# Chapter 02. Working with Sequence

**Python Programming for Bioinformatics** 

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- Create a Biological Sequence
- Basic Operations for Sequences
- Functions Applying for Sequences
- Biological Operations for Sequences
- Modifiable Sequences
- Biological Functions for Strings





# **CREATE A BIOLOGICAL SEQUENCE**

# Introduction to Bio.Seq.Seq

# **Bio.Seq.Seq**

### (Bio.Seq.Seq is a Read-only Object)



#### **GenBank**

	FOGAZ	Z78533 740 bp DNA linear PLN 30-NOV-2006
	DEFINITION	C.irapeanum 5.8S rRNA gene and ITS1 and ITS2 DNA.
	ACCESSION	Z78533
	VERSION	Z78533.1 GI:2765658
	KEYWORDS	5.8S ribosomal RNA; 5.8S rRNA gene; internal transcribed spacer;
		ITS1; ITS2.
	SOURCE	Cypripedium irapeanum
	ORGANISM	Cypripedium irapeanum
		Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
		Spermatophyta; Magnoliophyta; Liliopsida; Asparagales; Orchidaceae;
		Cypripedioideae; Cypripedium.
	REFERENCE	1
	AUTHORS	Cox,A.V., Pridgeon,A.M., Albert,V.A. and Chase,M.W.
	TITLE	Phylogenetics of the slipper orchids (Cypripedioideae:
		Orchidaceae): nuclear rDNA ITS sequences
	JOURNAL	Unpublished
	REFERENCE	2 (bases 1 to 740)
	AUTHORS	Cox, A.V.
	TITLE	Direct Submission
	JOURNAL	Submitted (19-AUG-1996) Cox A.V., Royal Botanic Gardens, Kew,
		Richmond, Surrey TW9 3AB, UK
	FEATURES	Location/Qualifiers
	Source	3 1740
1	<b>/</b>	/organ1sm="Cypr1ped1um 1rapeanum"
		/mol_type="genomic_DNA"
		/db_xref="taxon:49/11"
		leature 1580
		/note="internal transcribed spacer 1"
		561
		/gene= 0.05 IRNA
		501
		/gene= 5.05 IKNA /muchust="5.95 milessenal DNa"
		551 240
		noto-"internal transcribed enacer 2"
		note- internal (lanselloed spacel 2
l	y	costage teaacciece eaageatcat teateagann eteesatasa
	61 (	costcosete satcceesee acceetatat traectcarc eeeeecatte circceteet
	21	accordant tettettere correctere accelerate accertites acciding
	81 0	concoranti topococcaa occatatoaa agcatcacco occaatooca tiototicco
	241 0	casaaccogg agoggogg tectetogog teccoatga attiteatga cictogoaaa
	801 0	Constant to the second se
	- × * *	The second

1) create a state a

## **Different Types of Sequences**

```
from Bio.Seq import Seq
                                                                          Output:
 2
   # DNA Sequence
 3
                                                                          ATCG
   dna_seq = Seq("ATCG") Legal Chars: 'ATCG'
   print(dna seq)
                                                                          UAGC
 6
                                                                          MELKILV
   # RNA Sequence
   rna_seq = Seq("UAGC") Legal Chars: 'UAGC'
                                                                          ATNNNCG
   print(rna seq)
 9
10
                                                                          ATC--GCA
11
   # Protein Sequence
                                 Legal Chars:
12
   protein seq = Seq("MELKILV")
                                 'ACDEFGHIKLMNPQRSTVWY'
   print(protein_seq)
13
14
   # Sequence with Unknown Letter 'N' (@GenBank or EMBL)
15
   unknown_seq = Seq("ATNNNCG")
16
   print(unknown seq)
17
                                                       Notice:
18
                                                          Although you can add any character to Seq,
   # Sequence with gap letter '-'
19
                                                          invalid characters can cause errors in the program
   gap seq = Seq("ATC--GCA")
20
                                                          when performing certain actions with biological meaning.
   print(gap_seq)
21
                                                          (e.g., Transcription, Translation...etc.)
```



### Construct Different Types of Sequences

- Write and Run the following codes on a Colab page called "SeqObjects.ipynb":

•	Cre	eat	e Different Types of Sequence
~	[17]	1	from Bio.Seg import Seg
		2	
		3	# DNA Sequence
		4	<pre>dna_seg = Seg("ATCG")</pre>
		5	print(dna_seq)
		6	
		7	# RNA Sequence
		8	<pre>rna_seg = Seg("UAGC")</pre>
		9	print(rna_seq)
		10	
		11	# Protein Sequence
		12	protein_seq = Seq("MELKILV")
		13	<pre>print(protein_seq)</pre>
		14	
		15	# Sequence with Unknown Letter 'N' (@GenBank or EMBL)
		16	unknown_seq = Seq("ATNNNCG")
		17	print (unknown_seq)
		18	
		19	# Sequence with gap letter '-'
		20	<pre>gap seq = Seq("ATCGCA")</pre>
		21	<pre>print(gap_seq)</pre>



# **BASIC OPERATIONS FOR SEQUENCES**



### Add Two Sequences

```
from Bio.Seq import Seq
 3 # Add two Nucleic Acid Sequences
   dna seq1 = Seq("AATC")
 4
 5 dna seq2 = Seq("CGAT")
 6 dna seq = dna seq1 + dna seq2
 7 print(dna seq)
8 print(dna seg + Seg("CG"))
   print(dna seg + "AGCG")
9
10
11 # Legal, but non-sense
   protein seg = Seg("MELKILV")
12
13 rna seq = Seq("UAGC")
14 print(protein seq + dna seq)
15 print(dna seg + rna seg)
```





### Add Sequences

- Write and Run the following codes on a Colab page called "SeqObjects.ipynb":

```
    Add

           from Bio.Seg import Seg
24]
           # Add two Nucleic Acid Sequences
            dna seq1 = Seq("AATC")
        5 dna seq2 = Seq("CGAT")
          dna seg = dna seg1 + dna seg2
          print(dna seg)
            print(dna seq + Seq("CG"))
        8
            print(dna seg + "AGCG")
        9
       10
       11
           # Legal, but non-sense
       12
           protein_seg = Seg("MELKILV")
       13
            rna seg = Seg("UAGC")
       14
            print (protein seq + dna seq)
       15
            print (dna seq + rna seq)
```





### Multiply Sequences with Integer

### **Multiply = Repeat**

```
1 from Bio.Seq import Seq
2
3 # Multiply with an Integer
4 dna_seq = Seq("AATC") * 3
5 print(dna_seq)
6
7 protein_seq = Seq("MELKILV") * 2
8 print(protein_seq)
```





### Multiply Sequences by Integers

- Write and Run the following codes on a Colab page called "SeqObjects.ipynb":



### (Solution <u>URL</u> of this Practice)



### **Compare Sequences**







### Comparing Sequences

- Write and Run the following codes on a Colab page called "SeqObjects.ipynb":

```
    Compare

            from Bio.Seq import Seq
        2
         seq1 = Seg("ATCG")
        4 seq2 = Seq("ATCG")
        5 seq3 = Seq("GGCC")
        7
          # Compare two Bio.Seq.Seq objects
        8 print(seq1 == seq2)
        9 print(seq1 != seq3)
       10
       11 # Compare Bio.Seq.Seq against a string
       12 print(seq1 == "ATCG")
       13 print(seq1 == "GGCC")
       14
           print(seq2 != "AATT")
```



### Slice a Sequence

### "Slicing" = The Way to Get a "Sub-Sequence"

-14 -13 -12 -11 -10 -9 -8 -7 -6 -5 -4 -3 -2 -1

0 1 2 3 4 5 6 7 8 9 10 11 12 13

# "ATCCGATGCACCAG"

Slicing of 10 ~ 13

#### Notice:

- Python is "0-based" (0~13)
- NCBI sequences are "1-based" (1~14)

### Slice a Sequence

```
from Bio.Seq import Seq
   my seg = Seg("GATCGATGGGCCTATATAGGATCGAAAATCGC")
                                                   С
 4
                                                   GATGGGCC
   # Slice on a Single Character
   print(my seq[2])
                                                   GCTGTAGTAAG
   print(my_seq[-1])
                                                   AGGCATGCATC
 8
                                                   TAGCTAAGAC
   # Slice on a Region of the Sequence
 9
                                                   CGCTAAAAGCTAGGATATATCCGGGTAGCTAG
   print(my seq[4:12])
10
11
  # Slice with start, stop, and step
12
   # To get the 1, 2, 3 codon positions of this DNA sequence
13
14 print(my seq[0::3])
15 print(my seq[1::3])
   print(my seq[2::3])
16
17
18 # Reverse the order of sequence
19 print(my_seq[::-1])
```



### • Slicing a Sequence

- Write and Run the following codes on a Colab page called "SeqObjects.ipynb":

•	Slicing	
0s		from Bio.Seq import Seq
	3 4	<pre>my_seq = Seq("GATCGATGGGCCTATATAGGATCGAAAAATCGC")</pre>
	5	<pre># Slice on a Single Character print(my seg[2])</pre>
	7	<pre>print(my_seq[-1])</pre>
	9 10	<pre># Slice on a Region of the Sequence print(my seg[4:12])</pre>
	11	+ Slice with start stop and step
	13	# To get the 1, 2, 3 codon positions of this DNA sequence
	14	<pre>print(my_seq[0::3]) print(my_seq[1::3])</pre>
	16 17	<pre>print(my_seq[2::3])</pre>
	18 19	<pre># Reverse the order of sequence print(my_seq[::-1])</pre>

(Solution <u>URL</u> of this Practice)



### Iterate a Sequence

### • Iteration = Take 1 letter a time from the Sequence





- Iterating a Sequence
  - Write and Run the following codes on a Colab page called "SeqObjects.ipynb":





# FUNCTIONS APPLYING FOR SEQUENCES



# Length of a Sequence







### • Get the Length of a Sequence

- Write and Run the following codes on a Colab page called "SeqObjects.ipynb":



(Solution <u>URL</u> of this Practice)



## Change the Case





### Chage the Case of Sequences

- Write and Run the following codes on a Colab page called "SeqObjects.ipynb":





## **Check for Containing**

```
from Bio.Seq import Seq
 2
 3
   # Create a Sequence
    my_dna = Seq("ATATGAAATTTGAAAA")
 4
 5
    # Check for containing by Bio.Seq.Seq
 6
    print(Seq("AAA") in my_dna)
 7
 8
    # Check for containing by regular strings
 9
    print("AAA" in my_dna)
10
```

- **1** Create a Sequence
- 2 Sequence <compare> Sequence
- **3** String <compare> Sequence



### Check for Containing

- Write and Run the following codes on a Colab page called "SeqObjects.ipynb":
- Check for Containing





# Non-overlapped Counting

```
1 from Bio.Seq import Seq
2
3 # Create a Sequence
4 my_seq = Seq("AAAATCCGCGATAGC")
5
6 # Non-overlapped Counting
7 print(my_seq.count("A"))
8 print(my_seq.count("AA"))
9 print(my_seq.count("AA", 2, -1)) # index= 2~-1
```

.count("A") AAAATCCGCGATAGC  $\rightarrow$  6 .count("AA") AAAATCCGCGATAGC  $\rightarrow 2$ .count("AA", 2, -1) AATCCGCGATAGC → **1** 

# Overlapped Counting

```
1 from Bio.Seq import Seq
2
3 # Create a Sequence
4 my_seq = Seq("AAAATCCGCGATAGC")
5
6 # Overlapped Counting
7 print(my_seq.count_overlap("A"))
8 print(my_seq.count_overlap("AA"))
9 print(my_seq.count_overlap("AA", 2, -1))
```

 $\begin{array}{c} \text{.count\_overlap("A")} \\ \hline A & \hline A$ 

$$\frac{\text{.count_overlap("AA")}}{\text{AAAATCCGCGATAGC}} \rightarrow 3$$

$$\begin{array}{c} \text{.count\_overlap("AA", 2, -1)} \\ 2 & & -1 \end{array}$$

# Count for GC%





- Write and Run the following codes on a Colab page called "SeqObjects.ipynb":
- Count for Occurrence

```
[13] 1 from Bio.Seq import Seq
            # Create a Sequence
            my seq = Seq("AAAATCCGCGATAGC")
         5
        6 # Non-overlapped Counting
         7 print(my seq.count("A"))
          print(my seq.count("AA"))
         8
         9
            print(my_seq.count("AA", 2, -1)) # index= 2~-1
        10
           # Overlapped Counting
       11
       12 print(my_seq.count_overlap("A"))
       13 print(my seq.count overlap("AA"))
       14 print(my_seq.count_overlap("AA", 2, -1))
       15
       16
            # Count for GC%
            gc_percent = float(my_seq.count("G") + my_seq.count("C"))/len(my_seq)
        17
            print("{0:.2%}".format(gc_percent))
        18
       19
           from Bio.SeqUtils import GC
       20
        21 gc_percent = GC(my_seq)
       22 print("{0:.2f}%".format(gc_percent))
```





# **Find Sequence Patterns**

# Find Forwardly: .find()





### Find Sequence Patterns Forwardly

- Write and Run the following codes on a Colab page called "SeqObjects.ipynb":

```
from Bio.Seq import Seq
     # Create a Sequence
     my seq = Seq("AAAATCCGCGATAGC")
 5
     # Find forward for the first occurrence
 6
     print("First Occurred:", my seq.find("CG"))
 8
     # Find forward for all the occurrence
 0
     occur = my seq.count("CG")
10
     head = 0
11
     while occur > 0:
12
      found_index = my_seq.find("CG", start=head, end=-1)
13
14
      print("Found at:", found index)
15
      occur -= 1
16
      head = found index + len("CG")
17
       print("occur: {} head: {}".format(occur, head))
```

# Find Sequence Patterns

# Find Backwardly: .rfind()





### Find Sequence Patterns Backwardly

- Write and Run the following codes on a Colab page called "SeqObjects.ipynb":

```
from Bio.Seg import Seg
 2
    # Create a Sequence
 3
    my seg = Seg("AAAATCCGCGATAGC")
 4
 5
    # Find backward for the first occurrence
 6
    print("Last Occurred:", my seq.rfind("CG"))
 7
 8
    # Find backward for all the occurrence
 9
    occur = my seq.count("CG")
10
11
    tail = Len(my seq)
    while occur > 0:
12
        found_index = my_seq.rfind("CG", start=0, end=tail)
13
        print("Found at:", found index)
14
15
        occur -= 1
16
        tail = found index
        print("occur: {} tail: {}".format(occur, tail))
17
```



# Find Sequence Patterns

### Check for Start Codons & Stop Codons

```
from Bio.Seg import Seg
 2
   # Create a Sequence
    mRNA = Seq("AUGGCCAUUGUAAUGGGCCGCUGAAAGGGUGCCCGAUAGUUG")
 5
    # Check for start codon "AUG" (Met/Methionine)
                                                                         Check Start Codon
    print(mRNA.startswith("AUG"))
 8
    # Check for start codon from specific index
                                                                         Check Start Codon
    print(mRNA.startswith("AUG", start=12))
10
                                                                         at Specific Location
11
12
    # Check for other possible start codons of E. coli
                                                                       Check all Start Codons
    print(mRNA.startswith(("AUG", "GUG", "UUG", "AUU", "CUG")))
13
14
    # Check for possible stop codons: "UAG" (amber), "UAA" (orche), "UGA" (opal)
15
    print(mRNA.endswith(("UAG", "UAA", "UGA")))
16
                                                                       Check all Stop Codons
    print(mRNA.endswith(("UAG", "UAA", "UGA"), end=-3))
17
```



- Check for Start Codons & Stop Codons
  - Write and Run the following codes on a Colab page called "SeqObjects.ipynb":

```
from Bio.Seg import Seg
 2
    # Create a Sequence
 3
    mRNA = Seg("AUGGCCAUUGUAAUGGGCCGCUGAAAGGGUGCCCGAUAGUUG")
 4
 5
    # Check for start codon "AUG" (Met/Methionine)
 6
    print(mRNA.startswith("AUG"))
 8
    # Check for start codon from specific index
 9
    print(mRNA.startswith("AUG", start=12))
10
11
12
    # Check for other possible start codons of E. coli
    print(mRNA.startswith(("AUG", "GUG", "UUG", "AUU", "CUG")))
13
14
    # Check for possible stop codons: "UAG" (amber), "UAA" (orche), "UGA" (opal)
15
    print(mRNA.endswith(("UAG", "UAA", "UGA")))
16
                                                          (Solution URL of this Practice)
    print(mRNA.endswith(("UAG", "UAA", "UGA"), end=-3))
17
```



Biological Meanings of Split & Join



Split a Sequence



# Join Sequences

Exons = [Seq('AU'), Seq('AUUGUAAUG'), Seq('GCUGAAAGGGUGCCCGAUAGUUG')]



mRNA1 =

AUAUUGUAAUGGCUGAAAGGGUGCCCGAUAGUUG

mRNA2 =

AUNNNAUUGUAAUGNNNGCUGAAAGGGUGCCCGAUAGUUG



- Write and Run the following codes on a Colab page called "SeqObjects.ipynb":

```
from Bio.Seq import Seq
 1
 2
 З
     # Create a Sequence
     Pre mRNA = Seg("AUGGCCAUUGUAAUGGGCCGCUGAAAGGGUGCCCGAUAGUUG")
 4
 5
 6
     # Split at Introns
     Introns = "GGCC"
 7
     Exons = Pre_mRNA.split(Introns)
 8
     print(Exons)
 9
10
11
     # Join all Exons with spacers
12
    spacer = Seq("")
13
     mRNA1 = spacer.join(Exons)
     print(mRNA1)
14
15
     spacer = Seq("N"*3)
16
     mRNA2 = spacer.join(Exons)
17
     print(mRNA2)
18
```



### Strip Spaces or Other Symbols

• What is "Strip"?



### ATGCGATTACG

- Why "Strip"?
  - Remove unwanted characters from both sides of the sequence.

Strip Spaces or Other Symbols

```
1 from Bio.Seq import Seq
2
3 # Strip spaces enter by users
4 my_seq = Seq(input("Please enter sequence: "))
5 print("Before Strip:", my_seq)
6 print("After Strip:", my_seq.strip(" "))
7
8 # Strip unwanted characters
9 my_seq = Seq("---ATGCGACCGA-")
10 print("Before Strip:", my_seq)
11 print("After Strip:", my_seq.strip("-"))
```

Please enter sequence: ATGGCGAG Before Strip: ATGGCGAG After Strip: ATGGCGAG

Before Strip: ---ATGCGACCGA-After Strip: ATGCGACCGA



### • Strip Spaces or Other Symbols

- Write and Run the following codes on a Colab page called "SeqObjects.ipynb":





#### (Solution <u>URL</u> of this Practice)

# **BIOLOGICAL OPERATIONS FOR SEQUENCES**



## Ungap a Sequence

# What is "Ungap"?

- Return a copy of the sequence without the gap character(s).





```
from Bio.Seg import Seg
 2
3
   # Ungapped a Sequence
    my seq = Seq("-ATA--TGAAAT-TTGAAAA")
 4
5
   print("Before Ungapped:", my seq)
    print("After Ungapped:", my_seq.ungap())
 6
    # Ungapped with specific character
8
    my_seq = Seq("CGGGTAG=AAAAAA")
 9
    print("Before Ungapped:", my seq)
10
   print("After Ungapped:", my_seq.ungap("="))
11
```

Before Ungapped: -ATA--TGAAAT-TTGAAAA After Ungapped: ATATGAAATTTGAAAA

Before Ungapped: CGGGTAG=AAAAAA After Ungapped: CGGGTAGAAAAAA



### • Ungap a Sequence

- Write and Run the following codes on a Colab page called "SeqObjects.ipynb":

```
from Bio.Seq import Seq
 2
 3
   # Ungapped a Sequence
    my_seg = Seg("-ATA--TGAAAT-TTGAAAA")
 4
   print("Before Ungapped:", my seq)
 5
 6
   print("After Ungapped:", my_seq.ungap())
 7
   # Ungapped with specific character
 8
    my_seq = Seq("CGGGTAG=AAAAAA")
9
   print("Before Ungapped:", my_seq)
10
   print("After Ungapped:", my_seq.ungap("="))
11
```



# • What is Transcription?

DNA coding strand (aka Crick strand, strand +1)

5' ATGGCCATTGTAATGGGCCGCTGAAAGGGTGCCCGATAG 3'

......

3' TACCGGTAACATTACCCGGCGACTTTCCCACGGGCTATC 5'

DNA template strand (aka Watson strand, strand -1)



5' AUGGCCAUUGUAAUGGGCCGCUGAAAGGGUGCCCGAUAG 3'

Single stranded messenger RNA

### Forward Transcription



### Backward Transcription



Backward Transcribe on Protein Sequence will be ignored

Backward Transcription of Protein Sequence: MAIVMGR



- Write and Run the following codes on a Colab page called "SeqObjects.ipynb":

```
1 from Bio.Seq import Seq
 2
3 # Create a Sequence
4 coding dna = Seq("ATGGCCATTGTAATGGGCCGCTGAAAGGGTGCCCGATAG") # 5'→3'
 5 template dna = coding dna.complement() # 3'→5'
   mRNA = coding dna.transcribe()
 6
 7
8 # Forward Transcription
9 print("5' {} 3' : Coding Strand".format(coding dna))
10 print("3' {} 5' : Template Strand".format(template dna))
11 print("5' {} 3' : mRNA".format(mRNA))
12
13 # Transcribe Protein Sequence
14 print("Transcribe Protein Sequence:", Seq("MAIVMGR").transcribe())
15
16 # Backward Transcription
17 print("5' {} 3' : mRNA".format(mRNA))
18 print("5' {} 3' : Backward Transcription".format(mRNA.back_transcribe()))
19
20 # Backward Transcription of Protein Sequence
21 print("Backward Transcription of Protein Sequence:", Seq("MAIVMGR").back_transcribe())
```





#### Translation Table (from NCBI)

1. The standard code 2. The vertebrate mitochondrial code 3. The yeast mitochondrial code 4. The mold, protozoan, and coelenterate mitochondrial code and the mycoplasma/spiroplasma code 5. The invertebrate mitochondrial code 6. The ciliate, dasycladacean and hexamita nuclear code 7. The kinetoplast code 9. The echinoderm and flatworm mitochondrial code 10. The euplotid nuclear code 11. The bacterial, archaeal and plant plastid code 12. The alternative yeast nuclear code 13. The ascidian mitochondrial code 14. The alternative flatworm mitochondrial code 15. The Blepharisma nuclear code 16. The chlorophycean mitochondrial code 21. The trematode mitochondrial code 22. The Scenedesmus obliguus mitochondrial code 23. The Thraustochytrium mitochondrial code 24. The Pterobranchia mitochondrial code 25. The candidate division SR1 and gracilibacteria code 26. The Pachysolen tannophilus nuclear code 27. The karyorelict nuclear code 28. The Condylostoma nuclear code 29. The Mesodinium nuclear code 30. The peritrich nuclear code 31. The Blastocrithidia nuclear code 33. The Cephalodiscidae mitochondrial code

• Syntax of .translate() Function

.translate(table='Standard', stop\_symbol='\*', to\_stop=False, cds=False)

- **<u>table</u>**: Specify which codon table to use (string or integer)
- **<u>stop\_symbol</u>**: Specify which character to represent the stop codon.
- **to\_stop**: Translation is **stopped** at the **first stop codon** or not.
- <u>cds</u>: Sequence is <u>started</u> with <u>start</u> codon, <u>ended</u> with <u>stop</u> codon, and is a <u>multiple</u> of 3 or not.

• Translate with DNA + Standard Table

```
from Bio.Seg import Seg
 1
 2
 3
    # Create a Sequence
    coding_dna = Seq("ATGGCCATTGTGATGGGCCGCTGAAAGGGTGCCCGATAG")
 4
    print("5' {} 3' : Coding Strand".format(coding_dna))
 5
                                                                                 Create a Sequence
 6
                                                5' ATGGCCATTGTGATGGGCCGCTGAAAGGGTGCCCGATAG 3' : Coding Strand
    mRNA = coding dna.transcribe()
 7
                                                5' AUGGCCAUUGUGAUGGGCCGCUGAAAGGGUGCCCGAUAG 3' : mRNA
    print("5' {} 3' : mRNA".format(mRNA))
 8
 9
    # Translate from DNA with the Standard Table
10
                                                               Translate with DNA + Standard Table
    polypeptides = coding_dna.translate()
11
                                                                    Translate (DNA, Standard): MAIVMGR*KGAR*
    print("Translate (DNA, Standard):", polypeptides)
12
```

### Other Operations of Translation

```
# Change the Symbol of Stop Codons
14
    polypeptides = coding_dna.translate(stop_symbol="@")
15
    print("Translate (DNA, Stop='@'):", polypeptides)
16
17
18
    # Translate until Stop Codons
    polypeptides = coding dna.translate(to stop=True)
19
    print("Translate (DNA, Until Stop):", polypeptides)
20
21
    # Translate from mRNA with the Standard Table
22
23
    polypeptides = mRNA.translate()
    print("Translate (RNA, Standard):", polypeptides)
24
```

#### Change the Symbol

Translate (DNA, Stop='@'): MAIVMGR@KGAR@

#### **Translate until Stop Codon**

Translate (DNA, Until Stop): MAIVMGR

#### Change the Symbol

Translate (RNA, Standard): MAIVMGR\*KGAR\*

### Translate with Other Table

```
# Translate with Table=2 (Vertebrate Mitochondrial)
26
    polypeptides = coding_dna.translate(table=2)
27
                                                                      Translate (DNA, Table=2): MAIVMGRWKGAR*
28
    print("Translate (DNA, Table=2):", polypeptides)
29
    # Translate with Table=2, GTG=Start Codon, Fully Coding Sequence (CDS)
30
    sub CDS = coding dna[9:]
                                                     5' GTGATGGGCCGCTGAAAGGGTGCCCGATAG 3' : Sub-Coding Sequence
31
    print("5' {} 3' : Sub-Coding Sequence".format(sub CDS))
32
    # polypeptides = sub CDS.translate(cds=True) # Error: GTG is not a Start Codon
33
    polypeptides = sub CDS.translate(table=2, cds=True)
34
                                                                        Translate (DNA, Table=2, CDS): MMGRWKGAR
35
    print("Translate (DNA, Table=2, CDS):", polypeptides)
```

### **Practice**

```
from Bio.Seg import Seg
     # Create a Sequence
     coding dna = Seq("ATGGCCATTGTGATGGGCCGCTGAAAGGGTGCCCGATAG")
    print("5' {} 3' : Coding Strand".format(coding dna))
    mRNA = coding dna.transcribe()
 7
     print("5' {} 3' : mRNA".format(mRNA))
 8
 9
     # Translate from DNA with the Standard Table
10
11
     polypeptides = coding dna.translate()
    print("Translate (DNA, Standard):", polypeptides)
12
13
14
    # Change the Symbol of Stop Codons
     polypeptides = coding dna.translate(stop symbol="@")
15
    print("Translate (DNA, Stop='@'):", polypeptides)
16
17
18
    # Translate until Stop Codons
     polypeptides = coding dna.translate(to stop=True)
19
    print("Translate (DNA, Until Stop):", polypeptides)
20
21
    # Translate from mRNA with the Standard Table
22
     polypeptides = mRNA.translate()
23
    print("Translate (RNA, Standard):", polypeptides)
24
25
    # Translate with Table=2 (Vertebrate Mitochondrial)
26
     polypeptides = coding dna.translate(table=2)
27
28
    print("Translate (DNA, Table=2):", polypeptides)
29
    # Translate with Table=2, GTG=Start Codon, Fully Coding Sequence (CDS)
30
31
    sub CDS = coding dna[9:]
    print("5' {} 3' : Sub-Coding Sequence".format(sub CDS))
32
    # polypeptides = sub_CDS.translate(cds=True) # Error: GTG is not a Start Codon
33
34
    polypeptides = sub CDS.translate(table=2, cds=True)
    print("Translate (DNA, Table=2, CDS):", polypeptides)
35
```

### Translation

Write and Run the left codes on a Colab page called "SeqObjects.ipynb":





# **MODIFIABLE SEQUENCES**

### **Problem of Modifying Sequence**



Error!

**TypeError**: 'Seq' object does not support item assignment

→ **Bio.Seq.Seq** is an "Immutable" object (Immutable = Not allow to change it partially)



- Test for the Problem of Bio.Seq.Seq Modification
  - Write and Run the following codes on a Colab page called "SeqObjects.ipynb":

```
# Create a Sequence
   from Bio.Seg import Seg
2
    my seq = Seq("GCCATTGTAATGGGCCGCTGAAAGGGTGCCCGA")
3
4
    # Mutation from "T" to "G" at Index = 5 → Error!!
5
    my_seq[5] = "G"
6
TypeError
                                    Traceback (most recent call last)
<ipython-input-37-19a818f9f44a> in <module>()
     5 # Mutation from "T" to "G" at Index = 5 → Error!!
----> 6 my_seq[5] = "G"
TypeError: 'Seq' object does not support item assignment
```



# Solution: Bio.Seq.MutableSeq

- About Bio.Seq.MutableSeq
  - A Sequence object that can be modified partially.

### Possible Operations

<u>Comparison</u>: ==, !=, <, <=, >, >=

Length: len()

Partially Change: seq[5] ='T'

Partially Delete: del seq[5]

**Concatenation (Add)**: seq = seq + "GCG"

**Duplication (Multiple)**: seq = seq \* 3

Insert: seq.insert(8,'G')

Remove: seq.remove('A')

Count: seq.count("ATG")

Reverse: seq.reverse()

<u>Complement</u>: seq.complement() seq.reverse\_complement()

# **Conversion of MutableSeq**

### $\textbf{String} \rightarrow \textbf{MutableSeq}$

str\_seq = "GCCATTGTA"
mutable\_seq = MutableSeq(str\_seq)

### $\textbf{MutableSeq} \rightarrow \textbf{String}$

mutable\_seq = MutableSeq("GCCATTGTA")

str\_seq = str(mutable\_seq)

### $\textbf{Seq} \rightarrow \textbf{MutableSeq}$

seq = Seq("GCCATTGTA")
mutable\_seq = MutableSeq(seq)
mutable\_seq = seq.tomutable()

### $\textbf{MutableSeq} \rightarrow \textbf{Seq}$

mutable\_seq = MutableSeq("GCCATTGTA")

seq = Seq(mutable\_seq)

seq = mutable\_seq.toseq()



### Example for Bio.Seq.MutableSeq

Write and Run the following codes on a Colab page called "SeqObjects.ipynb":



# **BIOLOGICAL FUNCTIONS FOR** STRINGS

### **Biological Functions for Strings**

### • What do these functions do?

 When you want to perform some Biological Operations (e.g., transcribe, translate...etc.) directly on a "string" instead of using Seq or MutableSeq objects.

### • What functions are available?

- complement()
- reverse\_complement()
- transcribe()
- back\_transcribe()
- translate()







### Biological Functions for Strings

- Write and Run the following codes on a Colab page called "SeqObjects.ipynb":

```
from Bio.Seq import complement, reverse_complement
    from Bio.Seq import transcribe, back transcribe, translate
 3
    # Create a Sequence
   str coding = "ATGGCCATTGTAATGGGCCGCTGAAAGGGTGCCCGATAG"
   str template = complement(str coding)
 6
   str template reverse = reverse complement(str coding)
   print("5' {} 3' : Coding Strand".format(str coding))
    print("3' {} 5' : Template Strand".format(str template))
    print("5' {} 3' : Template Strand".format(str template reverse))
10
11
    # Transcription
12
   str mRNA = transcribe(str coding)
13
    print("5' {} 3' : mRNA".format(str mRNA))
14
    print("5' {} 3' : Backward Transcription".format(back transcribe(str mRNA)))
15
16
    # Translation
17
    str protein = translate(str coding, table=2, stop symbol="@")
18
    print("Protein Sequence:", str_protein)
19
```

5' ATGGCCATTGTAATGGGCCGCTGAAAGGGTGCCCGATAG 3' : Coding Strand 3' TACCGGTAACATTACCCGGCGACTTTCCCACGGGCTATC 5' : Template Strand 5' CTATCGGGCACCCTTTCAGCGGCCCATTACAATGGCCAT 3' : Template Strand 5' AUGGCCAUUGUAAUGGGCCGCUGAAAGGGUGCCCGAUAG 3' : mRNA 5' ATGGCCATTGTAATGGGCCGCTGAAAGGGTGCCCGATAG 3' : Backward Transcription Protein Sequence: MAIVMGRWKGAR@





- Parse the FASTA file of E. Coli
  - Please go to the URL: https://bit.ly/3wHQ6lu
  - Read the requirements listed on the above URL.
  - Create your own Colab page and start to implement the requirements.
  - When you finished, click "Share" button at the upper-right.
  - Click "Get Link" > "Copy Link" from the pop-up window.
  - Send your page's link to Google Classroom as the result of this homework.





- Create a Sequence
  - dna\_seq = Seq("ATCG")

#### Basic Operations

- Add: Seq("AATC") + Seq("CGAT")
- Multiple: Seq("AATC") \* 3
- Compare: Seq("ATCG") != Seq("GGCC")
- Slicing: Seq("GATCGA")[0:4:2]
- Iterate:
  - for letter in my\_seq:
  - for index, letter in enumerate(my\_seq):

#### Functions Applying for Sequences

- Length: len(my\_seq)
- Case: my\_seq.lower(), my\_seq.upper()
- Contain: Seq("AAA") in my\_dna
- Count:
  - Non-overlapped: my\_seq.count("AA")
  - Overlapped: my\_seq.count\_overlap("AA")
  - GC%: Bio.SeqUtils.GC(my\_seq)
- Find: find(), rfind(), startswith(), endswith()
- Split & Join:
  - Pre\_mRNA.split("GGCC")
  - "".join([Seq('AU'), Seq('AUUGUT')])
- Strip: my\_seq.strip("-")





#### Biological Operations

- Ungap: my\_seq.ungap("-")
- Transcription:
  - coding\_dna.transcribe()
  - protein.back\_transcribe()
- Translation: coding\_dna.translate(table=2, stop\_symbol="@")

#### Bio.Seq.MutableSeq

- Create: mutable\_seq = MutableSeq(my\_seq)
- Mutation: mutable\_seq[5] = "G"
- Reverse: mutable\_seq.reverse()

#### Biological Functions for Strings

- complement()
- reverse\_complement()
- transcribe()
- back\_transcribe()
- translate()

