**Final Practical Report**

**Group number: 2**

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**Summary:** In this final assignment, 3 original scripts (and many more) were written to support the aim of locating genes and reconstructing phylogenic trees from 5 unknown genomes that were assigned to the group.

1. Summary of your five genomes incl families, species, chromosomes, genome length.

|  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- |
| **File name** | **Family** | **Species** | **Chromosomes** | **Genome length** | **Organism type** |
| 16.fa.txt | Planctomycetaceae | *R. baltica* SH1 | ? | 7149689 | Prokaryotic |
| 20.fa.txt | Thermotogaceae | *T. maritima* strain Tma100 | Complete Genome | 1869610 | Prokaryotic |
| 29.fa.txt | Saccharomycetaceae | *S.cerevisiae* S288C | IX | 439888 | Eukaryotic |
| 44.fa.txt | Leuconostocaceae | *L. gelidum* JB7 | Complete Genome | 1893500 | Prokaryotic |
| 47.fa.txt | Neisseriaceae | *N. meningitidis* alpha710 | Complete Genome | 2242948 | Prokaryotic |

Table 1. Summary of genomes.

2. The results of applying your three Python scripts to your five genomes:

1. The GC, nucleotide and dinucleotide frequencies for the five genomes & amino acid and diamino acid frequencies for your predicted proteins (output of your ORF finder) using your statistics tool.

The python script stat\_tool\_1.py was used to evaluate both GC, nucleotide, and dinucleotide frequencies for the 5 genomes.

|  |  |  |  |  |
| --- | --- | --- | --- | --- |
| genome: 16.fa.txt  GC-content: 0.5548  #A = 1589939/7149689  #C = 1981614/7149689  #T = 1592923/7149689  #G = 1985204/7149689  #N = 9/7149689  #AA = 359096/7149689  #AC = 421691/7149689  #AT = 387766/7149689  #AG = 329767/7149689  #CA = 489129/7149689  #CC = 388521/7149689  #CT = 330792/7149689  #CG = 704454/7149689  #TA = 105255/7149689  #TC = 543628/7149689  #TT = 359547/7149689  #TG = 493028/7149689  #GA = 544841/7149689  #GC = 559056/7149689  #GT = 423352/7149689  #GG = 389509/7149689 | genome: 20.fa.txt  GC-content: 0.4625  #A = 504109/1869610  #C = 426011/1869610  #T = 500833/1869610  #G = 438657/1869610  #N = 0/1869610  #AA = 119138/1869610  #AC = 100392/1869610  #AT = 112580/1869610  #AG = 129961/1869610  #CA = 110613/1869610  #CC = 80193/1869610  #CT = 127257/1869610  #CG = 92039/1869610  #TA = 67219/1869610  #TC = 160752/1869610  #TT = 117679/1869610  #TG = 113839/1869610  #GA = 165101/1869610  #GC = 68766/1869610  #GT = 101972/1869610  #GG = 85445/1869610 | genome: 29.fa.txt  GC-content: 0.3890  #A = 134339/439888  #C = 85465/439888  #T = 134423/439888  #G = 85661/439888  #N = 0/439888  #AA = 34423/439888  #AC = 23399/439888  #AT = 38074/439888  #AG = 26070/439888  #CA = 28585/439888  #CC = 14956/439888  #CT = 25992/439888  #CG = 13236/439888  #TA = 31581/439888  #TC = 27250/439888  #TT = 34410/439888  #TG = 28797/439888  #GA = 27377/439888  #GC = 17163/439888  #GT = 23563/439888  #GG = 14961/439888 | genome: 44.fa.txt  GC-content: 0.3668  #A = 600998/1893500  #C = 346142/1893500  #T = 598020/1893500  #G = 348339/1893500  #N = 0/1893500  #AA = 159766/1893500  #AC = 105514/1893500  #AT = 190601/1893500  #AG = 88325/1893500  #CA = 138243/1893500  #CC = 54062/1893500  #CT = 87166/1893500  #CG = 58970/1893500  #TA = 149211/1893500  #TC = 94947/1893500  #TT = 159111/1893500  #TG = 138835/1893500  #GA = 96986/1893500  #GC = 83918/1893500  #GT = 105227/1893500  #GG = 54283/1893500 | genome: 47.fa.txt  GC-content: 0.5169  #A = 540971/2242948  #C = 577262/2242948  #T = 542592/2242948  #G = 582122/2242948  #N = 0/2242948  #AA = 136187/2242948  #AC = 116880/2242948  #AT = 136676/2242948  #AG = 97600/2242948  #CA = 140876/2242948  #CC = 121349/2242948  #CT = 96573/2242948  #CG = 195774/2242948  #TA = 83819/2242948  #TC = 125051/2242948  #TT = 136887/2242948  #TG = 143111/2242948  #GA = 126462/2242948  #GC = 191292/2242948  #GT = 118731/2242948  #GG = 122793/2242948 |

Table 2. Nucleotide contents of genomes.

Diamino acid frequencies included as attached files, diamino\_out\_\*.txt.

* 1. Present the formulas you used for the GC content and frequency calculation.

The formula used for GC content was (seq.count(“C”) + seq.count(“G”))/len(seq). The formula used for frequency calculation was seq.count(nucl)/len(seq), with nucl being an iterative element for a defined set of nucleotides/ dinucleotides.

* 1. If applicable use charts for the results you obtained.

Fig. 1. Nucleotide frequencies of genomes.

Fig. 2. Dinucleotide frequencies of genomes.

1. The predicted ORFs in your five genomes by your ORF predictor (include some examples and some statistics, e.g.

i. Which basic assumption on ORFs did you apply?

The script orf\_finder.py was written to fulfil the conditions given below, with an example output file called out\_orf16.fasta.

The basic assumption of an Open Reading Frame (ORF), is that it is a stretch of DNA with a start codon on the 5’ end, and a stop codon on the 3’ end, which has the possibility to encode a protein coding sequence[[1]](#endnote-1). Therefore, our ORF finder searches for sequences flanked by a start codon and stop codon, across all 6 reading frames; 3 per strand, forward and reverse. We also assumed that prokaryotic ORFs would be in general longer than 300bp[[2]](#endnote-2), while eukaryotic ORFs are more complex due to splicing and therefore assigned a length between 100-1500bp.

The initial gene turnout was extremely high, as such an additional condition was included in the ORF finder script to remove overlapping genes; ie., a shorter ORF nested within a longer. This resulted in a more realistic number of genes, except for the eukaryotic genome. We attribute the inaccuracy of the method to the lack of accounting for splicing and introns within the eukaryotic genome.

The code was written with several nested for-loops. First, the script is verified to be accounting for three different reading frames. Each reading frame was scanned for an ATG and then, starting from there and ending at a stop-sequence, with the found ORF added to a list. By implementing a variable (t2), which is set to the iterating variable after appending an ORF, the overlaps can removed. The script will always take the longer sequence, e.g. the first ATG to the next stop-sequence. The same was done for the reverse strand, where the sequence was first translated into its complement and then mirrored. Then the same script was used as above, as it reads in the sequence in 5´→3´.

ii. How many ORFs per genome did you predict?

|  |  |  |
| --- | --- | --- |
| **File** | **Gene count from reference** | **Gene count from ORF finder** |
| 16.fa.txt | 7406 | 11,420 |
| 20.fa.txt | 5806 | 2,725 |
| 29.fa.txt | 232 | 3,507 |
| 44.fa.txt | 1967 | 1,739 |
| 47.fa.txt | 2104 | 3,847 |

Table 3. ORFs counted using ORF finder compared against reference (UniProt).

iii. Assess the accuracy of your ORF predictor by comparing its predictions to a reference. Specify the used reference, accuracy measure, criteria for defining true positives, false positives, false negatives, etc..

For accessing the performance, the F1-score was used. As a reference, the proteomes of the organisms were downloaded from Uniprot. For the query sequence, the ORFs of the organisms were translated with Biopython into amino acid sequences. Then, the true positives, false positives and the false negatives were calculated by comparing the protein-coding genes of the reference and query proteome. The comparisons were compiled into the following statistics.

|  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- |
|  |  | Predicted Class | | **F1-score** | 0.0418 |
|  | 16.fa.txt | P | N | **Len Pred** | 11420 |
| Actual Class | P | 391 | 6880 | **Len Ref** | 7271 |
| N | 11029 | NIL |  |  |

|  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- |
|  |  | Predicted Class | | **F1-score** | 0.4274 |
|  | 20.fa.txt | P | N | **Len Pred** | 2725 |
| Actual Class | P | 978 | 874 | **Len Ref** | 1852 |
| N | 1747 | NIL |  |  |

|  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- |
|  |  | Predicted Class | | **F1-score** | 0.2519 |
|  | 29.fa.txt | P | N | **Len Pred** | 730 |
| Actual Class | P | 119 | 96 | **Len Ref** | 215 |
| N | 611 | NIL |  |  |

|  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- |
|  |  | Predicted Class | | **F1-score** | 0.7202 |
|  | 44.fa.txt | P | N | **Len Pred** | 1739 |
| Actual Class | P | 1273 | 523 | **Len Ref** | 1795 |
| N | 466 | NIL |  |  |

|  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- |
|  |  | Predicted Class | | **F1-score** | 0.0899 |
|  | 47.fa.txt | P | N | **Len Pred** | 3847 |
| Actual Class | P | 263 | 1742 | **Len Ref** | 2001 |
| N | 3584 | NIL |  |  |

Tables 3 – 7. Analysis scores and F1 scores for metric comparisons.

iv. Which additional improvements would be possible to increase the accuracy of your ORF predictor?

The first improvement that comes to mind will involve improving the specificity of the ORF predictor, by utilising more information available in the literature involving transcription and translation. This could include requiring that ORFs have upstream promoter sequences like the Pribnow box (TATAAT)[[3]](#endnote-3), or Shine-Delgarno sequence (AGGAGG)[[4]](#endnote-4). We briefly tried implementing these specifications, which resulted in a drastic reduction in the number of ORFs found, much lower than the reference. The suspected cause is the script being overly specific with the upstream length that the promoter sequences should be located at. Given more time a script could have been developed to allow for a larger range of such upstream lengths.

Another commonly used method to increase the accuracy of the finder is the Hidden Markov Model (HMM) architecture, found in the hugely popular GENSCAN[[5]](#endnote-5), which overcomes the complexity of splicing in eukaryotic genomes.

1. The distance matrix for your genomes computed using your third script with the output of your first script.
   1. Which distance method did you use and why?

The distances matrices were derived using the script dist\_matr\_3.py, producing \*.grimm files as output.

A variety of distance based methods were applied, the first was based on the pure distance between the various GC percentage values of the gene sets, the second based on distances between amino acid frequencies, and the third based between diamino acid frequencies. We sought to include a variety of distance calculation methods to compare the impact it may have on resulting trees.

* 1. Construct and present a phylogenetic tree based on the calculated distance matrix. Which tree building method did you use and why?

The trees were constructed using Belvu, using the Scoredist distance correction method, resulting in a rooted tree.

* 1. Compare shortly this tree with the trees you created during the various practicals.

We include a tree produced using the distance calculated with dinucleotide frequencies, normalized by multiplying all values by 1000, to adjust for the initial extremely miniscule values.



Fig. 3. Phylogenic tree produced using dinucleotide frequencies.

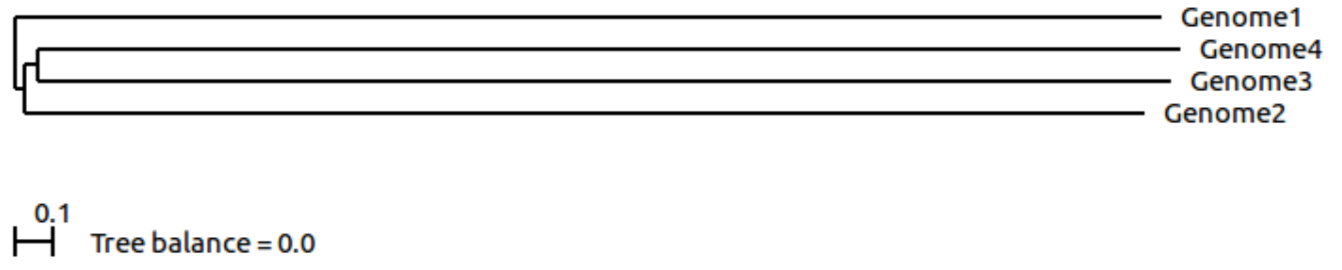


Fig. 5. Phylogenic tree produced using GRIMM-inferred distances.

Compared to the tree produced in Practical 5 with GRIMM, the basic branching structure is almost identical, except for the addition of genome 5, as we’ve included the eukaryotic genome (29.fa.txt) in this phylogenic reconstruction.

1. **References**

   Zvelebil, M., and Baum, J. (2008). Understanding bioinformatics (New York: Garland Science). [↑](#endnote-ref-1)
2. Burge, C. B. and Karlin, S. (1998) Finding the genes in genomic DNA. *Curr. Opin. Struct. Biol.* **8,** 346-354. [↑](#endnote-ref-2)
3. Pribnow, D. (1975). Nucleotide sequence of an RNA polymerase binding site at an early T7 promoter. Proceedings Of The National Academy Of Sciences *72*, 784-788. [↑](#endnote-ref-3)
4. Malys, N. Mol Biol Rep (2012) 39: 33. https://doi.org/10.1007/s11033-011-0707-4. [↑](#endnote-ref-4)
5. Burge, C. and Karlin, S. (1997) Prediction of complete gene structures in human genomic DNA. *J. Mol. Biol.***268,**78-94. [↑](#endnote-ref-5)