# motmot

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Models Of Trait Macroevolution On Trees (MOTMOT) is an R package that allows for testing of models of trait evolution (Thomas and Freckleton 2012).

- Tree transformation models estimated using Maximum likelihood: Brownian motion, Pagel's Lambda, Delta, Kappa, Ornstein-Uhlenbeck (OU), Acceleration-Deaceleration (ACDC), and estimating lambda alongside other models
- Rate heterogeneous models of evolution. Fit models in which the rate of evolution differs in clades selected a priori (Thomas, Freckleton, and Székely 2006, O'Meara et al. (2006)), and models with no a-priori shift locations (Thomas and Freckleton 2012)
- TimeSlice fit models in which all rates change at a specific time(s) by tested all times or those selected by the user
- Nested Shift mode Fit models models in which the ancestral BM rate switches to a 'nested' rate within a monophyletic clade in the phylogeny
- Bayesian estimation of tree transformation models

### Introduction

## Node labels:

## ## 2, 2, 2, 2, 2, ...

## Rooted; includes branch lengths.

For these examples we will use anolis data available from motmot. A time-calibrated phylogeny of anolis species ("anolis.tree"), and various trait and biogeographical trait data ("anolis.data")

```
library(motmot.2.0, quietly=T)
data(anolis.tree)
data(anolis.data)
names(anolis.data)
## [1] "Species"
                                      "ecomorph"
                      "Island_type"
                                                     "geo_ecomorph"
## [5] "Female_SVL"
                      "Male_SVL"
attach(anolis.data)
anolis.tree
##
## Phylogenetic tree with 165 tips and 164 internal nodes.
##
## Tip labels:
```

For the first part of the tutorial we will use the continuous trait data: male snout-ventral length 'Male\_SVL'.

A\_occultus, A\_darlingt, A\_monticol, A\_bahoruco, A\_dolichoc, A\_henderso, ...

We will construct a matrix of just these data, and check if we have missing data

```
male.length <- matrix(Male_SVL, dimnames=list(rownames(anolis.data)))
any(is.na(male.length[,1]))</pre>
```

```
## [1] TRUE
```

We do. So we will remove these data from the male.length data, and log the trait data

```
complete.male.length <- complete.cases(male.length)
missing.species <- rownames(male.length)[!complete.male.length]
male.length <- as.matrix(male.length[complete.male.length, ])
male.length <- log(male.length)</pre>
```

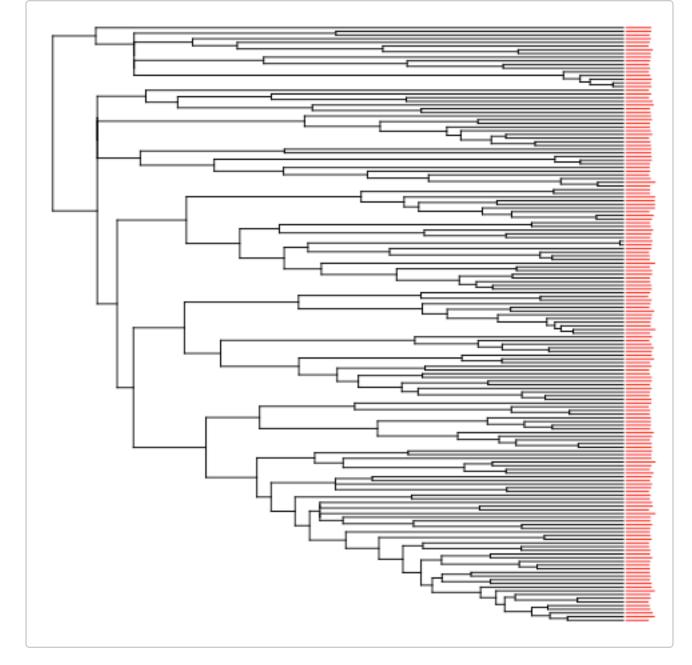
Finally, we will 'prune' the species from the tree using 'drop.tip' from APE. Do our species from the data and tree now match?

```
phy <- drop.tip(anolis.tree, missing.species)
name.check(phy, male.length)

## [1] "OK"</pre>
```

They do. We can now plot our tree and data using the "traitData.plot" function

```
traitData.plot(y=male.length, phy)
```



traitData showing the realtive male snout ventral length at the tips

# Models of trait evolution

We can now test various models of evolution using our trait data.

#### **Brownian motion**

To start we will fit a simple Brownian motion model to the data

```
bm.ml <- transformPhylo.ML(phy, y=male.length, model="bm")
bm.ml</pre>
```

```
## $brownianVariance
## [,1]
## [1,] 0.001917681
##
## $logLikelihood
## [,1]
## [1,] -17.35662
##
## $root.state
## [1] 4.120935
##
```

```
## $AIC

## [,1]

## [1,] 38.71324

##

## $AICc

## [,1]

## [1,] 38.78872
```

#### Pagel's lambda

We can also fit models to test Pagel's lambda

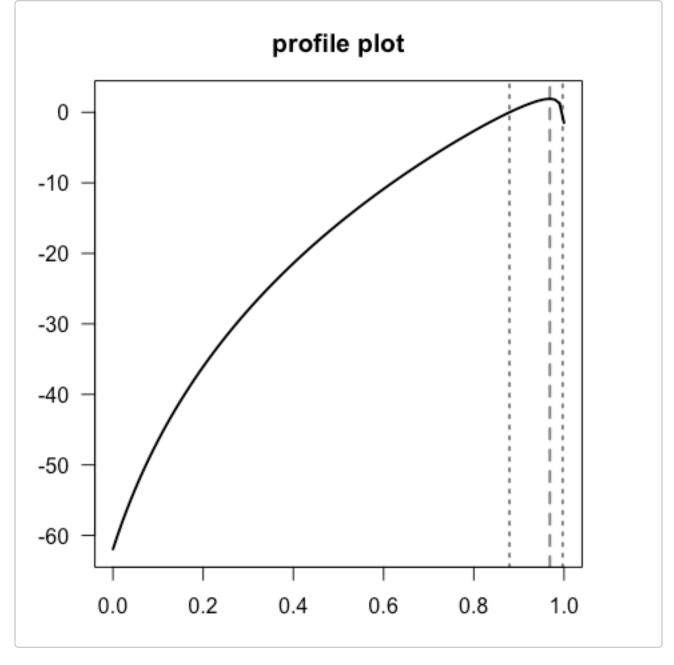
```
lambda.ml <- transformPhylo.ML(phy, y=male.length, model="lambda")
lambda.ml</pre>
```

```
## $MaximumLikelihood
## [1] -13.95609
##
## $Lambda
##
         MLLambda LowerCI
                              UpperCI
## [1,] 0.9685786 0.879023 0.9970324
##
## $brownianVariance
##
               [,1]
## [1,] 0.001675265
##
## $root.state
## [1] 4.122186
##
## $AIC
## [1] 33.91218
##
## $AICc
## [1] 34.06408
```

We can see from these results the CI of lambda is < 1, but there is still a large phylogenetic signal is these data

A new feature in motmot allows for plotting of the likelihood profile for the branch-transformation parameter, in this case Pagel's lambda

```
lambda.ml <- transformPhylo.ML(phy, y=male.length, model="lambda", profilePlot=T)
```



profile plot of ML estimation for Pagel's lambda

We can now compare the fit of the BM and Lambda models. Lambda has higher likelihood, but it also has more parameters. We can test whether this is a significant improvement. First we will use the chi-squared distribution. The models differ in one degree of freedom: BM has 2 parameters (brownian variance, root state) and lambda has those two plus the value of lambda. We can use the R function pchisq to obtain a p value, and see that lambda is not a superior fit to these male length data

```
p.value <- 1 - pchisq(lambda.ml$MaximumLikelihood - bm.ml$logLikelihood, 1)
p.value</pre>
```

```
## [,1]
## [1,] 0.06517542
```

However, there is a large Akaike Information Criterion (AICc) difference between the two models: BM has a higher AICc compared to Lambda. The differce (4.724636) is >4 which is tradtionally seen as indication of a superior fit (Burnham and Anderson 2003).

```
bm.ml$AICc- lambda.ml$AICc

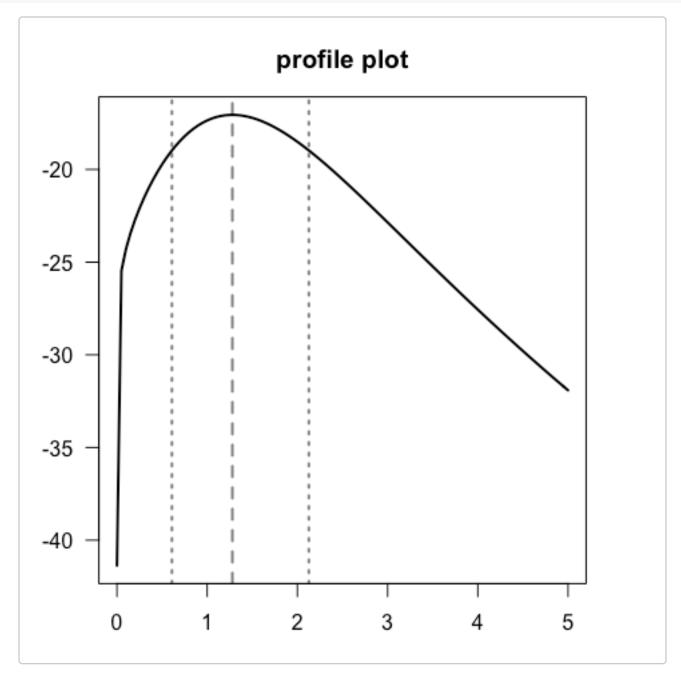
## [,1]
## [1,] 4.724636
```

The parameters, brownian variance, root state, Maximum likelihoods, AIC, and AICc can be obtained for a number of models in motmot.

#### **Delta**

Delta indicates a slow or increase in the rate of trait evolution through time; a value of 1 is equivalent to Brownian motion, < 1 indicates a slow-down, and > 1 is difficult to interpret (greater change near the present). Here we find a MLE of 1.27 but the CI spans < 1

```
delta.ml <- transformPhylo.ML(phy, y=male.length, model="delta", profilePlot=T)</pre>
```



profile plot to estimate delta

#### delta.ml

```
## $MaximumLikelihood
## [1] -17.05532
##
## $Delta
##
        MLDelta
                  LowerCI UpperCI
## [1,] 1.27924 0.6092799 2.127126
##
## $brownianVariance
##
             [,1]
## [1,] -17.05532
##
## $root.state
## [1] 4.127836
```

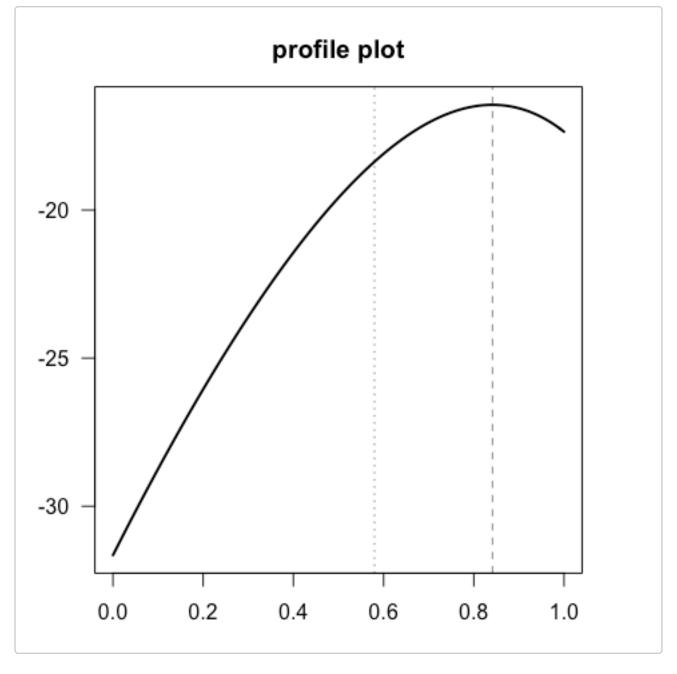
```
##
## $AIC
## [1] 40.11064
##
## $AICc
## [1] 40.26254
```

#### Kappa

Kappa is used as a measure of punctuated evolution and spans values of 0-1. 1 is equivalent to BM, and 0 indicates trait change occurs at events of speciation. Here the is some evidence of punctuated evolution, but the CI spans a value greater than one (as seen in the warning message)

```
kappa.ml <- transformPhylo.ML(phy, y=male.length, model="kappa", profilePlot=T)

## Warning in transformPhylo.ML(phy, y = male.length, model = "kappa",
## profilePlot = T): Confidence limits fall outside the parameter bounds -
## consider changing lowerBound and/or upperBound</pre>
```



profile plot to estimate kappa

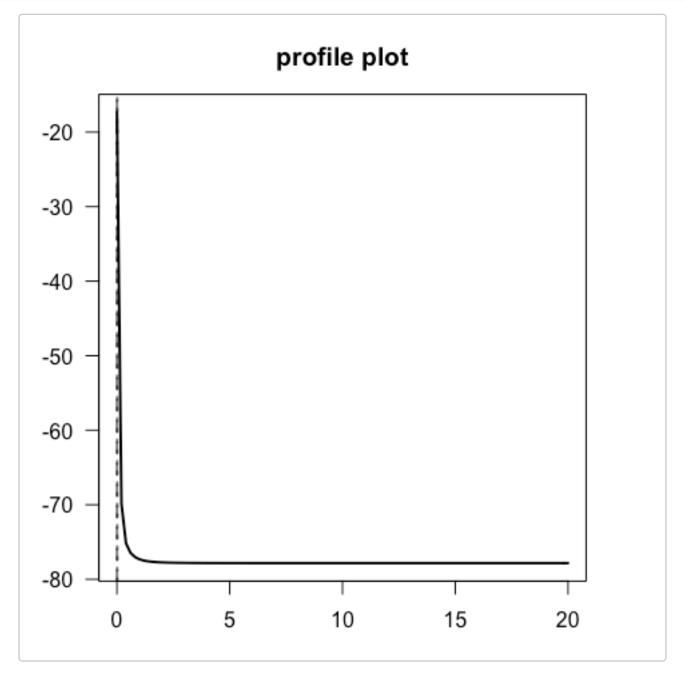
#### **Ornstein-Uhlenbeck**

The OU model allows for modelling of attraction to a optimum value (alpha), and psi fits a acceleration-deacceleration model to assess to the relative contributions of speciation and gradual evolution to a trait's evolutionary rate

```
ou.ml <- transformPhylo.ML(phy, y=male.length, model="OU", profilePlot=T)

## [1] "Warning - different start values produces different OU likelihoods"

## Warning in transformPhylo.ML(phy, y = male.length, model = "OU",
## profilePlot = T): Confidence limits fall outside parameter bounds -
## consider changing lowerBound and/or upperBound</pre>
```



profile plot to estimate alpha

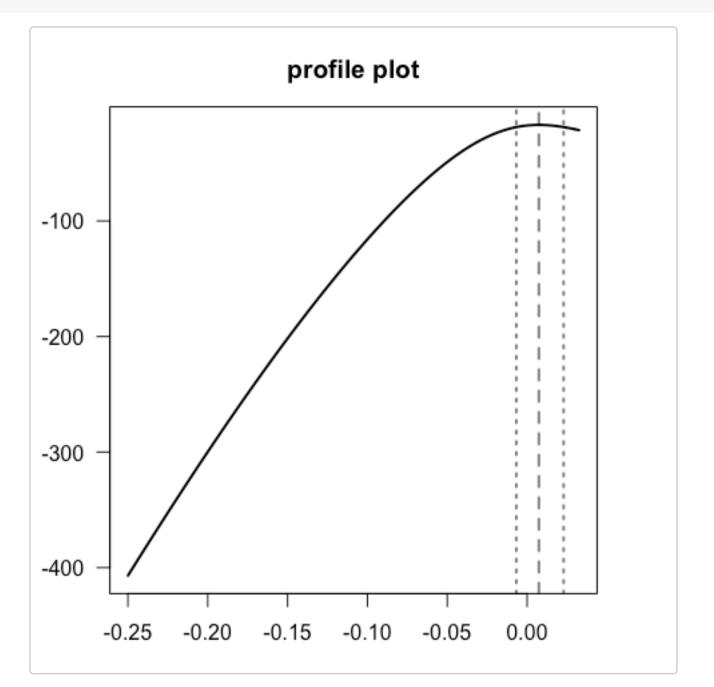
ou.ml

```
## $brownianVariance$brownianVariance
##
                [,1]
## [1,] 0.002206995
##
## $brownianVariance$logLikelihood
##
             [,1]
## [1,] -16.84778
##
##
## $root.state
## [1] 4.124566
##
## $AIC
## [1] 39.69557
##
## $AICc
## [1] 39.84747
```

#### **ACDC**

A new addition to MOTMOT is the ACDC model (Harmon et al. 2010; Blomberg, Jr, and Ives 2003). This model allows for exponential changes in the rate of evolution in the history of a clade. If the upperBound value is < 0, this is equivalent to the 'Early Burst' model fit in geiger

```
acdc.ml <- transformPhylo.ML(phy, y=male.length, model="ACDC", profilePlot=T)</pre>
```



```
acdc.ml
## $MaximumLikelihood
## [1] -16.84778
##
## $acdc
                          LowerCI
##
             MLacdc
                                      UpperCI
## [1,] 0.007384034 -0.006756934 0.02288467
##
## $brownianVariance
##
                [,1]
## [1,] 0.001117947
##
## $root.state
## [1] 4.124568
##
## $AIC
## [1] 39.69557
##
## $AICc
## [1] 39.84747
```

There is little evidence here of exponential decreases or increases in the rate of trait evolution - the acdc exponential parameter is close to 0 (0.007). We can see this is not a significant improvement on BM

```
p.value.2 <- 1 - pchisq(acdc.ml$MaximumLikelihood - bm.ml$logLikelihood , 1)
p.value.2

## [,1]
## [1,] 0.4756421</pre>
```

#### Estimate lambda alongside models

One way to deal with 'noisy' data is to estimate Pagel's lambda alongside a parameter of interest. In motmot, lambda can be estimated alongside the delta, kappa, OU, psi, and ACDC models. Here we look at example using ACDC. The model is fit with same function. 'transformPhyo.ML', but with the argument 'lambdaEst' set to TRUE

```
acdc.ml.lambda <- transformPhylo.ML(y=male.length, phy, model="ACDC", lambdaEst=T)
# original ACDC model
acdc.ml</pre>
```

```
##
## $brownianVariance
##
## [1,] 0.001117947
##
## $root.state
## [1] 4.124568
##
## $AIC
## [1] 39.69557
##
## $AICc
## [1] 39.84747
# ACDC model plus lambda
acdc.ml.lambda
## $MaximumLikelihood
## [1] -13.89171
##
## $acdc
##
              MLacdc
                         LowerCI
                                    UpperCI
## [1,] -0.003593448 -0.0176518 0.01111201
##
## $brownianVariance
##
## [1,] 0.002101792
##
## $root.state
## [1] 4.121082
##
## $Lambda
## [1] 0.9583652
##
```

We can see lambda is > 1, and this has affected the parameter estimation (slightly). However, the improvement in the model fit is not significant compared to the ACDC model without lambda, or the null BM model

## \$AIC

## \$AICc

##

## [1] 37.78342

## [1] 38.16804

```
# p value of the ACDC and ACDC+lambda models. No significant improvement
1 - pchisq(acdc.ml.lambda$MaximumLikelihood - acdc.ml$MaximumLikelihood , df=1)
## [1] 0.08555553
```

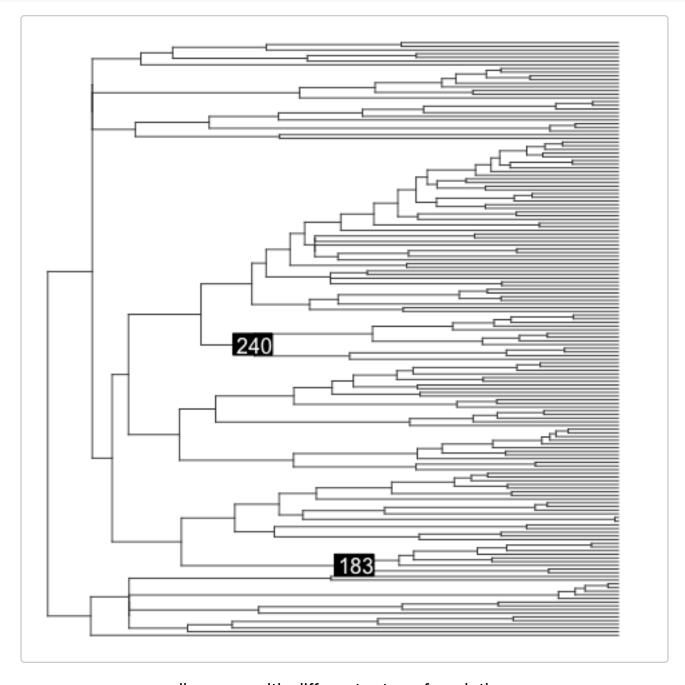
```
# p value of the BM and ACDC+lambda model comparison. No significant improvement
1 - pchisq(acdc.ml.lambda$MaximumLikelihood - bm.ml$logLikelihood, df=2)
```

# Rate heterogeneous models of evolution

#### rate heterogeneity selected a priori

MOTMOT can test models of evolution in which pre-defined clades can vary in the rate of evolution. Here we fit a model in which the nodes descending from nodes 183 and 240 have a seperate rate of evolution. We can visualise these nodes on the phylogeny

```
plot(phy, show.tip.label=F, no.margin=T, edge.col="grey20")
nodelabels(c(183, 240), c(183, 240), bg="black", col="white")
```



lineages with different rates of evolution

We then fit the motmot model, again using the function transformPhylo.ML. We use the argument "model=clade". This fits the non-censored model of O'Meara et al. (2006).

```
cladeRate.ml <- transformPhylo.ML(male.length, phy=phy, model="clade", nodeIDs=c(183, 240))</pre>
```

```
## $MaximumLikelihood
## [1] -17.1441
##
## $Rates
##
                MLRate LowerCI UpperCI
        node
## [1,] 183 0.7853043 0.3223851 2.47607
## [2,] 240 0.8119501 0.3927771 2.01348
##
## $brownianVariance
##
               [,1]
## [1,] 0.001969926
##
## $root.state
## [1] 4.123463
##
## $AIC
## [1] 42.28819
##
## $AICc
## [1] 42.54297
```

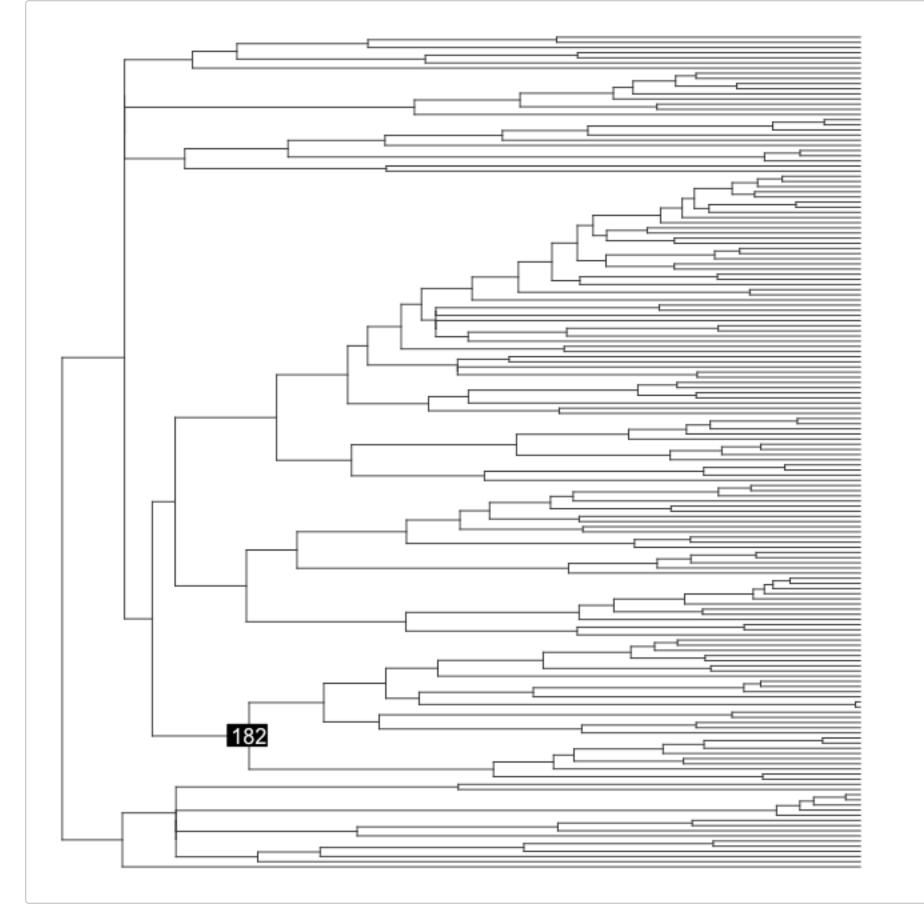
These results indicate that the two clades tend to have a lower rate of evolution compared to the background rate. However, the CIs indicate these decreases may not be robust

#### rate heterogeneity with no a priori information

We can also fit rate heterogeneous models without specifying where we expect shifts on the tree. We can use the arguments "model="tm1"" and "model="tm2""; these models fit 'traitMedusa' models in which all nodes are tested for rate increases or decreases. It is possible to exclude small nodes using the argument 'minCladeSize'. As well as allowing clade differences in rate, the "tm2" also allows for branch-based increases or decreases in rate.

We will fit the "tm2" model that allows for clade- and branch-specific changes in rate. This uses the familiar function 'transformPhylo.ML'. As the 'traitMedusa' is computationally-intensive (fitting models to all nodes, optimising increasing numbers of rates) we will fit the 'tm2' to a subset of the phylogeny and data using the APE function 'extract.clade'

```
plot(phy, show.tip.label=F, no.margin=T, edge.col="grey20")
nodelabels(182, 182, bg="black", col="white")
```



#### the subset of the tree

```
phy.clade <- extract.clade(phy, 182)
male.length.clade <- as.matrix(male.length[match(phy.clade$tip.label,
rownames(male.length)),])</pre>
```

We can now fit the 'tm2' algorithm. The output shows the log-likelihood, AIC, AICc, rate type (branch of clade), for the best-fitting model at each stage. This starts with the BM model, and then one shift model, two shift model, etc.,

```
# not run
# tm1.ml <- transformPhylo.ML(y=male.length.clade, phy=phy.clade, model="tm1",
minCladeSize=2, nSplits=3)
# trait.medusa.tm1.summary <- traitMedusaSummary(tm1.ml, cutoff=2, AICc=T)
tm2.ml <- transformPhylo.ML(y=male.length.clade, phy=phy.clade, model="tm2",
minCladeSize=2, nSplits=3)</pre>
```

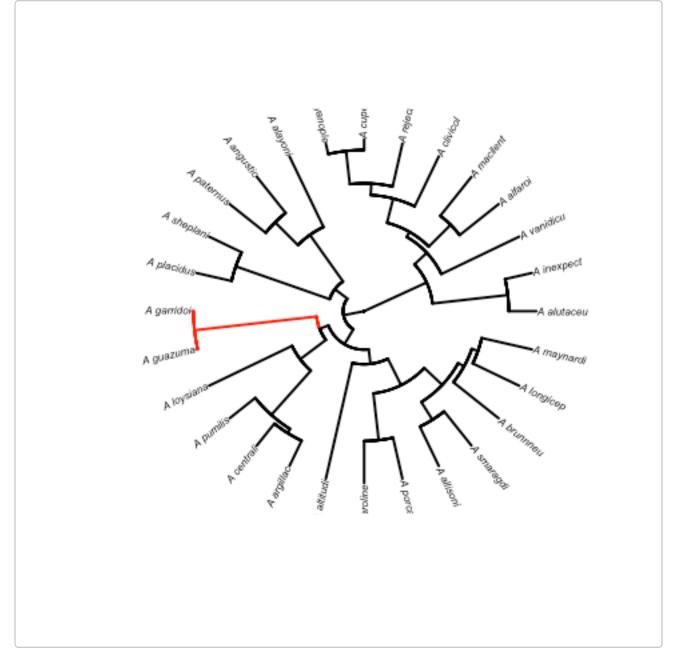
```
##
##
   BM model
         node shiftPos lnL n.params AIC
##
## BM
              1 -0.2838554 2 4.567711 5.047711
##
##
  Shift 1
         node shiftPos lnL n.params
                                        AIC
                                                 AICc rate.1
## shift.1
          45
               clade 3.6472 3 -1.294399 -0.2943992 8.603397
##
  Shift 2
##
         node shiftPos
                       lnL n.params
                                          AIC
                                                  AICc
                                                        rate.1
           39 clade 6.109191 5 -2.218382 0.5088912 7.278441
## shift.2
##
           rate.2
## shift.2 0.137804
##
## Shift 3
         node shiftPos lnL n.params AIC
                                               AICc rate.1
          31
                             7 -3.257608 2.342392 6.68661
## shift.3
                clade 8.628804
##
            rate.2
                     rate.3
## shift.3 0.1261663 0.00541799
```

We can now summarise the results of these data using 'traitMedusaSummary' and plotting the shifts on the phylogeny using 'plotPhylo.motmot'. These results show an increase at node 45 that we can visualise on the phylogeny.

```
trait.medusa.tm2.summary <- traitMedusaSummary(tm2.ml, cutoff=2, AICc=T)
trait.medusa.tm2.summary</pre>
```

```
## $ModelFit
             lnL n.params
                                AIC
##
                                          AICc
## shift.1 3.6472 3 -1.294399 -0.2943992
##
## $Rates
    node shiftPos
                            MLRate
                                            LowerCI
                                                            UpperCI
## 1
         clade 8.60339667795593 1.71610149858682 156.04985185668
##
## $optimalTree
##
## Phylogenetic tree with 28 tips and 27 internal nodes.
##
## Tip labels:
## A_alutaceu, A_inexpect, A_vanidicu, A_alfaroi, A_macilent, A_clivicol, ...
## Node labels:
## 2, 2, 2, 2, 2, ...
## Rooted; includes branch lengths.
```

```
colour_motmot <- plotPhylo.motmot(phy=phy.clade,
traitMedusaObject=trait.medusa.tm2.summary, reconType = "rates", type = "fan", cex=0.5,
edge.width=2)</pre>
```



the subset of the tree

Thomas and Freckleton (2012) showed the tm2 algorithm has a high type-one error rate. One way to ameriolate this is to estimate the level a one shift is supported when we know BM is the true model. For example, we could simulate 1000 BM datasets on the tree, estimate a single shift using the tm2 algorithm, and calculating the difference between the AlCcs for each BM and one shift model. We can these use this difference to estimate the AlCc 'penalty' the is needed to reduce the tm2 type-one error rate to 0.05. We could use this penalty in the 'cutoff' argument of the traitMedusaSummary argument.

This is shown but not run in the code below

# Time-slice model

A new addition to motmot is a Maximum likelihood model that allows for heterogeneous rates in different times of evolution. These models are seperate from the models that allow for heterogeneous rates among lineages, as modelled by the 'traitMedusa' algorithms.

The 'timeSlice' model is implemented using the 'transformPhylo.ML' function, using the argument model = 'timeSlice'. The function allows for two seperate models of evolution. In one, it is possible to test shifts

in evolution at times selected a priori. Alternatively, the fit of models can be tested at a range of different times, and the function will return the best-fitting model

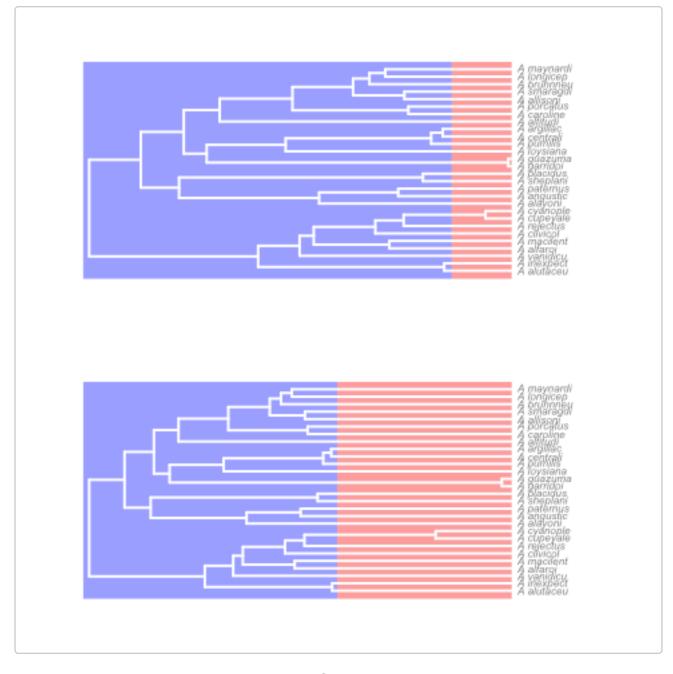
First we will test for a shift in the rate of evolution 10 million years ago.

```
timeSlice.10.ml <- transformPhylo.ML(y=male.length.clade, phy=phy.clade,
model="timeSlice", splitTime=c(10))</pre>
```

```
## [1] "BM model"
##
           lnL
                       AIC
                                  AICc
                                           sigma.sq
                                                      anc.state
## -0.28385540 4.56771081 5.04771081
                                         0.00185807
                                                     3.84948140
## [1] "shiftModel"
##
            lnL
                                      AICc
                                                           anc.state
                         AIC
                                               sigma.sq
                                            0.001006388
##
   2.946487446 2.107025107
                              3.846155542
                                                         3.860015288
##
                       rate2 time.split1
          rate1
   0.692072821 2.944767947 10.0000000000
```

We can use the function 'timeSliceSummary' to plot and summarise the results. The output summarises the best model according to AICc values. This function automatically plots the original tree showing the location of shift(s), and the colours show the relative rates in each time slice. The second plot below shows the same tree and colours, but with the branch lengths scaled to the ML optimised rates

```
outputSummary <- timeSliceSummary(timeSlice.10.ml, cutoff=0.001, cex.tip=0.5,
phylo.width=2, colour.ramp=c("blue", "red"))</pre>
```



We can also see other summarise information, such as the CI for each rate estimate.

```
outputSummary$RatesCI

## rates LCI UCI

## rate1 0.6920728 0.1873338 2.115629

## rate2 2.9447679 0.9633095 10.877403
```

Rather than testing the overall fit of each model, we can fit models to all times. The function automatically tests for all 1 Ma shifts between the age of the tree - 10 Ma, and the present + 10 Ma. We can specify a number of shifts we would like to test for. Here we will test for up to 3 shifts. The model will test one shift, save it, search for a second, save those two, etc...

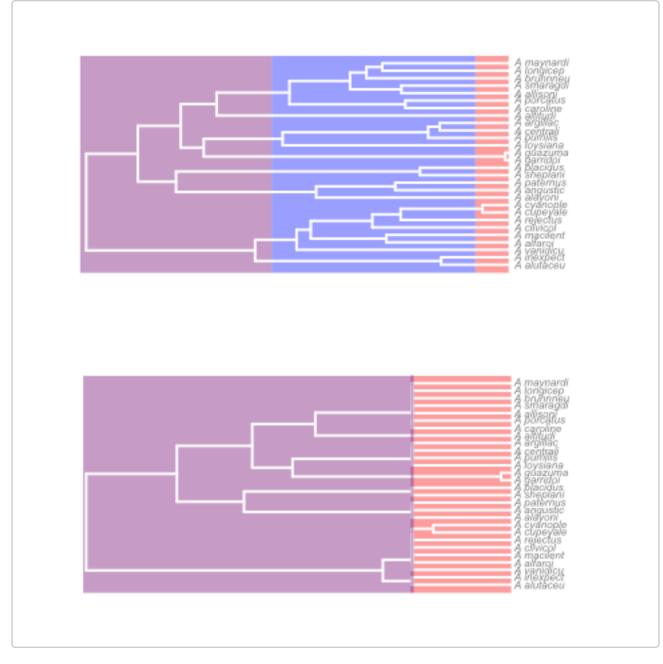
Here will modify the boundary age argument so all split times are tested between 65-5 Myrs, using the 'boundaryAge' argument. As we are not tested set times we need to set the number of splits to test using 'nSplits' - we will allow up to 3 splits

```
timeSlice.ml <- transformPhylo.ML(y=male.length.clade, phy=phy.clade,
model="timeSlice", nSplits=3, boundaryAge=5)</pre>
```

```
## [1] BM model
##
           lnL
                        AIC
                                   AICc
                                            sigma.sq
                                                       anc.state
                4.56771081
## -0.28385540
                            5.04771081
                                         0.00185807
                                                      3.84948140
## [1] shift 1
##
                          AIC
                                      AICc
            lnL
                                                sigma.sq
                                                            anc.state
    4.509949148 -1.019898297
                               0.719232138
                                            0.001070818
##
                                                          3.858225596
##
         rates1
                       rates2
                               time.split1
    0.613247119 3.633770015
                               5.545642000
##
   [1] shift 2
##
##
             lnL
                            AIC
                                         AICc
                                                    sigma.sq
                                                                  anc.state
    7.7641301455 -5.5282602909 -2.8009875637
                                                0.0006475698 3.8306699812
##
##
          rates1
                         rates2
                                       rates3
                                                 time.split1
                                                               time.split2
    3.4703057295 0.0000000100
                                 5.8887066391
                                                5.5456420000 39.5456420000
##
   [1] shift 3
##
##
             lnL
                            AIC
                                         AICc
                                                    sigma.sq
                                                                  anc.state
##
    7.9487076095 -3.8974152190
                                 0.1025847810
                                                0.0006734181
                                                              3.8417362745
##
                                                               time.split1
          rates1
                         rates2
                                        rates3
                                                      rates4
##
    0.0000000100
                  3.4550077536
                                 0.0000000100
                                                5.6540409952
                                                              5.5456420000
##
     time.split2
                   time.split3
## 39.5456420000 65.5456420000
```

And summarise the results. We can selected the cutoff AICc improvement needed to justify selecting the next model. Here we use the arbitary cut-off value of 1. We could test this formally by estimating the correct AICc value needed to reduced type-error > 5% by using BM simulated data (an example using the tm2 is shown above)

```
outputSummary <- timeSliceSummary(timeSlice.ml, cutoff=1, cex.tip=0.5, phylo.width=2,
colour.ramp=c("blue", "red"))</pre>
```



timeSlice plot

The output looks odd as the middle time period has be collapsed to zero

### Nested models of evolution

We can also tested models of nested evolution in which an ancestral model of BM evolution changes to a alternative model (EB, OU, kappa, delta, psi) within the phylogeny (Puttick, In Review)

Here we can show an example of BM -> OU and BM -> ACDC at node 44 of the phylogeny. However, neither of these is significantly better than BM

```
bm.model <- transformPhylo.ML(male.length.clade, phy=phy.clade, model="bm")
nested.acdc <- transformPhylo.ML(male.length.clade, phy=phy.clade, model="ACDC",
nodeIDs=c(44))

## Warning in transformPhylo.ML(male.length.clade, phy = phy.clade, model =
## "ACDC", : Confidence limits fall outside the current parameter bounds -
## consider changing lowerBound and/or upperBound

nested.ou <- transformPhylo.ML(male.length.clade, phy=phy.clade, model="OU",
nodeIDs=c(44))</pre>
```

## Warning in transformPhylo.ML(male.length.clade, phy = phy.clade, model =

## "OU", : Confidence limits fall outside parameter bounds - consider changing

```
## lowerBound and/or upperBound

1 - pchisq(nested.acdc$MaximumLikelihood - bm.model$logLikelihood, 1)

## [,1]
## [1,] 0.05740847

1 - pchisq(nested.ou$MaximumLikelihood - bm.model$logLikelihood, 1)

## [,1]
```

# Bayesian estimation of tree transformation models

## [1,] 0.361424

The function 'transformPhylo.MCMC' allows for the estimation of model parameters using Bayesian statistics. Models of lambda, delta, kappa, OU, ACDC, and psi can currently be modelled using transformPhylo.MCMC

The model allows for a pre-optimisation step. The model we test 30 (default) different deviations for the acceptance proposal distribution in order for the model to achieve an acceptance of around 0.44. This is done by default in the model but can be turned off by setting 'opt.accept.rate=FALSE'

We will run an MCMC chain of 2000 generations to estimate Pagel's lambda and discarding the first 10% ('200 generations ('burn.in = 0.1'). All the models use a 'uniform' prior for each of the parameters. For lambda, this is a uniform distribution between 0 and 1, meaning we think all potential values are equally likely

```
set.seed(20) # set seed so run will be identical - for example use only
lambda.mcmc <- transformPhylo.MCMC(y=male.length.clade, phy=phy.clade, model="lambda",
mcmc.iteration=2000, burn.in=0.1)</pre>
```

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## optimising acceptance ratio fine-tune
##
     running
acceptance attempt 0.463 best acceptance 0 best SD 0.448
   finished fine.tune
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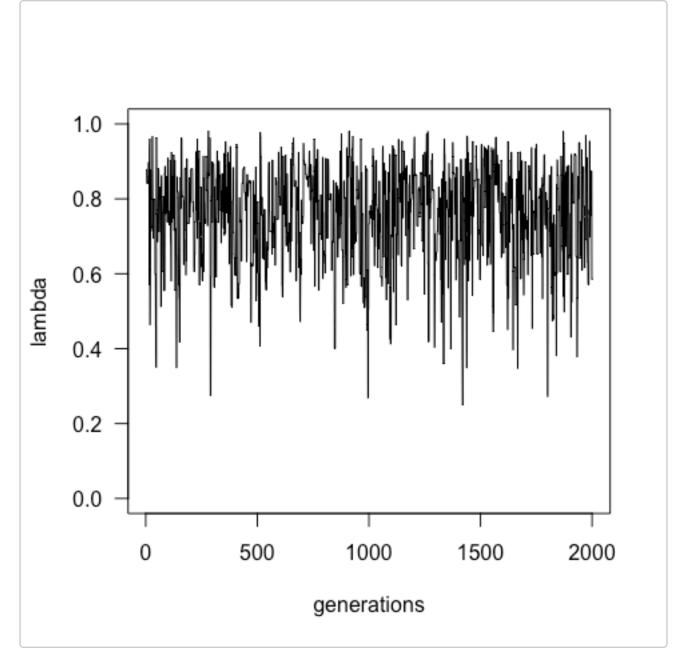
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## $median
##
      Lambda
## 0.7731313
##
## $`95.HPD`
## lower 95% HPD upper 95% HPD
       0.4845588 0.9449442
##
##
## $ESS
##
     Lambda
## 494.6253
##
## $acceptance.rate
## [1] 0.4708495
```

We can know check the posterior estimate of lambda and convergence of the model. The median and 95 Highest Posterior Density (HPD) is output by the model. Some convergene diagnostics are output as standard: Effective Sample Size (ESS) and acceptance rate. We aim for an ESS of at least 200 and an acceptance rate around 0.44

```
lambda.mcmc[1:4]
## $median
      Lambda
##
## 0.7731313
##
## $`95.HPD`
## lower 95% HPD upper 95% HPD
       0.4845588 0.9449442
##
##
## $ESS
##
     Lambda
## 494.6253
##
## $acceptance.rate
## [1] 0.4708495
```

Our lambda median value is 0.77 but there is a large 95% HPD (0.48-0.95). The ESS and acceptance rate look ok. We can also plot the trace from the MCMC chain

```
\label{eq:plot} \textbf{plot}(lambda.mcmc\$mcmc.chain, type="l", ylim=c(0, 1), xlab="generations", ylab="lambda", las=1)
```



MCMC trace for Pagel's lambda

We would hope this is flatish trace around the stable posterior distribution - it looks ok (if not great)

## References

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