Traineeship Part 1 NCBI-DPP9_continued_adapted

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Traineeship Part 1: Data collection using NCBI eUtils and esummary CONTIN-UED/ADAPTED - DPP9

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Loading required packages

```
[1]: # pip3 install --user eutils
import MySQLdb as my
from eutils import Client
from Bio import Entrez
import csv
```

UCSC connection for ncbiRefSeq search (mRNA/RNA transcripts with hg19 coordinates)

```
[3]: print("\nLoading currently available accession numbers from NCBI RefSeq table...

→")

print("="*50)

print("\nTranscript variant accession numbers: ")

accList = []

### Save data to csv file

with open('results-transcripts-UCSC.csv', mode='w') as result_transcripts:

result_writer = csv.writer(result_transcripts,delimiter=';')
```

```
result_writer.
 →writerow(["chromosome", "start", "end", "strand", "gene", "exonCount", "accession"])
    for row in result:
        transcript = row[1]
        print(transcript)
        accList.append(transcript)
        starts = str(row[9])[2:-2]
        ends = str(row[10])[2:-2]
        starts1 = starts.split(",")
        ends1 = ends.split(",")
        j = 0
        for i in starts1:
            result_writer.
\rightarrowwriterow([row[2],i,ends1[j],row[3],row[12],row[8],row[1]])
print("\nSearch results: {}\n".format(no_rows))
### Close csv file
result_transcripts.close()
```

Loading currently available accession numbers from NCBI RefSeq table...

```
Transcript variant accession numbers: NM_139159.5 NR_158699.2 NM_001365987.2 NR_164163.1 Search results: 4
```

Personal API-key for NCBI search

```
[4]: eclient = Client(api_key="8ecce891e7fa036ff84bccc7c74e5138dc09")
```

1) Entrez Nucleotide Search - mRNA Transcript Variants

```
[5]: ### Creating query

mRNAtranscripts = []

transcriptmRNA_esearch = eclient.esearch(db='nucleotide',

term='(DPP9[gene] AND "Homo sapiens"[Primary Organism] AND

→refseq[filter]) NOT biomol_genomic[PROP]')

print("\nLoading currently available ids from Entrez nucleotide...")

print("="*50)
```

```
print("\nTranscript variant ids: ")
    print(transcriptmRNA_esearch.ids)
    for item in transcriptmRNA_esearch.ids:
        mRNAtranscripts.append(item)
    print("\nSearch results: {}\n".format(transcriptmRNA_esearch.count))
    Loading currently available ids from Entrez nucleotide...
    _____
    Transcript variant ids:
    [1370476185, 1034610004, 1034610002, 768004630, 768004626, 768004622, 768004618,
    768004616, 578833714, 1677498370, 1677499978, 1700660497]
    Search results: 12
[]: ### Esummary for retrieving information
    Entrez.email = "iris.raes@hotmail.com"
     ### For each id in mRNAtranscripts
     ### Save data to csv file
    with open('results-nucleotide.csv', mode='w') as result_nucleotide:
        result_writer = csv.writer(result_nucleotide,delimiter=';')
        result_writer.
     writerow(["transcript_id", "description", "transcript_variant", "accession", "length_in_bp"])
        counter = 1
        for ids in mRNAtranscripts:
            handle = Entrez.esummary(db="nucleotide", id=ids)
            record = Entrez.read(handle)
            handle.close()
            ### Write info to csv file, row by row
            splittedtitle = record[0]["Title"].split(",")
            print(splittedtitle)
            result_writer.
     →writerow([record[0]["Id"],splittedtitle[0],splittedtitle[1],record[0]["AccessionVersion"],r
            counter += 1
     ### Close csv file
    result_nucleotide.close()
    2) dbVar Search - Pathogenic Copy Number Variation in Human
```

```
[6]: ### Creating query

CNV = []

CNV_esearch = eclient.esearch(db='dbVar',

term='DPP9[All Fields] AND ("Homo sapiens"[Organism] AND "copy

→number variation"[Variant Type] AND "Pathogenic"[clinical_interpretation])')
```

Loading currently available ids from dbVar...

```
dbVar ids:
```

[49623411, 49353191, 49353005, 49350830, 49349701, 49349293, 49345450, 49344315, 48468240, 48466558, 48466447, 48453939, 45807136, 17813982, 17813734, 3740775, 3739972, 3738955, 3738954, 3738649, 1212838, 1137112]

```
[]: ### Esummary for retrieving information
    Entrez.email = "iris.raes@hotmail.com"
    ### For each id in CNV
     ### Save data in csv file
    with open('results-CNV-dbVar.csv', mode='w') as result_CNV:
        result writer = csv.writer(result CNV,delimiter=';')
        result writer.
     →writerow(["CNV_variant_id", "variant_region_id", "type", "study_ID", "clinical_assertion", "Chr_
         counter = 1
        for ids in CNV:
            handle = Entrez.esummary(db="dbVar", id=ids)
             record = Entrez.read(handle)
            handle.close()
             varregid = record['DocumentSummarySet']['DocumentSummary'][0].get('SV')
             types = record['DocumentSummarySet']['DocumentSummary'][0].
      →get('dbVarVariantTypeList')
             studyid = record['DocumentSummarySet']['DocumentSummary'][0].get('ST')
             clinicalassertion = record['DocumentSummarySet']['DocumentSummary'][0].
      →get('dbVarClinicalSignificanceList')
      →record['DocumentSummarySet']['DocumentSummary'][0]['dbVarPlacementList'] !=

      → []:
                 Chr_1 =_
      →record['DocumentSummarySet']['DocumentSummary'][0]['dbVarPlacementList'][0].

    get('Chr')

                 assembly1 = 
      →record['DocumentSummarySet']['DocumentSummary'][0]['dbVarPlacementList'][0].
```

```
start1 =
 →record['DocumentSummarySet']['DocumentSummary'][0]['dbVarPlacementList'][0].
 end1 = 1
 →record['DocumentSummarySet']['DocumentSummary'][0]['dbVarPlacementList'][0].
 Chr_2 = 
 →record['DocumentSummarySet']['DocumentSummary'][0]['dbVarPlacementList'][1].

    get('Chr')
           assembly2 = 1
 →record['DocumentSummarySet']['DocumentSummary'][0]['dbVarPlacementList'][1].
 →record['DocumentSummarySet']['DocumentSummary'][0]['dbVarPlacementList'][1].

→get('Chr_start')
           end2 = 1
→record['DocumentSummarySet']['DocumentSummary'][0]['dbVarPlacementList'][1].

    get('Chr_end')
       ### Write info to csv file, row by row
       result_writer.
 →writerow([ids,varregid,types,studyid,clinicalassertion,Chr_1,assembly1+":
 \rightarrow"+start1+"-"+end1,Chr 2,assembly2+":"+start2+"-"+end2])
       counter += 1
### Close csv file
result_CNV.close()
```

3) dbVar Search - Insertions in Human

```
Loading currently available ids from dbVar...

-----
dbVar ids:
```

[49597698, 49580472, 48530760, 48377645, 48377627, 47753859, 47564069, 47178696, 46791711, 45897195, 45896455, 45807279, 36885535, 24618684, 24516168, 24501143, 17814018, 17813982, 14212055, 14211117, 14209696, 13414404, 11399938, 8023314, 7738722, 7694891, 7590450, 7474153, 6477950, 6451851, 6354196, 5661470, 5431842, 5195919, 1297001, 1028299, 286824, 285317, 284926, 40396]

```
[]: ### Esummary for retrieving information
    Entrez.email = "iris.raes@hotmail.com"
    ### For each id in insertion
    ### Save data to csv file
    with open('results-insertion-dbVar.csv', mode='w') as result_insertion:
        result_writer = csv.writer(result_insertion,delimiter=';')
        result_writer.
     →writerow(["insertion_variant_id", "variant_region_id", "type", "study_ID", "Chr_1", "assembly1",
        counter = 1
        for ids in insertion:
            handle = Entrez.esummary(db="dbVar", id=ids)
            record = Entrez.read(handle)
            handle.close()
            varregid = record['DocumentSummarySet']['DocumentSummary'][0].get('SV')
            types = record['DocumentSummarySet']['DocumentSummary'][0].
     studyid = record['DocumentSummarySet']['DocumentSummary'][0].get('ST')
     →record['DocumentSummarySet']['DocumentSummary'][0]['dbVarPlacementList'] !=,,
     □ :
                Chr_1 = 
     →record['DocumentSummarySet']['DocumentSummary'][0]['dbVarPlacementList'][0].

    get('Chr')

                assembly1 = 
     →record['DocumentSummarySet']['DocumentSummary'][0]['dbVarPlacementList'][0].
     start1 =
     →record['DocumentSummarySet']['DocumentSummary'][0]['dbVarPlacementList'][0].
     end1 = 1
     →record['DocumentSummarySet']['DocumentSummary'][0]['dbVarPlacementList'][0].

    get('Chr_end')

                Chr_2 = 
     →record['DocumentSummarySet']['DocumentSummary'][0]['dbVarPlacementList'][1].

    get('Chr')

                assembly2 = 
     →record['DocumentSummarySet']['DocumentSummary'][0]['dbVarPlacementList'][1].
```

4) dbVar Search - Inversions in Human

Loading currently available ids from dbVar...

dbVar ids:

[48377627, 47178696, 46791711, 45807289, 45807279, 36885535, 24618684, 24516168, 24501143, 17814018, 17813982, 5195919, 1297001, 1028299]

```
counter = 1
   for ids in inversion:
       handle = Entrez.esummary(db="dbVar", id=ids)
        record = Entrez.read(handle)
       handle.close()
        varregid = record['DocumentSummarySet']['DocumentSummary'][0].get('SV')
        types = record['DocumentSummarySet']['DocumentSummary'][0].
 →get('dbVarVariantTypeList')
        studyid = record['DocumentSummarySet']['DocumentSummary'][0].get('ST')
 →record['DocumentSummarySet']['DocumentSummary'][0]['dbVarRemappedAssemblyList']_
 →!= []:
            assembly1 =
→record['DocumentSummarySet']['DocumentSummary'][0]['dbVarRemappedAssemblyList'][0]
            assembly2 = 
 →record['DocumentSummarySet']['DocumentSummary'][0]['dbVarRemappedAssemblyList'][0]
        ### Write info to csv file, row by row
        result writer.
 →writerow([ids,varregid,types,studyid,"Chr19",assembly1,"Chr19",assembly2])
       counter += 1
### Close csv file
result_inversion.close()
```

5) dbVar Search - Short Tandem Repeats in Human (seems to be less important)

```
[9]: ### Creating query

STR = []

STR_esearch = eclient.esearch(db='dbVar',

term='DPP9[All Fields] AND ("Homo sapiens"[Organism] AND "short

tandem repeat"[Variant Type])')

print("\nLoading currently available ids from dbVar...")

print("="*50)

print("dbVar ids: ")

print(STR_esearch.ids)

for item in STR_esearch.ids:

STR.append(item)

print("\nSearch results: {}\n".format(STR_esearch.count))
```

35727324, 35726686, 35726677, 35726669, 35726663, 35726639, 30349921]

35728601, 35728076, 35727391, 35727380, 35727364, 35727355, 35727352, 35727332,

```
[]: ### Esummary for retrieving information
    Entrez.email = "iris.raes@hotmail.com"
    ### For each id in STR
    ### Save data to csv file
    with open('results-STR-dbVar.csv', mode='w') as result_STR:
        result_writer = csv.writer(result_STR,delimiter=';')
        result_writer.
     →writerow(["STR_variant_id","variant_region_id","type","study_ID","Chr_1","assembly1","Chr_2
        counter = 1
        for ids in STR:
            handle = Entrez.esummary(db="dbVar", id=ids)
            record = Entrez.read(handle)
            handle.close()
            varregid = record['DocumentSummarySet']['DocumentSummary'][0].get('SV')
            types = record['DocumentSummarySet']['DocumentSummary'][0].
     →get('dbVarVariantTypeList')
            studyid = record['DocumentSummarySet']['DocumentSummary'][0].get('ST')
            if
     →record['DocumentSummarySet']['DocumentSummary'][0]['dbVarPlacementList'] !=
                Chr_1 =_
     →record['DocumentSummarySet']['DocumentSummary'][0]['dbVarPlacementList'][0].

    get('Chr')
                assembly1 = 
     →record['DocumentSummarySet']['DocumentSummary'][0]['dbVarPlacementList'][0].
     start1 =
     →record['DocumentSummarySet']['DocumentSummary'][0]['dbVarPlacementList'][0].
     end1 =
     →record['DocumentSummarySet']['DocumentSummary'][0]['dbVarPlacementList'][0].

    get('Chr_end')
                Chr_2 = 
     →record['DocumentSummarySet']['DocumentSummary'][0]['dbVarPlacementList'][1].

    get('Chr')

                assembly2 = 1
     →record['DocumentSummarySet']['DocumentSummary'][0]['dbVarPlacementList'][1].
     start2 =
     →record['DocumentSummarySet']['DocumentSummary'][0]['dbVarPlacementList'][1].
```

```
end2 = □

→record['DocumentSummarySet']['DocumentSummary'][0]['dbVarPlacementList'][1].

→get('Chr_end')

### Write info to csv file, row by row

result_writer.writerow([ids,varregid,types,studyid,Chr_1,assembly1+":

→"+start1+"-"+end1,Chr_2,assembly2+":"+start2+"-"+end2])

###

counter += 1

### Close csv file

result_STR.close()
```

6) ClinVar Search - Genetic Variations in Human

```
handle.close()
      title = record['DocumentSummarySet']['DocumentSummary'][0].get('title')
      accession = record['DocumentSummarySet']['DocumentSummary'][0].
types = record['DocumentSummarySet']['DocumentSummary'][0].
description =
→record['DocumentSummarySet']['DocumentSummary'][0]['clinical_significance'].

→get('description')
      protein_change = record['DocumentSummarySet']['DocumentSummary'][0].

¬record['DocumentSummarySet']['DocumentSummary'][0]['variation_set'][0]['variation_loc']

→!= []:
          Chr_1 =_
→record['DocumentSummarySet']['DocumentSummary'][0]['variation_set'][0]['variation_loc'][0].

    get('chr')
          assembly1 = \square
→record['DocumentSummarySet']['DocumentSummary'][0]['variation_set'][0]['variation_loc'][0].
start1 =
→record['DocumentSummarySet']['DocumentSummary'][0]['variation_set'][0]['variation_loc'][0].
⇔get('start')
→record['DocumentSummarySet']['DocumentSummary'][0]['variation_set'][0]['variation_loc'][0].

    get('stop')
→record['DocumentSummarySet']['DocumentSummary'][0]['variation_set'][0].
try:
                  Chr_2 = 
→record['DocumentSummarySet']['DocumentSummary'][0]['variation_set'][0]['variation_loc'][1].

    get('chr')
                  assembly2 = 
→record['DocumentSummarySet']['DocumentSummary'][0]['variation_set'][0]['variation_loc'][1].
start2 =
→record['DocumentSummarySet']['DocumentSummary'][0]['variation_set'][0]['variation_loc'][1].

    get('start')
                  end2 = 
→record['DocumentSummarySet']['DocumentSummary'][0]['variation_set'][0]['variation_loc'][1].

    get('stop')
              except:
                  assembly2 = "not applicable"
                  start2 = "X"
                  end2 = "X"
```