

Typical workflow when working with sequence data (e.g., genomes)

- 1. Download sequence data from NCBI
- 2. Save on local harddrive
- 3. Do analysis on locally stored sequence data
- 4. Repeat from step 3
 - to fix bugs
 - to run a different analysis

We can do the initial download in Biopython

```
# retrieve record from Entrez
handle = Entrez.efetch(db="nucleotide", id="KT220438", rettype="gb", \
                                                               retmode="text")
# read from the handle using regular Python, not SeqIO.read()!
gb file contents = handle.read()
handle.close()
# the entire genbank file is now in the variable `gb file contents`
print(gb file contents)
                                    1701 bp cRNA linear VRL 20-JUL-2015
LOCUS
           KT220438
           Influenza A virus (A/NewJersey/NHRC 93219/2015(H3N2)) segment 4
DEFINITION
           hemagglutinin (HA) gene, complete cds.
           KT220438
ACCESSION
           KT220438.1 GI:887493048
VERSION
KEYWORDS
SOURCE
           Influenza A virus (A/New Jersey/NHRC 93219/2015(H3N2))
           Influenza A virus (A/New Jersey/NHRC 93219/2015(H3N2))
 ORGANISM
           Viruses; ssRNA viruses; ssRNA negative-strand viruses;
            Orthomyxoviridae; Influenzavirus A.
           1 (hages 1 + 0.1701)
```

We can do the initial download in Biopython

```
# retrieve record from Entrez
handle = Entrez.efetch(db="nucleotide", id="KT220438", rettype="qb", \
                                                                retmode="text")
# read from the handle using regular Python, not SegIO.read()!
gb file contents = handle.read()
handle.close()
# write out to disk
with open("KT220438.gb", "w") as out handle:
    out handle.write(gb file contents) # writes everything at once
# now we can read back in and process with Biopython
with open("KT220438.gb", "r") as in handle:
    record = SeqIO.read(in handle, format="gb") # use SeqIO to parse
```

Features in genbank files

```
Location/Oualifiers
FEATURES
                     1..1701
     source
                     /organism="Influenza A virus (A/New
                     Jersey/NHRC 93219/2015(H3N2))"
                     /mol type="viral cRNA"
                     /strain="A/NewJersey/NHRC 93219/2015"
                     /serotype="H3N2"
                     /isolation source="nasopharyngeal swab"
                     /host="Homo sapiens"
                     /db xref="taxon:1682360"
                     /segment="4"
                     /lab host="MDCK"
                     /country="USA: New Jersey"
                     /collection date="17-Jan-2015"
                     1..1701
     gene
                     /gene="HA"
     CDS
                     1..1701
                     /gene="HA"
                     /function="receptor binding and fusion protein"
                     /codon start=1
                     /product="hemagglutinin"
                     /protein id="AKQ43545.1"
                     /db xref="GI:887493049"
                     /translation="MKTIIALSYILCLVFAOKIPGNDNSTATLCLGHHAVPNGTIVKT
                     ITNDRIEVTNATELVONSSIGEICDSPHOILDGENCTLIDALLGDPOCDGFONKKWDL
                     FVERSKAYSNCYPYDVPDYASLRSLVASSGTLEFNNESFNWTGVTQNGTSSACIRRSS
                     SSFFSRLNWLTHLNYTYPALNVTMPNNEQFDKLYIWGVHHPGTDKDQIFLYAQSSGRI
                     TVSTKRSOOAVIPNIGSRPRIRDIPSRISIYWTIVKPGDILLINSTGNLIAPRGYFKI
                     RSGKSSIMRSDAPIGKCKSECITPNGSIPNDKPFONVNRITYGACPRYVKHSTLKLAT
                     GMRNVPEKOTRGIFGAIAGFIENGWEGMVDGWYGFRHONSEGRGQAADLKSTQAAIDQ
```

TNCKT.NRT.TCKTNEKEHOTEKEESEVECRTODLEKVVEDTKTDT.WSVNAET.T.VAT.ENO

The source feature

```
1..1701
/organism="Influenza A virus (A/New
Jersey/NHRC_93219/2015(H3N2))"
/mol_type="viral cRNA"
/strain="A/NewJersey/NHRC_93219/2015"
/serotype="H3N2"
(...)
```

- identifies the biological source of the specified span of the sequence; mandatory
- more than one source key per sequence is allowed
- every entry/record will have, as a minimum, either a single source key spanning the entire sequence or multiple source keys, which together, span the entire sequence

The gene feature

gene 1..1701 /gene="HA"

 region of biological interest identified as a gene and for which a name has been assigned

The CDS feature

```
CDS 1..1701
/gene="HA"
/function="receptor binding and fusion protein"
/codon_start=1
/product="hemagglutinin"
(...)
```

- coding sequence
- sequence of nucleotides that corresponds with the sequence of amino acids in a protein (location includes stop codon)

The mat_peptide feature

```
mat_peptide 49..1035
/gene="HA"
/product="HA1"
```

- mature peptide or protein coding sequence
- coding sequence for the mature or final peptide or protein product following post-translational modification
- the location does not include the stop codon (unlike the corresponding CDS)

Influenza HA has one CDS but two mature peptides

```
1..1701
CDS
                 /gene="HA"
                 /function="receptor binding and fusion protein"
                 /codon start=1
                 /product="hemagglutinin"
                 /protein id="AKQ43545.1"
                 (\ldots)
                 49..1035
mat peptide
                 /gene="HA"
                 /product="HA1"
                 1036..1698
mat peptide
                 /gene="HA"
                 /product="HA2"
```

Many more feature types exist

- centromere
- exon
- gap
- LTR
- mRNA
- tRNA

Full list with explanations at:

http://www.insdc.org/files/feature_table.html#7.2

Working with features in Biopython

Features are stored as a list in Biopython genbank records

```
In [1]: print(record.features)
Out[1]: [SeqFeature(FeatureLocation(ExactPosition(0),
ExactPosition(1701), strand=1), type='source'),
SeqFeature(FeatureLocation(ExactPosition(0),
ExactPosition(1701), strand=1), type='gene'),
SeqFeature(FeatureLocation(ExactPosition(0),
ExactPosition(1701), strand=1), type='CDS'),
SeqFeature(FeatureLocation(ExactPosition(48),
ExactPosition(1035), strand=1), type='mat peptide'),
SeqFeature(FeatureLocation(ExactPosition(1035),
ExactPosition(1698), strand=1), type='mat peptide')]
```

Features are stored as a list in Biopython genbank records

```
In [1]: for feature in record.features:
            print(feature)
Out[1]: type: source
        location: [0:1701](+)
        qualifiers:
            Key: collection date, Value: ['17-Jan-2015']
            Key: country, Value: ['USA: New Jersey']
            Key: db xref, Value: ['taxon:1682360']
            (\ldots)
        type: gene
        location: [0:1701](+)
        qualifiers:
            Key: gene, Value: ['HA']
        type: CDS
        location: [0:1701](+)
        qualifiers:
            Key: codon start, Value: ['1']
            Key: db vref Value: ['CT:887493049']
```

Features have member variables type, location, and qualifiers

```
In [1]: print("Type:", feature.type)
        print("\nLocation:", feature.location)
        print("\nQualifiers:", feature.qualifiers)
Out[1]: Type: source
        Location: [0:1701](+)
        Qualifiers: {'organism': ['Influenza A virus (A/
New Jersey/NHRC 93219/2015(H3N2))'], 'lab host':
['MDCK'], 'strain': ['A/NewJersey/NHRC 93219/2015'],
'db xref': ['taxon:1682360'], 'host': ['Homo sapiens'],
'segment': ['4'], 'isolation source': ['nasopharyngeal
swab'], 'collection date': ['17-Jan-2015'], 'mol type':
['viral cRNA'], 'serotype': ['H3N2'], 'country': ['USA:
Now Torcow 11
```

The qualifiers are stored as a dictionary

```
In [1]: print(feature.qualifiers["organism"])
Out[1]: ['Influenza A virus (A/New Jersey/
NHRC 93219/2015(H3N2))']
Note: The qualifiers dictionary returns a list! We need to extract the first
element of the list if we want to just get the contents string.
In [2]: organism = feature.qualifiers["organism"][0]
         print(organism)
Out[2]: Influenza A virus (A/New Jersey/
NHRC 93219/2015(H3N2))
```