PCA_analysis

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First we load the required packages including the mulTree package which will allow us to read back in the relavent information from the MCMCglmm models we ran previously. We will also require the paran package for the PCA test, caper, phytools and MCMCglmm to handle the phylogeny objects and related functions and SIBER and ggplot2 to creat the ellipses and plot the overlaps. As in previous scripts we will also use some costume functions.

```
library(caper)
## Loading required package: ape
## Loading required package: MASS
## Loading required package: mvtnorm
library(phytools)
## Loading required package: maps
## Loading required package: rgl
library(MCMCglmm)
## Loading required package: Matrix
## Attaching package: 'Matrix'
## The following object is masked from 'package:phytools':
##
##
       expm
## Loading required package: coda
library(mulTree)
## Loading required package: hdrcde
## This is hdrcde 3.3
## Loading required package: snow
library(paran)
library(SIBER)
library(ggplot2)
source("Demography_functions.R")
## Welcome to popdemo! This is version 1.3-0
## Use ?popdemo for an intro, or browseVignettes('popdemo') for vignettes
## Citation for popdemo is here: doi.org/10.1111/j.2041-210X.2012.00222.x
## Development and legacy versions are here: github.com/iainmstott/popdemo
```

Now we upload the previous data including both the life history metrics we calulated and also the body size, IUCN and other data.

Next we upload the matching distribution of phylogenies calculated from the previous Phylogeny_construction and Pop_metric_calulation scripts.

```
axis_trees <- read.tree("axis_analysis_phylo.tre")</pre>
```

Log10 the non index based metrics

And mean center the data

We also need to make a multree object that holds both the data and the multiphylo object and from which we will calulate the residuals from each of the modeel

Read back in the the MCMCglmm model outputs

For each of the life history metrics we read back in the information we need from the 100 MCMCglmm models we ran. We will then calculate the residuals form these which will be then used for the PCA analysis. We will also look at the random terms of phylogentic effect, population level variance (species) and the residual term.

Age at first reproduction

```
La_models <- read.mulTree("la_run")</pre>
summary(La_models)
                          Estimates(mode hdr) lower.CI(2.5) lower.CI(25)
                                    0.19747134 -1.50849226 -0.36692148
## (Intercept)
## mass_g
                                    0.59885721 0.38364820 0.52569656
## matrix_size
                                    0.06146852 -0.01051303 0.03715236
## phylogenetic.variance
                                    2.03307751 0.92585738
                                                                1.57129554
## residual.variance
                                    0.29103448
                                                  0.17437952
                                                                0.24779145
##
                          upper.CI(75) upper.CI(97.5)
## (Intercept)
                            0.76238085
                                            1.9104997
                            0.67352076
                                             0.8166299
## mass_g
## matrix size
                            0.08761325
                                             0.1367536
## phylogenetic.variance 2.54291734
                                             3.8375481
## residual.variance
                            0.34484428
                                             0.4648104
## attr(,"class")
## [1] "matrix" "mulTree"
Now we calculate the proportion of variance between phylogenetic, population and residual variance.
la_var <- read.mulTree("la_run", extract = "VCV")</pre>
#phylogenetic signal
la_phlyo <- list()</pre>
#population level variation
la_spec <- list()</pre>
#residual
la_unit <- list()</pre>
#extrace the random terms from across the models
for(i in 1:length(names(la_var))){
  la_phlyo[[i]] <- la_var[[1]][,1]</pre>
  la_spec[[i]] <- la_var[[1]][,2]</pre>
  la_unit[[i]] <- la_var[[1]][,3]</pre>
  }
la_phlyo <- unlist(la_phlyo)</pre>
la_spec <- unlist(la_spec)</pre>
la_unit <- unlist(la_unit)</pre>
la_prop_phlyo <- la_phlyo/(la_phlyo + la_spec + la_unit)</pre>
la_prop_spec <- la_spec/(la_phlyo + la_spec + la_unit)</pre>
la_prop_residuals <- la_unit/(la_phlyo + la_spec + la_unit)</pre>
#Phylogenetic signal
hdr(la_prop_phlyo)$mode
## [1] 0.8936751
#Population level variance
hdr(la_prop_spec)$mode
```

Mean Reporductive Rate

```
repo_models <- read.mulTree("mean_repo_rate_run")</pre>
summary(repo_models)
##
                          Estimates(mode hdr) lower.CI(2.5) lower.CI(25)
## (Intercept)
                                    -0.3821796 -1.9569241
                                                                 -0.8931401
                                    -0.3269163
                                                                 -0.4028667
## mass g
                                                   -0.5526083
                                    -0.2602267
                                                                 -0.3039558
## matrix size
                                                   -0.3864657
## phylogenetic.variance
                                     1.3403244
                                                    0.4631857
                                                                  0.9726163
## residual.variance
                                     0.3852503
                                                    0.2280839
                                                                  0.3277378
                          upper.CI(75) upper.CI(97.5)
## (Intercept)
                              0.0856870
                                            1.02958316
                             -0.2466111
                                           -0.09573132
## mass_g
## matrix_size
                             -0.2177105
                                            -0.13615120
## phylogenetic.variance
                              1.8029584
                                            2.99722645
## residual.variance
                              0.4587044
                                             0.61796005
## attr(,"class")
## [1] "matrix" "mulTree"
Now we calculate the proportion of variance between phylogenetic, population and residual variance.
repo_var <- read.mulTree("mean_repo_rate_run", extract = "VCV")</pre>
repo_phlyo <- list()</pre>
repo_spec <- list()</pre>
repo_unit <- list()</pre>
for(i in 1:length(names(repo_var))){
  repo_phlyo[[i]] <- repo_var[[1]][,1]</pre>
  repo_spec[[i]] <- repo_var[[1]][,2]</pre>
  repo_unit[[i]] <- repo_var[[1]][,3]</pre>
repo_phlyo <- unlist(repo_phlyo)</pre>
repo_spec <- unlist(repo_spec)</pre>
repo_unit <- unlist(repo_unit)</pre>
repo_prop_phlyo <- repo_phlyo/(repo_phlyo + repo_spec + repo_unit)</pre>
repo_prop_spec <- repo_spec/(repo_phlyo + repo_spec + repo_unit)</pre>
```

Mean Reporductive Rate not at the stabel state distribution

```
repo nst models <- read.mulTree("mean repo rate nst run")</pre>
summary(repo_nst_models)
##
                          Estimates(mode hdr) lower.CI(2.5) lower.CI(25)
                                   0.20587434 -1.9870391 -0.50269567
## (Intercept)
                                  -0.01381948 -0.2666529 -0.09755688
## mass_g
## matrix_size
                                  -0.18379335
                                                 -0.2820732 -0.21757472
## phylogenetic.variance
                                   3.02741680
                                                  1.3126973
                                                               2.25227288
## residual.variance
                                   0.35185781
                                                   0.1496883
                                                               0.27277345
##
                          upper.CI(75) upper.CI(97.5)
## (Intercept)
                           0.90939664
                                            2.3361914
## mass_g
                            0.07770733
                                            0.2478651
## matrix_size
                           -0.15030098
                                           -0.0860611
                                            6.5209695
## phylogenetic.variance 4.00080256
## residual.variance
                            0.42884686
                                            0.5987686
## attr(,"class")
## [1] "matrix" "mulTree"
Now we calculate the proportion of variance between phylogenetic, population and residual variance.
repo_nst_var <- read.mulTree("mean_repo_rate_nst_run", extract = "VCV")</pre>
repo_nst_phlyo <- list()</pre>
repo_nst_spec <- list()
repo_nst_unit <- list()</pre>
for(i in 1:length(names(repo_nst_var))){
  repo_nst_phlyo[[i]] <- repo_nst_var[[1]][,1]</pre>
```

```
repo_nst_spec[[i]] <- repo_nst_var[[1]][,2]</pre>
  repo_nst_unit[[i]] <- repo_nst_var[[1]][,3]</pre>
  }
repo_nst_phlyo <- unlist(repo_nst_phlyo)</pre>
repo_nst_spec <- unlist(repo_nst_spec)</pre>
repo_nst_unit <- unlist(repo_nst_unit)</pre>
repo_nst_prop_phlyo <- repo_nst_phlyo/(repo_nst_phlyo + repo_nst_spec + repo_nst_unit)
repo_nst_prop_spec <- repo_nst_spec/(repo_nst_phlyo + repo_nst_spec + repo_nst_unit)</pre>
repo_nst_prop_residuals <- repo_nst_unit/(repo_nst_phlyo + repo_nst_spec + repo_nst_unit)
#Phylogenetic signal
hdr(repo_nst_prop_phlyo)$mode
## [1] 0.9289587
#Population level variance
hdr(repo_nst_prop_spec)$mode
## [1] 0.08177648
#Residual term
hdr(repo_nst_prop_residuals)$mode
## [1] 0.009848131
Next calculate the residuals from the allometric model for mean reporductive rate not at the stable state.
repo_nst_resids <- mul_resids(mul_output = repo_nst_models,</pre>
                         mul data = pop multree,
                         Y_data_col = c("mean_repo_rate")
```

Standard deviation of mxlx

```
mxlxsd_models <- read.mulTree("mxlxsd_logged_10_run")</pre>
summary(mxlxsd models)
##
                         Estimates (mode hdr) lower.CI(2.5) lower.CI(25)
## (Intercept)
                                  -0.1635107 -1.65326374 -0.63001884
                                               -0.43282306 -0.27444772
## mass_g
                                  -0.1924576
## matrix_size
                                   0.1277956
                                               -0.03112926
                                                             0.07252093
                                   1.1087434
                                                0.30288752
                                                              0.75206045
## phylogenetic.variance
## residual.variance
                                   0.3783354
                                                0.16403320
                                                              0.29920713
##
                         upper.CI(75) upper.CI(97.5)
## (Intercept)
                            0.2940519
                                          1.21802905
## mass_g
                           -0.1085906
                                          0.05379147
                                          0.28327683
## matrix_size
                            0.1801876
                                          2.85798056
                            1.5470773
## phylogenetic.variance
## residual.variance
                            0.4665128
                                          0.66387366
## attr(,"class")
## [1] "matrix" "mulTree"
```

Now we calculate the proportion of variance between phylogenetic, population and residual variance.

```
mxlxsd_var <- read.mulTree("mxlxsd_logged_10_run", extract = "VCV")</pre>
mxlxsd_phlyo <- list()</pre>
mxlxsd_spec <- list()</pre>
mxlxsd_unit <- list()</pre>
for(i in 1:length(names(mxlxsd var))){
  mxlxsd_phlyo[[i]] <- mxlxsd_var[[1]][,1]</pre>
  mxlxsd_spec[[i]] <- mxlxsd_var[[1]][,2]</pre>
  mxlxsd_unit[[i]] <- mxlxsd_var[[1]][,3]</pre>
mxlxsd_phlyo <- unlist(mxlxsd_phlyo)</pre>
mxlxsd_spec <- unlist(mxlxsd_spec)</pre>
mxlxsd_unit <- unlist(mxlxsd_unit)</pre>
mxlxsd_prop_phlyo <- mxlxsd_phlyo/(mxlxsd_phlyo + mxlxsd_spec + mxlxsd_unit)
mxlxsd_prop_spec <- mxlxsd_spec/(mxlxsd_phlyo + mxlxsd_spec + mxlxsd_unit)</pre>
mxlxsd_prop_residual <- mxlxsd_unit/(mxlxsd_phlyo + mxlxsd_spec + mxlxsd_unit)</pre>
#Phylogenetic signal
hdr(mxlxsd prop phlyo)$mode
## [1] 0.6702649
#Population level variance
hdr(mxlxsd_prop_spec)$mode
## [1] 0.1697088
#Residual term
hdr(mxlxsd_prop_residual)$mode
## [1] 0.1641848
Next calculate the residuals from the allometric model for mxlxsd
mxlxsd_resids <- mul_resids(mul_output = mxlxsd_models,</pre>
                          mul_data = pop_multree,
                          Y_data_col = c("mxlxsd")
```

Generation Time

```
## phylogenetic.variance
                                    3.02776553
                                                  1.685256044
                                                                 2.50942926
## residual.variance
                                    0.07255637 -0.001590761
                                                                 0.04181470
                          upper.CI(75) upper.CI(97.5)
##
## (Intercept)
                           1.268474269
                                            2.61018832
## mass_g
                           0.658243780
                                            0.79282154
                          -0.002310125
                                            0.05672399
## matrix size
## phylogenetic.variance 3.694971664
                                            5.18105881
## residual.variance
                                            0.17293323
                           0.105109278
## attr(,"class")
## [1] "matrix" "mulTree"
Now we calculate the proportion of variance between phylogenetic, population and residual variance.
gen_time_var <- read.mulTree("gen_time_run", extract = "VCV")</pre>
gen_time_phlyo <- list()</pre>
gen_time_spec <- list()</pre>
gen_time_unit <- list()</pre>
for(i in 1:length(names(gen_time_var))){
  gen_time_phlyo[[i]] <- gen_time_var[[1]][,1]</pre>
  gen_time_spec[[i]] <- gen_time_var[[1]][,2]</pre>
  gen_time_unit[[i]] <- gen_time_var[[1]][,3]</pre>
gen_time_phlyo <- unlist(gen_time_phlyo)</pre>
gen time spec <- unlist(gen time spec)</pre>
gen_time_unit <- unlist(gen_time_unit)</pre>
gen_time_prop_phlyo <- gen_time_phlyo/(gen_time_phlyo + gen_time_spec + gen_time_unit)</pre>
gen time prop spec <- gen time spec/(gen time phlyo + gen time spec + gen time unit)
gen_time_prop_residual <- gen_time_unit/(gen_time_phlyo + gen_time_spec + gen_time_unit)</pre>
#Phylogenetic signal
hdr(gen_time_prop_phlyo)$mode
## [1] 0.9690505
#Population level variance
hdr(gen_time_prop_spec)$mode
## [1] 0.01624795
#Residual term
hdr(gen_time_prop_residual)$mode
## [1] 0.02174675
Next calculate the residuals from the allometric model for generation time
gen_time_resids <- mul_resids(mul_output = gen_time_models,</pre>
                         mul_data = pop_multree,
                         Y_data_col = c("gen_time")
```

Life expectancy conditional on reaching sexual maturity

```
M_rep_lif_exp_models <- read.mulTree("M_rep_lif_exp_run")</pre>
summary(M_rep_lif_exp_models)
                                                     Estimates (mode hdr) lower.CI(2.5) lower.CI(25)
## (Intercept)
                                                                         0.31580239 -1.439073835 -0.26028814
                                                                         0.58795350 0.386374260
                                                                                                                                  0.51712046
## mass_g
## matrix_size
                                                                        0.05466408 -0.049172640 0.02086333
## phylogenetic.variance
                                                                        2.24980397 1.100305459
                                                                                                                                  1.78280214
                                                                         0.07502929 -0.005531941
## residual.variance
                                                                                                                                  0.03867796
##
                                                     upper.CI(75) upper.CI(97.5)
## (Intercept)
                                                           0.9091395
                                                                                          2.0895027
                                                            0.6529346
                                                                                            0.7840525
## mass_g
## matrix_size
                                                           0.0939637
                                                                                           0.1641865
## phylogenetic.variance
                                                           2.7868984
                                                                                           4.0508399
## residual.variance
                                                           0.1141998
                                                                                            0.1933473
## attr(,"class")
## [1] "matrix" "mulTree"
Now we calculate the proportion of variance between phylogenetic, population and residual variance.
M_rep_lif_exp_var <- read.mulTree("M_rep_lif_exp_run", extract = "VCV")</pre>
M_rep_lif_exp_phlyo <- list()</pre>
M_rep_lif_exp_spec <- list()</pre>
M_rep_lif_exp_unit <- list()</pre>
for(i in 1:length(names(M_rep_lif_exp_var))){
    M_rep_lif_exp_phlyo[[i]] <- M_rep_lif_exp_var[[1]][,1]</pre>
    M_rep_lif_exp_spec[[i]] <- M_rep_lif_exp_var[[1]][,2]</pre>
    M_rep_lif_exp_unit[[i]] <- M_rep_lif_exp_var[[1]][,3]</pre>
    }
M_rep_lif_exp_phlyo <- unlist(M_rep_lif_exp_phlyo)</pre>
M_rep_lif_exp_spec <- unlist(M_rep_lif_exp_spec)</pre>
M_rep_lif_exp_unit <- unlist(M_rep_lif_exp_unit)</pre>
M_rep_lif_exp_prop_phlyo <- M_rep_lif_exp_phlyo/(M_rep_lif_exp_phlyo + M_rep_lif_exp_spec + M
M_rep_lif_exp_prop_spec <- M_rep_lif_exp_spec/(M_rep_lif_exp_phlyo + M_rep_lif_exp_spec + M_rep_lif_ex
M_rep_lif_exp_prop_residuals <- M_rep_lif_exp_unit/(M_rep_lif_exp_phlyo + M_rep_lif_exp_spec + M_rep_l
#Phylogenetic signal
hdr(M_rep_lif_exp_prop_phlyo)$mode
## [1] 0.9303974
#Population level variance
hdr(M_rep_lif_exp_prop_spec)$mode
## [1] 0.01849731
#Residual term
hdr(M_rep_lif_exp_prop_residuals)$mode
```

```
## [1] 0.05573421
```

Next calculate the residuals from the allometric model for life expectancy conditional on reaching sexual maturity

```
M_rep_lif_exp_resids <- mul_resids(mul_output = M_rep_lif_exp_models,</pre>
                         mul_data = pop_multree,
                         Y_data_col = c("M_rep_lif_exp")
)
```

Gini index

```
gini_models <- read.mulTree("gini_logged_run")</pre>
summary(gini_models)
                         Estimates (mode hdr) lower.CI(2.5) lower.CI(25)
##
## (Intercept)
                                 -0.42794181
                                                -2.0464228
                                                             -0.9689459
## mass g
                                 -0.19599691
                                                -0.4244735
                                                             -0.2745817
## matrix_size
                                 -0.09997641
                                                -0.2324744
                                                            -0.1452908
## phylogenetic.variance
                                  1.79736608
                                                 0.9172384
                                                             1.4300211
## residual.variance
                                  0.25793064
                                                 0.1276131
                                                              0.2086544
                         upper.CI(75) upper.CI(97.5)
## (Intercept)
                          0.11276323
                                          1.19790403
## mass_g
                          -0.11831885
                                          0.03280006
## matrix_size
                          -0.05474589
                                          0.03184347
                         2.19371282
                                          3.26168320
## phylogenetic.variance
                                          0.44078630
## residual.variance
                           0.31350150
## attr(,"class")
## [1] "matrix" "mulTree"
```

Now we calculate the proportion of variance between phylogenetic, population and residual variance.

```
gini_var <- read.mulTree("gini_logged_run", extract = "VCV")</pre>
gini_phlyo <- list()</pre>
gini_spec <- list()</pre>
gini_unit <- list()</pre>
for(i in 1:length(names(gini_var))){
  gini_phlyo[[i]] <- gini_var[[1]][,1]
  gini_spec[[i]] <- gini_var[[1]][,2]</pre>
  gini_unit[[i]] <- gini_var[[1]][,3]
gini_phlyo <- unlist(gini_phlyo)</pre>
gini_spec <- unlist(gini_spec)</pre>
gini_unit <- unlist(gini_unit)</pre>
gini_prop_phlyo <- gini_phlyo/(gini_phlyo + gini_spec + gini_unit)</pre>
gini_prop_spec <- gini_spec/(gini_phlyo + gini_spec + gini_unit)</pre>
gini_prop_residuals <- gini_unit/(gini_phlyo + gini_spec + gini_unit)</pre>
```

Standard deviation of mortality rates

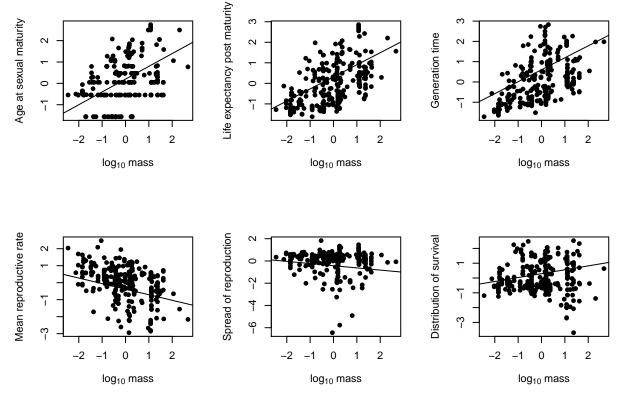
```
surv_sd_models <- read.mulTree("surv_sd_logged_run")</pre>
summary(surv_sd_models)
                          Estimates (mode hdr) lower.CI(2.5) lower.CI(25)
## (Intercept)
                                    0.2921592 -1.17657819 -0.1883418
## mass_g
                                    0.2633087
                                                  0.03469292
                                                                 0.1864593
## matrix_size
                                   -0.3503796 -0.50536667 -0.4040569
## phylogenetic.variance
                                    1.1640853
                                                 0.38884786
                                                                 0.8386304
                                                                 0.1882762
## residual.variance
                                    0.2433491
                                                  0.09921485
                          upper.CI(75) upper.CI(97.5)
## (Intercept)
                             0.7439494
                                            1.7093005
## mass_g
                             0.3445201
                                             0.5007707
## matrix_size
                                            -0.1995487
                            -0.2995992
## phylogenetic.variance
                                             2.8398280
                             1.6285460
## residual.variance
                             0.3083719
                                             0.4565608
## attr(,"class")
## [1] "matrix" "mulTree"
Now we calculate the proportion of variance between phylogenetic, population and residual variance.
surv_sd_var <- read.mulTree("surv_sd_logged_run", extract = "VCV")</pre>
surv_sd_phlyo <- list()</pre>
surv_sd_spec <- list()</pre>
surv_sd_unit <- list()</pre>
for(i in 1:length(names(surv_sd_var))){
  surv_sd_phlyo[[i]] <- surv_sd_var[[1]][,1]</pre>
  surv_sd_spec[[i]] <- surv_sd_var[[1]][,2]</pre>
```

```
surv_sd_unit[[i]] <- surv_sd_var[[1]][,3]</pre>
surv_sd_phlyo <- unlist(surv_sd_phlyo)</pre>
surv_sd_spec <- unlist(surv_sd_spec)</pre>
surv_sd_unit <- unlist(surv_sd_unit)</pre>
surv_sd_prop_phlyo <- surv_sd_phlyo/(surv_sd_phlyo</pre>
                                        + surv_sd_spec
                                        + surv_sd_unit)
surv_sd_prop_spec <- surv_sd_spec/(surv_sd_phlyo</pre>
                                       + surv_sd_spec
                                       + surv_sd_unit)
surv_sd_prop_residuals <- surv_sd_unit/(surv_sd_phlyo</pre>
                                            + surv_sd_spec
                                            + surv_sd_unit)
#Phylogenetic signal
hdr(surv_sd_prop_phlyo)$mode
## [1] 0.7196334
#Population level variance
hdr(surv_sd_prop_spec)$mode
## [1] 0.107074
#Residual term
hdr(surv_sd_prop_residuals)$mode
## [1] 0.2247204
Next calculate the residuals from the allometric model for the standard deviation
surv_sd_resids <- mul_resids(mul_output = surv_sd_models,</pre>
                          mul_data = pop_multree,
                          Y_data_col = c("surv_sd")
)
```

Plotting allometry model output

Using the outputs for the life history metrics lets plot them out.

```
summary(La_models)[2])
#M_rep_lif_exp
plot(pop_multree$data$M_rep_lif_exp ~ pop_multree$data$mass_g,
     pch = 16,
     xlab = expression('log'[10]*" mass"),
     ylab = "Life expectancy post maturity")
abline(summary(M_rep_lif_exp_models)[1],summary(M_rep_lif_exp_models)[2])
#generation time
plot(pop_multree$data$gen_time ~ pop_multree$data$mass_g, pch = 16,
          xlab = expression('log'[10]*" mass"),
          ylab = "Generation time")
abline(summary(gen_time_models)[1],summary(gen_time_models)[2])
##mean repo rate
plot(pop_multree$data$mean_repo_rate_stable_state ~ pop_multree$data$mass_g, pch = 16,
          xlab = expression('log'[10]*" mass"),
          ylab = "Mean reproductive rate")
abline(summary(repo_models)[1],summary(repo_models)[2])
#Gini
plot(pop_multree$data$gini ~ pop_multree$data$mass_g, pch = 16,
          xlab = expression('log'[10]*" mass"),
          ylab = "Spread of reproduction")
abline(summary(gini_models)[1],summary(gini_models)[2])
#SD of survival
plot(pop_multree$data$surv_sd ~ pop_multree$data$mass_g, pch = 16,
     xlab = expression('log'[10]*" mass"),
     ylab = "Distribution of survival")
abline(summary(surv_sd_models)[1],summary(surv_sd_models)[2])
```

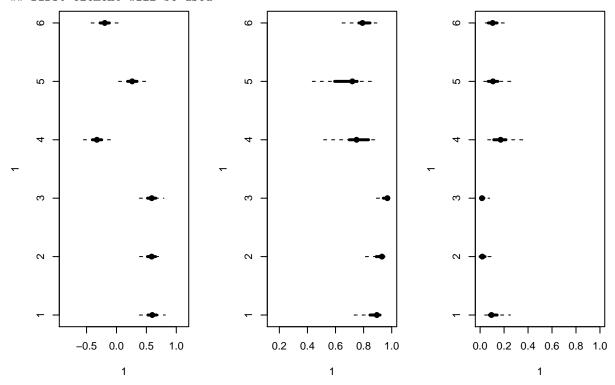


Lets also plot out the model slope coefficients into a table.

```
##Allometric scaling
par(mfrow=c(1,3))
scaling_list <- list( La_B = La_models$mass_g,</pre>
                       Sur_B = M_rep_lif_exp_models$mass_g,
                       T_B = gen_time_models$mass_g,
                       Repo_B = repo_models$mass_g,
                       life_shape_B = surv_sd_models$mass_g,
                       gini_B = gini_models$mass_g
MultiDisPlot(scaling_list)
##phylogentic signals
phy_var_list <- list( La_B = la_prop_phlyo,</pre>
                       Sur_B = M_rep_lif_exp_prop_phlyo,
                       T_B = gen_time_prop_phlyo,
                       Repo_B = repo_prop_phlyo,
                       life_shape_B = surv_sd_prop_phlyo,
                       gini_B = gini_prop_phlyo
MultiDisPlot(phy_var_list)
##population level variance
species_var_list <- list(</pre>
                       La_B = la_prop_spec,
```

```
Sur_B = M_rep_lif_exp_prop_spec,
    T_B = gen_time_prop_spec,
    Repo_B = repo_prop_spec,
    life_shape_B = surv_sd_prop_spec,
    gini_B = gini_prop_spec
    )
MultiDisPlot(species_var_list, xlim = c(0,1))
```

Warning in if (xlim == "auto") $\{: \text{ the condition has length} > 1 \text{ and only the}$ ## first element will be used



Now lets create a dataset of these residuals for each PCA analysis. This includes, the main analysis, the analysis using mean reporductive rate with the population not at its stable state distribution, the analysis using the standard deviation of mxlx curve as a measure of the Gini index and the analysis with generation time removed.

```
M_repo_nst = repo_nst_resids,
                         M_suv = M_rep_lif_exp_resids,
                         gini_r = gini_resids
#PCA dataset using the standard deviation of the mxlx curve
predicted_data_mxlxsd <- data.frame(</pre>
                         SD_mort = surv_sd_resids,
                         La_r = La_resids,
                         gen_r = gen_time_resids,
                         M_repo = repo_resids,
                         M_suv = M_rep_lif_exp_resids,
                         mxlxsd = mxlxsd_resids
#PCA dataset without generation time.
predicted_data_noT <- data.frame(</pre>
                          SD_mort = surv_sd_resids,
                         La_r = La_resids,
                         M_repo = repo_resids,
                         M_suv = M_rep_lif_exp_resids,
                         gini = gini_resids
```

PCA

Now we run a PCA for each of the datasets of residuals from the life history allometric models. We use Horn's Parallel Analysis of Principal Components to test for the number of axis to retain. Note that for ease of interpretation the sign of some axis was reversed in the corresponding supplementary table 3 so that the fast slow axis alway went from fast on the left to slow on the right.

First the main analysis

```
pca_res <- prcomp(predicted_data)</pre>
pca_res
## Standard deviations (1, .., p=6):
## [1] 1.5214900 1.1418273 0.8361469 0.6983264 0.4966644 0.2823536
##
## Rotation (n \times k) = (6 \times 6):
                 PC1
                                        PC3
                                                    PC4
                                                                PC5
##
                            PC2
## SD_mort -0.3373610 -0.5056419 -0.64858897 -0.30533698
                                                         0.30602653
## La_r
          -0.5012398  0.2074782  0.12294518  0.59463208
                                                         0.54472281
## gen_r
          ## M_repo
           0.1458494 -0.5755156 -0.14330980 0.69111082 -0.26505288
## M_suv
           -0.4812860 0.1440881 -0.14512090 0.09118089 -0.72576868
## gini_r
           0.2705314 \quad 0.5776088 \quad -0.72267129 \quad 0.21604545 \quad 0.01336949
##
                 PC6
## SD mort -0.1515365
## La_r
          -0.2007131
           0.8013147
## gen_r
## M_repo
           0.2812202
           -0.4375936
## M suv
## gini_r
          0.1551539
```

```
horn_res <- paran(predicted_data)</pre>
## Using eigendecomposition of correlation matrix.
## Computing: 10% 20% 30% 40% 50% 60% 70% 80% 90% 100%
##
##
## Results of Horn's Parallel Analysis for component retention
## 180 iterations, using the mean estimate
## -----
## Component Adjusted Unadjusted Estimated
             Eigenvalue Eigenvalue Bias
## ----
             ______
             2.601002
                        2.804730
## 1
                                    0.203727
            1.380085 1.481356
                                    0.101271
## -----
## Adjusted eigenvalues > 1 indicate dimensions to retain.
## (2 components retained)
Next the analysis with reporductive rate with the population not at the stable state distribution,
pca_nst <- prcomp(predicted_data_M_repo_nst)</pre>
pca_nst
## Standard deviations (1, .., p=6):
## [1] 1.5247503 1.1510163 0.8861361 0.7921989 0.4446327 0.3047726
## Rotation (n \times k) = (6 \times 6):
##
                   PC1
                             PC2
                                         PC3
                                                   PC4
## SD_mort -0.3812794 -0.2234415 0.713723340 -0.4832885 0.21567840
## La_r -0.4898175 0.1879261 -0.515947355 -0.1201502 0.56296273
## gen_r
            -0.5181094 0.2802035 0.035606085 0.1384867 0.02155379
## M_repo_nst -0.1788388 -0.6792153 -0.455790782 -0.4051659 -0.29258542
## M_suv -0.4480816 0.3229370 0.004226023 -0.0225824 -0.74060487
           0.3369081 0.5202133 -0.123961409 -0.7537561 -0.04460302
## gini_r
##
                   PC6
## SD_mort
            -0.1233449
         -0.3566481
## La_r
## gen_r
            0.7950709
## M_repo_nst 0.2217486
## M suv
          -0.3819831
## gini_r
            0.1742611
horn_nst <- paran(predicted_data_M_repo_nst)</pre>
##
## Using eigendecomposition of correlation matrix.
## Computing: 10% 20% 30% 40% 50% 60% 70% 80% 90% 100%
##
## Results of Horn's Parallel Analysis for component retention
## 180 iterations, using the mean estimate
##
```

```
## Component Adjusted Unadjusted Estimated
      Eigenvalue Eigenvalue Bias
## -----
                     2.759187
## 1
             2.560805
                                   0.198382
            1.305770 1.410377 0.104607
## 2
## Adjusted eigenvalues > 1 indicate dimensions to retain.
## (2 components retained)
Next the analysis with generation time not included.
pca_mxlxsd <- prcomp(predicted_data_mxlxsd)</pre>
pca_mxlxsd
## Standard deviations (1, .., p=6):
## [1] 1.5510226 1.1188432 0.9079943 0.6290138 0.4439734 0.2063080
##
## Rotation (n \times k) = (6 \times 6):
              PC1 PC2
                                   PC3
                                                PC4
## SD_mort -0.2541265 -0.58578261 -0.63423830 -0.19751156 0.3849226
## La_r -0.4827626 -0.11207474 0.40710084 0.59465655 0.4703647
## gen r -0.5579208 -0.04688062 0.01038699 -0.04873374 -0.3433282
## M_repo 0.2690637 -0.55268899 -0.09269274 0.58351718 -0.5214535
## M_suv -0.4406493 -0.24037911 0.30866469 -0.38899450 -0.4261037
## mxlxsd 0.3528700 -0.52806530 0.57274510 -0.33643537 0.2433912
                PC6
## SD_mort 0.05332925
## La r
         0.11742655
## gen_r -0.75244233
## M_repo 0.03378873
## M_suv
          0.56558915
## mxlxsd -0.31010446
horn_mxlxsd <- paran(predicted_data_mxlxsd)</pre>
##
## Using eigendecomposition of correlation matrix.
## Computing: 10% 20% 30% 40% 50% 60% 70% 80% 90% 100%
##
##
## Results of Horn's Parallel Analysis for component retention
## 180 iterations, using the mean estimate
##
## Component Adjusted Unadjusted
            Eigenvalue Eigenvalue
                                  Bias
## -----
## 1
             2.711875 2.911102
                                   0.199226
            1.333164 1.437719 0.104555
## 2
## -----
## Adjusted eigenvalues > 1 indicate dimensions to retain.
## (2 components retained)
```

Next the analysis with generation time not included.

```
pca_noT <- prcomp(predicted_data_noT)</pre>
pca_noT
## Standard deviations (1, .., p=5):
## [1] 1.2902907 1.1185509 0.8359158 0.6884132 0.4932431
##
## Rotation (n \times k) = (5 \times 5):
##
                 PC1
                           PC2
                                     PC3
                                                PC4
## SD_mort -0.508465012 -0.3638112 0.6467465 -0.33951272 -0.27487234
       0.001115311 -0.6034195 0.1458530 0.75663818 0.20520538
## M_repo
          -0.508048272 0.3288245 0.1320249 0.02472698 0.78467791
## M_suv
          0.440907500 0.4700652 0.7228146 0.24598171 -0.04088118
## gini
horn_noT <- paran(predicted_data_noT)</pre>
##
## Using eigendecomposition of correlation matrix.
## Computing: 10% 20% 30% 40% 50% 60% 70% 80% 90% 100%
##
## Results of Horn's Parallel Analysis for component retention
## 150 iterations, using the mean estimate
## -----
## Component
            Adjusted
                        Unadjusted
                                    Estimated
##
             Eigenvalue Eigenvalue
                                    Bias
             1.757631
                                     0.176491
## 1
                        1.934123
             1.394326
                       1.466940
                                    0.072614
##
## Adjusted eigenvalues > 1 indicate dimensions to retain.
## (2 components retained)
```

PCA plots

Note that for ease of interpretation the sign of some axis was reversed so that the fast slow axis goes from fast on the left to slow on the right.

```
results <- pca_res
results$rotation[,"PC1"] <- -results$rotation[,"PC1"]
results$x[,"PC1"] <- -results$x[,"PC1"]
results$rotation[,"PC2"] <- results$rotation[,"PC2"]
results$x[,"PC2"] <- results$x[,"PC2"]</pre>
```

We then set up the colors we will give to each of the taxinommic groups, the color of the PC arrows and the life history symbols

```
result <- results
loadings <- as.data.frame(result$rotation)</pre>
```

```
loadings[,"col"]=c("gray50",
                    "gray50",
                    "gray50",
                    "gray50",
                    "gray50",
                    "gray50"
loadings$LHT=rownames(loadings)
loadings$LHT=c("surv_sd",
                "La",
                "gen_time",
                "mean_repo_rate",
                "M_suv"
                ,"gini_r"
loadings$LHTexpr <- list(</pre>
                           expression(sigma),
                           expression("L"[alpha]),
                           expression("T"),
                           expression(phi),
                           expression(Rep["e"]),
                           expression("G")
arrowThickness=2.9
sizeArrowLetters=1
scalingArrows=2.5
scalingLetters=2.9
class_match <- vector()</pre>
species_match <- vector()</pre>
loads_taxa <- data.frame(results$x)</pre>
for(i in 1:length(loads_taxa[,1])){
   class_match[i] <- as.vector(pop_multree$data[i,"taxa_name"])</pre>
   species_match[i] <- as.vector(pop_multree$data[i,"species"])</pre>
   }
#combine the pca results data with the external data
pca_data <- data.frame(loads_taxa,</pre>
                        class_match = as.vector(pop_multree$data[,"taxa_name"]),
                        species_match = as.vector(pop_multree$data[,"species"]),
                        PCA_moblist = as.vector(pop_multree$data[,"mode_of_life"]),
                        PCA_met = as.vector(pop_multree$data[,"met_rate"]),
                        PCA_repo = as.vector(pop_multree$data[,"repo_output"]),
                        PCA_iucn = as.vector(pop_multree$data[,"IUCN"]),
```

```
PCA_met_type = as.vector(pop_multree$data[,"met_type"]),
                         animal = as.vector(pop_multree$data[,"species"])
mam_col <- rgb(0, 136, 170,
                max = 255)
bird_col <- rgb(255,153,85,
                 max = 255)
rep_col <- rgb(147,172,147,
                max = 255)
fish_col <- rgb(135,205,222,
                 max = 255)
sponge_col <- rgb(211,95,141,</pre>
                   max = 255)
coral_col <- rgb(153,85,255,</pre>
                  max = 255)
gast_col <- rgb(255,170,238,
                 max = 255)
biv_col <- rgb(205,135,222,
                max = 255)
shark_col <- rgb(85,0,212,
                  max = 255)
```

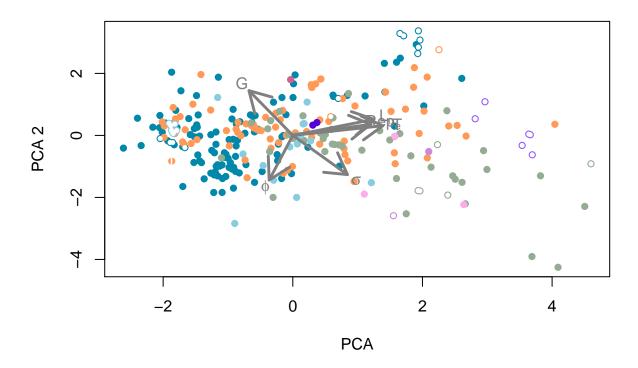
Now lets plot out the two retained axis from our PCA.

```
plot(pca_data[,1],
    pca_data[,2],
    pch=16,
     cex = 0.1,
     col = "white",
     xlab= "PCA",
     ylab= "PCA 2")
points(pca_data[pca_data$class_match == "Mammalia",1],
       pca_data[pca_data$class_match == "Mammalia",2],
       pch=16,
       col = mam_col)
points(pca_data[pca_data$class_match == "Aves",1],
       pca_data[pca_data$class_match == "Aves",2],
       pch=16,
       col = bird_col)
points(pca_data[pca_data$class_match == "Reptilia",1],
```

```
pca_data[pca_data$class_match == "Reptilia",2],
       pch=16,
       col = rep_col)
points(pca_data[pca_data$class_match == "Actinopterygii",1],
       pca_data[pca_data$class_match == "Actinopterygii",2],
       pch=16,
       col = fish_col)
points(pca_data[pca_data$class_match == "Gastropoda",1],
       pca_data[pca_data$class_match == "Gastropoda",2],
       pch=16,
       col = gast_col)
points(pca_data[pca_data$class_match == "Demospongiae",1],
       pca_data[pca_data$class_match == "Demospongiae",2],
       pch=16,
       col = sponge_col)
points(pca_data[pca_data$class_match == "Anthozoa",1],
       pca_data[pca_data$class_match == "Anthozoa",2],
       pch=16,
       col = coral_col)
points(pca_data[pca_data$class_match == "Bivalvia",1],
       pca_data[pca_data$class_match == "Bivalvia",2],
       pch=16,
       col = biv_col)
points(pca_data[pca_data$class_match == "Elasmobranchii",1],
       pca_data[pca_data$class_match == "Elasmobranchii",2],
       pch=16,
       col = shark_col)
###And lets add Humans
points(pca_data[pca_data$species_match == "Homo_sapiens",1],
       pca_data[pca_data$species_match == "Homo_sapiens",2],
       pch= 16,
       col = "white",
       cex = 0.7)
##and other points
points(pca_data[pca_data$species_match == "Elephas_maximus",1],
       pca_data[pca_data$species_match == "Elephas_maximus",2],
       pch= 16,
       col = "white",
       cex = 0.7)
points(pca_data[pca_data$species_match == "Fulmarus_glacialis",1],
       pca_data[pca_data$species_match == "Fulmarus_glacialis",2],
       pch= 16,
       col = "white",
       cex = 0.7
```

```
points(pca_data[pca_data$species_match == "Tympanuchus_cupido",1],
       pca_data[pca_data$species_match == "Tympanuchus_cupido",2],
       pch= "T",
       col = "white")
points(pca_data[pca_data$species_match == "Gyps_coprotheres",1],
       pca_data[pca_data$species_match == "Gyps_coprotheres",2],
       pch= 16,
       col = "white",
       cex = 0.7
points(pca_data[pca_data$species_match == "Crocodylus_johnsoni",1],
       pca_data[pca_data$species_match == "Crocodylus_johnsoni",2],
       pch= 16,
       col = "white",
       cex = 0.7)
points(pca_data[pca_data$species_match == "Urocitellus_armatus",1],
       pca_data[pca_data$species_match == "Urocitellus_armatus",2],
       pch= 16,
       col = "white",
       cex = 0.7
points(pca_data[pca_data$species_match == "Paramuricea_clavata",1],
       pca_data[pca_data$species_match == "Paramuricea_clavata",2],
       pch= 16,
       col = "white",
       cex = 0.7)
points(pca_data[pca_data$species_match == "Oncorhynchus_tshawytscha",1],
       pca_data[pca_data$species_match == "Oncorhynchus_tshawytscha",2],
       pch= 16,
       col = "white",
       cex = 0.7
points(pca_data[pca_data$species_match == "Mya_arenaria",1],
       pca_data[pca_data$species_match == "Mya_arenaria",2],
       pch=16,
       col = "white",
       cex = 0.7)
points(pca_data[pca_data$species_match == "Clemmys_guttata",1],
       pca_data[pca_data$species_match == "Clemmys_guttata",2],
       pch= 16,
       col = "white",
       cex = 0.7)
arrows(x0=0,
       y0=0,
       x1=loadings[,1]*scalingArrows,
       y1=loadings[,2]*scalingArrows,
       col="black",
```

```
lwd=2)
arrows(x0=0,
       v^{0=0}.
       x1=loadings[,1]*scalingArrows,
       y1=loadings[,2]*scalingArrows,
       col=as.character(loadings$col),
       lwd=arrowThickness)
text(loadings[1,"PC1"]*scalingLetters-.0,
     loadings[1,"PC2"]*scalingLetters,
     loadings$LHTexpr[[1]],
     col = loadings$col[1],
     cex=sizeArrowLetters)
text(loadings[2,"PC1"]*scalingLetters-.0,
     loadings[2,"PC2"]*scalingLetters,
     loadings$LHTexpr[[2]],
     col = loadings$col[2],
     cex=sizeArrowLetters)
text(loadings[3,"PC1"]*scalingLetters+.0,
     loadings[3,"PC2"]*scalingLetters,
     loadings$LHTexpr[[3]],
     col = loadings$col[3],
     cex=sizeArrowLetters)
text(loadings[4,"PC1"]*scalingLetters-.0,
     loadings[4,"PC2"]*scalingLetters,
     loadings$LHTexpr[[4]],
     col = loadings$col[4],
     cex=sizeArrowLetters)
text(loadings[5,"PC1"]*scalingLetters-.0,
     loadings[5,"PC2"]*scalingLetters,
     loadings$LHTexpr[[5]],
     col = loadings$col[5],
     cex=sizeArrowLetters)
text(loadings[6,"PC1"]*scalingLetters+.0,
     loadings[6,"PC2"]*scalingLetters,
     loadings$LHTexpr[[6]],
     col = loadings$col[6],
     cex=sizeArrowLetters)
```



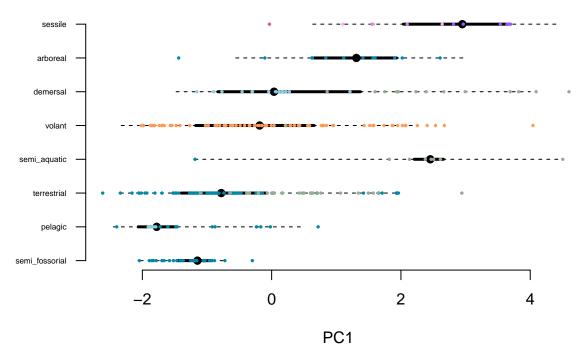
Mode of life analysis analysis

Using PC1 we test how different modes of life are distributed across the portion of life histroy space associated with the fast-slow continuum.

First lets plot out how differnt modes of life are distributed across PC1.

```
#put the mode-of-life data into a formate that can be plotted with the
#MultiDisPlot function
PCA moblist <- list(semi fossorial = pca data[pca data$PCA moblist == "semi fossorial",1],
                     pelagic = pca_data[pca_data$PCA_moblist == "pelagic",1],
                     terrestrial = pca_data[pca_data$PCA_moblist == "terrestrial",1],
                     semi_aquatic = pca_data[pca_data$PCA_moblist == "semi_aquatic",1],
                     volant = pca_data[pca_data$PCA_moblist == "volant",1],
                     demersal = pca_data[pca_data$PCA_moblist == "demersal",1],
                     arboreal = pca_data[pca_data$PCA_moblist == "arboreal",1],
                     sessile = pca_data[pca_data$PCA_moblist == "sessile",1]
#use the MultiDisPlot function to plot out the groups
#with credibility intervals
MultiDisPlot(PCA_moblist,
             yaxt = "n",
             xlab = "PC1"
             ylab = ""
             bty = "n")
#Plot the mode of life groups on y axis
tick_lables <- names(PCA_moblist)</pre>
```

```
axis(2,
             1:length(tick_lables),
            labels = tick_lables,
             cex.axis = 0.5)
#To plot the species on we need to convert the names into ther associated tick
mob_match <- as.vector(pca_data$PCA_moblist)</pre>
for(i in 1:(length(tick_lables))){
mob_match[mob_match == tick_lables[i]] <- i</pre>
mob_match <- as.numeric(mob_match)</pre>
pca_data$mob_match <- mob_match</pre>
#Plot the points onto the figure.
points(pca_data[pca_data$class_match == "Mammalia", "mob_match"] ~ pca_data[pca_data$class_match == "Mammalia", "mob_match"] ~ pca_data
                  col = mam col, pch = 20, cex = 0.5
points(pca_data[pca_data$class_match == "Aves", "mob_match"] ~ pca_data[pca_data$class_match == "Aves", 1
                  col = bird_col, pch = 20, cex = 0.5)
points(pca_data[pca_data$class_match == "Reptilia", "mob_match"] ~ pca_data[pca_data$class_match == "Rep
                  col = rep_col, pch = 20, cex = 0.5)
points(pca_data[pca_data$class_match == "Actinopterygii", "mob_match"] ~ pca_data[pca_data$class_match ==
points(pca_data[pca_data$class_match == "Gastropoda", "mob_match"] ~ pca_data[pca_data$class_match == "G
points(pca_data[pca_data$class_match == "Demospongiae", "mob_match"] ~ pca_data[pca_data$class_match == "Demospongiae", "mob_match == "Demospongiae", "mob_
points(pca_data[pca_data$class_match == "Anthozoa", "mob_match"] ~ pca_data[pca_data$class_match == "Ant
points(pca_data[pca_data$class_match == "Bivalvia", "mob_match"] ~ pca_data[pca_data$class_match == "Biv
```



Now lets test whether these differences of mode of life on the fast slow axis are differnt from each other whilst accounting for phylogentic relationship and population level variation. We do this using a MCMCglmm model both seperately for terrestrial groups and aquatic groups.

First we run the aquatic groups. To avoid long runs for this script we run it for one tree over two chains.

```
#subset to aquatic species
aquatic_res <- pca_data[pca_data$PCA_moblist == "sessile"</pre>
                         pca_data$PCA_moblist == "demersal"
                         | pca_data$PCA_moblist == "pelagic",]
#We set pelagic species as our base level.
aquatic_res$PCA_moblist <- factor(aquatic_res$PCA_moblist,</pre>
                                    levels = c("pelagic",
                                               "sessile",
                                               "demersal"))
#set a prior
prior < -list(R = list(V = 1/2, nu=0.002),
            G = list(G1=list(V = 1/2,
                              n = 1,
                              alpha.mu=rep(0,1),
                              alpha. V = diag(1)*10^3,
                      G1=list(V = 1/2,
                              n = 1,
                              alpha.mu=rep(0,1),
                              alpha. V= diag(1)*10^3)))
#run the analysis
#Chain 1
aquatic_pc1_c1 <- MCMCglmm(PC1 ~ PCA_moblist,</pre>
                      data = aquatic_res,
```

```
random=~animal + species_match,
                     pedigree = axis_trees[[1]],
                     prior = prior,
                     nitt = c(1100000),
                     burnin = 100000,
                     thin = 500,
                     verbose = F)
#Chain 2
aquatic_pc1_c2 <- MCMCglmm(PC1 ~ PCA_moblist,</pre>
                     data = aquatic_res,
                     random=~animal + species_match,
                     pedigree = axis_trees[[1]],
                     prior = prior,
                     nitt = c(1100000),
                     burnin = 100000,
                     thin = 500.
                     verbose = F)
#Check the chains converge
gelman.diag(mcmc.list(aquatic_pc1_c1$Sol,
                      aquatic_pc1_c2$Sol))
## Potential scale reduction factors:
##
                       Point est. Upper C.I.
##
## (Intercept)
                                1
                                            1
## PCA_moblistsessile
                                1
                                            1
## PCA_moblistdemersal
                                1
                                            1
##
## Multivariate psrf
gelman.diag(mcmc.list(aquatic_pc1_c1$VCV,
                      aquatic_pc1_c2$VCV))
## Potential scale reduction factors:
##
                 Point est. Upper C.I.
##
                                1.02
                       1.01
## animal
## species_match
                       1.00
                                  1.01
                       1.00
## units
                                  1.02
## Multivariate psrf
##
## 1
#summary
summary(aquatic_pc1_c1)
##
## Iterations = 100001:1099501
## Thinning interval = 500
## Sample size = 2000
```

```
##
   DIC: 142.0327
##
##
   G-structure: ~animal
##
##
          post.mean 1-95% CI u-95% CI eff.samp
##
              2.652 7.35e-06
                                 6.242
##
##
                  ~species_match
##
##
                 post.mean 1-95% CI u-95% CI eff.samp
                                                   2000
                    0.7488 3.795e-06
                                         1.613
##
  species_match
##
##
    R-structure: ~units
##
##
         post.mean 1-95% CI u-95% CI eff.samp
            0.4106 0.2177
                             0.6687
                                          1872
## units
##
   Location effects: PC1 ~ PCA_moblist
##
##
##
                       post.mean 1-95% CI u-95% CI eff.samp pMCMC
## (Intercept)
                         -1.0520 -3.7804
                                            1.4420
                                                         2027 0.399
                                                         2079 0.049 *
## PCA_moblistsessile
                                  0.1755
                                             5.4938
                          2.6764
## PCA moblistdemersal
                          1.6700
                                   0.1668
                                             3.2321
                                                         2000 0.036 *
## ---
## Signif. codes: 0 '***' 0.001 '**' 0.05 '.' 0.1 ' ' 1
#Calculate phyogentic signal, population level variation and residual variation.
aquatic_pc1_vcv_phylo <- aquatic_pc1_c1$VCV[,1]</pre>
aquatic_pc1_vcv_spec <- aquatic_pc1_c1$VCV[,2]</pre>
aquatic_pc1_vcv_units <- aquatic_pc1_c1$VCV[,3]</pre>
#Phyogentic signal
aquatic_pc1_phylo <- aquatic_pc1_vcv_phylo/</pre>
                            c(aquatic_pc1_vcv_phylo
                              + aquatic_pc1_vcv_spec
                              + aquatic_pc1_vcv_units)
#Population level variation
aquatic_pc1_spec <- aquatic_pc1_vcv_spec/</pre>
                            c(aquatic_pc1_vcv_phylo
                              + aquatic_pc1_vcv_spec
                              + aquatic_pc1_vcv_units)
#Residual
aquatic_pc1_units <- aquatic_pc1_vcv_units/</pre>
                            c(aquatic_pc1_vcv_phylo
                              + aquatic_pc1_vcv_spec
                              + aquatic_pc1_vcv_units)
hdr(aquatic_pc1_phylo)$mode
```

[1] 0.7285542

```
hdr(aquatic_pc1_spec)$mode
## [1] 0.1366372
hdr(aquatic_pc1_units)$mode
## [1] 0.09454943
Now lets do the same for terrestrial species.
##terrestiral species
ter_res <- pca_data[pca_data$PCA_moblist == "terrestrial"</pre>
                     pca_data$PCA_moblist == "arboreal"
                     pca_data$PCA_moblist == "volant"
                     pca_data$PCA_moblist == "semi_aquatic"
                     pca_data$PCA_moblist == "semi_fossorial",]
ter_res$PCA_moblist <- factor(ter_res$PCA_moblist,</pre>
                                levels = c("terrestrial",
                                            "arboreal",
                                            "volant",
                                            "semi_aquatic",
                                            "semi_fossorial"))
#set a prior
prior < -list(R = list(V = 1/2, nu=0.002),
            G = list(G1=list(V = 1/2,
                              n = 1,
                              alpha.mu=rep(0,1),
                              alpha.V= diag(1)*10^3),
                      G1=list(V = 1/2,
                              n = 1
                              alpha.mu=rep(0,1),
                              alpha. V= diag(1)*10^3)))
ter_pc1_c1 <- MCMCglmm(PC1 ~ PCA_moblist,</pre>
                     data = ter_res,
                     random=~animal + species_match,
                     pedigree = axis_trees[[1]],
                    prior = prior,
                    nitt = c(1100000),
                     burnin = 100000,
                     thin = 500,
                     verbose = F)
ter_pc1_c2 <- MCMCglmm(PC1 ~ PCA_moblist,</pre>
```

data = ter_res,

prior = prior,
nitt = c(1100000),

random=~animal + species_match,
pedigree = axis_trees[[1]],

```
burnin = 100000,
                    thin = 500,
                    verbose = F)
#Check the chains converge
gelman.diag(mcmc.list(ter_pc1_c1$Sol,
                      ter_pc1_c2$Sol))
## Potential scale reduction factors:
##
##
                             Point est. Upper C.I.
## (Intercept)
                                      1
                                              1.02
## PCA_moblistarboreal
                                              1.00
                                      1
## PCA_moblistvolant
                                              1.00
                                      1
## PCA_moblistsemi_aquatic
                                    1
                                              1.00
## PCA_moblistsemi_fossorial
                                    1
                                              1.01
## Multivariate psrf
##
## 1
gelman.diag(mcmc.list(ter_pc1_c1$VCV,
                     ter_pc1_c2$VCV))
## Potential scale reduction factors:
##
##
                Point est. Upper C.I.
## animal
                         1
                                 1.01
                                 1.00
## species match
                         1
## units
                                 1.00
## Multivariate psrf
##
## 1
summary(ter_pc1_c1)
##
  Iterations = 100001:1099501
## Thinning interval = 500
## Sample size = 2000
##
## DIC: 319.4351
##
## G-structure: ~animal
##
##
         post.mean 1-95% CI u-95% CI eff.samp
## animal 10.62
                       6.018
                                16.23
                                          2000
##
##
                 ~species_match
##
##
                post.mean 1-95% CI u-95% CI eff.samp
## species_match 0.1346 6.456e-07 0.3288
##
```

```
##
    R-structure: ~units
##
         post.mean 1-95% CI u-95% CI eff.samp
##
            0.1762
                     0.1314
                               0.2165
                                           2000
## units
##
   Location effects: PC1 ~ PCA moblist
##
##
##
                              post.mean 1-95% CI u-95% CI eff.samp pMCMC
## (Intercept)
                                0.57368 -4.64811 6.41147
                                                                2000 0.844
## PCA_moblistarboreal
                                0.23139 -0.63376 1.15915
                                                                1913 0.616
## PCA_moblistvolant
                                0.05750 -2.06626 2.14367
                                                                2000 0.950
## PCA_moblistsemi_aquatic
                                                                2000 0.966
                                0.03251 -1.76657 1.72798
## PCA_moblistsemi_fossorial -0.40138 -1.66672 0.96700
                                                                2000 0.555
ter_pc1_vcv_phylo <- ter_pc1_c1$VCV[,1]</pre>
ter_pc1_vcv_spec <- ter_pc1_c1$VCV[,2]</pre>
ter_pc1_vcv_units <- ter_pc1_c1$VCV[,3]</pre>
ter_pc1_phylo <- ter_pc1_vcv_phylo/</pre>
  c(ter_pc1_vcv_phylo + ter_pc1_vcv_spec + ter_pc1_vcv_units)
ter_pc1_spec <- ter_pc1_vcv_spec/</pre>
  c(ter_pc1_vcv_phylo + ter_pc1_vcv_spec + ter_pc1_vcv_units)
ter_pc1_units <- ter_pc1_vcv_units/</pre>
  c(ter_pc1_vcv_phylo + ter_pc1_vcv_spec + ter_pc1_vcv_units)
hdr(ter_pc1_phylo)$mode
## [1] 0.9760544
hdr(ter_pc1_spec)$mode
## [1] 0.0006235144
hdr(ter_pc1_units)$mode
## [1] 0.01518916
```

Metabolic rate analysis

Next we analysis how metabolic rate is associated with the fast slow continuum by regressing mass soecific metabolic rate against PC1.

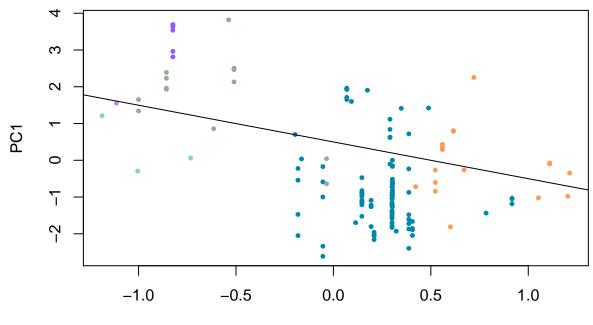
```
n = 1,
                              alpha.mu=rep(0,1),
                              alpha.V= diag(1)*10^3),
                      G1=list(V = 1/2,
                              n = 1,
                              alpha.mu=rep(0,1),
                              alpha. V= diag(1)*10^3)))
met_mod_ch1 <- MCMCglmm(PC1 ~ log10(PCA_met) ,</pre>
                    data = pca_data_met,
                    random=~animal + species_match,
                    pedigree = axis_trees[[1]],
                    prior = prior_met,
                    nitt = c(1100000),
                    burnin = 100000,
                    thin = 500,
                    verbose = F)
met_mod_ch2 <- MCMCglmm(PC1 ~ log10(PCA_met) ,</pre>
                    data = pca_data_met,
                    random=~animal + species_match,
                    pedigree = axis_trees[[1]],
                    prior = prior_met,
                    nitt = c(1100000),
                    burnin = 100000,
                    thin = 500,
                    verbose = F)
#Check the chains converge
gelman.diag(mcmc.list(met_mod_ch1$Sol,
                      met_mod_ch2$Sol))
## Potential scale reduction factors:
##
##
                  Point est. Upper C.I.
## (Intercept)
                            1
                                    1.00
## log10(PCA_met)
                                    1.01
                            1
## Multivariate psrf
##
## 1
gelman.diag(mcmc.list(met_mod_ch1$VCV,
                      met_mod_ch2$VCV))
## Potential scale reduction factors:
##
##
                 Point est. Upper C.I.
## animal
                          1
                                      1
## species_match
                           1
                                      1
## units
                           1
                                      1
```

```
##
## Multivariate psrf
##
## 1
summary(met_mod_ch1)
##
    Iterations = 100001:1099501
##
## Thinning interval = 500
##
   Sample size = 2000
##
##
  DIC: 221.3889
##
##
   G-structure: ~animal
##
##
          post.mean 1-95% CI u-95% CI eff.samp
##
  animal
               7.18
                       3.191
                                 11.45
                                           2000
##
##
                  ~species_match
##
                 post.mean 1-95% CI u-95% CI eff.samp
##
##
  species_match
                  0.06473 5.938e-09 0.2389
##
##
   R-structure: ~units
##
##
         post.mean 1-95% CI u-95% CI eff.samp
## units
          0.2072
                   0.1556
                              0.2645
##
##
    Location effects: PC1 ~ log10(PCA_met)
##
##
                  post.mean 1-95% CI u-95% CI eff.samp pMCMC
                    0.59897 -3.43038 4.11072
                                                   2000 0.753
## (Intercept)
## log10(PCA_met) -1.02322 -1.88377 -0.09075
                                                   2000 0.022 *
## Signif. codes: 0 '***' 0.001 '**' 0.05 '.' 0.1 ' ' 1
met_mod_phylo <- met_mod_ch1$VCV[,1]</pre>
met_mod_spec <- met_mod_ch1$VCV[,2]</pre>
met_mod_units <- met_mod_ch1$VCV[,3]</pre>
met_mod_phylo_H <- met_mod_phylo/c(met_mod_phylo + met_mod_spec + met_mod_units)</pre>
met_mod_spec_H <- met_mod_spec/c(met_mod_phylo + met_mod_spec + met_mod_units)</pre>
met_mod_units_H <- met_mod_units/c(met_mod_phylo + met_mod_spec + met_mod_units)</pre>
hdr(met_mod_phylo_H)$mode
## [1] 0.9711183
hdr(met_mod_spec_H)$mode
## [1] 0.0002180417
hdr(met_mod_units_H)$mode
```

[1] 0.02678225

Lets also plot out how metabolic rate changes with a species position on the fast slow continum.

```
plot(log10(pca_data_met$PCA_met),
     pca_data_met$PC1,
     pch = 16,
     col = "white",
     cex = 0.1,
     xlab = "Log10 Mass specific metabolic rate (W/g)",
    ylab = "PC1")
points(log10(pca_data_met[pca_data_met$class_match == "Mammalia", "PCA_met"]),
       pca_data_met[pca_data_met$class_match == "Mammalia","PC1"],
       col = mam_col, pch = 20, cex = 0.8)
points(log10(pca_data_met[pca_data_met$class_match == "Aves", "PCA_met"]),
       pca_data_met[pca_data_met$class_match == "Aves", "PC1"],
       col = bird_col,
       pch = 20,
       cex = 0.8)
points(log10(pca_data_met[pca_data_met$class_match == "Reptilia", "PCA_met"]),
       pca_data_met[pca_data_met$class_match == "Reptilia","PC1"],
       col = rep_col,
       pch = 20,
       cex = 0.8)
points(log10(pca_data_met[pca_data_met$class_match == "Actinopterygii", "PCA_met"]),
       pca_data_met[pca_data_met$class_match == "Actinopterygii","PC1"],
       col = fish_col,
       pch = 20,
       cex = 0.8)
points(log10(pca_data_met[pca_data_met$class_match == "Gastropoda", "PCA_met"]),
       pca_data_met[pca_data_met$class_match == "Gastropoda","PC1"],
       col = gast_col,
       pch = 20,
       cex = 0.8)
points(log10(pca_data_met[pca_data_met$class_match == "Demospongiae", "PCA_met"]),
       pca_data_met[pca_data_met$class_match == "Demospongiae","PC1"],
       col = sponge_col,
       pch = 20,
       cex = 0.8)
points(log10(pca_data_met[pca_data_met$class_match == "Anthozoa", "PCA_met"]), pca_data_met[pca_data_met
points(log10(pca_data_met[pca_data_met$class_match == "Bivalvia", "PCA_met"]), pca_data_met[pca_data_met
abline(hdr(met_mod_ch1$Sol[,1])$mode, hdr(met_mod_ch1$Sol[,2])$mode)
```



Log10 Mass specific metabolic rate (W/g)

Reporductive productivity analysis

As reporductive output is only defined by numbers of offspring in matrix models we also tested if reporductive productivity, the amount of reporductive mass produced per year, would be more strongly associated with the fast-slow continuum as represented by PC1, then we found in the main PCA.

```
pca_data_repo <- na.omit(pca_data[,c("PC1",</pre>
                                       "species_match",
                                       "class_match",
                                       "animal",
                                       "PCA_repo")])
#set a prior
prior_repo <-list(R = list(V = 1/2, nu=0.002),
            G = list(G1=list(V = 1/2,
                              n = 1,
                              alpha.mu=rep(0,1),
                              alpha. V= diag(1)*10^3),
                      G1=list(V = 1/2,
                              n = 1,
                              alpha.mu=rep(0,1),
                              alpha. V= diag(1)*10^3)))
egg_size_ch1 <- MCMCglmm(PC1 ~ PCA_repo,
                      data = pca_data_repo,
                      random=~animal + species_match,
                      pedigree = axis_trees[[1]],
                      prior = prior_repo,
                      nitt = c(11000000),
```

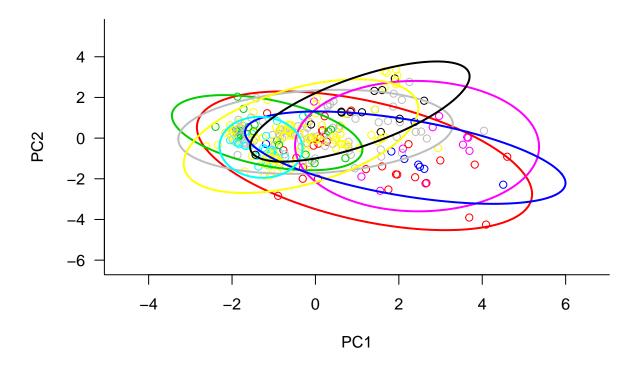
```
burnin = 1000000,
                     thin = 5000,
                     verbose = F)
egg_size_ch2 <- MCMCglmm(PC1 ~ PCA_repo,
                     data = pca_data_repo,
                     random=~animal + species_match,
                     pedigree = axis_trees[[1]],
                    prior = prior_repo,
                     nitt = c(11000000),
                     burnin = 1000000,
                     thin = 5000,
                     verbose = F)
gelman.diag(mcmc.list(egg_size_ch1$Sol,
                      egg_size_ch2$Sol))
## Potential scale reduction factors:
##
               Point est. Upper C.I.
##
                               1.00
## (Intercept)
                      1
                                1.01
## PCA_repo
                        1
## Multivariate psrf
##
## 1
gelman.diag(mcmc.list(egg_size_ch1$VCV,
                      egg_size_ch2$VCV))
## Potential scale reduction factors:
##
                 Point est. Upper C.I.
## animal
                                  1.00
                         1
## species_match
                                  1.00
## units
                                  1.01
##
## Multivariate psrf
## 1
summary(egg_size_ch1)
##
## Iterations = 1000001:10995001
## Thinning interval = 5000
## Sample size = 2000
##
## DIC: 361.6741
##
## G-structure: ~animal
##
          post.mean 1-95% CI u-95% CI eff.samp
## animal
          9.556 4.981 14.31
                                          2000
```

```
##
##
                  ~species_match
##
##
                 post.mean 1-95% CI u-95% CI eff.samp
##
  species_match
                    0.1341 1.628e-10 0.3386
##
##
   R-structure:
                 ~units
##
##
        post.mean 1-95% CI u-95% CI eff.samp
##
  units
           0.1999 0.1537
                               0.245
                                         1831
##
   Location effects: PC1 ~ PCA_repo
##
##
##
              post.mean 1-95% CI u-95% CI eff.samp pMCMC
                                               2000 0.675
## (Intercept)
                  0.9176 -3.1126
                                    5.5581
## PCA_repo
                 -0.7736 -1.4770 -0.0721
                                               2000 0.027 *
## ---
## Signif. codes: 0 '***' 0.001 '**' 0.05 '.' 0.1 ' ' 1
egg_size_phylo <- egg_size_ch1$VCV[,1]/(egg_size_ch1$VCV[,1] +
                                         egg_size_ch1$VCV[,2] +
                                         egg_size_ch1$VCV[,3])
egg_size_species <- egg_size_ch1$VCV[,2]/(egg_size_ch1$VCV[,1] +
                                         egg_size_ch1$VCV[,2] +
                                         egg_size_ch1$VCV[,3])
egg_size_units <- egg_size_ch1$VCV[,3]/(egg_size_ch1$VCV[,1] +
                                         egg_size_ch1$VCV[,2] +
                                         egg_size_ch1$VCV[,3])
hdr(egg_size_phylo)$mode
## [1] 0.9725956
hdr(egg_size_species)$mode
## [1] 0.0007247752
hdr(egg_size_units)$mode
## [1] 0.01931435
```

Ellipses

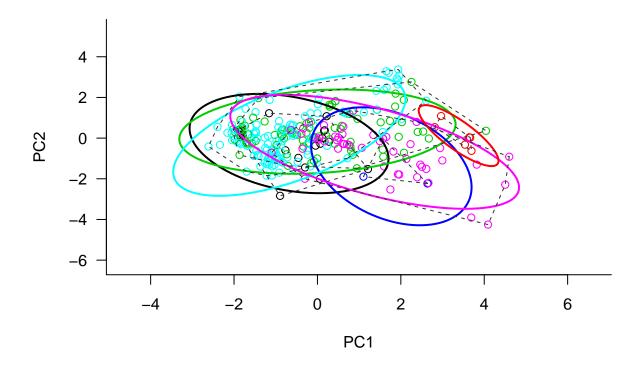
To test how different groupings are associated with PCA space we fit a series of ellipses using the SIBER package. Whilst SIBER is developed for fitting Baysian ellipses for stable isotope data it can be applied in the same way. TO get the figures to the standard seen in the paper I used inkscape.

Mode of life Ellipse

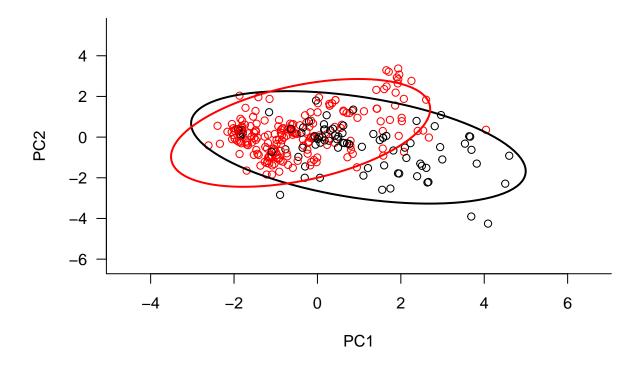


Taxinomic groupings Ellipse

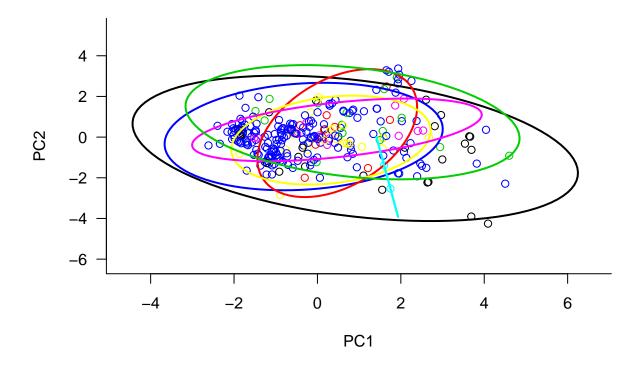
```
sidpca <- data.frame(iso1 = siber_pca_data$PC1,</pre>
                      iso2 = siber_pca_data$PC2,
                      group = as.numeric(siber_pca_data$class_match),
                      community = rep(1,length(siber_pca_data$class_match)))
siber.plots <- createSiberObject(sidpca)</pre>
#plot for taxa
plotSiberObject(siber.plots,
                   ax.pad = 2,
                  hulls = F,
                   community.hulls.args,
                   ellipses = T,
                   group.ellipses.args,
                   group.hulls = T,
                   group.hull.args,
                   bty = L'',
                   iso.order = c(1,2),
                   xlab = "PC1",
                  ylab = "PC2"
```



Thermal groupings Ellipse



IUCN Ellipse



Ellipse overlap

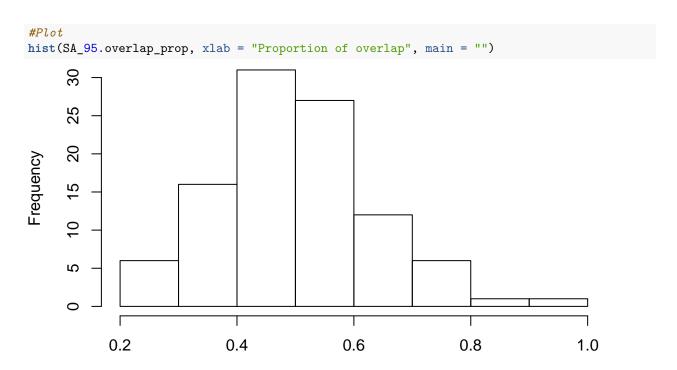
Using a Baysian resampling approach we see how much each of the elliplies overlap with each other and plot these out. These calculations require jags. For more see the SIBER package

Mode of life ellipse overlap

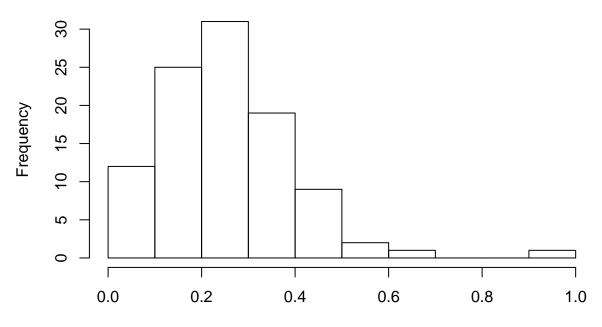
```
group.ML <- groupMetricsML(siber.plots)</pre>
group.MLmob <- groupMetricsML(siber.mob)</pre>
# options for running jags
parms <- list()</pre>
parms$n.iter <- 2 * 10^4 # number of iterations to run the model for
parms$n.burnin <- 1 * 10^3 # discard the first set of values
parms$n.thin <- 10
                       # thin the posterior by this many
parms$n.chains <- 2</pre>
                             # run this many chains
# define the priors
priors <- list()</pre>
priors$R <- 1 * diag(2)</pre>
priors$k <- 2
priors$tau.mu <- 1.0E-3</pre>
ellipses.posterior_mob <- siberMVN(siber.mob, parms, priors)</pre>
```

```
##
      Resolving undeclared variables
##
      Allocating nodes
##
  Graph information:
      Observed stochastic nodes: 28
##
##
      Unobserved stochastic nodes: 3
##
      Total graph size: 43
## Initializing model
##
  Compiling model graph
##
      Resolving undeclared variables
      Allocating nodes
##
##
  Graph information:
      Observed stochastic nodes: 21
##
##
      Unobserved stochastic nodes: 3
##
      Total graph size: 36
##
## Initializing model
##
## Compiling model graph
##
      Resolving undeclared variables
##
      Allocating nodes
## Graph information:
##
      Observed stochastic nodes: 82
      Unobserved stochastic nodes: 3
##
##
      Total graph size: 97
##
## Initializing model
##
## Compiling model graph
##
      Resolving undeclared variables
##
      Allocating nodes
##
  Graph information:
##
      Observed stochastic nodes: 12
##
      Unobserved stochastic nodes: 3
##
      Total graph size: 27
##
## Initializing model
##
  Compiling model graph
##
      Resolving undeclared variables
      Allocating nodes
##
## Graph information:
##
      Observed stochastic nodes: 95
##
      Unobserved stochastic nodes: 3
##
      Total graph size: 110
##
## Initializing model
##
## Compiling model graph
##
      Resolving undeclared variables
      Allocating nodes
##
## Graph information:
##
      Observed stochastic nodes: 8
```

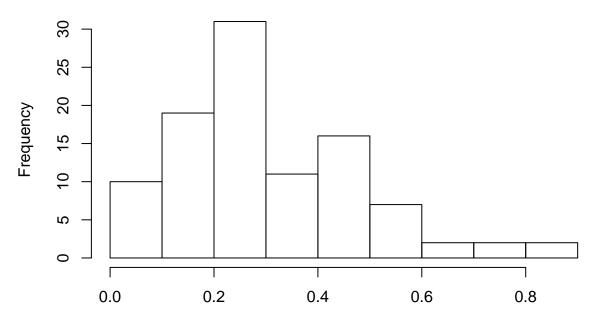
```
##
      Unobserved stochastic nodes: 3
##
      Total graph size: 23
##
## Initializing model
##
## Compiling model graph
      Resolving undeclared variables
      Allocating nodes
##
## Graph information:
##
      Observed stochastic nodes: 27
##
      Unobserved stochastic nodes: 3
##
      Total graph size: 42
##
## Initializing model
##
## Compiling model graph
##
      Resolving undeclared variables
##
      Allocating nodes
## Graph information:
##
      Observed stochastic nodes: 12
##
      Unobserved stochastic nodes: 3
##
      Total graph size: 27
##
## Initializing model
# The first ellipse is referenced using a character string representation where
# in "x.y", "x" is the community, and "y" is the group within that community.
# So in this example: community 1, group 1
#ellipse group numbers
ellipse_sessile <- "1.1"
ellipse_arboreal <- "1.2"
ellipse_benthic <- "1.3"
ellipse_volant <- "1.4"
ellipse_semiaquatic <- "1.5"
ellipse_terrestrial <- "1.6"
ellipse_pelagic <- "1.7"
ellipse_semifossorial <- "1.8"
#####sessile - arboreal
SA_95.overlap <- bayesianOverlap(ellipse_sessile,
                                   ellipse_arboreal,
                                   ellipses.posterior_mob,
                                   draws = 100,
                                   p.interval = 0.95,
                                   n = 100)
SA_95.overlap_prop <- vector()</pre>
for(i in 1:length(SA_95.overlap$)){
SA_95.overlap_prop[i] <- SA_95.overlap[i]/min(SA_95.overlap[i,1:2])
}
```



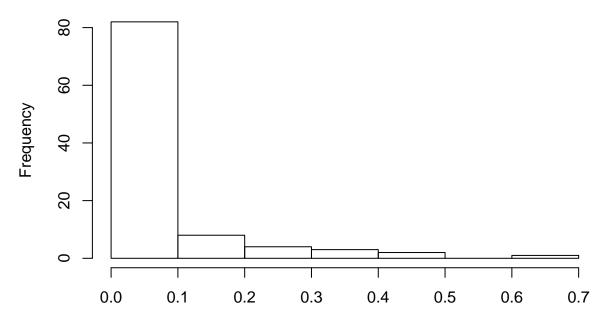
Sessile – Arboreal Overlap 200.00 0.25 0.50 0.75 1.00 Overlap



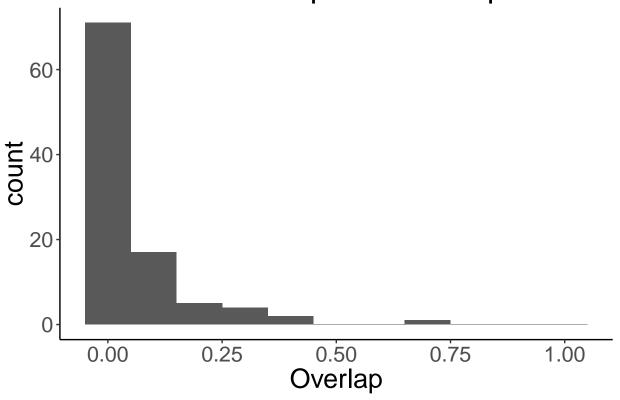
Sessile – Demersal Overlap 30100.00 0.25 0.50 0.75 1.00 Overlap

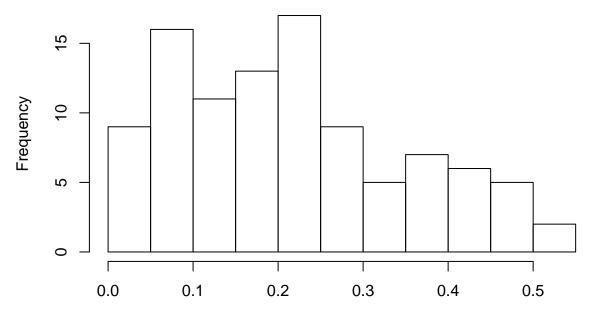


Sessile – Volant Overlap 201500.00 0.25 0.50 0.75 1.00 Overlap

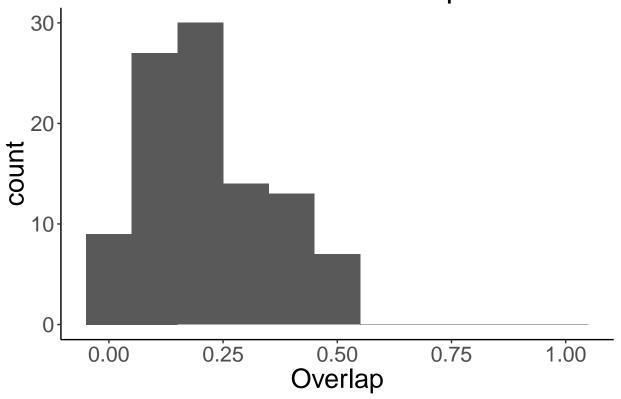


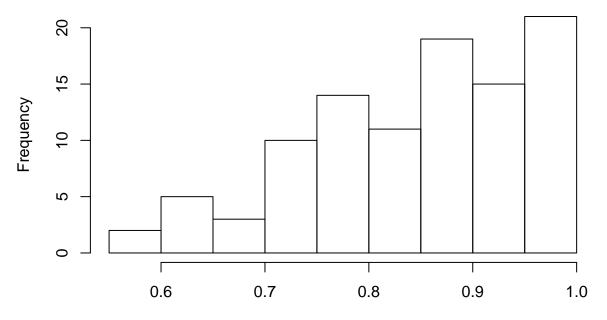
Sessile – Semiaquatic Overlap



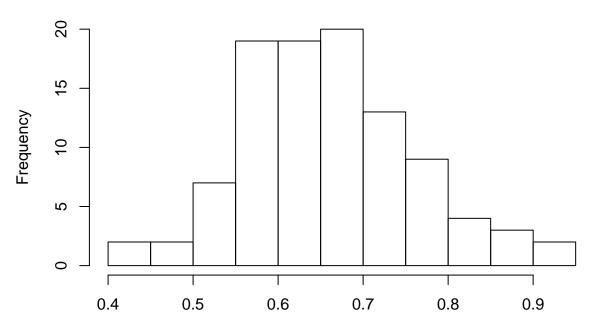


Sessile - terrestrial Overlap

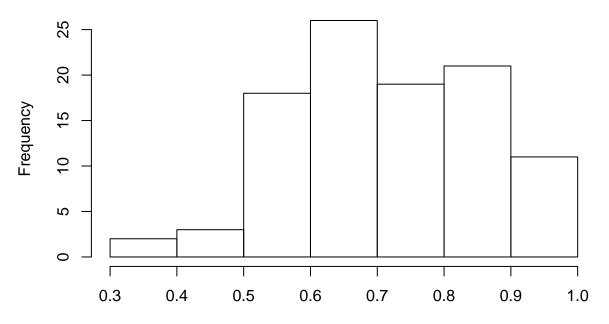




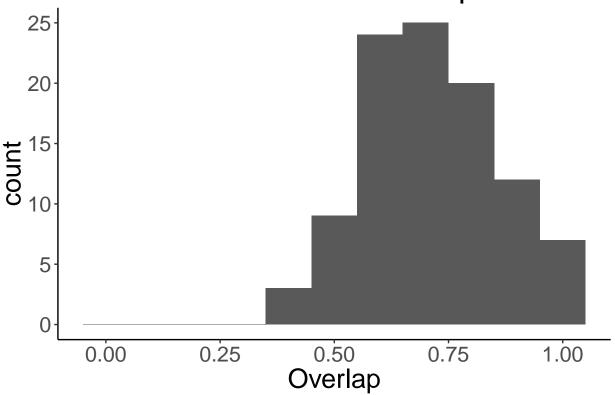
Sessile – pelagic Overlap 30100.00 0.25 0.50 0.75 1.00 Overlap

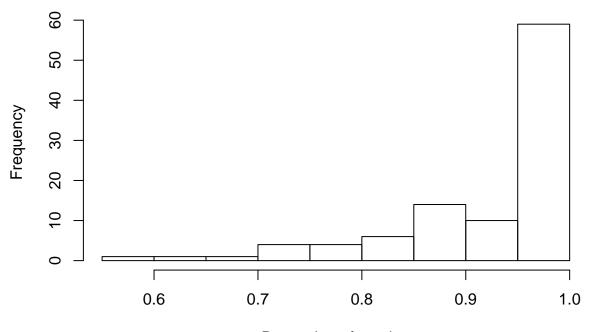


Sessile – semifossorial Overlap 30100.00 0.25 0.50 0.75 1.00 Overlap

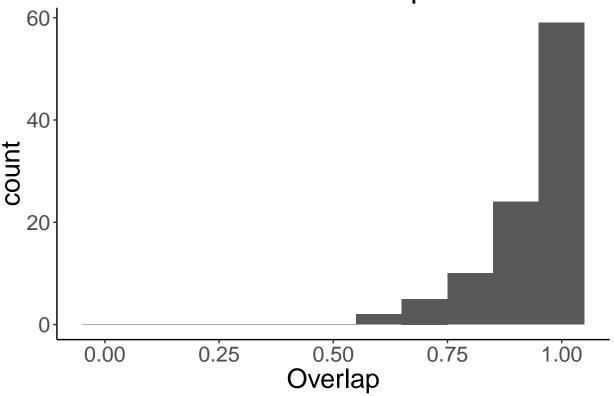


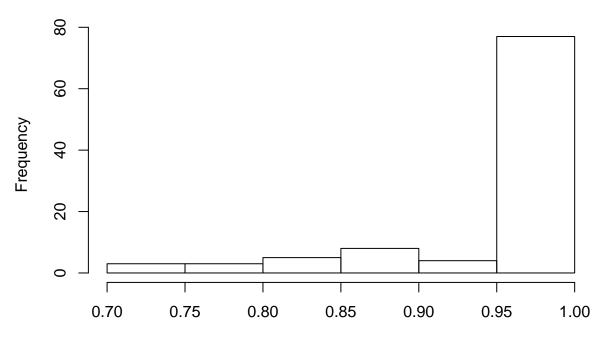
Arboreal – demersal Overlap



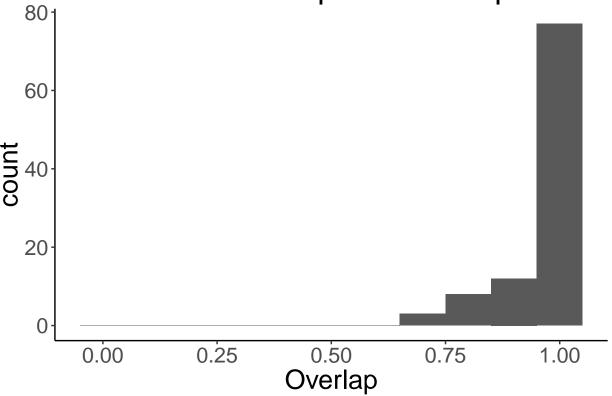


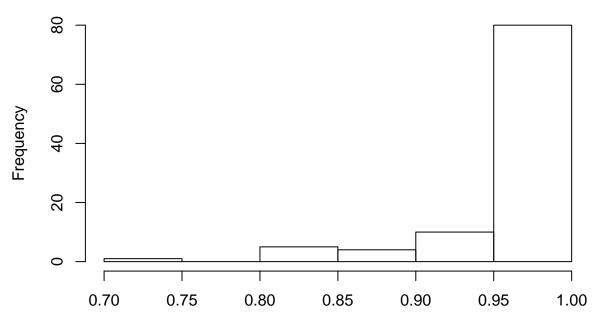
Arboreal - volant Overlap



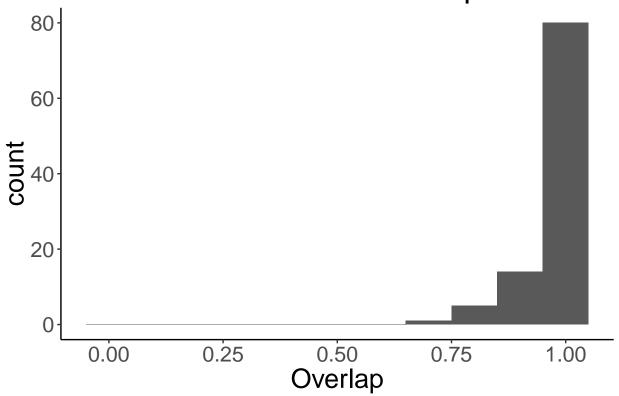


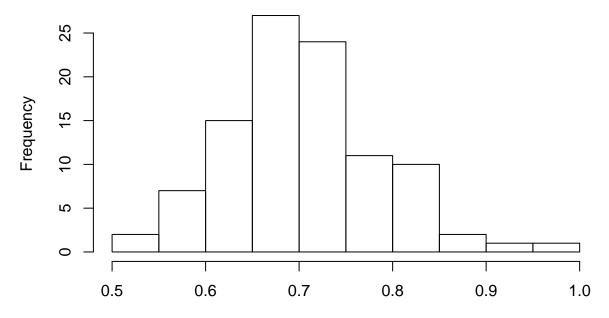
Arboreal – semiaquatic Overlap



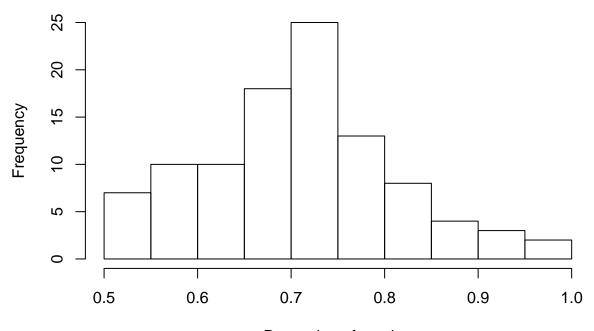


Arboreal – terrestrial Overlap

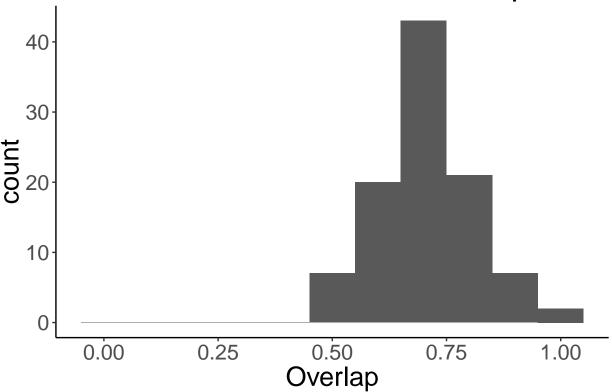


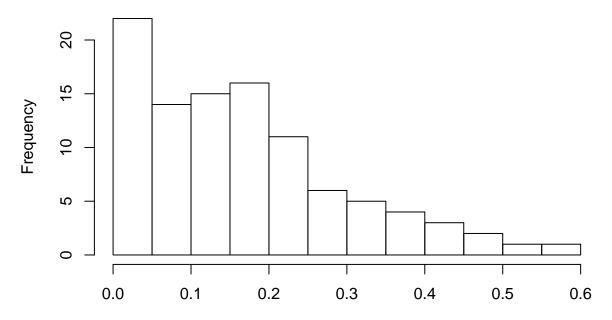


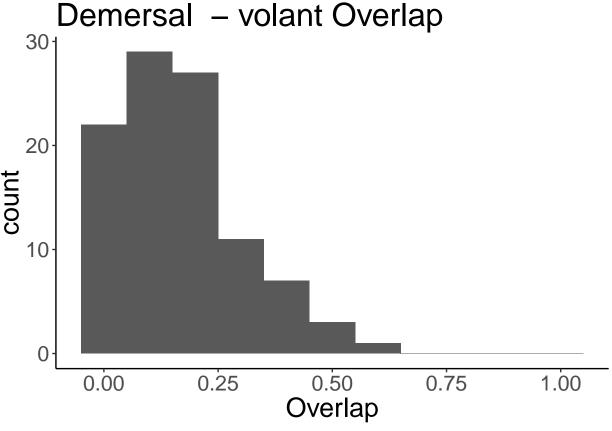
Arboreal – pelagic Overlap 504020100.00 0.25 0.50 0.75 1.00 Overlap

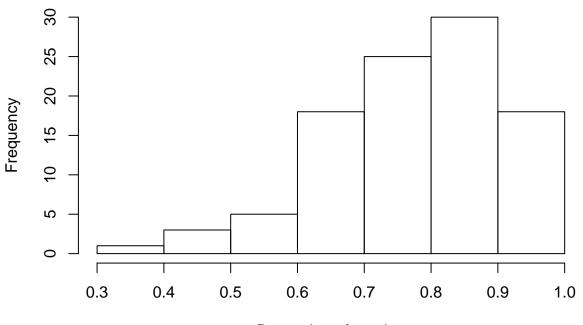


Arboreal – semifossorial Overlap

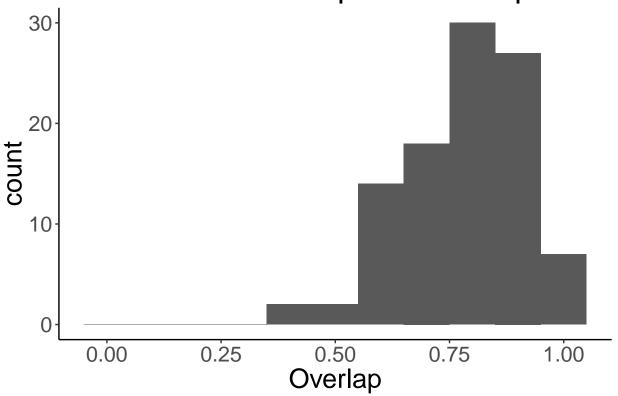


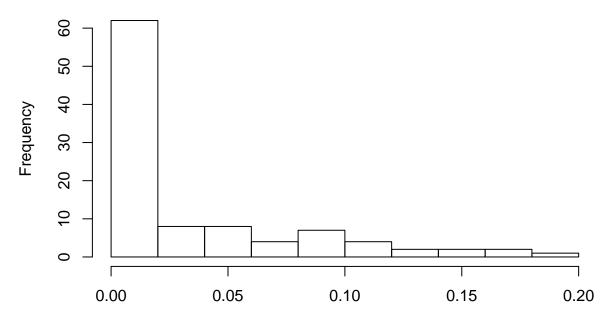




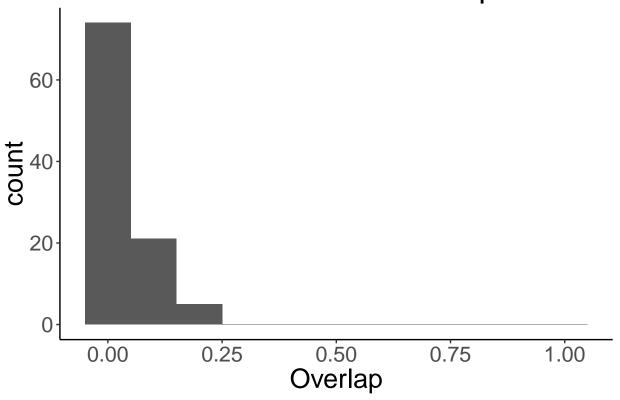


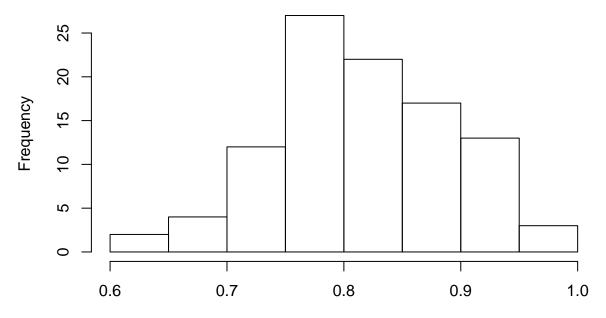
Demersal – semiaquatic Overlap



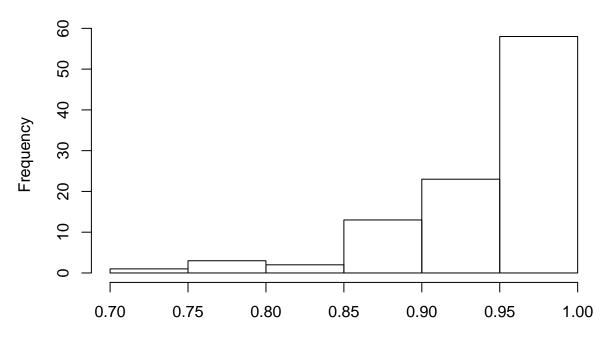


Demersal – terrestrial Overlap

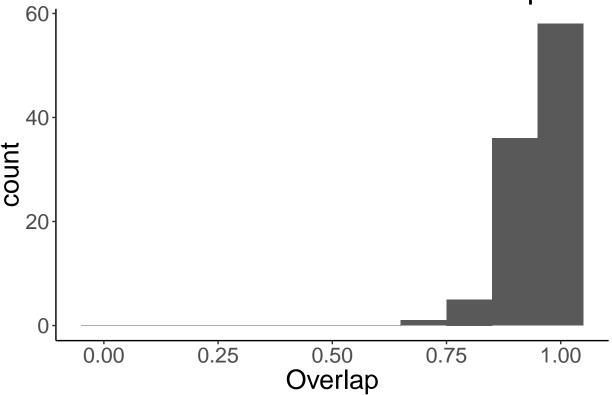


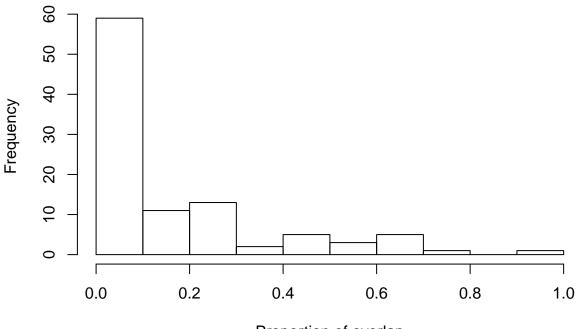


Demersal – pelagic Overlap 50 40 40 10 0.00 0.25 0.50 0.75 1.00

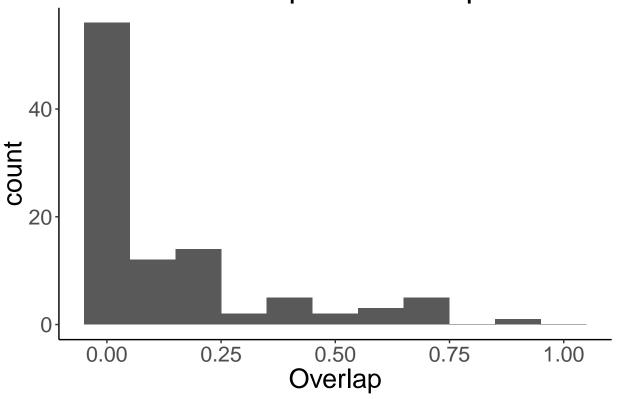


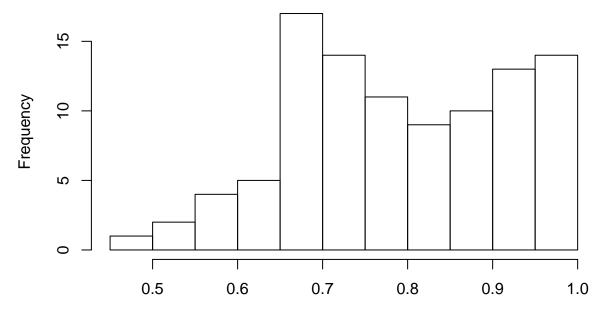
Demersal – semifossorial Overlap



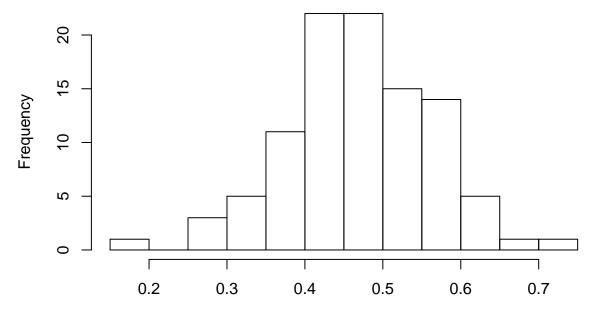


Volant – semiaquatic Overlap

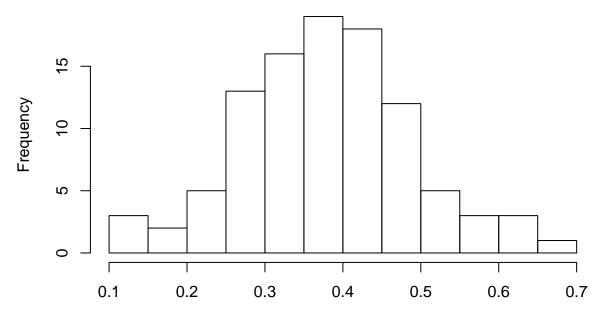




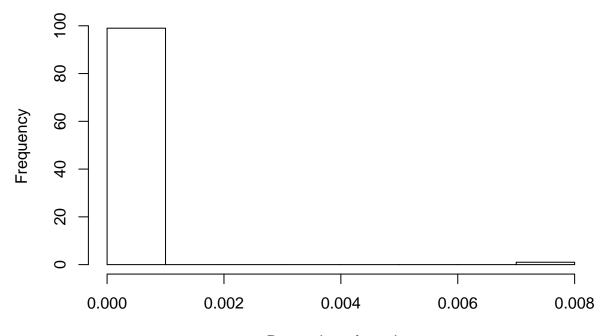
volant – terrestrial Overlap 30 10 0.00 0.25 0.50 0.75 1.00 Overlap



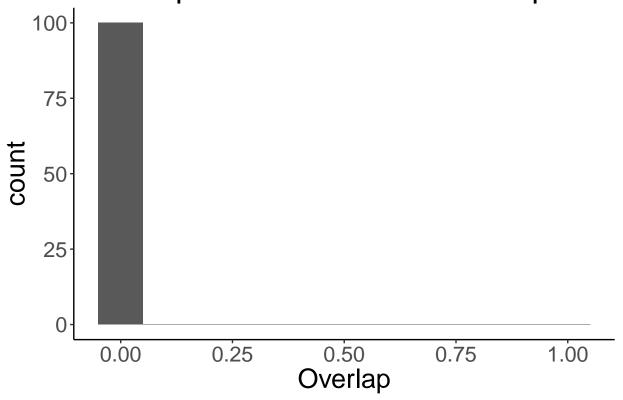
Volant – pelagic Overlap 30100.00 0.25 0.50 0.75 1.00 Overlap

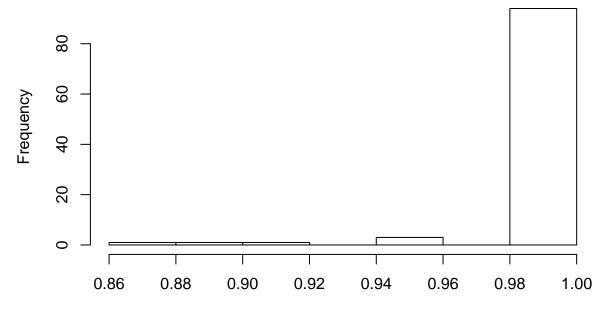


Demersal – semiaquatic Overlap 30 10 0.00 0.25 0.50 0.75 1.00

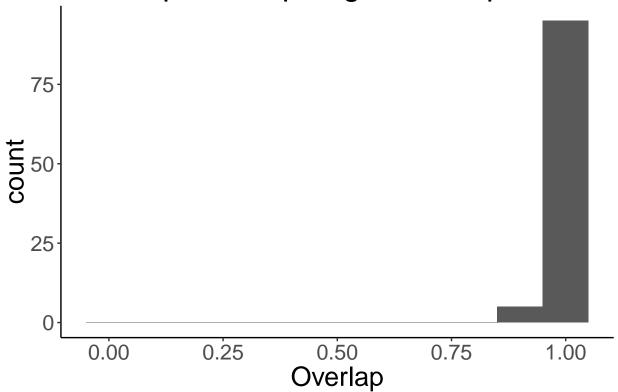


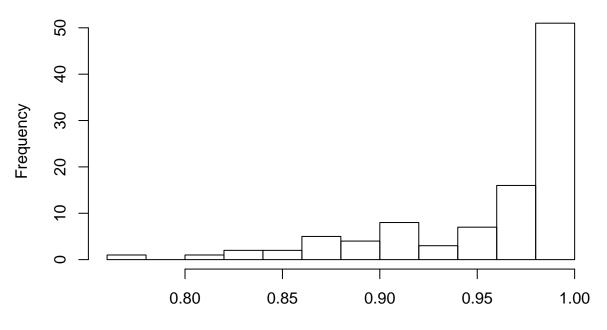
Semiaquatic – terrestrial Overlap



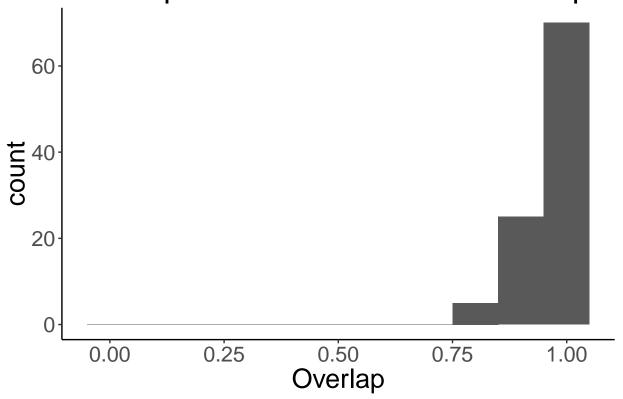


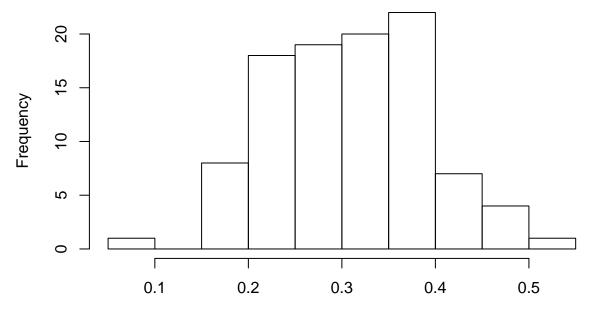
Semiaquatic – pelagic Overlap



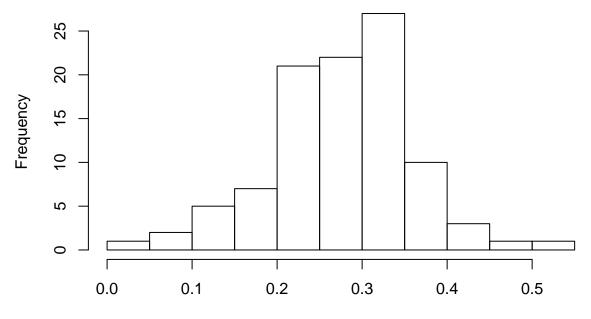


Semiaquatic - semifossorial Overlap

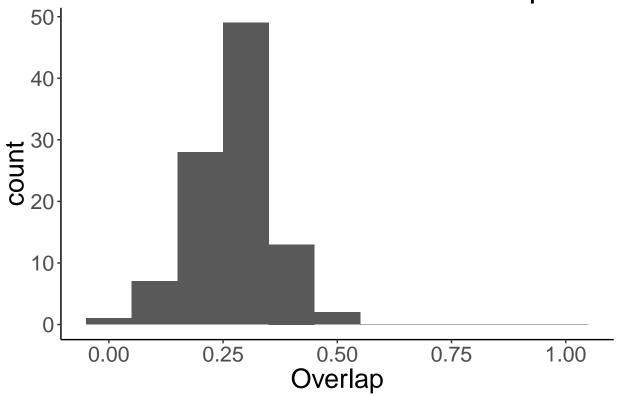


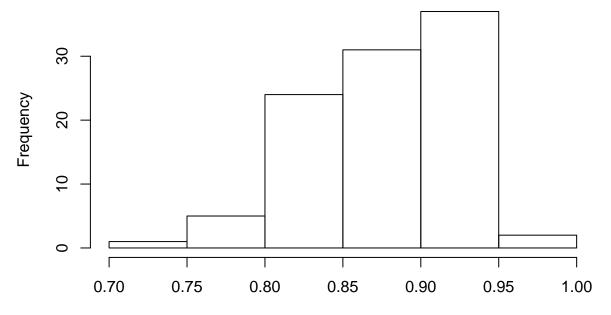


terrestrial – pelagic Overlap 30100.00 0.25 0.50 0.75 1.00 Overlap

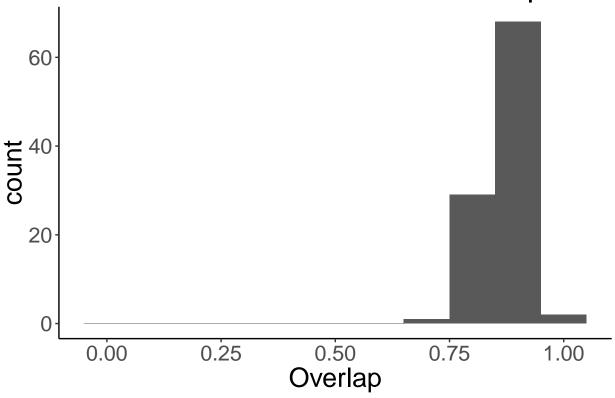


terrestrial - semifossorial Overlap





terrestrial - semifossorial Overlap

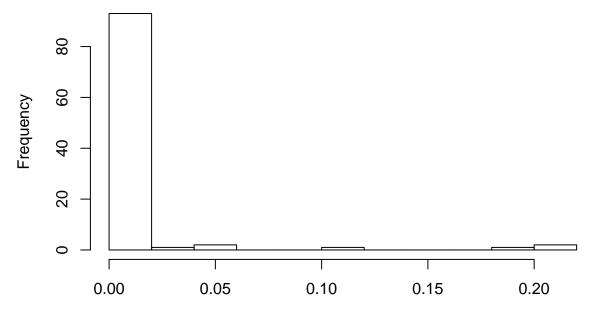


Ellipse overlap calculations for taxa

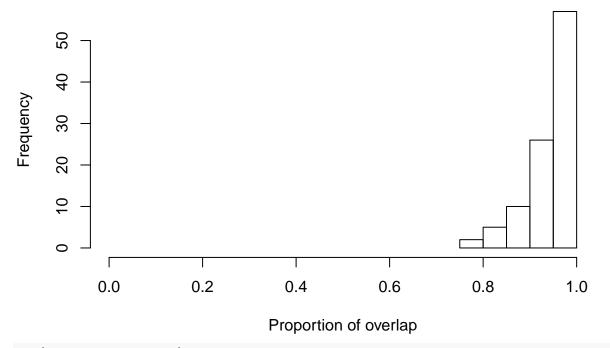
```
group.MLtaxa <- groupMetricsML(siber.plots)</pre>
group.MLmob <- groupMetricsML(siber.mob)</pre>
# options for running jags
parms <- list()</pre>
parms$n.iter <- 2 * 10^4 # number of iterations to run the model for
parms$n.burnin <- 1 * 10^3 # discard the first set of values
                        # thin the posterior by this many
parms$n.thin <- 10
parms$n.chains <- 2</pre>
                              # run this many chains
# define the priors
priors <- list()</pre>
priors R \leftarrow 1 * diag(2)
priors$k <- 2</pre>
priors$tau.mu <- 1.0E-3</pre>
ellipses.posterior <- siberMVN(siber.plots,</pre>
                                  parms,
                                  priors)
```

```
## Compiling model graph
##
      Resolving undeclared variables
##
      Allocating nodes
## Graph information:
##
      Observed stochastic nodes: 21
##
      Unobserved stochastic nodes: 3
##
      Total graph size: 36
##
## Initializing model
##
  Compiling model graph
##
      Resolving undeclared variables
      Allocating nodes
##
## Graph information:
##
      Observed stochastic nodes: 81
##
      Unobserved stochastic nodes: 3
##
      Total graph size: 96
##
## Initializing model
##
##
  Compiling model graph
##
      Resolving undeclared variables
      Allocating nodes
##
## Graph information:
      Observed stochastic nodes: 3
##
      Unobserved stochastic nodes: 3
##
##
      Total graph size: 18
##
## Initializing model
##
## Compiling model graph
##
      Resolving undeclared variables
##
      Allocating nodes
## Graph information:
##
      Observed stochastic nodes: 121
##
      Unobserved stochastic nodes: 3
##
      Total graph size: 136
##
## Initializing model
##
  Compiling model graph
##
      Resolving undeclared variables
      Allocating nodes
##
## Graph information:
##
      Observed stochastic nodes: 48
##
      Unobserved stochastic nodes: 3
##
      Total graph size: 63
##
## Initializing model
##
## Compiling model graph
##
      Resolving undeclared variables
##
      Allocating nodes
## Graph information:
```

```
Observed stochastic nodes: 6
##
      Unobserved stochastic nodes: 3
##
##
      Total graph size: 21
##
## Initializing model
# The first ellipse is referenced using a character string representation where
\# in "x.y", "x" is the community, and "y" is the group within that community.
# So in this example: community 1, group 1
#Actinopteryqii
ellipse_Actinopterygii <- "1.1"
#Anthozoa
ellipse_Anthozoa <- "1.2"
#Aves
ellipse_Aves <- "1.3"
\#Gastropoda
ellipse_Gastropoda <- "1.7"
#Mammalia
ellipse_Mammalia <- "1.8"
#Reptilia
ellipse_Reptilia <- "1.9"
####fish - coral
AAn_95.overlap <- bayesianOverlap(ellipse_Actinopterygii,
                                   ellipse_Anthozoa,
                                   ellipses.posterior,
                                   draws = 100,
                                   p.interval = 0.95,
                                   n = 100
AAn_95_overlap_prop <- vector()
for(i in 1:length(AAn_95.overlap$overlap)){
AAn_95_overlap_prop[i] <- AAn_95.overlap$overlap[i]/min(AAn_95.overlap[i,1:2])
}
hist(AAn_95_overlap_prop, xlab = "Proportion of overlap", main = "")
```

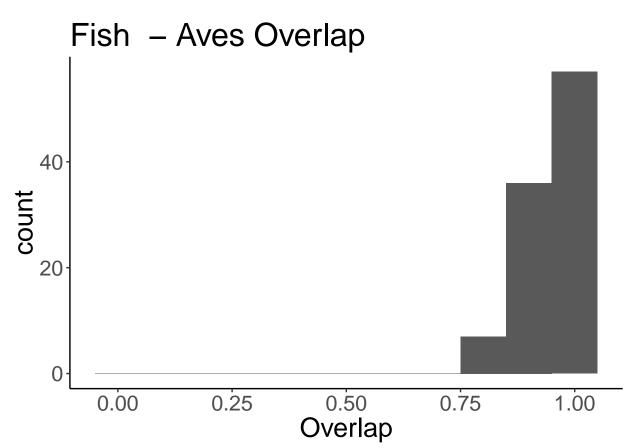


Fish – Coral Overlap 75250.00 0.25 0.50 0.75 1.00 Overlap



hdr(AAv_95_overlap_prop)

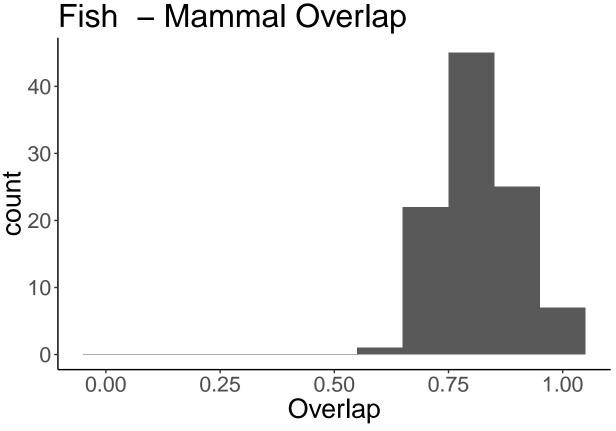
```
## $hdr
            [,1]
                      [,2]
                                 [,3]
                                          [,4]
##
## 99% 0.7754661 1.0334730
                                   NA
                                            NA
## 95% 0.8120950 0.8156312 0.8401105 1.029851
## 50% 0.9656204 1.0142572
                                  NA
                                            NA
##
## $mode
## [1] 0.9962782
##
## $falpha
                              50%
##
          1%
                    5%
## 0.6094313 1.0703473 5.9849876
myplot_Aav = ggplot(data.frame(Overlap = AAv_95_overlap_prop),
                    aes(Overlap)) +
                    geom_histogram(binwidth=0.1) +
                    ggtitle("Fish - Aves Overlap") +
                    scale_x_continuous(limits = c(-0.05, 1.05))
myplot_Aav + theme_bw() + theme(panel.border = element_blank(),
                                 panel.grid.major = element_blank(),
                                panel.grid.minor = element_blank(),
                                 axis.line = element_line(colour = "black"),
                                 text = element_text(size=20))
```



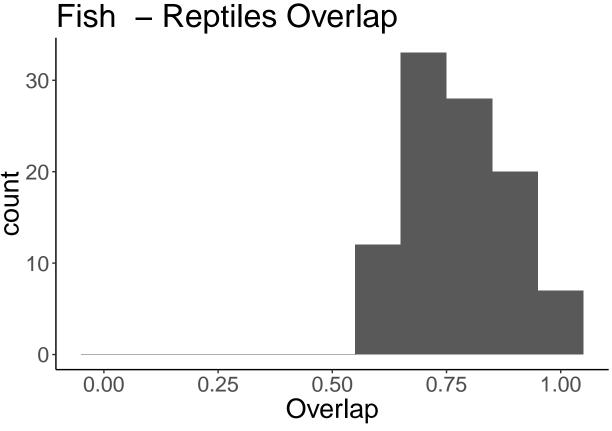
```
####fish - gastropod
AG95.overlap <- bayesianOverlap(ellipse_Actinopterygii,
                                   ellipse_Gastropoda,
                                   ellipses.posterior,
                                   draws = 100,
                                   p.interval = 0.95,
                                   n = 100)
AG_95_overlap_prop <- vector()
for(i in 1:length(AG95.overlap$)){
AG_95_overlap_prop[i] <- AG95.overlap$overlap[i]/min(AG95.overlap[i,1:2])
}
myplot_AG = ggplot(data.frame(Overlap = AG_95_overlap_prop),
                    aes(Overlap)) +
                    ggtitle("Fish - Gastropod Overlap") +
                    geom_histogram(binwidth=0.1) +
                    scale_x_continuous(limits = c(-0.05, 1.05))
myplot_AG + theme_bw() + theme(panel.border = element_blank(),
                               panel.grid.major = element_blank(),
                               panel.grid.minor = element_blank(),
                               axis.line = element line(colour = "black"),
                               text = element_text(size=20))
```

Fish – Gastropod Overlap 201500.00 0.25 0.50 0.75 1.00 Overlap

```
##fish - mammal
AM95.overlap <- bayesianOverlap(ellipse_Actinopterygii,
                                   ellipse_Mammalia,
                                   ellipses.posterior,
                                   draws = 100,
                                   p.interval = 0.95,
                                   n = 100)
Am_95_overlap_prop <- vector()</pre>
for(i in 1:length(AM95.overlap$)){
Am_95_overlap_prop[i] <- AM95.overlap$overlap[i]/min(AM95.overlap[i,1:2])
}
myplot_AM = ggplot(data.frame(Overlap = Am_95_overlap_prop),
                    aes(Overlap)) +
                    ggtitle("Fish - Mammal Overlap") +
                    geom_histogram(binwidth=0.1) +
                    scale_x_continuous(limits = c(-0.05, 1.05))
myplot_AM + theme_bw() + theme(panel.border = element_blank(),
                               panel.grid.major = element_blank(),
                               panel.grid.minor = element_blank(),
                               axis.line = element_line(colour = "black"),
                               text = element text(size=20))
```

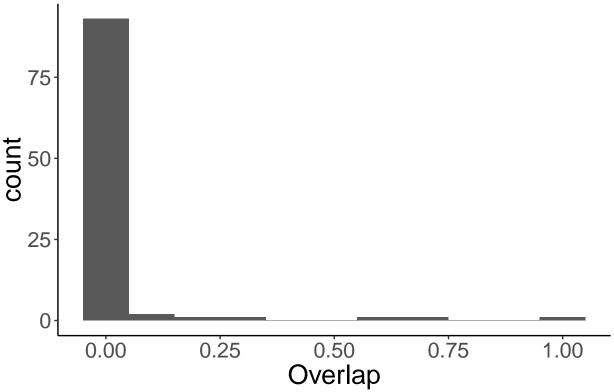


```
###fish and reptiles
AR95.overlap <- bayesianOverlap(ellipse_Actinopterygii,
                                   ellipse_Reptilia,
                                   ellipses.posterior,
                                   draws = 100,
                                   p.interval = 0.95,
                                   n = 100)
AR_95_overlap_prop <- vector()
for(i in 1:length(AR95.overlap$)){
AR_95_overlap_prop[i] <- AR95.overlap$overlap[i]/min(AR95.overlap[i,1:2])
}
myplot_AR = ggplot(data.frame(Overlap = AR_95_overlap_prop),
                    aes(Overlap)) +
                    ggtitle("Fish - Reptiles Overlap") +
                    geom_histogram(binwidth=0.1) +
                    scale_x_continuous(limits = c(-0.05, 1.05))
myplot_AR + theme_bw() + theme(panel.border = element_blank(),
                               panel.grid.major = element_blank(),
                               panel.grid.minor = element_blank(),
                               axis.line = element_line(colour = "black"),
                               text = element text(size=20))
```



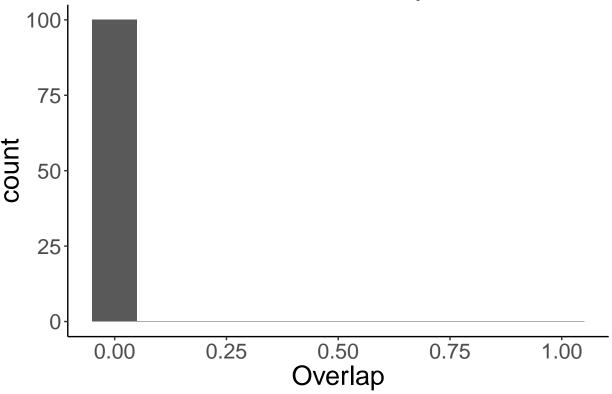
```
###coral-gastropod
AnG95.overlap <- bayesianOverlap(ellipse_Anthozoa,</pre>
                                   ellipse_Gastropoda,
                                   ellipses.posterior,
                                   draws = 100,
                                   p.interval = 0.95,
                                   n = 100)
AnG95_overlap_prop <- vector()</pre>
for(i in 1:length(AnG95.overlap$)){
AnG95_overlap_prop[i] <- AnG95.overlap[i]/min(AnG95.overlap[i,1:2])
}
myplot_AnG = ggplot(data.frame(Overlap = AnG95_overlap_prop),
                    aes(Overlap)) +
                    ggtitle("Coral - Gastropods Overlap") +
                    geom_histogram(binwidth=0.1) +
                    scale_x_continuous(limits = c(-0.05, 1.05))
myplot_AnG + theme_bw() + theme(panel.border = element_blank(),
                                panel.grid.major = element_blank(),
                                panel.grid.minor = element_blank(),
                                axis.line = element_line(colour = "black"),
                                text = element_text(size=20))
```

Coral - Gastropods Overlap

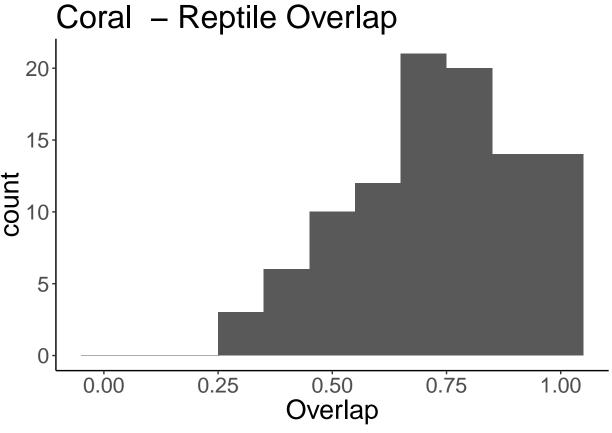


```
##coral mammal
AnM95.overlap <- bayesianOverlap(ellipse_Anthozoa,</pre>
                                    ellipse_Mammalia,
                                    ellipses.posterior,
                                    draws = 100,
                                    p.interval = 0.95,
                                    n = 100)
AnM95_overlap_prop <- vector()</pre>
for(i in 1:length(AnG95.overlap$overlap)){
AnM95_overlap_prop[i] <- AnM95.overlap$overlap[i]/min(AnM95.overlap[i,1:2])
}
myplot_AnM = ggplot(data.frame(Overlap = AnM95_overlap_prop),
                    aes(Overlap)) +
                    ggtitle("Coral - Mammal Overlap") +
                    geom_histogram(binwidth=0.1) +
                    scale_x_continuous(limits = c(-0.05, 1.05))
myplot_AnM + theme_bw() + theme(panel.border = element_blank(),
                                 panel.grid.major = element_blank(),
                                 panel.grid.minor = element_blank(),
                                 axis.line = element_line(colour = "black"),
                                 text = element_text(size=20))
```

Coral - Mammal Overlap



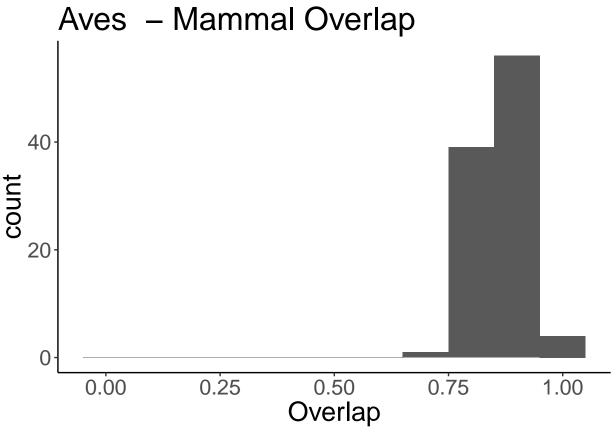
```
##coral reptile
AnR95.overlap <- bayesianOverlap(ellipse_Anthozoa,</pre>
                                   ellipse_Reptilia,
                                   ellipses.posterior,
                                   draws = 100,
                                   p.interval = 0.95,
                                   n = 100)
AnR95_overlap_prop <- vector()</pre>
for(i in 1:length(AnR95.overlap$)){
AnR95_overlap_prop[i] <- AnR95.overlap$overlap[i]/min(AnR95.overlap[i,1:2])
}
myplot_AnR = ggplot(data.frame(Overlap = AnR95_overlap_prop),
                    aes(Overlap)) +
                    ggtitle("Coral - Reptile Overlap") +
                    geom_histogram(binwidth=0.1) +
                    scale_x_continuous(limits = c(-0.05, 1.05))
myplot_AnR + theme_bw() + theme(panel.border = element_blank(),
                                panel.grid.major = element_blank(),
                                panel.grid.minor = element_blank(),
                                axis.line = element_line(colour = "black"),
                                text = element_text(size=20))
```



```
###bird - gastropd
AvG95.overlap <- bayesianOverlap(ellipse_Aves,</pre>
                                    ellipse_Gastropoda,
                                    ellipses.posterior,
                                    draws = 100,
                                    p.interval = 0.95,
                                    n = 100)
AvG95_overlap_prop <- vector()</pre>
for(i in 1:length(AvG95.overlap$overlap)){
AvG95_overlap_prop[i] <- AvG95.overlap$overlap[i]/min(AvG95.overlap[i,1:2])</pre>
}
myplot_AvG = ggplot(data.frame(Overlap = AvG95_overlap_prop),
                    aes(Overlap)) +
                    ggtitle("Aves - Gastropod Overlap") +
                    geom_histogram(binwidth=0.1) +
                    scale_x_continuous(limits = c(-0.05, 1.05))
myplot_AvG + theme_bw() + theme(panel.border = element_blank(),
                                 panel.grid.major = element_blank(),
                                 panel.grid.minor = element_blank(),
                                 axis.line = element_line(colour = "black"),
                                 text = element_text(size=20))
```

Aves – Gastropod Overlap 25201500.00 0.25 0.50 0.75 1.00 Overlap

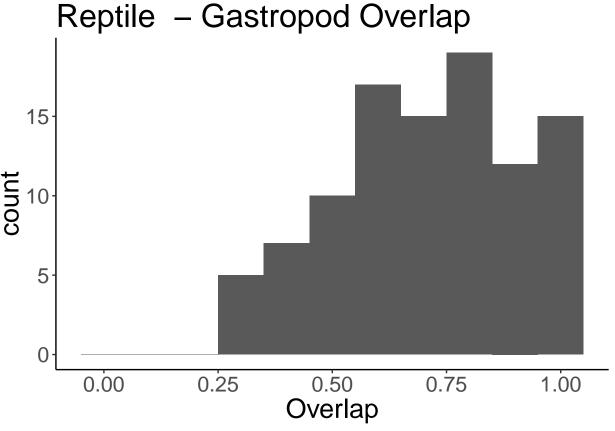
```
####aves mammal
AvM95.overlap <- bayesianOverlap(ellipse_Aves,</pre>
                                   ellipse_Mammalia,
                                   ellipses.posterior,
                                   draws = 100,
                                   p.interval = 0.95,
                                   n = 100)
AvM95_overlap_prop <- vector()</pre>
for(i in 1:length(AvM95.overlap$)){
AvM95_overlap_prop[i] <- AvM95.overlap[i]/min(AvM95.overlap[i,1:2])
}
myplot_AvM = ggplot(data.frame(Overlap = AvM95_overlap_prop),
                    aes(Overlap)) +
                    ggtitle("Aves - Mammal Overlap") +
                    geom_histogram(binwidth=0.1) +
                    scale_x_continuous(limits = c(-0.05, 1.05))
myplot_AvM + theme_bw() + theme(panel.border = element_blank(),
                                panel.grid.major = element_blank(),
                                panel.grid.minor = element_blank(),
                                axis.line = element_line(colour = "black"),
                                text = element_text(size=20))
```



```
##### Aves reptile
AvRR95.overlap <- bayesianOverlap(ellipse_Aves,</pre>
                                    ellipse_Reptilia,
                                    ellipses.posterior,
                                    draws = 100,
                                    p.interval = 0.95,
                                    n = 100)
AvR95_overlap_prop <- vector()</pre>
for(i in 1:length(AvRR95.overlap$)){
AvR95_overlap_prop[i] <- AvRR95.overlap$overlap[i]/min(AvRR95.overlap[i,1:2])</pre>
}
myplot_AvR = ggplot(data.frame(Overlap = AvR95_overlap_prop),
                    aes(Overlap)) +
                    ggtitle("Aves - Reptile Overlap") +
                    geom_histogram(binwidth=0.1) +
                    scale_x_continuous(limits = c(-0.05, 1.05))
myplot_AvR + theme_bw() + theme(panel.border = element_blank(),
                                 panel.grid.major = element_blank(),
                                 panel.grid.minor = element_blank(),
                                 axis.line = element_line(colour = "black"),
                                 text = element_text(size=20))
```

Aves – Reptile Overlap 40 30 10 0.00 0.25 0.50 0.75 1.00 Overlap

```
##### reptile gastropod
RG95.overlap <- bayesianOverlap(ellipse_Reptilia,
                                    ellipse_Gastropoda,
                                    ellipses.posterior,
                                    draws = 100,
                                    p.interval = 0.95,
                                    n = 100)
RG95_overlap_prop <- vector()
for(i in 1:length(RG95.overlap$overlap)){
RG95_overlap_prop[i] <- RG95.overlap$overlap[i]/min(RG95.overlap[i,1:2])</pre>
}
myplot_RG = ggplot(data.frame(Overlap = RG95_overlap_prop),
                    aes(Overlap)) +
                    ggtitle("Reptile - Gastropod Overlap") +
                    geom_histogram(binwidth=0.1) +
                    scale_x_continuous(limits = c(-0.05, 1.05))
myplot_RG + theme_bw() + theme(panel.border = element_blank(),
                               panel.grid.major = element_blank(),
                               panel.grid.minor = element_blank(),
                               axis.line = element line(colour = "black"),
                                text = element_text(size=20))
```



```
##### Aves coral
AvAn95.overlap <- bayesianOverlap(ellipse_Aves,</pre>
                                    ellipse_Anthozoa,
                                    ellipses.posterior,
                                    draws = 100,
                                    p.interval = 0.95,
                                    n = 100)
AvAn95_overlap_prop <- vector()</pre>
for(i in 1:length(AvAn95.overlap$overlap)){
AvAn95_overlap_prop[i] <- AvAn95.overlap$overlap[i]/min(AvAn95.overlap[i,1:2])</pre>
}
myplot_AvAn = ggplot(data.frame(Overlap = AvAn95_overlap_prop),
                    aes(Overlap)) +
                    ggtitle("Aves - Coral Overlap") +
                    geom_histogram(binwidth=0.1) +
                    scale_x_continuous(limits = c(-0.05, 1.05))
myplot_AvAn + theme_bw() + theme(panel.border = element_blank(),
                                  panel.grid.major = element_blank(),
                                  panel.grid.minor = element_blank(),
                                  axis.line = element_line(colour = "black"),
                                  text = element_text(size=20))
```

Aves – Coral Overlap Toology Toology

0.50

Overlap

0.75

1.00

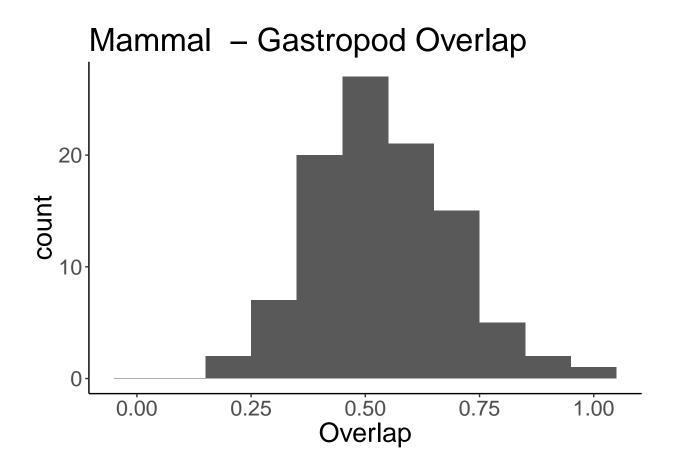
0.25

0.00

```
##### reptile mammal
RM95.overlap <- bayesianOverlap(ellipse_Mammalia,
                                   ellipse_Reptilia,
                                   ellipses.posterior,
                                   draws = 100,
                                   p.interval = 0.95,
                                   n = 100)
RM95_overlap_prop <- vector()
for(i in 1:length(RG95.overlap$)){
RM95_overlap_prop[i] <- RM95.overlap$overlap[i]/min(RM95.overlap[i,1:2])</pre>
}
myplot_RM = ggplot(data.frame(Overlap = RM95_overlap_prop),
                    aes(Overlap)) +
                    ggtitle("Reptile - Mammal Overlap") +
                    geom_histogram(binwidth=0.1) +
                    scale_x_continuous(limits = c(-0.05, 1.05))
myplot_RM + theme_bw() + theme(panel.border = element_blank(),
                               panel.grid.major = element_blank(),
                               panel.grid.minor = element_blank(),
                               axis.line = element_line(colour = "black"),
                               text = element_text(size=20))
```

Reptile – Mammal Overlap 30 10 0.00 0.25 0.50 0.75 1.00

```
##### mammal gastropod
MG95.overlap <- bayesianOverlap(ellipse_Mammalia,
                                   ellipse_Gastropoda,
                                   ellipses.posterior,
                                   draws = 100,
                                   p.interval = 0.95,
                                   n = 100)
MG95_overlap_prop <- vector()
for(i in 1:length(RG95.overlap$overlap)){
MG95_overlap_prop[i] <- MG95.overlap$overlap[i]/min(MG95.overlap[i,1:2])
}
myplot_MG = ggplot(data.frame(Overlap = MG95_overlap_prop),
                    aes(Overlap)) +
                    ggtitle("Mammal - Gastropod Overlap") +
                    geom_histogram(binwidth=0.1) +
                    scale_x_continuous(limits = c(-0.05, 1.05))
myplot_MG + theme_bw() + theme(panel.border = element_blank(),
                               panel.grid.major = element_blank(),
                               panel.grid.minor = element_blank(),
                               axis.line = element_line(colour = "black"),
                               text = element_text(size=20))
```



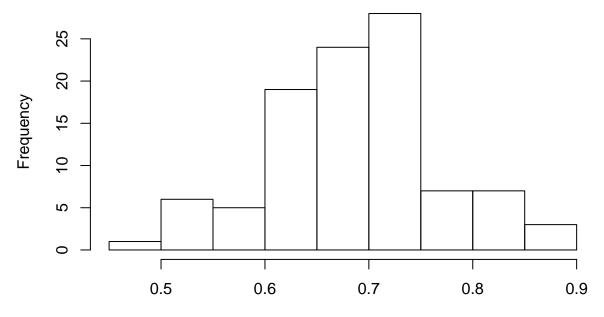
IUCN overlap calculations for mode of life

```
group.ML <- groupMetricsML(siber.plots)</pre>
group.ML_iucn <- groupMetricsML(siber.iucn)</pre>
# options for running jags
parms <- list()</pre>
parms$n.iter <- 2 * 10^4 # number of iterations to run the model for
parms$n.burnin <- 1 * 10^3 # discard the first set of values
                       # thin the posterior by this many
parms$n.thin <- 10
parms$n.chains <- 2</pre>
                             # run this many chains
# define the priors
priors <- list()</pre>
priors$R <- 1 * diag(2)</pre>
priors$k <- 2</pre>
priors$tau.mu <- 1.0E-3</pre>
ellipses.posterior_iucn <- siberMVN(siber.iucn, parms, priors)</pre>
```

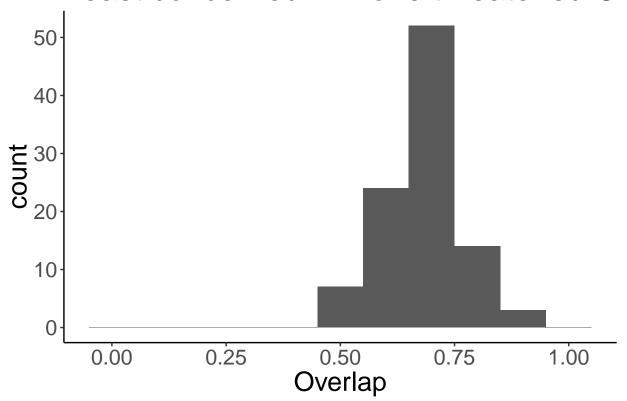
Compiling model graph

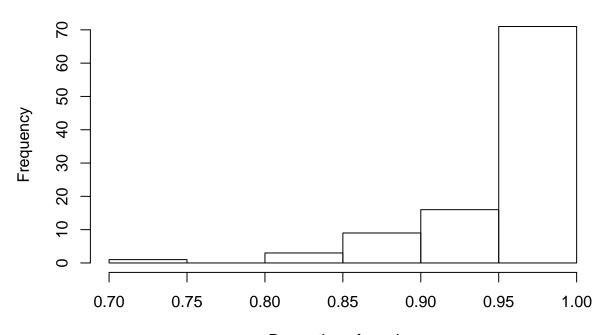
```
##
      Resolving undeclared variables
##
      Allocating nodes
##
  Graph information:
      Observed stochastic nodes: 31
##
##
      Unobserved stochastic nodes: 3
      Total graph size: 46
##
## Initializing model
##
##
   Compiling model graph
      Resolving undeclared variables
##
      Allocating nodes
##
  Graph information:
      Observed stochastic nodes: 191
##
##
      Unobserved stochastic nodes: 3
##
      Total graph size: 206
##
## Initializing model
##
## Compiling model graph
##
      Resolving undeclared variables
##
      Allocating nodes
## Graph information:
##
      Observed stochastic nodes: 10
##
      Unobserved stochastic nodes: 3
##
      Total graph size: 25
##
## Initializing model
##
  Compiling model graph
##
      Resolving undeclared variables
##
      Allocating nodes
##
   Graph information:
##
      Observed stochastic nodes: 21
##
      Unobserved stochastic nodes: 3
##
      Total graph size: 36
##
## Initializing model
##
  Compiling model graph
##
      Resolving undeclared variables
##
      Allocating nodes
## Graph information:
      Observed stochastic nodes: 14
##
##
      Unobserved stochastic nodes: 3
##
      Total graph size: 29
##
## Initializing model
##
##
  Compiling model graph
##
      Resolving undeclared variables
##
      Allocating nodes
## Graph information:
##
      Observed stochastic nodes: 16
```

```
##
      Unobserved stochastic nodes: 3
##
      Total graph size: 31
##
## Initializing model
##
## Compiling model graph
      Resolving undeclared variables
      Allocating nodes
##
## Graph information:
##
      Observed stochastic nodes: 2
##
      Unobserved stochastic nodes: 3
      Total graph size: 17
##
##
## Initializing model
# The first ellipse is referenced using a character string representation where
# in "x.y", "x" is the community, and "y" is the group within that community.
# So in this example: community 1, group 1
#ellipse group numbers
ellipse_NA <- "1.1"
ellipse_CE <- "1.2"
ellipse_E <- "1.3"
ellipse_LC <- "1.4"
ellipse_LR <- "1.5"
ellipse NT <- "1.6"
ellipse_V <- "1.7"
#####LC - NT
LC_NT_95.overlap <- bayesianOverlap(ellipse_LC,</pre>
                                    ellipse_E,
                                    ellipses.posterior_iucn,
                                    draws = 100,
                                    p.interval = 0.95,
                                    n = 100)
LC_NT_95.overlap_prop <- vector()</pre>
for(i in 1:length(LC_NT_95.overlap$overlap)){
LC_NT_95.overlap_prop[i] <- LC_NT_95.overlap$overlap[i]/min(LC_NT_95.overlap[i,1:2])
}
hist(LC_NT_95.overlap_prop, xlab = "Proportion of overlap", main = "")
```

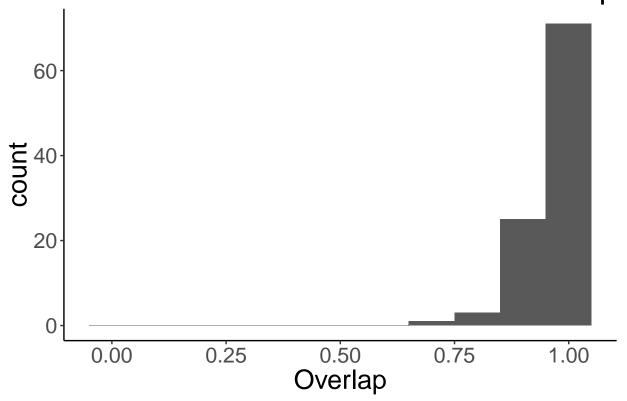


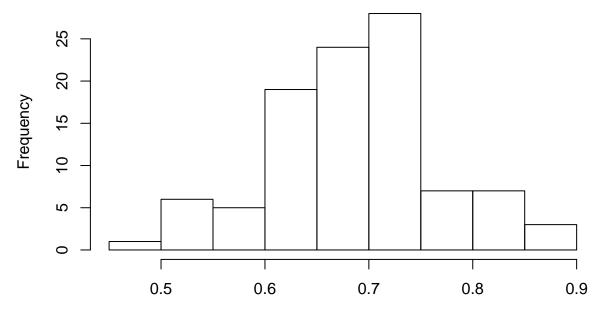
Least concerned - None threatened O



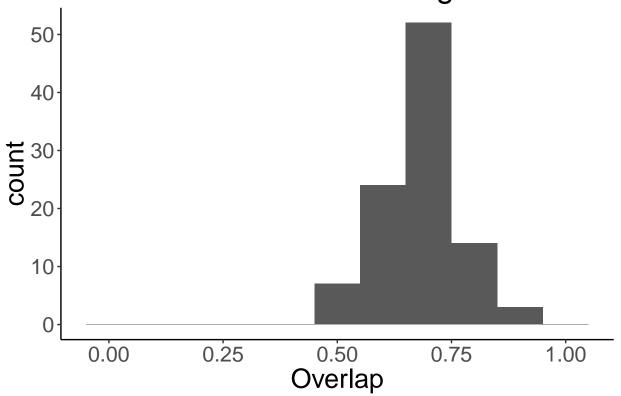


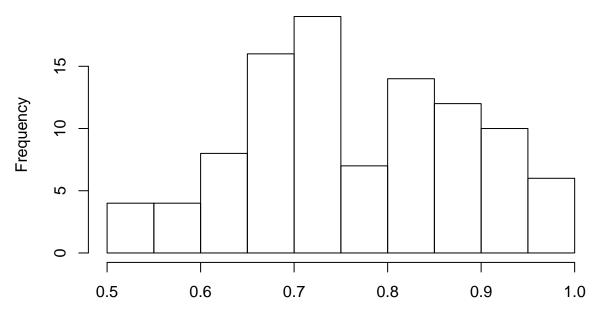
Least concerned - Vulnerable Overlap



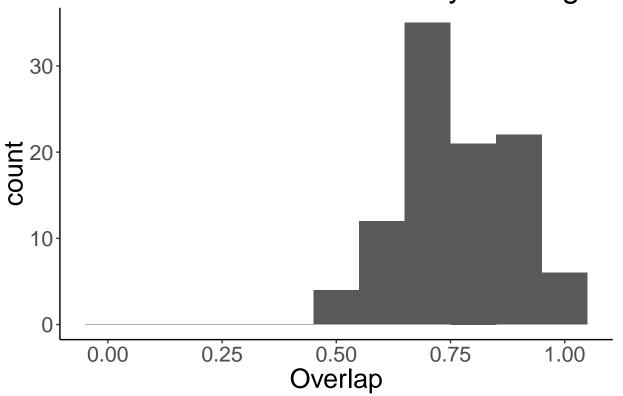


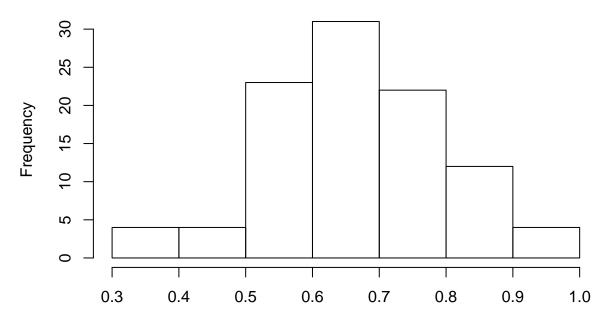
Least concerned - Endangered Overla



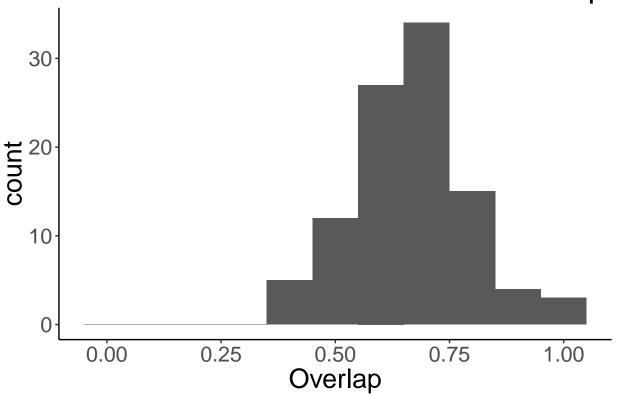


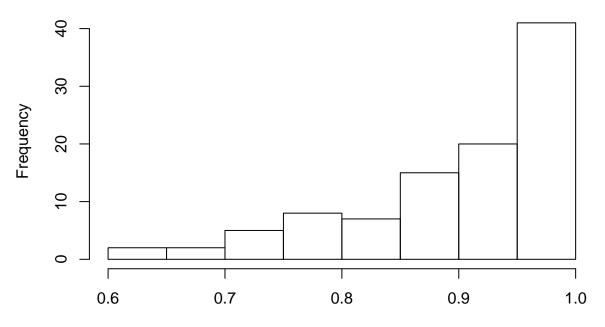
Least concerned - Critically endangere



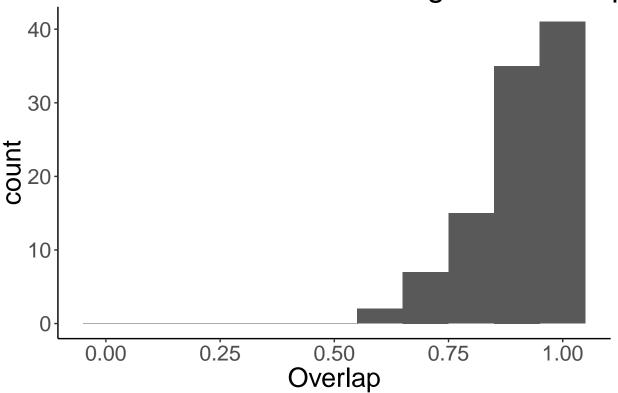


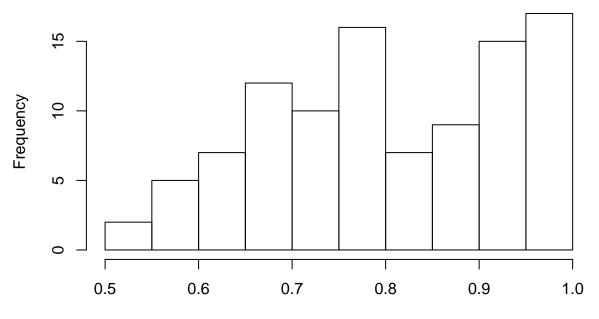
Near Theatened - Vulnerable Overlap



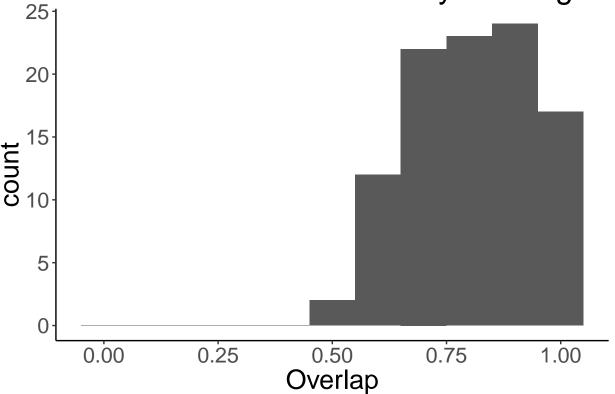


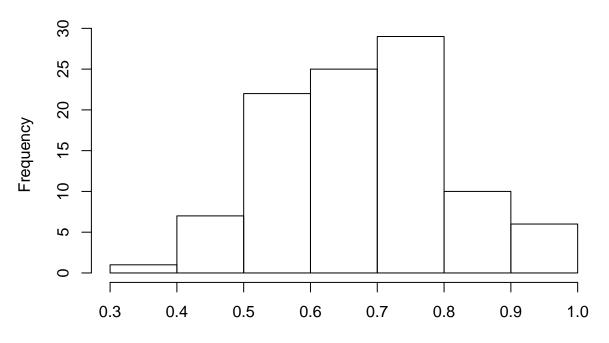
Near Theatened - Endangered Overlag



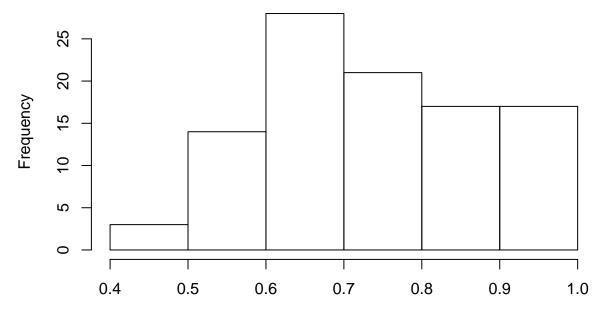


Near Theatened - Critically Endangere

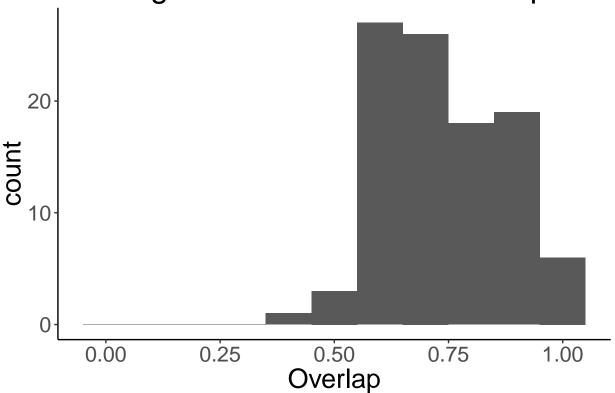


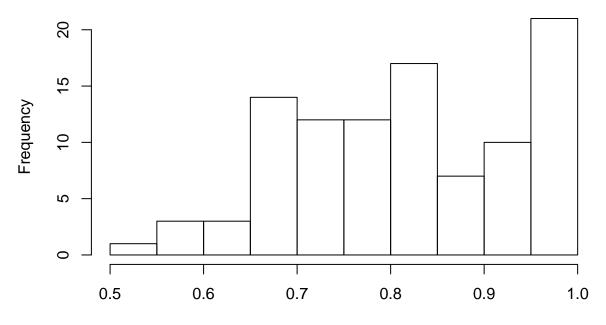


Endangered – Critically Endangered O

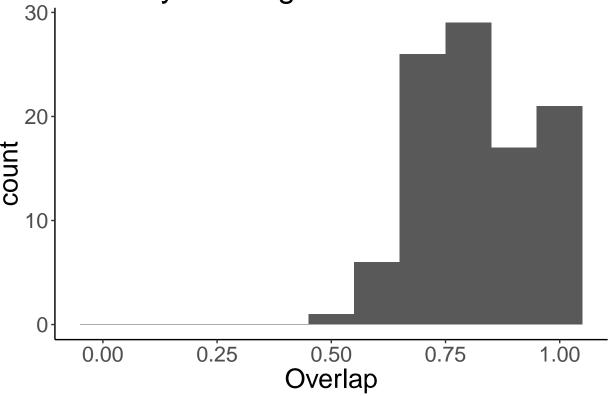


Endangered – Vulnerable Overlap





Critically Endangered – Vulnerable Ove



Juvinal Survival versus SD of mortality

##

```
Iterations = 10001:109951
    Thinning interval = 50
##
    Sample size = 2000
##
##
    DIC: -386.2264
##
##
    G-structure: ~animal
##
          post.mean 1-95% CI u-95% CI eff.samp
##
            0.08128 0.03129 0.1342
##
   animal
##
##
                   ~species
##
           post.mean 1-95% CI u-95% CI eff.samp
##
           0.01015 0.003956 0.01644
  species
                                            1753
##
##
    R-structure: ~units
##
         post.mean 1-95% CI u-95% CI eff.samp
##
          0.01115 0.008687 0.01352
## units
##
##
   Location effects: prop_la ~ surv_sd
##
               post.mean 1-95% CI u-95% CI eff.samp pMCMC
##
## (Intercept)
                  0.4605
                           0.1300 0.7807
                                                2000 0.005 **
## surv_sd
                 -4.6707 -6.2276 -2.9508
                                                2000 <5e-04 ***
## ---
## Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
j_surv_vcv_phylo <- j_surv$VCV[,1]</pre>
j_surv_vcv_spec <- j_surv$VCV[,2]</pre>
j_surv_vcv_units <- j_surv$VCV[,3]</pre>
j_surv_phylo <- j_surv_vcv_phylo/c(j_surv_vcv_phylo + j_surv_vcv_spec + j_surv_vcv_units)</pre>
j_surv_spec <- j_surv_vcv_spec/c(j_surv_vcv_phylo + j_surv_vcv_spec + j_surv_vcv_units)</pre>
j_surv_units <- j_surv_vcv_units/c(j_surv_vcv_phylo + j_surv_vcv_spec + j_surv_vcv_units)</pre>
hdr(j_surv_phylo)$mode
## [1] 0.7940034
hdr(j_surv_spec)$mode
## [1] 0.07884778
hdr(j_surv_units)$mode
```

[1] 0.107091