

Bioinformatique

Manipulations de séquences ADN, ARN, protéines,...



: à compléter (différents formats, bases de données ?)

Compter les nucléotides d'une séquence ADN

```
<sxh python; title : Counting_DNA_Nucleotides-01.py> #!/usr/bin/env python # -*- coding: utf-8 -*-  
""" On dispose d'un exemple de chaîne ADN (constituée des symboles 'A', 'C', 'G', 'T') Le programme  
utilise plusieurs techniques pour donner les nombres d'occurrences respectifs des différentes bases  
""" adn =  
"AGCTTTTCATTCTGACTGCAACGGGCAATATGTCTCTGTGTGGATTAAAAAAGAGTGTCTGATAGCAGC"
```

utilisation d'une liste et de la méthode .count() bases=["A","C","G","T"] for base in bases:

```
print adn.count(base),
```

print

Variante : for c in 'ACGT':

```
print adn.count(c),
```

print

variante un peu moins lisible out = [] for c in 'ACGT':

```
out.append(str(adn.count(c)))
```

print(' '.join(out))

utilisation de la technique "list comprehension" count=[adn.count(c) for c in 'ACGT'] for val in count:

```
print val,
```

print

autre "list comprehension", avec impression formatée → version "one line" print "%d %d %d %d" % tuple([adn.count(X) for X in "ACGT"])

count "à la main", sans utilisation de fonctions/librairie ACGT = "ACGT" count = [0,0,0,0] for c in adn:

```
for i in range(len(ACGT)):  
    if c == ACGT[i]:
```

```
count[i] +=1
```

for val in count:

```
print val,
```

print

count "à la main", avec .index() ACGT = "ACGT" count = [0,0,0,0] for c in adn:

```
count[ACGT.index(c)] += 1
```

for val in count:

```
print val,
```

print

utilisation de la librairie collections from collections import defaultdict ncount = defaultdict(int) for c in adn:

```
ncount[c] += 1
```

print ncount['A'], ncount['C'], ncount['G'], ncount['T']

collections.Counter from collections import Counter for k,v in sorted(Counter(adn).items()): print v, print

avec un dictionnaire freq = {'A': 0, 'C': 0, 'G': 0, 'T': 0} for c in adn:

```
freq[c] += 1
```

print freq['A'], freq['C'], freq['G'], freq['T']

avec un dictionnaire et count(), impression différente dico={} for base in bases:

```
dico[base]=adn.count(base)
```

for key,val in dico.items():

```
print "{} = {}".format(key, val)
```

</sxh>

Trouver un motif

+ lecture de fichier

<sxh python; title : Finding_a_Protein_Motif-01.py> #!/usr/bin/env python # -*- coding: utf-8 -*- """ La description complète et les caractéristiques d'une protéine particulière peuvent être obtenues via l'ID "uniprot_id" de la "UniProt database", en insérant la référence dans ce lien :

http://www.uniprot.org/uniprot/uniprot_id

On peut aussi obtenir la séquence peptidique au format FASTA via le lien :

http://www.uniprot.org/uniprot/uniprot_id.fasta """

```
from Bio import SeqIO from Bio import Expasy from Bio import SeqIO
```

```
dic = {"UUU":"F", "UUC":"F", "UUA":"L", "UUG":"L",
```

```
"UCU":"S", "UCC":"S", "UCA":"S", "UCG":"S",
"UAU":"Y", "UAC":"Y", "UAA":"STOP", "UAG":"STOP",
"UGU":"C", "UGC":"C", "UGA":"STOP", "UGG":"W",
"CUU":"L", "CUC":"L", "CUA":"L", "CUG":"L",
"CCU":"P", "CCC":"P", "CCA":"P", "CCG":"P",
"CAU":"H", "CAC":"H", "CAA":"Q", "CAG":"Q",
"CGU":"R", "CGC":"R", "CGA":"R", "CGG":"R",
"AUU":"I", "AUC":"I", "AUA":"I", "AUG":"M",
"ACU":"T", "ACC":"T", "ACA":"T", "ACG":"T",
"AAU":"N", "AAC":"N", "AAA":"K", "AAG":"K",
"AGU":"S", "AGC":"S", "AGA":"R", "AGG":"R",
"GUU":"V", "GUC":"V", "GUA":"V", "GUG":"V",
"GCU":"A", "GCC":"A", "GCA":"A", "GCG":"A",
"GAU":"D", "GAC":"D", "GAA":"E", "GAG":"E",
"GGU":"G", "GGC":"G", "GGA":"G", "GGG":"G", }
```

```
aminoacids = ".join(sorted(list(set([v for k,v in dic.items() if v <> "STOP"])))) print aminoacids
```

```
# UniProt Protein Database access IDs proteins = ['A2Z669', 'B5ZC00', 'P07204_TRBM_HUMAN',
'P20840_SAG1_YEAST']
```

```
handle = Expasy.get_sprot_raw(proteins[0]) seq_record = SeqIO.read(handle, "swiss") handle.close()
print print seq_record
```

```
</sxh>
```

Références

- [Using biological databases to teach evolution and biochemistry](#)
- [Rosalind](#), plateforme d'apprentissage de la programmation en bioinformatique
- [GenBank](#)
- [Biopython](#)
- <https://en.wikipedia.org/wiki/Bioinformatics>
- https://en.wikipedia.org/wiki/Open_Bioinformatics_Foundation
- https://en.wikipedia.org/wiki/FASTA_format
- https://en.wikipedia.org/wiki/List_of_open-source_bioinformatics_software
- <http://www.amberbiology.com/>, "Python For The Life Sciences. A gentle introduction to Python for life scientists" (à paraître)

- références sur la lecture de fichiers :

- http://www.uniprot.org/help/programmatic_access#id_mapping_python_example
- <http://www.python-simple.com/python-biopython/Lecture-ecriture-sequences.php>

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