

## ***Biopython lesson 2***

**\* What? Does it tell ?**

**ATGGCCATTGTAATGGGCCGCTGAAAGGGTGCCCGATAG** DNA sequence **protein encoder one gene coding** This DNA sequence is part of

**transcription and translation** process as a result **amino acid to the chain** turns into .

So one result is taken :

ATGGCCATTGTAATGGGCCGCTGAAAGGGTGCCCGATAG (DNA) ↓ Transcription (mRNA synthesis )

AUGGCCAUUGUAAUGGGCCGCUGAAAGGGUGCCCGAUAG (mRNA) ↓ Translation (Amino acid sequence ) MAIVMGR\*KGAR\*

Here **\* sign STOP codon** nu ( stop codon ) indicates .

# STOP Codons What is it ?

Protein synthesis **codons ( triplets) nucleotide groups )** through management Genetic in the code **3 types of STOP codons** there is :

•**UAA ( Ochre )**

•**UAG ( Amber )**

•**UGA ( Opal )**

these codons one ribosome by when read , **protein synthesis stands .**

## STOP Codon How Is it happening ?

Your DNA sequence let's see :

ATGGCCATTGTAATGGGCCGCTGAAAGGGTGCCCGATAG (DNA)

AUGGCCAUUGUAAUGGGCCGCUGAAAGGGUGCCCGAUAG (mRNA)

Here **The UAG codon is** a STOP codon .

**UAG ("Amber Stop Codon ")** is called and **protein synthesis stops .**

### Summary

✓ \* **STOP codon means .**

✓ **UAA, UAG, UGA protein synthesis stopping are codons .**

✓ Of these codons one As for **the ribosome , protein completes and him/her free leaves .**

That is , the \* **symbol indicates protein synthesis. there that it's over show**

**Genetic Code : Standard Code and Vertebrates Mitochondrial Code Between Differences**

Genetic code , **DNA and or in RNA codons ( triplets) nucleotides )** which one amen to acids suitable that he came certain who rules is a collection of .

**Different organisms and organelles ( e.g. , mitochondria ) various genetic from codes use can can .**

Genetics in **Biopython** code tables **transl\_table** with is shown .

The most very use done two code :

- ❖ **Standard Genetic Code (The Standard Code, transl\_table =1)**
- ❖ **Vertebrates Mitochondrial Code (The Vertebrate Mitochondrial Code, transl\_table =2)**

**Home Differences**

Feature	Standard Code (Table 1)	Mitochondrial Code (Table 2)
Start Codon	AUG (M)	AUG, AUA, AUU (M)
Stop Codons	UAA, UAG, UGA	UAA, UAG (but UGA = W)
UGA codon	STOP	Tryptophan (W)
AUA codon	Isoleucine (I)	Methionine (M)

## 1. Standard Genetic Code ( transl\_table =1)

All cell in the nucleus ( nucleus genome ) use done main code This is it .

Bacteria , archaea and nuclear to the genome possessor which **animal , plant and mushrooms** by use is being done .

✦ **Features :**

✓ **Start codon :** AUG (Methionine, M)

✓ **Stop codons :** UAA, UAG, UGA

✓ All other codons **standard amen acids encodes** .

## 2. Vertebrates Mitochondrial Code ( transl\_table =2)

In mitochondria different evolution way because according to **mitochondrial DNA ( mtDNA )** from nuclear DNA different genetic from the code use does .

✦ **Features :**

✓ **Start codons :** AUA, AUG, AUU ( all encode Methionine, M )

✓ **Stop codons is different :**

• **UGA stop codon not , Tryptophan (W) coding !**

• **Stop codons only UAA and UAG .**

## Biopython with Usage

You Biopython with **various translation from the tables use can you can** .

### Standard Code Translation (Table 1)

```
from Bio.Seq import Seq dna = Seq ("ATGGCCATTGTAATGGGCCGCTGAAAGGGTGCCCGATAG") protein = dna.translate (table=1) print(protein)
```

✦ **Exit :**

markdown

MAIVMGR\*KGAR\*

Here \* STOP codon shows .

## Conclusion

✓ **Standard code** cell in the core use ✓ Mitochondrial

**code** in mitochondria use is done and some differences there is :

• **UGA encodes W (Tryptophan), not STOP .**

• **encodes Methionine (M) , in nuclear DNA and Isoleucine (I).**

• **Stop codons It's just UAA and UAG .**

### Mitochondrial Code Translation (Table 2)

```
protein_mito = dna.translate (table=2) print( protein_mito )
```

✦ **Difference :**

If the UGA codon If there is , **it will be W (Tryptophan) instead of STOP .**

## Mitochondrial Code Translation Result : MAIVMGRWKGAR\* What Does that mean ?

Mitochondrial genetic from the code (Vertebrate Mitochondrial Code) use by doing broadcast Have you done it ?

```
print( coding_dna.translate (table="Vertebrate Mitochondrial"))
```

Output :

MAIVMGRWKGAR\*

```
#back-transcription method
messenger_rna=Seq("AUGGCCAUUGUAAUGGGCCGCUGAAAGGGUGCCCGAUAG")
print(messenger_rna)
print(messenger_rna.back_transcribe())
print(messenger_rna.translate())
#ATGGCCATTGTAATGGGCCGCTGAAAGGGTGTCCCGATAG burdan ede bilersen...
```

```
AUGGCCAUUGUAAUGGGCCGCUGAAAGGGUGCCCGAUAG
ATGGCCATTGTAATGGGCCGCTGAAAGGGTGTCCCGATAG
MAIVMGR*KGAR*
```

### 1. Standard Code with Difference What is it ?

Previous standard genetic in the code output so was :

MAIVMGR\*KGAR\*

But **mitochondrial code** result :

MAIVMGRWKGAR\*

Here **main difference W (Tryptophan) amine It is acidic .**

## 2. The UGA Codon Encodes

Standard UGA **STOP** codon in the code where for **protein early stands** .

But **vertebrates mitochondrial in the code** UGA is not **STOP** , instead **Tryptophan (W)** encodes .

**DNA sequence :**

ATGGCCATTGTAATGGGCCGCTGAAAGGGTGCCCGATAG (DNA)

AUGGCCAUUGUAAUGGGCCGCUGAAAGGGUGCCCGAUAG ( mRNA )

**Standard code translation :**

MAIVMGR\*KGAR\* (UGA STOP )

**Mitochondrial code translation :**

MAIVMGRWKGAR\* (UGA → W)

Here UGA is no longer a **STOP** codon No , it encodes **Tryptophan (W)**

```
#veya
coding_dna=Seq("ATGGCCATTGTAATGGGCCGCTGAAAGGGTGCCCGATAG")
print(coding_dna)
print(coding_dna.translate())
```

```
ATGGCCATTGTAATGGGCCGCTGAAAGGGTGCCCGATAG
MAIVMGR*KGAR*
```

```
print(coding_dna.translate(table="Vertebrate Mitochondrial"))
#veya
coding_dna.translate(table=2)
```

```
MAIVMGRWKGAR*
Seq('MAIVMGRWKGAR*')
```

## 3. Conclusion

✓ **Vertebrate mitochondrial code** UGA codon **Like STOP** no , like **Tryptophan (W)** is reading .

✓ For this reason standard code received **Instead of STOP** in the result , **W (Tryptophan)** appears .

✓ Other codons the same remains .

To him/her according to also the output becomes **MAIVMGRWKGAR\*** .

## Deletion and addition of stop codons

```
print(coding_dna.translate())  
#Seq("MAIVMGR*KGAR*")  
print(coding_dna.translate(to_stop=True))
```

```
MAIVMGR*KGAR*  
MAIVMGR
```

```
print(coding_dna.translate(table=2))
```

```
MAIVMGRWKGAR*
```

```
print(coding_dna.translate(table=2, to_stop=True))
```

```
MAIVMGRWKGAR
```

This code **BioPython** from the library use by doing **DNA sequence ( coding\_dna )** amen to **acids translates** ( translates) does ).



```
print( coding_dna.translate ())
```

→ **Standard genetic DNA sequence with code** amen to acids turns into .

**Output :**

MAIVMGR\*KGAR\*

**Asterisk (\*) indicates STOP codons shows .**

STOP codon , protein synthesis where it stands It is a place .

```
print( coding_dna.translate ( to_stop =True))
```

→ using to\_stop =True when done , from **the STOP codon then coming part is thrown away .**

**Output :**

MAIVMGR

This is where **the protein is completely stopped. to the ground until which part shows .**

Alternative Genetic Code (Table=2)

```
print( coding_dna.translate (table=2))
```

→ Genetic code table=2 is selected (Vertebrate Mitochondrial Code)

Output :

MAIVMGRWKGAR\*

Here UGA encodes Tryptophan (W) instead of STOP .

```
print( coding_dna.translate (table=2, to_stop =True))
```

→ Here mitochondrial with code to STOP converts .

Output :

MAIVMGRWKGAR

the STOP codon next part is thrown away .

```
print(coding_dna.translate())
#Seq( "MAIVMGR*KGAR*")
print(coding_dna.translate(to_stop=True))
print(coding_dna.translate(table=2))
print(coding_dna.translate(table=2, to_stop=True))
```

MAIVMGR\*KGAR\*  
MAIVMGR  
MAIVMGRWKGAR\*  
MAIVMGRWKGAR

🔍 STOP Codon (\*) Why Is it visible ?

DNA triplet with codons is read and some codons protein synthesis that it's over ( STOP codon ).

The main STOP codons are:

- TAA, TAG, TGA ( Standard in the code )
- TGA → W (Tryptophan) ( Mitochondrial in the code )

🔥 If using to\_stop =True if not , STOP codons like \* at the exit it seems .

🔥 If to\_stop =True added if we do , from the STOP codon next part is thrown away .

```
coding_dna.translate(table=2, stop_symbol="@")

Seq( 'MAIVMGRWKGAR@' )
```

## Full encoder sequence (CDS – Complete Coding Sequence) and standard missing start codons

**What is CDS (Coding Sequence) ?**

**CDS is a complete protein encoder genetic is a sequence :**

- ✓ **Its length is divisible by 3** ( each one one amen acid encoder threesome from codons consists of ).
- ✓ **With start codon (ATG) begins** ( protein synthesis start point ).
- ✓ **With stop codon (TAA, TAG, TGA) ends** ( protein synthesis stop point ).
- ✓ **STOP codon inside does not exist** ( if if so , protein unfinished remains ).

**This kind of sequence if any , translate() method with comfortable in the way to protein turns into .**

## **Standard Missing Start Codon What is it ?**

**In bacteria some genes instead of ATG ( Methionine ) other with codons to begin can .**

**For example , in E. coli K12 , the " yaaX " gene has a GTG ( Valine ) instead of ATG . begins .**

## **Standard Missing Start Codon Problem with**

**If so one translate() method in sequence use if k , the previous codons other amen acids with start over .**

**Standard ATG gives Methionine ( M) , but some bacteria GTG and or TTG codons as start use does .**

**Bacteria for specific genetic from the code** use when you do :

```
print( coding_dna.translate (table=11)) # Bacteria and Archaea for genetic code
```

**At this time some non-standard start codons such as ATG will be read .**

## **Conclusion**

- ✦ **A complete one encoder sequence (CDS) if any , translate() method correct will work .**
- ✦ **In bacteria and some in organisms standard missing start codons use is being done .**
- ✦ **Bacteria genes correct to read for table=11 like suitable genetic codes should be selected .**

```
[70]: from Bio.Seq import Seq
gene=Seq("GTGAAAAGATGCAATCTATCGTACTCGCAC TTTCCCTGGTTCTGGTCGCTCCCATGGCA"
        "GCACAGGCTGCGGAAATTACGTTAGTCCCGTCAGTAAAATTACAGATAGGCGATCGTGAT"
        "AATCGTGGCTATTACTGGGATGGAGGTCAC TGGCGCGACCACGGCTGGTGGAAACAACAT"
        "TATGAATGGCGAGGCAATCGCTGGCACCTACACGGAECGCCGCCACCGCCGCCACCAT"
        "AAGAAAGCTCCTCATGATCATCACGGCGGTCATGGTCCAGGCAAACATCACCGCTAA")
print(gene)
```

```
GTGAAAAGATGCAATCTATCGTACTCGCAC TTTCCCTGGTTCTGGTCGCTCCCATGGCAGCACAGGCTGCGGAAATTACGTTAGTCCCGTCAGTAAAATTACAGATAGGCGATCGTGATAATCGTGGCTATTACTGGGATGGAGGTCAC TGGCGCGACCACGGCTGGTGGAAACAACATTATGAATGGCGAGGCAATCGCTGGCACCTACACGGAECGCCGCCACCGCCGCCACCAT AAGAAAGCTCCTCATGATCATCACGGCGGTCATGGTCCAGGCAAACATCACCGCTAA
```

```
[72]: print(gene.translate(table="Bacterial"))
```

```
VKKMQSIVLALS LVLVAPMAAQAAEITLVPSVKLQIGDRDNRGYYWDGGHWRDHGWWKQHYEWRGNRWHLHGPPPPRHKKAPHDHHGGHGP GKHR*
```

```
[74]: gene.translate(table="Bacterial", to_stop=True)
```

```
[74]: Seq('VKKMQSIVLALS LVLVAPMAAQAAEITLVPSVKLQIGDRDNRGYYWDGGHWRDH...HHR')
```

```
[76]: gene.translate(table="Bacterial", cds=True)
```



```
[76]: Seq('MKKMQSIVLALS LVLVAPMAAQAAEITLVPSVKLQIGDRDNRGYYWDGGHWRDH...HHR')
```

```
[84]: print(gene)
print(gene.translate(table="Bacterial"))
print(gene.translate(table="Bacterial", to_stop=True))
print(gene.translate(table="Bacterial", cds=True))
```

```
GTGAAAAGATGCAATCTATCGTACTCGCAC TTTCCCTGGTTCTGGTCGCTCCCATGGCAGCACAGGCTGCGGAAATTACGTTAGTCCCGTCAGTAAAATTACAGATAGGCGATCGTGATAATCGTGGCTATTACTGGGATGGAGGTCAC TGGCGCGACCACGGCTGGTGGAAACAACATTATGAATGGCGAGGCAATCGCTGGCACCTACACGGAECGCCGCCACCGCCGCCACCAT AAGAAAGCTCCTCATGATCATCACGGCGGTCATGGTCCAGGCAAACATCACCGCTAA
VKKMQSIVLALS LVLVAPMAAQAAEITLVPSVKLQIGDRDNRGYYWDGGHWRDHGWWKQHYEWRGNRWHLHGPPPPRHKKAPHDHHGGHGP GKHR*
VKKMQSIVLALS LVLVAPMAAQAAEITLVPSVKLQIGDRDNRGYYWDGGHWRDHGWWKQHYEWRGNRWHLHGPPPPRHKKAPHDHHGGHGP GKHR
MKKMQSIVLALS LVLVAPMAAQAAEITLVPSVKLQIGDRDNRGYYWDGGHWRDHGWWKQHYEWRGNRWHLHGPPPPRHKKAPHDHHGGHGP GKHR
```

```
[100]: print (gene.translate(table="Bacterial"))  
#Seq('VKKMQSIVLALSLVLVAPMAAQAAEITLVPSVKLQIGDRDNRGYYWDGGHWRDH...HR*',ProteinAlpabet())  
  
VKKMQSIVLALSLVLVAPMAAQAAEITLVPSVKLQIGDRDNRGYYWDGGHWRDHGWKQHYEWRGNRWHLHGPPPPRHHKKAPHDHHGGHGP GKHHR*
```

```
[104]: gene.translate(table="Bacterial", to_stop=True)  
Seq('VKKMQSIVLALSLVLVAPMAAQAAEITLVPSVKLQIGDRDNRGYYWDGGHWRDH...HHR')  
#Bakterial genetik kodda GTG etibarlı başlanğıc kodonudur və  
#normal olaraq Valini kodlasa da, başlanğıc kodonu kimi istifadə edilərsə,  
#metionin kimi tərcümə edilməlidir.
```

```
[104]: Seq('VKKMQSIVLALSLVLVAPMAAQAAEITLVPSVKLQIGDRDNRGYYWDGGHWRDH...HHR')
```

```
[106]: gene.translate(table="Bacterial", cds=True)  
Seq('MKKMQSIVLALSLVLVAPMAAQAAEITLVPSVKLQIGDRDNRGYYWDGGHWRDH...HHR')
```

```
[106]: Seq('MKKMQSIVLALSLVLVAPMAAQAAEITLVPSVKLQIGDRDNRGYYWDGGHWRDH...HHR')
```

```
[112]: #Translation tables
#Standart translation cadvali va Vertebrate Mitochondrial DNT üçün translation cadvali.
from Bio.Data import CodonTable
standard_table = CodonTable.unambiguous_dna_by_name["Standard"]
mito_table = CodonTable.unambiguous_dna_by_name["Vertebrate Mitochondrial"]
print(standard_table)
print(mito_table)
```

This table **genetic code** is the schedule and **mRNA codons** which one amen to **acids** that it has become shows .

Genetic code in the table every one amen acid **one** literally representat **being** :

Table 1 Standard, SGC0

	T	C	A	G	
T	TTT F	TCT S	TAT Y	TGT C	T
T	TTC F	TCC S	TAC Y	TGC C	C
T	TTA L	TCA S	TAA Stop	TGA Stop	A
T	TTG L(s)	TCG S	TAG Stop	TGG W	G
C	CTT L	CCT P	CAT H	CGT R	T
C	CTC L	CCC P	CAC H	CGC R	C
C	CTA L	CCA P	CAA Q	CGA R	A
C	CTG L(s)	CCG P	CAG Q	CGG R	G
A	ATT I	ACT T	AAT N	AGT S	T
A	ATC I	ACC T	AAC N	AGC S	C
A	ATA I	ACA T	AAA K	AGA R	A
A	ATG M(s)	ACG T	AAG K	AGG R	G
G	GTT V	GCT A	GAT D	GGT G	T
G	GTC V	GCC A	GAC D	GGC G	C
G	GTA V	GCA A	GAA E	GGA G	A
G	GTG V	GCG A	GAG E	GGG G	G

Table 2 Vertebrate Mitochondrial, SGC1

	T	C	A	G	
T	TTT F	TCT S	TAT Y	TGT C	T
T	TTC F	TCC S	TAC Y	TGC C	C
T	TTA L	TCA S	TAA Stop	TGA W	A
T	TTG L	TCG S	TAG Stop	TGG W	G
C	CTT L	CCT P	CAT H	CGT R	T
C	CTC L	CCC P	CAC H	CGC R	C
C	CTA L	CCA P	CAA Q	CGA R	A
C	CTG L	CCG P	CAG Q	CGG R	G
A	ATT I(s)	ACT T	AAT N	AGT S	T
A	ATC I(s)	ACC T	AAC N	AGC S	C
A	ATA M(s)	ACA T	AAA K	AGA Stop	A
A	ATG M(s)	ACG T	AAG K	AGG Stop	G
G	GTT V	GCT A	GAT D	GGT G	T
G	GTC V	GCC A	GAC D	GGC G	C
G	GTA V	GCA A	GAA E	GGA G	A
G	GTG V(s)	GCG A	GAG E	GGG G	G

Symbol	Amino acid	Full name
<b>D</b>	Aspartic acid	Aspartate (Asp)
<b>E</b>	Glutamic acid	Glutamate (Glu)
<b>K</b>	Lysine	Lysine (Lys)
<b>Q</b>	Glutamine	Glutamine (Gln)
<b>H</b>	Histidine	Histidine (His)
<b>R</b>	Arginine	Arginine (Arg)
<b>W</b>	Tryptophan	Tryptophan (Trp)
<b>L</b>	Leysin	Leucine (Leu)
<b>F</b>	Phenylalanine	Phenylalanine (Phe)
<b>S</b>	Cool	Serine (Ser)
<b>A</b>	Alanine	Alanine (Ala)
<b>M</b>	Methionine	Methionine (Met)
<b>V</b>	Valine	Valine (Val)



❖ **unambiguous\_dna\_by\_name** what does it ?

Biopython in the library **CodonTable.unambiguous\_dna\_by\_name** method **uncertain codons non-** ( i.e., not completely accurate ) **genetic code tables to call for use is being done .**

This means that that **only standard codons use is being and "ambiguous"** ( i.e. indefinite ( bases ) **There is no** . For example :

- A, T, G, C** The bases are unambiguous .
- Ambiguous bases** while **N, Y, R , etc.** like certain not done nucleotides ( for example , **N = A/T/G/C** ) **can** ).

**from Bio.Data import CodonTable**

```
standard_table = CodonTable.unambiguous_dna_by_name ["Standard"]  
mito_table = CodonTable.unambiguous_dna_by_name ["Vertebrate Mitochondrial"]
```

✓ This code two different genetic code schedule calls :

Standard genetic code ) – all in living things , mainly nuclear DNA for use is being done .  
Vertebrate Mitochondrial code ) – in mitochondria use is being and some different codon have meanings .

- ❖ **L(s) → Start codon be knowing leucine**
- ❖ **M(s ) → Start codon be knowing methionine**

- Normally **AUG codon For Methionine (M)** use is being and beginning codon (start codon ) exit does .
- However some in bacteria **L(s) and or M(s)** beginning codon be can .

- **TAA, TAG, TGA → are Stop codons (translation stops).**
- **ATG → Methionine (M) and is also a Start codon.**

One **codon , 3 nucleotides consisting of one is a group** and one amen to acid suitable comes . For example :

- **AUG → Methionine (M)**
- **UUU → Phenylalanine (F)**
- **UGA → Stop ( translate) stops )**

" Ambiguous" genetics codes What is it ?

Uncertain (ambiguous) codons which genetic codes are also used to be done can . This for ambiguous\_dna\_by\_name There is a method .

For example :

python

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```
from Bio.Data import CodonTable
```

```
ambiguous_table = CodonTable.ambiguous_dna_by_name ["Standard"]
```

```
print( ambiguous_table )
```

This version uncertain bases are also accepted ( for example , N, Y, R ) codes ).

✓ "Unambiguous" means that only from bases A, T, G, C use is done .

✓ unambiguous\_dna\_by\_name exactly and fully determined done genetic code tables to call for use is being done .

```
[ ]: #stop ve start codonlari...
      mito_table.stop_codons
      ['TAA', 'TAG', 'AGA', 'AGG']
      mito_table.start_codons
      ['ATT', 'ATC', 'ATA', 'ATG', 'GTG']
      mito_table.forward_table["ACG"]
      'T'
```

```
[114]: print(mito_table.stop_codons)
```

```
['TAA', 'TAG', 'AGA', 'AGG']
```

```
[116]: print(mito_table.start_codons)
```

```
['ATT', 'ATC', 'ATA', 'ATG', 'GTG']
```

```
[118]: print(mito_table.forward_table)
```

```
{'TTT': 'F', 'TTC': 'F', 'TTA': 'L', 'TTG': 'L', 'TCT': 'S', 'TCC': 'S', 'TCA': 'S', 'TCG': 'S', 'TAT': 'Y', 'TAC': 'Y', 'TGT': 'C', 'TGC': 'C', 'TGA': 'W', 'TGG': 'W', 'CTT': 'L', 'CTC': 'L', 'CTA': 'L', 'CTG': 'L', 'CCT': 'P', 'CCC': 'P', 'CCA': 'P', 'CCG': 'P', 'CAT': 'H', 'CAC': 'H', 'CAA': 'Q', 'CAG': 'Q', 'CGT': 'R', 'CGC': 'R', 'CGA': 'R', 'CGG': 'R', 'ATT': 'I', 'ATC': 'I', 'ATA': 'M', 'ATG': 'M', 'ACT': 'T', 'ACC': 'T', 'ACA': 'T', 'ACG': 'T', 'AAT': 'N', 'AAC': 'N', 'AAA': 'K', 'AAG': 'K', 'AGT': 'S', 'AGC': 'S', 'GTT': 'V', 'GTC': 'V', 'GTA': 'V', 'GTG': 'V', 'GCT': 'A', 'GCC': 'A', 'GCA': 'A', 'GCG': 'A', 'GAT': 'D', 'GAC': 'D', 'GAA': 'E', 'GAG': 'E', 'GGT': 'G', 'GGC': 'G', 'GGA': 'G', 'GGG': 'G'}
```

```
[ ]:
```

# Seq objects comparison making

```
[128]: #Seq obyektlərinin müqayisə edilməsi
```

```
from Bio.Seq import Seq
seq1 = Seq("ACGT")
"ACGT" == seq1
print("ACGT"==seq1)
print(seq1=="ACGT")
```

```
True
```

```
True
```

## Sequence unknown which sequences

Some bioinformatics in files ( e.g. , **GenBank** and **EMBL** ) sometimes **complete genomic DNA sequence is not given** , on the contrary , **only configuration ( structure ) information exists** . This means that that the **sequence length known** , but specific as which one consisting of letters ( **nucleotides** ) where not shown .

What is this ? for use is it ?

- All to show the genome in full necessary not – just length to know enough does .
- Information big when to memory savings to do for use to be done can .
- Sequence yet known not and either investigation is in process . So , Seq (None, length=1000) is a sequence created it happens , but inside is empty and only 1000 nucleotides consisting of where It is known .

```
from Bio.Seq import Seq
seq_unknown = Seq(None, length=1000) # 1000 uzunluğunda, lakin içi boş olan bir Seq obyekti
print("seq_unknown")
print(len("seq_unknown")) # 1000 çap edəcək
```

```
seq_unknown
```

```
11
```

# MAF (Multiple Alignment Format)

- **MAF (Multiple Alignment Format)** is a format used in bioinformatics to compare multiple genome sequences .
- **the alignment of DNA sequences** of different organisms .
- It is used to find genetic evolution, mutations, and orthologous genes.

## Structure Explanation

Every in line different one genomics of the organism There is information .

Here **human (hg38)** , **chimpanzee (panTro4)** , **macaque (rheMac3)** , **mouse (mm10)** , **rat (rn5)** , **dog (canFam3)** and **opossum (monDom5)** genomes given .

Every of the line format is like this :

```
#MAF (Multiple Alignment Format)
s hg38.chr7      117512683 36 + 159345973 TTGAAAACCTGAATGTGAGAGTCAGTCAAGGATAGT
s panTro4.chr7   119000876 36 + 161824586 TTGAAAACCTGAATGTGAGAGTCACTCAAGGATAGT
s rheMac3.chr3   156330991 36 + 198365852 CTGAAATCCTGAATGTGAGAGTCAATCAAGGATGGT
s mm10.chr6      18207101 36 + 149736546 CTGAAAACCTAAGTAGGAGAATCAACTAAGGATAAT
s rn5.chr4       42326848 36 + 248343840 CTGAAAACCTAAGTAGGAGAGACAGTTAAAGATAAT
s canFam3.chr14  56325207 36 + 60966679 TTGAAAAACTGATTATTAGAGTCAATTAAGGATAGT
s monDom5.chr8   173163865 36 + 312544902 TTAAGAACTGGAAATGAGGGTTGAATGACAACTT
```

Every of the line format is like this :

s < genome name > < chromosome > < start position > < length > < strand > < total length > <DNA sequence >

**For example let's see :**

s hg38.chr7 117512683 36 + 159345973 TTGAAAACCTGAATGTGAGAGTCAGTCAAGGATAGT

- s → sequence information ( this sell sequence that shows )

- hg38.chr7 → Human genome , chromosome **7**

- 117512683 → This part **chromosome on it beginning position**

- 36 → **Sequence length** (36 nucleotides )

- + → **Strand direction** ( front strand "+" and either reverse strand "-")

- 159345973 → This chromosome general length

- TTGAAAACCTGAATGTGAGAGTCAGTCAAGGATAGT → **DNA sequence**

This is different in types of the same DNA region comparison shows .

- **Human** and **chimpanzee** genomes very is similar .
- **Mouse and rat** in their genomes some There are differences .
- **Opossum** while more very differs .

All this information what for is it necessary ?

- **Genetic evolution to understand** ( which species more is it close ?)
- **Evolution in the process which one mutations head gave ?**
- **Diseases genetic the reasons to investigate**
- **Which genes protected and functional that to understand**
- **This kind of many multiple alignment studies genetic in analyses , especially also evolution , diseases genetic reasons and gene functions to understand for use is being done .**



# MutableSeq

MutableSeq , collection order to change , elements additional did to delete opportunity gives .

MutableSeq objects , usually in Python programming data structures with while working use done and elements to be changed opportunity giving function This type of functions , data on made of changes directly give birth in itself change to do opportunity recognizes .

MutableSeq in Python , in general collections.abc in the module compilation done one abstract This class is a collection class in Python . and lists representation to do for use done one main is a class . MutableSeq , a list representation does and elements on directly change to do opportunity gives ( additional create , delete , update etc. ).

```
[160]: #MutableSeq objects
from Bio.Seq import Seq
my_seq = Seq("GCCATTGTAATGGGCCGCTGAAAGGGTGCCCGA")
from Bio.Seq import MutableSeq
mutable_seq = MutableSeq(my_seq)
MutableSeq('GCCATTGTAATGGGCCGCTGAAAGGGTGCCCGA')
```

```
[160]: MutableSeq('GCCATTGTAATGGGCCGCTGAAAGGGTGCCCGA')
```

```
[162]: mutable_seq[5]="C"
```

```
[164]: print(mutable_seq)
```

```
GCCATCGTAATGGGCCGCTGAAAGGGTGCCCGA
```

```
[166]: #remove funksiyasi T nukleotidini cixaracaq.
mutable_seq.remove("T")
print(mutable_seq)
```

```
GCCACGTAATGGGCCGCTGAAAGGGTGCCCGA
```

```
[170]: #reverse funksiyasi
mutable_seq.reverse()
print(mutable_seq)
```

```
AGCCCGTGGGAAAGTCGCCGGGTAATGCACCG
```

```
[180]: #Stringlerle islemek...  
from Bio.Seq import reverse_complement, transcribe, back_transcribe, translate  
my_string = "GCTGTTATGGGTCGTTGGAAGGGTGGTCGTGCTGCTGGTTAG"  
print(reverse_complement(my_string))  
print(transcribe(my_string))  
print(back_transcribe(my_string))  
print(translate(my_string))
```

```
CTAACCAGCAGCACGACCACCCTTCCAACGACCCATAACAGC  
GCUGUUAUGGGUCGUUGGAAGGGUGGUCGUGCUGCUGGUUAG  
GCTGTTATGGGTCGTTGGAAGGGTGGTCGTGCTGCTGGTTAG  
AVMGRWKGGRAAG*
```

```
[182]: #Alt ardıcılıqların tapılması
from Bio.Seq import Seq, MutableSeq
seq = Seq("GCCATTGTAATGGGCCGCTGAAAGGGTGCCCGA")
print(seq.index("ATGGGCCGC"))
print(seq.index(b"ATGGGCCGC"))
print(seq.index(bytearray(b"ATGGGCCGC"))))
print(seq.index(Seq("ATGGGCCGC"))))
print(seq.index(MutableSeq("ATGGGCCGC"))))

9
9
9
9
9
```

**1.Seq ("GCCATTGTAATGGGCCGCTGAAAGGGTGCCCGA") :** This is Biopython library Seq from the class use by doing one genetic The sequence (GCCATTGTAATGGGCCGCTGAAAGGGTGCCCGA) represents does .

**2.seq.index ("ATGGGCCGC") :** This is Seq the sequence "ATGGGCCGC" in the object beginning finds the index of . The output will be 9 , because the sequence "ATGGGCCGC" seg from position 9 in the facility begins .

**3.seq.index ( b"ATGGGCCGC ") :** Here the sequence "ATGGGCCGC" in bytes format given ( i.e. b"ATGGGCCGC "). Biopython also accepts bytes format does and this is index 9 returns because this sequence also seq from position 9 in the facility begins .

**4.seq.index ( bytearray ( b"ATGGGCCGC ")) :** **This is** the index of the sequence "ATGGGCCGC" byte array in format bytearray , bytes can be changed knowing is version and this is index 9 returns .

**5.seq.index ( Seq ("ATGGGCCGC")) :** Here sequence "ATGGGCCGC". Seq to the object turning seg at the facility This is again index 9 . gives , because Seq objects also adaptable knows .

**6.seq.index ( MutableSeq ("ATGGGCCGC")) :** Here sequence "ATGGGCCGC". MutableSeq to the object turning seg at the facility This is index 9 . returns because MutableSeq also Seq to the object similar in the way works and adaptable knows .

### Result :

These operations shows that , Biopython , various given compare formats (string, bytes, bytearray , Seq , MutableSeq ) and their indexes to find for very is flexible . index() method given sequence representation who of the object of the type dependent regardless of the sequence located index the value returns . In this case , all this operations of the sequence "ATGGGCCGC" seg in the facility index as 9 returns .

```
[198]: for index, sub in seq.search(["CC", "GGG", "CC"]):  
        print(index, sub)  
seq.find("ACTG")  
seq.find("CC")  
seq.rfind("CC")  
#GCCATTGTAATGGGCCGCTGAAAGGGTGCCCGA
```

```
1 CC  
11 GGG  
14 CC  
23 GGG  
28 CC  
29 CC
```

```
[198]: 29
```

This code fragment , **Biopython** library's search() method how that it works This method shows that the given certain sequences (sub- sequences ) one genetic in sequence ( Seq in the object ) which in positions located to find for use is being done .

Code meaning Let's explain :

**1.seq.search ("CC", "GGG", "CC") :**

- The search() method returns the string within the sequence ( seq ). every a sub- sequence ( this in the case of ["CC", "GGG", "CC"] list sequences ) are searched .
- This method , given sequences inside where positions and every finds a sub- sequence .

**2.for index, sub in seq.search ("CC", "GGG", "CC"):**

- Here for loop , search() method turned the result every a sub- sequence and his/her found index taking works .
- index, sub- sequence started index position shows .
- sub is the found sub- sequence .

**3.print(index, sub) :**

- Every of the found sub- sequence index and sub- sequence himself to the screen print does .

**Result :**

This operation seg ["CC", "GGG", "CC"] in the sequence looking for and found every position and sub- sequence print does . In the output :

- CC sub- sequence at indices 1, 14 and 28 found .
- GGG sub- sequence at indices 11 and 23 found .