# 22BIO201 Intelligence of Biological Systems 1

### Lab Sheet 3

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### **Problem 1: Generate a Random DNA Sequence**

**Description:** Create a random DNA string with letters from the whole alphabet A, C, G, and T. First make a list of random letters and then join all those letters to a string. Also write another function to count the number of bases in the random sequence and measure the CPU time for large such DNA strings. (Hint: use import random, import time)

# Reference

Illustrating Python via Bioinformatics Examples, Hans Petter Langtangen, Geir Kjetil Sandve, https://hplgit.github.io/bioinf-py/doc/pub/html/main\_bioinf.html

### **Problem 2: Compute the Hamming Distance Between Two Strings**

We say that position i in k-mers  $p_1 \dots p_k$  and  $q_1 \dots q_k$  is a mismatch if  $p_i \neq q_i$ . For example, CGAAT and CGGAC have two mismatches. The number of mismatches between strings p and q is called the Hamming distance between these strings and is denoted HammingDistance(p, q).

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### **Hamming Distance Problem**

Compute the Hamming distance between two DNA strings.

Given: Two DNA strings.

Return: An integer value representing the Hamming distance.

### **Sample Dataset**

**GGGCCGTTGGT** 

**GGACCGTTGAC** 

### **Sample Output**

3

Visit  $\underline{\text{http://rosalind.info/problems/ba1G/}}$ . Solve the problem. Use the sample dataset given in the site.

# **Problem 3: Find Patterns Forming Clumps in a String**

Given integers L and t, a string Pattern forms an (L, t)-clump inside a (larger) string Genome if there is an interval of Genome of length L in which Pattern appears at least t times.

For example, TGCA forms a (25,3)-clump in the following *Genome*: gateageataagggteeeTGCAATGCATGACAAGCCTGCAgttgttttac

### **Clump Finding Problem**

Find patterns forming clumps in a string.

Given: A string *Genome*, and integers k, L, and t.

Return: All distinct k-mers forming (L, t)-clumps in *Genome*.

#### **Pseudocode:**

```
ClumpFinding(Genome,k,L,t)

for i←0 to|Genome|-L

count←0 for all kmers in Genome(i,L)

for j←0 to L- k

kmer=Genome(i+j,L)

count(kmer)=count(kmer)+1

for all kmers in count

if count(kmer)>=t and kmer has not been outputted

output kmer
```

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# **Sample Dataset**

CGGACTCGACAGATGTGAAGAATGTGAAGACTGAGTGAAGAGAAGAGAAAAAATGTACGAACATGTACGACATATGTCCTATGGC

5 75 4

# **Sample Output**

CGACA GAAGA AATGT

Visit <a href="http://rosalind.info/problems/ba1E/">http://rosalind.info/problems/ba1E/</a> . Solve the problem. Use the sample dataset given in the site.

### **Problem 4: Find a Position in a Genome Minimizing the Skew**

Define the **skew** of a DNA string *Genome*, denoted *Skew*(*Genome*), as the difference between the total number of occurrences of 'G' and 'C' in *Genome*. Let *Prefix<sub>i</sub>* (*Genome*) denote the **prefix** (i.e., initial substring) of *Genome* of length *i*. For example, the values of *Skew*(*Prefix<sub>i</sub>* ("CATGGGCATCGGCCATACGCCCATGGGCATCGGCCATACGCC")) are:

### **Minimum Skew Problem**

Find a position in a genome minimizing the skew.

Given: A DNA string Genome.

**Return:** All integer(s) i minimizing  $Skew(Prefix_i (Text))$  over all values of i (from 0 to |Genome|).

### Sample Dataset

CCTATCGGTGGATTAGCATGTCCCTGTACGTTTCGCCGCGAACTAGTTCA CACGGCTTGATGGCAAATGGTTTTTCCGGCGACCGTAATCGTCCACCGA G

**Sample Output** 

53 97

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