

## Load libraries

```
In [3]: import numpy as np                                #NumPy
        from os import listdir                            #List items in directory
        from Bio.Align import PairwiseAligner             #Used to align sequences (my 1
        import matplotlib.pyplot as plt                  #Data visualization (for the p
        from scipy.cluster.hierarchy import dendrogram   #Plot a dendrogram/phylogeneti
        from scipy.cluster.hierarchy import linkage       #Clustering required to by den
        from matplotlib.pyplot import figure             #Used to adjust the figure siz
        from Bio import AlignIO                           #Used to import an aligned fas
```

## Create a FASTA node

```
In [4]: class FASTA():
        def __init__(self, name, sequence, animal):
            self.name = name
            self.sequence = sequence
            self.animal = animal
```

## Load FASTA files while also combining the data into 1 file

### Soy Leghemoglobin Data

```
In [5]: leg_data = []    # Basic data set
        combined = open("Alignment\leg_data_unaligned.fasta", "w")

        for filename in listdir("Raw Data\Soy Leghemoglobin"):
            file = open("Raw Data\Soy Leghemoglobin\\"+filename, "r")
            name = file.readline()[1:]
            combined.write(">" + name)

            sequence = ""
            for line in file.read().split("\n"):
                if not line == "":
                    sequence += line
            combined.write(sequence+"\n\n")

            leg_data.append(FASTA(name, sequence, "Soy"))

            file.close()
        combined.close()
```

### Various Myoglobin Data

```
In [6]: myo_data = []
        combined = open("Alignment\myo_data_unaligned.fasta", "w")

        for filename in listdir("Raw Data\Various Myoglobin"):
            file = open("Raw Data\Various Myoglobin\\"+filename, "r")
            name = file.readline()[1:]
            combined.write(">" + name)

            sequence = ""
            for line in file.read().split("\n"):
                if not line == "":
                    sequence += line
            combined.write(sequence+"\n\n")
```

```

myo_data.append(FASTA(name, sequence, filename[:filename.index(" ")]))

file.close()
combined.close()

```

## Build a Phylogenetic Tree using a Distance Matrix

```

In [7]: def build_tree(data, labels = None):
        # Create a distance matrix from the sequences
        dists = []
        for i in range(len(data)-1):
            dists_i = []
            for j in range(i+1, len(data)):
                alignment = PairwiseAligner().align(data[i].sequence, data[j].sequence)
                dists_i.append(alignment.score)
            dists.append(dists_i)
        # Fill repetitive half of distance matrix
        dists.append([])
        dists = np.flip(dists).tolist()
        for i in range(int((len(dists)+1)/2)):
            dist_save = dists[i]
            dists[i] = dists[i] + [0] + np.flip(dists[len(dists)-i-1]).tolist()
            if i!=len(dists)-i-1:
                dists[len(dists)-i-1] = dists[len(dists)-i-1] + [0] + np.flip(dist_save).to
        # Build tree
        linkage_matrix = linkage(dists)
        no_labels = False
        if labels == None:
            no_labels = True
        dendrogram(linkage_matrix, no_labels = no_labels, labels=labels, orientation="right")
        plt.show()

```

## Leghemoglobin data set

```

In [8]: figure(figsize=(10,6))
        build_tree(leg_data)
        #Very large numbers, hence they are super similar

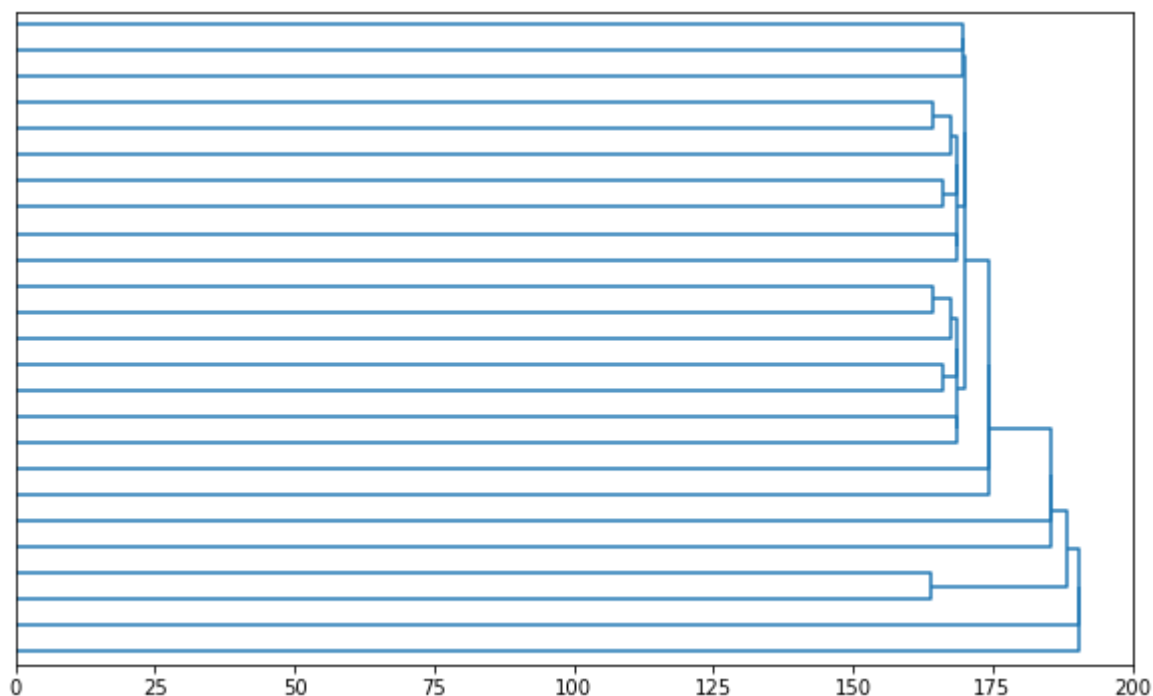
```

C:\Users\truon\anaconda3\lib\site-packages\numpy\core\\_asarray.py:83: VisibleDeprecationWarning: Creating an ndarray from ragged nested sequences (which is a list-or-tuple of lists-or-tuples-or ndarrays with different lengths or shapes) is deprecated. If you meant to do this, you must specify 'dtype=object' when creating the ndarray

```

return array(a, dtype, copy=False, order=order)

```



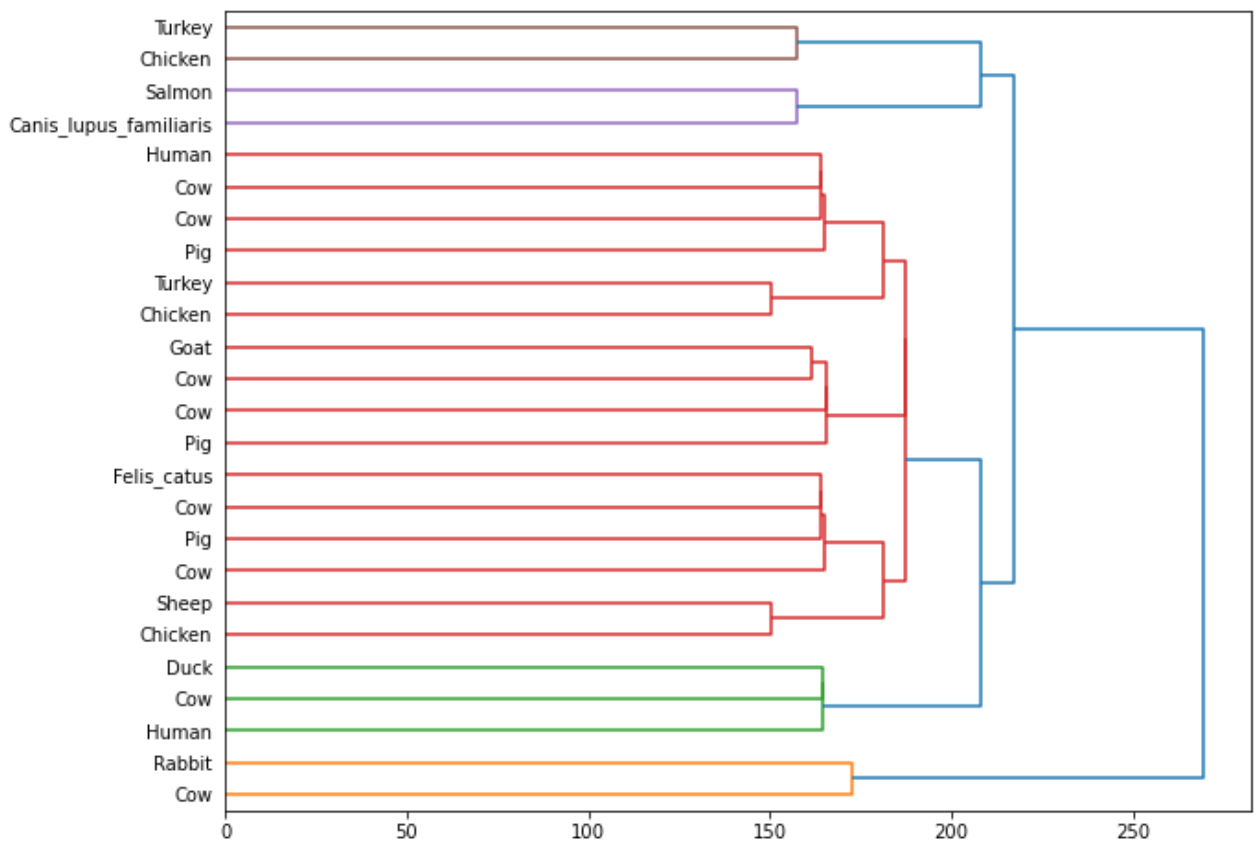
## Myoglobin data set

Gather animals for figure label

```
In [9]: myo_labels = []  
        for entry in myo_data:  
            myo_labels.append(entry.animal)
```

Plot figure

```
In [10]: figure(figsize=(10,8))  
         build_tree(myo_data, myo_labels)  
         # No correlation, hence there isn't really a difference
```



## Combination of both sets

### Combine data

```
In [11]: all_data = myo_data + leg_data
```

### Remove outliers (Comment to view original results)

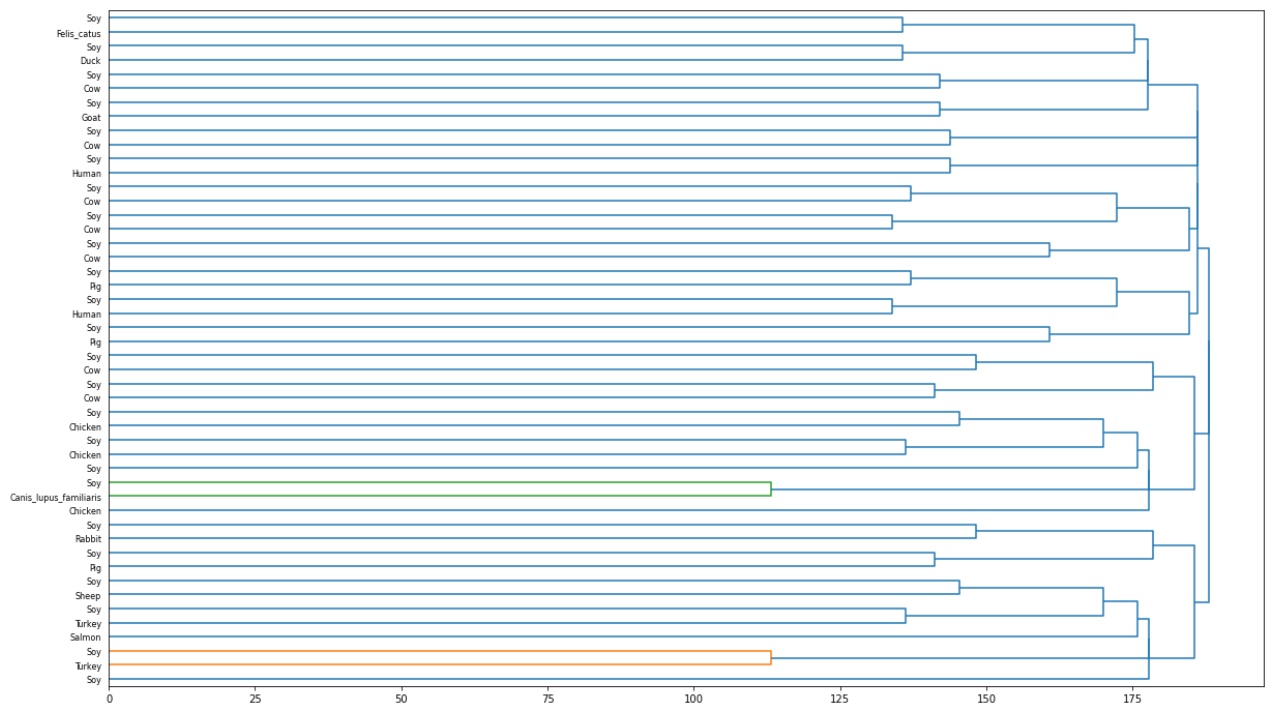
```
In [12]: all_data = np.delete(all_data, [4,45]).tolist()
```

### Gather animals for figure label

```
In [13]: all_labels = []
for entry in all_data:
    all_labels.append(entry.animal)
```

### Plot figure

```
In [14]: figure(figsize=(20,12))
build_tree(all_data, all_labels)
#Generally random/low differences; roughly every soy's closest neighbor is non-soy
```



**\*Align each combined file extrenally (ClustalOmega:**  
<https://www.ebi.ac.uk/Tools/msa/clustalo/>)

Align X\_data\_unaligned.fasta and save results onto X\_data\_aligned.clustal

Create a combination both of files

```
In [15]: file = open("Alignment\combined_data_unaligned.fasta", "w")
leg = open("Alignment\leg_data_unaligned.fasta", "r")
myo = open("Alignment\myo_data_unaligned.fasta", "r")

file.write(leg.read())
file.write(myo.read())

leg.close()
myo.close()
file.close()
```

**Import extrenal alignment of sequences (should be included with project)**

**Leghemoglobin data**

```
In [42]: leg_data_aligned = AlignIO.read(open("Alignment\leg_data_aligned.clustal"), "clustal")
for entry in leg_data_aligned:
    print(entry.seq+" |(Soy)" + entry.id)
# Very clean alignment (low indels), with the exception of the first 2 insertions

MGAFTEKQEALVNSSF EAFKANLP HHSVVFNS ILEKAPAAKNMFS FLGDAVDPKNPKLAGHAEKLFGLVRDSAVQLQTKGLVVADAT
LGPIHTQKGVTDLQFAVVKEALLKTIKEAVGDKWSEELSNPWEVAYDEIAAAIKKAMAIGSLV |(Soy)V00451.1
MGAFTEKQEALVNSSF EAFKANLP HHSVVFNS ILEKAPAAKNMFS FLGDAVDPKNPKLAGHAEKLFGLVRDSAVQLQTKGLVVADAT
LGPIHTQKGVTDLQFAVVKEALLKTIKEAVGDKWSEELSNWEVAYDEIAAAIKKAMAIGSLV |(Soy)KHN37872.1
-GAFTEKQDALVSSSF EAFKANIPQYSVVFNS ILEKAPAAKDLFSLANGVDPTNP KLTGHA EKLFALVRDSAGQLKASGTVVADAA
LGSIH AQKAVTNPEFV-VKEALLKTIKEAVGDKWSEELSSAWEVAYDELA AAIKKAF----- |(Soy)pir|GPSYC2
-VAFTEKQDALVSSSF EAFKANIPQYSVVFYTS ILEKAPAAKDLFSLANGVDPTNP KLTGHA EKLFALVRDSAGQLKASGTVVADAA
LGSVHAQKAVTDPQFVVVKEALLKTIKAAVGDKWSEELSR AWEVAYDELA AAIKKA----- |(Soy)pdb|1FSL|A
-VAFTEKQDALVSSSF EAFKANIPQYSVVFYTS ILEKAPAAKDLFSLANGVDPTNP KLTGHA EKLFALVRDSAGQLKASGTVVADAA
```

```

LGSVHAQKAVTDPQFVVVKEALLKTIKAAVGDKWSDELSRAEVAYDELA AAIKKA----- |(Soy)pdb|1FSL|B
MVAFTEKQDALVSSSF EAFKANIPQYSVVFYTSILEKAPAAKDLFSFLANGVDPTNP KLTGHA EKL FALVRDSAGQLKASGTVVADAA
LGSVHAQKAVTDPQFVVVKEALLKTIKAAVGDKWSDELSRAEVAYDELA AAIKKA----- |(Soy)V00453.1
MVAFTEKQDALVSSSF EAFKANIPQYSVVFYTSILEKAPAAKDLFSFLANGVDPTNP KLTGHA EKL FALVRDSAGQLKASGTVVADAA
LGSVHAQKAVTDPQFVVVKEALLKTIKAAVGDKWSDELSRAEVAYDELA AAIKKA----- |(Soy)NP_001235928.1
MVAFTEKQDALVSSSF EAFKANIPQYSVVFYTSILEKAPAAKDLFSFLANGVDPTNP KLTGHA EKL FALVRDSAGQLKASGTVVADAA
LGSVHAQKAVTDPQFVVVKEALLKTIKAAVGDKWSDELSRAEVAYDELA AAIKKA----- |(Soy)KHN37870.1
MVAFTEKQDALVSSSF EAFKANIPQYSVVFYTSILEKAPAAKDLFSFLANGVDPTNP KLTGHA EKL FALVRDSAGQLKASGTVVADAA
LGSVHAQKAVTDPQFVVVKEALLKTIKAAVGDKWSDELSRAEVAYDELA AAIKKA----- |(Soy)sp|P02238|LGBA_SOY
BN
MVAFTEKQDALVSSSF EAFKANIPQYSVVFYTSILEKAPAAKDLFSFLANGVDPTNP KLTGHA EKL FALVRDSAGQLKASGTVVADAA
LGSVHAQKAVTDPQFVVVKEALLKTIKAAVGDKWSDELSRAEVAYDELA AAIKKA----- |(Soy)CAA23731.1
MGAFTEKQDALVSSSF EAFKANIPQYSVVFYTSILEKAPAAKDLFSFLANGVDPTNP KLTGHA EKL FALVRDSAGQLKASGTVVADAA
LVSIIHAQKAVTDPQFVVVKEALLKTIKAAVGGNWSDELSSA EVAYDELA AAIKKA----- |(Soy)V00452.1
MGAFTEKQDALVSSSF EAFKANIPQYSVVFYTSILEKAPAAKDLFSFLANGVDPTNP KLTGHA EKL FALVRDSAGQLKASGTVVADAA
LVSIIHAQKAVTDPQFVVVKEALLKTIKAAVGGNWSDELSSA EVAYDELA AAIKKA----- |(Soy)NP_001345001.1
MGAFTEKQDALVSSSF EAFKANIPQYSVVFYTSILEKAPAAKDLFSFLANGVDPTNP KLTGHA EKL FALVRDSAGQLKASGTVVADAA
LVSIIHAQKAVTDPQFVVVKEALLKTIKAAVGGNWSDELSSA EVAYDELA AAIKKA----- |(Soy)sp|P02235|LGB1_SOY
BN
MGAFTEKQDALVSSSF EAFKANIPQYSVVFYTSILEKAPAAKDLFSFLANGVDPTNP KLTGHA EKL FALVRDSAGQLKASGTVVADAA
LVSIIHAQKAVTDPQFVVVKEALLKTIKAAVGGNWSDELSSA EVAYDELA AAIKKA----- |(Soy)KHN37871.1
MGAFTEKQDALVSSSF EAFKANIPQYSVVFYTSILEKAPAAKDLFSFLANGVDPSNP KLTGHA EKL FGLVRDSAGQLKANGTVVADAA
LGSIIHAQKAITDPQFVVVKEALLKTIKAAVGDKWSDELSSA EVAYDELA AAIKKA----- |(Soy)NM_001248319.3
MGAFTEKQDALVSSSF EAFKANIPQYSVVFYTSILEKAPAAKDLFSFLANGVDPSNP KLTGHA EKL FGLVRDSAGQLKANGTVVADAA
LGSIIHAQKAITDPQFVVVKEALLKTIKAAVGDKWSDELSSA EVAYDELA AAIKKA----- |(Soy)NP_001235248.2
MGAFTEKQDALVSSSF EAFKANIPQYSVVFYTSILEKAPAAKDLFSFLANGVDPSNP KLTGHA EKL FGLVRDSAGQLKANGTVVADAA
LGSIIHAQKAITDPQFVVVKEALLKTIKAAVGDKWSDELSSA EVAYDELA AAIKKA----- |(Soy)KHN00941.1
MGAFTEKQDALVSSSF EAFKANIPQYSVVFYTSILEKAPAAKDLFSFLANGVDPSNP KLTGHA EKL FGLVRDSAGQLKANGTVVADAA
LGSIIHAQKAITDPQFVVVKEALLKTIKAAVGDKWSDELSSA EVAYDELA AAIKKA----- |(Soy)AAA33980.1
MGAFTEKQDALVSSSF EAFKANIPQYSVVFYTSILEKAPAAKDLFSFLANGVDPSNP KLTGHA EKL FGLVRDSAGQLKANGTVVADAA
LGSIIHAQKAITDPQFVVVKEALLKTIKAAVGDKWSDELSSA EVAYDELA AAIKKA----- |(Soy)sp|P02236|LGB2_SOY
BN
MGAFTEKQDALVSSSF EAFKANIPQYSVVFYTSILEKAPAAKDLFSFLANGVDPSNP KLTGHA EKL FGLVRDSAGQLKANGTVVADAA
LGSIIHAQKAITDPQFVVVKEALLKTIKAAVGDKWSDELSSA EVAYDELA AAIKKA----- |(Soy)sp|P02236.2|LGB2_S
OYBN
MGAFTEKQDALVSSSF EAFKANIPQYSVVFYTSILEKAPAAKDLFSFLANGVDPSNP KLTGHA EKL FGLVRDSAGQLKANGTVVADAA
LGSIIHAQKAITDPQFVVVKEALLKTIKAAVGDKWSDELSSA EVAYDELA AAIKKA----- |(Soy)J01301.1
MGAFTEKQDALVSSSF EAFKANIPQYSVVFYTSILEKAPAAKDLFSFLANGVDPTNP KLTGHA EKL FGLVRDSAGQLKASGTVVIDAA
LGSIIHGQKAITDPQFVVVKEALLKTIKAAVGDKWSDELSSA EVAYDELA AAIKKA----- |(Soy)NP_001235423.1
MGAFTEKQDALVSSSF EAFKANIPQYSVVFYTSILEKAPAAKDLFSFLANGVDPTNP KLTGHA EKL FGLVRDSAGQLKASGTVVIDAA
LGSIIHAQKAITDPQFVVVKEALLKTIKAAVGDKWSDELSSA EVAYDELA AAIKKA----- |(Soy)V00454.1
MGAFTEKQDALVSSSF EAFKANIPQYSVVFYTSILEKAPAAKDLFSFLANGVDPTNP KLTGHA EKL FGLVRDSAGQLKASGTVVIDAA
LGSIIHAQKAITDPQFVVVKEALLKTIKAAVGDKWSDELSSA EVAYDELA AAIKKA----- |(Soy)CAA23732.1
MGAFTEKQDALVSSSF EAFKANIPQYSVVFYTSILEKAPAAKDLFSFLANGVDPTNP KLTGHA EKL FGLVRDSAGQLKASGTVVIDAA
LGSIIHAQKAITDPQFVVVKEALLKTIKAAVGDKWSDELSSA EVAYDELA AAIKKA----- |(Soy)sp|P02237.2|LGB3_S
OYBN

```

## Myoglobin data

```

In [17]: myo_data_aligned = AlignIO.read(open("Alignment\myo_data_aligned.clustal"), "clustal")
for entry in myo_data_aligned:
    for fasta in myo_data:
        if fasta.name.split(" ")[0] == entry.id:
            print(entry.seq+" |"+fasta.animal)
            break
# Very clean alignment (mostly mismatches, less indels)

```

```

MVLSAADKGNVKAAGKVGGHAAEYGAELERMFLSFPTTKTYFPHFDL-----SHGSAQVKGHGAKVAAALTKAVEHLDLPGALS
ELSDLHAHKLRLVDPVNFKLLSHSLVLTSLASHLPDSFTPAVHASLDKFLANVSTVLT SKYR----- |Cow
----MANYDMVLQCWEPVEADYNHGGVLVSR LFAEHPETLTLFPKFAGIAAG-DLSGNAAVA AHGATVLRKL GELLNARGDHAATLK
SLATTHANKHKIPLKNFTLITNICKVMGEKAGL--DEAGQEALRQVMGVIIADINVTYME LGFAG |Salmon
MGLSDQEWQQVLTIWGKVEADIAGHGHEVLMRLFHDHPETLDRFDKFKGLKTPDQMKGSEDLKKHGATVLTQLGKILKQKGNHESELK
PLAQTHATKHKIPVKYLEFISEVIIKVIAEKHAADFGADSQAMKKALELFRNDMASKYKEFGFQG |Chicken
MGLSDQEWQQVLTIWGKVEADIAGHGHEVLMRLFHDHPETLDRFDKFKGLKTPDQMKGSEDLKKHGATVLTQLGKILKQKGNHESELK

```

[illegible]

## All data

```
In [18]: combined_data_aligned = AlignIO.read(open("Alignment\combined_data_aligned.clustal"), "
i=0
for entry in combined_data_aligned:
    is_leg = True
    for fasta in myo_data:
        if fasta.name.split(" ")[0] == entry.id:
            print(entry.seq+" |"+fasta.animal)
            is_leg = False
            i+=1
            break
    if is_leg:
        print(entry.seq+" |(Soy)" +entry.id)
# There is a very noticable segregation between Leghemoglobin and hemoglobin
```

MGAFTEKQEALVNSSFEAFKANLPHHSVFFNSILEKAPAAKNMFSFLGDAVDP----KNPKLAGHAEKLFGLVRDSAVQLQTKGLVV  
A-DATLGPHTQKGV-TDLQFAVVEKALLKTIKEAVGDKWSEELSNPWE----VAYDEIAAAIKKAMAIGSLV | (Soy)V00451.1  
MGAFTEKQEALVNSSFEAFKANLPHHSVFFNSILEKAPAAKNMFSFLGDAVDP----KNPKLAGHAEKLFGLVRDSAVQLQTKGLVV

A-DATLGPIHTQKGV-TDLQFAVWKEALLKTIKEAVGDKWSEELSNAWE----VAYDEIAAAIKKAMAIGSLV |(Soy)KHN3787  
2.1  
-GAFTEKQDALVSSSF EAFKANIPQYSVVFYNSILEKAPAAKDLFSFLANGVDP----TNPKLTGHA EKLFALVRDSAGQLKASGTVV  
A-DAALGSIHAQKAV-TNPEFV-VKEALLKTIKEAVGDKWSELSNAWE----VAYDEIAAAIKKAF----- |(Soy)pir|GPS  
YC2  
-VAFTEKQDALVSSSF EAFKANIPQYSVVFYTSILEKAPAAKDLFSFLANGVDP----TNPKLTGHA EKLFALVRDSAGQLKASGTVV  
A-DAALGSVHAQKAV-TDPQFVVVKEALLKTIKAAVGD KWSDEL SRAWE----VAYDEIAAAIKKA----- |(Soy)pdb|1FSL  
|A  
-VAFTEKQDALVSSSF EAFKANIPQYSVVFYTSILEKAPAAKDLFSFLANGVDP----TNPKLTGHA EKLFALVRDSAGQLKASGTVV  
A-DAALGSVHAQKAV-TDPQFVVVKEALLKTIKAAVGD KWSDEL SRAWE----VAYDEIAAAIKKA----- |(Soy)pdb|1FSL  
|B  
MVAFTEKQDALVSSSF EAFKANIPQYSVVFYTSILEKAPAAKDLFSFLANGVDP----TNPKLTGHA EKLFALVRDSAGQLKASGTVV  
A-DAALGSVHAQKAV-TDPQFVVVKEALLKTIKAAVGD KWSDEL SRAWE----VAYDEIAAAIKKA----- |(Soy)V00453.1  
MVAFTEKQDALVSSSF EAFKANIPQYSVVFYTSILEKAPAAKDLFSFLANGVDP----TNPKLTGHA EKLFALVRDSAGQLKASGTVV  
A-DAALGSVHAQKAV-TDPQFVVVKEALLKTIKAAVGD KWSDEL SRAWE----VAYDEIAAAIKKA----- |(Soy)NP\_00123  
5928.1  
MVAFTEKQDALVSSSF EAFKANIPQYSVVFYTSILEKAPAAKDLFSFLANGVDP----TNPKLTGHA EKLFALVRDSAGQLKASGTVV  
A-DAALGSVHAQKAV-TDPQFVVVKEALLKTIKAAVGD KWSDEL SRAWE----VAYDEIAAAIKKA----- |(Soy)KHN3787  
0.1  
MVAFTEKQDALVSSSF EAFKANIPQYSVVFYTSILEKAPAAKDLFSFLANGVDP----TNPKLTGHA EKLFALVRDSAGQLKASGTVV  
A-DAALGSVHAQKAV-TDPQFVVVKEALLKTIKAAVGD KWSDEL SRAWE----VAYDEIAAAIKKA----- |(Soy)sp|P0223  
8|LGBA\_SOYBN  
MVAFTEKQDALVSSSF EAFKANIPQYSVVFYTSILEKAPAAKDLFSFLANGVDP----TNPKLTGHA EKLFALVRDSAGQLKASGTVV  
A-DAALGSVHAQKAV-TDPQFVVVKEALLKTIKAAVGD KWSDEL SRAWE----VAYDEIAAAIKKA----- |(Soy)CAA2373  
1.1  
MGAFTEKQDALVSSSF EAFKANIPQYSVVFYNSILEKAPAAKDLFSFLANGVDP----TNPKLTGHA EKLFALVRDSAGQLKNGTGVV  
A-DAALVSIHAQKAV-TDPQFVVVKEALLKTIKEAVGGNWSDEL SNAWE----VAYDEIAAAIKKA----- |(Soy)V00452.1  
MGAFTEKQDALVSSSF EAFKANIPQYSVVFYNSILEKAPAAKDLFSFLANGVDP----TNPKLTGHA EKLFALVRDSAGQLKNGTGVV  
A-DAALVSIHAQKAV-TDPQFVVVKEALLKTIKEAVGGNWSDEL SNAWE----VAYDEIAAAIKKA----- |(Soy)NP\_00134  
5001.1  
MGAFTEKQDALVSSSF EAFKANIPQYSVVFYNSILEKAPAAKDLFSFLANGVDP----TNPKLTGHA EKLFALVRDSAGQLKNGTGVV  
A-DAALVSIHAQKAV-TDPQFVVVKEALLKTIKEAVGGNWSDEL SNAWE----VAYDEIAAAIKKA----- |(Soy)sp|P0223  
5|LGB1\_SOYBN  
MGAFTEKQDALVSSSF EAFKANIPQYSVVFYTSILEKAPAAKDLFSFLANGVDP----TNPKLTGHA EKLFALVRDSAGQLKNGTGVV  
A-DAALVSIHAQKAV-TDPQFVVVKEALLKTIKEAVGGNWSDEL SNAWE----VAYDEIAAAIKKA----- |(Soy)KHN3787  
1.1  
MGAFTEKQDALVSSSF EAFKANIPQYSVVFYTSILEKAPAAKDLFSFLANGVDP----SNPKLTGHA EKLFGLVRDSAGQLKANGTVV  
A-DAALGSIHAQKAI-TDPQFVVVKEALLKTIKEAVGDKWSELSNAWE----VAYDEIAAAIKKAF----- |(Soy)NM\_00124  
8319.3  
MGAFTEKQDALVSSSF EAFKANIPQYSVVFYTSILEKAPAAKDLFSFLANGVDP----SNPKLTGHA EKLFGLVRDSAGQLKANGTVV  
A-DAALGSIHAQKAI-TDPQFVVVKEALLKTIKEAVGDKWSELSNAWE----VAYDEIAAAIKKAF----- |(Soy)NP\_00123  
5248.2  
MGAFTEKQDALVSSSF EAFKANIPQYSVVFYTSILEKAPAAKDLFSFLANGVDP----SNPKLTGHA EKLFGLVRDSAGQLKANGTVV  
A-DAALGSIHAQKAI-TDPQFVVVKEALLKTIKEAVGDKWSELSNAWE----VAYDEIAAAIKKAF----- |(Soy)KHN0094  
1.1  
MGAFTEKQDALVSSSF EAFKANIPQYSVVFYTSILEKAPAAKDLFSFLANGVDP----SNPKLTGHA EKLFGLVRDSAGQLKANGTVV  
A-DAALGSIHAQKAI-TDPQFVVVKEALLKTIKEAVGDKWSELSNAWE----VAYDEIAAAIKKAF----- |(Soy)AAA3398  
0.1  
MGAFTEKQDALVSSSF EAFKANIPQYSVVFYTSILEKAPAAKDLFSFLANGVDP----SNPKLTGHA EKLFGLVRDSAGQLKANGTVV  
A-DAALGSIHAQKAI-TDPQFVVVKEALLKTIKEAVGDKWSELSNAWE----VAYDEIAAAIKKAF----- |(Soy)sp|P0223  
6|LGB2\_SOYBN  
MGAFTEKQDALVSSSF EAFKANIPQYSVVFYTSILEKAPAAKDLFSFLANGVDP----SNPKLTGHA EKLFGLVRDSAGQLKANGTVV  
A-DAALGSIHAQKAI-TDPQFVVVKEALLKTIKEAVGDKWSELSNAWE----VAYDEIAAAIKKAF----- |(Soy)sp|P0223  
6.2|LGB2\_SOYBN  
MGAFTEKQDALVSSSF EAFKANIPQYSVVFYTSILEKAPAAKDLFSFLANGVDP----SNPKLTGHA EKLFGLVRDSAGQLKANGTVV  
A-DAALGSIHAQKAI-TDPQFVVVKEALLKTIKEAVGDKWSELSNAWE----VAYDEIAAAIKKAF----- |(Soy)J01301.1  
MGAFTEKQDALVSSSF EAFKANIPQYSVVFYTSILEKAPAAKDLFSFLANGVDP----TNPKLTGHA EKLFGLVRDSAGQLKASGTVV  
I-DAALGSIHQKAI-TDPQFVVVKEALLKTIKEAVGDKWSELSNAWE----VAYDEIAAAIKKAF----- |(Soy)NP\_00123  
5423.1  
MGAFTEKQDALVSSSF EAFKANIPQYSVVFYTSILEKAPAAKDLFSFLANGVDP----TNPKLTGHA EKLFGLVRDSAGQLKASGTVV  
I-DAALGSIHAQKAI-TDPQFVVVKEALLKTIKEAVGDKWSELSNAWE----VAYDEIAAAIKKA----- |(Soy)V00454.1  
MGAFTEKQDALVSSSF EAFKANIPQYSVVFYTSILEKAPAAKDLFSFLANGVDP----TNPKLTGHA EKLFGLVRDSAGQLKASGTVV  
I-DAALGSIHAQKAI-TDPQFVVVKEALLKTIKEAVGDKWSELSNAWE----VAYDEIAAAIKKAF----- |(Soy)CAA2373  
2.1  
MGAFTEKQDALVSSSF EAFKANIPQYSVVFYTSILEKAPAAKDLFSFLANGVDP----TNPKLTGHA EKLFGLVRDSAGQLKASGTVV



```

I-DAALGSIHAQKAI-TDPQFVVVKEALLKTIKEAVGDKWSDELSSAWE----VAYDELAIAIKKAF----- |(Soy)sp|P0223
7.2|LGB3_SOYBN
MVL-SAADKGNVKAAGWKGVGHAAEYGAELERMFLSFPTTKTYFPHFDL-----SHGSAQVKGHGAKVAAAL---TKAVEHLDDL
GALSELSDLHAHKLVRDPVNFKLLSHSLVTLASHLPSDFTPAVHASLDKFLANVSTVLT SKYR----- |Cow
-----MANYDMVLQCWEPVEADYNNHGGVLVSLFAEHPELTLFPKFAGIAAG-DLSGNAAVAAGHATVLRKL---GELLNARGDHA
ATLKS LATTHANKHKIPLKNFTLITNIICKVMGEKAGL--DEAGQEALRQVMGVIIADINVTYME LGFAG--- |Salmon
MGL-SDQEWQVLTIIWGKVEADIAGHGHEVLMRLFHDHPETLDRFDKFKGLKTPDQMKGSEDLKKHGATVLTQL---GKILKQKGNHE
SELKPLAQTHATKHKIPVKYLEFISEVIIKVIKHAADFAGDSQAAMKKALELFRNDMASKYKEFGFQG--- |Chicken
MGL-SDQEWQVLTIIWGKVEADIAGHGHEVLMRLFHDHPETLDRFDKFKGLKTPDQMKGSEDLKKHGATVLTQL---GKILKQKGNHE
SELKPLAQTHATKHKIPVKYLEFISEVIIKVIKHAADFAGDSQAAMKKALELFRNDMASKYKEFGFQG--- |Chicken
MGL-SDQEWQVLTIIWGKVEADIAGHGHEVLMRLFHDHPETLDRFDKFKGLKTPDQMKGSEDLKKHGATVLTQL---GKILKQKGNHE
SELKPLAQTHATKHKIPVKYLEFISEVIIKVIKHAADFAGDSQAAMKKALELFRNDMASKYKEFGFQG--- |Chicken
MGL-SDQEWQVLTIIWGKVEADIAGHGHEVLMRLFHDHPETLDRFDKFKGLKTPDQMKGSEDLKKHGATVLTQL---GKILKQKGNHE
SELKPLAQTHATKHKIPVKYLEFISEVIIKVIKHAADFAGDSQAAMKKALELFRNDMASKYKEFGFQG--- |Turkey
MGL-SDQEWQVLTIIWGKVEADIAGHGHEVLMRLFHDHPETLDRFDKFKGLKTPDQMKGSEDLKKHGATVLTQL---GKILKQKGNHE
SELKPLAQTHATKHKIPVKYLEFISEVIIKVIKHAADFAGDSQAAMKKALELFRNDMASKYKEFGFQG--- |Turkey
MGL-SDQEWQVLTIIWGKVEADLAGHGHA VLMRLFQDHPETLDRFEKFKGLKTPDQMKGSEDLKKHGATVLTQL---GKILKQKGNHE
AELKPLAQTHATKHKIPVKYLEFISEVIIKVIKHAADFAGDSQAAMKKALELFRNDMASKYKEFGFQG--- |Duck
MGL-SDGEWQLVLNAWGKVEADVAGHGQEV LIRLFTGHPETLEKFDKFKHLKTEAEMKASEDLKKHGNTVLTAL---GGILEKKKGHHE
AEVKHLAESHANKHKIPVKYLEFISDAIIHVLHAKHPSDFGADAQAAMSKALELFRNDMAAQYKVLGFQG--- |Sheep
MGL-SDGEWQLVLNAWGKVEADVAGHGQEV LIRLFTGHPETLEKFDKFKHLKTEAEMKASEDLKKHGNTVLTAL---GGILKKKGHHE
AEVKHLAESHANKHKIPVKYLEFISDAIIHVLHAKHPSDFGADAQAAMSKALELFRNDMAAQYKVLGFQG--- |Goat
MGL-SDGEWQLVLNAWGKVEADVAGHGQEV LIRLFTGHPETLEKFDKFKHLKTEAEMKASEDLKKHGNTVLTAL---GGILKKKGHHE
AEVKHLAESHANKHKIPVKYLEFISDAIIHVLHAKHPSDFGADAQAAMSKALELFRNDMAAQYKVLGFHG--- |Cow
MGL-SDGEWQLVLNAWGKVEADVAGHGQEV LIRLFTGHPETLEKFDKFKHLKTEAEMKASEDLKKHGNTVLTAL---GGILKKKGHHE
AEVKHLAESHANKHKIPVKYLEFISDAIIHVLHAKHPSDFGADAQAAMSKALELFRNDMAAQYKVLGFHG--- |Cow
MGL-SDGEWQLVLNAWGKVEADVAGHGQEV LIRLFTGHPETLEKFDKFKHLKTEAEMKASEDLKKHGNTVLTAL---GGILKKKGHHE
AEVKHLAESHANKHKIPVKYLEFISDAIIHVLHAKHPSDFGADAQAAMSKALELFRNDMAAQYKVLGFHG--- |Cow
MGL-SDGEWQLVLNAWGKVEADVAGHGQEV LIRLFTGHPETLEKFDKFKHLKTEAEMKASEDLKKHGNTVLTAL---GGILKKKGHHE
AEVKHLAESHANKHKIPVKYLEFISDAIIHVLHAKHPSDFGADAQAAMSKALELFRNDMAAQYKVLGFHG--- |Cow
MGL-SDGEWQLVLNAWGKVEADVAGHGQEV LIRLFTGHPETLEKFDKFKHLKTEAEMKASEDLKKHGNTVLTAL---GGILKKKGHHE
AEVKHLAESHANKHKIPVKYLEFISDAIIHVLHAKHPSDFGADAQAAMSKALELFRNDMAAQYKVLGFHG--- |Cow
MGL-SDGEWQLVLNAWGKVEADVAGHGQEV LIRLFTGHPETLEKFDKFKHLKTEAEMKASEDLKKHGNTVLTAL---GGILKKKGHHE
AEVKHLAESHANKHKIPVKYLEFISDAIIHVLHAKHPSDFGADAQAAMSKALELFRNDMAAQYKVLGFHG--- |Cow
MGL-SDGEWQLVLNAWGKVEADVAGHGQEV LIRLFTGHPETLEKFDKFKHLKTEAEMKASEDLKKHGNTVLTAL---GGILKKKGHHE
AEVKHLAESHANKHKIPVKYLEFISDAIIHVLHAKHPSDFGADAQAAMSKALELFRNDMAAQYKVLGFHG--- |Cow
MGL-SDGEWQLVLNIWGKVETDLAGHGQEV LIRLFKNHPETLDFDKFKHLKTEDEMKGSEDLKKHGNTVLTAL---GGILKKKGHHE
AELKPLAQSHATKHKIPVKYLEFISDAIIQVLQSKHSGDFHADTEAAMKKALELFRNDIAAKYKELGFQG--- |Canis_lupus_f
amiliaris
MGL-SDGEWQLVLNVWGKVETDLAGHGQEV LISLFKGHPETLEKFEKFKHLKTEDEMKGSEDLKKHGSTVLTAL---GGILKKKGQHE
AELKPLAQSHATKHKIPVKYLEFISEAIIHVLQSKHPHDFGTDAQAAMRKALELFRNDIAAKYKELGFQG--- |Felis_catus
MGL-SDAEWQLVLNVWGKVEADLAGHGQEV LIRLFTHTPETLEKFDKFKHLKSEDEMKASEDLKKHGNTVLTAL---GAILKKKGHHE
AEIKPLAQSHATKHKIPVKYLEFISEAIIHVLHSHKHPGDFGADAQAAMSKALELFRNDIAAQYKELGFQG--- |Rabbit
MGL-SDGEWQLVLNVWGKVEADIPGHGQEV LIRLFKGHPETLEKFDKFKHLKSEDEMKASEDLKKHGATVLTAL---GGILKKKGHHE
AEIKPLAQSHATKHKIPVKYLEFISECIIQVLQSKHPGDFGADAQAAMNKALELFRKDMASNYKELGFQG--- |Human
MGL-SDGEWQLVLNVWGKVEADIPGHGQEV LIRLFKGHPETLEKFDKFKHLKSEDEMKASEDLKKHGATVLTAL---GGILKKKGHHE
AEIKPLAQSHATKHKIPVKYLEFISECIIQVLQSKHPGDFGADAQAAMNKALELFRKDMASNYKELGFQG--- |Human
MGL-SDGEWQLVLNVWGKVEADVAGHGQEV LIRLFKGHPETLEKFDKFKHLKSEDEMKASEDLKKHGNTVLTAL---GGILKKKGHHE
AELTPLAQSHATKHKIPVKYLEFISEAIIQVLQSKHPGDFGADAQAAMSKALELFRNDMAAKYKELGFQG--- |Pig
MGL-SDGEWQLVLNVWGKVEADVAGHGQEV LIRLFKGHPETLEKFDKFKHLKSEDEMKASEDLKKHGNTVLTAL---GGILKKKGHHE
AELTPLAQSHATKHKIPVKYLEFISEAIIQVLQSKHPGDFGADAQAAMSKALELFRNDMAAKYKELGFQG--- |Pig
MGL-SDGEWQLVLNVWGKVEADVAGHGQEV LIRLFKGHPETLEKFDKFKHLKSEDEMKASEDLKKHGNTVLTAL---GGILKKKGHHE
AELTPLAQSHATKHKIPVKYLEFISEAIIQVLQSKHPGDFGADAQAAMSKALELFRNDMAAKYKELGFQG--- |Pig

```

## Split data 80/20

### Leghemoglobin data

```

In [19]: leg_seqs = []
         for entry in leg_data_aligned:
             leg_seqs.append(str(entry.seq))

         leg_train = leg_seqs[:int(0.8*len(leg_seqs))]

```

```
leg_test = leg_seqs[int(0.8*len(leg_seqs)):]

#Unalign test set (luckily all the deletes are at the end)
for i, test in enumerate(leg_test):
    leg_test[i] = test[:test.index('- ')]
```

## Myoglobin data (Hand pick test values)

```
In [20]: myo_seqs = []
        for entry in myo_data_aligned:
            myo_seqs.append(str(entry.seq))

        selection = [5, 15, 2, 18, 24] # Cow, human, chicken, pig, turkey
        myo_train = np.delete(myo_seqs, selection).tolist()
        myo_test = np.array(myo_data)[selection]
        myo_test_labels = ["Cow", "Human", "Chicken", "Pig", "Turkey"]

        #Extract sequences out of FASTA node
        for i, test in enumerate(myo_test):
            myo_test[i] = test.sequence
```

## Myoglobin data (Hand pick only cow test sequences)

```
In [21]: cow_selection = range(4,12)
        cow_test = []
        for i in cow_selection:
            cow_test.append(myo_data[i].sequence)

        cow_selection = [0, 10, 11, 12, 13, 14, 15, 16]
        cow_aligned = np.array(myo_seqs)[cow_selection]
```

## HMM from scratch

### Initialize final variables

```
In [22]: E = 0.01
        ins_E = 0.01
        num_proteins = 20
```

### Base State

```
In [23]: class Base_State(object):
        def __init__(self):
            self.next = {} #{ "name": (probability, node_object) }
```

### Normal State

```
In [24]: class State(Base_State):
        def __init__(self):
            super().__init__()
            self.attributes = {
                "F": E, "L": E, "I": E, "M": E,
                "V": E, "A": E, "T": E, "P": E,
                "S": E, "D": E, "E": E, "N": E,
                "K": E, "Q": E, "H": E, "Y": E,
                "C": E, "W": E, "R": E, "G": E
            }
```

## Insert State

```
In [25]: class InsertState(Base_State):
        def __init__(self):
            super().__init__()
            self.next["insert"] = (ins_E, self)
```

## Delete State

```
In [26]: class DeleteState(Base_State):
        def __init__(self):
            super().__init__()
```

## HMM Class (Main)

```
In [27]: class HMM:
        def __init__(self):
            self.root = State()
            self.stop = State()

        # Train the model
        def fit(self, data):
            curr_state = self.root
            curr_del_state = None
            for i in range(len(data[0])):
                # Get all the chars in the column
                aas = {}
                for seq in data:
                    if seq[i] in aas.keys():
                        aas[seq[i]] += 1
                    else:
                        aas[seq[i]] = 1

                # Calculate the remaining probability to distribute among Normal State's At
                att_prob = 1
                not_included = num_proteins - len(aas.keys()) #Count # of aa not include
                has_gap = 0
                if '-' in aas.keys(): #Gap is NOT an aa
                    not_included += 1
                    has_gap = aas['-']
                att_prob -= not_included * E #aa not included error tak
                att_prob /= (len(data) - has_gap) #Divide among each # of se

                # Create states
                next_state = State()
                next_state_prob = 1 - ins_E #Probability of reaching the normal state
                next_delete = None
                has_delete = False
                for aa in aas:
                    if not aa == '-':
                        next_state.attributes[aa] = att_prob * aas[aa] #Db: calculated cor
                    else:
                        next_delete = DeleteState()
                        curr_state.next["delete"] = (1/len(data) * aas[aa], next_delete) #
                        next_state_prob -= 1/len(data) * aas[aa] # Upd
                    if curr_del_state is not None: # Lin
                        curr_del_state.next["delete"] = (1/len(data) * aas[aa], next_dele
                        curr_del_state.next["normal"] = (next_state_prob + ins_E, next_st
                        curr_del_state = next_delete
                has_delete = True
```

```

curr_state.next["normal"] = (next_state_prob, next_state)

# If state no delete state was created with an active delete state,
# update curr_del_state to None and link to new state.
if not has_delete and curr_del_state is not None:
    curr_del_state.next["normal"] = (next_state_prob+ins_E, next_state)
    curr_del_state = None

#Add insert state
ins_state = InsertState()
ins_state.next["normal"] = (1-ins_E, next_state)
curr_state.next["insert"] = (ins_E, ins_state)

#Iterate to next state
curr_state = next_state

# Link to stop node (no need to range(-1) in loop)
ins_state = InsertState()
ins_state.next["normal"] = (1-ins_E, self.stop)

curr_state.next["insert"] = (ins_E, ins_state)
curr_state.next["normal"] = (1-ins_E, self.stop)

if curr_del_state is not None:
    curr_del_state.next["normal"] = (1, self.stop)

# Get the Viterbi Score
def predict(self, sequence):
    # Initialize the table with initial probability (Rows: normal = 0, insert = 1,
    table = []
    for i in range(3):
        table.append([1/3])

    delete = False
    # Initialize states
    states = [None, None, None] #Again: normal = 0, insert = 1, delete = 2)
    states[0] = self.root #Normal
    states[1] = states[0].next["insert"][1]
    states[2] = None
    if "delete" in states[0].next.keys():
        states[2] = states[0].next["delete"][1]
        delete = True

    # Fill table
    reached_end = False
    for x in range(len(sequence)):
        next_state = None

        # If reached end of Linked List and insertion is needed
        if reached_end:
            table[0].append(table[0][x] * ins_E)
            table[1].append(table[0][x] * ins_E)
            table[2].append(table[0][x] * ins_E) # Not possible, but we need some
        else:
            next_state = states[0].next["normal"][1]

        # Calculate value for the normal state row
        temp = []
        for i in range(3):
            if i == 2 and (delete or states[i] == None): # In case delete sta
                break

```

```

        temp.append(table[i][x] * states[i].next["normal"][0] * next_state.
normal_value = max(temp)
table[0].append(normal_value)

# Calculate value for the insert state row
temp = []
for i in range(2): # An insert in a delete (delete->insert) is not po
    temp.append(table[i][x] * states[i].next["insert"][0])
insert_value = max(temp)
table[1].append(insert_value)

# Calculate value for the delete state row
# Also: A delete in an insert (insert->delete) is not possible: would j
temp = []
delete_value = table[0][x] * ins_E # Psuedo-delete state if delete
if "delete" in states[0].next: # If delete state exists
    temp.append(table[0][x] * states[0].next["delete"][0])

    if not states[2] == None and "delete" in states[2].next.keys() and
        temp.append(table[2][x] * states[2].next["delete"][0])
    else:
        temp.append(table[2][x] * ins_E) # Psuedo-delete state
        delete_value = max(temp)
table[2].append(delete_value)

#Iterate nodes
if reached_end or len(next_state.next) == 0: # Sequence is Longer than t
    reached_end = True
else:
    # Handle delete state. If there is a delete, has to skip an iteration b
    if "delete" in next_state.next.keys():
        states[2] = next_state.next["delete"][1]
        delete = True
    else:
        if delete:
            delete = False
        else:
            states[2] = None
    #Iterate normal state
    if isinstance(next_state, DeleteState):
        states[1] = None
    else:
        states[1] = next_state.next["insert"][1]
        states[0] = next_state

# Get Viterbi score
score = max(
    table[0][-1],
    table[1][-1],
    table[2][-1]
)

return score

```

## Train Model and Test

```

In [28]: leg_hmm = HMM()
         leg_hmm.fit(leg_train)

```

```
myo_hmm = HMM()  
myo_hmm.fit(myo_train)
```

## Test Leghemoglobin HMM

```
In [29]: for test in leg_test:  
         score = leg_hmm.predict(test)  
         print("Soy\t"+str(score))  
         # Base scores to sompare with below
```

```
Soy      5.30033873848049e-20  
Soy      5.614113567991492e-29  
Soy      7.457788463719895e-27  
Soy      4.101783655045943e-27  
Soy      4.101783655045943e-27
```

## Test Myohemoglobin HMM

```
In [30]: for test, label in zip(myo_test, myo_test_labels):  
         score = myo_hmm.predict(test)  
         print(label+"\t"+str(score))  
         # Base scores to sompare with below
```

```
Cow      5.746529244269801e-29  
Human    2.404579744105982e-35  
Chicken  7.613327645334508e-43  
Pig      4.477347112747736e-31  
Turkey  7.613327645334508e-43
```

## Use Leghemoglobin test set on Myohemoglobin HMM

```
In [31]: for test in leg_test:  
         score = myo_hmm.predict(test)  
         print("Soy\t"+str(score))  
         # Bad scores
```

```
Soy      4.4437712440684957e-262  
Soy      6.7600268408003205e-264  
Soy      6.76002684080032e-262  
Soy      6.7600268408003205e-264  
Soy      6.7600268408003205e-264
```

## Use Myohemoglobin test set on Leghemoglobin HMM

```
In [32]: for test, label in zip(myo_test, myo_test_labels):  
         score = leg_hmm.predict(test)  
         print(label+"\t"+str(score))  
         # Bad scores
```

```
Cow      2.829675374206272e-281  
Human    2.829675374206272e-281  
Chicken  2.187259222762203e-283  
Pig      2.2971304687806525e-280  
Turkey  2.187259222762203e-283
```

Extra analysis: Cow Myohemoglobin test (359 lines of code up to here)

## Use Cow only Myohemoglobin test set on Leghemoglobin HMM

```
In [33]: for test in cow_test:  
         score = leg_hmm.predict(test)
```

```
print("Cow\t"+str(score))
# Bad scores
```

```
Cow      5.929022975528739e-267
Cow      2.829675374206272e-281
Cow      2.829675374206272e-281
Cow      2.829675374206272e-281
Cow      2.829675374206272e-281
Cow      2.829675374206272e-281
Cow      2.829675374206272e-281
Cow      2.829675374206272e-281
```

## Train model with Cow sequences only and test with *all* Leghemoglobin sequences

```
In [34]: cow_hmm = HMM()
cow_hmm.fit(cow_aligned)
for entry in leg_data:
    score = cow_hmm.predict(entry.sequence)
    print("Soy\t"+str(score))
# Bad scores
```

```
Soy      8.891572360514127e-253
Soy      8.891572360514127e-253
Soy      1.2760140602112642e-266
Soy      1.7963806148048626e-266
Soy      1.8228772288732357e-265
Soy      1.276014060211264e-264
Soy      1.2790445936042654e-275
Soy      1.8228772288732357e-265
Soy      1.8228772288732357e-265
Soy      1.8228772288732357e-265
Soy      1.276014060211264e-264
Soy      1.276014060211264e-264
Soy      8.958808362748201e-253
Soy      1.2760140602112642e-266
Soy      1.2760140602112642e-266
Soy      1.2790445936042654e-275
Soy      1.7963806148048626e-268
Soy      1.7963806148048626e-268
Soy      1.8228772288732357e-265
Soy      1.2760140602112642e-266
Soy      1.276014060211264e-264
Soy      1.2760140602112642e-266
Soy      1.2760140602112642e-266
Soy      1.7963806148048626e-268
Soy      1.2760140602112642e-266
```

## Test *all* myhemoglobin sequences on Leghemoglobin model

```
In [35]: for entry in myo_data:
#Formating
    t = ""
    if len(entry.animal) < 12:
        t = "\t"
        if len(entry.animal) < 8:
            t += "\t"

#Calucation here
    score = leg_hmm.predict(entry.sequence)
    print(entry.animal+"\t"+t+str(score))
# Soy is most similar to Salmon and flawed Cow
```

```
Canis_lupus_familiaris 2.829675374206272e-281
```

Chicken	2.187259222762203e-283
Chicken	2.187259222762203e-283
Chicken	2.187259222762203e-283
Cow	5.929022975528739e-267
Cow	2.829675374206272e-281
Cow	2.829675374206272e-281
Cow	2.829675374206272e-281
Cow	2.829675374206272e-281
Cow	2.829675374206272e-281
Cow	2.829675374206272e-281
Cow	2.829675374206272e-281
Duck	1.753963170733009e-281
Felis_catus	2.829675374206272e-281
Goat	2.829675374206272e-281
Human	2.829675374206272e-281
Human	2.829675374206272e-281
Pig	2.2971304687806525e-280
Pig	2.2971304687806525e-280
Pig	2.2971304687806525e-280
Rabbit	3.8729782571052977e-283
Salmon	2.919629763048611e-269
Sheep	3.8729782571052977e-283
Turkey	2.187259222762203e-283
Turkey	2.187259222762203e-283

## Test *all* myhemoglobin sequences on Cow model

In [36]:

```

for entry in myo_data:
    #Formating
    t = ""
    if len(entry.animal) < 12:
        t = "\t"
        if len(entry.animal) < 8:
            t += "\t"

    #Calucation here
    score = cow_hmm.predict(entry.sequence)
    print(entry.animal+"\t"+t+str(score))
# Notice: one flawed Cow and Salmon

```

Canis_lupus_familiaris	2.0711225011059564e-64
Chicken	1.0356012715263339e-91
Chicken	1.0356012715263339e-91
Chicken	1.0356012715263339e-91
Cow	1.0774767495508747e-200
Cow	3.4790542839395505e-22
Cow	1.3916217135758214e-22
Cow	3.4790542839395505e-22
Cow	1.3916217135758214e-22
Cow	3.4790542839395505e-22
Cow	3.4790542839395505e-22
Cow	3.4790542839395505e-22
Cow	3.4790542839395505e-22
Duck	1.2726626584187439e-94
Felis_catus	1.4711700905980877e-62
Goat	1.6297455850862154e-29
Human	1.4901012337395633e-63
Human	1.4901012337395633e-63
Pig	2.1608344133386394e-52
Pig	2.1608344133386394e-52
Pig	2.1608344133386394e-52
Rabbit	3.365530858479962e-53
Salmon	8.929123017613468e-281
Sheep	1.425291884884441e-28



Turkey  
Turkey

1.0356012715263339e-91  
1.0356012715263339e-91

## Export Train/Test set to compare with HMM given from Lab (393 lines of code up to here)

### Leghemoglobin Training Set

```
In [37]: file = open("Export\export_leg_train.fa", "w")
for i, train in enumerate(leg_train):
    file.write(">Sequence (Soy) "+str(i+1)+"\n")
    file.write(train+"\n")
file.close()
```

### Leghemoglobin Test Set

```
In [38]: file = open("Export\export_leg_test.fa", "w")
for i, train in enumerate(leg_test):
    file.write(">Sequence (Soy) "+str(i+1)+"\n")
    file.write(train+"\n")
file.close()
```

### Myoglobin Test Set

```
In [39]: file = open("Export\export_myo_train.fa", "w")
for i, train in enumerate(myo_train):
    file.write(">Sequence "+str(i+1)+"\n")
    file.write(train+"\n")
file.close()
```

### Myoglobin Test Set

```
In [40]: file = open("Export\export_myo_test.fa", "w")
for i, (train, name) in enumerate(zip(myo_test, myo_test_labels)):
    file.write(">"+name+"\n")
    file.write(train+"\n")
file.close()
```

### Cow only Dataset

```
In [41]: file = open("Export\export_cow_test.fa", "w")
for i, train in enumerate(cow_test):
    file.write(">"+str(i)+"\n")
    file.write(train+"\n")
file.close()
```

In [ ]: