Introduction to Bioinformatics for Computer Scientists

Lecture 10

Plan for next lectures

- Today (Alexis):
 - More on Models
 - Data Structures for unrooted Phylogenetic Trees
 - Implementing and Optimizing Likelihood Calculations
 - Parallel Likelihood Calculations
- Lecture 11 (Pierre): Discrete Operations on Trees

Protein Substitution Models

- The GTR *Q* matrix for protein data has 189 free parameters instead of just 5 (DNA)
- Estimating **189** rate parameters is difficult, time-consuming, and may lead to overparameterizing the model
- Instead, empirical models such as JTT, LG, WAG, MTMAM, etc. are used
- The *Q* matrices are obtained by jointly optimizing model parameters on a large collection of reference alignments
- The models differ with respect to:
 - the amount of data used to obtain them
 - the type of data on which the models have been optimized
 - e.g., dedicated models for HIV, FLU, Mammals
 - the numerical optimization methods used
- Examples of general models:

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- LG: Le & Gascuel: "An Improved General Amino Acid Replacement Matrix"
- WAG: Whelan & Goldman: "A General Empirical Model of Protein Evolution
- Derived from Multiple Protein Families Using a Maximum-Likelihood Approach"

Rate Heterogeneity among Sites

Gene



- Among-site rate heterogeneity
 - Biological phenomenon
 - \rightarrow different sites/columns evolve at different speeds
 - Need to accommodate this in our models

Γ-Distribution



Γ-Distribution



Discrete Γ-Distribution



An Abstract View of Γ



This is the integral of the likelihood we approximate via discretization

$$LnL(i) = log(\frac{1}{4} * (L_0 + L_1 + L_2 + L_3))$$

Log likelihood All Γ rates have equal probability $_{8}$ at site i

An Abstract View of Γ



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An Abstract View of Γ



4 times more floating point operations

Γ Model of Rate Heterogeneity with 4 discrete rates



Mixture Models

- The of rate heterogeneity is a simple example of so-called mixture models
- From Wikipedia: "In statistics, a mixture model is a probabilistic model for representing the presence of subpopulations within an overall population, without requiring that an observed data set should identify the sub-population to which an individual observation belongs. Formally a mixture model corresponds to the mixture distribution that represents the probability distribution of observations in the overall population."
- The Γ model gives us 4 discrete evolutionary rates over which we integrate (add) the likelihood for each site, without assigning a specific rate to a specific site

Mixture Models

- We can also imagine to integrate the likelihood over a set of
 - distinct *Q* matrices
 - distinct base frequencies
 - or combinations thereof
- The LG protein substitution model is an example thereof:
- It uses 4 distinct empirical Q matrices and 4 distinct sets of base frequencies π over which we integrate just like for the Γ model

An example



Taken from: "Measuring Service Reliability Using Automatic Vehicle Location Data" \rightarrow bus service reliability

Heterotachous Models



Heterotachous Models



Heterotachous Models



Multi-gene or whole-genome alignment



Multi-gene or whole-genome alignment



Multi-gene or whole-genome alignment







Joint branch length estimate









Models and Parameters

- If we add an additional parameter to a model, the likelihood will become better
- However, this does not mean anything, as
 - We might be over-parameterizing
 - The key question is if the more complex model yields a different tree topology
- So, how do we determine the best-fit model for a given dataset?

Nested models

- A particular model is said to be nested within a more complex model only if constraining parameter values of the later yields the former!
- So, the model can only be constrained in one direction to determine if its nested!
- If I need to constrain both models for which I intend to assess nesting, they are not nested.
- Example: The F81 (equal rates, unequal stationary frequencies) and K2P (2 distinct rates, equal stationary frequencies) models are not nested within each other.

 \rightarrow This is because fixing the parameter values of either model does not yield the other model

• However, they are both nested within GTR

Model Testing

- If models are nested we can use a likelihood ratio test
- Model A is nested in model B if parameters in model A are a subset of the parameters in model B
- For instance: the *Jukes Cantor (JC)* model is nested in the *General Time Reversible (GTR)* model of nucleotide substitution
- LR = P(D|A) / P(D|B) = L(A) / L(B)
- $\Delta = ln(LR^2) = 2 (ln(L(A)) ln(L(B)))$
- Compare Δ to x^2 distribution with $k_A k_B$ degrees of freedom to determine if the Δ is significant or not
- The degrees of freedom difference is the difference in the number of free parameters in the models
- How many free parameters do the *JC* and *GTR* models have?

Model Testing

- If models are nested we can use a likelihood ratio test
- Model A is nested in model B if parameters in model A are a subset of the parameters we are only allowed to compare
- For instance likelihoods on the same data D! General Time Reversible (G of nucleotide substitution
- LR = P(D|A) / P(D = L(A) / L(B))
- $\Delta = ln(LR^2) = 2 (ln(L(A)) ln(L(B)))$
- Compare Δ to x^2 distribution with $k_A k_B$ degrees of freedom to determine if the Δ is significant or not
- The degrees of freedom difference is the difference in the number of free parameters in the models
- How many free parameters do the *JC* and *GTR* models have?

What if Models are not nested?

- One can use other criteria such as
 - Akaike Information Criterion (AIC)
 - Bayesian Information Criterion (BIC)
- I will spare you the details, but the basic idea always is:
 - Compute likelihood of alternative models
 - Penalize the more parameter-rich models

Outline

- More on Models
- Data Structures for unrooted Phylogenetic Trees
- Implementing and Optimizing Likelihood Calculations
- Parallel Likelihood Calculations

Data Structures for unrooted Trees

- Unrooted trees with dynamically changing virtual roots need a dedicated tree data structure
- Why can the virtual root positions change dynamically?
- If we apply a topological move (NNI, SPR, TBR) will we have to re-compute all conditional likelihood vectors?

Memory Organization: Conditional Likelihood Vectors with an Unrooted View



Memory Organization: Conditional Likelihood Vectors with a Rooted View



Memory Organization: CLVs with a Rooted View



Memory Organization: Ancestral Vectors with a Rooted View

New Virtual Root


Memory Organization: Tip Vectors



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Optimization of Likelihood Calculations

- Use SSE3 & AVX vector intrinsics
- Also: GPUs, FPGAs, the Intel Xeon PHI
- Special implementations (why?) for computing CLVs:





Detect identical patterns and omit second computation







Floating Point Numbers

• Machine numbers are an imperfect mapping of the infinite real numbers to a finite number of machine values!



Floating Point Arithmetics: The Root of All Evil

- Computational science mostly relies on floating-point intensive codes
- How do we verify these codes?
- We stand on shaky grounds
- Scientists using those codes assume that there are no bugs
- Double precision arithmetics required for certain applications
- Who knows what de-normalized floating point numbers are?
 - \rightarrow Please have a look at:

J. Björndalen, O. Anshus: "Trusting floating point benchmarks-are your benchmarks really data-independent?" Applied Parallel Computing. State of the art in Scientific Computing 2010; pp 178-188, Springer.

and at my micro-benchmark at: https://github.com/stamatak/denormalizedFloatingPointNumbers

Floating Point Arithmetics: The Root of All Evil

- Computational science mostly reli
- How do we verify these codes?
- We stand on shaky grounds

Why is this relevant when Talking about Maximum Likelihood?

- Scientists using those codes assume that there are no bugs
- Double precision arithmetics required for certain applications
- Who knows what de-normalized floating point numbers are?
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Post-order Traversal



Values in conditional likelihood vectors get smaller and smaller as we move to the root



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P[i] = f(Q[i], R[i])



P[i] = f(Q[i], R[i])



Loop Level Parallelism

virtual root





Parallel Post-order Traversal

Only need to synchronize at the root \rightarrow MPI_Reduce() to calculate: $\Sigma \log(I_i)$





Classic Fork-Join with Busy-Wait



Synchronizations in RAxML with Pthreads

- RAxML Pthreads for a run time of about 10 seconds on 16 cores/threads
- 404 taxa 7429 sites: **194,000** Barriers
- 1481 taxa 1241 sites: **739,000** Barriers
- A paper on performance of alternative PThreads barrier implementations:

S.A. Berger, A. Stamatakis: "Assessment of Barrier Implementions for Fine-Grain Parallel Regions on Current Multi-core Architectures", *IEEE Cluster* 2010.

Classic Fork-Join with Busy-Wait



Classic Fork-Join with Busy-Wait



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Problems start with partitioned datasets!



Parallel Performance Problems

- They all start with partitioned datasets!
- How do we distribute partitions to processors?
- How do we calculate parameter changes?
- How much time does our broadcast take?
- Goal: Keep all processors busy all the time
 - \rightarrow minimize communication and synchronization!

Example

Blue Gene Red Gene

Sequence 1 Sequence 5







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G0	G1	G2	G3

P0 P1







- The multiprocessor job scheduling problem in phylogenetics
 - Problem when #partitions >> #cores
 - Tested per-site (cyclic/modulo) data distribution versus per partition data distribution
 - We used the Longest Processing Time (LPT) heuristics for assigning partitions to processors
 - 25 taxa, 220,000 sites, 100 genes
 - GAMMA model
 - naïve: 613 secs
 - LPT: **550** secs
 - CAT model
 - naïve: 298 secs
 - LPT: **127** secs
 - Larger protein dataset under Γ model of rate heterogeneity: 10-fold performance improvement!

J. Zhang, A. Stamatakis: "The Multi-Processor Scheduling Problem in Phylogenetics", 11th IEEE HICOMB workshop (in conjunction with IPDPS 2012).

LPT heuristics for multi-processor scheduling

- Sort jobs (partitions) by processing length (partition length) in decreasing order
- Remove a job (partition) from the sorted list and assign it to the processor with the earliest end time (the smallest sum of partition lengths)
- Repeat until the sorted list is empty
- Upper bound: 4/3 1/(3p) * OPT, where p is the number of processors
- Graham, R. L.: "Bounds on Multiprocessing Timing Anomalies". *SIAM Journal on Applied Mathematics* 17 (2): 416–429, 1969.
- Remark: LPT works surprisingly well (see our paper on the phylogenetic problem where we also tested other heuristics)

Partitioned Branch Lengths & other parameters



separate estimate of Q-Matrix alpha-shape Branch Lengths separate estimate of Q–Matrix alpha–shape Branch Lengths separate estimate of Q-Matrix alpha-shape Branch Lengths

Load-Balance II



- Assume 10 branches
- Each branch requires 10 Newton-Raphson Iterations
- Each NR Iteration requires a synchronization via a reduction operation
- One branch/partition at a time: 100 sync. points, less work (only one partition) per sync. point
- All branches concurrently: 10 sync. points, more work per sync. point
- Branches will need distinct number of operations
- Add convergence state → bit vector

Org1 AC GT Org2 AC TT









 00
 00
 01



In this example: 4 instead of 7 sync points!



A. Stamatakis, M. Ott: "Load Balance in the Phylogenetic Likelihood Kernel". Proceedings of ICPP 2009, Vienna, Austria, September 2009.



Alternative MPI parallelization



time

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Execution

Alternative MPI parallelization



Execution time

ExaML

- New code implementing this new parallelization scheme
- https://github.com/stamatak/ExaML
- A. Stamatakis, A. J. Aberer: "Novel Parallelization Schemes for Large-Scale Likelihood-based Phylogenetic Inference", accepted for publication at *IPDPS 2013*, Boston, USA, 2013.
- Up to 3 times faster than RAxML-Light (2012) on large, partitioned datasets
- Tested with up to 1536 cores on our cluster at HITS
- Future developments
 - 20,000,000 CPU hors on SuperMUC for
 - Improving scalability
 - Implementing fault tolerance
 - Execute 1KITE tree inferences
 - Further details \rightarrow ask Andre Aberer