

# Biopython

MBV-INFx410

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## Object oriented programming

- Biopython is object-oriented
- Some knowledge helps understand how biopython works
- OOP is a way of organizing data and methods that work on them in a coherent package
- OOP helps structure and organize the code

## Classes and objects

- A class:
  - is a user defined type
  - is a mold for creating objects
  - specifies how an object can contain and process data
  - represents an abstraction or a template for how an object of that class will behave
- An object is an instance of a class
- All objects have a type – shows which class they were made from

## Attributes and methods

- Classes specify two things:
  - Attributes – data holders
  - Methods – functions for this class
- Attributes are variables that will contain the data that each object will have
- Methods are functions that an object of that class will be able to perform

## Fake class and object example

- Class: MyCup
- MyCup has:
  - attribute contents
  - method heat
- An object of the class MyCup is created like this:
  - mycup = MyCup("Water")
    - Here: the attribute contents is assigned the value "Water"
- Find out what the content is (access attribute):
  - mycup.contents – will report the contents
- Heat contents (use method):
  - mycup.heat() – will heat contents, in this case "Water"

## Summary

- An object has to be instantiated, i.e. created, to exist
- Every object has a certain type, i.e. is of a certain class
- The class decides which attributes and methods an object of that class has
- Attributes and methods are accessed using . after the object variable name

## Biopython

- Package that assists with processing biological data
- Consists of several modules – some with common operations, some more specialized
- Website: [biopython.org](http://biopython.org)

## Working with sequences

- Biopython has many ways of working with sequence data
- Focus on:
  - Alphabet package
  - Seq class
  - SeqRecord class
  - SeqIO package
- Other useful classes for working with alignments, blast searches and results etc are also available, not covered today

## Class Alphabet

- Every sequence needs an alphabet
  - CCTTGGCC – DNA or protein?
- Biopython contains several alphabets
  - DNA
  - RNA
  - Protein
  - the three above with IUPAC codes
  - ...and others
- Can all be found in Bio.Alphabet package

## Alphabet example

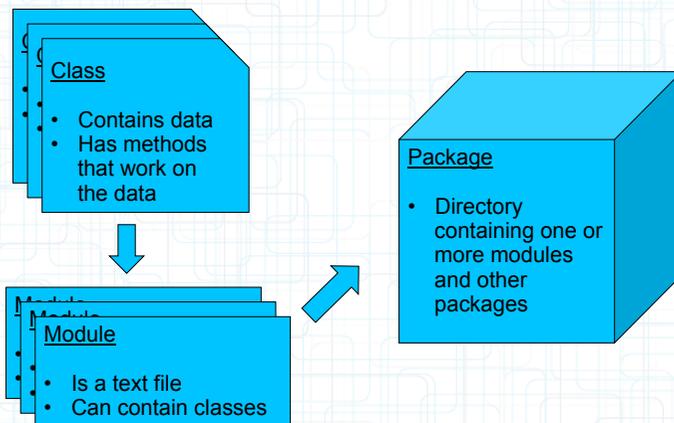
```
>>> import Bio.Alphabet
>>> Bio.Alphabet.ThreeLetterProtein.letters
['Ala', 'Asx', 'Cys', 'Asp', 'Glu', 'Phe', 'Gly', 'His', 'Ile',
'Lys', 'Leu', 'Met', 'Asn', 'Pro', 'Gln', 'Arg', 'Ser', 'Thr',
'Sec', 'Val', 'Trp', 'Xaa', 'Tyr', 'Glx']
>>> from Bio.Alphabet import IUPAC
>>> IUPAC.IUPACProtein.letters
'ACDEFGHIKLMNPQRSTVWY'
>>> IUPAC.unambiguous_dna.letters
'GATC'
>>>
```

NOTE: have to import Alphabets to use them

Can now print all of the common three letter abbreviations

Can work with both ambiguous and unambiguous sequences

## Packages, modules and classes



## Packages, modules and classes

- What happens here?

```
>>> from Bio.Alphabet import IUPAC
>>> IUPAC.IUPACProtein.letters
```
- Bio and Alphabet are packages
  - packages contain other packages and modules
- IUPAC is a module
  - a module is a file with python code
  - a module can contain 0 to many classes
- IUPAC module contains class IUPACProtein and other classes specifying alphabets
- IUPACProtein class specifies objects that have the attribute letters

## Class Seq

- Represents one sequence with its alphabet

```
newseq = Seq(string, alphabet)
```

- Has attributes that keeps the string and the alphabet

- Methods:

- newseq.translate()
- newseq.transcribe()
- newseq.complement()
- newseq.reverse\_complement()
- ...

## Using Seq

```
>>> from Bio.Seq import Seq
>>> import Bio.Alphabet
>>> seq = Seq("CCGGGTT", Bio.Alphabet.IUPAC.unambiguous_dna)
>>> seq
Seq('CCGGGTT', IUPACUnambiguousDNA())
>>> seq.transcribe()
Seq('CCGGGUU', IUPACUnambiguousRNA())
>>> seq.translate()
Seq('PG', IUPACProtein())
>>> seq = Seq("CCGGGUU", Bio.Alphabet.IUPAC.unambiguous_rna)
>>> seq.transcribe()
Traceback (most recent call last):
  File "<stdin>", line 1, in <module>
  File "/site/VERSIONS/python-2.6.2/lib/python2.6/site-packages/Bio/Seq.py",
line 830, in transcribe
    raise ValueError("RNA cannot be transcribed!")
ValueError: RNA cannot be transcribed!
>>> seq.translate()
Seq('PG', IUPACProtein())
>>>
```

Import classes  
Create object

Use methods

New object, different alphabet

Alphabet dictates which  
methods make sense

## Seq as a string

- Most string methods work on Seqs
- If string is needed, do str(seq)

```
>>> seq = Seq('CCGGGTTAACGTA', Bio.Alphabet.IUPAC.unambiguous_dna)
>>> seq[5]
Seq('CCGGG', IUPACUnambiguousDNA())
>>> len(seq)
13
>>> seq.lower()
Seq('ccgggttaacgta', DNAAlphabet())
>>> print seq
CCGGGTTAACGTA
>>> mystring = str(seq)
>>> print mystring
CCGGGTTAACGTA
>>> type(seq)
<class 'Bio.Seq.Seq'>
>>> type(mystring)
<type 'str'>
>>>
```

Slicing

Length of Seq

Lower case

Printing

Convert as needed

Can check what class  
or type an object is from

## SeqRecord

- Seq contains the sequence and alphabet
- But sequences often come with a lot more
- SeqRecord = Seq + metadata
- Main attributes:
  - id – name or identifier
  - seq – Seq object containing the sequence

# SeqRecord example

```
>>> seq
Seq('CCGGGTTAACGTA', IUPACUnambiguousDNA())
>>> from Bio.SeqRecord import SeqRecord
>>> seqRecord = SeqRecord(seq, id='001')
>>> seqRecord
SeqRecord(seq=Seq('CCGGGTTAACGTA', IUPACUnambiguousDNA()),
id='001', name=<unknown name>, description=<unknown description>,
dbxrefs=[])
```

Existing seq object

SeqRecord is a class found inside the Bio.SeqRecord module

Using existing seq object to create a SeqRecord with an identifier

Several other attributes that for now don't have any value

# SeqRecord attributes

- From the biopython webpages:

Main attributes:

- id** - Identifier such as a locus tag (string)
- seq** - The sequence itself (Seq object or similar)

Additional attributes:

- name** - Sequence name, e.g. gene name (string)
- description** - Additional text (string)
- dbxrefs** - List of database cross references (list of strings)
- features** - Any (sub)features defined (list of SeqFeature objects)
- annotations** - Further information about the whole sequence (dictionary)  
Most entries are strings, or lists of strings.
- letter\_annotations** - Per letter/symbol annotation (restricted dictionary). This holds Python sequences (lists, strings or tuples) whose length matches that of the sequence. A typical use would be to hold a list of integers representing sequencing quality scores, or a string representing the secondary structure.

# SeqRecords in practice...

```
>>> from Bio.SeqRecord import SeqRecord
>>> from Bio.Seq import Seq
>>> from Bio.Alphabet import DNAAlphabet

>>> seqRecord = SeqRecord(Seq('GCAGCCTCAAACCCAGCTG',
... DNAAlphabet), id = 'NM_005368.2', name = 'NM_005368',
... description = 'Myoglobin var 1',
... dbxrefs = ['GeneID:4151', 'HGNC:6915'])

>>> seqRecord
SeqRecord(seq=Seq('GCAGCCTCAAACCCAGCTG',
<class 'Bio.Alphabet.DNAAlphabet'>), id='NM_005368.2',
name='NM_005368', description='Myoglobin var 1',
dbxrefs=['GeneID:4151', 'HGNC:6915'])
```

Import necessary classes

Create object, assign values to attributes

Print object

# The SeqIO package

- How to get sequences in and out of files
- Retrieves sequences as SeqRecords, can write SeqRecords to files
- Reading:
  - SeqIO.parse(filehandle, format)
  - returns a generator that gives SeqRecords
- Writing:
  - SeqIO.write(SeqRecord(s), filehandle, format)
- Note: both input and output files can contain several sequences

## SeqIO formats

- List: <http://biopython.org/wiki/SeqIO>
- Some examples:
  - fasta
  - genbank
  - several fastq-formats
  - ace
- Note: a format might be readable but not writable depending on biopython version

## Reading a file

- SeqIO.parse returns a SeqRecord iterator
- An iterator will give you the next element the next time it is called
- Useful because if a file contains many records, we avoid putting all into memory all at once

```
from Bio import SeqIO
fh = open("example.fasta", "r")
for record in SeqIO.parse(fh,"fasta") :
    print record.id
fh.close()
```

## Parsing fasta files

- In python interactive shell:

```
>>> from Bio import SeqIO
>>> fh = open("mb.fsa", "r")
>>> for record in SeqIO.parse(fh, "fasta"):
...     print record.id
...     print record.seq[:10]
...
NM_005368.2
GCAGCCTCAA
XM_001081975.2
CCTCTCCCA
NM_001164047.1
TAGCTGCCCA
>>>
```

Import modules, open the file

Per element in file:

Print the identifier id  
Print the first ten sequence letters

## convert.py

- Goal: convert from genbank to fasta
- Create script file:
  - Import SeqIO
  - Open the file
  - For each entry (record) in file
    - print record
  - Close file

## convert.py – read sequences

```
import sys
from Bio import SeqIO
```

Need to import SeqIO, otherwise methods not available!

```
# Open the input file
fh = open(sys.argv[1], "r")
```

Opening file, as before

```
for record in SeqIO.parse(fh, "genbank"):
```

Per genbank record in file

```
    # Print the entire record
    print record
fh.close()
```

Each genbank entry is printed

Closing file, as before

Try it out with mb.gbk!

## convert.py results 1

ID: NM\_005368.2

Name: NM\_005368

Description: Homo sapiens myoglobin (MB), transcript variant 1, mRNA.

Number of features: 11

/comment=REVIEWED REFSEQ: This record has been curated by NCBI staff. The reference sequence was derived from BU585249.1, BQ956082.1 and BC014547.1.

On Mar 4, 2004 this sequence version replaced gi:4885476.

Summary: This gene encodes a member of the globin superfamily and is expressed in skeletal and cardiac muscles. The encoded protein is a haemoprotein contributing to intracellular oxygen storage and transcellular facilitated diffusion of oxygen. At least three alternatively spliced transcript variants encoding the same protein have been reported. [provided by RefSeq, Jul 2008].

.....

## convert.py take 2

- Modification: get only sequence and id information
- Fasta description line consists of id and description
- Print
  - The id
  - The description
  - The sequence

## convert.py take 2

```
import sys
from Bio import SeqIO
```

```
# Open the input file
fh = open(sys.argv[1], "r")
```

```
for record in SeqIO.parse(fh, "genbank"):
```

```
    # Print only id, description and sequence
```

```
    print record.id
    print record.description
    print record.seq
```

```
fh.close()
```

Select only the attributes that you actually want! Available since the parser has read it in the file

## convert.py results 2

```
NM_005368.2
Homo sapiens myoglobin (MB), transcript variant 1, mRNA.
GCAGCCTCAAACCCCAGCTGTTGGGGCCAGGACACCC
AGTGAGCCCATACTTGCTCTTTTTGTCTTCTTCAGACTG
CGCCATGGGGCTCAGCGACGGGGAATGGCAGTTGGTG
CTGAACGTCTGGGGGAAGGTGGAGGCTGACATCCCAG
GCCATGGGCAGGAAGTCCATCAGGCTCTTTAAGGGT
CACCCAGAGACTCTGGAGAAGTTTGACAAGTTCAAGCA
.....
XM_001081975.2
PREDICTED: Macaca mulatta myoglobin, transcript variant 1 (MB), mRNA.
CCTCTCCCCACCCCAGCCCTGGCCGCTTGGCTGGAAG
CTCTGCGAGGACAGCTGGGGAGAAGGGGAGCTGTGAC
TGCGCCATGGGGCTCAGCGACGGGGAATGGCAGTTGG
.....
```

## Other available info

- Description:
  - seqRecord.description
- Annotations:
  - seqRecord.annotations.keys()
- Taxonomy:
  - seqRecord.annotations['taxonomy']

## Writing files

- Note: sequences is here a list containing several SeqRecords
- Can write any iterable containing SeqRecords to a file
- By specifying format, we specify what information to print out – no need to specify what we want to write out

```
from Bio import SeqIO
sequences = ... # add code here
output_handle = open("example.fasta", "w")
SeqIO.write(sequences, output_handle, "fasta")
output_handle.close()
```

## convert.py take 3

- Modification: write output to file
  - Open outfile
  - Per record,
    - Write it to file in fasta format
  - Close input file
  - Close output file

## convert.py take 3

```
import sys
from Bio import SeqIO

# Open the input file
fh = open(sys.argv[1], "r")
# Open the output file
fo = open("mb.fsa", "w")

for record in SeqIO.parse(fh, "genbank"):
    # Use SeqIO to write properly
    # formatted record
    SeqIO.write(record, fo, "fasta")

# ...and closing files
fh.close()
fo.close()
```

Open both input and output file

Per entry in the gbk file

Write out record in fasta format

Close both input and output file

## name\_w\_organism.py

- Starting point: genbank file
- Goal: print fasta file with description lines that begin with organism name
- Process:
  - Read in fasta sequences as SeqRecords
  - Open output handle
  - Per fasta sequence:
    - figure out the organism name
    - change description line
    - print to output file
  - Close files

## name\_w\_organism.py

```
import sys
from Bio import SeqIO

# Open the input file
fh = open(sys.argv[1], "r")
# Open the output file
fo = open(sys.argv[2], "w")

for record in SeqIO.parse(fh, "genbank"):
    # using SeqRecord annotation dictionary
    # to get the correct name
    organism = record.annotations['organism']
    species_name = organism.split()[1]
    # adding it onto the record id
    record.id = species_name + "_" + record.id
    SeqIO.write(record, fo, "fasta")
fh.close()
fo.close()
```

## Tips and hints

- Always comment your code – easier to understand later
- Never write lots of code without testing while writing – makes for less code to debug
- Always test on input where you know what the results should be
- If it went too easy, too well or too fast: it is probably wrong!

## Learning more

- Recommended book:  
Sebastian Bassi:  
Python for Bioinformatics
- [www.python.org](http://www.python.org)
  - has lots of documentations  
and beginner tutorials
- Google

