1.1        crayon_1.4.2     lattice_0.20-45
## [64] pillar_1.6.4          igraph_1.2.7     codetools_0.2-18
## [67] fastmatch_1.1-3       glue_1.4.2       evaluate_0.14
## [70] ggimage_0.3.0          pdftools_3.1.1   qpdf_1.1
## [73] ggfunk_0.0.6           vctrs_0.3.8      tweenr_1.0.2
## [76] cellranger_1.1.0      Rttf2pt1_1.3.9   gtable_0.3.0
## [79] purrr_0.3.4            polyclip_1.10-0  tidyR_1.1.4
## [82] xfun_0.26              ggforce_0.3.3    rsvg_2.2.0
## [85] tidytree_0.3.9         viridisLite_0.4.0 tibble_3.1.5
## [88] ellipsis_0.3.2

```

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