

Genome 559

Intro to Statistical and Computational Genomics

Lecture 17b-18b:
Biopython
Larry Ruzzo

Biopython

What is Biopython?

How do I get it to run on my computer?

What can it do?

Biopython

Biopython is tool kit, not a program—a set of Python modules useful in bioinformatics

Features include:

- Sequence class (can transcribe, translate, invert, etc)
- Parsing files in different database formats
- Interfaces to progs/DBs like Blast, Entrez, PubMed
- Code for handling alignments of sequences
- Clustering algorithms, etc, etc.

Useful tutorials at <http://biopython.org>

Making Biopython available on your computer

Runs on Windows, MaxOSX, and Linux

<http://biopython.org/>

Look for download/install instructions

May require “Admin” privileges

1.53 is latest; works with Python 2.5 -2.6 (not 3?)

Use in the lab

(you shouldn't need this if you install biopython at home)

Try

```
import Bio
```

If it fails, try:

```
import sys
```

```
for p in sys.path: # if curious, here's where
```

```
    print p           # Python looks for imports
```

```
sys.path.append('/Library/Python/  
2.5/site-packages')
```

```
import Bio
```

Sequence class

```
>>> from Bio.Seq import Seq # sequence class
>>> myseq = Seq("AGTACACTGGT")
>>> myseq.alphabet
Alphabet()
>>> myseq.tostring()
'AGTACACTGGT'
```

More functionality than a plain string

```
>>> myseq
Seq('AGTACACTGGT', Alphabet())
>>> myseq.complement()
Seq('TCATGTGACCA', Alphabet())
>>> myseq.reverse_complement()
Seq('ACCAGTGTACT', Alphabet())
```

A sequence in a specified alphabet

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

Transcribe/Translate

```
>>> from Bio import Transcribe
>>> mydna = Seq("GATCGATGGGCCTATATAGGATCGAAAATCGC",
... IUPAC.unambiguous_dna)
>>> myrna = mydna.transcribe()
>>> print myrna
Seq('GAUCGAUGGCCUAUAUAGGAUCGAAAAUCGC', IUPACUnambiguousRNA())
>>> myprot = myrna.translate()
Seq('DRWAYIGSKI', ExtendedIUPACProtein())
>>> s2 = Seq("AUGGCCAUUGUAAUGGGCCGCUGAAAGGGUGCCCGAUAG",
... IUPAC.unambiguous_rna)
>>>s2.translate()
Seq('MAIVMGR*KGAR*', HasStopCodon(IUPACProtein(), '*'))
# also possible to reverse transcribe, etc etc
```

Parsing a database format

FASTA database file named "ls_orchid.fasta":

```
>gi|2765658|emb|Z78533.1|CIZ78533 C.irapeanum 5.8S rRNA gene
CGTAACAAGGTTCCGTAGGTGAACCTGCGGAAGGATCATTGATGAGACCCTGGAATAAACGATCGAGTG
AATCCGGAGGACCGGTGTACTCAGCTCACCGGGGGATTGCTCCCGTGGTGACCCCTGATTGTTGTTGGG
.....
from Bio import SeqIO
handle = open("ls_orchid.fasta")
for seqrec in SeqIO.parse(handle, "fasta") :
    print seqrec.id
    print seqrec.seq
    print len(seqrec.seq)
handle.close()

gi|2765658|emb|Z78533.1|CIZ78533
Seq('CGTAACAAGGTTCCGTAGGTGAACCTGCGGAAGGATCATTGATGAGACCCTGGAATAAACGATCGAGTG ...',
     SingleLetterAlphabet())
740
...
```

Exercise I

Modify the example above to print the GC% of each sequence, too.

Solution I

```
from Bio import SeqIO
handle = open("ls_orchid.fasta")
for seqrec in SeqIO.parse(handle, "fasta"):
    print seqrec.id
    s = seqrec.seq
    print s
    print len(s),
    na = s.count('A')
    nc = s.count('C')
    ng = s.count('G')
    nt = s.count('T')
    print "GC%=", (ng+nc)*100.0/(na+nc+ng+nt)
handle.close()
```

Q1: there's also a Biopython func to calc gc%; can you find it?

Q2: Why did I not use $(G+C)/\text{len}(s)$?

Parses GenBank Format, too

```
from Bio import SeqIO
handle = open("ls_orchid.gbk")
for seqrec in SeqIO.parse(handle, "genbank") :
    print seqrec.id
    print repr(seqrec.seq)
    print len(seqrec)
handle.close()
```

This should give:

```
Z78533.1
Seq
('CGTAACAAGGTTCCGTAGGTGAACCTGCGGAAGGATCATTGATGAGACC GTGG .
..CGC', IUPACAmbiguousDNA())
740
...
```

Exercise 2

Change above example to save the records in a list called seqrecs

Solution 2

```
from Bio import SeqIO
handle = open("ls_orchid.gbk")
seqrecs = []
for seqrec in SeqIO.parse(handle, "genbank") :
    seqrecs.append(seqrec)
    print seqrec.id
    print repr(seqrec.seq)
    print len(seqrec)
handle.close()
```

This should give:

```
Z78533.1
Seq
('CGTAACAAGGTTCCGTAGGTGAACCTGCAGAAGGATCATTGATGAGACCGTGG.
..CGC', IUPACAmbiguousDNA())
740
...
```

Feature Tables

```
>>>seqrecs[0]
SeqRecord(seq=Seq('CGTA...CGC', IUPACAmbiguousDNA()),
id='Z78533.1', name='Z78533', description='C.irapeanum
5.8S rRNA gene and ITS1 and ITS2 DNA.', dbxrefs=[])

>>> print seqrecs[0]
ID: Z78533.1
Name: Z78533
Description: C.irapeanum 5.8S rRNA gene and ITS1 and ITS2 DNA.

Number of features: 5
/sequence_version=1
/source=Cypripedium irapeanum
/taxonomy=['Eukaryota', ..., 'Cypripedium']
/keywords=['5.8S ribosomal RNA', ... 'ITS2']
/references=[<Bio.SeqFeature.Reference instance...>]
/accessions=['Z78533']
/data_file_division=PLN
/date=30-NOV-2006
/organism=Cypripedium irapeanum
/gi=2765658

Seq('CGTAACAAGGTTCCGTAGGTGA...CGC', IUPACAmbiguousDNA())
```

Extracting Features

(Lists of objects with dicts of lists of lists of dicts of ...Oh my!)

```
>>> seqrecs[0].annotations  
{'sequence_version': 1, 'source': 'Cypripedium  
irapeanum', 'taxonomy': ['Eukaryota', ... ... ... ...]}
```

it's a dictionary! What keys does it have?

```
>>> seqrecs[0].annotations.keys()  
['sequence_version', 'source', 'taxonomy', 'keywords',  
 'references', 'accessions', 'data_file_division', 'date',  
 'organism', 'gi']  
# grab one dict entry
```

```
>>> seqrecs[0].annotations['keywords']  
['5.8S ribosomal RNA', '5.8S rRNA gene', 'internal  
transcribed spacer', 'ITS1', 'ITS2']
```

#It's a list! We can index into it...

```
>>> seqrecs[0].annotations['keywords'][1]  
'5.8S rRNA gene'
```

Searching GenBank

```
# This example & next require internet access

from Bio import GenBank
gilist = GenBank.search_for("Opuntia AND rpl16")
# (that's RPL-sixteen, not RP-one-one-six)

# gilist will be a list of all of the GenBank
# identifiers that match our query:
print gilist
['6273291', '6273290', '6273289', '6273287',
 '6273286', '6273285', '6273284']
```

Searching GenBank

```
ncbidict = GenBank.NCBIDictionary("nucleotide", "genbank")
gbrecord = ncbidict[gilist[0]]
print gbrecord
```

```
LOCUS      AF191665 902 bp DNA PLN 07-NOV-1999
DEFINITION Opuntia marenae rpl16 gene; chloroplast gene for
           chloroplast product, partial intron sequence.
ACCESSION  AF191665
VERSION    AF191665.1 GI:6273291
...
...
```

Exercise 3: What kind of a thing is “gbrecord”? Is there other stuff hidden with it like annotations or feature tables? How do I access it?

Solution 3

```
>>> type(gbrecord)
<type 'str'>

# Aha, it's just a plain string.

>>> gbrecord
'LOCUS          AY851612                      892 bp      DNA
linear    PLN 10-APR-2007\nDEFINITION  Opuntia subulata
rpl16 gene, intron; chloroplast.\nACCESSION
AY851612\nVERSION      AY851612.1  GI:
57240072\nKEYWORDS      .\nSOURCE      chloroplast
Austrocylindropuntia subulata\n  ...  ...  ...  ...  ...  ...  '
```

Can we get Biopython to parse it?

To parse a string

```
>>>SeqIO.parse(gbrecord, "genbank")
Traceback (most recent call last): blah blah blah...
```

```
# Oops, a string isn't a handle...
```

Turn a string
into a handle

```
>>> import cStringIO
>>> SeqIO.parse(cStringIO.StringIO(gbrecord), "genbank")
<generator object at 0x5254b8> ## Oops, need loop
>>> for rec in SeqIO.parse(cStringIO.StringIO(gbrecord),
... "genbank"):
...     print rec
...
ID: AY851612.1
Name: AY851612
Description: Opuntia subulata rpl16 gene, intron;
chloroplast.
Number of features: 3
/sequence_version=1
/source=chloroplast Austrocylindropuntia subulata ...
```

(Some) Other Capabilities

AlignO

consensus

PSSM (weight matrix)

BLAST

both local and internet

Entrez EUtils

including GenBank and PubMed

Other Databases

SwissProt, Prosite, ExPASy, PDB

How would I use Biopython?

Biopython is not a program itself; it's a collection of tools for Python bioinformatics programming

When doing bioinformatics, keep Biopython in mind

Browse the documentation; become familiar with its capabilities

Use help(), type(), dir() & other built-in features to explore

You might prefer it to writing your own code for:

- Defining and handling sequences and alignments
- Parsing database formats
- Interfacing with databases

You don't have to use it all! Pick out one or two elements to learn first

Exercises

Many!

As one suggestion, look at the “Cookbook” section of the tutorial. Figure out how to read my hem6.txt Phylip alignment & make a WMM from it.

Feel free to do something with one of the other pieces instead.