

## ▼ Question 1

Write a program to find the anti-complimentary sequence of an input DNA sequence

```
def complement(seq):
    seqr=""
    for i in range(len(seq)):
        if seq[i] == "A":
            seqr=seqr+"T"
        if seq[i] == "T":
            seqr=seqr+"A"
        if seq[i] == "G":
            seqr=seqr+"C"
        if seq[i] == "C":
            seqr=seqr+"G"
    return seqr

def reverse(seq):
    return seq[::-1]

def first():
    print("Give the input sequence : ")
    seq=input()
    print("Complement of the given sequence: "+complement(seq))
    print("Anti complement/ reverse complement of given sequence: "+reverse(complement(seq)))

first()

Give the input sequence :
AAGGAAAC
Complement of the given sequence: TTCCTTTG
Anti complement/ reverse complement of given sequence: GTTTCCTT
```

## ▼ Question 2

Intrinsic terminators have two prominent structural features:

- (1) a sequence of nucleotides that includes an inverted repeat (i.e., the sequence 5'-CGGATG|CATCCG-3' contains an inverted repeat centered at the "|" because "5'-CGGATG-3'" reads "5'-CATCCG-3'" on its complementary strand), and
- (2) a run of roughly six uracils immediately following the inverted repeat. Write a function to identify intrinsic terminators from the sequence.

```
def finding(seq):
    req="UUUUUU"
    if (seq.find(req) != -1):
        seq=seq[: seq.find(req)]
        return seq
    else:
        return False

def finds(x,seq):
    result=[]
    for i in range(len(seq)):
        if seq[i]==x:
            result.append(i)
    return result

def truth(x,y,seq):
    if (seq[x]=="T" and seq[y]=="A") or (seq[x]=="A" and seq[y]=="T") or (seq[x]=="C"
        return True
    else:
        return False

def cond(seq):
    n=len(seq)
    if seq[n-1]=="A":
        index=finds("T",seq)
    elif seq[n-1]=="T":
        index=finds("A",seq)
    elif seq[n-1]=="C":
        index=finds("G",seq)
    elif seq[n-1]=="G":
        index=finds("C",seq)
    #print(index)
    for i in index:
        if (n-1-i)%2!=0:
            #print(i)
            x=i
            y=n-1

    while(x<y):
        cond=0
        if truth(x,y,seq):
            #print(x,y)
            x=x+1
            y=y-1
        else:
            cond=1
            x=y+1
    if cond==0:
        #print("here i am")
        seq=seq[i:x]
        return seq
    break
```

```

    return -1

#fixing length by 6
def second():
    print("Give the sequence: ")
    seq=input()
    sub_seq=finding(seq)
    #print(sub_seq)
    #print(len(sub_seq))
    if sub_seq:
        x=cond(sub_seq)
        if x== -1:
            print("No intrinsic terminator")
        else:
            print("The intrinsic terminator is: "+x)
    else:
        print("No intrinsic terminator")

print("How many times do you want to verify for intrinsic terminator: ")
n=int(input())
for i in range(n):
    second()

    How many times do you want to verify for intrinsic terminator:
    3
    Give the sequence:
    AAGGAAACGTTT
    No intrinsic terminator
    Give the sequence:
    AAGGAAACGTTTUUUUUU
    The intrinsic terminator is: AAAC
    Give the sequence:
    AAGGAACTUUUUUU
    No intrinsic terminator

```

## ▼ Question 3

Given DNA sequence, arrange the codons in the decreasing order of frequency.

```

def identify_codons(seq):
    codons=[]
    while (len(seq)>2):
        codons.append(seq[:3])
        seq=seq[3:]
    return codons

```

```
def codons_x(codons):
    Codon_count= {}
    for element in codons:
        if element in Codon_count:
            Codon_count[element] += 1
        else:
            Codon_count[element] = 1
    Codon_list= sorted(Codon_count.items(),key=operator.itemgetter(1),reverse=True)
    return Codon_list
#for key, value in Codon_list.items():
    # print(key, ' : ', value)

import operator
def main():
    print("Give the nucleotide sequence ")
    seq=input()
    print("The codons identified in the sequence are: ")
    codons= identify_codons(seq)
    codons=codons_x(codons)
    print("(codon,frequency)")
    for i in codons:
        print(i)
```

```
main()
```

```
Give the nucleotide sequence
ACTACTGGG
The codons identified in the sequence are:
(codon,frequency)
('ACT', 2)
('GGG', 1)
```

## ▼ Question 4

Given two strings  $s$  and  $t$  of equal length, the Hamming distance between  $s$  and  $t$ , denoted  $dH(s,t)$ , is the number of corresponding symbols that differ in  $s$  and  $t$ . Write a function to compute  $dH(s,t)$

```
def distance(seq1,seq2):
    res=0
    for i in range(len(seq1)):
        if seq1[i]!=seq2[i]:
            res+=1
    return res

def hamming():
    print("Give the first sequence ")
    seq1=input()
```

```
print("Give the second sequence ")
seq2=input()
print("The hamming distance is "+ str(distance(seq1,seq2)))
```

```
hamming()
```

```
Give the first sequence
AGCT
Give the second sequence
ATCG
The hamming distance is 2
```

## ▼ Question 5

Given Two protein strings s and t in FASTA format (each of length at most 100 aa), write a program to find the optimal global alignment score (use match score, mismatch and gap penalties as variables)

```
def needlmen(seq1,seq2):
    gap=-1
    mismatch=-1
    match=1
    n= len(seq1)
    m= len(seq2)
    res = [ [ 0 for i in range(m+1) ] for j in range(n+1) ]
    #Initialization
    count=0
    for i in range(n+1):
        res[i][0]=count
        count-=1
    count=0
    for i in range(m+1):
        res[0][i]=count
        count-=1
    print(res)
    for i in range(1,n+1):
        for j in range(1,m+1):
            x=res[i-1][j]+gap
            y=res[i][j-1]+gap
            if seq1[i-1]==seq2[j-1]:
                z=res[i-1][j-1]+match
            else:
                z=res[i-1][j-1]+mismatch
            res[i][j]=max(x,y,z)
    print(res)
    return res[n][m]
```

```
print(needlmen("ACTG","ACTG"))
```

```
[[0, -1, -2, -3, -4], [-1, 0, 0, 0, 0], [-2, 0, 0, 0, 0], [-3, 0, 0, 0, 0], [-4, 0, 0, 0, 0],
[0, -1, -2, -3, -4], [-1, 1, 0, -1, -2], [-2, 0, 2, 1, 0], [-3, -1, 1, 3, 2],
4
```

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```
def alignment():
    print("Give the sequence 1: ")
    seq1=input()
    print("Give the sequence 2: ")
    seq2=input()
    print("The optimal global alignment score is: "+str(needleman(seq1,seq2)))
```

```
alignment()
```

```
Give the sequence 1:
gactt
Give the sequence 2:
att
[[0, -1, -2, -3], [-1, 0, 0, 0], [-2, 0, 0, 0], [-3, 0, 0, 0], [-4, 0, 0, 0],
[0, -1, -2, -3], [-1, -1, -2, -3], [-2, 0, -1, -2], [-3, -1, -1, -2], [-4, -2, 0, -1],
The optimal global alignment score is: 1
```

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