4-Translation

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1 How to transcribe and translate a DNA sequence

First, we need to import the required modules:

```
[1]: from Bio import SeqIO from Bio.SeqIO.FastaIO import as_fasta
```

Then, we can open the file using with open(<filename>) and parse its content with the SeqIO.read() function:

OBS! SeqIO.read() only works on single records. If we want to read a file with several sequences, we use the function SeqIO.parse() instead.

When we read a fasta file in Biopython, we get an object of class SeqRecord, that we stored in the record variable, and what you can see after printing are the various attributes that this class has:

```
[3]: print(record)
```

```
ID: NC_000913.3:105305-106456

Name: NC_000913.3:105305-106456

Description: NC_000913.3:105305-106456 ftsZ [organism=Escherichia coli str. K-12 substr. MG1655] [GeneID=944786] [chromosome=]

Number of features: 0
```

•

Seq('ATGTTTGAACCAATGGAACTTACCAATGACGCGGTGATTAAAGTCATCGGCGTC...TAA')

```
We can look more into attributes and methods by using dir():
```

```
'__delattr__',
'__dict__',
'__dir__',
'__doc__',
'__eq__',
'__format__',
'__ge__',
'__getattribute__',
'__getitem__',
'__getstate__',
'__gt__',
'__hash__',
'__init__',
'__init_subclass__',
'__iter__',
'__le__',
'__len__',
'__lt__',
'__module__',
'__ne__',
'__new__',
'__radd__',
'__reduce__',
'__reduce_ex__',
'__repr__',
'__setattr__',
'__sizeof__',
'__str__',
'__subclasshook__',
'__weakref__',
'_per_letter_annotations',
'_seq',
'_set_per_letter_annotations',
'_set_seq',
'annotations',
'count',
'dbxrefs',
'description',
'features',
'format',
'id',
'islower',
'isupper',
'letter_annotations',
'lower',
'name',
'reverse_complement',
```

```
'seq',
'translate',
'upper']
```

To check if the parsing worked, let's take a look at one attribute in particular: the **seq** attribute, where the gene sequence is stored:

[5]: print(record.seq)

It worked! But we are looking only at a part of the information that we have. The seq attribute is not the only useful one. In this tutorial, we want to create a new fasta file. In this file type, we have a description as a header and then the sequence, this way:

```
[6]: print(f">{record.description}")
print(record.seq)
```

>NC_000913.3:105305-106456 ftsZ [organism=Escherichia coli str. K-12 substr. MG1655] [GeneID=944786] [chromosome=]

In the example above, we printed the description attribute of the record object as a header, and then the seq attribute. The seq attribute consists of an object of class Seq that includes a string representing the gene sequence. There are several interesting methods that you can apply on a Seq object; for example, you can search for specific sub-sequences inside your sequence. Let's search for the start codon (ATG):

```
[7]: record.seq.find("ATG")
```

[7]: 0

The cell above returned position 0, since, of course, the start codon should be at the start of the sequence. We can also count the number of times a sub-sequence is present in a sequence. For example, let's do that for the stop codon TAA:

```
[8]: record.seq.count("TAA")
```

[8]: 14

It returned 14! How is that possible? Do we have 14 stop codons? Of course not! The sequence is searched in the sequence string without taking into account the codon frames. In this case, because we have a gene, codons start at every third possition, but if we were looking at a full genome, for example, we might have genes overlapping over different codon frames, so this can be useful. We can look into all the methods that are available for the Seq class by using dir():

```
[9]: dir(record.seq)
```

```
[9]: ['__abstractmethods__',
       __add__',
       '__array_ufunc__',
        __bytes__',
       '__class__',
      '__contains__',
      '__delattr__',
       '__dict__',
      '__dir__',
       '__doc__',
        __eq__',
       '__format__',
       __ge__',
       __getattribute__',
       __getitem__',
      '__getstate__',
       '__gt__',
        __hash__'
       __imul__',
      '__init__',
       '__init_subclass__',
       '__iter__',
       '__le__',
```

```
'__len__',
'__lt__',
'__module__',
'__mul__',
'__ne__',
'__new__',
'__radd__',
'__reduce__',
'__reduce_ex__',
'__repr__',
'__rmul__',
'__setattr__',
'__sizeof__',
'__slots__',
'__str__',
'__subclasshook__',
'__weakref__',
'_abc_impl',
'_data',
'back_transcribe',
'complement',
'complement_rna',
'count',
'count_overlap',
'defined',
'defined_ranges',
'endswith',
'find',
'index',
'islower',
'isupper',
'join',
'lower',
'lstrip',
'replace',
'reverse_complement',
'reverse_complement_rna',
'rfind',
'rindex',
'rsplit',
'rstrip',
'split',
'startswith',
'strip',
'transcribe',
'translate',
'ungap',
```

'upper']

Here you can see many other useful attributes and, more commonly, methods. Some of these methods are shared with strings, such as split, strip or startwith. However, there are others that are unique to Seq objects. For this tutorial, we will try some of them to transform the DNA sequence into protein. First, the DNA sequence can be converted to RNA by using the transcribe() function:

```
[10]: RNA_seq = record.seq.transcribe() # Transcribe (DNA > RNA) the sequence print(RNA_seq) # Print the RNA sequence
```

Then, we can easily convert the DNA or RNA sequence into protein by using the translate() function:

```
[11]: protein_seq = RNA_seq.translate() # Overwrite the DNA sequence with the translation (DNA > RNA > protein)
print(protein_seq) # Print the protein sequence
```

MFEPMELTNDAVIKVIGVGGGGGNAVEHMVRERIEGVEFFAVNTDAQALRKTAVGQTIQIGSGITKGLGAGANPEVGRNA
ADEDRDALRAALEGADMVFIAAGMGGGTGTGAAPVVAEVAKDLGILTVAVVTKPFNFEGKKRMAFAEQGITELSKHVDSL
ITIPNDKLLKVLGRGISLLDAFGAANDVLKGAVQGIAELITRPGLMNVDFADVRTVMSEMGYAMMGSGVASGEDRAEEAA
EMAISSPLLEDIDLSGARGVLVNITAGFDLRLDEFETVGNTIRAFASDNATVVIGTSLDPDMNDELRVTVVATGIGMDKR
PEITLVTNKQVQQPVMDRYQQHGMAPLTQEQKPVAKVVNDNAPQTAKEPDYLDIPAFLRKQAD*

The asterisk at the end represents the **stop codon**. If we don't want the asterisk, we have to specify that the translation should stop at the stop codon:

MFEPMELTNDAVIKVIGVGGGGGNAVEHMVRERIEGVEFFAVNTDAQALRKTAVGQTIQIGSGITKGLGAGANPEVGRNA ADEDRDALRAALEGADMVFIAAGMGGGTGTGAAPVVAEVAKDLGILTVAVVTKPFNFEGKKRMAFAEQGITELSKHVDSL ITIPNDKLLKVLGRGISLLDAFGAANDVLKGAVQGIAELITRPGLMNVDFADVRTVMSEMGYAMMGSGVASGEDRAEEAA EMAISSPLLEDIDLSGARGVLVNITAGFDLRLDEFETVGNTIRAFASDNATVVIGTSLDPDMNDELRVTVVATGIGMDKR PEITLVTNKQVQQPVMDRYQQHGMAPLTQEQKPVAKVVNDNAPQTAKEPDYLDIPAFLRKQAD

OBS! The translate function can be used directly on DNA sequences, but here we used the transcribe() function first to test more methods.

Now, we can save the output to a fasta file by using the **as_fasta()** function that we imported in the beginning:

There are alternatives to using this function; for example, you can use the **SeqIO.write()** function and specify the output file type:

```
[17]: out_file = "inputs/tutorial5/ftsZ.faa"

protein_record = record

protein_record.seq = protein_seq

with open(out_file, "w") as faa:

SeqIO.write(protein_record, faa, "fasta") # Write amino acid sequence to⊔

→ fasta with SeqIO.write
```

Last, we can check the content of the newly-created file:

```
>NC_000913.3:105305-106456 ftsZ [organism=Escherichia coli str. K-12 substr. MG1655] [GeneID=944786] [chromosome=]
MFEPMELTNDAVIKVIGVGGGGGGNAVEHMVRERIEGVEFFAVNTDAQALRKTAVGQTIQI
GSGITKGLGAGANPEVGRNAADEDRDALRAALEGADMVFIAAGMGGGTGTGAAPVVAEVA
KDLGILTVAVVTKPFNFEGKKRMAFAEQGITELSKHVDSLITIPNDKLLKVLGRGISLLD
AFGAANDVLKGAVQGIAELITRPGLMNVDFADVRTVMSEMGYAMMGSGVASGEDRAEEAA
EMAISSPLLEDIDLSGARGVLVNITAGFDLRLDEFETVGNTIRAFASDNATVVIGTSLDP
DMNDELRVTVVATGIGMDKRPEITLVTNKQVQQPVMDRYQQHGMAPLTQEQKPVAKVVND
NAPQTAKEPDYLDIPAFLRKQAD
```

We managed to translate the sequence and save it to a new file! Perhaps you want to try to do the same now.

1.1 So, what do I do now?

Download a file containing the DNA sequence of a gene from NCBI and try to translate the sequence yourself. You can also save the translated version to a new file if you want, and you can play around with other attributes of the SeqRecord class; for example, by modifying the description. You can use the cell below.

[]: # Write your code here