CSE 142, Autumn 2018 Programming Assignment #7: DNA (20 points) Due Tuesday, November 20th, 9:00 PM

Special thanks to UW CSE professor Martin Tompa for his help with the development of this assignment.

This assignment focuses on arrays and file/text processing. Turn in a file named DNA.java. You will also need the two input files dna.txt and ecoli.txt from the course web site. Save these files in the same folder as your program.

The assignment involves processing data from genome files. Your program should work with the two given input files. If you are curious (this is not required), the National Center for Biotechnology Information publishes many other bacteria genome files. The last page tells you how to use your program to process other published genome files.

Background Information About DNA:

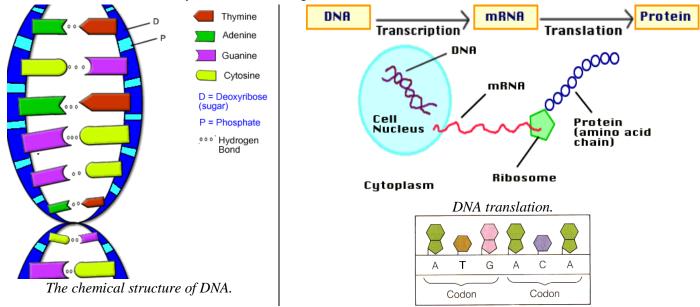
Note: This section explains some information from the field of biology that is related to this assignment. It is for your information only; you do not need to fully understand it to complete the assignment.

Deoxyribonucleic acid (DNA) is a complex biochemical macromolecule that carries genetic information for cellular life forms and some viruses. DNA is also the mechanism through which genetic information from parents is passed on during reproduction. DNA consists of long chains of chemical compounds called *nucleotides*. Four nucleotides are present in DNA: Adenine (A), Cytosine (C), Guanine (G), and Thymine (T). DNA has a double-helix structure (see diagram below) containing complementary chains of these four nucleotides connected by hydrogen bonds.

Certain regions of the DNA are called genes. Most genes encode instructions for building proteins (they're called "protein-coding" genes). These proteins are responsible for carrying out most of the life processes of the organism.

Nucleotides in a gene are organized into *codons*. Codons are groups of three nucleotides and are written as the first letters of their nucleotides (e.g., TAC or GGA). Each codon uniquely encodes a single amino acid, a building block of proteins.

The process of building proteins from DNA has two major phases called *transcription* and *translation*, in which a gene is replicated into an intermediate form called *mRNA*, which is then processed by a structure called a *ribosome* to build the chain of amino acids encoded by the codons of the gene.



The sequences of DNA that encode proteins occur between a *start codon* (which we will assume to be ATG) and a *stop codon* (which is any of TAA, TAG, or TGA). Not all regions of DNA are genes; large portions that do not lie between a valid start and stop codon are called *intergenic DNA* and have other (possibly unknown) functions. Computational biologists examine large DNA data files to find patterns and important information, such as which regions are genes. Sometimes they are interested in the percentages of mass accounted for by each of the four nucleotide types. Often high percentages of Cytosine (C) and Guanine (G) are indicators of important genetic data.

For more information, visit the Wikipedia page about DNA: http://en.wikipedia.org/wiki/DNA

Program Overview:

In this assignment, you will read an input file containing named sequences of nucleotides and produce information about them. For each nucleotide sequence, your program should **count** the occurrences of each of the four nucleotides (A, C, G, and T). The program should also compute the **mass percentage** occupied by each nucleotide type, rounded to one digit past the decimal point. Next, the program should report the **codons** (trios of nucleotides) present in each sequence and predicts whether or not the sequence is a **protein-coding** gene. For our purposes, a protein-coding gene is a string that:

- begins with a valid start codon (ATG)
- ends with a valid *stop codon* (one of the following: TAA, TAG, or TGA)
- contains at least 5 total codons (including its initial start codon and final stop codon)
- Cytosine (C) and Guanine (G) combined account for at least 30% of its total mass

(Note that these are approximations for our assignment, not exact constraints used in computational biology.)

The DNA input data consists of line pairs. The first line is the name of the nucleotide sequence, and the second is the nucleotide sequence itself. Each character in a sequence of nucleotides will be A, C, G, T, or a dash character, "-". The nucleotides in the input can be either upper or lowercase.

Input file dna.txt (partial):

```
cure for cancer protein
ATGCCACTATGGTAG
captain picard hair growth protein
ATGCCAACATGGATGCCcGATAtGGATTGA
bogus protein
CCATt-AATGATCa-CAGTt
...
```

The **dash** "-" **characters** represent "junk" or "garbage" regions in the sequence. For most of the program they should be ignored in your computations, though they do contribute to the total mass of the sequence as described later.

Program Behavior:

Your program should begin with an introduction and a prompt for input and output file names. You may assume the user will type the name of an existing input file that is in the proper format. Your program should read the input file to process its nucleotide sequences and output the results into the given output file. Notice that the nucleotide sequence is output in all uppercase and that the nucleotide counts and mass percentages are shown in the order A, C, G, T. Also notice that a given codon (such as GAT) might occur more than once in the same sequence.

Log of execution (user input underlined):

```
This program reports information about DNA nucleotide sequences that may encode proteins.
Input file name? dna.txt
Output file name? output.txt
```

Output file output. txt after above execution (partial):

```
Region Name: cure for cancer protein
Nucleotides: ATGCCACTATGGTAG
Nuc. Counts: [4, 3, 4, 4]
Total Mass%: [27.3, 16.8, 30.6, 25.3] of 1978.8
Codons List: [ATG, CCA, CTA, TGG, TAG]
Is Protein?: YES
Region Name: captain picard hair growth protein
Nucleotides: ATGCCAACATGGATGCCCGATATGGATTGA
Nuc. Counts: [9, 6, 8, 7]
Total Mass%: [30.7, 16.8, 30.5, 22.1] of 3967.5
Codons List: [ATG, CCA, ACA, TGG, ATG, CCC, GAT, ATG, GAT, TGA]
Is Protein?: YES
Region Name: bogus protein
Nucleotides: CCATT-AATGATCA-CAGTT
Nuc. Counts: [6, 4, 2, 6]
Total Mass%: [32.3, 17.7, 12.1, 29.9] of 2508.1
Codons List: [CCA, TTA, ATG, ATC, ACA, GTT]
Is Protein?: NO
```

Implementation Guidelines, Hints, and Development Strategy:

The main purpose of this assignment is to demonstrate your understanding of arrays and array traversals with for loops. Therefore, you should use arrays to store the various data for each sequence. In particular, your nucleotide counts, mass percentages, and codon lists should all be stored using arrays. Additionally you should use arrays and for loops to transform the data from one form to another as follows:

- from the original nucleotide sequence string to nucleotide counts;
- from nucleotide counts to mass percentages; and
- from the original nucleotide sequence string to codon triplets.

These transformations are summarized by the following diagram using the "cure for cancer" protein data:

```
      Nucleotides:
      "ATGCCACTATGGTAG"

      What is computed
      Output to file

      Counts:
      {4, 3, 4, 4}

      Nuc. Counts:
      [4, 3, 4, 4]

      Mass %:
      {27.3, 16.8, 30.6, 25.3}
      Total Mass%:
      [27.3, 16.8, 30.6, 25.3] of 1978.8

      Codons:
      {ATG, CCA, CTA, TGG, TAG}
      Codons List:
      [ATG, CCA, CTA, TGG, TAG]

      Is protein?:
      YES
```

Recall that you can print any array using the method Arrays.toString. For example:

To compute **mass percentages**, use the following as the mass of each nucleotide (grams/mol). The dashes representing "junk" regions are excluded from many parts of your computations, but they *do* contribute mass to the total.

Adenine (A): 135.128
Cytosine (C): 111.103
Guanine (G): 151.128
Thymine (T): 125.107
Junk (-): 100.000

For example, the mass of the sequence ATGG-AC is (135.128 + 125.107 + 151.128 + 151.128 + 100.000 + 135.128 + 111.103) or 908.722. Of this, 270.256 (29.7%) is from the two Adenines; 111.103 (12.2%) is from the Cytosine; 302.256 (33.3%) is from the two Guanines; 125.107 (13.8%) is from the Thymine; and 100.000 (11.0%) is from the "junk" dash.

We suggest that you start this program by writing the code to read the input file. Try writing code to simply read each protein's name and sequence of nucleotides and print them. Read each line from the input file using Scanner's nextLine method. This will read an entire line of input and return it as a String.

Next, write code to pass over a nucleotide sequence and count the number of As, Cs, Gs, and Ts. You can use string's charAt method to get individual characters. Put your counts into an array of size 4. To map between nucleotides and array indexes, you may want to write a method that converts a single character (i.e. A, C, T, G) into an index (i.e. 0 to 3).

Once you have the counts working correctly, you can convert your counts into a new array of percentages of mass for each nucleotide using the preceding nucleotide mass values. If you've written code to map between nucleotide letters and array indexes, it may also help you to look up mass values in an array such as the following:

```
double[] masses = {135.128, 111.103, 151.128, 125.107};
```

You may store your mass percentages already rounded to one digit past the decimal or you can round when printing the mass percentages array using printf. If you choose to store the percentages pre-rounded, use Math.round as follows:

```
double num = 1.6666667;
double rounded = Math.round(num * 10.0) / 10.0;
System.out.print("the answer is " + rounded);  // the answer is 1.7
```

Remember that the "junk" dashes do contribute mass to the total. For other parts of your program you may want to remove dashes from the input; consider using the replace method on the nucleotide string to eliminate these characters.

After computing mass percentages, you must break apart the sequence into codons and examine each codon. You may wish to review methods of the String class presented in Chapters 3 and 4, such as substring, charAt, indexOf, replace, toUpperCase, and toLowerCase.

We also suggest that you first get your program working correctly printing its output to the *console* before you save the output to a file. Once you have your program printing correct output to the console, save the output to a file by using a PrintStream as described in Section 6.4 of the textbook.

You may assume that the input file exists, is readable, and contains valid input. (In other words, you should not re-prompt for input or output file names.) You may assume that each sequence's number of nucleotides (without dashes) will be a multiple of 3, although the nucleotides on a line might be in either uppercase or lowercase or a combination. Your program should overwrite any existing data in the output file (this is the default PrintStream behavior).

Style Guidelines:

For this assignment you are required to have the following **four class constants**:

- one for the **minimum number of codons** a valid protein must have, as an integer (default of 5)
- a second for the **percentage** of mass from C and G in order for a protein to be valid, as an integer (default of 30)
- a third for the number of **unique nucleotides** (4, representing A, C, G, and T)
- a fourth for the number of **nucleotides per codon** (3)

For full credit it should be possible to change the first two constant values (minimum codons and minimum mass percentage) to change your program's behavior for evaluating protein validity. The other two constants won't ever be changed but are still useful to make your program more readable. Refer to these constants in your code and do not refer to the bare number for these values (4 or 3) directly. You may use additional constants if they make your code clearer.

We will grade your method structure strictly on this assignment. Use at least **four nontrivial methods** besides main. These methods should use parameters and returns, including arrays, when appropriate. The methods should be well-structured and avoid redundancy. No one method should do too large a share of the overall task. The textbook's case study at the end of Chapter 7 is a good example of a larger program with methods that pass arrays as parameters.

In particular, we require that you have a method that can be called exactly once to print all file output for a given nucleotide sequence (nucleotides, counts, %, is it a protein, etc.). In other words, all output to the file should be done through one method called on each nucleotide sequence from the input. Your other methods should do the computations to gather information to be passed to this output method.

Your main method should be a concise summary of the overall program. It is okay for main to contain some code such as println statements. But main should not perform too large a share of the overall work itself, such as examining each character of an input line. Also avoid "chaining," when many methods call each other without ever returning to main.

We will also check strictly for redundancy on this assignment. If you have a very similar piece of code that is repeated several times in your program, eliminate the redundancy by creating a method, by using for loops over the elements of arrays, and/or by factoring if/else code as described in section 4.3 of the textbook.

Since arrays are a key component of this assignment, part of your grade comes from using arrays properly. For example, you should reduce redundancy by using **traversals** over arrays (for loops over the array's elements). This is preferable to writing out a separate statement for each array element (a statement for element [0], then another for [1], then for [2], etc.). Also carefully consider how arrays should be passed as parameters and/or returned from methods as you are decomposing your program. Recall that arrays use *reference semantics* when passed as parameters, meaning that an array passed to a method can be modified by that method and the changes will be seen by the caller.

You are limited to features in Chapters 1 through 7 of the textbook. Follow all previous style guidelines for indentation, names, variables, types, line lengths, and comments (at the beginning of your program, on each method, and on complex sections of code).

Additional Input Files (Optional):

If you would like to generate additional input files to test your program, you can create them from actual NCBI genetic data. The following web site has many data files that contain complete genomes for viruses:

ftp://ftp.ncbi.nlm.nih.gov/genomes/Viruses/

The site contains many directories with names of organisms. After entering a directory, you can find and save a genome file (a file whose name ends with .fna) and a protein table (a file whose name ends with .ptt). On the course web site we will provide you with a program to convert these .fna and .ptt files into input files suitable for your homework.