Odds and Ends

Last day of class is usually a mix of trying to find gaps or fill in holes.

I discussed phylogenetic tree building.

```
git clone https://github.com/biodataprog/GEN220_2022_examples.git
cd GEN220_2022_examples/Trees
module load muscle
module load fasttree
module load hmmer/3
module load IQ-TREE/2.1.1
module load trimal


# build an alignment of sequences already identified as homologs
# previously I had started with MET12 (S. cerevisiae) enzyme
more  MET12.fa # single sequence
ls -l MET12.hit_seqs.fasta # the collection of homologs for MET12 in a few yeast fungi
# denovo multiple alignment - writes in multi-fasta format
muscle -in MET12.hit_seqs.fasta -out MET12.hit_seqs.fasaln
# denovo multiple alignment - writes in multi-fasta format - writes in Clustal format
muscle -in MET12.hit_seqs.fasta -out MET12.hit_seqs.fasaln.clw -clw
# trim sequences - using automated parameters - see http://trimal.cgenomics.org/trimal for
trimal -automated1 -in MET12.hit_seqs.fasaln -out MET12.hit_seqs.mfa.trim


# build a tree w fastree (FastTreeMP uses multiple processors, FastTree uses 1 processor on
FastTreeMP <  MET12.hit_seqs.fasaln >  MET12.hit_seqs.tre
# build a tree with IQ-TREE2 - ultrafast bootstrap and first determine optimal number of pro
iqtree2 -s MET12.hit_seqs.fasaln -nt AUTO -bb 1000 -alrt 1000
```

Some links * Muscle - Multiple alignment tool * TrimAl - alignment trimming
tool * HMMER - HMMER - Hidden Markov Model for biosequence analyses. *
FastTree - Fast Phylogenetic Tree construction * IQ-TREE - Phylogenetic Tree
construction * RAxML; a tutorial * iTOL - Tree visualization (web-based) tool *
FigTree - Tree visualization (can run on HPCC if you have X11 enabled: `module
load figtree; figtree`) * ggtree - R package for Tree rendering

I also showed how to use HMMER and hmmalign

```
module load hmmer/3


# build an HMM from a multiple alignment
hmmbuild MET12.hmm MET12.hit_seqs.fasaln
```

This is a little circular I am searching the HMM back against the original
sequences, but if you wanted to instead search this HMM against a database of
proteins (eg swissprot or your collection of proteins from species)

```
module load hmmer/3
```

```
# domtbl is the result file which has columns of data that are parseable instead of more com
hmmsearch -E 1e-3 --domtblout MET12.search.domtbl MET12.hmm DATABASE > MET12.search.hmmsear

# to align a set of proteins back to an HMM (which is instead of doing a denovo multiple al
hmmalign MET12.hmm MET12.hit_seqs.fasta  > MET12.hit_seqs.stk
# convert the stockholm format to multifasta
esl-reformat afa MET12.hit_seqs.stk > MET12.hit_seqs.hmmalign.fasaln
# convert the stockholm format to clustal
esl-reformat clustal MET12.hit_seqs.stk > MET12.hit_seqs.hmmalign.fasaln
```