A Rough Guide to CladeAge

Bayesian estimation of clade ages based on probabilities of fossil sampling

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1 Introduction

1.1 Summary

CladeAge is an add-on package for the Bayesian software BEAST 2 (Bouckaert $et\ al.$, 2014) which allows time calibration of phylogenetic trees based on probability densities for clade ages, calculated from a model of constant diversification and fossil sampling.

1.2 Background

In Bayesian node dating, phylogenies are commonly time calibrated through the specification of calibration densities on nodes representing clades with known fossil occurrences. Unfortunately, the optimal shape of these calibration densities is usually unknown and they are therefore often chosen arbitrarily, which directly impacts the reliability of the resulting age estimates. CladeAge overcomes this limitation by calculating optimal calibration densities for clades with fossil records, based on estimates for diversification rates and the sampling rate for fossils. This rate characterizes the frequency, at which fossils are preserved along branches of a phylogeny, and only fossils that are ultimately sampled (and published) by researchers are used for this measure.

1.3 Reference

A manuscript describing the CladeAge model is accepted (pending revision) for publication in Systematic Biology, and a preprint can be found on BioRxiv:

Matschiner M, Musilová Z, Barth JMI, Starostová Z, Salzburger W, Steel M, Bouckaert R (2016) Bayesian node dating based on probabilities of fossil sampling supports trans-Atlantic dispersal of cichlid fishes, BioRxiv preprint, doi: http://dx.doi.org/10.1101/038455.

1.4 Contact

If this guide still leaves questions you can contact the authors, or consult the BEAST user list on http://groups.google.com/group/beast-users.

2 CladeAge calibration densities

CladeAge calculates probability densities for the age of origin of a clade, given the age of its oldes fossil, based on an assumed model of constant diversification and fossil sampling. Under the further assumption that the age of origin of a clade is equivalent to the divergence time between this clade and its sister lineage, these probabilities can be used as calibration densities for Bayesian divergence-time estimation. The performance of CladeAge calibration densities for divergence-time estimation has been tested extensively with simulated and empirical datasets (Matschiner et al., 2016). The CladeAge model includes four required parameters (the first occurrence age, net diversification rate, turnover rate, and sampling rate) and one optional parameter (the sampling gap), which are described below. Note that ranges can be specified for all parameters to account for uncertainties in their estimates.

2.1 CladeAge model parameters: the first occurrence age

The first occurrence age of a clade is the age of the oldest fossil that can be reliably assigned to this clade. For example, the oldest currently known fossil taxon that can be assigned to cetaceans (whales) is †*Himalayacetus subathuensis* from the Indian Subathu Formation (Benton et al., 2015). Within this formation, the fossil was found in a layer characterized by the occurrence of the foraminifer Nummulites atacicus (Bajpai & Gingerich, 1998), which allows correlation with the Shallow Benthic Zone (SBZ) 8 (Vandenberghe et al., 2012). The ages of the boundaries of SBZ8 are not directly defined (Vandenberghe et al., 2012), but SBZ 8 is usually considered to correspond to nannoplankton zones NP10-11, and these are known to span 56.0-53.5 Ma (Vandenberghe et al., 2012). Thus, the first occurrence age of cetaceans is 56.0-53.5 Ma.

As becomes obvious from this example, the identification of the first occurrence age from the systematic, paleontological, and geological literature is not always straightforward. However, some excellent starting points for this investigation are Benton et al. (2015), the Fossil Calibration Database (http://fossilcalibrations.org), and the Paleobiology Database (https://paleobiodb.org). For conversions to absolute ages, the book The Geologic Time Scale 2012 (Gradstein, 2012) is highly recommended, but should be complemented with the latest version of the International Chronostratigraphic Chart, published by the International Commission on Stratigraphy (http://stratigraphy.org/index.php/ics-chart-timescale).

Similar to †*Himalayacetus subathuensis*, nearly all fossil ages are associated with uncertainties that often can be considered uniform within lower and upper boundaries. By specifying these boundaries as input for CladeAge, the uncertainty in first occurrence ages can (and should) be accounted for.

2.2 CladeAge model parameters: the diversification rates

To estimate the total phylogenetic history of a clade prior to the preservation of its oldest fossil record, CladeAge implements a birth-death model of constant diversification. Thus, the model includes both a speciation rate λ and an extinction rate μ . However, as diversification rates are more easily estimated, and more commonly reported in the literature as net diversification rate $\lambda - \mu$ and turnover rate μ/λ , CladeAge is written to accept as input net diversification rate and turnover rate, rather than speciation rate and extinction rate (internally, it then uses the specified net diversification rate and turnover rate to calculate speciation and extinction rates). If diversification rates for the clade of interest have not previously been reported in the literature, these can be estimated either from the fossil record alone or from separate phylogenies, using software such as BAMM (Rabosky, 2014), RPANDA (Morlon et al., 2015), MEDUSA (Alfaro et al., 2009), TRiPS (Starrfelt & Liow, 2016), PyRate (Silvestro et al., 2014), or paleotree (Bapst, 2012).

2.3 CladeAge model parameters: the sampling rate

Probability densities calculated with CladeAge are based on estimates of the sampling rate ψ of fossils, assuming that sampling is a constant Poisson process. The concept of a sampling rate of fossils is commonly used in the paleontological literature, and is considered to include all processes that lead to the identification and scientific description of a fossil. Thus, the sampling rate encompasses deposition of an organism in the sediment, fossilisation, survival of geological processes, outcropping of the rock formation including the fossil, discovery, correct taxonomic identification, and publication of the fossil find.

Estimates of sampling rates can be obtained by a variety of methods, including the freqRat method of Foote & Raup (1996), the survivorship analysis of Foote (2001), and the analysis of waiting times between sampling events, introduced by (Solow & Smith, 1997). All these methods have in common that a very large number of fossils, and ideally full occurrence data, i.e., information for each known fossil of a given group, is needed for reliable sampling rate estimates. Thus, for most studies aiming to time-calibrate phylogenies, the estimation of sampling rates for the investigated group is beyond the scope of the analysis. Fortunately however, sampling rates have already been estimated and published for many groups and can readily be used in CladeAge. For example, Foote et al. (1999) estimated the sampling rate of Late Cretaceous mammals as 0.03-0.06, Alba et al. (2001) found the sampling rate of Iberian mammals of the Neogene to be around 0.8, and Foote & Raup (1996) determined the sampling rate of European Jurassic bivalves to be around 0.41 (all values given are per lineage and per million years).

Previously, sampling rates have often been published on genus level instead of species level, and not per million years, but for shorter or longer time bins (Foote & Raup, 1996; Foote & Sepkoski, 1999; Friedman & Brazeau, 2011). In order to use these estimates in CladeAge, they must be transformed into species-level

estimates per million years (assuming that fossil ages are also specified in million years in BEAST). One way to transform genus level estimates into species level estimates is to calculate the average number of extant species per genus in the investigated group, assuming that this ratio also holds for fossil members of the same group. If rates have been given for time bins other than a million years, the rate conversion can be performed e.g. with the function Prob2sRate of the R package paleotree (Bapst, 2012). A compilation of published species-level sampling rate estimates for diverse groups of organisms can be found in Supplementary Table S1 of Matschiner $et\ al.\ (2016)$.

2.4 CladeAge model parameters: the sampling gap

As an optional parameter, a sampling gap can be specified. The sampling gap represents the time period at the very beginning of a clade's history, for which one can assume that the clade's preservation potential was negligible, effectively setting the sampling rate ψ to zero for this period. There are several reasons why one might assume the existence of a sampling gap: For example, the populations sizes of a clade might have been very small shortly after its origination, which could have reduced its potential to become preserved in the fossil record. Similarly, if the population has been geographically limited to a small region, which is more likely at the beginning of a clade's history, this would also lower its preservation potential. One might also assume that clades require time to evolve morphological features that allow fossils to be assigned to these clades. As the sampling rate ψ includes all processes from the fossilisation of an organism to its discovery and correct taxonomic assignment, ψ would also be zero for the time before the emergence of recognisable morphological features of a clade.

Whether or not a sampling gap should be used in CladeAge analyses is debatable, as one might argue that over time periods of millions of years, as are usually considered in divergence-time estimation, the times required to reach substantial population sizes and geographic distributions, or to evolve recognisable morphological features, are negligible. In fact, the latter may even go simultaneously with clade origination according to the theories of punctuated equilibrium and key innovations (Pennell et al., 2013; Rabosky et al., 2013). Note that a sampling gap was not used in combination with simulated datasets in Matschiner et al. (2016) and the performance of CladeAge with specified sampling gaps has not yet been tested.

2.5 Units

Note that the units used for calibrations should match those used for other parts of the model, so if you specify the parameters of the CladeAge model in millions of years, the clock prior should be in million of years as well. Obviously, the tree will then be interpreted in millions of years as well.

3 Applying CladeAge calibration densities

3.1 Which fossils to use

CladeAge calibration densities describe the probabilities for the age of a clade, given the age of its first occurrence. Thus, **only the oldest fossil of a clade should be used** for the specification of CladeAge calibration densities, and younger fossils of the same clade should be ignored. Obviously, there should be morphological evidence that the putatively oldest fossil record of a clade is in fact a member of that clade. Based on morphology, the general assignment of fossils to clades (regardless of the fossil's position within the clade) is usually more robust than determining whether or not the fossil is a member of the crown or stem group of that clade. Therefore, CladeAge calibration densities are designed to be applied to the origin of a clade regardless of whether its oldest fossil is part of the stem group or the crown group of the clade.

3.2 Which clades to use

In practice, it may not always be clear which clades in a phylogeny should be used for time calibration: If a fossil represents the earliest record of not only one clade, but of multiple nested clades, CladeAge calibration densities could be used to constrain the age of origin of all these clades, only of the most inclusive of these clades, or only of the least inclusive clade. Furthermore, if two sister clades both possess a fossil record, these fossils could be used to constrain the ages of both of the two clades. However, as the ages of the two clades are necessarily linked by their simultaneous divergence, two time constraints would effectively be placed on one and the same node. Instead, it may seem more intuitive to use only the older of the two fossils for time calibration and disregard the younger fossil. Thus, a number of schemes could be chosen to select clades for calibration (illustrated in Fig. 2 of Matschiner et al. 2016). The performance of these calibration schemes was tested in Matschiner et al. (2016) with a wide range of simulated datasets. Based on the results of these analyses, we recommend a calibration scheme according to which fossils are used to strictly constrain all clades, for which these fossils represent the oldest known record (see Fig. 1 on the next page). This means that some fossils may be used to constrain the ages of more than one of several nested clades, particularly if groups with poor fossil records are investigated. It also means that constraints on clade ages should be applied irrespective of the fossil record of the sister group of the clade. Thus, even if the sister group has older fossil records, the oldest fossil of a clade should still be used as a constraint. A positive aspect of this is that sister groups do not need to be known before the analysis, unlike in traditional node-dating approaches.

Thus, CladeAge calibration densities should be applied to all clades that have a fossil record, and bias would be introduced if only clades with particularly old (or young) fossils would be used. There are two exceptions to this rule:

- 1. For clades that are recovered in phylogenies, but are not supported by morphological synapomorphies, the sampling rate is effectively reduced to zero, as it would be impossible to assign fossils directly to these clades based on morphology. Indirectly, these clades might still have a fossil record as it may be possible to assign fossils to subclades that are characterized by morphological synapomorphies. Nevertheless, these clades should not be constrained with fossils assigned to subclades (see Matschiner et al., 2016, for a more detailed discussion of this case).
- 2. While it is not necessary to know the sister group of calibrated clades, the sister group should still be included in the dataset. If the true sister group of a calibrated clade was instead missing from a phylogeny, the origin of the branch leading to the calibrated clade would be older than the clade itself, which would violate assumptions made when using CladeAge probability densities as calibration densities in BEAST 2. Thus, in this case, the age of the clade should not be constrained. To ensure that a clade's sister group is included in the phylogeny even if the identity of the sister group is not known, one could simply include representatives of all possible sister groups in the dataset. Missing sister groups can be avoided completely if so-called diversity trees (Alfaro et al., 2009) are used for the analysis, where each extant species of a given group can be assigned to one of the tips of the phylogeny.

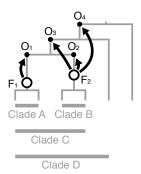
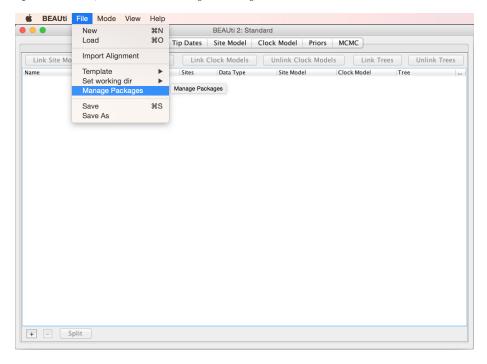


Figure 1: Calibration scheme to be used with CladeAge calibration densities. When using CladeAge calibration densities, fossils should be used as constraints for the age of origin of all clades, for which the fossil represents the oldest known record. Here, fossils F_1 and F_2 , represented by white circles, are the oldest fossils of clades A and B, respectively, and no fossils are known outside of these two clades. As F_2 is older than F_1 it also represents the oldest record of clades C and D. Thus, F_1 should be used to constrain the age of clade A (marked with a black dot and the label O_1), and F_2 should be used to constrain the age of origin of clades B, C, and D (marked with O_2 , O_3 , and O_4). Note that since clades A and B are sister lineages, O_1 and O_2 are identical in age.

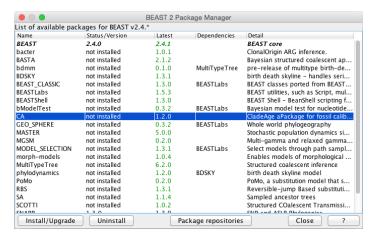
4 Installing CladeAge

4.1 Using BEAUti

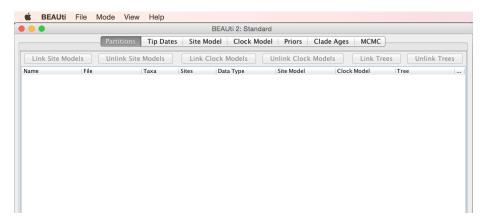
Like other add-on packages for BEAST 2, CladeAge can be installed through the package manager that's built into BEAUti, a utility tool that comes with the BEAST 2 download (see http://beast2.org for details). To install CladeAge, open BEAUti, and select *Manage Packages* from the File menu:



Then select the CA package from the list and click the Install/Upgrade button:



Restart BEAUti, and a new panel for *Clade Ages* should have appeared, as you can see from the buttons in the top row of the BEAUti window:



The *Clade Ages* panel is where you can specify rate estimates for speciation, extinction, and fossil sampling, and CladeAge calibration densities will be automatically calculated based on these parameters (see below).

4.2 Using the command line

Alternatively, you can use the command-line utility addonmanager to install CladeAge (or other packages for BEAST 2), which is useful for cluster computers without GUI access. To see the available options, type

PATH_TO_BEAST/bin/addonmanager

in a console window (make sure to replace *PATH_TO_BEAST* with the actual path to the BEAST 2 directory). You should see the following output:

```
Usage: addonmanager [-list] [-add <NAME>] [-del <NAME>] [-useAppDir]
        [-dir <DIR>] [-help]
-list List available add-ons
-add Install the <NAME> add-on
-del Uninstall the <NAME> add-on
-useAppDir Use application (system wide) installation directory. Note this requires writing rights to the application directory. If not specified, the user's BEAST directory will be used.
-dir Install/uninstall add-on in directory <DIR>. This overrides the useAppDir option
-help Show help
```

To install the CladeAge package with addonmanager, use

PATH_TO_BEAST/bin/addonmanager -add CA

4.3 Manual installation

If for some reason installation with BEAUti or addonmanager is not possible, you can also install the CladeAge package manually. To do so, download the latest version of the compressed CladeAge package (named CA.addon.vX.X.x.zip, where X.X.X is a version number) from https://github.com/CompEvol/CBAN/releases. Uncompress the package, and place the directory CA in one of the following locations, depending on your operating system (make sure to replace USERNAME with your user name, and X with the version number of BEAST 2).

 $Windows: \verb"Users\username" \verb| Beast\2.X|$

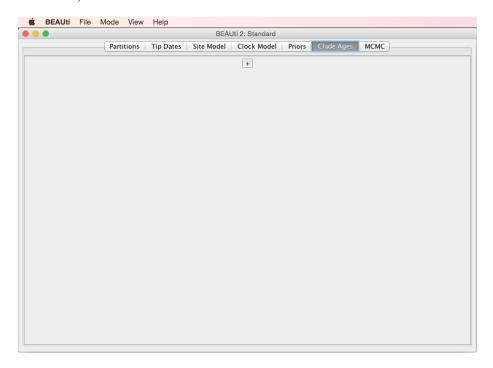
 Mac : /Users/USERNAME/Library/Application Support/BEAST/2.X

Linux: /home/USERNAME/.beast/2.X

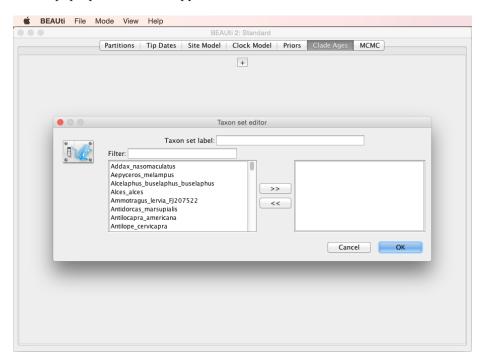
5 Specifying CladeAge calibration densities

5.1 Using BEAUti

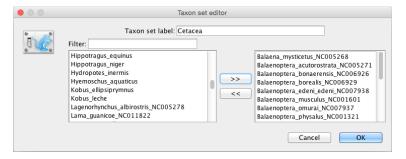
To specify a new CladeAge calibration density in BEAUti, open the *Clade Ages* tab (after you imported a sequence alignment with *Import Alignment* in the *File* menu).



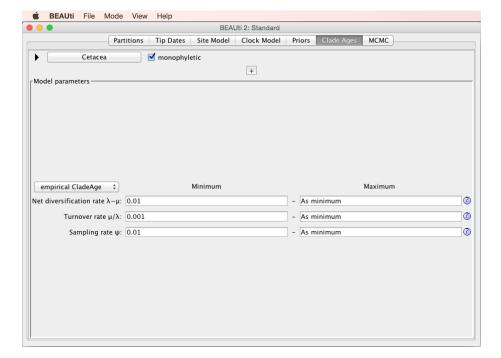
To specify a taxon set click on the little '+' button in the middle of the window. A new pop-up window will appear:



The dataset used in this example is an alignment of mitochondrial genome sequences of 183 species of cetartiodactyls (even-toed ungulates, including cetaceans) (Hassanin et al., 2012). To constrain the age of origin of cetaceans with the fossil †Himalayacetus subathuensis (see 2.1), a taxon set comprising all cetaceans first needs to be specified. To do so, provide a taxon set label (here Cetacea), select taxa that are part of this clade, and click the '>>' button to move these into the ingroup.



Click OK.



The Clade Ages panel should then look similar to this:

Here, you can specify estimates for the net diversification rate, the turnover rate, and the sampling rate. If both minima and maxima are specified for these parameters, CladeAge will account for the uncertainty of in the rate estimates. If only a minimum is specified, it will be considered as a point estimate without uncertainty. The default values for these parameters should not be trusted, instead you should identify reasonable estimates for all three parameters from the literature or based on separate analyses (see 2.2 and 2.3).

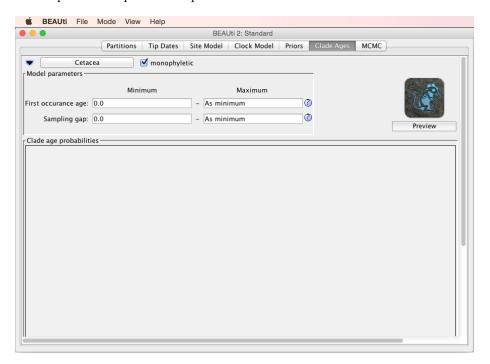
The specified rates for net diversification, turnover, and sampling will be used for all CladeAge calibration densities specified through BEAUti, thus assuming that all clades included in the dataset share the same characteristics of diversification and preservation. This assumption may be unrealistic for large phylogenies including highly divergent groups. To specify different rates for different clades, the XML file produced by BEAUti can be edited as described in 5.2.

If you click on the question marks to the right of the input fields, more information will be provided in a pop-up window. The drop-down menu just above *Net diversification rate* indicates the type of CladeAge calibration density. The *empirical CladeAge* densities should be fine for all purposes (but if you're interested in the alternative options, please see 6).

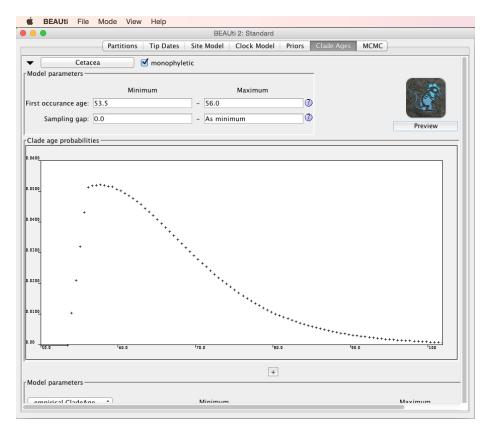
At the top left of the panel, you'll see a field with the specified name of the clade (here *Cetacea*). The checkbox to the right of this field indicates that by default clades used for fossil calibrations are expected to be monophyletic. If

you uncheck this box, the topology of the taxa included in the clade will not be constrained. However, clades for which monophyly is questionable should not be constrained in the first place, as the assignment of fossils to these clades is unlikely to be reliable if the clade itself is not monophyletic. Thus, we advice to leave the *monophyletic* checkbox set.

To specify the first occurrence age (and the sampling gap) for this clade, click on the little triangle to the left of the field with the clade name (*Cetacea*). A new part of the panel will open:



Enter the minimum and maximum for the first occurrence age. If no maximum is provided, the minimum will be used as a point estimate, assuming that the fossil age is known exactly. To ignore the sampling gap, just leave the minimum and maximum fields as they are. Again, more information about the two parameters can be displayed by clicking on the question marks to the right of the input fields. To see the shape of the CladeAge calibration density calculated on the basis of all specified parameters, click the Preview button below the CladeAge icon. The density shown in the figure on the next page is based on a first occurrence age between 56.0 and 53.5 million years, and more or less arbitrarily chosen point estimates for net diversification rate (0.04 per million year), turnover rate (0.5), and sampling rate (0.05 per million year).



To specify calibration densities for additional clades, click the triangle at the top left again to collapse the information for the first calibration density, and click little '+' button in the middle of the window once again. Once you have specified constraints for all clades, don't forget to set all other options for BEAST 2 analyses in the panels Site Model and Clock Model and check the prior distributions in the Priors panel. Then, set the length of the MCMC in the MCMC panel and save the XML file for BEAST 2 (more information on these settings can be found, e.g., in the Divergence Dating tutorial available from https://beast2.org/tutorials/).

5.2 Manual specification in the XML

Obviously, BEAUti is just one way to produce XML input files for BEAST 2. You could also prepare XML files manually by modifying the examples provided in the *examples* folder that comes as part of the BEAST 2 download, or you could use a script such as BEASTmasteR (https://github.com/nmatzke/BEASTmasteR; a similar script is provided here: https://github.com/mmatschiner/Introgression-Tutorial). However, if the XML is prepared in any other way than with BEAUti, CladeAge calibration densities will also have to be added to the XML code manually.

The following code block, which is placed inside the *posterior* distribution element, specifies a CladeAge calibration density in the XML file. As you can see, changing the minimum and maximum for any of the CladeAge model parameters is rather straightforward - for example, to change the minimum and maximum sampling rate, just replace 0.05 after minSamplingRate and maxSamplingRate with another value.

```
<distribution id="fossilCalibrations" spec="util.CompoundDistribution">
 <distribution id="Cetacea.fossilprior"</pre>
     spec="beast.math.distributions.FossilPrior"
     monophyletic="true" tree="@Tree.t:id">
   <fossilDistr id="FossilCallibration.0"</pre>
       spec="beast.math.distributions.FossilCalibration">
     <parameter id="RealParameter.0" name="minOccuranceAge">53.5</parameter>
     <parameter id="RealParameter.01" name="maxOccuranceAge">56.0</parameter>
     <parameter id="minDivRate" name="minDivRate">0.04</parameter>
     <parameter id="maxDivRate" name="maxDivRate">0.04</parameter>
     <parameter id="minTurnoverRate" name="minTurnoverRate">0.5</parameter>
     <parameter id="minSamplingRate" name="minSamplingRate">0.05</parameter>
     <parameter id="maxSamplingRate" name="maxSamplingRate">0.05</parameter>
     <parameter id="RealParameter.02" name="minSamplingGap">0.0</parameter>
     <parameter id="RealParameter.03" name="maxSamplingGap">0.0</parameter>
   </fossilDistr>
   <taxonset id="Cetacea" spec="TaxonSet">
     <taxon id="Balaena_mysticetus_NC005268" spec="Taxon"/>
     <taxon id="Balaenoptera_acutorostrata_NC005271" spec="Taxon"/>
     <taxon id="Balaenoptera_bonaerensis_NC006926" spec="Taxon"/>
     ... (a list of all taxon ids included in this clade) ...
     <taxon id="Tursiops_truncatus_NC012059" spec="Taxon"/>
   </taxonset>
 </distribution>
</distribution>
```

To specify multiple CladeAge calibration densities, one could simply copy paste the above code block several times, each time replacing the species ids in the taxonset element, depending on which species are part of the calibrated clade. When doing so, one should also take care to change the id of the taxonset element (which says Cetacea in the code block example). Additionally, the id of the distribution element that says Cetacea fossilprior in the example should be changed as well as the id of the fossil Distr element (which is Fossil Callibration. 0 in the example code). Furthermore the ids of the parameter elements (which are RealParameter.0, RealParameter.01, minDivRate, etc. in the example) will need to be modified in copied blocks, and of course parameter values may need to be adjusted. It does not matter what the ids are replaced with (as they will not be referred to by other elements), they should be changed only for the reason that the same ids are not used for multiple elements (which would cause BEAST 2 to stop the run). Also note that the tree id Tree.t:id to which the 'Cetacea.fossilprior' distribution element refers to will need to be changed to match the actual tree id specified in the XML.

6 Types of CladeAge calibration density distributions

If you've been wondering what the difference is between the different types of CladeAge calibration density distributions available through BEAUti (4.1), the following descriptions may be helpful. However, note that the so-called *empirical CladeAge* distributions which are used by default have been thoroughly tested and there's currently no reason to use any other types of CladeAge distributions instead. So, the below descriptions are given mostly for the sake of completeness and can safely be skipped.

What all types of CladeAge calibration density distributions have in common is that they are based on calculations of the clade age probability density for 100 time points between the minimum first occurrence age of the clade and a maximal time point that is predetermined based on a quick approximation (details given in Matschiner *et al.*, 2016). However, the difference between the types of CladeAge distributions lies in the way in which these 100 calculated densities are turned into probability density distributions.

- Empirical CladeAge distributions: In the default empirical CladeAge distributions, the calculated probability densities are used directly for the respective time points. Probability densities for times in between these time points are interpolated from the probability densities of the two neighbouring time points, using a linear regression. For all times older than the oldest sampled time point (the tail of the distribution), probability densities are approximated by an exponential distribution that is calculated on the basis of the two oldest time points, so that the probability densities calculated for these two time points lie on this exponential distribution. Finally, empirical CladeAge distributions are scaled so that the total probability mass becomes 1. Empirical CladeAge distributions are the only available distribution type when a sampling gap (see 2.4) is specified.
- Fitted CladeAge distribution: CladeAge allows the fitting of what we call fitted CladeAge distributions to the calculated probability densities. These distributions are similar to truncated lognormal distributions, except that the parameterisation is simplified. The probability density function of an fitted CladeAge distribution is as follows:

$$f_u = \begin{cases} 0, & \text{if } u \le 0\\ \frac{c}{u+s} \times e^{\frac{-(\ln(u+s)-m)^2}{w}}, & \text{if } u > 0, \end{cases}$$

where $u = t_o - t_f$ and parameters c, s, m, and w (all ≥ 0) can be optimised so that f_u fits the calculated probability densities and its integral sums to 1. We have chosen to introduce this type of distribution, because we found truncated lognormal distributions to provide a near-perfect fit to the calculated probability densities, however the parameterisation of lognormal

distributions makes analytical handling unnecessarily complicated. While providing the same fit as truncated lognormal distributions, the simplified parameterisation of *fitted CladeAge distributions* facilitates analytical solution of distribution parameters, as well as analytical integration of the distribution.

Knowing the integral of a distribution is necessary in order to analytically account for uncertainty in the age of first occurrence. The integral of an *fitted CladeAge distribution* is

$$\int_0^a f_u \, \mathrm{d}u = \begin{cases} 0, & \text{if } a \le 0\\ -\frac{c\sqrt{\pi w}}{2} \times \mathrm{erf}(\frac{m - \ln(a + s)}{w}), & \text{if } a > 0, \end{cases}$$

where erf is the error function, which CladeAge calculates via a numerical approximation with a maximal error of 1.2×10^{-7} (Press *et al.*, 2007).

The analytical solution of the parameters of an fitted CladeAge distribution is possible when probability densities of only 4 instead of 100 time points are considered, and the fit of the distribution is only slightly worse if these four time points are carefully selected a priori. This massively reduces computation time not only by the limited number of time points for which probability densities must be calculated, but also because the analytical solution of distribution parameters is much faster than the distribution fitting through minimisation of root mean square deviations. As a result, the time needed for the calculation of fitted CladeAge distribution parameters is on the order of 0.005 seconds. Of course, if prior distributions are calculated just once before a time-calibration analysis, it hardly matters whether this calculation takes 0.005 s or 1 s. However, the fast analytical calculation of distribution parameters in would allow repeatedly recalculating these parameters during a BEAST run without substantial delay. In this case, diversification rates λ and μ would not need to be specified by the user. As the BEAST MCMC estimates these parameters anyway, these estimates could be used directly for fitted CladeAge distributions. Whenever the BEAST estimates for λ and μ change during the MCMC run, fitted CladeAge distributions are recalculated with the updated estimates. This feature is experimental and run-time recalculation of fitted CladeAge distributions has not been made available yet.

• Standard distributions: Besides empirical and fitted CladeAge distributions, CladeAge allows the fitting of commonly used distribution types against the calculated probability densities. Available distributions include exponential, lognormal, normal and gamma distributions. These distributions are scaled during the fitting process, however the scaling factor is not reported and should be ignored when these distributions are used as priors in Bayesian analyses, otherwise the probability mass of the distribution would not sum to 1.

References

- Alba DM, Agustí J, Moyà-Solà S (2001) Completeness of the mammalian fossil record in the Iberian Neogene. *Paleobiology*, **27**, 79–83.
- Alfaro ME, Santini F, Brock CD *et al.* (2009) Nine exceptional radiations plus high turnover explain species diversity in jawed vertebrates. *Proc. Natl. Acad. Sci. USA*, **106**, 13410–13414.
- Bajpai S, Gingerich PD (1998) A new Eocene archaeocete (Mammalia, Cetacea) from India and the time of origin of whales. *Proc. Natl. Acad. Sci. U.S.A.*, **95**, 15464–15468.
- Bapst DW (2012) paleotree: an R package for paleontological and phylogenetic analyses of evolution. *Method Ecol. Evol.*, **3**, 803–807.
- Benton MJ, Donoghue MJ, Asher RJ et al. (2015) Constraints on the timescale of animal evolutionary history. Palaeontol. Electron., 18.1.1FC, 1–106.
- Bouckaert R, Heled J, Kühnert D *et al.* (2014) BEAST 2: a software platform for Bayesian evolutionary analysis. *PLOS Comput. Biol.*, **10**, e1003537.
- Foote M (2001) Inferring temporal patterns of preservation, origination, and extinction from taxonomic survivorship analysis. *Paleobiology*, **27**, 602–630.
- Foote M, Raup DM (1996) Fossil preservation and the stratigraphic ranges of taxa. *Paleobiology*, **22**, 121–140.
- Foote M, Sepkoski JJ Jr (1999) Absolute measures of the completeness of the fossil record. *Nature*, **398**, 415–417.
- Foote M, Hunter JP, Janis CM, Sepkoski JJ Jr (1999) Evolutionary and preservational constraints on origins of biologic groups: divergence times of eutherian mammals. *Science*, **283**, 1310–1314.
- Friedman M, Brazeau MD (2011) Sequences, stratigraphy and scenarios: what can we say about the fossil record of the earliest tetrapods? *Proc. R. Soc. B*, **278**, 432–439.
- Gradstein FM (2012) The Geologic Time Scale 2012. Elsevier, Amsterdam, Netherlands.
- Hassanin A, Delsuc F, Ropiquet A *et al.* (2012) Pattern and timing of diversification of Cetartiodactyla (Mammalia, Laurasiatheria), as revealed by a comprehensive analysis of mitochondrial genomes . *C R Biol*, **335**, 32–50.
- Matschiner M, Musilová Z, Barth JMI et al. (2016) Bayesian node dating based on probabilities of fossil sampling supports trans-Atlantic dispersal of cichlid fishes. BioRxiv preprint.

- Morlon H, Lewitus E, Condamine FL et al. (2015) RPANDA: an R package for macroevolutionary analyses on phylogenetic trees. Method Ecol. Evol., 7, 589–597.
- Pennell MW, Harmon LJ, Uyeda JC (2013) Is there room for punctuated equilibrium in macroevolution? *Trends Ecol. Evol.*
- Press WH, Teukolsky SA, Vetterling WT, Flannery BP (2007) Numerical Recipes: The Art of Scientific Computing. Cambridge University Press, New York, USA.
- Rabosky DL (2014) Automatic detection of key innovations, rate shifts, and diversity-dependence on phylogenetic trees. *PLOS ONE*, **9**, e89543.
- Rabosky DL, Santini F, Eastman J et al. (2013) Rates of speciation and morphological evolution are correlated across the largest vertebrate radiation. Nat. Commun., 4, 1958.
- Silvestro D, Salamin N, Schnitzler J (2014) PyRate: a new program to estimate speciation and extinction rates from incomplete fossil data. *Method Ecol. Evol.*, 5, 1126–1131.
- Solow AR, Smith W (1997) On fossil preservation and the stratigraphic ranges of taxa. *Paleobiology*.
- Starrfelt J, Liow LH (2016) How many dinosaur species were there? Fossil bias and true richness estimated using a Poisson sampling model. *Phil. Trans. R. Soc. B*, **371**, 20150219.
- Vandenberghe N, Hilgen FJ, Speijer RP (2012) The Paleogene period. In: *The Geologic Time Scale 2012* (ed. Gradstein FM), pp. 855–921. Elsevier, Amsterdam, Netherlands.