

## Introduction to Bioinformatics

### Problem Set 8: Tables and Motif Discovery Through PSSM's

#### Tables and the [ ] notation

1. Create a table of the form letter[number], where letter[1] = "A", letter[2] = "B", etc.
2. Make and display an x \* y multiplication table for x and y, each going from 1 to 15. The [] function (in the List/Tables menu) will be essential here. You can use it to define elements of a two-dimensional table, as shown by example below:

The screenshot shows three steps in the PSSM software interface:

- DEFINE row = 1**: A window where the variable 'row' is defined with the value '1'.
- DEFINE column = "A"**: A window where the variable 'column' is defined with the value '"A"'. Below this, a larger window shows the definition of 'excel-table' as a function of 'row' and 'column', with the value '47' displayed.
- DISPLAY-TABLE excel-table**: A window where the 'excel-table' is displayed.

3. Make and display a table containing information about the organisms known to BioBIKE. The row labels should be the names of the organisms. The column labels should genome size, number of genes, and GC-fraction.
4. Determine the frequency of each dinucleotide in the genome of the cyanobacterium ss120. Put the values in a table and then display it. The following function will be useful:

The screenshot shows the function **ALL-DNA-SEQUENCES OF-LENGTH** with the value '2' entered in the input field.

5. The [ ] notation can be used for other types of data besides tables. Think of a[b] as "a sub b". Define a list of some sort, and use the [ ] function to pull out its third element. Try the same thing with a string or a sequence.
6. Use tables within a loop that counts every instance of 13-mers (13-nucleotide sequences) within the che12 genome. Sort the resulting table, sorting the most abundant 13-mers to be at the top of the list, and display it. Compare your list to the sequences found by Gomathi et al.

#### Using PSSM's to find sequence motifs

7. In last week's lab you found motifs of iron-sulfur proteins – now use it!
  - 7a. Redo the steps to obtain the motifs, and identify a motif with the characteristic C\*\*C\*\*C\*\*\*C pattern.
  - 7b. Extract the alignment of sequences from that motif (output by MOTIFS-IN). One of the FIRST, SECOND,... functions will be useful for this purpose, once you know which of the functions is the one you want.

- 7c.** With that alignment in hand, run APPLY-PSSM-TO the PROTEINS-OF any bacterium, using the alignment to fill in the box governed by the WITH-PSSM-FROM option. The output will be of the form: (protein amino-acid-coordinate FORWARD score).
  - 7d.** Extract the proteins from the list produced by APPLY-PSSM-TO. The FIRST function will be useful, if you use the IN-EACH pre-option.
  - 7e.** Display the descriptions of the proteins. Are any/all iron-sulfur proteins?
- 8.** If your research project is protein-oriented, use MOTIFS-IN and APPLY-PSSM-TO to look for instances of your favorite protein in different phage genomes.