Biol5705

Module: Gene Sequence Analysis

Assignment 2

Email results to morgan.g.i.langille@gmail.com by start of class next week.

PSI-BLAST

- Access PSI-BLAST from the BLAST page http://www.ncbi.nlm.nih.gov/BLAST/
 - Paste the gi number "4505641" in the query box.
 - Use the default parameters, except for choosing the RefSeq database, setting Expect
 Threshold (E-value) to 0.1, and setting maximum target sequences to 1000. Ensure that
 the PSI-BLAST Threshold is 0.005. Run PSI-BLAST.
 - The hits are divided into two sections. The hits with better statistical significance than the e-value threshold, 0.005, are listed first. Those with e-values worse than threshold, but have an e-value better than that selected on the query page, 0.1, are listed further down the page. Hits with e-values better than the threshold are used in forming the profile that will be used in subsequent PSI-BLAST iterations.
 - 1) What is the name of the guery sequence?
 - 2) What is the length of the query sequence?
 - 3) How many sequences are worse then your PSI-BLAST Threshold (but better than your Expect Threshold)?
 - Run PSI-BLAST iteration 2 with default selected sequences (above psi-blast threshold)
 - Note that newly found sequences are marked high-lighted in yellow.
 - Keep running PSI-BLAST until you find a very distant homolog from Bacillus smithii (note: try using ctrl-F to search for organism name, instead of manually scanning).
 - 4) What iteration of PSI-BLAST did you find the B.smithii homolog?
 - 5) What is the name of the homolog
 - 6) What is the e-value and identity of the hit?

Genome Alignment with Mauve

- Download 4 genomes files:
- http://morganlangille.com/teaching/biol5705/assignment_2_genomes.zip
- Download Mauve (http://gel.ahabs.wisc.edu/mauve/download.php)

Start Mauve by clicking on "Mauve" file
Align your 4 genomes using "progressive mauve" with default parameters (this will take awhile)

- 7) Export your alignment as an image (Tools->Export->Export Image). Put the alignment picture embedded with your answers or send as separate attachment with your answer document.
- 8) What genome contains a very large inversion?

Search for PLES_53841 by name using View->Go to->Find Features

- 9) What genome is this gene in? And what is the gene name? Does this gene appear to be an insertion or deletion?
- 10) What is the name of the flanking genes (appearing directly on the left and right) of PLES_53841? Are these conserved (appearing in other genomes) or variable (not present in some of the genomes)?