

Using Simulations To Examine Temporal Discrepancy Between the First Appearance Times of Ancestors and Descendants

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Du and Alemseged 2019: Temporal Evidence and Ancestor-Descendant Relationships

In an incomplete fossil record, the order of appearance in the fossil record may be a poor reflection of phylogeny. Additional complications arise when morphologically-delimited taxa are persistently found over geological durations, including when they ‘persist through’ a speciation event that produces a differentiated descendant morphotaxon. In such cases, the ‘ancestral’ morphotaxa will be extant at the same point in time (but perhaps not the same location) as their ‘descendant’ morphotaxa, and thus the incomplete record may preserve traces of the ‘descendant’ before its own ‘ancestor’. But how often should we expect to see this in the real fossil record? How unlikely is an age ‘discrepancy’ (as I will term it) of a given size?

Du and Alemseged (2019, hereafter D&A) attempt to answer this question, with specific reference to whether the *A. sediba* morphotaxon could possibly be the ancestor to *Homo*, whose first supposed fossil occurrences appear to post-date its own occurrence by a considerable duration (for the hominin fossil record). Their finding is that they overall find the observed ancestor-descendant discrepancy to be of an unlikely duration. This document is an investigation of that claim. I’m not an anthropologist, and I hold no opinion whatsoever to the actual data, so I am going to largely accept their statements about dates and taxon identity, and mainly interrogate their methods.

D&A present two lines of arguments to evaluate the supposed ancestor-descendant hypothesis, given the temporal discrepancy:

- a) A model describing the probability of a discrepancy of a given magnitude, under a certain set of assumptions. (Described more below.) This model finds a very low probability of a discrepancy as large as the observed discrepancy, and consistently finds such under a range of conditions.
- b) An empirical dataset of the geologic age difference between the ‘first found’ fossils for 28 previously-hypothesized ancestor-descendant pairs from the paleoanthropology literature. In other words, if each species was only ever a single find - the first reported collection of that find with no subsequent collections- what would the age difference be between those two species? Only one A-D pair has a discrepancy (with the descendant’s first-find being geologically dated older than the first-find of the descendant), and that discrepancy is very small relative to the gap between *Homo* and *A. sediba*.

I will critique these arguments in reverse order.

The Metadata of Published Reports of Ancestor-Descendant Pairs

My initial concern upon reading the paper was that ancestor-descendant pairs as reported from the literature are often a limited subset of what might be true. As D&A describe in their discussion, this might particularly impact their conclusion because temporal information is (nearly always) taken into account when someone decides to suggest something is ancestral to something else. Thus, they use the first-finds of each taxon instead, based on the idea that the hypothesized ancestor-descendant relationships might be based on later, roughly independent temporal information.

I rather like the attempt using first-finds to avoid the issue of a taxon's whole temporal range being non-independent of it being placed as an ancestor or descendant, but I'm not sure its enough.

First, does first-finds remove the bias? I am not entirely convinced, because when people make arguments for ancestor-descendant pairs, I think that's more likely to take into account temporal information from their earliest collections, not their latest collections. I mean, in the invert paleo world, newly published finds often change a taxon's temporal range - but that information might take decades for someone else to notice and take that into account in their interpretations. Thus, the first find and the original geochronological interpretation of that find is most influential. I would probably like to know (a) what's the geologic age difference if we restricted each pair to looking at the most recent collections from each species, and/or (b) what's the distribution of age difference look like if we took all the collections from each taxon in these ancestor-descendant pairs, and used a Monte Carlo approach to sample a single collection at random from each taxon, and look at the distribution in difference in age for all these artificial 'one-collection' taxa. (This is similar to Alroy's 1996 analysis of Cope's Rule, where he sampled subsampled species pairs from genera to examine ancestor-descendant trends in body size change.) A full examination would compare these distributions of randomized age differences, to the age differences in the actual, currently-accepted first appearance times for those taxa.

Second, while I absolutely believe that ancestor-descendant relationships is something we should talk about and consider, but I'm not convinced of the utility of previously published pairs as a baseline for what we should expect of ancestor-descendant pairs at large. For example, in my 2017 paper with Melanie Hopkins on the finely resolved record of late Cambrian ptercephaliid trilobites, we found support for up-to 16 ancestor-descendant pairs, when really only 9 pairs had been proposed previously in the literature (including several pairs where which taxon was the ancestor, and which was the descendant, was uncertain, due to uncertainty in the first appearance dates). Now, no single phylogenetic hypothesis (the method used produces samples containing hundreds of possible trees) contained all 16 pairs, but a single tree often had higher numbers of pairs than had been previously proposed. So, my expectation is that for most groups, the inclination to label specific ancestor-descendant pairs has probably been overly conservative (one exception may be ammonites or planktonic foraminifera, where systematics is still largely very traditional rather than based on quantitative analysis of character matrices, and large numbers of taxa are placed as stages along long ancestor-descendant sequences of anagenesis). Maybe hominids are like those groups, and workers have been less conservative than they should have been, but maybe not, and my expectation is that thus the ancestor-descendant pairs reported for any given group is an extremely conservative subset of those, probably over-representing those that are the most consistent with being ancestor-descendant pairs in terms of their morphological and stratigraphic relationships. Overall, this adds up to an overwhelming bias against proposing ancestors that appear later than their descendants, a bias that probably leads to us underestimating the true occurrence of ancestors occurring latter than their descendants.

Third, I would still have problems with comparison because this is so unlike the ancestor-descendant pair we are most interested in - *A. sediba* and *Homo*. Well, the first collection of *Homo* would be, **well**, us recognizing that we are extant human beings (first fossil collection would be - well, I really have no idea, but its probably pretty geologically recent). Thus, that pair has no discrepancy as evaluated by the first-find approach. One taxon is also a species known from a single very time-constrained set of collections, and the other is a genus, that becomes relatively diverse (encompassing multiple species or sub-species or proto-species or whatever you want to call them. . .). Under a standard birth-death-sampling model, we would expect that such increased lineage diversity makes sampling *Homo* later much more likely later than earlier, even if there is a long 'fuse' interval with a very incomplete record. (Note though that this dynamic, admittedly, strengthens their case for their probabilistic model.) I think really we would need to identify a specific *Homo* species, and talk about that species-pair specifically rather than the genus itself.

Du & Alemseged's Model of Range Overlap and Ancestor-Descendant Temporal Discrepancy

In my data analysis courses, I tell students that models are just sets of assumptions about the world. We may describe those assumptions as mathematical relationships, but really any model is just a bundle of assumptions,

and models derive all their explanatory power from the strength and specificity of those assumptions. So what are D&A assumptions? Three of these are explicitly laid out:

- 1) The two taxa in question are assumed to have true ranges (the time between the true time of origination and extinction) of equal duration. In this case, the two taxa are both assumed to have 0.97 Ma (mega-annum) ranges. That's an average taken from Robinson et al - its obviously not true for *Homo*, which obviously has a longer duration than 1 Ma. It isn't clear from D&A what the choice of 0.97 Ma or inferring equal length ranges has on their findings.
- 2) The fossil record is assumed to have uniform sampling, across space and time, and across both the ancestor and descendant. As I already indicated above, the fact *Homo* encompasses multiple species would effectively violate this assumption, but the if that effect was taken into account, it would make it even more unlikely for the record to preserve *Homo* earlier in geologic time, and thus the bias works in agreement with D&A's findings. This is a common assumption of sampling models for the fossil record, and while it is often critized (see Steven Holland's work in particular), D&A far and away exceed the typical defense against this criticism by presenting evidence that hominin.
- 3) Sampling events (collections, occurrences, fossil horizons - whatever you want to call them!) are independent in time and across lineages. This is a common assumption of sampling models for the fossil record, and there aren't many cases where I would argue for the opposite.

Others are not so explicit:

- 4) This is a direct ancestor-descendant relationship. There are no additional, 'unknown unknown' taxa with similar sampling rates. Often, when we find ancestor-descendant pairs in the fossil record, we are probably not finding direct ancestor-descendant pairs (this morphotaxon is *directly* descended from this other morphotaxon) but rather we might be missing a few completely un-sampled, un-observed morphotaxa in between. As Foote (1996) recognized, ancestor-descendant relationships are probably relatively common in the fossil record, especially indirect ancestor-descendant relationships. Now, recognizing the possibility for 'unseen' lineages would decrease the probability of finding larger A-D discrepancies, so this bias works against D&A's test (and thus strengthens their findings).

From this, they derive the probability that any one occurrence of a descendant would occur before any one occurrence of an ancestor. Interestingly, and surprisingly to me, they find that sampling rate doesn't actually impact this probability - a well-sampled fossil record and poorly-sampled fossil record are expected to have very similar probabilities for a given ancestor-descendant discrepancy, for two taxa that have identical durations, and share the same uniform sampling rate (whatever that rate happens to be).

Some properties of this model are unclear in the draft, such as 'what if taxon range is varied?'. So let's consider scenarios under this model ourselves. In doing this, I will avoid relying on or referring to the supplemental R script provided by D&A, so that I am sure I understand all the relevant details.

This model mainly has three major variables related to intervals of various length: the duration of the range for each taxon (T_R , which D&A set to 0.97 Ma), the observed minimum stratigraphic overlap between the two taxa, which is a fixed variable of interest and is essentially the A-D discrepancy, or difference in age between the descendant and the ancestor (T_d , which D&A set to 0.8 Ma, the apparent discrepancy between *A. sediba* and *Homo*), and the true, total amount of overlap between the ancestor and the descendant (both of which, note well, have equally long ranges), and is treated as a random variable in D&A's model (T_o).

Several additional variables are worth noting. With respect to D&A's general notation and figures, the age of the ancestor's horizon is H_A , the age of the descendant's horizon is H_D , and the offset between an descendant's true time of origination and its time of sampling is X_D , while X_A is designated as the offset between the descendant's and ancestor's horizons, minus the minimum discrepancy (T_d). X_A and X_D are thus treated as random variables in this model, with minimums at 0. For the descendant's horizon to post-date the ancestor's horizon by a gap as large as T_d , and for all other assumptions to hold, $T_R > T_o > T_d$, and thus putting bounds on T_R and T_d defines the range of possible values for T_o . Additionally, $X_A + X_D = T_o - T_d$, $H_A - H_D > T_o$ and $X_D \leq X_A$ (D&A state greater than, but I do not comprehend any reason for why they cannot be equal and for all other conditions to still hold).

Equation 1 describes the probability of one fossil being found in a horizon within the region of overlap defined by $T_o - T_d$, and equation 2 describes the joint probability for two fossils (one for the ancestor, one for the descendant) from this limited region (referred to as $\Pr(\text{endA_endD})$). This is used in equation 3 (which is derived through a number of steps, during which the sampling rate drops out) to find the probability of $X_A > X_D$. By alternatively setting X_D or X_A equal to a random variable τ and iterating over that variable, D&A determine that the probability of $X_A > X_D$ is $1/2$, regardless of whether X_A or X_D is set equal to τ . This leads them to state in equation 4 that $1/2$ is also the probability of $H_A - H_D > T_d$, when conditioned on only sampling horizons from the region of ‘excess’ overlap (D&A continually refer to this ‘excess overlap’ interval, or pair of intervals, as ‘the black region(s)’, in reference to their figure 2). This conditioned probability as $\Pr(H_A - H_D > T_d \mid \text{endA_endD})$.

In equation 5, they use the joint probability of endA_endD , from equation 2, to remove the conditioning from $\Pr(H_A - H_D > T_d \mid \text{endA_endD})$, and thus obtain the probability of $H_A - H_D > T_d$. This probability is discontinuous, with the probability always being zero when $T_d > T_o$.

For example, when $\{r\} T_R = 0.97$ and $\{r\} T_d = 0.8$ (as in D&A’s analyses), and (to take a specific example), $\{r\} T_o = 0.9$, then the probability of $H_A - H_D > T_d$ is:

```
((T_o - T_d)^2) / (2*(T_R^2))
```

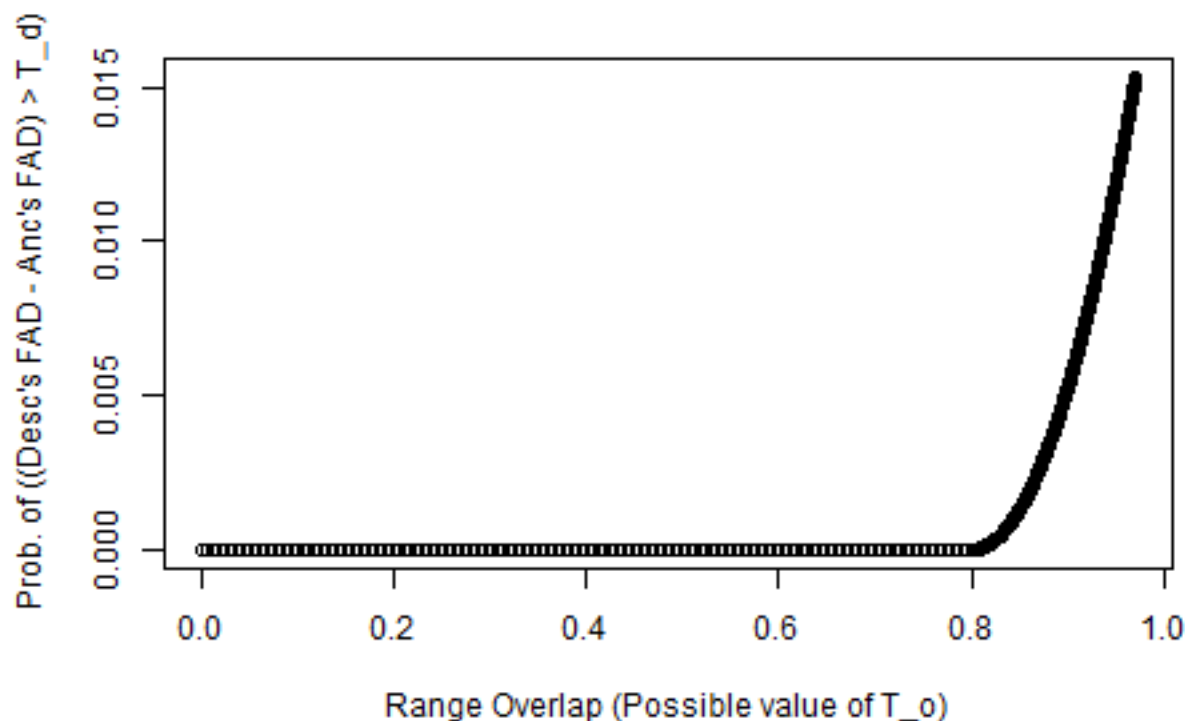
```
## [1] 0.04782655
```

Or, if $\{r\} T_R = 1$, $\{r\} T_d = 0.1$, and $\{r\} T_o = 0.5$, then the probability of $H_A - H_D > T_d$ is:

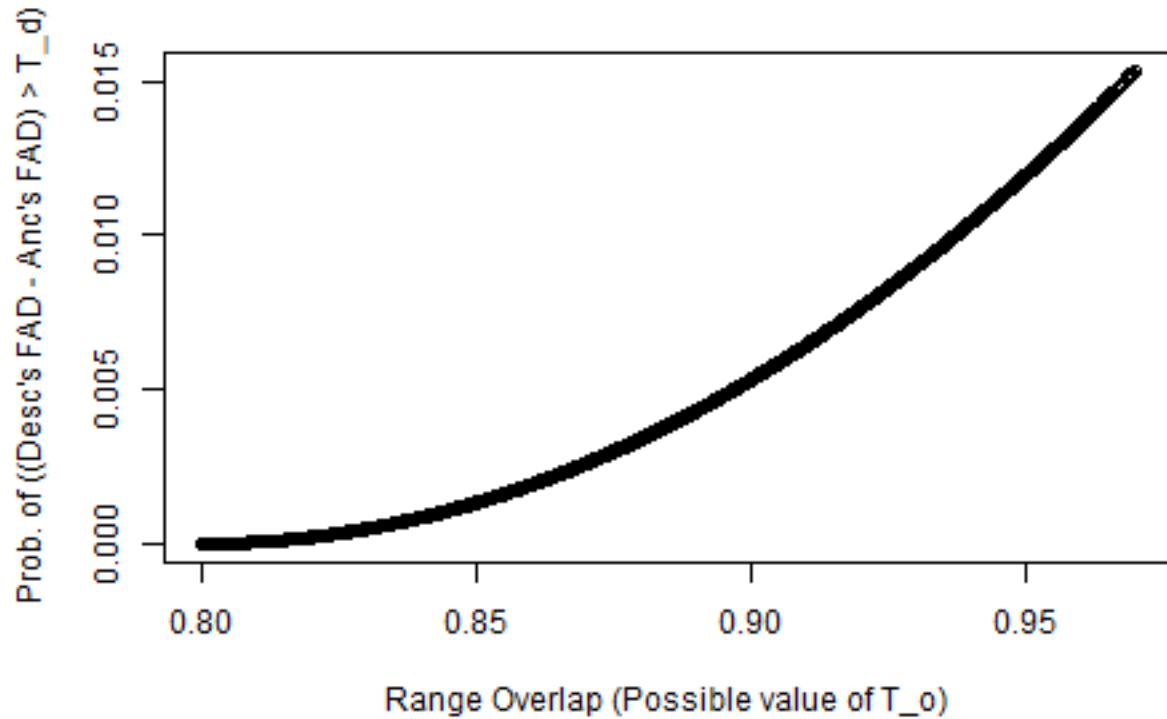
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((T_o - T_d)^2) / (2*(T_R^2))
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## [1] 0.04782655
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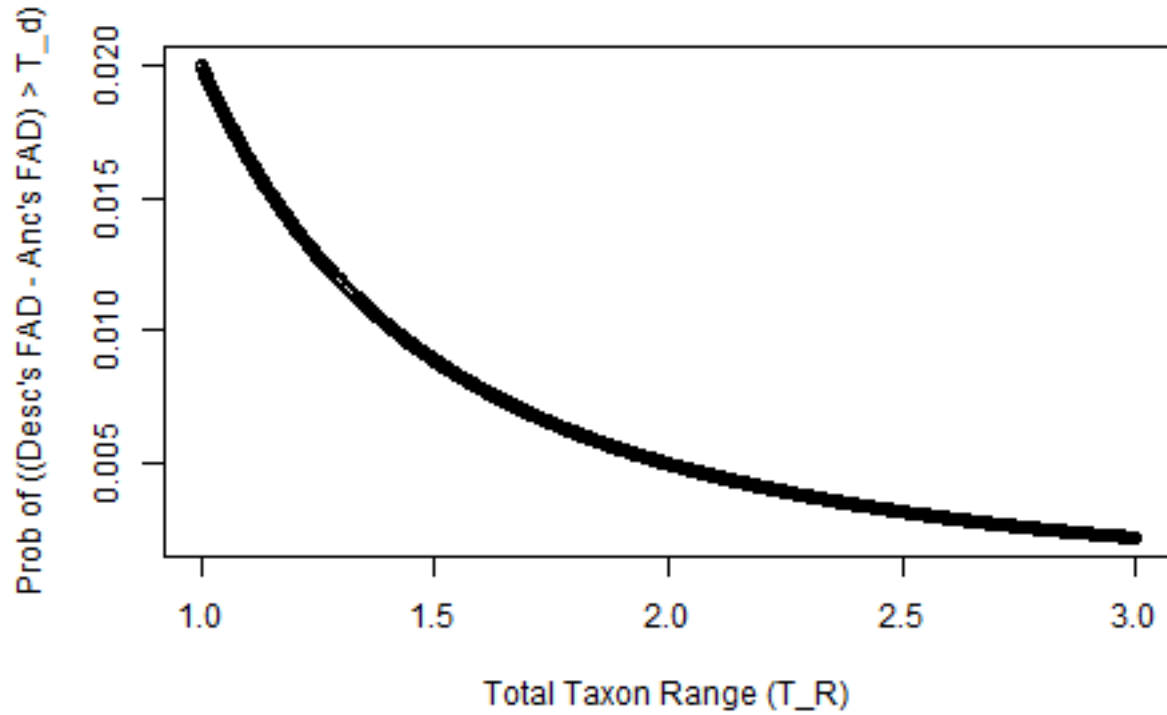
We can also use this equation to recreate their figure 3:



That was almost too easy. But note that most of this graph (just like D&A's Figure 3 - look at their horizontal axes) is at 0 probability, as the values of T_o include the range of 0 Ma to 0.8 Ma, even though $T_d = 0.8$ and thus its impossible for an ancestor to occur after its descendant. Its curious why this non-informative interval with *a priori* zero probability is even shown, so let's look at just the informative region.



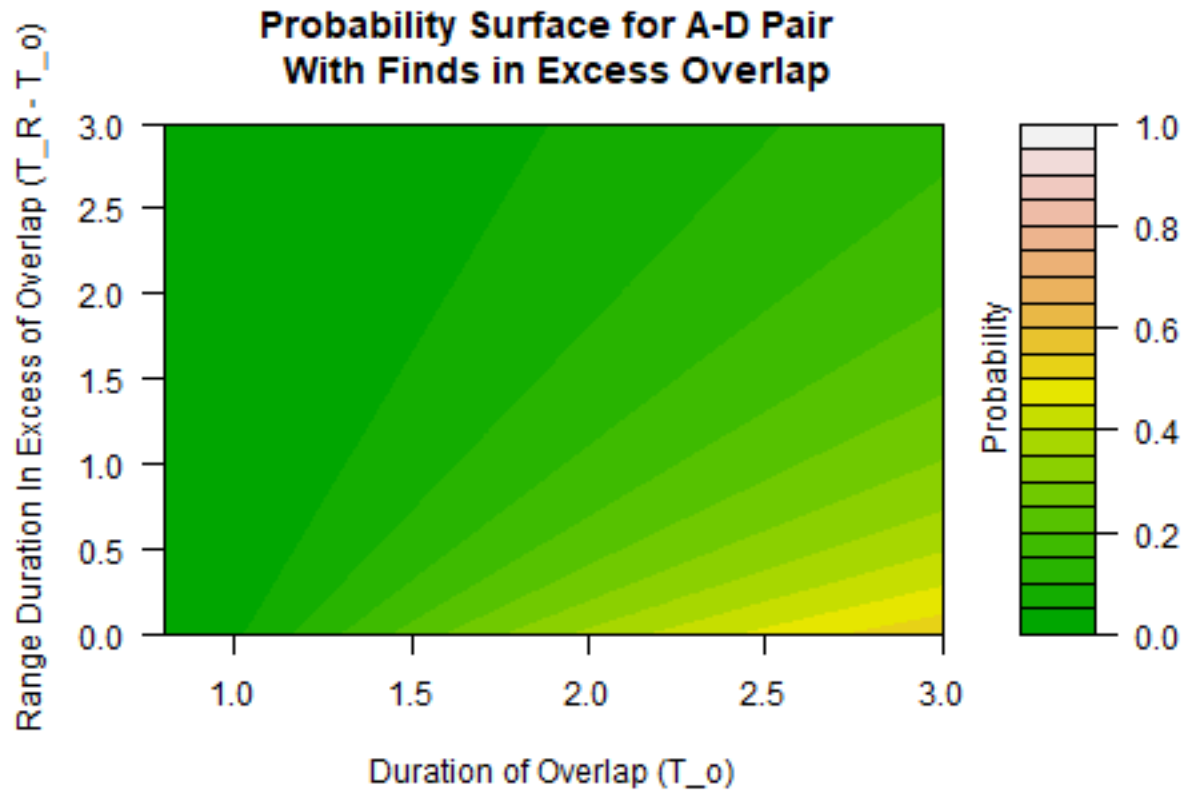
Now we can do things like allowing T_R to vary, instead of T_o - what if we assume the true overlap between *Homo* and *A. sediba* is just slightly more than the observed minimum overlap ($T_o = 1$, which would mean that *Homo* originated from *A. sediba* at most 20 Ka before *Homo* was first sampled in the fossil record), but we don't know how long the individual ranges of each taxon is, but otherwise put a hard cap on their maximum ranges at 3 million years?



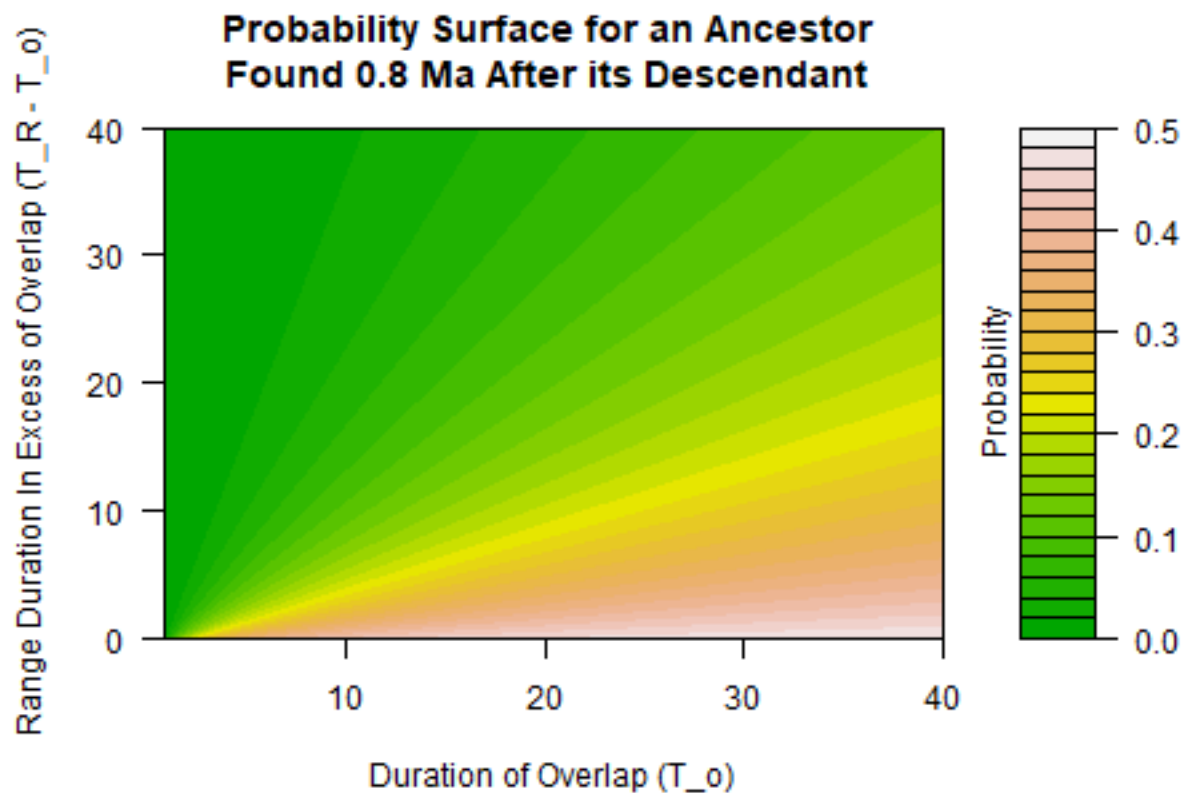
We can see that giving larger ranges relative to the overlap doesn't at all help the case for descendants to occur before their ancestors, because it means more geologic time during which only the ancestor existed, but wasn't sampled.

So, in figure 3 (or its recreation above), where only T_o is varied, we can see that the probability never gets better than 0.015 (1.5%), which leads us to wonder - what set of parameters would produce a high probability for the descendant to predate the ancestor? Presumably the hard cap on this probability is 0.5; this is theoretically true (in what scenario of uniform sampling could an ancestor be less likely to be found first than its descendant?), but also mathematically true as $\Pr(H_A - H_D > T_d \mid \text{endA_endD})$ is $1/2$, and thus the actual determining part of the model is the probability of fossil finds being in the 'end-caps' excess overlap' of the taxon ranges, given by equation 2 ($\Pr(\text{endA_endD})$).

So when is $\Pr(\text{endA_endD})$ closest to 1? Let's hold $T_d = 0.8$ constant, but vary T_o and T_R , and look at the values of $\Pr(\text{endA_endD})$ we obtain. Because $T_R > T_o > T_d$, we will vary T_o from the value of T_d to 3 Ma (extremely unrealistic for our case study, as it probably would require *A. sediba* to still be extant, but this is just an exploration - calm down!), and vary the amount of time in the fossil taxon ranges that is not part of their overlap (T_{R_no} , where $T_R - T_o = T_{R_no}$), with T_{R_no} ranging from 0 to (again) a maximum 3 Ma (which isn't just implausible, but patently impossible in our case as the fossil record in quest isn't from six million years ago...).



And now we can see that the probability of sampling from those regions of excess overlap is only going to be relatively high when (a) the ranges of the ancestor and the descendant are nearly perfectly overlapping ($T_{R_no} \sim 0$, or $T_o \sim T_R$), and (b) when the ranges are long relative to the discrepancy between the descendant and the ancestor. And, let's expand those maxima to 80 million years just to illustrate the point ($T_d * 50h0 = 40$) and look at a very similar plot but for the probability of observing a descendant predating its ancestor by a discrepancy as long as T_d , $\Pr(H_A - H_D > T_d)$. We'll rescale the color scale (probability) to a range of 0 to 0.5, as we know that 0.5 is the maximum limit in this case.



Thus, under the model presented by D&A and its set of assumptions, we can see that ancestors post-dating their descendants by even what might be small geological intervals (e.g. less than a million years) is only *likely* when geologic durations are long relative to the length of the discrepancy in the ages of appearance (at least 3-5 times T_d , the discrepancy), and the descendant's range overlaps for about a quarter of its range with its ancestor.

Now, there are hundreds, or possibly thousands, of putative ancestor-descendant relationships that exist in the paleontological literature. This agrees with our theoretical understanding of diversification and the incompleteness of the fossil record, which suggests that there should sampled ancestors, and that their number may not be great but should not be negligible (Foote, 1996). However, D&A's model implies we should rarely expect ancestors to occur after their descendants. But, are these reasonable assumptions? What does a realistic birth-death-sampling simulation tell us?

Simulation Tests

Simulation Structure, Parameters, Conditions

Here, I will attempt to compare D&A's model to a birth-death-sampling simulation, obtain 10,000 simulated fossil records that meet certain conditions (i.e. having more than one taxon), and parameterize these simulation to closely as possible match the parameters D&A worked with in their study. A birth-death-sampling model is simply a model of lineages, with events occurring as if a series of independent Poisson processes, with morphotaxa 'budding' off from each other at some rate (the rate of origination, or the *birth rate*), going extinct at some rate (the rate of extinction, or *death rate*), and being sampled at particular instances in time (the *sampling rate*). These processes are presumed to be constant with uniform rates, that do not vary over time. That's almost certainly false when humanity involved, but let's just ignore that for the sake of doing the

simulations at all. A birth-death-sampling model is also known as a BDS model, a birth-death-serial-sampling model, or (most commonly) a fossilized-birth-death model (Heath et al., 2014), and is exactly the sort of model now becoming increasingly and widely applied to paleontological datasets using tip-dating approaches in Bayesian phylogenetics.

Handily, the extinction rate is widely known to be the inverse of the mean *true* species duration. Note the ‘true’ - the observed species durations will always be smaller than the true ranges, due to incomplete sampling at both ends (and thus in better sampled records, more sampling means ranges that look more like the ‘true’ ranges.) However, we’ll ignore that effect of sampling and assume extinction rates is $1/0.97$, the inverse of the mean species duration used by D&A. (This literally means that if we had a hundred species at time A, we’d expect about half of them to be extinct 0.97 Ma later - so this is a **very** high extinction rate! The average for marine invertebrates is 0.1 per lineage, per Ma - implying an average extinction rate of 10 Ma.) Even more handily, we know that for most groups in the fossil record, the long-term origination and extinction rates are generally nearly equal (Stanley, 1976; Sepkoski, 1998; Marshall, 2016). We don’t really understand why that is (its partly a mathematical artifact, but also partly *not*, and doesn’t agree with estimates of speciation and extinction rates from molecular phylogenies), but we can probably safely assume that the origination rate for hominins has, in the long term, been pretty similar to the rate of extinction, given that only ones species has managed to survive to today. A per-lineage sampling rate isn’t really touched upon by D&A, as the parameter does not matter to their calculations, and I would hesitate to try to estimate such myself, so we can set that also equal to the extinction rate for now.

Birth-death-sampling simulations are notorious for having wildly variable results (e.g. most that start with a single species go extinct immediately), and thus we need to set constraints on what simulation output we’ll accept as a viable run or not (non-viable runs are thrown out, and the simulation is run again, until the required number of acceptable runs is produced). We will require at least 10 taxa sampled in each simulated fossil record (but no more than a 100), and at least 1 extant taxon in each output run. Most runs will likely have dozens of sampled species, even with these seemingly low limits.

We will also not treat simply surviving to time 0 as ‘being sampled’ - that way, when we look at differences in sampling times, its always fossil finds, modeled as under a uniform Poisson process - exactly as expected by D&A’s own model.

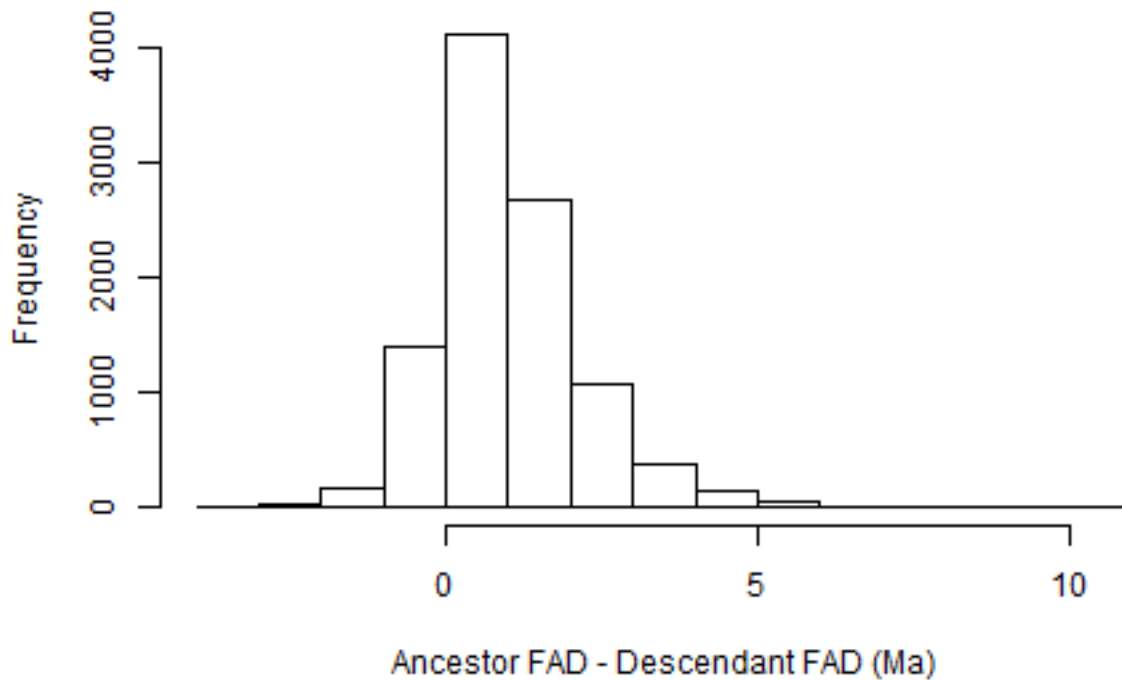
Simulation Patterns

Once we run the simulations as described above, we can explore our 10,000 simulated fossil records. In our case, there are two central variables of interest that we can look at as summary statistics, and examine the relationships between them, to evaluate how well D&A’s model holds up in these more realistic (yet still simulated) scenarios.

The first of these variables of interest is The difference in age between the first sampled appearance of the ancestor and the first sampled appearance of the descendant, for every sampled pair of ancestor-descendant taxa. For all those pairs in which descendants predate their ancestor, there will be a negative temporal offset, which is the temporal difference between descendant and ancestor labeled T_d by D&A. Note that this is not limited to only direct ancestor-descendants, but includes ancestor-descendants where there are unsampled intermediates as well. In order to properly avoid statistical artifacts due to accidentally including data from two separate pairs involving the same ancestor, or artifacts brought on by ignoring taxa with more than one descendant, only a single such ancestor-descendant pair was sampled from each tree. This sampling was doing at random from all possible ancestor-descendant trees in a given simulated fossil record.

The distribution of age offsets looks, in same ways, what we would expect: dominated by cases of the ancestor preceding the descendant (positive values):

Difference in Age Between Ancestor and Descendant



Note however that a significant portion are less than zero, and thus represent ancestor-descendant discrepancy - cases where the ancestor's first appearance is younger than the descendant. How many, in fact:

```
sum(AD_diff_all>0)/length(AD_diff_all)
```

```
## [1] 0.8414
```

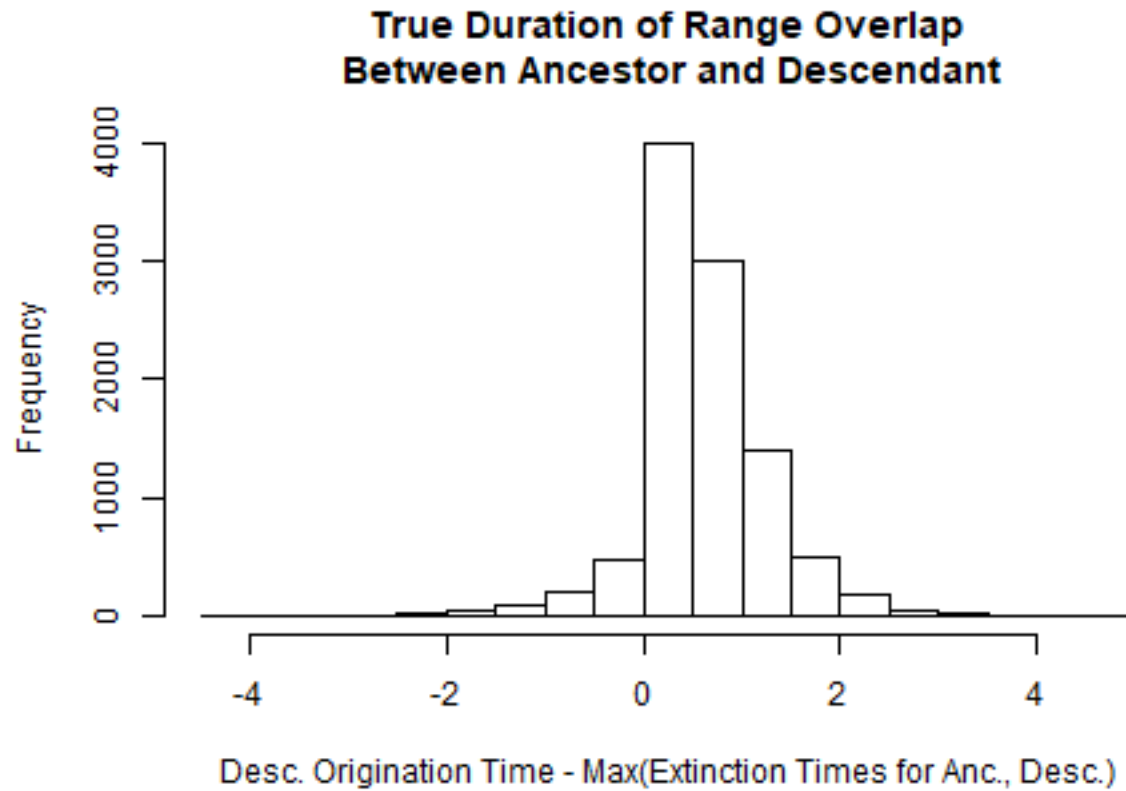
One thought that might occur is that there might be different patterns for ancestor-descendant relationships where the ancestor or descendant are still extant. (The mathematical rules of entirely extinct taxa in birth-death-sampling models are actually usually a lot more tractable, as extant taxa forces us to consider the wonderfully complicated world of censored and truncated exponential distributions). We can record that information and look at the proportion of our returned offsets that came from those that either had extant members, as opposed to being entirely extinct:

```
sum(isExtant_all)/length(AD_diff_all)
```

```
## [1] 0.2441
```

As we go forward, we will make sure to compare the entire simulation output with what we obtain from the all-extinct and still-extant sets, to make sure there is no divergence from the whole dataset.

The second variable of interest is the amount of true overlap between the complete durations of the ancestor and the descendant (T_o in D&A's model), for each of the randomly-selected observed ancestor-descendant pairs we obtained temporal offsets from above. This is quite easy to calculate - note that in many cases the amount of overlap is small (and in the case of indirect ancestor-descendant relationships, may be non-existent, leading to the seemingly negative overlap values, as the two sampled taxa never were co-extant).



As with the temporal offset values, we may find that pairs with extant members or not might differ.

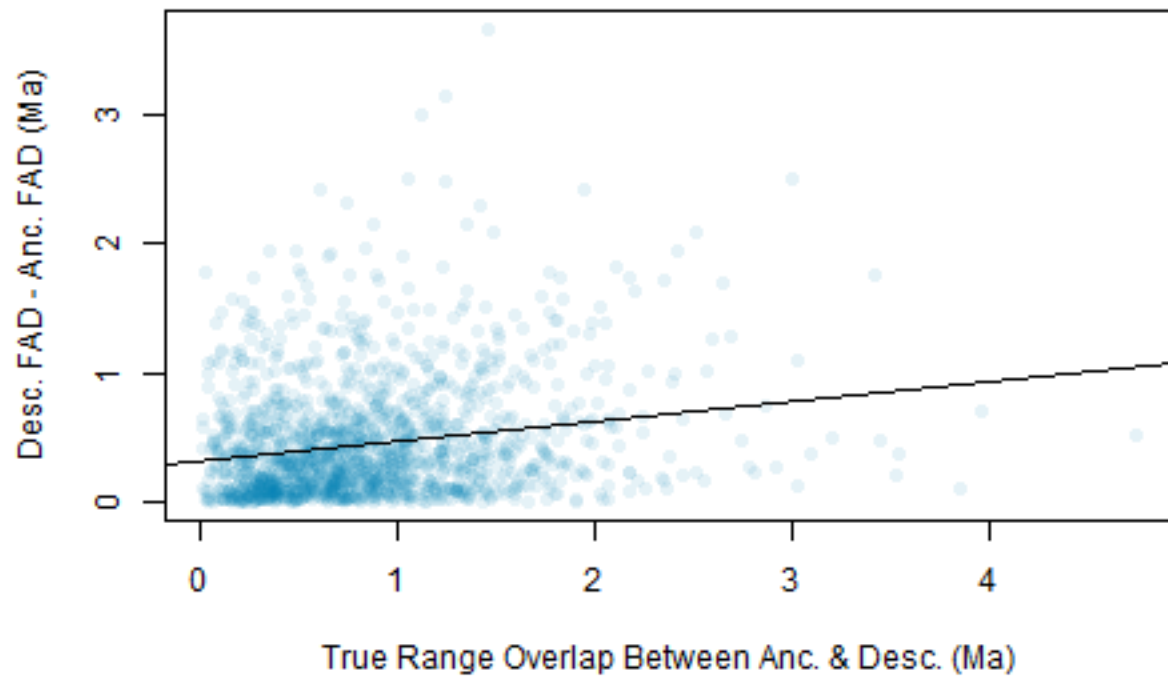
Comparing Simulation Results to Du and Alemseged's Model Expectations

As discussed above, one of the central expectations of D&A's model is that the magnitude of ancestor-descendant age discrepancy will depend greatly on the amount of true overlap between ancestors and descendants. In either case, we will need to filter the data.

For example, we could filter on those pairs that have discrepancies (the pairs where the descendant is sampled first):

```
## Warning: In lm.fit(x, y, offset = offset, singular.ok = singular.ok, ...) :
## extra argument 'lty' will be disregarded
```

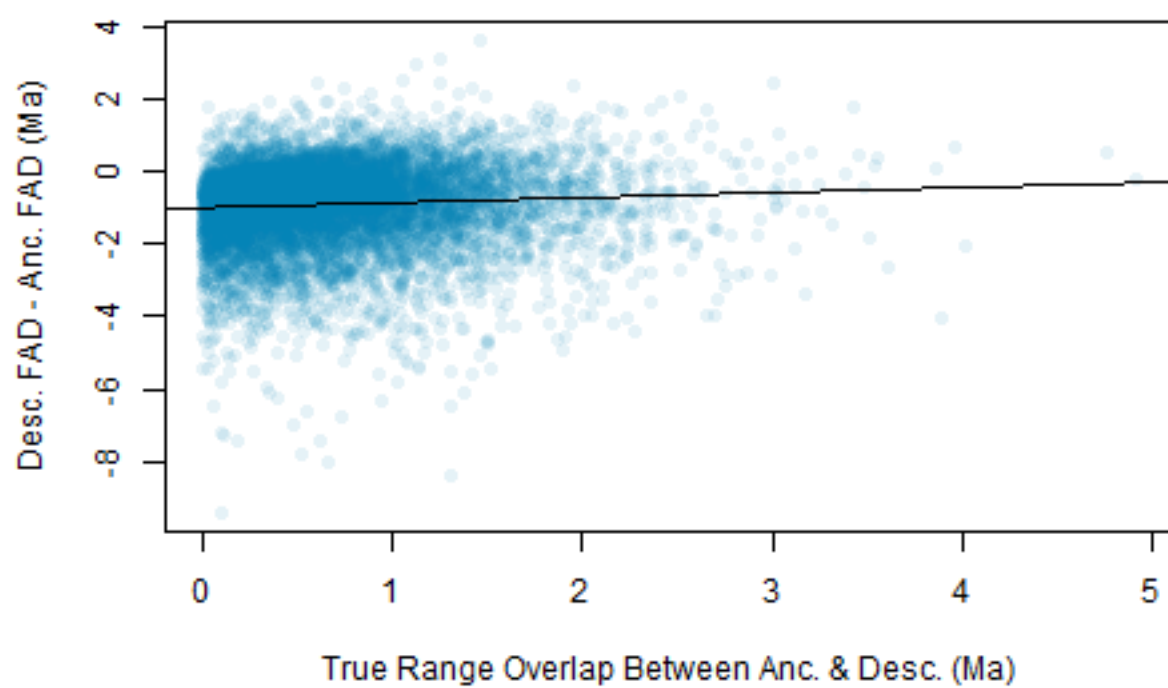
Temporal Offset For Pairs Where Desc. Appears First

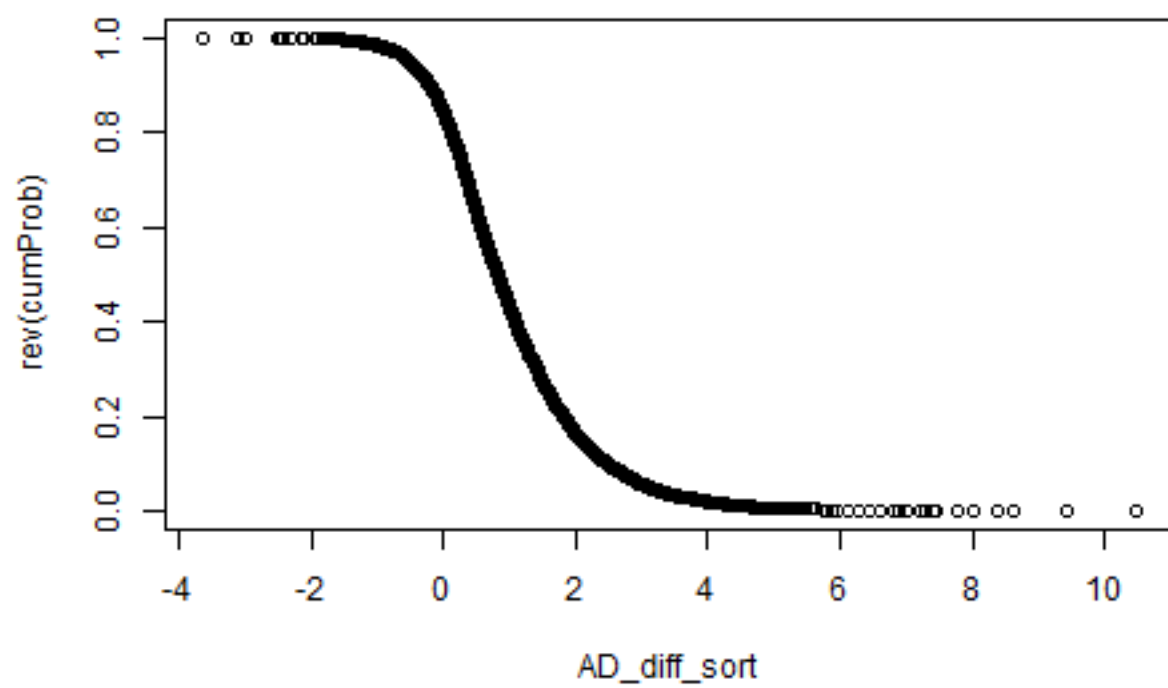


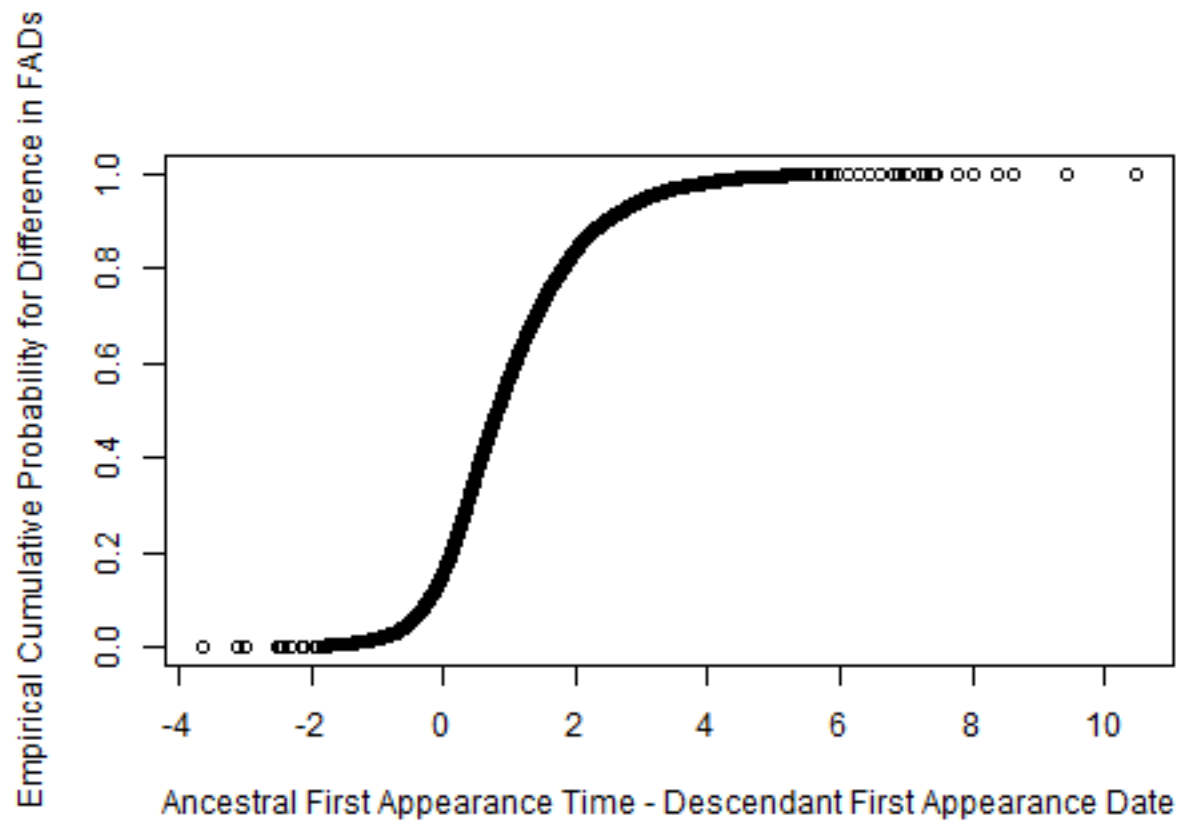
The dashed line here represents the trend line of a linear regression (not that the relationship between these two variables is very linear). Note that we are here looking at the difference between the descendant

Or, we could examine the temporal offset for the pairs (again, negated) where there is actual overlap between the ancestor and the descendant:

Temporal Offset For Pairs with Range Overlap







```
## [1] 0.1586
## [1] 0.0182
## [1] 0.0014
```