R script for Comparative phylogenomic analyses of SNP versus full locus datasets: insights and recommendations for researchers

Prepared by Jacob S. Suissa 2023-08-30

Overview

This is an R Markdown document for the Cornell SIPS Plant Biology Phylogenetic Journal Club project exploring how SNP vs. locus data and missing data affect phylogeny topology, branch length, nodal support, node dating, and downstream phylogenetic comparative methods.

Load libraries

First load the libraries

library(tidyverse)
library(ape)
library(phytools)
library(geiger)
library(ggdist)
library(ggpubr)
library(ggsci)
library(treeio)
library(RColorBrewer)

Import data

Read in all the trees and make sure to subset the locus and the SNPS files

```
alltreeFiles <-
  list.files(
    path = path,
    pattern = "*all.raxml.support",
    full.names = TRUE,
    recursive = FALSE
  )
alltree_list <- list()</pre>
for (i in 1:length(alltreeFiles)) {
  a <-
    paste(gsub("\\..*", "", basename(alltreeFiles[i])), ".tree", sep = "")
 tree <- read.tree(alltreeFiles[i])</pre>
  alltree_list[[a]] <- tree</pre>
}
#Same thing as above but for SNPS
snptreeFiles <-</pre>
  list.files(
    path = path,
    pattern = "*variant.raxml.support",
    full.names = TRUE,
    recursive = FALSE
  )
snptree list <- list()</pre>
for (i in 1:length(snptreeFiles)) {
    paste(gsub("\\..*", "", basename(snptreeFiles[i])), ".tree", sep = "")
 tree <- read.tree(snptreeFiles[i])</pre>
  snptree_list[[a]] <- tree</pre>
}
```

Analyze the data

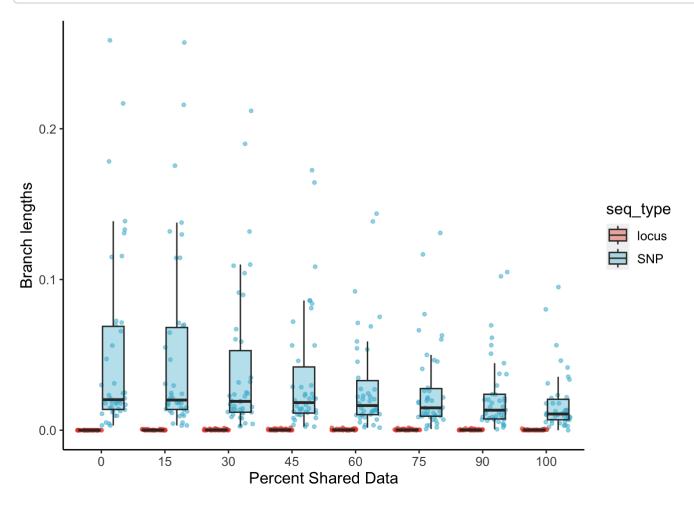
Extract all of the edge lengths and node label data from the trees

First make simple boxplots. Start with a boxplots of edgelength

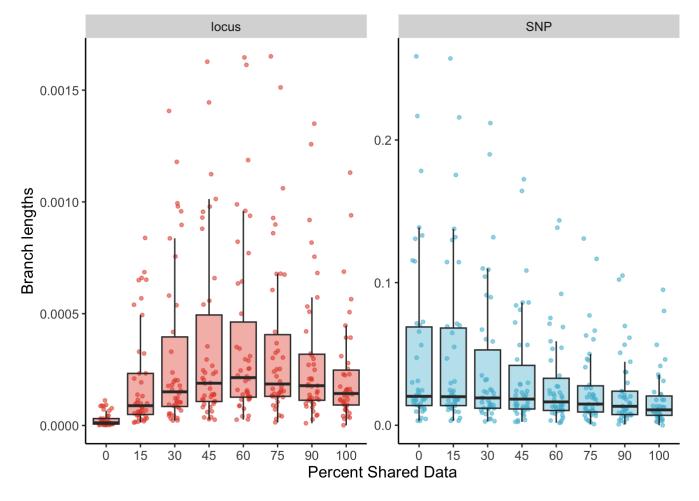
```
Order<-c( "0", "15", "30", "45", "60", "75", "90", "100")
```

edgePlot<- ggplot(totTree1.dat, aes(y=edgeLength, x=factor(filter, levels = Order))) +ge
om_point(aes(color=seq_type, alpha=0.5), size= 1, show.legend = FALSE, position=positio
n_jitterdodge())+ geom_boxplot(aes(fill=seq_type), alpha=0.4, outlier.colour = NA)+ xlab
("Percent Shared Data")+ ylab("Branch lengths")+ scale_fill_manual(values= pal_npg("nr
c", alpha = 0.7)(2))+ scale_color_manual(values= pal_npg("nrc", alpha = 0.6)(2)) + theme
(text = element_text(size=12)) + theme(panel.grid.major = element_blank(), panel.grid.mi
nor = element_blank(), panel.background = element_blank(), axis.line = element_line(colou
r = "black"))</pre>

edgePlot



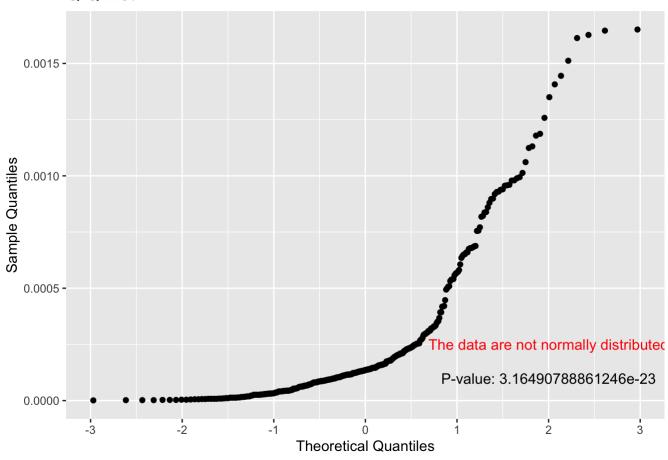
Plot branch lengths differently from edge lengths



Test the normality of both locus branch length datasets

```
#Locus data
R script for Comparative phylogenomic analyses of SNP versus full locus datasets: insights and recommendations for researchers
  filter(seq type == "locus", .preserve = TRUE)
# Shapiro-Wilk Test for Normality
shapiro.test result <- shapiro.test(loc.dat$edgeLength)</pre>
shapiro_p_value <- shapiro.test_result$p.value</pre>
# Q-Q Plot
qq plot <- qqnorm(loc.dat$edgeLength, plot.it = FALSE)</pre>
qq_plot_data <- data.frame(Theoretical = qq_plot$x, Sample = qq_plot$y)
qq_plot <- ggplot(qq_plot_data, aes(Theoretical, Sample)) +</pre>
  geom point() +
  labs(x = "Theoretical Quantiles", y = "Sample Quantiles") +
  ggtitle("Q-Q Plot") +
  annotate("text", x = 2, y = 0.0001, label = paste("P-value:", shapiro_p_value)) +
  annotate("text", x = 2, y = 0.00025, label = ifelse(shapiro p value < 0.01, "The data
are not normally distributed", "The data are normally distributed"), color = ifelse(shap
iro p value < 0.01, "red", "black"))</pre>
# Display the Q-Q plot with annotations
print(qq plot)
```

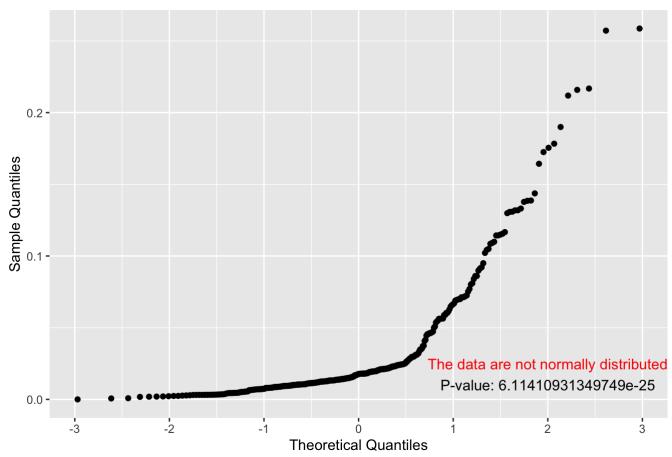




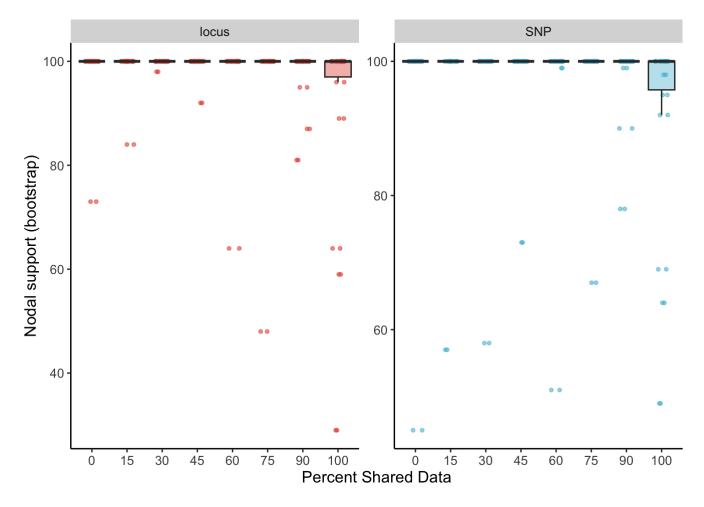
Test the normality of both snp branch length datasets

```
#SNP data
SNP.dat <- totTree1.dat%>%
  filter(seq type == "SNP", .preserve = TRUE)
# Shapiro-Wilk Test for Normality
shapiro.test result <- shapiro.test(SNP.dat$edgeLength)</pre>
shapiro_p_value <- shapiro.test_result$p.value</pre>
# Q-Q Plot
qq plot <- qqnorm(SNP.dat$edgeLength, plot.it = FALSE)</pre>
qq_plot_data <- data.frame(Theoretical = qq_plot$x, Sample = qq_plot$y)
qq_plot <- ggplot(qq_plot_data, aes(Theoretical, Sample)) +</pre>
  geom point() +
 labs(x = "Theoretical Quantiles", y = "Sample Quantiles") +
  ggtitle("Q-Q Plot") +
  annotate("text", x = 2, y = 0.01, label = paste("P-value:", shapiro_p_value)) +
  annotate("text", x = 2, y = 0.025, label = ifelse(shapiro p value < 0.01, "The data ar
e not normally distributed", "The data are normally distributed"), color = ifelse(shapir
o p value < 0.01, "red", "black"))</pre>
# Display the Q-Q plot with annotations
print(qq plot)
```





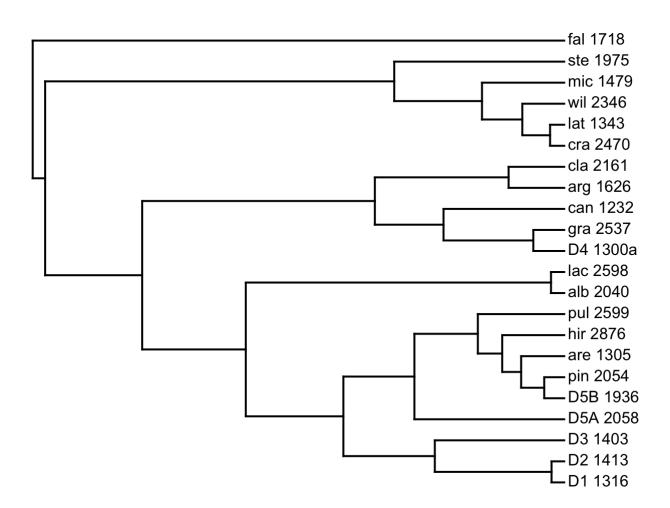
Make boxplots of nodal support



Explore relative branch lengths across each raxml tree. Bring in the BEAST data and explore patterns of node age across the different data and filtering types

Plot one of the beast trees to visualize

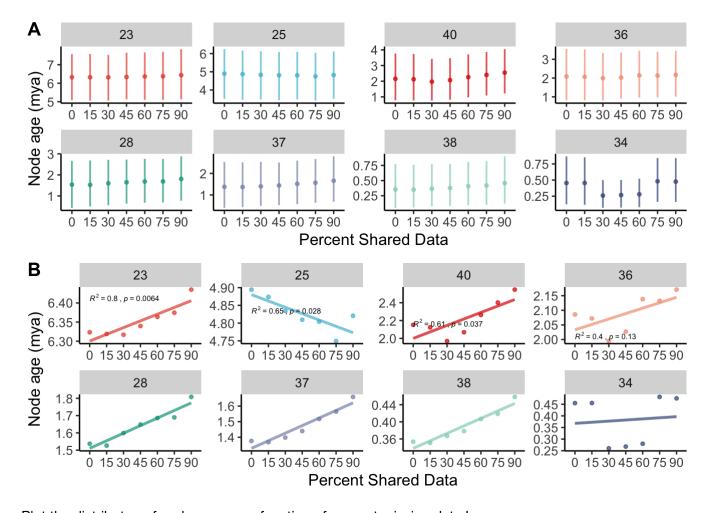
```
plotTree(allBeastTree_list[["30P_all_MCC.tre"]]@phylo)
```



Extract all of the ages for the tree

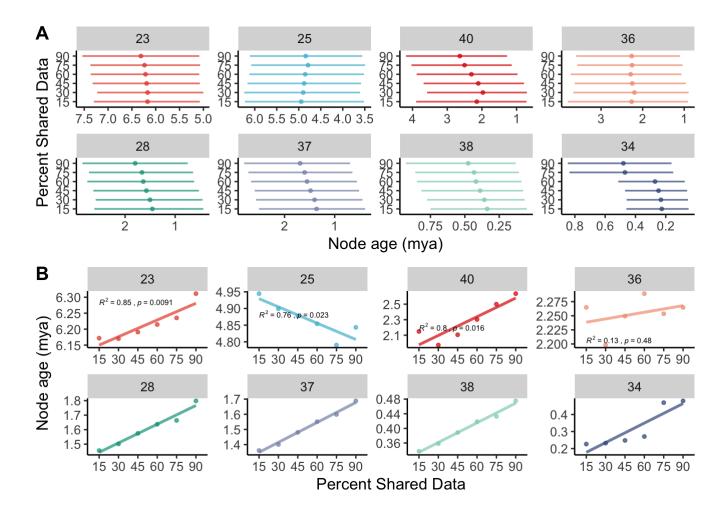
Make a new columns where we specifically denote the percent missing data, and whether its variant or full locus Plot the distributon of node ages as a function of percent missing data SNPS

```
snp_only <- totBeastTree1.dat%>%
 filter(seq type == "SNP", .preserve = TRUE )%>%
 filter(!is.na(early), .preserve = TRUE)%>%
 mutate(node = as.numeric(node))%>%
 filter(node == "23" | node == "25" | node == "40" | node == "36" | node == "28" | node ==
"37" | node == "38" | node == "34" )%>%#Select a 8 nodes
 mutate(across(node, factor, levels=c("23" , "25" , "40" , "36" , "28" , "37"
     , "34")))
 m <-ggplot(snp_only, aes(x= filtered, y=CAheight_mean, group = node )) +geom_point(aes</pre>
(color= as.character(node) ), position = position_jitter(height = 0, width = 0, seed =
1) , size=1) + facet_wrap(~node, nrow = 2, scales = "free" )+ guides(color="none") +geo
m segment( aes(y=early, yend=late, x=filtered, xend= filtered, color= as.character(nod
e)), position = position_jitter(height = 0, width = 0, seed = 1), size= 0.6) + guides(co
lor=FALSE) + ylab("Node age (mya)")+ xlab("Percent Shared Data")+ scale_color_manual(val
ues= pal npg("nrc", alpha = 0.7)(8))+ theme(text = element text(size=12)) + theme(pane
l.grid.major = element_blank(), panel.grid.minor = element_blank(),panel.background = el
ement_blank(), axis.line = element_line(colour = "black"))+facet_wrap(~node, nrow = 2, s
cales = "free" )
#Then plot just the mean and run a linear regression!
 n <-ggplot(snp only, aes(x= filtered, y=CAheight mean, group = node )) +geom point(aes
(color= as.character(node) ), position = position jitter(height = 0, width = 0, seed =
1) , size=1) + facet wrap(~node, nrow = 2, scales = "free" )+ guides(color="none")+ geo
m smooth(method = "lm", aes(color = as.character(node) ), se =FALSE )+
bel = paste(..rr.label.., ..p.label.., sep = "~`,`~")), size=2) + guides(color=FALSE) +
ylab("Node age (mya)")+ xlab("Percent Shared Data")+ scale color manual(values= pal npg
("nrc", alpha = 0.7)(8))+ theme(text = element_text(size=12)) + theme(panel.grid.major
= element blank(), panel.grid.minor = element blank(),panel.background = element blank
(), axis.line = element line(colour = "black"))+facet wrap(~node, nrow = 2, scales = "fr
ee")
ggarrange(m, n ,
         labels = c("A", "B"),
          nrow = 2, widths = 8, heights = 5)
```



Plot the distributon of node ages as a function of percent missing data Locus

```
all_only <- totBeastTree1.dat%>%
 filter(seq type == "locus", .preserve = TRUE )%>%
 filter(!is.na(early), .preserve = TRUE)%>%
 mutate(node = as.numeric(node))%>%
 filter(node == "23" | node == "25" | node == "40" | node == "36" | node == "28" | node ==
"37" | node == "38" | node == "34" )%>%#Select a 8 nodes
 mutate(across(node, factor, levels=c("23" , "25" , "40" , "36" , "28" , "37"
     , "34")))
 1<- ggplot(all_only, aes(y= filtered, x=CAheight_mean )) +geom_point(aes(color= as.cha</pre>
racter(node)),
                 position = position_jitter(height = 0, width = 0, seed = 1) , size=1)
+geom_segment( aes(x=early, xend=late, y=filtered, yend= filtered, color= as.character(n
ode)), position = position jitter(height = 0, width = 0, seed = 1), size= 0.6)+scale x r
everse() + guides(color=FALSE) + xlab("Node age (mya)")+ ylab("Percent Shared Data")+ sc
ale_color_manual(values= pal_npg("nrc", alpha = 0.7)(8))+ facet_wrap(~node, nrow = 2, sc
ales = "free" )+ theme(text = element text(size=12)) + theme(panel.grid.major = element
blank(), panel.grid.minor = element_blank(),panel.background = element_blank(), axis.lin
e = element line(colour = "black"))
       ggplot(all_only, aes(x= filtered, y=CAheight_mean, group = node )) +geom_point(a
es(color= as.character(node) ), position = position_jitter(height = 0, width = 0, seed =
1) , size=1) + facet_wrap(~node, nrow = 2, scales = "free" )+ guides(color="none")+ geo
m smooth(method = "lm", aes(color = as.character(node) ), se =FALSE )+
bel = paste(..rr.label.., ..p.label.., sep = "~`,`~")), size=2) + guides(color=FALSE) +
ylab("Node age (mya)")+ xlab("Percent Shared Data")+ scale color manual(values= pal npg
("nrc", alpha = 0.7)(8))+ theme(text = element text(size=12)) + theme(panel.grid.major
= element_blank(), panel.grid.minor = element_blank(),panel.background = element_blank
(), axis.line = element line(colour = "black"))+facet wrap(~node, nrow = 2, scales = "fr
ee")
ggarrange(1, p ,
         labels = c("A", "B"),
          nrow =2, widths = 8, heights = 5)
```



Phylogenetic comparative methods

Use trees from the 45% filtered trees to explore the impact of SNP vs. locus datasets and the use of phylograms vs. chronograms in PCMs.

First create a funciton that trims data and puts it into the right format for stochastic character mapping

Run stochastic character mapping on the randomized data

```
#import data
snp_mock.dat<-read.csv(path2)

#bring in the three trees of interest

locus_45raxml.tre<-read.tree(path3)
snp_45raxml.tre<-read.tree(path4)

locus_45beast.tre<-read.beast(path5)

locus_45beast.tre<- as.phylo(locus_45beast.tre)
snp_45beast.tre<-read.beast(path6)
snp_45beast.tre<-as.phylo(snp_45beast.tre)

trees.list <- c(locus_45raxml.tre,snp_45raxml.tre,locus_45beast.tre, snp_45beast.tre )

trees.list.name <- c("locus_45raxml","snp_45raxml","locus_45beast", "snp_45beast" )</pre>
```

Loop through each tree and run a stochastic character map and density map

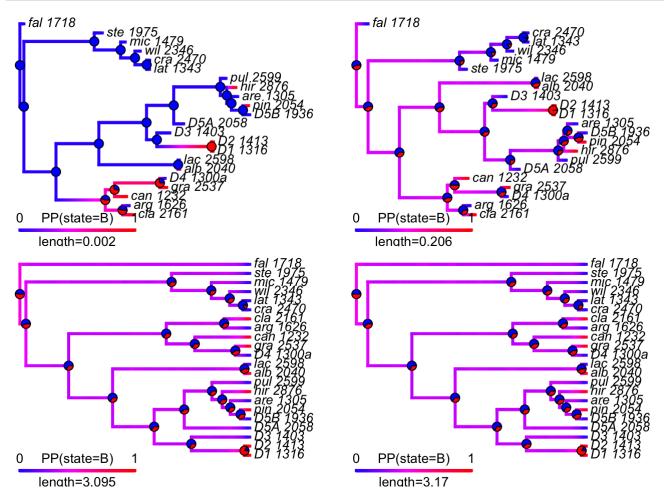
```
mockdat.mode<-setNames(snp mock.dat$State,snp mock.dat$Tip)</pre>
mockdat.mode<-as.factor(mockdat.mode) #This is essential now for some rason...
simmap list <- list()</pre>
dmap list <-list()</pre>
summary list<-list()</pre>
for (i in 1:length(trees.list)) {
  a <-make.simmap(tree = trees.list[i], x =mockdat.mode, model = "ARD", nsim = 1000, Q=
"empirical" )
  tree file <- paste0( trees.list.name[i])</pre>
  simmap list[[tree file]] <- a</pre>
  simmap list[[i]] <- a</pre>
  b <- densityMap(a,plot=FALSE)</pre>
  dmap_list[[tree_file]] <- b</pre>
  c<- summary(a)</pre>
  summary list[[tree file]]<-c</pre>
}
```

```
## make.simmap is sampling character histories conditioned on
## the transition matrix
##
## O =
##
         Α
## A -0.4719 0.4719
## B 0.4719 -0.4719
## (estimated using likelihood);
## and (mean) root node prior probabilities
## pi =
## A
## 0.5 0.5
## make.simmap is sampling character histories conditioned on
## the transition matrix
##
## Q =
##
                       В
## A -25.55625 25.55625
## B 41.28740 -41.28740
## (estimated using likelihood);
## and (mean) root node prior probabilities
## pi =
##
   Α
## 0.5 0.5
## make.simmap is sampling character histories conditioned on
## the transition matrix
##
## Q =
                       В
## A -2.240632 2.240632
## B 4.063175 -4.063175
## (estimated using likelihood);
## and (mean) root node prior probabilities
## pi =
##
   Α
## 0.5 0.5
## make.simmap is sampling character histories conditioned on
## the transition matrix
##
## Q =
##
                       В
            Α
## A -2.352308 2.352308
## B 4.300970 -4.300970
## (estimated using likelihood);
## and (mean) root node prior probabilities
## pi =
## A
## 0.5 0.5
```

```
par(mfrow = c(2, 2)) # Set up a 2x2 grid layout for the plots

for (i in 1:length(dmap_list)) {
    # Extract the file name
    file_name <- names(dmap_list)[i]

plot(dmap_list[[file_name]])+ nodelabels(node = as.numeric(row.names(summary_list[[file_name]]$ace)), pie=summary_list[[file_name]]$ace, piecol = c("blue", "red"), cex=0.7 )
}</pre>
```



Run ancestral character estimation on a continuous character

Simulate a continuous character based on brownian motion on the 45 filtered SNP Beast tree

```
# Set the parameters for the Brownian motion simulation
sigma <- 1  # Trait evolutionary rate
nsim <- 1  # Number of simulations

# Simulate the continuous trait using Brownian motion
sim_data <- fastBM(snp_45beast.tre, sigma = sigma, nsim = nsim)</pre>
```

Loop through and make cont maps

```
#Make anc recons TEST
fitBM<-anc.ML(snp_45beast.tre,sim_data,model="BM")</pre>
BM<-contMap(snp_45beast.tre,sim_data, method="user",anc.states=fitBM$ace,plot=FALSE)
#Make them for all the trees
fit_list <-list()</pre>
contmap_list <- list()</pre>
for (i in 1:length(trees.list)) {
  a <-anc.ML(tree = trees.list[[i]], x =sim_data, model = "BM" )</pre>
  tree file <- paste0( trees.list.name[i])</pre>
  fit_list[[tree_file]] <- a</pre>
  fit_list[[i]] <- a</pre>
  b<-contMap(trees.list[[i]],sim_data, method="user",anc.states=a$ace,plot=FALSE)
  b <-setMap(b, brewer.pal(11, "Spectral"))</pre>
  contmap_list[[tree_file]] <- b</pre>
}
```

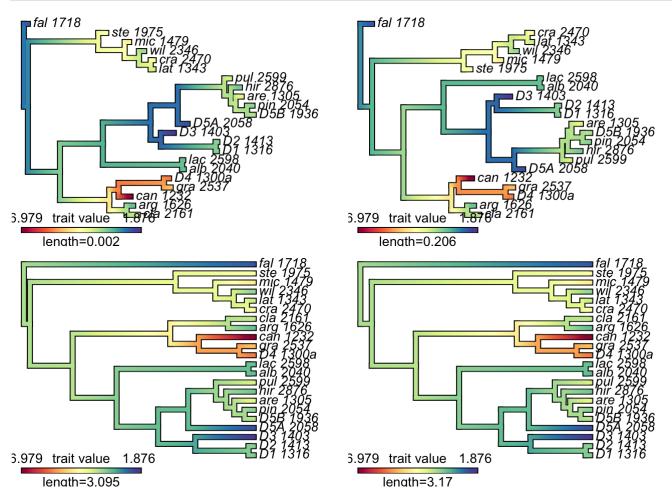
Plot the contmaps and save everything

```
par(mfrow = c(2, 2)) # Set up a 2x2 grid layout for the plots

for (i in 1:length(dmap_list)) {

    # Extract the file name
    file_name <- names(contmap_list)[i]

    plot(contmap_list[[file_name]])
}</pre>
```



#Done! If you. have any questions please email Jacob Suissa at jsuissa@utk.edu (mailto:jsuissa@utk.edu)