

# Project 1: DNA Analysis

## 1 Introduction

You have recently been employed by Rhonda Labs, a leading research institute working to uncover the secrets of rare genetic diseases. Your first task is to write a program that analyzes DNA from various species and individuals, compares their similarities, and performs analyses on the given sequences.

Before you dive into the project, let's review a bit of biology to help you understand what you will be working with. DNA, or deoxyribonucleic acid, is the molecule that carries genetic information in almost all living organisms. It acts like a set of instructions that tells cells how to function, grow, and reproduce.

DNA is made up of four chemical bases:

- Adenine (A)
- Cytosine (C)
- Guanine (G)
- Thymine (T)

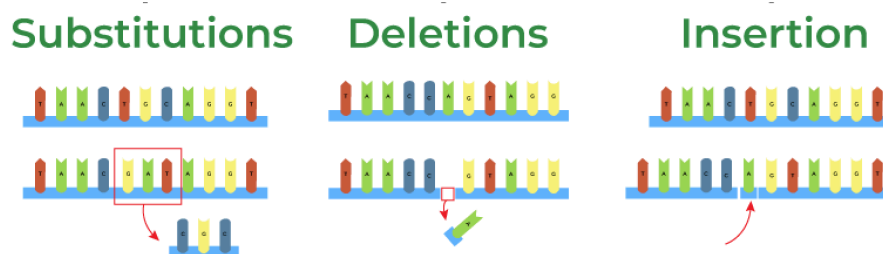


Figure 7.1: Types of DNA mutations

These bases pair together (A with T, C with G) to form the DNA double helix, which is the overall structure of DNA. A DNA strand is made up of a sequence of these base pairs. Every organism has its own unique DNA sequence, though many organisms share similarities, especially related species. Sometimes, DNA sequences can change, leading to mutations, such as:

- Substitutions (one base is swapped for another)
- Insertions (extra bases are added)
- Deletions (bases are removed)

For this task, the steps are outlined and divided into individual questions. By the end of the project, you will have developed a cohesive program that a user can interact with by inputting their DNA sequences. Below is a breakdown of the steps:

1. Check whether your DNA strands are composed of valid bases

2. Develop functions to compare DNA strands and assess their similarity
3. Develop a function that compares two DNA sequences and identifies all types of mutations between them
4. Transcribe DNA to RNA and compute the reverse complement of a DNA strand
5. Find open reading frames within a DNA strand
6. Final step: make it more accessible and user-friendly!

## 2 Assignment

**Warning: You are not allowed to use global variables for this project.**

All function names, return types, and parameters must precisely match those shown. You may not use pass by reference or otherwise modify the function prototypes. You are welcome to create additional functions that may help streamline your code. You must well comment your code, and follow good formatting practices. You should create assert statements for each function to test your code in VS Code before moving to CodeRunner.

### 2.1 Question 1: isValidBase()

One of the first steps when working with DNA sequences is to check whether the data is valid or corrupted. You must ensure the sequences you are working with consist only of valid DNA bases: A, C, G, and T. You will write a function, `isValidBase()`, that checks if a character is a valid DNA base.

<b>Function:</b> <code>isValidBase(char)</code>	<code>bool isValidBase(char base)</code>
<b>Purpose:</b>	The function will determine whether a given character is a valid DNA base. The function should not print anything.
<b>Parameters:</b>	<code>char base</code> - The character to validate.
<b>Return Value:</b>	The function should return true if the character is a valid base (A, C, G, or T) and false otherwise.
<b>Error handling/ Boundary conditions:</b>	<ul style="list-style-type: none"> <li>- The function should be case-sensitive, e.g., 'A' is a valid base, but 'a' is not.</li> <li>- <b>Note:</b> True is represented by 1 and false by 0 when you cout boolean variables.</li> </ul>

For Question 1, develop and validate your solution in VS Code. Once you are happy with your solution, go to coderunner on Canvas and paste only the `isValidBase()` to the answer box!

#### Sample Runs 2.1.1

Function call	Expected return value
<code>isValidBase('A')</code>	true
<code>isValidBase('X')</code>	false
<code>isValidBase('a')</code>	false
<code>isValidBase('T')</code>	true

### 2.2 Question 2: isValidStrand()

Next, write the `isValidStrand()` function that checks if a string contains only valid DNA bases.

<b>Function:</b> isValidStrand(string)	<b>bool</b> isValidStrand(string strand)
<b>Purpose:</b>	The function will determine whether a given string consists only of valid DNA bases. The function should not print anything.
<b>Parameters:</b>	<b>string strand</b> - The DNA strand to validate.
<b>Return Value:</b>	The function should return true if the string is a valid DNA strand and false otherwise.
<b>Error handling/ Boundary conditions:</b>	<ul style="list-style-type: none"> <li>- The input string is only considered valid if it consists only of A, C, T, and G bases.</li> <li>- If the string is empty, then it should not be considered a valid DNA strand, and your function should return false.</li> <li>- <b>Hint:</b> The function should make use of your isValidBase function.</li> </ul>

For Question 2, develop and validate your solution in VS Code. Once you are happy with your solution, go to coderunner on Canvas and paste the isValidBase() and isValidStrand() to the answer box!

#### Sample Runs 2.2.1

Function call	Expected return value
isValidStrand("ATGCTTCAA")	true
isValidStrand("CTTZ")	false
isValidStrand("")	false
isValidStrand("ATCG")	true

### 2.3 Question 3: strandSimilarity()

Comparing DNA sequences allows researchers to identify similarities and differences that may be significant in understanding genetic relationships or disease mechanisms. In this step, you'll develop functions to compare DNA strands and assess their similarity.

For two DNA strands of equal length, the similarity score is calculated based on the number of matching bases at corresponding positions using the following formula:

$$\text{Similarity} = \frac{\text{total matches}}{\text{total positions}}$$

**Example:** Let's compare the following two DNA strands:

Position	Strand 1	Strand 2
1	G	G
2	A	T
3	T	T
4	C	C
5	A	A
6	G	A

Table 7.11: Positions of two DNA strands

The total number of matches is 4 out of 6 positions, resulting in a similarity score of  $\frac{4}{6} = 0.667$ .

The function strandSimilarity() compares two strands position by position, counting the number of positions where the bases are identical. This provides a direct measure of how similar the two sequences are.

<b>Function:</b> strandSimilarity(string, string)	<b>double</b> strandSimilarity(string strand1, string strand2)
<b>Purpose:</b>	The function will find the similarity between two DNA strands. The function should not print anything.
<b>Parameters:</b>	<b>string strand1</b> - The first DNA strand to compare. <b>string strand2</b> - The second DNA strand to compare.
<b>Return Value:</b>	The function should return the similarity score between the two strands.
<b>Error handling/ Boundary Condition:</b>	- The parameters should be two strings of equal length. If they are not equal in length, your function should return 0. - You may assume that the input to strandSimilarity() will always be a valid strand, i.e., you do not have to account for arbitrary strings.

For Question 3, develop and validate your solution in VS Code. Once you are happy with your solution, go to coderunner on Canvas and paste the **strandSimilarity()** and any helper function(s) you used to the answer box!

#### Sample Runs 2.3.1

Function call	Expected return value
strandSimilarity("AGGT" , "CTGA")	0.25
strandSimilarity("CCTT" , "CCTT")	1
strandSimilarity("ATG" , "AAATTT")	0
strandSimilarity("CTGTAGAGCT" , "TAGCTACCAT")	0.2

## 2.4 Question 4: bestStrandMatch()

In bestStrandMatch(), the strands can be different lengths, therefore, you'll want to compare overlapping sections and calculate similarity scores at each position, and to do that you'll slide the shorter strand along the longer strand. The maximum score across all positions indicates the best alignment between the two strands.

<b>Function:</b> bestStrandMatch(string, string)	<b>int</b> bestStrandMatch(string input_strand, string target_strand)
<b>Purpose:</b>	The function will find the best similarity between two DNA strands. The function should print out the best similarity score.
<b>Parameters:</b>	<b>string input_strand</b> - The input DNA strand to be checked against the target_strand (length greater than or equal to the target strand) <b>string target_strand</b> - The target DNA strand.
<b>Return Value:</b>	If the parameters are valid, returns an <b>int</b> representing the starting index of the substring in the input strand where the best alignment with target strand occurs.

<b>Error handling/ Boundary conditions:</b>	<ul style="list-style-type: none"> <li>- If the input strand is shorter than the target strand, the function returns -1 as the alignment index and prints out "Best similarity score: 0.0".</li> <li>- This function should make use of the <code>strandSimilarity()</code> function.</li> <li>- You may assume that the input to <code>bestStrandMatch()</code> will always be a valid DNA sequence, i.e., you do not have to account for arbitrary strings.</li> </ul>
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For Question 4, develop and validate your solution in VS Code. Once you are happy with your solution, go to coderunner on Canvas and paste the `bestStrandMatch()` and any helper function(s) you used to the answer box!

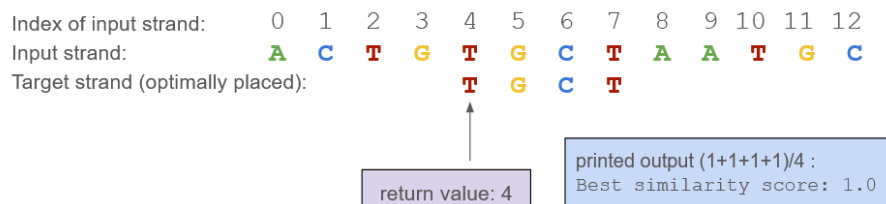


Figure 7.2: An illustration of `bestStrandMatch()` between two example strands

#### Sample Runs 2.4.1

Function call	Expected return value	Expected printed value
<code>bestStrandMatch("GATCAGT", "TCA")</code>	2	Best similarity score: 1.0
<code>bestStrandMatch("AACCTGAC", "ACT")</code>	1	Best similarity score: 0.666667
<code>bestStrandMatch("CTG", "CCCC")</code>	-1	Best similarity score: 0.0
<code>bestStrandMatch("ATCGTA", "TTCGAT")</code>	0	Best similarity score: 0.5

## 2.5 Question 5: Mutation Detection

Now, you want to be able to make deeper observations between DNA sequences. To do this, you will create a function that compares two DNA sequences and identifies all types of mutations between them. The function should align the sequences based on the best possible match and then process the sequences character by character, printing out mutations as they are detected.

Your function should be able to identify the following mutations:

- Substitution: When bases at the same position differ
- Insertion: When an extra base is present in the target strand
- Deletion: When a base from the input strand is missing in the target strand

The function should determine the longest of the two strands and use the `bestStrandMatch()` function to optimally align them. Upon alignment, the function should print out the best alignment index. After alignment, it should compare the sequences character by character to identify mutations. It detects substitutions

when bases at the same aligned position differ, deletions when extra bases are present in the input strand but not in the target strand, and insertions when bases are present in the target strand but missing from the input strand.

<b>Function:</b> identifyMutations(string, string)	<b>void</b> identifyMutations(string input_strand, string target_strand)
<b>Purpose:</b>	The function compares two DNA sequences to identify all types of mutations between them. It aligns the sequences based on the best possible match and processes them character by character, printing out any mutations as they are detected.
<b>Parameters:</b>	<b>string</b> input_strand - The input strand to be checked against the target <b>string</b> target_strand - The target strand
<b>Return Value:</b>	N/A
<b>Error handling/ Boundary conditions:</b>	-You may assume that the input and target strands are both valid DNA strands. - If no mutations are found, the function outputs "No mutations found."

For Question 5, develop and validate your solution in VS Code. Once you are happy with your solution, go to coderunner on Canvas and paste **identifyMutations()** and any helper function(s) to the answer box!

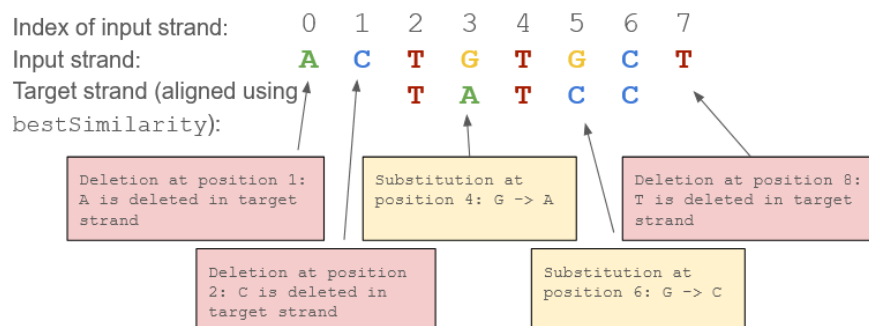


Figure 7.3: An illustration of identifyMutation() between two example strands

### Sample Runs 2.5.1

Note: "Best Similarity Score:..." print line should come from your bestStrandMatch() function.

Function call	Expected printed value
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identifyMutations("GGG", "TAGGGTA")	Best similarity score: 1 Best alignment index: 2 Insertion at position 1: T is inserted in target strand Insertion at position 2: A is inserted in target strand Insertion at position 6: T is inserted in target strand Insertion at position 7: A is inserted in target strand
identifyMutations("AACCGG", "AACCGG")	Best similarity score: 1 Best alignment index: 0 No mutations found.
identifyMutations("TA", "TAGG")	Best similarity score: 1 Best alignment index: 0 Insertion at position 3: G is inserted in target strand Insertion at position 4: G is inserted in target strand
identifyMutations("TAGG", "TA")	Best similarity score: 1 Best alignment index: 0 Deletion at position 3: G is deleted in target strand Deletion at position 4: G is deleted in target strand
identifyMutations("AGTCACG", "AGCTACA")	Best similarity score: 0.571429 Best alignment index: 0 Substitution at position 3: T -> C Substitution at position 4: C -> T Substitution at position 7: G -> A

## 2.6 Question 6: DNA Sequence Transformations

In this part, you will simulate the transcription process by converting a DNA sequence into an RNA sequence. This involves replacing every occurrence of thymine ('T') with uracil ('U') in the DNA strand.

<b>Function:</b> transcribedDNAtoRNA(string)	<b>void</b> transcribedDNAtoRNA(string strand)
<b>Purpose:</b>	The function will transcribe a DNA sequence to RNA and print the RNA sequence to the console. The function will replace all occurrences of 'T' with 'U'.
<b>Parameters:</b>	<b>string</b> strand - The DNA sequence to be transcribed.
<b>Return Value:</b>	N/A
<b>Error handling/ Boundary conditions:</b>	- You may assume that the input DNA strand is a valid DNA sequence.

For Question 6, develop and validate your solution in VS Code. Once you are happy with your solution, go to coderunner on Canvas and paste **transcribedDNAtoRNA()** and any helper function(s) to the answer box!

### Sample Runs 2.6.1

Function call	Expected printed value
transcribeDNAtoRNA("ATCGTACG")	AUCGUACG
transcribeDNAtoRNA("TTAA")	UUAA
transcribeDNAtoRNA("ACCG")	ACCG
transcribeDNAtoRNA("T")	U

## 2.7 Question 7: Reverse Complement of a DNA Sequence

In this part, you will write a function to compute the reverse complement of a DNA strand. The reverse complement is obtained by reversing the DNA sequence and then replacing each base with its complement:

- 'A' is complemented by 'T'
- 'T' is complemented by 'A'
- 'C' is complemented by 'G'
- 'G' is complemented by 'C'

<b>Function:</b> reverseComplement(string)	<b>void</b> reverseComplement(string strand)
<b>Purpose:</b>	The function will compute the <b>reverse</b> complement for a DNA sequence and print the result to the console.
<b>Parameters:</b>	<b>string strand</b> - The DNA sequence for which the reverse complement will be computed.
<b>Return Value:</b>	N/A
<b>Error handling/ Boundary condition:</b>	You may assume that the input DNA strand is a valid DNA sequence.

For Question 7, develop and validate your solution in VS Code. Once you are happy with your solution, go to coderunner on Canvas and paste the `reverseComplement()` and any helper function(s) you used to the answer box!

### Sample Runs 2.7.1

Function call	Expected printed value
reverseComplement("ATCGTACG")	CGTACGAT
reverseComplement("AAACCC")	GGGTTT
reverseComplement("AGCT")	AGCT
reverseComplement("A")	T

## 2.8 Question 8: Extracting Reading Frames

When working with DNA sequences, it is often necessary to focus on specific regions called open reading frames (ORF). These DNA regions are read in groups of three bases, called codons, which are later translated into amino acids during the process of protein synthesis. An ORF is a continuous sequence of DNA that begins with a start codon (ATG) and ends with a stop codon (TAA, TAG, or TGA).



Your task is to identify valid protein-coding regions in a DNA sequence. A valid ORF starts with the ATG codon and ends with one of the stop codons (TAA, TAG, or TGA), with the number of bases between the start and stop codons being divisible by 3.

<b>Function:</b> getCodingFrames(string)	<b>void</b> getCodingFrames(string strand)
<b>Purpose:</b>	The function will print out complete reading frames.
<b>Parameters:</b>	<b>string strand</b> - The DNA strand from which to extract reading frames.
<b>Return Value:</b>	N/A
<b>Error handling/ Boundary condition:</b>	- If no reading frames are found, the function should print "No reading frames found." - You may assume that the input DNA strand is a valid DNA sequence. - <b>Note:</b> There could be multiple ORF within a single DNA strand.

For Question 8, develop and validate your solution in VS Code. Once you are happy with your solution, go to coderunner on Canvas and paste the **getCodingFrames()** and any helper function(s) you used to the answer box!

#### Sample Runs 2.8.1

Function call	Expected printed value
getCodingFrames("ATGCGGTAA")	ATGCGGTAA
getCodingFrames("AAACCC")	No reading frames found.
getCodingFrames("ATGCGTAGCTAAATGGGGTAG")	ATGCGTAGCTAA ATGGGGTAG

## 2.9 Question 9: Tying It All Together

You have successfully written functions to help your research team analyze DNA sequences! However, now you want to make it more accessible and user-friendly. For this question, you will create a main function that allows a user to interact with your program by providing their DNA sequences. Your main function should present the user with a menu containing the following options:

1. Calculate the similarity between two sequences of the same length
2. Calculate the best similarity between two sequences of either equal or unequal length
3. Identify mutations
4. Transcribe DNA to RNA
5. Find the complement of a DNA sequence
6. Extract reading frames
7. Exit

Your menu should run on a loop, continually offering the user each option until they choose to exit. Be sure to use the functions you wrote in questions 1 through 6 as needed.

**Note:** Your main function should account for any user input that isn't a valid sequence. If user input is not a valid sequence, your program should print **"Invalid input. Please enter a valid sequence."** until

the user enters a valid sequence. Additionally, functions that require strings of the same length should have their inputs validated for matching lengths before being called in the user menu. If the strands are not of the same size, the program should display **"Error: Input strands must be of the same length."** and return to the menu.

**For Question 9, develop and validate your solution in VS Code. Once you are happy with your solution, go to coderunner on Canvas and paste you entire program to the answer box!**

#### Sample Run 2.9.1

```
--- DNA Analysis Menu ---
1. Calculate the similarity between two sequences of the same length
2. Calculate the best similarity between two sequences of either equal or unequal length
3. Identify mutations
4. Transcribe DNA to RNA
5. Find the reverse complement of a DNA sequence
6. Extract reading frames
7. Exit
Please enter your choice (1 - 7):
8
Invalid input. Please select a valid option.
--- DNA Analysis Menu ---
1. Calculate the similarity between two sequences of the same length
2. Calculate the best similarity between two sequences of either equal or unequal length
3. Identify mutations
4. Transcribe DNA to RNA
5. Find the reverse complement of a DNA sequence
6. Extract reading frames
7. Exit
Please enter your choice (1 - 7):
7
Exiting program.
```

#### Sample Run 2.9.2

```
--- DNA Analysis Menu ---
1. Calculate the similarity between two sequences of the same length
2. Calculate the best similarity between two sequences of either equal or unequal length
3. Identify mutations
4. Transcribe DNA to RNA
5. Find the reverse complement of a DNA sequence
6. Extract reading frames
7. Exit
Please enter your choice (1 - 7):
1
Enter the first DNA sequence:
ATC
Enter the second DNA sequence:
AG
Error: Input strands must be of the same length.
--- DNA Analysis Menu ---
1. Calculate the similarity between two sequences of the same length
2. Calculate the best similarity between two sequences of either equal or unequal length
3. Identify mutations
```

```
4. Transcribe DNA to RNA
5. Find the reverse complement of a DNA sequence
6. Extract reading frames
7. Exit
Please enter your choice (1 - 7):
```

```
Exiting program.
```

### Sample Run 2.9.3

```
--- DNA Analysis Menu ---
1. Calculate the similarity between two sequences of the same length
2. Calculate the best similarity between two sequences of either equal or unequal length
3. Identify mutations
4. Transcribe DNA to RNA
5. Find the reverse complement of a DNA sequence
6. Extract reading frames
7. Exit
Please enter your choice (1 - 7):
```

```
1
```

```
Enter the first DNA sequence:
```

```
AT
```

```
Enter the second DNA sequence:
```

```
AG
```

```
Similarity score: 0.5
```

```
--- DNA Analysis Menu ---
1. Calculate the similarity between two sequences of the same length
2. Calculate the best similarity between two sequences of either equal or unequal length
3. Identify mutations
4. Transcribe DNA to RNA
5. Find the reverse complement of a DNA sequence
6. Extract reading frames
7. Exit
Please enter your choice (1 - 7):
```

```
Exiting program.
```

### Sample Run 2.9.4

```
--- DNA Analysis Menu ---
1. Calculate the similarity between two sequences of the same length
2. Calculate the best similarity between two sequences of either equal or unequal length
3. Identify mutations
4. Transcribe DNA to RNA
5. Find the reverse complement of a DNA sequence
6. Extract reading frames
7. Exit
Please enter your choice (1 - 7):
```

```
2
```

```
Enter the first DNA sequence:
```

```
AGTC
```

```
Enter the second DNA sequence:
```

ATCG

Best similarity score: 0.25

--- DNA Analysis Menu ---

1. Calculate the similarity between two sequences of the same length
2. Calculate the best similarity between two sequences of either equal or unequal length
3. Identify mutations
4. Transcribe DNA to RNA
5. Find the reverse complement of a DNA sequence
6. Extract reading frames
7. Exit

Please enter your choice (1 - 7):

7

Exiting program.

### Sample Run 2.9.5

--- DNA Analysis Menu ---

1. Calculate the similarity between two sequences of the same length
2. Calculate the best similarity between two sequences of either equal or unequal length
3. Identify mutations
4. Transcribe DNA to RNA
5. Find the reverse complement of a DNA sequence
6. Extract reading frames
7. Exit

Please enter your choice (1 - 7):

3

Enter the first DNA sequence:

ATTTG

Enter the second DNA sequence:

ATCTG

Best similarity score: 0.8

Best alignment index: 0

Substitution at position 3: T -> C

--- DNA Analysis Menu ---

1. Calculate the similarity between two sequences of the same length
2. Calculate the best similarity between two sequences of either equal or unequal length
3. Identify mutations
4. Transcribe DNA to RNA
5. Find the reverse complement of a DNA sequence
6. Extract reading frames
7. Exit

Please enter your choice (1 - 7):

7

Exiting program.

### Sample Run 2.9.6

--- DNA Analysis Menu ---

1. Calculate the similarity between two sequences of the same length
2. Calculate the best similarity between two sequences of either equal or unequal length
3. Identify mutations
4. Transcribe DNA to RNA

```

5. Find the reverse complement of a DNA sequence
6. Extract reading frames
7. Exit
Please enter your choice (1 - 7):
4
Enter the DNA sequence to be transcribed:
ATTC
The transcribed DNA is: AUUC
--- DNA Analysis Menu ---
1. Calculate the similarity between two sequences of the same length
2. Calculate the best similarity between two sequences of either equal or unequal length
3. Identify mutations
4. Transcribe DNA to RNA
5. Find the reverse complement of a DNA sequence
6. Extract reading frames
7. Exit
Please enter your choice (1 - 7):
7
Exiting program.

```

#### Sample Run 2.9.7

```

--- DNA Analysis Menu ---
1. Calculate the similarity between two sequences of the same length
2. Calculate the best similarity between two sequences of either equal or unequal length
3. Identify mutations
4. Transcribe DNA to RNA
5. Find the reverse complement of a DNA sequence
6. Extract reading frames
7. Exit
Please enter your choice (1 - 7):
5
Enter the DNA sequence:
ATTCG
The reverse complement is: CGAAT
--- DNA Analysis Menu ---
1. Calculate the similarity between two sequences of the same length
2. Calculate the best similarity between two sequences of either equal or unequal length
3. Identify mutations
4. Transcribe DNA to RNA
5. Find the reverse complement of a DNA sequence
6. Extract reading frames
7. Exit
Please enter your choice (1 - 7):
7
Exiting program.

```

#### Sample Run 2.9.8

```

--- DNA Analysis Menu ---
1. Calculate the similarity between two sequences of the same length
2. Calculate the best similarity between two sequences of either equal or unequal length

```

```

3. Identify mutations
4. Transcribe DNA to RNA
5. Find the reverse complement of a DNA sequence
6. Extract reading frames
7. Exit
Please enter your choice (1 - 6):
6
Enter the DNA sequence:
ATGCGGTAA
The extracted reading frames are:
ATGCGGTAA
--- DNA Analysis Menu ---
1. Calculate the similarity between two sequences of the same length
2. Calculate the best similarity between two sequences of either equal or unequal length
3. Identify mutations
4. Transcribe DNA to RNA
5. Find the reverse complement of a DNA sequence
6. Extract reading frames
7. Exit
Please enter your choice (1 - 7):
7
Exiting program.

```

### Sample Run 2.9.9

```

--- DNA Analysis Menu ---
1. Calculate the similarity between two sequences of the same length
2. Calculate the best similarity between two sequences of either equal or unequal length
3. Identify mutations
4. Transcribe DNA to RNA
5. Find the reverse complement of a DNA sequence
6. Extract reading frames
7. Exit
Please enter your choice (1 - 7):
6
Enter the DNA sequence:
AGCTTTAA
The extracted reading frames are:
No reading frames found.
--- DNA Analysis Menu ---
1. Calculate the similarity between two sequences of the same length
2. Calculate the best similarity between two sequences of either equal or unequal length
3. Identify mutations
4. Transcribe DNA to RNA
5. Find the reverse complement of a DNA sequence
6. Extract reading frames
7. Exit
Please enter your choice (1 - 7):
7
Exiting program.

```

### Sample Run 2.9.10

--- DNA Analysis Menu ---

1. Calculate the similarity between two sequences of the same length
2. Calculate the best similarity between two sequences of either equal or unequal length
3. Identify mutations
4. Transcribe DNA to RNA
5. Find the reverse complement of a DNA sequence
6. Extract reading frames
7. Exit

Please enter your choice (1 - 7):

1

Enter the first DNA sequence:

ATRBAD

Invalid input. Please enter a valid sequence.

Enter the first DNA sequence:

ATTA

Enter the second DNA sequence:

ATTAx

Invalid input. Please enter a valid sequence.

Enter the second DNA sequence:

ATCG

Similarity score: 0.5

--- DNA Analysis Menu ---

1. Calculate the similarity between two sequences of the same length
2. Calculate the best similarity between two sequences of either equal or unequal length
3. Identify mutations
4. Transcribe DNA to RNA
5. Find the reverse complement of a DNA sequence
6. Extract reading frames
7. Exit

Please enter your choice (1 - 7):

7

Exiting program.