### **June 6, 2016 Notes**

- Discovered and fixed two bugs involving an uninitialized selection vector and Mphi values
- Said bug fixes resolved the issue of the program crashing around iteration 2500
- Despite the fixed bugs resolving that issue, log likelihood values still plummet to values eventually reaching 10e-23 which causes the not a number error, causing the MCMC algorithm to halt. <sup>1 2</sup>
- Due to Newton failing to execute my job request, I had to run the R script on my laptop so I was only able to run it once so more testing is needed but for the meantime it appears it takes until around iteration 42000 for the program to crash. 3 4

### TODO:

Work on psuedocode Remove redundancies Determine if Sphi prior is being used Improve variabbe names and documentation Document the R scripts

<sup>&</sup>lt;sup>1</sup>mikeg: June 10 2016 –Is this a typo? Usually it is  $\exp -300$  that starts to push the limit of numerical precision. Further, it is unclear to me why this would cause a problem unless you are later converting this number back to a log scale. Is it necessary to exponentiate and then take the log? ANSWER: 10E-23 was user defined.

<sup>&</sup>lt;sup>2</sup>mikeg: June 10 2016 –This is confusing given the previous statement. It sounds like you didn't really resolve the underlying issue. ANSWER: Problem is now fixed in that the code run, but it seems likely that the bug is still there.

<sup>&</sup>lt;sup>3</sup>mikeg: June 10 2016 – why is newton failing to execute your job requests? How and when will you solve this issue?

<sup>&</sup>lt;sup>4</sup>aland: June 27 2016 – The problem comes from the .SGE scripts. Jeremy Rogers had them tailored to how his directories were set up and where he wanted files to go. Fixing this is pretty low on my priority list since Gauley handles pretty much everything I need to run.

### June 7, 2016

- Removed exit statements from FONSEParameter.cpp, PANSEParameter.cpp, and RFPParameter.cpp in order to eliminate warnings when running the check in R
- Downloaded all of the necessary files to my Gauley account.
- Running FONSE with simulated data and b = 0.001 led to "nan" being reached much more quickly than on my laptop, around iteration 5500. <sup>1</sup>
- Said issue may be a cause of R not working correctly on Gauley as compared to my laptop. Multiple errors and missing packages were encountered. These errors made it impossible to check any plots.
- mikeg: If it strange you're running into such errors since gauley is used to develop the code base. Are they really errors or, since you execute code, warnings? What are the error codes? When discussing errors, it is good to always include some output using the 'quote' or similar environment in LaTeX. Do other people run into these same errors?
- Two new Items for the TODO list: Continue troubleshooting R on Gauley and alter Jeremy's .sge scripts so that they no longer email him when I run a job on Newton.

<sup>&</sup>lt;sup>1</sup>mikeg: June 10 2016 –is this based on one run on each machine? It's more useful to be more quantitative and descriptive

### June 8, 2016

- Changed the .sge scripts for Newton so that Jeremy no longer receives an email upon the completion of a job.
- Changed a function in FONSEModel.cpp to match it's corresponding function in ROCModel.cpp. In particular, I changed a long series of divisions to on division and a series of multiplications because multiplication is a faster operation. <sup>1</sup>
- Fixed a typo in the documentation of ROCModel.cpp
- Worked out some kink experienced while running R scripts on Gauley. In particular, finally made it so that the plots actually showed up and worked.
- Formulated the hypothesis that the reason we are experiencing a drop in log likelihood comes from a bug involving either deltaM, deltaOmega, or phi after observing the trace plots. <sup>2</sup>

<sup>&</sup>lt;sup>1</sup>mikeg: June 10 2016 –include the function name. Any information on speed up?

<sup>&</sup>lt;sup>2</sup>mikeg: June 10 2016 –Include plots in your notes, especially if you refer to them, and explain more clearly why you think its behavior indicates this.

## June 13, 2016

- Fixed an error in FONSEModel.cpp involving a couple of missing semicolons
- Fixed an error in the .tex file for the notes where using the hyperref package kept the notes from compiling
- After running FONSE again, with simulated data and b = 0.001, it seems the errors in log likelihood are no longer deterministic. Values still plummet, and the traces suggest that both mutation and selection values may be the cause. Further testing and observation is required. <sup>1</sup>
- Found a difference between FONSEModel and ROCModel where ROC would take mutation prior into account and FONSE wouldn't. This might part of the issues causing log likelihood values to plummet, but testing is required.
- Uploaded both the new .tex file and its corresponding pdf just in case there's another error that doesn't keep me from compiling on my machine but might keep another from compiling on a different machine.

<sup>&</sup>lt;sup>1</sup>mikeg: June 16, 2016 - If you run the code in serial and set the random number seed, does the behavior become repeatable?

<sup>&</sup>lt;sup>2</sup>mikeg: June 16, 2016 - In general, you should refrain from making copies of files (e.g. labNotes → Lab\_Notes). Instead, rely on the fact that you are using version control and can always revert back or merge with an earlier version of a file.

## June 14, 2016

- The addition of the mutation prior into FONSE seems to have stopped the rapid descent of the Log Likelihood values. After running it once with 1000 samples and 1 thining, the log likelihood stayed somewhat stable after starting low and rising to about -20000. The vaules were also similar when running with 1000 samples and 10 thining. More extensive testing is required to ensure that the plummeting problem is finally resolved. <sup>1</sup>
- Although the log likelihood values are starting to come together, there is an issue observed in both the mutation and selection traces where certain codons are not being accepted.
- Conducted further comparisons on FONSE and ROC to remove redundancies and semantic oversight.

<sup>&</sup>lt;sup>1</sup>mikeg: June 16, 2016 - I am glad that the prior seems to have fixed this problem. I don't understand, however, why it did so. Were the  $\Delta M$  values diverging from the true value substantially? Note also that one can relax the effect of the prior by making its  $\sigma_{\Delta} M$  value large. In fact at the  $\lim_{\sigma_{\Delta} M \to \infty}$  the system essentially behaves as if it has a flat prior (which is proportional to 'no prior').

## June 15, 2016

- After running FONSE overnight, I feel it's safe to conclude that the plummeting log likelihood problem is resolved although there have been issues with multiple, sometimes all but a few, codons with acceptance rates of 0 as backed up by both the output and the trace plots of both mutation and selection, although this is again non deterministic. Running it on Gauley produced far more codons with no impact than when I ran it on my laptop so the non determinism may come from memory or operating system differences. <sup>1</sup>
- There was a problem with Gauley involving the machine randomly restarting without being prompted. It doesn't seem like a major problem and the effects seem to be confined to the processes being run on it at the time being prematurely terminated. I'll be keeping an eye for any unespected behavior from Gauley in case the problem ends up being reoccuring.
- Studied the model to try and deduce how the simple inclusion of the mutation priors resolved the plummeting problem. So far it still remains fairly unclear and more studying is necessary.

<sup>&</sup>lt;sup>1</sup>mikeg: June 16, 2016 - This shouldn't happen since we are using an MCMC whose step sizes adapt based on the acceptance ratio of the previous samples. (Of course, it would be good to make sure this adaptive option is set.) We saw similar 'non-acceptance' behavior with Gabe and/or Jeremy in the past month or two. It had to deal with the parameters not being correctly referenced or defined. Cedric will remember the details.

### June 16, 2016

- Ran FONSE with b = 0.001, samples = 1000, and Thinning = 10 3 separate times. <sup>1</sup>
- Run 1<sup>2</sup>
  - Log likelihood settled around -9600
  - AA's L and R had acceptance rates of 0
  - stdDevSynthesisRate posterior estimate settled around 1.4, .05 for the proposal width
  - Trace of the expected values of phi goes to 0.
  - All codons for L in the mutation traces stop changing and fall out around sample 100 <sup>3</sup>
  - R falls out around sample 300
  - It's the same story for both L and R in the selection traces except it takes about half as many samples for both to fall out

### • Run 2 4

- Log likelihood settled around -14000
- AA's L and R still had acceptance rates of 0<sup>5</sup>
- stdDevSynthesisRate posterior estimate settles around 1.9, .045 for the proposal width
- Trace of the log likelihood didn't level off as much. It seemed to still be rising.
- expected phi trace was the same as Run 1
- L falls out around the same place in Mutation as Run 1; R falls out twice as fast.
- Both fall out twice as fast in the selection traces compared to Run 1.

### • Run 3

- Log likelihood settled around -20000
- AA's L and R still had acceptance rates of 0
- stdDevSynthesisRate posterior estimate settled around 2.5, .04 for the proposal width
- There was a more pronounced level-off in the log likelihood trace, much like Run 1
- expected phi trace leveled off but slightly above 0
- Both L and R had mutation and selection traces similar to Run 1
- Also worked on documenting FONSE by taking documentation from similar parts in ROC and making the necessary alterations before putting it into FONSE

<sup>&</sup>lt;sup>1</sup>mikeg: June 18, 2016- Where all three of these runs carried out with the same set of genes? If not, then LLik(data||parmeters) surfaces, although similiar, will differ between the runs. '

<sup>&</sup>lt;sup>2</sup>mikeg: June 18, 2016 - Per my previous request, please include the relevant figures. I recommend organizing figures by putting them in a separate folders such as "Figures/2016/June/16". You should also name them in a consistent manner such as "FONSE\_run\_1\_trace\_LLik.png"

<sup>&</sup>lt;sup>3</sup>mikeg: June 18, 2016 - what does 'fall out' mean?

<sup>&</sup>lt;sup>4</sup>mikeg: June 18, 2016 - again, figures should be included to help illustrate and document the behavior you describe.

<sup>&</sup>lt;sup>5</sup>mikeg: June 18, 2016 - Have you described this behavior to Cedric? We've seen similar behavior before.

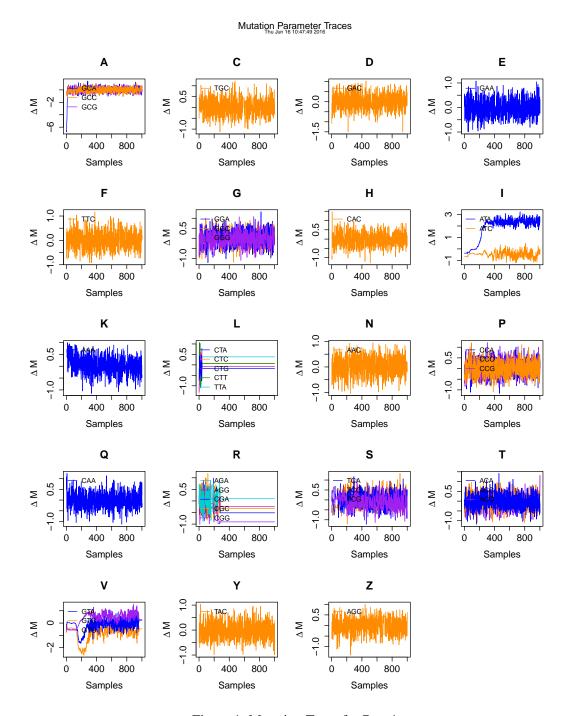


Figure 1: Mutation Trace for Run 1

### С D Α Ε 900 GCA GCC GCG TGC Δη 300 4 ۲۷ Δη 800 400 800 400 400 800 400 800 0 0 0 Samples Samples Samples Samples F G Н I CAC GGA 200 Δη η ηΔ ۲۷ -300 -0.02 400 800 400 800 400 800 400 800 0 Samples Samples Samples Samples Κ L Ν Ρ 150 100 CTA CTC CTG CTT η -10 η۷ η۷ 40 20 -30 0 800 400 800 400 800 400 800 400 0 Samples Samples Samples Samples T Q R S AGA AGG CGA -50 μV 9 Δ CGC CGG -25 -250 -10 400 400 800 800 400 800 800 Ó 400 Samples Samples Samples Samples ٧ Υ Z 250 GTA GTC GTG AGC -50 ηΔ Δη -15 Δη 100 -150-30 0 800 400 400 800 400 800 0 Samples Samples Samples

Selection Parameter Traces

Figure 2: Selection Trace for Run 1

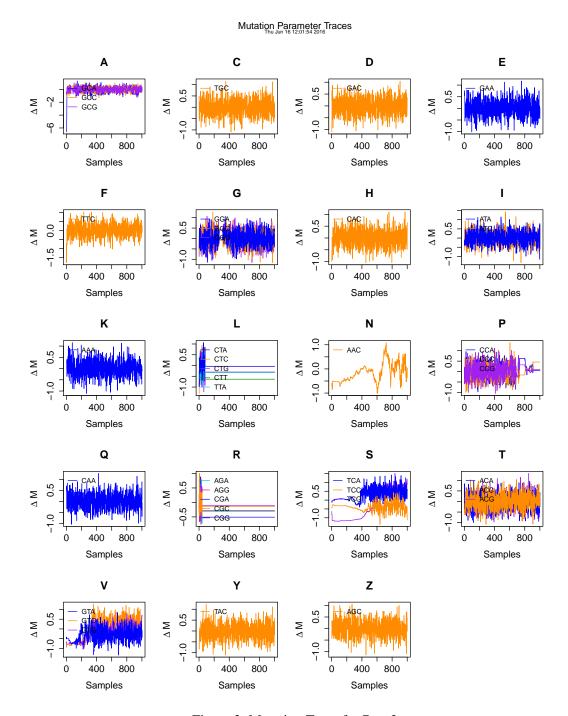


Figure 3: Mutation Trace for Run 2

### D Α С Ε 40 -100 GCA GCC TGC 20 Δη -30 -500 0 -60 800 400 800 400 800 400 800 400 0 0 0 Samples Samples Samples Samples F G Н I 80 -50 -10 ηΔ ۲Δ η۷ ηΔ 9 -250-30 0 400 800 400 800 400 800 400 800 Samples Samples Samples Samples Κ L Ν Ρ CTA CTC CTG -1.0 -100 η η۷ η۷ -250 -3.0 -20 400 800 800 400 800 400 800 400 0 Samples Samples Samples Samples Q R S T AGA AGG CGA CGC CGG TCA TCC TCG -100 -0.5 μV Δ -300 -1.5 400 ó 800 400 400 800 800 400 800 Samples Samples Samples Samples Υ ٧ Z 80 GTA GTC GTG ۲۷ Δη 9 0 -80 800 400 400 800 400 800 0 Samples Samples Samples

Selection Parameter Traces

Figure 4: Selection Trace for Run 2

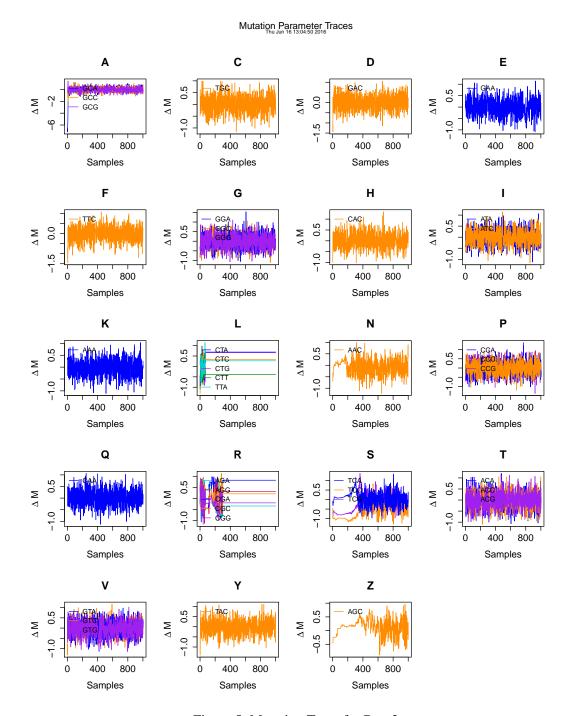


Figure 5: Mutation Trace for Run 3

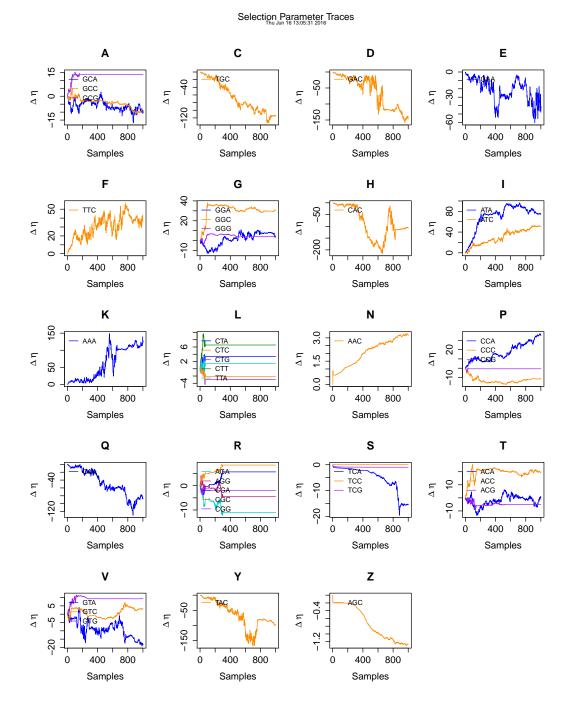


Figure 6: Selection Trace for Run 3

## June 21, 2016

• Most of today was spent resolving a problem with RStudio crashing. Both Cedric and I were experiencing this problem so we troubleshooted the issue and discovered that the issue was in a change that had been made in the R-only section of the code in Parameter.cpp. The issue was a "+1" had been added while referencing an element of an array, causing the program to attempt to reference out of the bounds of the array causing a segmentation fault.

# June 22, 2016

• Added documentation to the FONSE R-scripts from home due to illness.

## June 23, 2016

### • TODO:

- Alter the code so that there is flexibility when someone invokes a prior i.e. specify whether or not a prior should be used.
- run the program with a uniform prior since it is proportional to no prior.
- Change from using a covariance matrix to the diagonal to see how that changes the issue where codons have acceptance rates of 0.
- Alternatively update the covariance matrix instead.
- Check to see if taking the mutation prior out of ROC causes the same phenomenon.
- Dump out the values of the genome step of MCMC so we can observe the traces per amino acid.
- Put in traces from when the code crashes into the lab notes.

### June 24, 2016

- As requested, I removed the prior from FONSE and ran the model twice in order to get more data and plots showing what's going wrong with the algorithm. <sup>1</sup> <sup>2</sup>
- Run 1: b = 0.001, samples = 1000, thinning = 1.
  - Log Likelihood values plummet once again. At the end of the MCMC loop the value had reached
    -1.8e-40. See Figure 1.<sup>3</sup>
  - Acceptance rates for most of the codons were highly suspect with most having acceptance rates of 1 and others having rates of 0.
  - Both mutation and selection traces seem consistent with the behavior of the other plots and parameters. See Figures 2 and 3.
- Run 2: b = 0.001, samples = 1000, thinning = 10. <sup>4</sup>
  - Log Likelihood values plummet once again. At the end of the MCMC loop the value had reached
    -7.1e-41. See Figure 4.
  - Acceptance rates for most of the codons were agian highly suspect with most having acceptance rates of 1 and others having rates of 0.
  - Both mutation and selection traces again seem consistent with the behavior of the other plots and parameters. See Figures 5 and 6.

<sup>1</sup>mikeg: 06/24/16 –

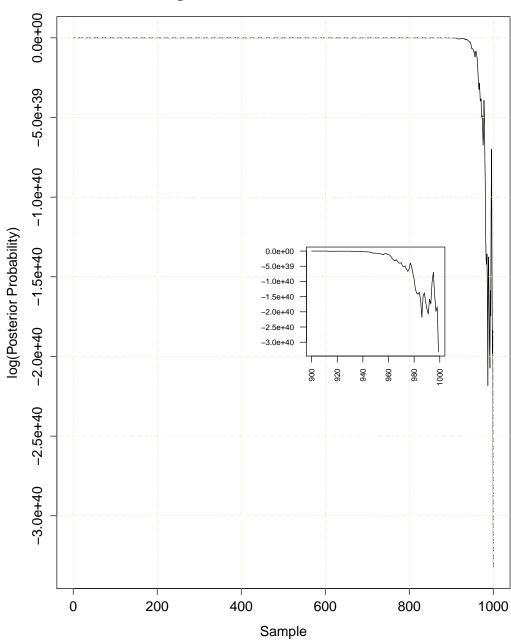
- Is this all you were able to get done today? If not, your notes should reflect this and, in general, be more detailed.
- What did the ROC runs show? If you were unable to run it, why is this?
- Your choice of file names for plots is good and organizing them by date is also good. However, you should also incorporate the year into your folder structure (e.g. Fig/2016/06/24).
- It appears that  $\Delta \eta$  also goes to unrealistic values, not just  $\Delta M$ .
- Please add the  $\sigma_{\phi}$  trace.
- There may be information in the earlier steps of the traces that we can't observe because the final plots cover such large ranges. Please generate and add plots that focus in on the earlier steps. We want to know if the code behaves normally at any point or if it is always climbing downwards.
- Where are the traces for the individual AA per our discussion yesterday?
- As requested, please respond to my previous notes created using \footnote{}. I'm particularly concerned with your inability to run things on newton.

<sup>2</sup>aland: 06/27/16 –

- Documentation was done while FONSE was running
- I hadn't been able to run ROC at this time. Cedric has since helped me and I'm now able to do so.
- I'm still in the process of implementing the traces of individual AA's. Both that and including plots that focus on earlier steps are on my TODO list.

<sup>&</sup>lt;sup>3</sup>mikeg: 06/24/16 – You should use the \label{} and \ref{} commands rather than labeling and referring to figures using hard coded numbers.

<sup>&</sup>lt;sup>4</sup>mikeg: 06/24/16 – instead of cutting and pasting the exact same text, you should clearly state what you want to reader to infer from this second set. That is, "the choice of thinning does not affect the model's behavior." I would not have expected it to and, instead, would have recommended you do a run with a different *b* value.



logPP: -4.15013321272666e+39

Figure 1: LogLikelihood trace for Run 1

### Α С D Ε GCA GCC GCG GAC TGC GAA Δ Σ Σ Δ -10000 -30000 -5e+11 -15000 400 800 400 800 400 800 0 400 800 0 0 0 Samples Samples Samples Samples F G Н I 1e+08 GGA GGC GGG ATA ATC CAC 10000 Δ ∑ ⊘ Δ Δ -5e+07 -20000 0 400 800 400 800 400 800 400 800 0 Samples Samples Samples Samples L Ρ Κ Ν 4e+19 6e+11 CTA CTC CTG CTT TTA CCA CCC CCG Δ Δ Δ Δ 0 -4e+19 -6e+11 က 400 800 800 400 800 400 400 800 0 Samples Samples Samples Samples Q R S T -4e+11 4e+11 AGA AGG CGA CGC CGG TCA TCC TCG ACA ACC ACG CAA 0.5 Σ Σ Δ Δ -4e+19 -30000 -0.5 400 400 800 400 800 800 800 400 Samples Samples Samples Samples ٧ Υ Z -5000 GTA GTC -1.5 ΔΔ Δ Δ -5e+11 -35000 -3.5 400 800 400 800 400 800 Samples Samples Samples

Mutation Parameter Traces

Figure 2: Mutation trace for Run 1

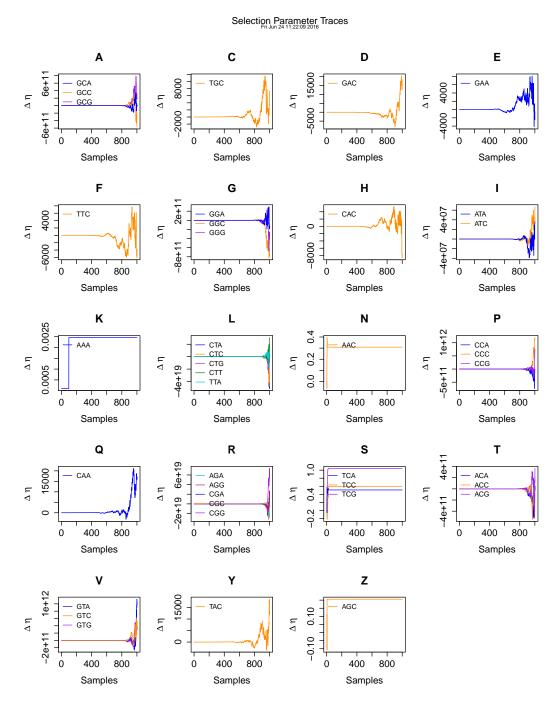
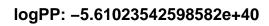


Figure 3: Selection trace for Run 1



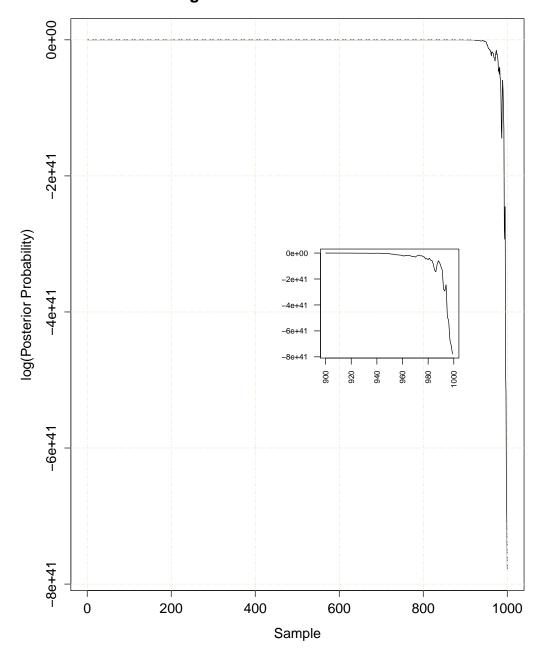


Figure 4: LogLikelihood trace for Run 2

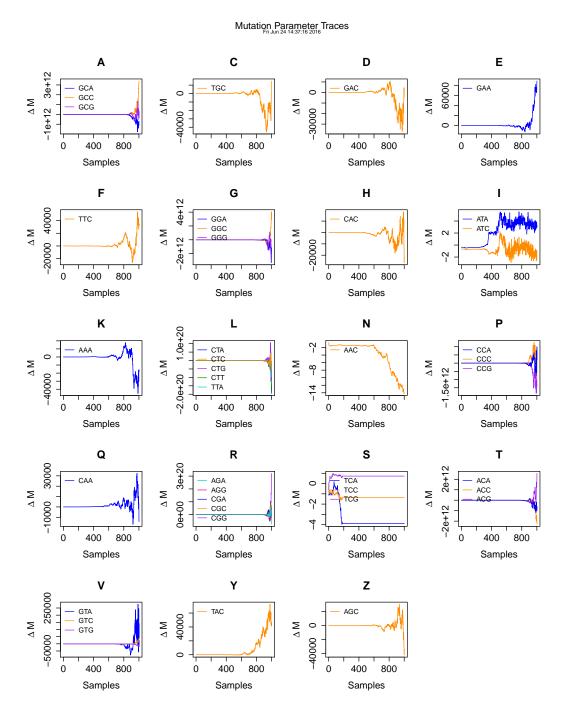


Figure 5: Mutation trace for Run 2

### Selection Parameter Traces С Α D Ε 1e+12 30000 GCA GCC GCG TGC GAC GAA η 0 Δ Δη -30000 -3e+12-60000 -20000 800 400 800 Ö 400 0 400 800 400 800 0 0 Samples Samples Samples Samples F G Н I -10000 30000 -0.010 0.010 -20000 60000 GGA GGC GGG TTC CAC ηΔ ۲Δ Δη Δη -1e+12 400 800 400 800 400 800 400 800 Samples Samples Samples Samples Κ L Ν Ρ 5.0e+19 30000 2.0 -5e+12 0e+00 CTA CTC CTG CTT TTA CCA CCC CCG AAA AAC 1.0 Δη ۲۷ Δη ηΔ -2.5e+20 -100000.0 400 800 400 800 400 800 800 0 400 0 Samples Samples Samples Samples S Т Q R -1.5e+20 1.5e+20 1e+12 AGA AGG CGA CGC CGG ACA ACC ACG CAA TCA TCC TCG -2.0 -0.5 η η Λ۵ Δ -30000 0 -4e+12 400 ó 400 800 800 800 400 800 400 Samples Samples Samples Samples ٧ Υ Z 40000 GTA GTC GTG AGC 1000 -20000 0 ηΔ ۲۷ Δη -20000 0 800 400 400 800 400 800 0

Figure 6: Selection trace for Run 2

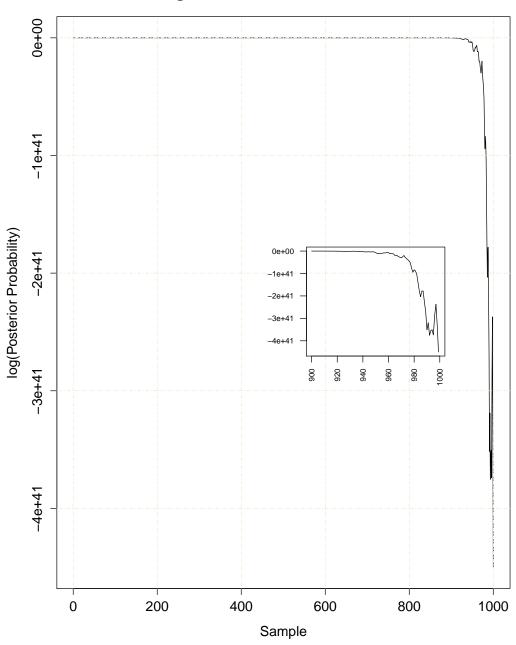
Samples

Samples

Samples

## June 27, 2016

- Worked on the implementation of adding plots of the earlier stages of the trace for the log likelihood.
- Worked on altering the .SGE scripts so I can run them on Newton
- Added further documentation to FONSE
- Ran FONSE and ROC both without the prior, both with samples = 1000 and thinning = 10.
- FONSE: b = 0.001
  - Log likelihood values plummeted once again, this time ending at -3.31e+41.
  - High acceptance rates for most of the AA's with over half of them being 1. Those that weren't high or 1 were 0.
- ROC with simulated data:
  - Log likelihood values quickly rose and settled around -810000
  - Very low acceptance rates for almost all AA's (some being moderate at around .35) with almost half being 0.



logPP: -5.78890248527369e+40

Figure 1: LogLikelihood trace for FONSE

#### logPP: -810535.952010936

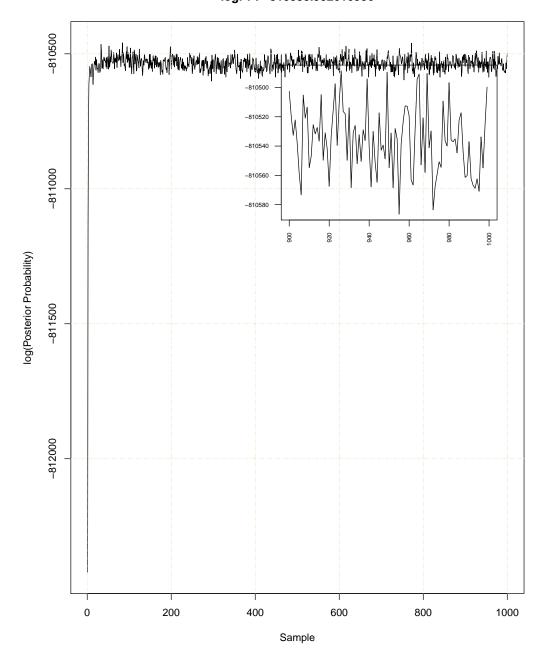


Figure 2: LogLikelihood trace for ROC

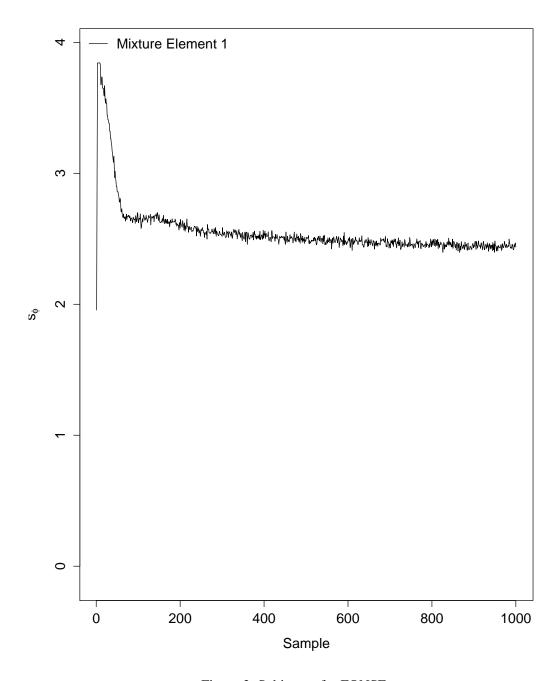


Figure 3: Sphi trace for FONSE

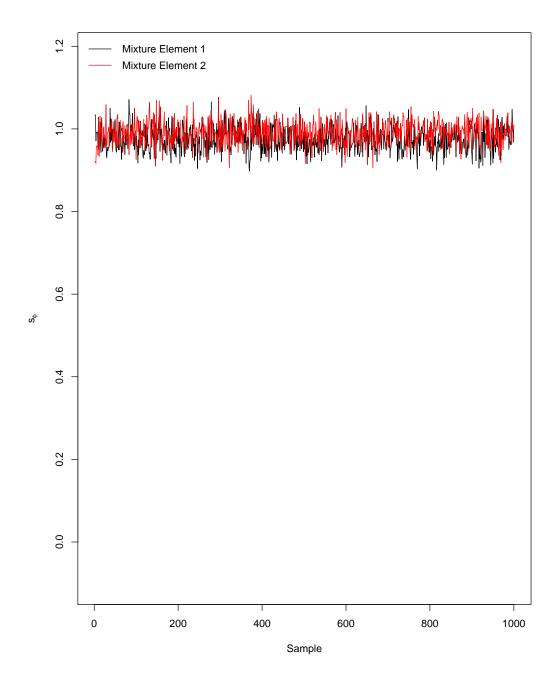


Figure 4: Sphi trace for ROC

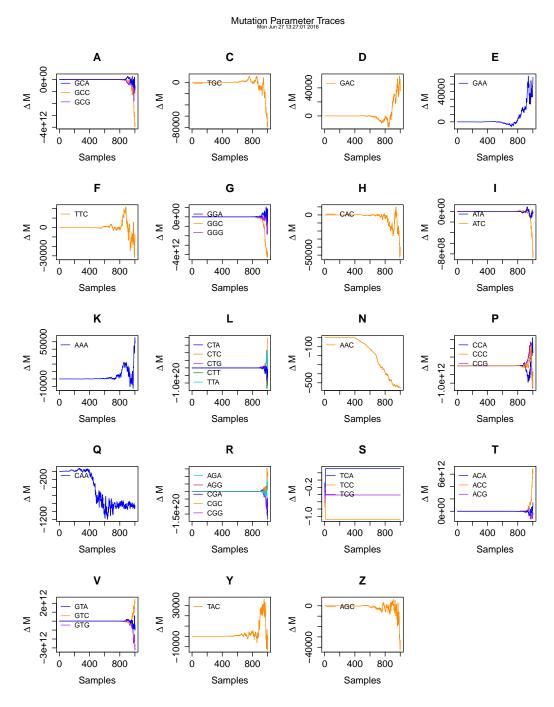


Figure 5: Mutation trace for FONSE

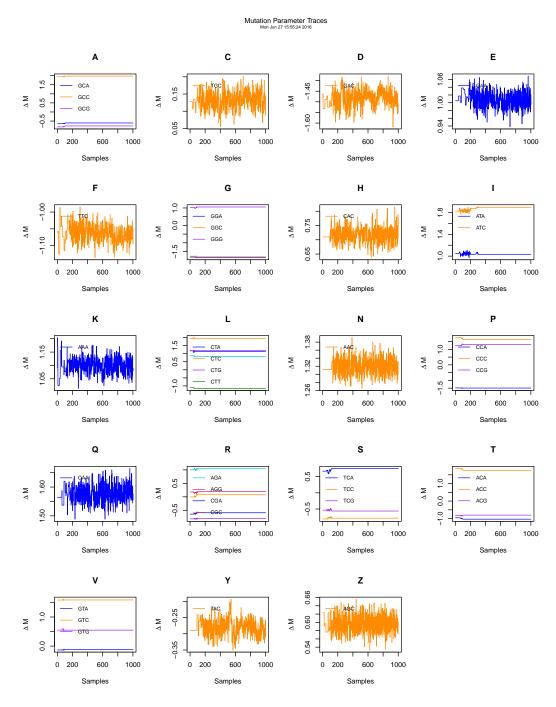


Figure 6: Mutation trace for ROC

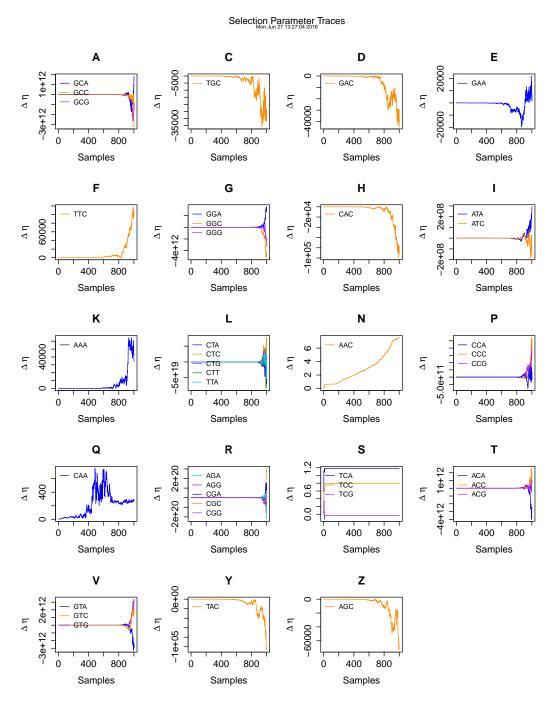


Figure 7: Selection trace for FONSE

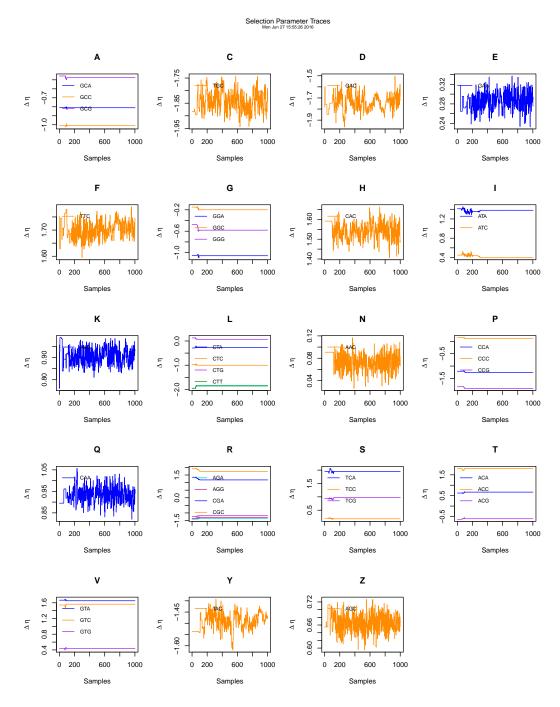


Figure 8: Selection trace for ROC

# June 29, 2016

- The main focus of today was on implementing the option to control what portion of the log likelihood trace is zoomed in. With this it will now be possible to observe the behavior of the log likelihood at the earlier stages instead of being able to observe just the drop.
- Ran FONSE with b = 0.001, samples = 1000, and thinning = 10 in order to get the zoomed in traces for each tenth of the trace. Looking at these it appears that the dropping starts fairly early and continues to drop at slightly increasing rates until the MCMC loop ends as shown in figures .

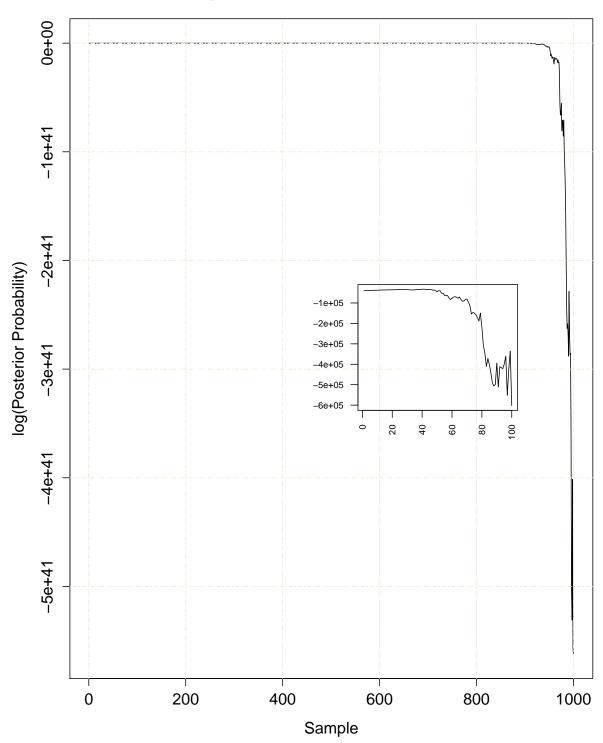


Figure 1: Log Likelihood trace with the zoom at 1-100

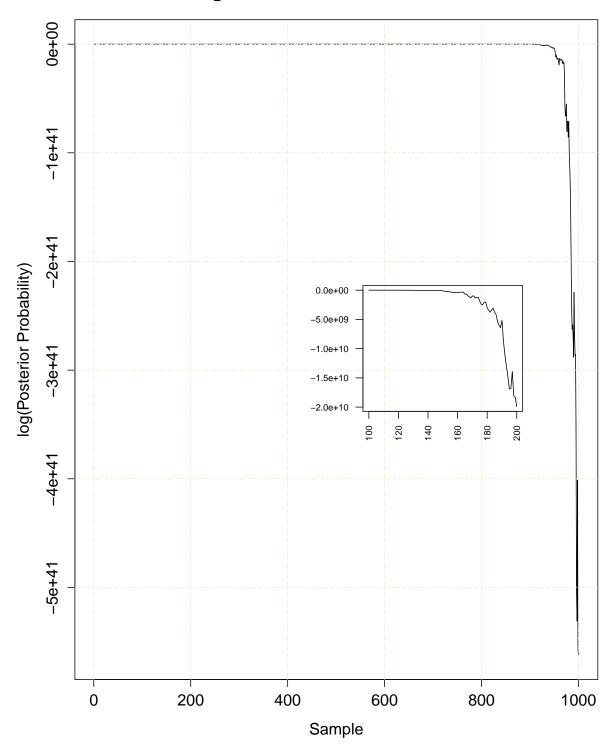


Figure 2: Log Likelihood trace with the zoom at 100-200

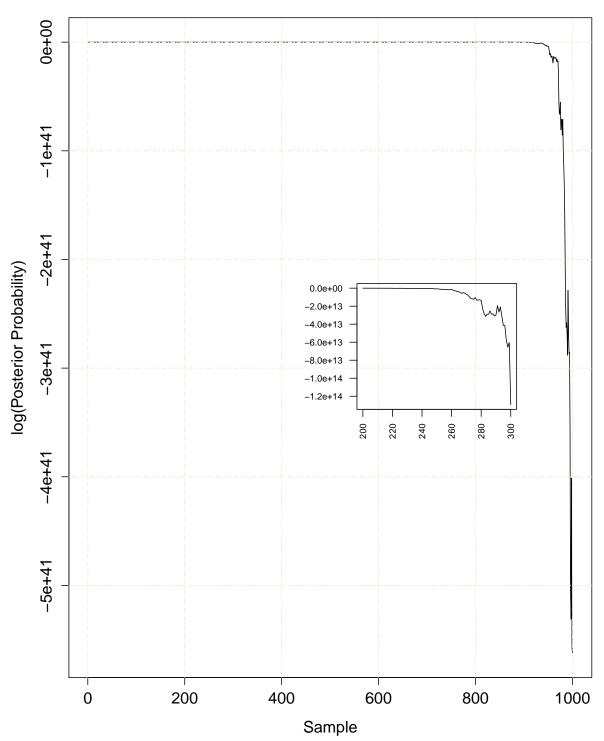


Figure 3: Log Likelihood trace with the zoom at 200-300

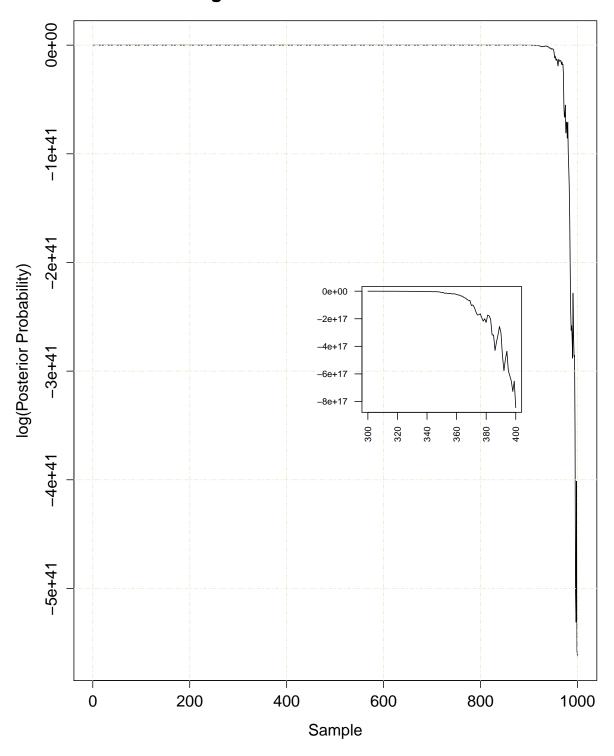


Figure 4: Log Likelihood trace with the zoom at 300-400

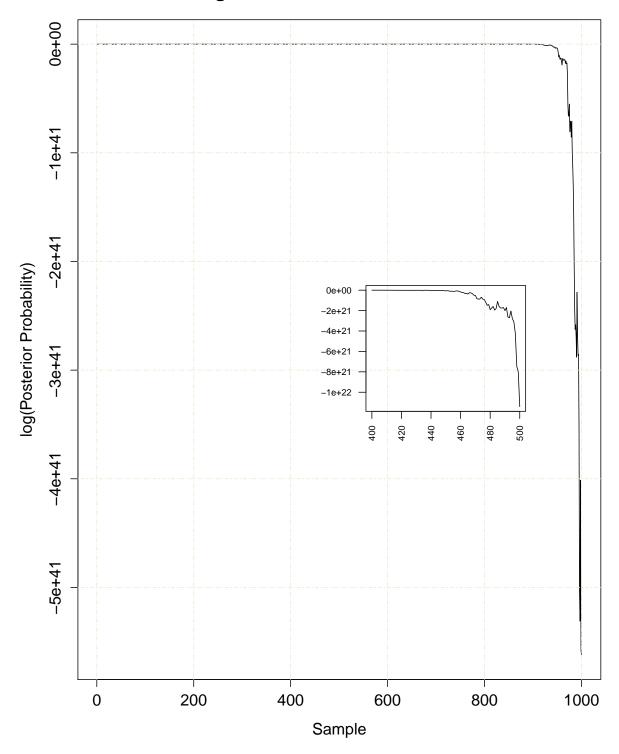


Figure 5: Log Likelihood trace with the zoom at 400-500

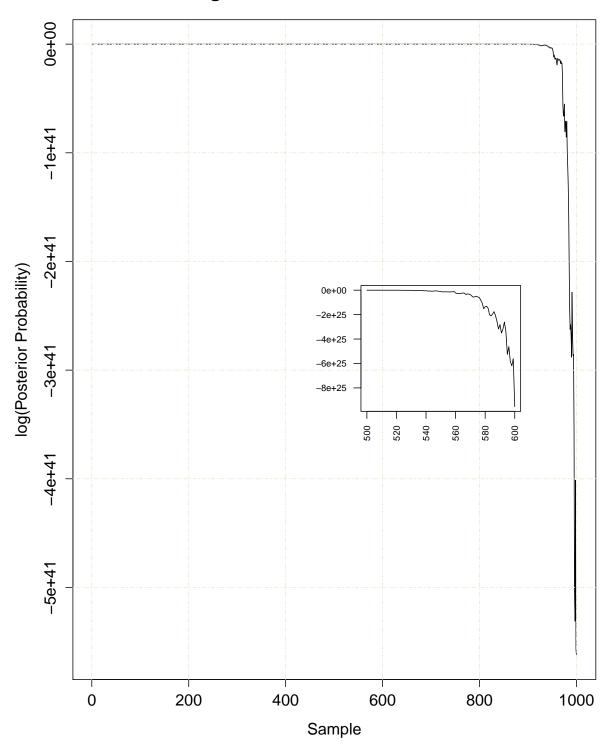


Figure 6: Log Likelihood trace with the zoom at 500-600

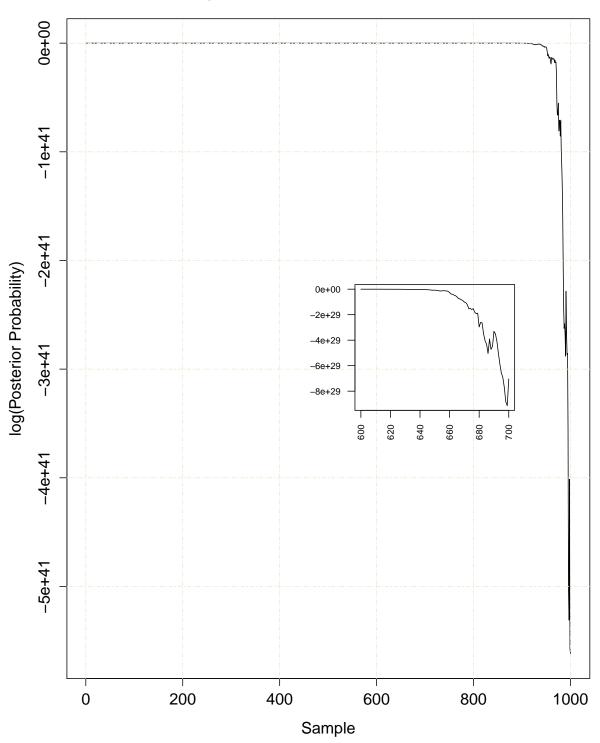


Figure 7: Log Likelihood trace with the zoom at 600-700

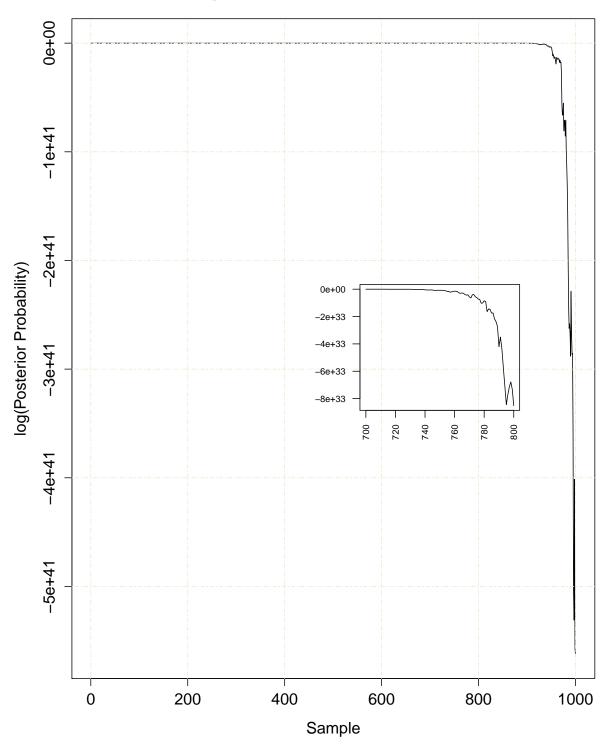


Figure 8: Log Likelihood trace with the zoom at 700-800

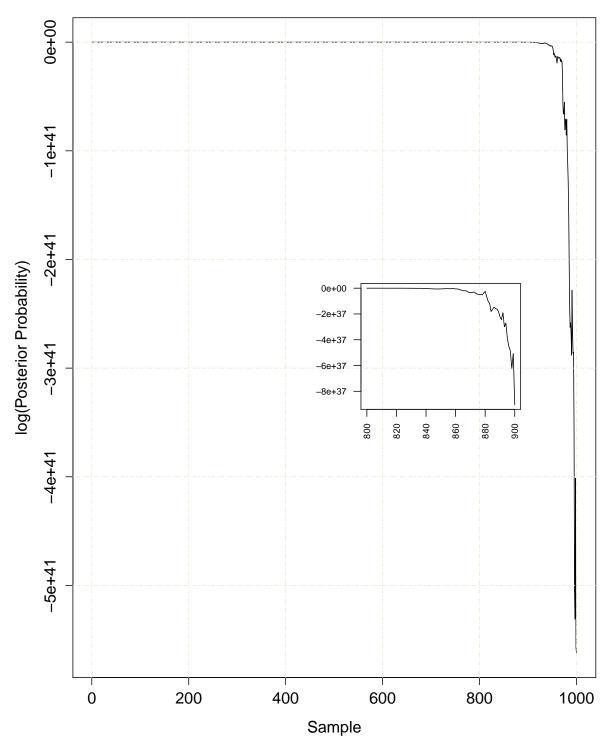


Figure 9: Log Likelihood trace with the zoom at 800-900

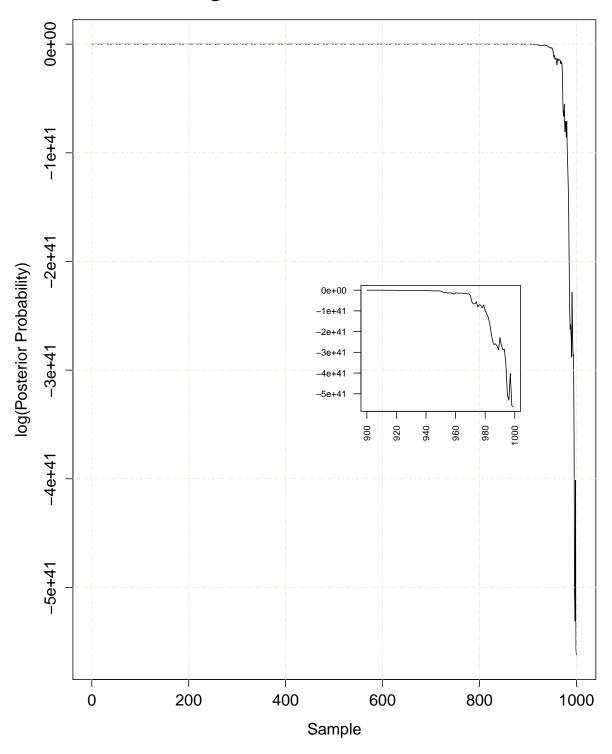


Figure 10: Log Likelihood trace with the zoom at 900-1000

## June 30, 2016

- While combing the code for the MCMC algorithm I noticed that for the large part, arrays were being used instead of vectors, and while iterating through both takes about the same amount of time and vectors have a slightly larger base overhead due to them keeping track of their own sizes, the arrays were being initialized inside of a loop by using calls to "new". Since this loop iterates "numGenes" times, every iteration caused six arrays to be declared with "new" and then deleted with "delete" at the end of the loop, leading to enormous overhead, so I ran FONSE on my laptop to get a control time and changed all of the arrays in the acceptRejectSynthesisRateLevelForAllGenes function in MCMCAlgorithm.cpp to vectors and reran FONSE twice.
- Run 1: b = 0.001, samples = 1000, thinning = 10 with arrays.
  - The MCMC loop for FONSE ran for 5708.03 seconds
  - I defined the rate as iterations/second and since the loop ran for 10000 iterations, the rate for the first was 1.75 iterations/second.
- Runs 2 & 3: same as run 1 but with vectors instead of arrays
  - The loop ran first for 3832.63 seconds then for 3859.51 seconds for an average of 3846.07
  - The rate consequently was 2.6 iterations per second
- Although one run with arrays and two with vectors isn't quite enough to confirm how much faster the algorithm for FONSE now is, initial results suggest that the algorithm is now 50% faster, but more testing is needed and will be done tomorrow.