Introduction to Python for Biologists Lecture 3: Biopython

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- Biopython is a toolkit
- Seq objects and their methods
- SeqRecord objects have data fields
- SeqIO to read and write sequence objects
- Direct access to GenBank with Entrez.efetch
- Working with BLAST results

Modules

- Python functions are divided into 3 sets
 - A small core set that are always available
 - Some built-in modules such as math and os that can be imported from the basic install (ie. >>> import math)
 - An extremely large number of optional modules that must be downloaded and installed before you can import them
 - Code that uses such modules is said to have "dependencies"
- The code for these modules are located in different places on the internet such as SourceForge, GitHub, and developer's own websites (Perl and R are better organized)
- Anyone can write new Python modules, and often several different modules are available that can do the same task

Download a file

 urllib() is a module that lets Python download files from the internet with the .urlretrieve method

>>> import urllib

>>>urllib.urlretrieve('http://biopython.org/SRC/biopyth
on/Tests/GenBank/NC_005816.fna', 'yp.fasta')



- Biopython is an integrated collection of modules for "biological computation" including tools for working with DNA/protein sequences, sequence alignments, population genetics, and molecular structures
- It also provides interfaces to common biological databases (ie. GenBank) and to some common locally installed software (ie. BLAST).
- Loosely based on BioPerl

Biopython Tutorial

• Biopython has a "Tutorial & Cookbook" : <u>http://biopython.org/DIST/docs/tutorial/Tutorial.html</u>

by: Jeff Chang, Brad Chapman, Iddo Friedberg, Thomas Hamelryck, Michiel de Hoon, Peter Cock, Tiago Antao, Eric Talevich, Bartek Wilczyński

from which, most of the following examples are drawn

Object Oriented Code

- Python uses the concept of Object Oriented Code.
- Data structures (known as classes) can contain complex and well defined forms of data, and they can also have built in methods
- For example, many classes of objects have a "print" method
- Complex objects are built from other objects

The Seq object

- The Seq object class is simple and fundamental for a lot of Biopython work. A Seq object can contain DNA, RNA, or protein.
- It contains a string (the sequence) and a defined alphabet for that string.
- The alphabets are actually defined objects such as IUPACAmbiguousDNA Or IUPACProtein
 - Which are defined in the Bio.Alphabet module
 - A Seq object with a DNA alphabet has some different methods than one with an Amino Acid alphabet

```
>>> from Bio.Seq import Seq
>>> from Bio.Alphabet import IUPAC
>>> my_seq = Seq('AGTACACTGGT', IUPAC.unambiguous_dna)
>>> my_seq
Seq('AGTACACTGGT', IUPAC.unambiguous_dna())
>>> print(my_seq)
AGTACACTGGT
```

Seq objects have string methods

- Seq objects have methods that work just like string objects
- You can get the len() of a Seq, slice it, and count() specific letters in it:

```
>>> my_seq = Seq('GATCGATGGGCCTATATAGGATCGAAAATCGC',
IUPAC.unambiguous_dna)
>>> len(my_seq)
32
>>> print(my_seq[6:9])
TGG
>>> my_seq.count("G")
9
```

Turn a Seq object into a string

- Sometimes you will need to work with just the sequence string in a Seq object using a tool that is not aware of the Seq object methods
- Turn a Seq object into a string with str()

```
>> my_seq
Seq('GATCGATGGGCCTATATAGGATCGAAAATCGC',
IUPACUnambiguousDNA())
>>> seq_string=str(my_seq)
>>> seq_string
'GATCGATGGGCCTATATAGGATCGAAAATCGC'
```

Seq Objects have special methods

- DNA Seq objects can .translate() to protein
 - With optional translation table and to_stop=True parameters

```
>>>coding_dna=Seq("ATGGCCATTGTAATGGGCCGCTGAAAGGGTGCCCGATAG",
IUPAC.unambiguous_dna)
```

```
>>> coding_dna.translate()
Seq('MAIVMGR*KGAR*', HasStopCodon(IUPACProtein(), '*'))
>>> print(coding_dna.translate(table=2, to_stop=True))
MAIVMGRWKGAR
```

Seq objects with a DNA alphabet have the reverse_complement() method:

```
>>> my_seq = Seq('TTTAAAATGCGGG', IUPAC.unambiguous_dna)
>>> print(my_seq.reverse_complement())
cccGCATTTTAAA
```

 The Bio.SeqUtils module has some useful methods, such as GC() to calculate % of G+C bases in a DNA sequence.

```
>>> from Bio.SeqUtils import GC
>>> GC(my_seq)
46.875
```

Protein Alphabet

- You could re-define my_seq as a protein by changing the alphabet, which will totally change the methods that will work on it.
 - ('G','A','T','C' are valid protein letters)

>>> from Bio.SeqUtils import molecular_weight
>>> my_seq
Seq('AGTACACTGGT', IUPACUnambiguousDNA())
>>> print(molecular_weight(my_seq))
3436.1957

```
>>> my_seq.alphabet = IUPAC.protein
>>> my_seq
Seq('AGTACACTGGT', IUPACProtein())
>>> print(molecular_weight(my_seq))
912.0004
```

SeqRecord Object

 The SeqRecord object is like a database record (such as GenBank). It is a complex object that contains a Seq object, and also annotation fields, known as "attributes".

.seq

.id

.name

.description

.letter_annotations

.annotations

.features

.dbxrefs

 You can think of attributes as slots with names inside the SeqRecord object. Each one may contain data (usually a string) or be empty.

SeqRecord Example

```
>>> from Bio.Seq import Seq
>>> from Bio.SeqRecord import SeqRecord
>>> test_seq = Seq('GATC')
>>> test_record = SeqRecord(test_seq, id='xyz')
>>> test_record.description= 'This is only a test'
>>> print(test record)
ID: xyz
Name: <unknown name>
Description: This is only a test
Number of features: 0
Seq('GATC', Alphabet())
>>> print(test_record.seq)
GATC
```

Specify fields in the SeqRecord object with a . (dot) syntax

SeqIO and FASTA files

- **SeqIO** is the all purpose file read/write tool for SeqRecords
 - SeqIO can read many file types: http://biopython.org/wiki/SeqIO
- SeqIO has .read() and .write() methods
 - (do not need to "open" file first)
- It can read a text file in FASTA format
- In Biopython, fasta is a type of SeqRecord with specific fields
 - Lets assume you have already downloaded a FASTA file from GenBank, such as: <u>NC_005816.fna</u>, and saved it as a text file in your current directory

```
>>> from Bio import SeqIO
```

```
>>> gene = SeqIO.read("NC_005816.fna", "fasta")
```

>>> gene.id

```
'gi|45478711|ref|NC_005816.1|'
```

```
>>> gene.seq
```

```
Seq('TGTAACGAACGGTGCAATAGTGATCCACACCCCAACGCCTGAAATCAGATCCAGG...CTG', SingleLetterAlphabet())
```

```
>>> len(gene.seq)
```

```
9609
```

Multiple FASTA Records in one file

- The FASTA format can store many sequences in one text file
- SeqIO.parse() reads the records one by one
- This code creates a list of SeqRecord objects:

Database as a FASTA file

• Entire databases of sequences (DNA or protein) can be downloaded as a single FASTA file (e.g. human proteins, *Drosophila* coding CDS, Uniprot UniRef50)

FTP directory /pub/databases/uniprot/uniref/uniref50/ at ftp.uniprot.org

07/22/2015	02:00PM	7,171	README
07/22/2015	02:00PM	4,422	uniref.xsd
07/22/2015	02:00PM	1,755	uniref50.dtd
07/22/2015	02:00PM	3,050,	098,524 <u>uniref50.fasta.gz</u>
07/22/2015	02:00PM	310	uniref50.release note

(not necessarily a good idea to keep 3 GB of data on your computer)

Grab sequence from FASTA file

• If you have a large local FASTA file, and a list of sequences ('my_gene_list.txt') that you want to grab:

SeqIO.write(test, output, 'fasta')

>>> output.close()

SeqIO for FASTQ

- FASTQ is a format for Next Generation DNA sequence data (FASTA + Quality)
- SeqIO can read (and write) FASTQ format files

Direct Access to GenBank

- BioPython has modules that can directly access databases over the Internet
- The Entrez module uses the NCBI Efetch service
- Efetch works on many NCBI databases including protein and PubMed literature citations
- The 'gb' data type contains much more annotation information, but rettype='fasta' also works
- With a few tweaks, this code could be used to download a list of GenBank ID's and save them as FASTA or GenBank files:

```
>>> print(record)
ID: EU490707.1
Name: EU490707
```

Description: Selenipedium aequinoctiale maturase K (matK) gene, partial cds; chloroplast.

Number of features: 3

/sequence version=1



These are sub-fields of the .annotations field

/source=chloroplast Selenipedium aequinoctiale

```
/taxonomy=['Eukaryota', 'Viridiplantae', 'Streptophyta', 'Embryophyta', 'Tracheophyta', 'Spermatophyta', 'Magnoliophyta', 'Liliopsida', 'Asparagales', 'Orchidaceae', 'Cypripedioideae', 'Selenipedium']
/keywords=['']
```

```
/references=[Reference(title='Phylogenetic utility of ycf1 in orchids: a plastid gene
more variable than matK', ...), Reference(title='Direct Submission', ...)]
```

```
/accessions=['EU490707']
```

```
/data_file_division=PLN
```

```
/date=15-JAN-2009
```

/organism=Selenipedium aequinoctiale

/gi=186972394

```
Seq('ATTTTTTACGAACCTGTGGAAATTTTTGGTTATGACAATAAATCTAGTTTAGTA...GAA', IUPACAmbiguousDNA())
```

BLAST

- BioPython has several methods to work with the popular NCBI BLAST software
- NCBIWWW.qblast() sends queries directly to the NCBI BLAST server. The query can be a Seq object, FASTA file, or a GenBank ID.

Parse BLAST Results

- It is often useful to obtain a BLAST result directly (local BLAST server or via Web browser) and then parse the result file with Python.
- Save the BLAST result in XML format
 - NCBIXML.read() for a file with a single BLAST result (single query)
 - NCBIXML.parse() for a file with multiple BLAST results (multiple queries)
 - >>> from Bio.Blast import NCBIXML
 - >>> handle = open("my_blast.xml")
 - >>> blast_record = NCBIXML.read(handle)
 - >>> for hit in blast_record.descriptions:
 - print hit.title
 - print hit.e

BLAST Record Object



View Aligned Sequence

>>> from Bio.Blast import NCBIXML
>>> handle = open("my_blast.xml")
>>> blast_record = NCBIXML.read(handle)
>>> for hit in blast_record.alignments:
 for hsp in hit.hsps:
 print hit.title
 print hit.title
 print hsp.expect
 print (hsp.query[0:75] + '...')
 print(hsp.match[0:75] + '...')
 print(hsp.sbjct[0:75] + '...')

Many Matches

- Often a BLAST search will return many matches for a single query (save as an XML format file)
- NCBIXML.parse() can return these as BLAST record objects in a list, or deal with them directly in a for loop.

Illumina Sequences

Illumina sequence files are usually stored in the FASTQ format.
 Similar to FASTA, but with an additional pair of lines for the quality annotation of each base.

@SRR350953.5 MENDEL_0047_FC62MN8AAXX:1:1:1646:938 length=152 NTCTTTTTCTTTCCTCTTTTGCCAACTTCAGCTAAATAGGAGCTACACTGATTAGGCAGAAACTTGATTAACAGGGCTTAAGGTA ACCTTGTTGTAGGCCGTTTTGTAGCACTCAAAGCAATTGGTACCTCAACTGCAAAAGTCCTTGGCCC +SRR350953.5 MENDEL 0047 FC62MN8AAXX:1:1:1646:938 length=152 +50000222C@@@@@22::::8888898989::::::<<<:<<<::<<::::<<<:<<!!!!!!GFEEGGGGGGGGII@IGDGBG GGGGGDDIIGIIEGIGG>GGGGGGGGGGGGGGGIIHIIBIIIGIIIHIIIGII @SRR350953.6 MENDEL 0047 FC62MN8AAXX:1:1:1686:935 length=152 NATTTTTACTAGTTTATTCTAGAACAGAGCATAAACTACTATTCAATAAACGTATGAAGCACTACTCACCTCCATTAACATGACGTT TTTCCCTAATCTGATGGGTCATTATGACCAGAGTATTGCCGCGGTGGAAATGGAGGTGAGTAGTG +SRR350953.6 MENDEL 0047 FC62MN8AAXX:1:1:1686:935 length=152 @@@@@@CIGIHIIDGIGIIIIHHIIHGHHIIHHIFIIIIIHIIIIIBIIIEIFGIIIFGFIBGDGGGGGGFIGDIFGADGAE @SRR350953.7 MENDEL_0047_FC62MN8AAXX:1:1:1724:932 length=152 NTGTGATAGGCTTTGTCCATTCTGGAAACTCAATATTACTTGCGAGTCCTCAAAGGTAATTTTTGCTATTGCCAATATTCCTCAGA +SRR350953.7 MENDEL 0047 FC62MN8AAXX:1:1:1724:932 length=152 GDDDIHIHIIII8GGGGGIIHHIIIGIIGIBIGIIIEIHGGFIHHIIIIIIGIIFIG

Get a file by FTP in Python

```
>>> from ftplib import FTP
>>> host="ftp.sra.ebi.ac.uk"
>>> ftp=FTP(host)
>>> ftp.login()
'230 Login successful.'
ftp.cwd('vol1/fastq/SRR020/SRR020192')
'250 Directory successfully changed.'
>>> ftp.retrlines('LIST')
               ftp 1777817 Jun 24 20:12 SRR020192.fastq.gz
-r--r-- 1 ftp
'226 Directory send OK.'
>>> ftp.retrbinary('RETR SRR020192.fastq.gz', \
open('SRR020192.fastq.gz', 'wb').write)
'226 Transfer complete.'
>>> ftp.quit()
'221 Goodbye.'
```

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