## Requesting data from Entrez in different formats

### We can request data as text or as XML

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```
handle = Entrez.efetch(db="nucleotide", id="KT220438", rettype="gb", \
                                                               retmode="text")
gb file contents = handle.read()
handle.close()
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
VERSION
           KT220438.1 GI:887493048
KEYWORDS
            Influenza A virus (A/New Jersey/NHRC 93219/2015(H3N2))
SOURCE
           Influenza A virus (A/New Jersey/NHRC 93219/2015(H3N2))
 ORGANISM
            Viruses; ssRNA viruses; ssRNA negative-strand viruses;
            Orthomyxoviridae; Influenzavirus A.
            1 (bases 1 to 1701)
REFERENCE
            Sitz, C.R., Thammavong, H.L., Balansay-Ames, M.S., Hawksworth, A.W.,
  AUTHORS
```

### We can request data as text or as XML

```
handle = Entrez.efetch(db="nucleotide", id="KT220438", rettype="gb", \
                                                                retmode="xml")
gb file contents = handle.read()
handle.close()
print(gb file contents)
<?xml version="1.0" ?>
<!DOCTYPE GBSet PUBLIC "-//NCBI//NCBI GBSeq/EN"</pre>
"https://www.ncbi.nlm.nih.gov/dtd/NCBI GBSeg.dtd">
<GBSet>
<GBSeq>
  <GBSeq locus>KT220438</GBSeq locus>
  <GBSeq length>1701</GBSeq length>
  <GBSeq strandedness>single</GBSeq strandedness>
  <GBSeq moltype>cRNA</GBSeq moltype>
  <GBSeq topology>linear</GBSeq topology>
  <GBSeq division>VRL</GBSeq division>
  <GBSeq update-date>20-JUL-2015</GBSeq update-date>
  <GBSeq create-date>20-JUL-2015</GBSeq create-date>
  <GBSeq definition>Influenza A virus (A/NewJersey/NHRC 93219/2015(H3N2))
```

#### Pros and cons of text and XML

#### Text:

- Easier to read for humans
- Requires special parser for each datatype

#### XML:

- Very hard to read for humans
- Can be parsed with a generic parser

### We parse text format with SeqIO.read()

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```
handle = Entrez.efetch(db="nucleotide", id="KT220438", rettype="qb", \
                                                                retmode="text")
record = SeqIO.read(in handle, format="gb") # use SeqIO.read() to parse
handle.close()
print(record)
ID: KT220438.1
Name: KT220438
Description: Influenza A virus (A/NewJersey/NHRC 93219/2015(H3N2)) segment 4
hemagglutinin (HA) gene, complete cds.
Number of features: 5
/data file division=VRL
/date=20-JUL-2015
/accessions=['KT220438']
/sequence version=1
/keywords=['']
/source=Influenza A virus (A/New Jersey/NHRC 93219/2015(H3N2))
/organism=Influenza A virus (A/New Jersey/NHRC 93219/2015(H3N2))
/taxonomy=['Viruses', 'ssRNA viruses', 'ssRNA negative-strand viruses',
'Orthomyxoviridae', 'Influenzavirus A']
/references=[Reference(title='GEISS Influenza Surveillance Response Program',
...), Reference(title='Direct Submission', ...)]
```

# We parse XML format with Entrez.parse()

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```
handle = Entrez.efetch(db="nucleotide", id="KT220438", rettype="gb", \
                                                               retmode="xml")
parsed = Entrez.parse(in handle) # use Entrez.parse() to parse
record = list(parsed)[0] # Need to convert into list and get 1st element
handle.close()
print(record) # Record contains nested dictionaries and lists
{'GBSeq locus': 'KT220438', 'GBSeq length': '1701', 'GBSeq strandedness':
'single', 'GBSeq moltype': 'cRNA', 'GBSeq topology': 'linear',
'GBSeq division': 'VRL', 'GBSeq update-date': '20-JUL-2015', 'GBSeq create-
date': '20-JUL-2015', 'GBSeq definition': 'Influenza A virus
(A/NewJersey/NHRC 93219/2015(H3N2)) segment 4 hemagglutinin (HA) gene, complete
cds', 'GBSeq primary-accession': 'KT220438', 'GBSeq accession-version':
'KT220438.1', 'GBSeq other-seqids': ['gb|KT220438.1|', 'gi|887493048'],
'GBSeq source': 'Influenza A virus (A/New Jersey/NHRC 93219/2015(H3N2))',
'GBSeq organism': 'Influenza A virus (A/New Jersey/NHRC 93219/2015(H3N2))',
'GBSeq taxonomy': 'Viruses; ssRNA viruses; ssRNA negative-strand viruses;
Orthomyxoviridae; Influenzavirus A', 'GBSeq references':
[{'GBReference reference': '1', 'GBReference position': '1..1701',
'GBReference authors': ['Sitz,C.R.', 'Thammavong,H.L.', 'Balansay-Ames,M.S.',
'Hawksworth, A.W.', 'Myers, C.A.', 'Brice, G.T.'], 'GBReference title': 'GEISS
Influenza Surveillance Response Program', 'GBReference journal':
```

# All information from parsed XML format can be accessed using dict & list methods

```
# extract all the features
features = record['GBSeq_feature-table']
# print feature key & location for all features
for feature in features:
    print(feature['GBFeature key'] + ": " + \
                        feature['GBFeature location'])
source: 1..1701
gene: 1..1701
CDS: 1..1701
mat peptide: 49..1035
mat peptide: 1036..1698
```

# All information from parsed XML format can be accessed using dict & list methods

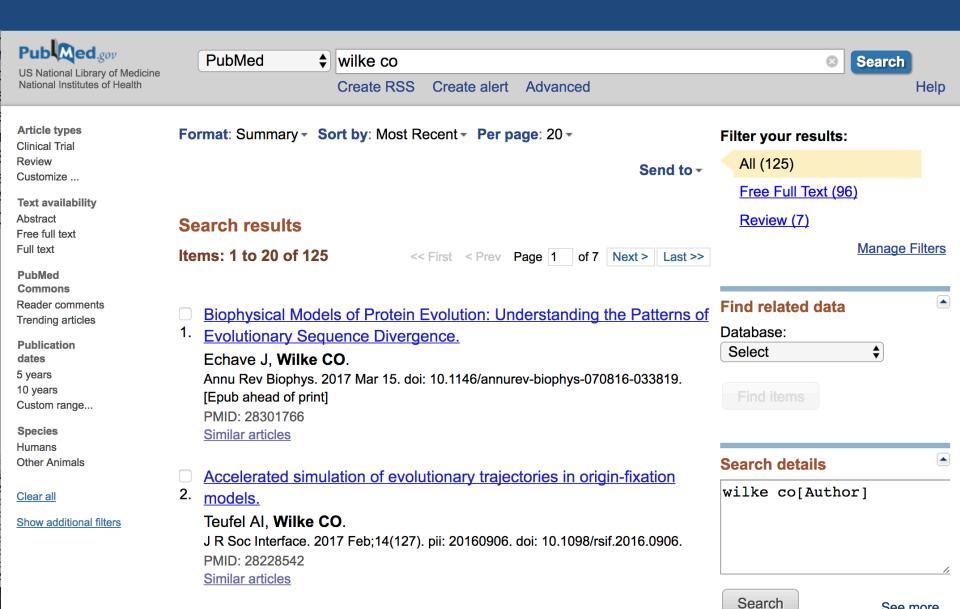
```
# extract all the features
features = record['GBSeq feature-table']
# print feature key & location for all features
for feature in features:
    print(feature['GBFeature key'] + ": " + \
                        feature['GBFeature location'])
source: 1..1701
gene: 1..1701
CDS: 1..1701
mat peptide: 49..1035
mat peptide: 1036..1698
```

# All information from parsed XML format can be accessed using dict & list methods

```
# extract all the features
features = record['GBSeq feature-table']
# print feature key & location for all features
for feature in features:
    print(feature['GBFeature key'] + ": " + \
                        feature['GBFeature location'])
source: 1..1701
gene: 1..1701
CDS: 1..1701
mat peptide: 49..1035
mat_peptide: 1036..1698
```

## Running searches through Entrez

## Example: Literature search using pubmed



### Example: Literature search using pubmed

#### We retrieve search results with efetch()

```
# For references, the file format is called "Medline"
from Bio import Medline
handle = Entrez.efetch(db="pubmed", id=pmid list,
                       rettype="medline", retmode="text")
records = Medline.parse(handle)
# Must not close handle yet!
for record in records:
    print(record['AU']) # author list
    print(record['TI']) # title
    print(record['SO']) # source (reference)
    print()
handle.close() # Close after all records have been processed
```

#### We retrieve search results with efetch()

```
['Echave J', 'Wilke CO']
Biophysical Models of Protein Evolution: Understanding the Patterns of
Evolutionary Sequence Divergence.
Annu Rev Biophys. 2017 Mar 15. doi: 10.1146/annurev-biophys-070816-033819.
['Teufel AI', 'Wilke CO']
Accelerated simulation of evolutionary trajectories in origin-fixation models.
J R Soc Interface. 2017 Feb; 14(127). pii: 20160906. doi:
10.1098/rsif.2016.0906.
['Lipsitch M', 'Barclay W', 'Raman R', 'Russell CJ', 'Belser JA', 'Cobey S',
'Kasson PM', 'Lloyd-Smith JO', 'Maurer-Stroh S', 'Riley S', 'Beauchemin CA',
'Bedford T', 'Friedrich TC', 'Handel A', 'Herfst S', 'Murcia PR', 'Roche B',
'Wilke CO', 'Russell CA']
Viral factors in influenza pandemic risk assessment.
Elife. 2016 Nov 11;5. pii: e18491. doi: 10.7554/eLife.18491.
['McWhite CD', 'Meyer AG', 'Wilke CO']
Sequence amplification via cell passaging creates spurious signals of positive
adaptation in influenza virus H3N2 hemagglutinin.
Virus Evol. 2016 Jul; 2(2). pii: vew026. Epub 2016 Oct 3.
['Spielman SJ', 'Wan S', 'Wilke CO']
A Comparison of One-Rate and Two-Rate Inference Frameworks for Site-Specific
dN/dS Estimation.
```