# Indian Institute of Technology Gandhinagar



## **Project-Assignment-1**

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Branch: Biological Sciences & Engineering

**Subject:** Biocomputing (BE623)

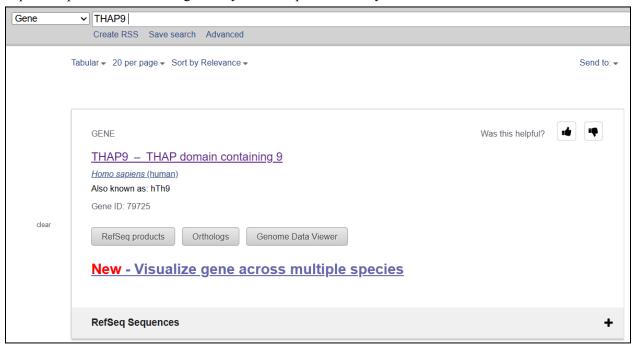
**Submission Date: 19/09/2025** 

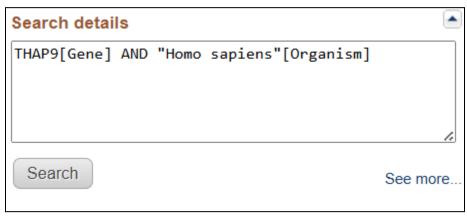
## **Question 1 (5 Points)**

Choose a gene of your interest from human (*Homo sapiens*) and perform the following: a) Mention the gene name and some description and search for this gene at NCBI. Refine the query using the methods discussed in the lab sessions and download the gene sequence as fasta file (Take screenshots to show the search) (1 Point)

**Gene Name:** THAP domain containing 9 (THAP9)

**Description:** THAP9 is a transposable element-derived gene that encodes the THAP9 protein, which is homologous to the Drosophila P-element transposase and can cut and paste DNA. It enables sequence-specific DNA binding activity and transposase activity.



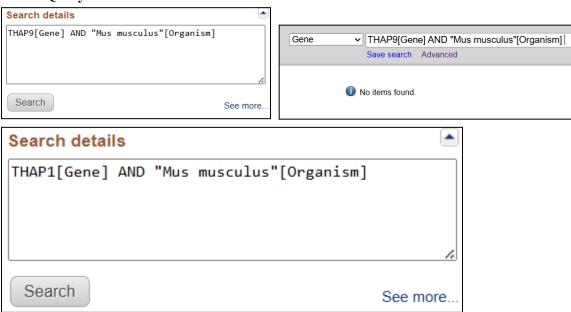


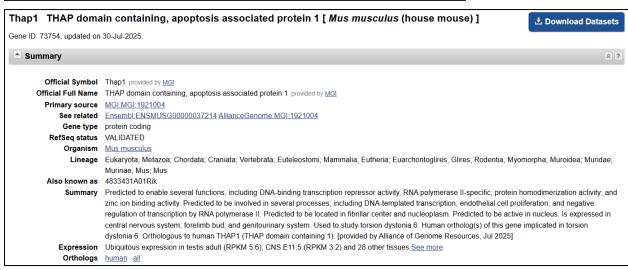


NC\_000004.12:82900735-82919969 Homo sapiens chromosome 4, GRCh38.p14 Primary Assembly GGAGCTAAAGTGGTCGTGATTCATGCTGTCGCGGGAACCCCGAAGGTGGGGCCCCACGTAACAAGAAGAT GACCCGAAGTTGCTCCGCAGTGGGCTGCAGCACCCGTGACACCGTGCTCAGCCGGGAGCGCGGCCTCTCC TTCCACCAGTGCGTATGGGAGCAGCCTCGAAGCCTTCGAACTCCCTGCGGGGCCCGGCGGGCCGTGGCGT GGCGTGGGGCGGGCCGGGCCGCACTGTGGGTCGCGCCGCGTGTGTGACGCGACGTGACGTGCGCAGCGT CTGAATAGCCGGTTCTCGCACCGCCTACCGCTTTAAGGAGAGTTACGCGAAAGGAGTCCTCCATTAATTG GTGGCGTCTTCCTGAGCACGAGTACAGATCAGTTTTCTCCTGTACTGCCTGACATTGCAAGTTCTTAAGT GGCTCGACAGTGGGGAAGAATTTCAGATCAGTCTTATCTGAACTCATATCCATGGTTCAGACTGGTATAG GTCATTCAGTATTCAGGGTACACTAGCAAGTAGAAAAGTGGGTAAAATAGATCCTCACAGACAAAAGGAG CAGATTTATGGAAAGATCAAAAAGAACTTTATGGTTGGAAAGGGGGGGTAGGAAGAGGGTAGTTAGGAAT GGCATGGGAAAATGTAAAGGATAGAAAGTGCAAATATAGAGGAGTGGAAAAGTACCTGAGAAACCATTTT TTTTTTTTTTTGTTTTCTATTTTTTAAATAGGGCTGATTCCTCCACCTAAATTGTTAGCAGTGTGAACCA ACTCATTGTGTATGTGAGACCATAATTGATTTGGGATTTTTCTTGTGATAGAGGTAGTTTTAAACTAGGG AGACATGAATTCCAATTTTAATTGAGTATTTACTGATGTCTGTACTTGGCTCTGGGTCTTGGTTTCTTCT TCTGTAAGAGGAATAAAAACCTGCCTTGTAGAATTTTGTAAAACCTAAAACGTAAGACTTTATAGCATATG GTCTCACTTTGTCGCCCAGGCTGGAGAGCAGTGGCACTATCACAGCTCAGCGCAGCCTCTATCTCCCAGG

b) Find the orthologous (similar) gene from Mouse (Mus musculus), Rat (Rattus norvegicus), Chimpanzee (Pan troglodytes) and Fruit fly (Drosophila melanogaster) and download the sequences as fasta file. (1 Point)

For Mouse(*Mus musculus*), we didn't find any gene named THAP9. So we searched for a gene named THAP1 in the Mouse, i.e, the orthologous of the THAP9 gene.

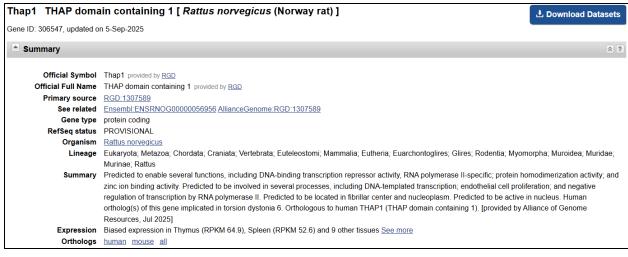




>NC 000074.7:26648197-26654179 Mus musculus strain C57BL/6J chromosome 8, GRCm39 CGGCGAGCCGAGTCCTCGCGAGAGGTGAGGCTGAGCTGCCTCCGCCATCTTGAAGATGGGAGACGGGCG ACAGCTGTGGTCCTTCTGCTAAAGCAAACCCCACAACGGACAGGGTAGTCACTCCCCCACCCCAACCCCC ACCCCACGGCGAGGTGATCGTCCCCGTAACTGCTGACCGACGCCACCGAGAGCGGCGAGCGTTATCAAGG CCGAGCGCGGCGACCCCGACGGCCCCTTCGCCTGCCTCCCGGGCCGAAGGGAAGTGTGGAGGGCCAGAA GGATGGTGCAGTCCTGCTCCGCCTACGGCTGCAAGAACCGCTACGACAAGGACAAGCCCGTCTCCTTCCA CGCGTGGCCGCGCGAGACCTGGAGGAGGCGGGGGGGGGCGCTCGGGCGCTCAGGGCGGCCGGACGGGCCGGG CGGCGTCTCCGTTGGTCTTGGCCTCCGCTGGGGGCTGAGACCGCGCCCTTCGCGCTCGCGGAGGGTTTC CGGCGGGCGGCAAGCGAGCGACGCCCAGAAGCCTCTCGCAGCCCTGCGTACCTCTTCGTCCCAAG CCGCGGCGTGGCCGTGGCACATAGGGAGGAAGTCGAGCTGGGAACGCACAGGGGGGAGGACAGGTTTC GGACCTCACTGTGGATGCCGGGAAGGAAGCAGGGCAGCTGACCGGGATGCCAGGGAGGACTCTTAAGTCC TAGCGTCAAAGAACCAACCTTTAGCATCCATGGGCTTTAGAGGAGGCTCTTGGTATTGTCGAGATGAACC CCTGTCTTTCAACACACAGGAAGAACGAGACGATAGCACACACTTGACTCTGTTGCCGTGGTGAAGGTTA ATGTGAGCGAATACCAGCTTTACCCCCACAGCCGTCAGGGCTCGTTCTTGTAGTGGGTGCAGCAATCAAG CTGAGTTCAATTCCCAGCAACCACATGGTGGCTCACAACCATCCGTAATGAGAAATGACGCCTTCTTCTG AATGTGACAATTTTTAGGCAGGTAAATTTGTTGTTACAAGTTTTATGTTAGGTTTTGGAGGAATTCCTGA CTGTATGCAGATGATAGCTAAGAAATCAGATCCACTGGTAAATTATGCAAGTTATCCATAGCAAAACGAT

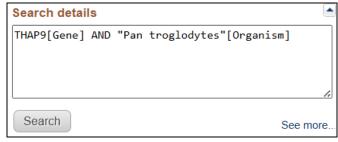
For Rat (*Rattus norvegicus*), we didn't find any gene named THAP9. So we searched for a gene named THAP1 in the Rat, i.e, the orthologous of the THAP9 gene.



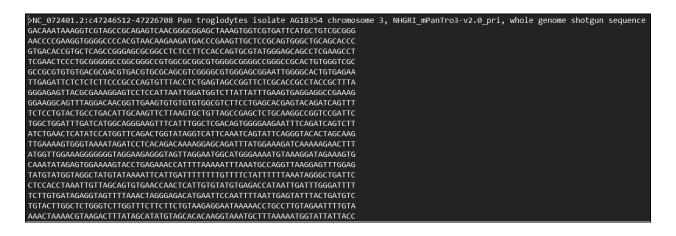


>NC\_086034.1:c72612690-72608096 Rattus norvegicus strain BN/NHsdMcwi chromosome 16, GRCr8, whole genome shotgun sequence GTTGAAGCTGCCTCCGCCATCTTGAAGATGGGAGACGAGCGACAGCTGTTGTCTTTCTGCTAAAGCAAAC ACCACAACGGACCGGGTGGTCACCCCCCACGGTGAGGTAATCGTCCCCATAACTGCTAACCAAAGCCACC GAAAGCGGCGACCGTATCAAAGCCGAGCGCGGCGCGCACCGACGGCCCCTTCACCTGCCTCCCGGGCAGAG GGGAAGTGTGGAGGGCCAGAAGGATGGTGCAGTCCTGTTCCGCCTACGGCTGCAAGAACCGGTACGATAA GGACAAGCCCGTCTCCTTCCACAAGTAGGTAACTGGCGTGCTCCGCGGCCGCGGCAGGGGCGCGGGGGG CCTGGAAGCCTCTCGCTGCCCTCCGGATTTCGTCCCAAGCCAAGCCGTGGAAATGTCCCATAGGGATAAA CTGGAACTAGAAAAGCATGGGAGGGAGACCTGGCTTCTGACCTCGCTGCTGCTGTTAGGAGGGAAGCAGG ATAGCTGAGTGTTGACATCTGACCAGGACTTTTAGACCTGGTTTCAAAGAACCAAGCTTTAAGGATCTGT GGGTTTTAGAGGTGACTCTTGGTATTGCCAAGATGAGCCCCAGTCTTTCAACACACGGTAGTAATGAAAC GATAGCATTCACTTTACTCAGATGCCCTGGGAAAGGTTGATGGAACTAATGTTAGCTTTCCCCCACAGCT GTCAGGCCTTGTTCTTGTGTGTGCAGCAAGCAAGAATCCAGGATGAAAAATACTCTCTTGAAAGAAGAAG GCAATAGAATTAACGTTACTGATGGTGATTATTAGCAATGCTAACGGTCCACAAAATGTGACATACTTTT

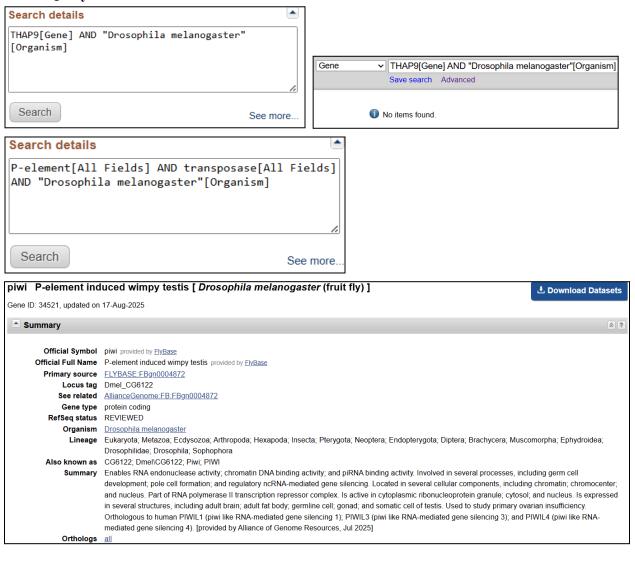
For the Chimpanzee (*Pan troglodytes*), we found a gene named THAP9, which is the orthologous of THAP9 in humans.







For the Fruit fly (*Drosophila melanogaster*), we didn't find any gene named THAP9. So we searched for a gene named piwi (P-element transposes) in the Fruit Fly, i.e, orthologous to THAP9 in humans.



>NT 033779.5:c10987420-10982205 Drosophila melanogaster chromosome 2L TCACACTGAGTCCAAAGCGTCGTTTTCAAAGTACTCTTTCAGTTTCCATTGTGAAGTTTTAAGTGATCGC GAGTGCCAAAAAGTAACAATGGCTGATGATCAGGGACGTGGACGCAGGCGTCCACTTAACGAAGATGATT AGCGGTAAAGCAAATTCCTTATAATTCCGTTATGTCGCAAACACAAACACAGCCATGTACATAGATTGTA AATTTGAGCTTGACTGCATAAAATCTTCTTGAATTTCTGTTGACCAAGCACTTTTTTCAAAGGGAATTCA TTAAACATATTTCCAAAAACGCTTCTTTCAATTTGAATTTCCAAGGAAATGTAAAGCATTATCTGTTTCG TTTCATTTATTTTTTTAATAAAAACTGCTAAAATAACCATCACCTGACAGGAAGGTGGCTGTGGCAAG TACATTTGATTCCTAGAAACTCTACGAGTAACGGGTATTTCTCCGATATTCATCCCAATTCTATACAACT TTGAGATGGCTTTCAGCATTGATATACTCTGAAATTGCAGGCAAAGCATACTACAAAAACTATTTTTTAC CATTTTTTTTCTGCTTCATCTTTGATTATGAGTAACAATATTTTCGTAGCTTGACCGTTAGTTTTATGAT TTAAAATCTGTCGTACTTTAGGTAGTTGTTTAAATGGCTCACATTTAGGAATCACAACTATTTGTTTAAA CCCCTTAATATTTTTCTTTTAATAATAAATAATAAACAGGTAATACTTTTTGATTGTTCATCGCGATTG TTTTCAATGTACAAAGTTGAGTCTTTCTGGTGTTTATTGCTGTCTCGTGTTTATTGCTGAATCTCGCACG ATCTCTCTCTTTCTATGTTATTTTTTCCATCGGCGTCAATTATTCATAACTGCCGCTAGGTGGAGCT AGTAACTTACATCATTTAATGCTTTCATGCAGCGGGTGAAAGTATTCAGAGGATCTTCATCAGGTGACCC GGCCCGACAGAGCGAAAGCCGTGGGGTGACCAATATGATTACCTGAATACCCGTCCGGCTGAGCTGGTAT CCAAGAAGGGAACCGATGGCGTCCCGGTCATGCTGCAGACGAACTTTTTCCGATTAAAAACCAAGCCGGA

## c) For each of these five sequences, extract the header and save in a separate file (1 Point)

`>/p' THAP9\_human.fasta THAP9\_chim panzee.fasta THAP1\_rat.fasta THAP1\_mouse.fasta piwi\_fruit\_fly.fasta > header\_files.txt

(base) intern@rosalind:~/Nabendu/Biocomputing\_Assignment/Project\_assignment\$ vi header\_files.txt

```
>NC_000004.12:82900735-82919969 Homo sapiens chromosome 4, GRCh38.p14 Primary Assembly
>NC_072401.2:c47246512-47226708 Pan troglodytes isolate AG18354 chromosome 3, NHGRI_mPanTro3-v2.0_pri, whole genome shot
gun seguence
>NC_086034.1:c72612690-72608096 Rattus norvegicus strain BN/NHsdMcwi chromosome 16, GRCr8, whole genome shotgun sequence
>NC_000074.7:26648197-26654179 Mus musculus strain C57BL/6J chromosome 8, GRCm39
>NT_033779.5:c10987420-10982205 Drosophila melanogaster chromosome 2L
```

So in this step, we extract five header lines and save them to a .txt file named header files.txt

## d) calculate the GC content of each sequence (2 Points)

```
cbase) Intern@rosalind:~/Nabendu/Biocomputing_Assignment/Project_assignm
sub(/[GC]/," "); total+=length($0)} END {print "GC% =", (gc/total)*100}'
GC% = 36.9743
(base) intern@rosalind:~/Nabendu/Piccomputing
                                                                                                                                  nt$ sed '/^>/d' THAP9_human.fasta | awk '{gc+=g
(base) intern@rosalind:~/Nabendu/Biocomputing_Assignment/Project_assignment$ sed '/^>/d' THAP9_chimpanzee.fasta | awk '{gc+=gsub(/[GC]/," "); total+=length($0)} END {print "GC% =", (gc/total)*100}'
 30\% = 36.8543
(base) intern@rosalind:~/Nabendu/Biocomputing_Assignment/Project_assignment$ sed '/^>/d' THAP1_rat.fasta | awk '{gc+=gsu b(/[GC]/," "); total+=length($0)} END {print "GC% =", (gc/total)*100}' GC% = 44.0261
(base) intern@rosalind:~/Nabendu/Biocomputing_Assignment/Project_assignment$ sed '/^>/d' THAP1_mouse.fasta | awk '{gc+=g sub(/[GC]/," "); total+=length($0)} END {print "GC% =", (gc/total)*100}'
GC% = 45.3953
(base) intern@rosalind:~/Nabendu/Biocomputing_Assignment/Project_assignment$ sed '/^>/d' piwi_fruit_fly.fasta | awk '{gc +=gsub(/[GC]/," "); total+=length($0)} END {print "GC% =", (gc/total)*100}'
```

Here, I have calculated the GC content of each sequence using sed and awk commands.

| Gene Name          | % GC content |
|--------------------|--------------|
| THAP9 (Human)      | 36.97        |
| THAP9 (Chimpanzee) | 36.85        |
| THAP1 (Rat)        | 44.03        |
| THAP1 (Mouse)      | 45.40        |
| piwi (Fruit Fly)   | 39.63        |

## **Question 2 (15 Points)**

For the same chosen gene:

a. Use the coding sequence (CDS) of this gene to translate to amino acid sequence. You may reduce the gene length to only 30 nucleotides for ease. (Hint: use Genetic code table and search/replace, Coding sequence can be downloaded from NCBI) (3 points)

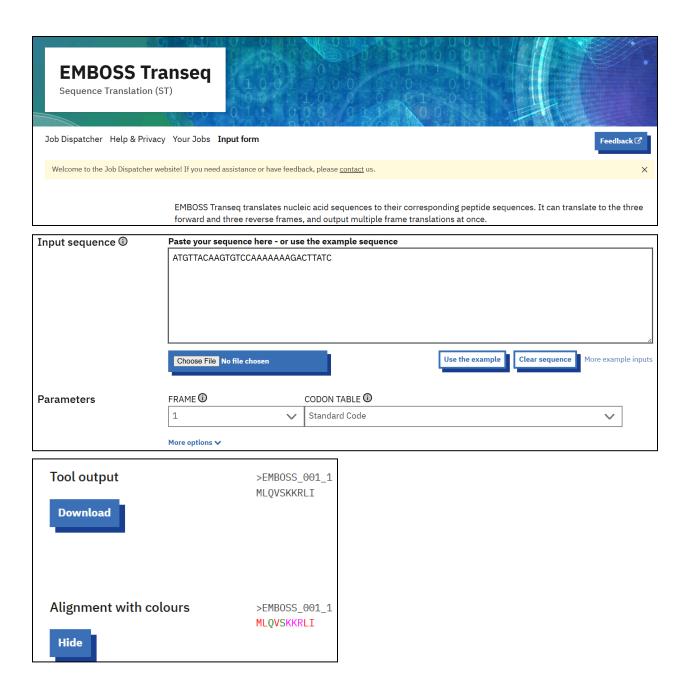
### **Coding sequence:**

(base) intern@rosalind:~/Nabendu/Biocomputing\_Assignment/Project\_assignment\$ sed '/^>/d' cds.fna | tr -d '\n' | cut -c1-30 ATGTTACAAGTGTCCAAAAAAAAAGACTTATC

So I got this sequence,

#### ATGTTACAAGTGTCCAAAAAAAAGACTTATC

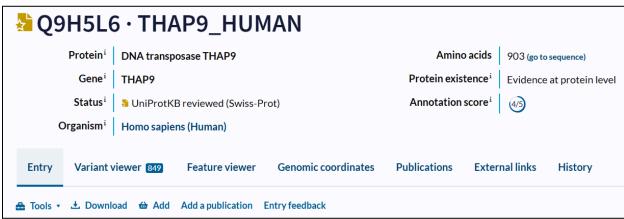
To translate it into an amino acid sequence, I use EMBOSS Transeq. It uses the standard genetic code to code the amino acids.

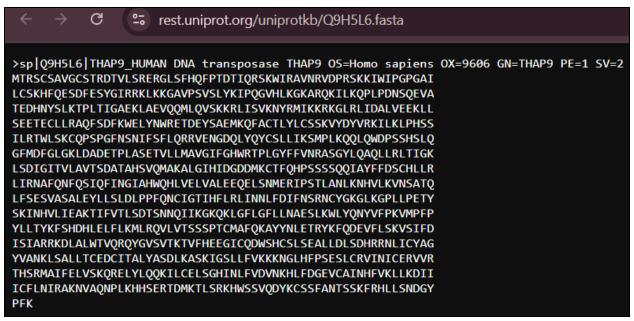


So we got these amino acid sequences: MLQVSKKRLI, from EMBOSS Transeq as an output.

b. Retrieve the protein sequence for this gene from UniProt in FASTA format and compare the sequence of gene translated by you with that retrieved from Uniprot. (2 points) (Hint: You may use bash commands to compare files or strings)

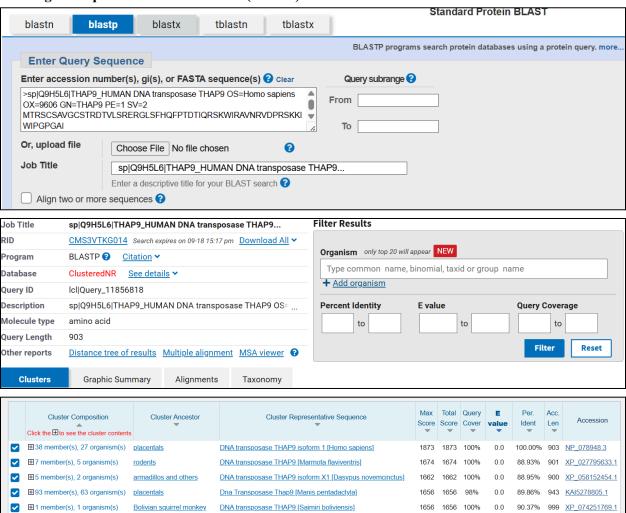






So from Uniprot, the first 10 amino acid sequences are MTRSCSAVGC. So it is not matched with the sequence translated by me. I think the probable reason may be the differences in the gene annotation format between NCBI and Uniprot.

## c. Use the protein sequence from Uniprot to run a BLASTP search at NCBI and download the top 5 homologous sequences as a FASTA file. (1 Point)



So I have downloaded the top 5 homologous sequences from running a blastp of THAP9.

## d. From this combined FASTA file, extract all the sequence IDs and save them in a separate file. (1 point)



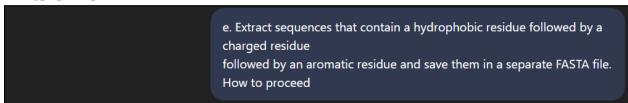
Here, first I print all the sequence IDs using awk, and then save them to a separate file named sequence id5.txt.

e. Extract sequences that contain a hydrophobic residue followed by a charged residue followed by an aromatic residue and save them in a separate FASTA file. (2 points)

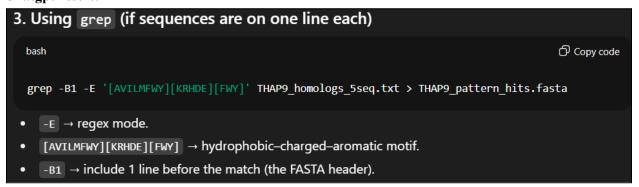
So hydrophobic residues in the protein sequence are Alanine(A), Valine(V), Isoleucine(I), Leucine(L), Methionine(M), Phenylalanine(F), Tyrosine(Y), and Tryptophan(W).

Charge residues are Arginine (R), Histidine(H), Lysine(K), Aspartic Acid(D), and Glutamic Acid(E). Aromatic residues are Phenylalanine(F), Tryptophan(W), Tyrosine(Y).

### **Chatgpt prompt:**



### **Chatgpt result:**



Using Chatgpt I know the use of -E, -B1, and how I write a hydrophobic-charged-aromatic motif. So, find these motifs using grep search.

KKGAVPSVSLYKIPQGVHLKGKARQKILKQPLPDNSQEVATEDHNYSLKTPLTIGAEKLAEVQQMLQVSKKRLISVKNYR MIKKRKGLRLIDALVEEKLLSEETECLLRAQFSDFKWELYNWRETDEYSAEMKQFACTLYLCSSKVYDYVRKILKLPHSS ILRTWLSKCQPSPGFNSNIFSFLQRRVENGDQLYQYCSLLIKSMPLKQQLQWDPSSHSLQGFMDFGLGKLDADETPLASE TVLLMAVGIFGHWRTPLGYFFVNRASGYLQAQLLRLTIGKLSDIGITVLAVTSDATAHSVQMAKALGIHIDGDDMKCTFQ HPSSSSQQIAYFFDSCHLLRLIRNAFQNFQSIQFINGIAHWQHLVELVALEEQELSNMERIPSTLANLKNHVLKVNSATQ LFSESVASALEYLLSLDLPPFQNCIGTIHFLRLINNLFDIFNSRNCYGKGLKGPLLPETYSKINHVLIEAKTIFVTLSDT SNNQIIKGKQKLGFLGFLLNAESLKWLYQNYVFPKVMPFPYLLTYKFSHDHLELFLKMLRQVLVTSSSPTCMAFQKAYYN LETRYKFQDEVFLSKVSIFDISIARRKDLALWTVQRQYGVSVTKTVFHEEGICQDWSHCSLSEALLDLSDHRRNLICYAG YVANKLSALLTCEDCITALYASDLKASKIGSLLFVKKKNGLHFPSESLCRVINICERVVRTHSRMAIFELVSKQRELYLQ

KKGAVPSVSLYKIPQGVHLKGKARQKILKQPLPDNSQEVASEDHNYSLRRPWTGGAEKLAEVQQMLQVSKRRLISVKNYR IIKKRKGLQLIDSLVEERLLSEETECLLRAQFSDFNWEMYNWRETAEYSTEMKQFACTLYLCSSKVYDYVRKILKLPHSS ILRTWLSRCKPSPGFNSNIFSFLQQRVENGDQIYQYCSLIIKGISLKEQLQWDPSSNSLQGFMDFGLGKLDADETPLASE TILLMAVGIFGHWRTPLGYFFVNRTSGYLQAQLLRLTIGKLSDIGITVLAVTSDATAHSVQMAKALGIHIDGDNMKCTFQ HPSSSSQQIAYFFDSCHLLRLIRNAFQNFQSIQFINGTAHWQHLVELVALGEQELSYMKGIPRKHANLKNHVLKMNCAAQ LFSESVASALEYLLSVGLPPFQNCIGTIHFLRLINNLFDVFNSRNCYGKGLKGPLLPETFSKINHVLIEAKTIFLTLSDT SNNQILKGKRKLGFLGFLLNAESLKWLYQNYVFPKVMPFPYLLTYKFSQDHLELFLKMLRQVLVPSSSFTCMAFQKAYHN LETRYRFQDEVFLSEVSIFDISVARRKDLALWTVQRQYGVRVMKSLFHKEDICQDWSNCSLSEALLDLSDQKRSLTCYAG YIATRLSAILTCEDCISALYASDLKASKIGSLLCVKKKNGLHFPSESLCQIINICEQVLRTHSRMVVFELVPKHRELYLQ

MTRSCSAVGCSTRDTVLSRERGLSFHQCVFPTDTIQRAKWIRAVNRVDPRSKKIWIPGPGAILCSKHFQESDFESYGIRR KLKKGAVPSVSLYKVLQAADLKGKARQKILKQPLPDNSQEVATEDHNYSLKTPLIIGAEKLAEVQQMLQVSKKRLISVKN YRLIKKRKGLRLIDALVEEKLLSEETEYLLRAQFSDFKWELYNWRETAEYSTEMKQFACTLYLCGSKVYDYVRKILKLPH SSILRTWLSKCKPSPGFNSHIFSFLQQRVENGDQLYQYCSLIIKGISLKQQLQWDPSSHRLQGFMDFGLGKLDADETPLA SETILLMAVGIFGHWRTPLGYFFVNRSSGYLQAQLLRLTIGKLSDIGITVLAVTSDATAHSVQMAKALGIRIDGDNVKCT FQHPSSSSQQIAYFFDSCHLLRLIRNAFQYFQSIQFINGIAHWQHLVELVALKEQELSSTERISSKFPKLKNHVLKMNCA AQLFSESVASALECLLSLGLPPFQNCIGTIHFLRLINNLFDVFNSRNYYGKGLKGPLLPETYNKINHVLIEAKTIFVTLS DTSNNQIIKGKRKLGFLGFLLNAESLKWLYQNYVFPKVMPFPYLLTYKFSQDHLELFLKMLRQVLVSSSSPTCMAFQKAY HNLETRYRFQDKAFLSEVSILDISIARRTDLALWTIQHEYGGSIIKALFHKEVICQDWSNCSLSEALLDLSDHRRNLTYY AGYIATKLTALLTCEECITALYASDLKAFKIGSLLCVKKKNGLHFPSESLCRIINICERVVRTHSRMGIHELVPKQMDLY

KKGAVPSVSLYKVLQGVHLKGKARQKILKQPLPDNSPEFATEDHNYSLKRPLTIGAEKLAEVQQMLQVSKKRLVSVKNYR MIKKRKGLRLIDALVEEKLLSEETECLLRAQFSDFKWELYNWRETAEYSTEMKQFACTLYLCSSKVYDYVRKILKLPHSS ILRTWLSKCKPSPGFNSNIFSFLQQRVENGDQLYQYCSLIIKGISLKEQLQWDPSSHHLQGFMDFGLGKLDADEMPLASE TILLMAVGVSGHWRTPLGYFFVNRASGYLQAQLLRLTIGKLSDIGITVLAVTSDATAHSVQMAKALGIHIDGDNMKCTFQ HPSSSSQQIAYFFDSCHLLRLIRNAFQNFQSIQFINGTAHWQHLVELVALEEQELANMERIPRKLANLKNHVLKMNFAAQ LFSESVASALECLLSLGLPPFQNCIGTIHFLRLINNLFDVFNSRNCYGKGLKGPLLPETFSKINRVLIEAKTIFVTLSDT SNNQIIKGKRKLGFLGFLLNAESLKWLYQNYVFPKVLPFPYLLTYKFSQDHLELFLRQVLVNSSNPTCLAFQKAYHN LETRYRLQDETFLSEVSILDISIARRTDLALRTVQRQYDVSIIKTLFHKEDFCQDWSNCSLSEALLDLSDHMRNLNCCAG YIANKLSALLTCEDCISALYASDLRASKIGSLLCVKKKNGLHFPSESLCRVINICERVLRTHSRMAVYELLPKQRELYLQ

MVGRELPPYFCIGKLQDKAQEHDTHLKGGAVKYTSSGLGPSDWLLRQNVACDWLQCRRRAFGEEVWESRASKARLSCQIK

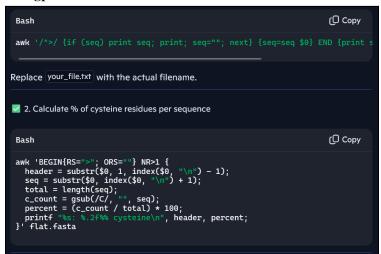
PLTVGAEKLAEVQQMLQVSKKRLISVKNYKMIKKRKGLPLIDALVEEKLLSEETECLLRAQFSGAASEAGGKPGDCAGLE AKCRKYFKEGVVHYVKCCWEIRWLSKCQPGPGFNSHIFSFLQRRVQNGDQLYQYCSLIIKSIPLKQQLQWDPSSHSLQGF MDFGLGKLDADETPLATETVLLMAVGVFGHWRTPLGYFFVNRASGCLQAQLLRLTIGKLSDIGITVLAVTSDATAHSVQM AKALGIHIDGDDMKCTFQHPSSCSQQIAYFFDSCHLLRLIRNAFQNFQSIQFINGIAHWQHLVELVALEEQELSNMERIP STPANLKNHILKVNCAAQLFSESVASALEYLLSLGLPPFQNCIGTIHFLRLINNLFDIFNSRNCYGKGLKGPLLPETYSK INHVLIEAKTIFVTLSDSSNNQIIQGKQKLGFLGFLLNAESLKWLYQNYVFPKVMPFPYLLTYKFSHDHLELFLKMLRQV LVTSSSPTCMAFQKAYHNLETRCTFQDEVFLSKVSIFDISVARRKDLALWAVQRQYGVSVTKTLFHKEGICQDWSNCSLS EALLDLSDNRRNLICYAGYVADKLSALLTCEDCITALYASDLKASKIGSLLFVKKKNGLHFPSESLCQVINICERVLRTH

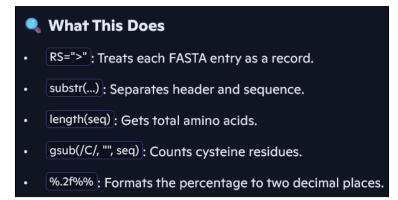
Here, I searched the motifs using grep search in the entire THAP9 5 homologs sequence and saved it to a text file named THAP9 hca.txt.

# f. Calculate the percentage of cysteine residues in each sequence and report it. (2 points) Chatgpt Prompt:



### **Chatgpt Result:**





In this question, I use chatgpt to understand how to proceed with the question step by step. So, for this, I know about the RS, substr(), and other things that I knew before.

```
(base) intern@rosalind:~/Nabendu/Biocomputing_Assignment/Project_assignment$ awk '/^>/ {if (seq) print seq; print; seq="
"; next} {seq=seq $0} END {print seq}' THAP9_homologs_5seq.txt > THAP9_oneline.fasta

(base) intern@rosalind:~/Nabendu/Biocomputing_Assignment/Project_assignment$ awk 'BEGIN{RS=">"; ORS=""} NR>1 {header = s
ubstr($0, 1, index($0, "\n") - 1); seq = substr($0, index($0, "\n") + 1); total = length(seq); cys_count = gsub(/c/, "",
seq); percent = (cys_count / total) * 100; printf "%s: %.2f% cysteine\n", header, percent;}' THAP9_oneline.fasta
NP_078948.3 DNA transposase THAP9 isoform 1 [Homo sapiens]: 2.65% cysteine
XP_027795633.1 DNA transposase THAP9 [Marmota flaviventris]: 2.77% cysteine
XP_088152454.1 DNA transposase THAP9 isoform X1 [Dasypus novemcinctus]: 2.66% cysteine
KAI5278805.1 Dna Transposase ThAP9 [Manis pentadactyla]: 2.75% cysteine
XP_074251769.1 DNA transposase THAP9 [Saimiri boliviensis]: 3.40% cysteine
```

| Sequence Header Name   | % Cysteine Residue |
|--|--------------------|
| NP_078948.3 DNA transposase THAP9 isoform 1 [Homo sapiens]             | 2.65               |
| XP_027795633.1 DNA transposase THAP9 [Marmota flaviventris]            | 2.77               |
| XP_058152454.1 DNA transposase THAP9 isoform X1 [Dasypus novemcinctus] | 2.66               |
| KAI5278805.1 Dna Transposase Thap9 [Manis pentadactyla]                | 2.75               |
| XP_074251769.1 DNA transposase THAP9 [Saimiri boliviensis]             | 3.40               |

In this question, I have calculated the percentage of cystine residue in each sequence using the awk command and printed the header-wise cysteine percentage in each sequence.

g. From the BLAST results, find the longest sequence and report its ID and its length. For this sequence, calculate the amino acid frequency and provide a distribution of each residue type. (4 points)

```
(base) intern@rosalind:~/Nabendu/Biocomputing_Assignment/Project_assignment$ awk '/^>/ {if (len > max) {max=len; id=hdr} hdr=$0; len=0} /^[^>]/ {len+=length($0)} END {if (len > max) {max=len; id=hdr} print id, max}' THAP9_homologs_5seq.txt >XP_074251769.1 DNA transposase THAP9 [Saimiri boliviensis] 999
```

First, I find the longest sequence and report its ID and length using awk command. So I find >XP\_074251769.1 DNA transposase THAP9 [Saimiri boliviensis] sequence length is higher i.e. 999.

## **Chatgpt Prompt:**

How I calculate amino acid frquency's of the longest sequence, solve using linux

Perfect! Let's do this **entirely in Linux** using command-line tools. I'll assume your file is a **multi-FASTA file** called sequences.fasta.

## **Chatgpt Result:**

```
bash

awk '/^>/{if(seq!=""){print header"\n"seq}; header=$0; seq=""} /^[^>]/ {seq=seq $0} END{print header bash

bash

Copy code

grep -v '^>' longest_seq.fasta | fold -w1 | sort | uniq -c | awk '{printf "%s\t%s\t%.2f\%\n",$2,$1
```

So, using chatgpt I learn how to calculate the amino acid frequency of the longest sequence. Then grep -v removes the header lines and leaves only the sequence lines, fold -w1 wraps the sequence to one character per line, sort is used to sorts the single-letter lines alphabetically so identical amino acids group together, uniq -c counts how many times each unique letter appears and prints lines.

```
header=$0; seq=""} /^[^>]/ {seq=seq $0} END{print header"\n"seq}' THAP9_homologs_5seq.txt | awk
header=$0; seq=""} /^[^>]/ {seq=seq $0} END{print header"\n"seq}' THAP9_homologs_5seq.txt | awk '/">XP_074251769.1/{print; getline;print}'

XP_074251769.1 DNA transposase THAP9 [Saimiri boliviensis]

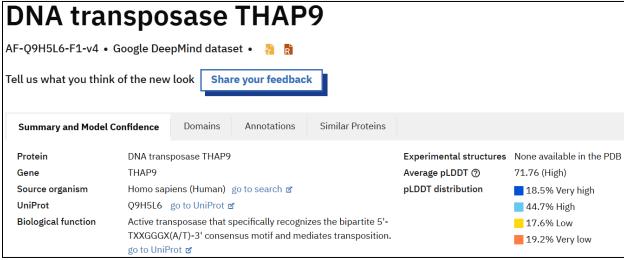
MVGRELPPYFCIGKLQDKAQEHDTHLKGGAVKYTSSGLGPSDMLLRQNVACDWLQCRRRAFGEEVWESRASKARLSCQIKVAAAESTGGAKVIVIPAFRGPEGGDPRNEKMTRSCSAVGC
STRDTVLSRERGLSFHQFPTDTIQRSKWIRAVNRVDPKSKKIWIPGPGAILCSKHFQESDFESYGIRRKLKKGTVPSVSLYKVPQGVHLKGKARQKILKQPLPINSQEVATEDHNYSLKT
PLTVGAEKLAEVQQMLQVSKKRLISVKNYKMIKKRKGLPLIDALVEEKLLSEETECLLRAQFSGAASEAGGKKPGDCAGLEAKCRKYPKGEVVHYVKCCWEIRMLSKCQPGPGFNSHIFSL
LQRRVQNGDQLYQVCSLIIKSIPLKQQLQWDPSSHSLQGFMDFGLGKLDADETPLATETVLLMAVGVFGHWRTPLGYFFVNRASGCLQAQLLRLTIGKLSDTGITVLAVTSDATAHSVQM
AKALGIHIDGDDMKCTFQHPSSCSQQIAYFFDSCHLLRLIRNAFQNFQSIQFINGIAHWQHLVELVALEEQELSNMERIPSTPANLKNHILKVNCAAQLFSESVASALEYLLSLGLPPFQ
NCIGTIHFLRLINNLFDIFNSRNCYGKGLKGPLLPETYSKINHVLIEAKTIFVTLSDSSNNQIIQGKQKLGFLGFLNAESLKWLYQNYVFPKVMPFPYLLTYKFSHDHLELFLKMLRQV
LVTSSSPTCMAFQKAYHNLETRCTFQDEVFLSKVSIFDISVARRKDLALWAVQRQYGVSVTKTLFHKEGICQDWSNCSLSEALLDLSDNRRNLICYAGYVADKLSALLTCEDCITALYAS
DLKASKIGSLLFVKKNNGLHFPSESLCQVINTCERVLRTHSRMAIFFELIPKQRELYLQQKILCELSGHIYLFVDLNKHLFDGEVCAINHFVKLLKDIIICFLNIRANNVAQNPLKHSERT
DMKTLSBKHWSSLPDYKCSSFANTNKFRHLLSNDGYPFK
   DMKTLSRKHWSSLPDYKCSSFANTNKFRHLLSNDGYPFK
 (base) intern@rosalind:~/Nabendu/Biocomputing_Assignment/Project_assignment$ awk '/^>/{if(seq!=""){print header"\n"seq} header=$0; seq=""} /^[^>]/ {seq=seq $0} END{print header"\n"seq}' THAP9_homologs_5seq.txt | awk '/^>XP_074251769.1/{print;getline;print}' > longest_seq.fasta
                                                   | awk '{printf "%s\t%s\t%.2f%%\n",$2,$1,($1/999)*100}' | sort
                                                           31%
                          ЗΠ
                                                     3.40%
                                                     4.10%
                          49
                                                     4.90%
                           50
                          59
31
58
77
121
                                                      3.10%
                                                     5.81%
                                                    7.71%
12.11%
                                                     4.10%
                          39
53
49
                                                     3.90%
                                                      5.31%
                          82
                          40
                          56
14
                                                     5.61%
                                                     1 Ц0%
```

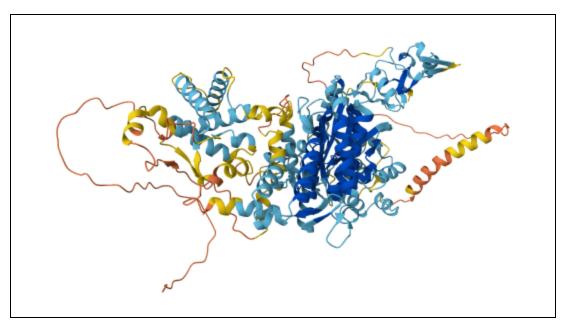
## **Question 3 (7 Points)**

a. Retrieve the 3D structure of the same protein from the RCSB PDB database in PDB format. If the PDB structure is not present, then download the PDB file of the AlphaFold model from AlphaFold database. (1 Points)

I don't find the THAP9 pdb structure in the RCSB PDB database. So I downloaded the PDB structure of THAP9 from the AlphaFold database.







## THAP9 pdb structure

```
HEADER
                                                 01-JUN-22
         ALPHAFOLD MONOMER V2.0 PREDICTION FOR DNA TRANSPOSASE THAP9 (Q9H5L6)
TITLE
COMPND
        2 MOLECULE: DNA TRANSPOSASE THAP9;
COMPND
COMPND
       3 CHAIN: A
SOURCE
         MOL ID: 1;
SOURCE
        2 ORGANISM SCIENTIFIC: HOMO SAPIENS;
        3 ORGANISM TAXID: 9606
SOURCE
REMARK
REMARK
        1 REFERENCE 1
                  JOHN JUMPER, RICHARD EVANS, ALEXANDER PRITZEL, TIM GREEN,
REMARK
        1 AUTH
REMARK
        1 AUTH 2 MICHAEL FIGURNOV, OLAF RONNEBERGER, KATHRYN TUNYASUVUNAKOOL,
        1 AUTH 3 RUSS BATES, AUGUSTIN ZIDEK, ANNA POTAPENKO, ALEX BRIDGLAND,
REMARK
        1 AUTH 4 CLEMENS MEYER, SIMON A A KOHL, ANDREW J BALLARD,
REMARK
REMARK
        1 AUTH 5 ANDREW COWIE, BERNARDINO ROMERA-PAREDES, STANISLAV NIKOLOV,
        1 AUTH 6 RISHUB JAIN, JONAS ADLER, TREVOR BACK, STIG PETERSEN,
REMARK
REMARK
        1 AUTH 7 DAVID REIMAN, ELLEN CLANCY, MICHAL ZIELINSKI,
REMARK
        1 AUTH 8 MARTIN STEINEGGER, MICHALINA PACHOLSKA, TAMAS BERGHAMMER,
REMARK
       1 AUTH 9 DAVID SILVER, ORIOL VINYALS, ANDREW W SENIOR,
           AUTH10 KORAY KAVUKCUOGLU, PUSHMEET KOHLI, DEMIS HASSABIS
REMARK
REMARK
        1 TITL
                  HIGHLY ACCURATE PROTEIN STRUCTURE PREDICTION WITH ALPHAFOLD
REMARK
        1 REF
                  NATURE
                                                V. 596
                                                         583 2021
                                  ISSN 0028-0836
REMARK
        1 REFN
REMARK
        1 PMID
                  34265844
REMARK
        1 DOI
                  10.1038/s41586-021-03819-2
REMARK
```

```
ATOM
                               31.867 21.904 -41.751 1.00 51.11
                MFT A
ATOM
                               30.655 22.682 -41.391 1.00 51.11
ATOM
                MFT A
                              30.962 23.625 -40.232 1.00 51.11
                                                                          C
ATOM
            CB
                MET A
                               29.480
                                      21.766 -41.004 1.00 51.11
ATOM
            0
                MET A
                               31.238 23.156 -39.134 1.00 51.11
                                                                          0
                              28.720 21.225 -42.217 1.00 51.11
ATOM
            CG
               MFT A
ATOM
                              27.357 20.134 -41.741 1.00 51.11
ATOM
            CE
                              26.627 19.795 -43.369 1.00 51.11
                MET A
MOTA
                              30.933 24.938 -40.463 1.00 64.93
         9
                THR A
                                                                          N
ATOM
                               31.023
                                      25.965 -39.407
        10
                THR A
                                                      1.00 64.93
ATOM
                              29.744 25.973 -38.572 1.00 64.93
        11 C
                THR A
ATOM
           CB
                THR A
                              31.213 27.361 -40.021 1.00 64.93
ATOM
                              28.651 26.085 -39.127 1.00 64.93
                                                                          c
ATOM
        14 CG2 THR A
                              32.648 27.579 -40.486 1.00 64.93
ATOM
        15
            OG1 THR A
                               30.363 27.495 -41.143 1.00 64.93
ATOM
        16
                ARG A
                              29.863 25.860 -37.247 1.00 81.72
                                                                          N
ATOM
                              28.713 25.896 -36.331 1.00 81.72
        17 CA
               ARG A
ATOM
                              28.223 27.341 -36.173 1.00 81.72
ATOM
                              29.097 25.276 -34.978 1.00 81.72
        19
           CB
               ARG A
                       3
                                                                          C
ATOM
        20
            0
                ARG A
                               29.037 28.251 -36.053 1.00 81.72
                                                                          0
                               29.545 23.809 -35.063 1.00 81.72
MOTA
        21
            CG
                ARG A
```

- b. Extract the following details of all C-alpha (CA) atoms of Alanine residues in chain A. (3 Points)
- Atom name, Residue name, Residue ID, Chain ID, x, y, z coordinates and save as a new alanine info.pdb

```
(base) intern@rosalind:~/Nabendu/Biocomputing_Assignment/Project_assignment$ awk '/^ATOM/ && $3=="CA" && $4=="ALA" && $5=="A" {print $3, $4, $5, $6, $7, $8, $9}' AF-Q9H5L6-F1-model_v4.pdb > alanine_info.pdb
```

(base) intern@rosalind:~/Nabendu/Biocomputing\_Assignment/Project\_assignment\$ vi alanine\_info.pdb

```
CA ALA A 7 20.025 32.322 -30.475
CA ALA A 41 16.432 31.839 -21.892
                                               385 11.344 -4.951 -6.142
CA ALA A
                                      CA ALA A 410 7.176 6.084 1.505
CA ALA A
                                      CA ALA A 425 -6.579 -14.274 10.684
CA ALA A
                                      CA ALA
                                            Α
CA ALA A
                                      CA ALA
                                            Α
CA ALA A
                                                   -33.386 -1.481 29.692
                                         ALA A
         140 51.818 7.502 -15.443
                                      CA ALA A
CA ALA A
                                      CA ALA A
CA ALA A
                                      CA ALA
                                             Α
CA ALA A
                                      CA ALA
                                             Α
CA ALA A
                                         ALA
                                             Α
CA ALA A
                                      CA ALA
                                            Α
CA ALA A
                                                   -16.070 3.018 -20.060
CA ALA A
                                      CA ALA
                                             Α
CA ALA A
                                      CA ALA
                                            Α
CA ALA A
                                         ALA
                                            Α
CA ALA A
                                      CA ALA A
                                                            10.269 9.881
CA ALA A
                                      CA ALA A
CA ALA A
                                     CA ALA A
```

So, in this question, I extracted the C-alpha atoms of Alanine residues in chain A. Also, print the atom name, residue name, residue id, chain id, x, y, z coordinates, and save it to a new file named alanine info.pdb.

- c. Using the extracted CA coordinates of Alanine, compute the pairwise distances between all alanine CA atoms in chain A. Report the results in the form of a table with the following format: (3 Points)
- Residue\_i | Residue\_j | Distance

### **Chatgpt prompt:**

```
how to calculate pairwise distances

Good question 3 — let's go step by step on how pairwise distances are calculated.
```

## **Chatgpt result:**

```
formula the table header
echo -e "Residue_i\tResidue_j\tDistance(Å)"

# Now run awk on your alanine_info.pdb file
awk '
# step 1: Collect only Alanine CA atoms in chain A
# $1=atom name, $2=residue name, $3=chain, $4=residue ID, $5/$6/$7 = x y z
$1=="CA" && $2=="ALA" && $3=="A" {
    res[++n] = $4  # store residue ID in array res[]
    x[n] = $5  # store x coordinate
    y[n] = $6  # store y coordinate
    z[n] = $7  # store z coordinate
}

# Step 2: After reading all lines, compute pairwise distances
END {
    # Loop over all unique pairs (i < j)</pre>
```

```
# Step 2: After reading all lines, compute pairwise distances
END {
    # Loop over all unique pairs (i < j)
    for (i=1; i<=n; i++) {
        for (j=i+1; j<=n; j++) {
            # Compute coordinate differences
            dx = x[i] - x[j]
            dy = y[i] - y[j]
            dz = z[i] - z[j]

# Step 3: Apply distance formula
        # d = sqrt(dx^2 + dy^2 + dz^2)
            dist = sqrt(dx*dx + dy*dy + dz*dz)

# Step 4: Print residue IDs and distance
            printf "%d\t%d\t%.3f\n", res[i], res[j], dist
        }
    }
}
' alanine_info.pdb</pre>
```

Echo -e tells echo to interpret \t as a tab character. So the header columns are tab-separated. In res[++n] = \$4, ++n increments n by 1 and returns the incremented value. res[n] stores the residue ID (from field \$4) into the res array, after the first match, n==1. x[n] = \$5; y[n] = \$6; z[n] = \$7 stores the x, y, z coordinates of that CA atom into parallel arrays. dx = x[i] - x[j] (and dy, dz) compute differences in coordinates between the two residues. Here we apply the 3-D Euclidean distance formula, dist = sqrt(dx\*dx + dy\*dy + dz\*dz).

(base) intern@rosalind:~/Nabendu/Biocomputing\_Assignment/Project\_assignment\$ vi pairwise\_dist.sh

```
#!/bin/bash
(
    echo -e "Residue_i\tResidue_j\tDistance|"
    awk '
    $1=="CA" && $2=="ALA" && $3=="A" {
        res[++n]=$4; x[n]=$5; y[n]=$6; z[n]=$7
    }
    END {
        for(i=1;i<=n;i++){
            for(j=i+1;j<=n;j++){
                  dx=x[i]-x[j]; dy=y[i]-y[j]; dz=z[i]-z[j]
                  dist=sqrt(dx*dx+dy*dy+dz*dz)
            printf "%d\t%d\t%.3f\n", res[i], res[j], dist
        }
    }
    ' alanine_info.pdb
)</pre>
```

(base) intern@rosalind:~/Nabendu/Biocomputing\_Assignment/Project\_assignment\$ bash pairwise\_dist.sh > pairwise\_dist.csv

| Residu | e_i | Residue_j | Distance |
|--------|-----|-----------|----------|
| 7      | 41  | 9.317     |          |
| 7      | 59  | 9.782     |          |
| 7      | 84  | 15.870    |          |
| 7      | 103 | 30.679    |          |
| 7      | 120 | 37.832    |          |
| 7      | 136 | 47.421    |          |
| 7      | 140 | 43.044    |          |
| 7      | 173 | 47.596    |          |
| 7      | 190 | 49.720    |          |
| 7      | 210 | 34.399    |          |
| 7      | 216 | 35.397    |          |
| 7      | 312 | 47.876    |          |

#### Github Code:

```
Nabendu@LAPTOP-DM1JT3BQ MINGW64 ~/Downloads/Project-1-Biocomputing (master)
$ git init
Initialized empty Git repository in C:/Users/Nabendu/Downloads/Project-1-Biocomputing/.git/
Nabendu@LAPTOP-DM1JT3BQ MINGW64 ~/Downloads/Project-1-Biocomputing (master)
$ git config user.email "nabendudas"
Nabendu@LAPTOP-DM1JT3BQ MINGW64 ~/Downloads/Project-1-Biocomputing (master)
$ git config user.name "nabendudas"
Nabendu@LAPTOP-DM1JT3BQ MINGW64 ~/Downloads/Project-1-Biocomputing (master)
$ git config user.email "nabendudas001@gmail.com"
Nabendu@LAPTOP-DM1JT3BQ MINGW64 ~/Downloads/Project-1-Biocomputing (master)
$ pwd
/c/Users/Nabendu/Downloads/Project-1-Biocomputing
Nabendu@LAPTOP-DM1JT3BQ MINGW64 ~/Downloads/Project-1-Biocomputing
Nabendu@LAPTOP-DM1JT3BQ MINGW64 ~/Downloads/Project-1-Biocomputing
warning: in the working copy of 'README.txt', LF will be replaced by CRLF the ne
```

```
Nabendu@LAPTOP-DMIJT3BQ MINGW64 ~/Downloads/Project-1-Biocomputing

Nabendu@LAPTOP-DMIJT3BQ MINGW64 ~/Downloads/Project-1-Biocomputing

warning: in the working copy of 'README.txt', LF will be replaced by CRLF the ne

xt time Git touches it

warning: in the working copy of 'data/THAP1_mouse.fasta', LF will be replaced by

CRLF the next time Git touches it

warning: in the working copy of 'data/THAP1_rat.fasta', LF will be replaced by

CRLF the next time Git touches it

warning: in the working copy of 'data/THAP9_chimpanzee.fasta', LF will be replaced

dby CRLF the next time Git touches it

warning: in the working copy of 'data/THAP9_chimpanzee.fasta', LF will be replaced

by CRLF the next time Git touches it

warning: in the working copy of 'data/THAP9_human.fasta', LF will be replaced

by CRLF the next time Git touches it

warning: in the working copy of 'results/AF-Q9H5L6-F1-model_v4.pdb', LF will be

replaced by CRLF the next time Git touches it

warning: in the working copy of 'results/THAP9_coding sequences.txt', LF will be

replaced by CRLF the next time Git touches it

warning: in the working copy of 'results/THAP9_homologs_5seq.txt', LF will be re

placed by CRLF the next time Git touches it

warning: in the working copy of 'results/THAP9_homologs_5seq.txt', LF will be re

placed by CRLF the next time Git touches it

warning: in the working copy of 'results/THAP9_homologs_5seq.txt', LF will be re

placed by CRLF the next time Git touches it

warning: in the working copy of 'results/Salanine_info.pdb', LF will be replaced

by CRLF the next time Git touches it

warning: in the working copy of 'results/emboss_transeq-I20250916-162342-0572-73

014202-plm.out', LF will be replaced by CRLF the next time Git touches it

warning: in the working copy of 'results/leader_files.txt', LF will be replaced

by CRLF the next time Git touches it

warning: in the working copy of 'results/longest_seq.fasta', LF will be replaced

by CRLF the next time Git touches it

warning: in the working copy of 'results/pairwise_dist.c
```

```
Nabendu@LAPTOP-DM1JT3BQ MINGW64 ~/Downloads/Project-1-Biocomputing (master)
$ git status
On branch master
No commits yet

Changes to be committed:
   (use "git rm --cached <file>..." to unstage)
        new file: README.txt
        new file: data/Project_1.pdf
        new file: data/THAP1_mouse.fasta
        new file: data/THAP2_chimpanzee.fasta
        new file: data/THAP9_chimpanzee.fasta
        new file: data/THAP9_homan_fasta
        new file: results/AF-Q9H5L6-F1-model_v4.pdb
        new file: results/AF-Q9H5L6-F1-model_v4.pdb
        new file: results/THAP9_coding sequences.txt
        new file: results/THAP9_homologs_5seq.txt
        new file: results/THAP9_homologs_5seq.txt
        new file: results/THAP9_homologs_5seq.txt
        new file: results/cds.fna
        new file: results/cds.fna
        new file: results/cds.fna
        new file: results/beader_files.txt
        new file: results/beader_files.txt
        new file: results/pairwise_dist.csv
        new file: results/pairwise_dist.csv
        new file: results/pairwise_dist.csv
        new file: results/pairwise_dist.csv
        new file: results/pairwise_dist.sh
```

```
Nabendu@LAPTOP-DM1JT3BQ MINGW64 ~/Downloads/Project-1-Biocomputing (master)

§ git commit -m "/c/Users/Nabendu/Downloads/Project-1-Biocomputing"
[master (root-commit) 5f0dc3c] C:/Users/Nabendu/Downloads/Project-1-Biocomputing
20 files changed, 10161 insertions(+)
create mode 100644 README.txt
create mode 100644 data/FhAP1_mouse.fasta
create mode 100644 data/THAP1_mouse.fasta
create mode 100644 data/THAP2_chimpanzee.fasta
create mode 100644 data/THAP9_chimpanzee.fasta
create mode 100644 data/THAP9_human.fasta
create mode 100644 data/THAP9_human.fasta
create mode 100644 results/AF-Q9H5L6-F1-model_v4.pdb
create mode 100644 results/AF-Q9H5L6-F1.png
create mode 100644 results/THAP9_coding sequences.txt
create mode 100644 results/THAP9_homologs_5seq.txt
create mode 100644 results/THAP9_homologs_5seq.txt
create mode 100644 results/CHAP9_homologs_5seq.txt
  create mode 100644 results/header_files.txt
create mode 100644 results/longest_seq.fasta
create mode 100644 results/pairwise_dist.csv
create mode 100644 results/sequence_id5.txt
  create mode 100644 scripts/pairwise_dist.sh
   git remote add origin https://github.com/nabendudas/Project-1-Biocomputing.git
                  @LAPTOP-DM1JT3BQ MINGW64 ~/Downloads/Project-1-Biocomputing (master)
$ git branch -M main
  Nabendu@LAPTOP-DM1JT3BQ MINGW64 ~/Downloads/Project-1-Biocomputing (main)
$ git push -u origin main
Enumerating objects: 25, done.
Counting objects: 100% (25/25), done.
Delta compression using up to 12 threads
Compressing objects: 100% (23/23), done.
Writing objects: 100% (25/25), 317.86 KiB | 3.15 MiB/s, done.
Total 25 (delta 4), reused 0 (delta 0), pack-reused 0 (from 0)
remote: Resolving deltas: 100% (4/4), done.
To https://github.com/nabendudas/Project-1-Biocomputing.git
           [new branch]
                                                                          main -> main
branch 'main' set up to track 'origin/main'.
```

| Project-1-Biocomputing Public                  |                                   | \$\hat{S}\$ ₽          | in                |
|--|-----------------------------------|------------------------|-------------------|
| ্বি main ▼ ু 1 Branch 🟷 0 Tags                 | Q Go to file                      | t Add file 🔻           | <> Code ▼         |
| nabendudas C:/Users/Nabendu/Downloads/Project- | -1-Biocomputing                   | 5f0dc3c · 20 hours ago | <b>™</b> 1 Commit |
| adata  | C:/Users/Nabendu/Downloads/Projec | t-1-Biocomputing       | 20 hours ago      |
| results  | C:/Users/Nabendu/Downloads/Projec | t-1-Biocomputing       | 20 hours ago      |
| scripts  | C:/Users/Nabendu/Downloads/Projec | t-1-Biocomputing       | 20 hours ago      |
| ☐ README.txt                                   | C:/Users/Nabendu/Downloads/Projec | t-1-Biocomputing       | 20 hours ago      |

Github Link: https://github.com/nabendudas/Project-1-Biocomputing/tree/main