

# Lab 1 - Gr. 12 - Bioinformatics (732A93)

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# Assignment 1

## Question 1: Hardy–Weinberg equilibrium

### Question 1.1

We define the following probability space

$$(\Omega, \mathcal{F}, P)$$

Where,

$$\Omega = \{(A, A), (A, a), (a, A), (a, a)\}$$

We also define a probability measure  $P$  such as

$$\begin{aligned} P(X, Y) \\ X, Y \in \{A, a\} \end{aligned}$$

So  $P(X, Y)$  is the probability that an allele is  $(X, Y)$ .

By definition,  $p$  is the proportion of  $A$ 's in the allele population and  $q$  is the proportion of  $a$ 's in the allele population, so:

$$P(A) = P(A, A) + P(A, a) = p$$

The same applies to  $P(a)$ .

$$P(a) = P(a, a) + P(a, A) = q$$

The fact that we assume random mating means that  $X$  and  $Y$  are *IID*, which entails the following:

$$P(A, A) = P(A) * P(A) = p^2$$

$$P(a, a) = P(a) * P(a) = q^2$$

$$P(A, a) = P(a, A) = P(A) * P(a) = pq$$

The probability of an allele of it being a heterozygote is:

$$P(A, a) + P(a, A) = 2pq$$

$$P(\Omega) = P(A, A) + P(a, a) + 2P(A, a) = p^2 + q^2 + 2pq = 1$$

Thus, we show that by random mating the proportion of  $A$ 's and  $a$ 's is the same in the offsprings and the Hardy-Weinberg equilibrium is obtained and can't deviate from it.

## Question 1.2

We now look at the MN blood group, that has two possible co-dominating alleles,  $L^M$  (denoted  $M$ ) and  $L^N$  (denoted  $N$ ). In a population of  $n = 1000$  Americans of Caucasian descent, the following genotype counts were observed:

- $n_{MM} = 357$  individuals were  $MM$  homozygotes;
- $n_{MN} = 485$  individuals were  $MN$  heterozygotes;
- $n_{NN} = 158$  individuals were  $NN$  homozygotes;

The relative proportion, obtained by dividing the genotype counts by the total population, are 0.357, 0.485 and 0.158 respectively. According to the Hardy–Weinberg principle, we expect these quantities to follow the proportions  $p^2$ ,  $2pq$  and  $q^2$ , where  $p$  and  $q$  are the proportions of  $N$  and  $M$  in the alleles. In formulas:

$$\begin{aligned} p &= \frac{n_{MM}}{n} + \frac{1}{2} \cdot \frac{n_{MN}}{n} = 0.357 + \frac{1}{2} \cdot 0.485 = 0.5995 \\ q &= \frac{n_{NN}}{n} + \frac{1}{2} \cdot \frac{n_{MN}}{n} = 0.158 + \frac{1}{2} \cdot 0.485 = 0.4005 \end{aligned} \tag{1}$$

With these empirical quantities we formulate what would the population look if it was on equilibrium and compare them with the real proportions ( $p_{MM} = 0.375$ ,  $p_{NN} = 0.158$ ,  $p_{MN} = 0.485$ ):

$$\begin{aligned} p_{MM}^0 &= p^2 = 0.3594002 \\ p_{NN}^0 &= q^2 = 0.1604002 \\ p_{MN}^0 &= 2pq = 0.4801995 \end{aligned}$$

Performing a chi-square goodness of fit test (`chisq.test()` function in R) results in  $p\text{-value} = 1$ . This result shows that there is not enough statistical evidence to reject the hypothesis that the population is in a Hardy-Weinberg equilibrium.

```
##  
## Chi-squared test for given probabilities  
##  
## data: c(0.357, 0.485, 0.158)  
## X-squared = 9.9938e-05, df = 2, p-value = 1
```

## Assignment 2 : Exploring a genomic sequence

Note that, by convention, a *coding strand* is used when displaying a DNA sequence. “A coding strand, is the segment within double-stranded DNA that runs from 5’ to 3’, and which is complementary to the antisense strand of DNA, or template strand, which runs from 3’ to 5” ([https://en.wikipedia.org/wiki/Sense\\_strand](https://en.wikipedia.org/wiki/Sense_strand)).

### Question 2.1

We go to the following link <https://www.ncbi.nlm.nih.gov/nuccore/CU329670>, scroll down to “Features”, click on “CDS” to see the first 5662 nucleotides of the sequence.

The protein product is: **RecQ type DNA helicase**

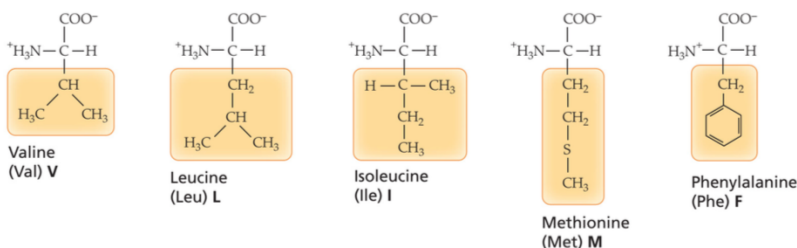
## Question 2.2

Note that proteins are made up of 20 different amino acids (linked together). Amino acids are molecules composed of an alpha carbon, a carboxyl group, an amino group and a side chain. The side chain is what makes an amino acid unique (see p. 5, Concepts in Bioinformatics and Genomics). The 20 different amino acids can be found in the picture below.

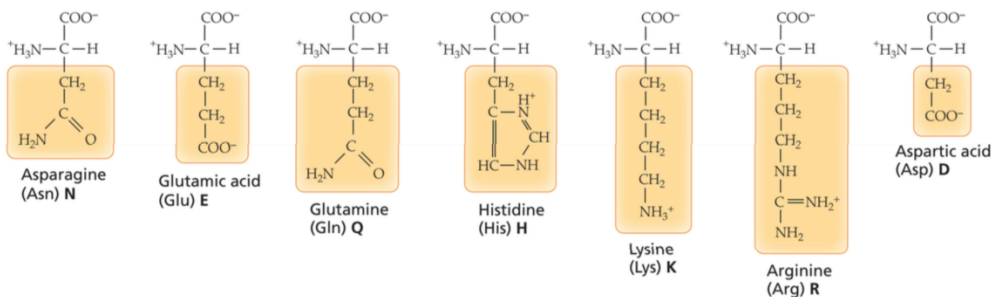
In our case, the first four amino acids are:

1. M (= **Methionine**, coded by ATG, representing the starting sequence)
2. V (= **Valine**)
3. V (= **Valine**)
4. A (= **Alanine**)

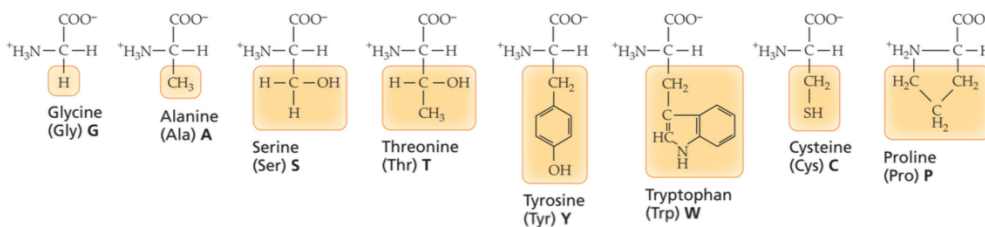
### Amino acids with hydrophobic side groups



### Amino acids with hydrophilic side groups



### Amino acids with side groups that are in between



## Question 2.3

Note that nucleotides code for amino acids. They are molecules composed of a sugar, a phosphate and a base. The part of nucleotides that distinguishes them is the base.

There are four possible bases: A (Adenine), C (Cytosine), G (Guanine), T (Thymine)

TABLE 1-1. SINGLE-LETTER ABBREVIATIONS USED FOR DNA NUCLEOTIDE SEQUENCES

ONE-LETTER ABBREVIATION	NUCLEOTIDE NAME	BASE NAME	CATEGORY
A	Adenosine monophosphate	Adenine	Purine
C	Cytidine monophosphate	Cytosine	Pyrimidine
G	Guanosine monophosphate	Guanine	Purine
T	Thymidine monophosphate	Thymine	Pyrimidine
N	Any nucleotide	Any base	NA
R	A or G	A or G	Purine
Y	C or T	C or T	Pyrimidine
— or *	—	—	Gap

Specific combinations of these nucleotides code for specific amino acids. In genetic code, three nucleotides (called codon) always code for one amino acid. E.g. ATG codes for met (Methionine) which is a starting point of a protein. Which codons code for which amino acids is shown in the overview below:

1st base	2nd base								3rd base
	T		C		A		G		
T	TTT	(Phe/F) Phenylalanine	TCT	(Ser/S) Serine	TAT	(Tyr/Y) Tyrosine	TGT	(Cys/C) Cysteine	T
	TTC		TCC		TAC		TGC		C
	TTA		TCA		TAA <sup>[B]</sup>	Stop (Ochre)	TGA <sup>[B]</sup>	Stop (Opal)	A
	TTG		TCG		TAG <sup>[B]</sup>	Stop (Amber)	TGG	(Trp/W) Tryptophan	G
C	CTT	(Leu/L) Leucine	CCT	(Pro/P) Proline	CAT	(His/H) Histidine	CGT	(Arg/R) Arginine	T
	CTC		CCC		CAC		CGC		C
	CTA		CCA		CAA		CGA		A
	CTG		CCG		CAG	(Gln/Q) Glutamine	CGG		G
A	ATT	(Ile/I) Isoleucine	ACT	(Thr/T) Threonine	AAT	(Asn/N) Asparagine	AGT	(Ser/S) Serine	T
	ATC		ACC		AAC		AGC		C
	ATA		ACA		AAA		AGA		A
	ATG <sup>[A]</sup>	(Met/M) Methionine	ACG		AAG	(Lys/K) Lysine	AGG	(Arg/R) Arginine	G
G	GTT	(Val/V) Valine	GCT	(Ala/A) Alanine	GAT	(Asp/D) Aspartic acid	GGT	(Gly/G) Glycine	T
	GTC		GCC		GAC		GGC		C
	GTA		GCA		GAA		GGA		A
	GTG		GCG		GAG	(Glu/E) Glutamic acid	GGG		G

*Saving the nucleotide sequence from GenBank:*

The complete nucleotide sequence of the coding strand from GenBank (<https://www.ncbi.nlm.nih.gov/nucleotide/329670.1?from=1&to=5662>) was saved in FASTA format as **2.3\_nucleotid-sequence.FASTA**. Note that only the last 12 characters AGCGACGACCAT actually correspond to the amino acids MVVA if the reverse complement is taken (see 2.4).

*Using backtrack to obtain the amino acid sequence from the protein sequence:*

We used backtrack to obtain the amino acid sequence corresponding to the protein sequence MVVA ([https://www.ebi.ac.uk/Tools/st/emboss\\_backtranseq/](https://www.ebi.ac.uk/Tools/st/emboss_backtranseq/)). After pasting MVVA and selecting “Schizosaccharomyces pombe (CAI equivalent)”, the following sequence is returned: **ATGGTCGTCGCT**

## Question 2.4

The above mentioned obtained coding strand sequence ATGGTCGTCGCT does not exist in the provided nucleotide sequence (that was saved in FASTA format as 2.3\_nucleotid-sequence.FASTA) since this sequence is not found when searching the file.

### Option 1

However, we can modify the displayed sequence to get what we are looking for. On GenBank, under “Display options”, one has to select both “Show sequence” and “**Show reverse complement**” (<https://www.ncbi.nlm.nih.gov/nuccore/CU329670.1?from=1&to=5662>). Afterwards, the displayed nucleotid sequence under “ORIGIN” starts with ATGGTCGTCGCT which codes for our amino acids MVVA.

### Option 2

Alternatively, one can take the last 12 characters from the original nucleotid sequence (the one in the FASTA file, i.e. the sequence before selecting “Show reverse complement” on GenBank). These characters are: AGCGACGACCAT. Copy this string and paste it here: <http://arep.med.harvard.edu/labgc/adnan/projects/Utilities/revcomp.html> Then click “**reverse complement**”. The result is again the sequence that we are looking for: ATGGTCGTCGCT which codes for our amino acids MVVA.

### Conclusion

**ATGGTCGTCGCT is exactly what we got in 2.3 using backtranseq** to obtain the amino acid sequence from the protein sequence MVVA. The only tricky thing here was that we needed to take the reverse complement of the correct characters from the nucleotid sequence (namely the last 12 characters).

## Question 2.5

*Number range that corresponds to these amino acids (protein sequence):*

In the saved FASTA file, the *last 12 characters* AGCGACGACCAT correspond to the amino acids MVVA (after taking the reverse complement). The number range corresponding to them is 5651 to 5662.

If we had selected “Show reverse complement” on GenBank before, then the *first 12 characters* ATGGTCGTCGCT would correspond to the amino acids MVVA. The number range would then be 1 to 12.

*Stop codon in the nucleotide sequence:*

Note that there are three possible stop codons: TAA, TAG, TGA. Still, it is not easy to identify the stop codon manually. However, one can use: <https://www.ncbi.nlm.nih.gov/orffinder> and paste the complete (reverse complement) nucleotide sequence in FASTA format. With the default parameters, this is automatically translated to the correct amino acid sequence. Also, after scrolling down a bit, one can see that the last stop happens at 5661. We find TC from 5661 to 5662 (and at 5662, we have the last nucleotid). From 5658 to 5660, we have TGA which is one of the above mentioned stop codons. We can conclude that the stop codon in the nucleotid sequence is hence TGA from 5658 to 5660 (looking at the reverse complement nucleotid sequence).

*Chromosome on which the genomic sequence lies:*

We can see on GenBank (in the header under the definition as well as in the features under source), that this genomic sequence lies on the chromosome 1 of schizosaccharomyces pombe.

## Assignment 3 : Exploring a genomic sequence

### Question 3.1

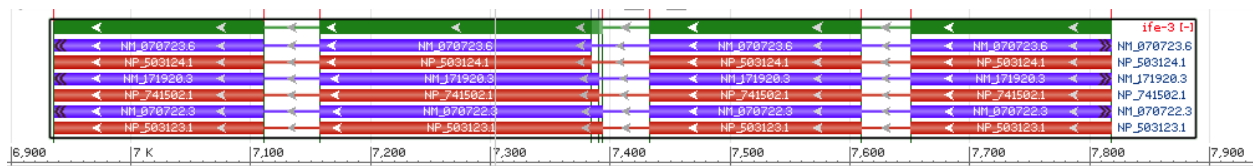
*Caenorhabditis elegans* is a free-living, nematode worm, which lives in soil environments all over the world. it is divided in two sexes:

- i) male
- ii) self-fertilizing hermaphrodite

Finally, it has all human sensations, such as taste and smell, despite the fact that it has no eyes. ([https://en.wikipedia.org/wiki/Caenorhabditis\\_elegans](https://en.wikipedia.org/wiki/Caenorhabditis_elegans))

In general C.elegans has a similar construction and characteristics as humans. It is also important that the genes, which are responsible for human's evolution, had similar ancestor with C.elegans. Therefore, instead of making experiments on humans, which most of the time is expensive and difficult, scientist can work with C.elegans. That is the reason why studying C.elegans biology is crucial for scientific field. ([http://www.people.ku.edu/~erikl/Lundquist\\_Lab/Why\\_study\\_C.\\_elegans.html](http://www.people.ku.edu/~erikl/Lundquist_Lab/Why_study_C._elegans.html))

### Question 3.2



This diagram shows exons and introns. We have 4 exons in our searching query. Our searching query found between 6529 and 8028. Our exons and introns:

- exon1: 6936 - 7110
- introns1: 7111 - 7157
- exon2: 7158 - 7393
- introns2: 7394 - 7432
- exon3: 7433 - 7609
- introns3: 7610 - 7650
- exon4: 7651 - 7818

### Question 3.3

We can find all the information needed from the summary :

Caenorhabditis elegans chromosome V

Sequence ID: [NC\\_003283.11](#) Length: 20924180 Number of Matches: 1

Range 1: 6529 to 8028 [GenBank](#) [Graphics](#)

▼ Next Match ▲ Previous Match

Score	Expect	Identities	Gaps	Strand
2771 bits(1500)	0.0	1500/1500(100%)	0/1500(0%)	Plus/Plus

The numbering of the sequences in the alignment is 1500. The identities section in the summary shows this result. In general, identities shows the number of identical bases between the query and the subject sequence.

The summary also contains information about the orientation both of the query and the subject sequence, which can be found in “Strand” section. In this case the the database genomic sequence progress has the same direction as the query sequence because both of them are plus and they are increasing.

Using <http://www.bioinformatics.nl/cgi-bin/emboss/revseq> , we reverse the query sequence, we run again the BLAST tool and we get the below summary :

#### Caenorhabditis elegans chromosome V

Sequence ID: [NC\\_003283.11](#) Length: 20924180 Number of Matches: 1

Range 1: 6529 to 8028 [GenBank](#) [Graphics](#)

▼ Next Match ▲ Previous Match

Score	Expect	Identities	Gaps	Strand
2771 bits(1500)	0.0	1500/1500(100%)	0/1500(0%)	Plus/Minus

It is clear from this summary that the reverse sequence has opposite direction from the subject sequence ( Plus / Minus in the “Strand” section ), while all the other information, such as Identities and Score, is the same.

### Question 3.4

Our query sequence is found at “*Caenorhabditis elegans chromosome V*” and its range is from 6528 to 8027. We can see chromosome name in the figure of question 3.3.

### Question 3.5

We will find sequences of exons by this code:

```
## gets sequence, starting index and ending index as a parameter
## split and return the sequence between start and index
get_exon = function(seq, start, end){
  org_seq_start = 6529
  fst_index = start - org_seq_start + 1
  return(substr(seq, fst_index, fst_index + (end-start)))
}

## read the query sequence
fasta_file = readLines("732A51_BioinformaticsHT2018_Lab01Ex03.fasta")
query_sequence = paste(fasta_file[-1], collapse = "")

## Starting and end indexes of exons.
exon1_start = 6936
exon1_end = 7110

exon2_start = 7158
exon2_end = 7393

exon3_start = 7433
exon3_end = 7609

exon4_start = 7651
exon4_end = 7818
```



```

## get exons with our split function
exon1 = get_exon(query_sequence, exon1_start, exon1_end)
exon2 = get_exon(query_sequence, exon2_start, exon2_end)
exon3 = get_exon(query_sequence, exon3_start, exon3_end)
exon4 = get_exon(query_sequence, exon4_start, exon4_end)

## save exons for protein analysis
writeLines(exon1, file("q3.5_exon1.txt"))
writeLines(exon2, file("q3.5_exon2.txt"))
writeLines(exon3, file("q3.5_exon3.txt"))
writeLines(exon4, file("q3.5_exon4.txt"))

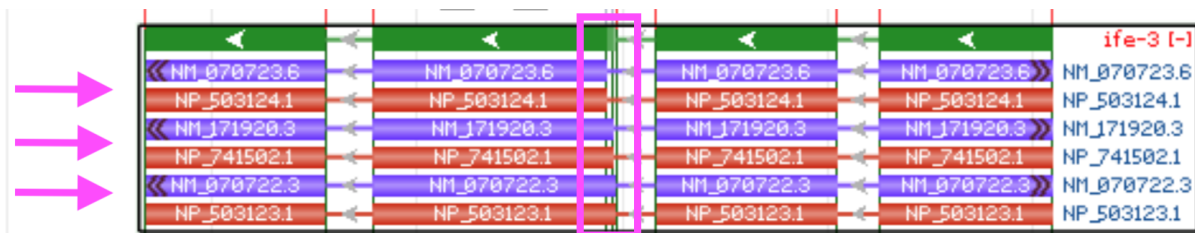
```

## Analysis

After getting whole DNA code from all exons, we can search it on blastx to find which protein it is and its protein sequence. We can also see the protein sequence in the screenshots of *Sequence Text View Tool* in blast nucleotide search for each exons.

### blastx overview of all exons

This is overview of all exons (result of pasting together the nucleotide strings of all the exons and using the resulting aggregated string for blastx). Note that there are 3 possible versions because there are 3 possible ranges for exon 2 (see following screenshot):



- NP\_503123.1

### Eukaryotic translation initiation factor 4E-3 [Caenorhabditis elegans]

Sequence ID: [NP\\_503123.1](#) Length: 251 Number of Matches: 1

► [See 2 more title\(s\)](#)

Range 1: 1 to 236 [GenPept](#) [Graphics](#)

▼ Next Match ▲ Previous Match

	Score	Expect	Method	Identities	Positives	Gaps	Frame
	489 bits(1260)	3e-178	Compositional matrix adjust.	236/236(100%)	236/236(100%)	0/236(0%)	-1
Query	756	MSTSVAENKALSASGDVNASDASVPPELLTRHPLQNRWALWYLKADRNKEWEDCLKMVSL				577	
Sbjct	1	MSTSVAENKALSASGDVNASDASVPPELLTRHPLQNRWALWYLKADRNKEWEDCLKMVSL				60	
Query	576	FDTVEDFWSLYNHIQSAGGLNWGSDYYLKFKEGIKPMWEDVNNVQGGRWLVVVVKQKLQRR				397	
Sbjct	61	FDTVEDFWSLYNHIQSAGGLNWGSDYYLKFKEGIKPMWEDVNNVQGGRWLVVVVKQKLQRR				120	
Query	396	TQLLDHYWLELLMAIVGEQFDEYGDYICGAVVNVVRQKGDVSLWTRDATRDDVNLRIQGV				217	
Sbjct	121	TQLLDHYWLELLMAIVGEQFDEYGDYICGAVVNVVRQKGDVSLWTRDATRDDVNLRIQGV				180	
Query	216	LKQKLSIPDTEILRYEVHKDSSARTSSTVKPRICLPAKDPAPVKEKGPAATTSPSN				49	
Sbjct	181	LKQKLSIPDTEILRYEVHKDSSARTSSTVKPRICLPAKDPAPVKEKGPAATTSPSN				236	

- NP\_741502.1

### Eukaryotic translation initiation factor 4E-3 [Caenorhabditis elegans]

Sequence ID: [NP\\_741502.1](#) Length: 250 Number of Matches: 1

► [See 1 more title\(s\)](#)

Range 1: 1 to 235 [GenPept](#) [Graphics](#)

▼ Next Match ▲ Previous Match

Score	Expect	Method	Identities	Positives	Gaps	Frame
483 bits(1243)	1e-175	Compositional matrix adjust.	235/236(99%)	235/236(99%)	1/236(0%)	-1
Query 756	MSTSVAENKALSASGDVNASDASVPPELLTRHPLQNRWALWYLKADRNKEWEDCLKMVSL					577
Sbjct 1	MSTSVAENKALSASGDVNASDASVPPELLTRHPLQNRWALWYLKADRNKEWEDCLKMVSL					60
Query 576	FDTVEDFWSLYNIQSAGGLNWGSDYYLFKEGIKPMWEDVNNVQGGRWLVVVDKQKLQRR					397
Sbjct 61	FDTVEDFWSLYNIQSAGGLNWGSDYYLFKEGIKPMWEDVNNVQGGRWLVVVDKQ LQRR					119
Query 396	TQLLDHYWLELLMAIVGEQFDEYGDYICGAVVNVVRQKGDVSLWTRDATRDDVNLRIQV					217
Sbjct 120	TQLLDHYWLELLMAIVGEQFDEYGDYICGAVVNVVRQKGDVSLWTRDATRDDVNLRIQV					179
Query 216	LKQKLSIPDTEILRYEVHKDSSARTSSTVKPRICLPAKDPAPVKEKGPAATTSPSN					49
Sbjct 180	LKQKLSIPDTEILRYEVHKDSSARTSSTVKPRICLPAKDPAPVKEKGPAATTSPSN					235

- NP\_503124.1

### Eukaryotic translation initiation factor 4E-3 [Caenorhabditis elegans]

Sequence ID: [NP\\_503124.1](#) Length: 248 Number of Matches: 1

► [See 1 more title\(s\)](#)

Range 1: 1 to 233 [GenPept](#) [Graphics](#)

▼ Next Match ▲ Previous Match

Score	Expect	Method	Identities	Positives	Gaps	Frame
478 bits(1231)	7e-174	Compositional matrix adjust.	233/236(99%)	233/236(98%)	3/236(1%)	-1
Query 756	MSTSVAENKALSASGDVNASDASVPPELLTRHPLQNRWALWYLKADRNKEWEDCLKMVSL					577
Sbjct 1	MSTSVAENKALSASGDVNASDASVPPELLTRHPLQNRWALWYLKADRNKEWEDCLKMVSL					60
Query 576	FDTVEDFWSLYNIQSAGGLNWGSDYYLFKEGIKPMWEDVNNVQGGRWLVVVDKQKLQRR					397
Sbjct 61	FDTVEDFWSLYNIQSAGGLNWGSDYYLFKEGIKPMWEDVNNVQGGRWLVVVDK QRR					117
Query 396	TQLLDHYWLELLMAIVGEQFDEYGDYICGAVVNVVRQKGDVSLWTRDATRDDVNLRIQV					217
Sbjct 118	TQLLDHYWLELLMAIVGEQFDEYGDYICGAVVNVVRQKGDVSLWTRDATRDDVNLRIQV					177
Query 216	LKQKLSIPDTEILRYEVHKDSSARTSSTVKPRICLPAKDPAPVKEKGPAATTSPSN					49
Sbjct 178	LKQKLSIPDTEILRYEVHKDSSARTSSTVKPRICLPAKDPAPVKEKGPAATTSPSN					233

As we can see from output of blastx, all exons are used for producing *Eukaryotic translation initiation factor 4E-3 [Caenorhabditis elegans]* protein by “NP\_503123.1”, “NP\_741502.1” and “NP\_503124.1” unique protein sequences.

In “NP\_503123.1” sequence, we use all exons as a whole. But we can see some missing amino acids in “NP\_741502.1” and “NP\_503124.1”. We can see easily at the figure of Question 3.2, that the lengths of the different exon2 alternatives differ. While “NP\_503123.1” uses the whole exon2, “NP\_741502.1” and “NP\_503124.1” do not use the whole exon2. We will see which amino acids are produced from which exon sequence in the continuation of this question.

After DNA code translation, blastx compared our protein sequence with a negative direction. While our query sequence order is decreasing, our subject sequence order is increasing.

exon 1

- `get_exon` function output:

```
## TTAAGGAGTTGGGGTGGCTGGAGAAGTTCCTGTAGCCTCCGTGCCGGGAT ...
```

- Sequence Text View Tool in blast nucleotide search

```
6841 CGAAAAACAGTTCATTTTCAAGACAACATTGAGACTGGGAGTACGGGGAAGCTCATTACGGTGAGAGGAATTGGTGAGATCTTTAGAATATGCTTAAG
6941 GAGTTGGGGTGGCTGGAGAAGTTCCTGTAGCCTCCGTGCCGGGATTCGATGGAGAAGTCGTTGCCGGCTGGTCCCTTTTCCTTCACTGGTGGCTGGATCCTT
7041 GGCTGGAAGACATATGCGTGGCTTGACAGTCGATGAGGTGCCGAGCCGACGAGTCCTTGTGAACCTCGTATCTGGAAATATTTACTTAGATAGCAAATAC
A P L C I R P K V T S S T R A S S D K H V E Y
```

- `blastx` - NP\_503123.1

### Eukaryotic translation initiation factor 4E-3 [Caenorhabditis elegans]

Sequence ID: [NP\\_503123.1](#) Length: 251 Number of Matches: 1

[▶ See 2 more title\(s\)](#)

Range 1: 195 to 236 [GenPept](#) [Graphics](#)

▼ Next Match ▲ Previous Match

Score	Expect	Method	Identities	Positives	Gaps	Frame
87.8 bits(216)	3e-23	Compositional matrix adjust.	42/42(100%)	42/42(100%)	0/42(0%)	-2
Query 174	YEVHKDSSARTSSTVKPRICLPKDPAPVKEKGPAATTSPSN	49				
	YEVHKDSSARTSSTVKPRICLPKDPAPVKEKGPAATTSPSN					
Sbjct 195	YEVHKDSSARTSSTVKPRICLPKDPAPVKEKGPAATTSPSN	236				

- `blastx` - NP\_741502.1

### Eukaryotic translation initiation factor 4E-3 [Caenorhabditis elegans]

Sequence ID: [NP\\_741502.1](#) Length: 250 Number of Matches: 1

[▶ See 1 more title\(s\)](#)

Range 1: 194 to 235 [GenPept](#) [Graphics](#)

▼ Next Match ▲ Previous Match

Score	Expect	Method	Identities	Positives	Gaps	Frame
87.8 bits(216)	3e-23	Compositional matrix adjust.	42/42(100%)	42/42(100%)	0/42(0%)	-2
Query 174	YEVHKDSSARTSSTVKPRICLPKDPAPVKEKGPAATTSPSN	49				
	YEVHKDSSARTSSTVKPRICLPKDPAPVKEKGPAATTSPSN					
Sbjct 194	YEVHKDSSARTSSTVKPRICLPKDPAPVKEKGPAATTSPSN	235				

- `blastx` - NP\_503124.1

### Eukaryotic translation initiation factor 4E-3 [Caenorhabditis elegans]

Sequence ID: [NP\\_503124.1](#) Length: 248 Number of Matches: 1

[▶ See 1 more title\(s\)](#)

Range 1: 192 to 233 [GenPept](#) [Graphics](#)

▼ Next Match ▲ Previous Match

Score	Expect	Method	Identities	Positives	Gaps	Frame
87.4 bits(215)	3e-23	Compositional matrix adjust.	42/42(100%)	42/42(100%)	0/42(0%)	-2
Query 174	YEVHKDSSARTSSTVKPRICLPKDPAPVKEKGPAATTSPSN	49				
	YEVHKDSSARTSSTVKPRICLPKDPAPVKEKGPAATTSPSN					
Sbjct 192	YEVHKDSSARTSSTVKPRICLPKDPAPVKEKGPAATTSPSN	233				

## exon 2

- `get_exon` function output:

```
## CTCAAAATCTCAGTATCCGGAATGCTCAATTTCTGCTTCAAAACCTGTCC ...
```

- Sequence Text View Tool in blast nucleotide search

```
TAAAAATTGTAATACTCTCAAAATCTCAGTATCCGGAATGCTCAATTTCTGCTTCAAAACCTGTCCGATGCGAAGATTGACATCATCGCGAGTAGCATC
R L I E T D P I S L K Q K L V Q G I R L N V D D R T A D
ACGAGTCCACAAGGAACCTTGTCACCCCTTTGACGAACATTACGACAGCTCCGCAGATGTAGTCTCCGTACTCGTCGAATTGCTCTCCAACAATAGCC
R T W L S V K D G K Q R V N V V A G C I Y D G Y E D F Q E G V I A
ATCAACAGCTCCAACAGTAGTCGAGCAATTGCGTTCTTCTCTGAAGCTTCTATGATTCATTGAATAAAATATATTTCTCAAAACGTACTTGCTTAT
M L L E L W Y H D L L Q T R R Q K D
```

- `blastx` - NP\_503123.1

### Eukaryotic translation initiation factor 4E-3 [Caenorhabditis elegans]

Sequence ID: [NP\\_503123.1](#) Length: 251 Number of Matches: 1

[▶ See 2 more title\(s\)](#)

Range 1: 116 to 193 [GenPept](#) [Graphics](#)

▼ Next Match ▲ Previous Match

Score	Expect	Method	Identities	Positives	Gaps	Frame
162 bits(410)	5e-52	Compositional matrix adjust.	78/78(100%)	78/78(100%)	0/78(0%)	-1
Query 236	KLQRRTQLLDHYWLELLMAIVGEQFDEYGDYICGAVVNVVRQKGDVSLWTRDATRDDVNL					57
Sbjct 116	KLQRRTQLLDHYWLELLMAIVGEQFDEYGDYICGAVVNVVRQKGDVSLWTRDATRDDVNL					175
Query 56	RIGQVLKQKLSIPDTEIL	3				
Sbjct 176	RIGQVLKQKLSIPDTEIL	193				

- `blastx` - NP\_741502.1

### Eukaryotic translation initiation factor 4E-3 [Caenorhabditis elegans]

Sequence ID: [NP\\_741502.1](#) Length: 250 Number of Matches: 1

[▶ See 1 more title\(s\)](#)

Range 1: 115 to 192 [GenPept](#) [Graphics](#)

▼ Next Match ▲ Previous Match

Score	Expect	Method	Identities	Positives	Gaps	Frame
161 bits(407)	2e-51	Compositional matrix adjust.	77/78(99%)	78/78(100%)	0/78(0%)	-1
Query 236	KLQRRTQLLDHYWLELLMAIVGEQFDEYGDYICGAVVNVVRQKGDVSLWTRDATRDDVNL					57
Sbjct 115	KLQRRTQLLDHYWLELLMAIVGEQFDEYGDYICGAVVNVVRQKGDVSLWTRDATRDDVNL					174
Query 56	RIGQVLKQKLSIPDTEIL	3				
Sbjct 175	RIGQVLKQKLSIPDTEIL	192				



- **blastx - NP\_503124.1**

### Eukaryotic translation initiation factor 4E-3 [Caenorhabditis elegans]

Sequence ID: [NP\\_503124.1](#) Length: 248 Number of Matches: 1

[▶ See 1 more title\(s\)](#)

Range 1: 115 to 190 [GenPept](#) [Graphics](#)

▼ Next Match ▲ Previous Match

Score	Expect	Method	Identities	Positives	Gaps	Frame
159 bits(402)	8e-51	Compositional matrix adjust.	76/76(100%)	76/76(100%)	0/76(0%)	-1
Query 230	QRRTQLLDHYWLELLMAIVGEQFDEYGDYICGAVVNVVRQKGDVSLWTRDATRDDVNLRI					51
Sbjct 115	QRRTQLLDHYWLELLMAIVGEQFDEYGDYICGAVVNVVRQKGDVSLWTRDATRDDVNLRI					174
Query 50	GQVLKQKLSIPDTEIL 3					
Sbjct 175	GQVLKQKLSIPDTEIL 190					

As we can see in the figure of **blastx - NP\_741502.1**, there is one unmatched amino acid. This means that if we want to produce the factor 4E-3 protein with the NP\_741502.1 sequence, we have to use an exon2 which is by 1 aminoacid (1 codons, 3 nucleotids) shorter.

Similarly, for **blastx - NP\_503124.1**, we found that 76 nucleotids out of 78 matched in exon2. If we want to produce factor 4E-3 protein by NP\_503124.1, we have to use an exon2 which is by 2 aminoacids (2 codons, 6 nucleotids) shorter.

#### exon 3

- **get\_exon** function output:

```
## TTGCTTATCGACAACAACCAACCAACGTCACCTTGAACGTTGTTGACGT ...
```

- **Sequence Text View Tool** in blast nucleotide search

```
7341 ATCAACAGCTCCAACAGTAGTGCAGCAATTGCGTTCTCTCTGAAGCTTCTATGATTCATTGAATAAAATATATTCTCAAAACGTAATTGCTTAT
M L L E L W Y H D L L Q T R R
7441 CGACAACAACCAACCAACGTCACCTTGACGTTGTTGACGTCCTCCACATTTGGCTTGATTCCTTCCTTGAACAAGTAATAATCGGATCCCCAGTTCAA
V V V L W R G G Q V N N V D E W M P K I G E K F L Y D S G W N L
7541 TCCTCCGGCAGAGTGAATGTGATTGTACAGCGACGAGAGTCCTCGACAGTGTGCGAAAGTGAACCATCTGGAAAAATCGATAAAGACGTATTTAAA
G G A S Q I H N Y L S W F D E V T D F L S V M
```

- **blastx - NP\_503123.1**

### Eukaryotic translation initiation factor 4E-3 [Caenorhabditis elegans]

Sequence ID: [NP\\_503123.1](#) Length: 251 Number of Matches: 1

[▶ See 2 more title\(s\)](#)

Range 1: 57 to 115 [GenPept](#) [Graphics](#)

▼ Next Match ▲ Previous Match

Score	Expect	Method	Identities	Positives	Gaps	Frame
128 bits(322)	5e-39	Compositional matrix adjust.	59/59(100%)	59/59(100%)	0/59(0%)	-1
Query 177	MVSLFDTVEDFWSLYNHIQSAGGLNWGSDYYLKFKEGIKPMWEDVNNVQGGRWLVVVDKQ					1
Sbjct 57	MVSLFDTVEDFWSLYNHIQSAGGLNWGSDYYLKFKEGIKPMWEDVNNVQGGRWLVVVDKQ					115

- blastx - NP\_741502.1

### Eukaryotic translation initiation factor 4E-3 [Caenorhabditis elegans]

Sequence ID: [NP\\_741502.1](#) Length: 250 Number of Matches: 1

[▶ See 1 more title\(s\)](#)

Range 1: 57 to 115 [GenPept](#) [Graphics](#)

▼ Next Match ▲ Previous Match

Score	Expect	Method	Identities	Positives	Gaps	Frame
128 bits(322)	5e-39	Compositional matrix adjust.	59/59(100%)	59/59(100%)	0/59(0%)	-1
Query 177	MVSLFDTVEDFWSLYNHIQSAGGLNWGSDYYLFKEGIKPMWEDVNNVQGGRWLVVVDKQ				1	
Sbjct 57	MVSLFDTVEDFWSLYNHIQSAGGLNWGSDYYLFKEGIKPMWEDVNNVQGGRWLVVVDKQ				115	

- blastx - NP\_503124.1

### Eukaryotic translation initiation factor 4E-3 [Caenorhabditis elegans]

Sequence ID: [NP\\_503124.1](#) Length: 248 Number of Matches: 1

[▶ See 1 more title\(s\)](#)

Range 1: 57 to 115 [GenPept](#) [Graphics](#)

▼ Next Match ▲ Previous Match

Score	Expect	Method	Identities	Positives	Gaps	Frame
128 bits(321)	7e-39	Composition-based stats.	59/59(100%)	59/59(100%)	0/59(0%)	-1
Query 177	MVSLFDTVEDFWSLYNHIQSAGGLNWGSDYYLFKEGIKPMWEDVNNVQGGRWLVVVDKQ				1	
Sbjct 57	MVSLFDTVEDFWSLYNHIQSAGGLNWGSDYYLFKEGIKPMWEDVNNVQGGRWLVVVDKQ				115	

exon 4

- get\_exon function output:

## CTCAGACAATCCTCCCATTCCTTGTTACGGTCAGCTTTCAAGTACCAGA ...

- Sequence Text View Tool in blast nucleotide search

```

7641 AATCTTCTA CTCAGACAATCCTCCCATTCCTTGTTACGGTCAGCTTTCAAGTACCAGAGCCAGCGATTCTGGAGGGGTGTCTGGTGAGAAGCTC
      K L C D E W E K N R D A K L Y W L A W R N Q L P H R T L L E
7741 TGGAGGAAGTGAAGCATCGGACGATTACATCGCCGGAAGCTGACAATGCTTTGTTTCCGCTACGGATGTGCTCAT TAGCTGAAAATAGGTAATATT
      P P V S A D S A N V D G S A S L A K N E A V S T S M
-----

```

- blastx - NP\_503123.1

### Eukaryotic translation initiation factor 4E-3 [Caenorhabditis elegans]

Sequence ID: [NP\\_503123.1](#) Length: 251 Number of Matches: 1

[▶ See 2 more title\(s\)](#)

Range 1: 1 to 56 [GenPept](#) [Graphics](#)

▼ Next Match ▲ Previous Match

Score	Expect	Method	Identities	Positives	Gaps	Frame
117 bits(293)	1e-34	Compositional matrix adjust.	56/56(100%)	56/56(100%)	0/56(0%)	-1
Query 168	MSTSVAENKALSASGDVNASDASVPPELLTRHPLQNRWALWYLKADRNKEWEDCLK				1	
Sbjct 1	MSTSVAENKALSASGDVNASDASVPPELLTRHPLQNRWALWYLKADRNKEWEDCLK				56	

- blastx - NP\_741502.1

### Eukaryotic translation initiation factor 4E-3 [Caenorhabditis elegans]

Sequence ID: [NP\\_741502.1](#) Length: 250 Number of Matches: 1

[▶ See 1 more title\(s\)](#)

Range 1: 1 to 56 [GenPept](#) [Graphics](#)

▼ Next Match ▲ Previous Match

Score	Expect	Method	Identities	Positives	Gaps	Frame
117 bits(292)	1e-34	Compositional matrix adjust.	56/56(100%)	56/56(100%)	0/56(0%)	-1
Query 168	MSTSVAENKALSASGDVNASDASVPPELLTRHPLQNRWALWYLKADRNKEWEDCLK					1
	MSTSVAENKALSASGDVNASDASVPPELLTRHPLQNRWALWYLKADRNKEWEDCLK					
Sbjct 1	MSTSVAENKALSASGDVNASDASVPPELLTRHPLQNRWALWYLKADRNKEWEDCLK					56

- blastx - NP\_503124.1

### Eukaryotic translation initiation factor 4E-3 [Caenorhabditis elegans]

Sequence ID: [NP\\_503124.1](#) Length: 248 Number of Matches: 1

[▶ See 1 more title\(s\)](#)

Range 1: 1 to 56 [GenPept](#) [Graphics](#)

▼ Next Match ▲ Previous Match

Score	Expect	Method	Identities	Positives	Gaps	Frame
117 bits(292)	1e-34	Compositional matrix adjust.	56/56(100%)	56/56(100%)	0/56(0%)	-1
Query 168	MSTSVAENKALSASGDVNASDASVPPELLTRHPLQNRWALWYLKADRNKEWEDCLK					1
	MSTSVAENKALSASGDVNASDASVPPELLTRHPLQNRWALWYLKADRNKEWEDCLK					
Sbjct 1	MSTSVAENKALSASGDVNASDASVPPELLTRHPLQNRWALWYLKADRNKEWEDCLK					56

## Question 3.6

From [https://www.wormbase.org/species/c\\_elegans/gene/WBGene00002061#0-3gi1-3](https://www.wormbase.org/species/c_elegans/gene/WBGene00002061#0-3gi1-3) and <https://www.ncbi.nlm.nih.gov/gene/178536> we can gain the below information:

First of all, the gene's name is ife-3 and it belongs to the gene class ife. It can be found in humans, monkeys, cows and other animals and insects. It is one of the five c. elegans homologs responsible for the protein eIF4E (Eukaryotic translation initiation factor 4E-3). Its ancestry is also related to Eukaryota, Metazoa, Ecdysozoa and other elegans. Moreover, except for c.elegans, which referred in previous parts of the exercise, BLASTP showed matches with some other species. For instance, the highest match (100%) is with c. brenneri and the lowest (73.7) with S. cerevisiae. Finally, from the gene's ontology perspective, ife - 3 participates in some molecular functions and biological functions such as RNA binding, protein metabolic process and gene expression.

## Appendix

```
knitr::opts_chunk$set(fig.width = 7, fig.height = 3, echo = FALSE,
                        warning = FALSE, message = FALSE)

library(dplyr)
library(tidyr)
library(magrittr)
library(kableExtra)

# -----
# Question 1.2
# -----

# The prob of one allele is the proportion of the homozygote + half the proportion of the heterozygote
p = 0.357 + 0.485 / 2
q = 0.158 + 0.485 / 2

# Expected values (Hardy-Weinberg equilibrium)
p_2 = p^2
q_2 = q^2
pq = 2*p*q

Xsq = chisq.test(c(0.357, 0.485, 0.158), p=c(p_2, pq, q_2))
Xsq
# Xsq$observed
# Xsq$expected

knitr::include_graphics("images/2.2_amino-acids.png")
knitr::include_graphics("images/2.3_nucleotides.png")
knitr::include_graphics("images/2.3_codons-aminoacids")
## gets sequence, starting index and ending index as a parameter
## split and return the sequence between start and index
get_exon = function(seq, start, end){
  org_seq_start = 6529
  fst_index = start - org_seq_start + 1
  return(substr(seq, fst_index, fst_index + (end-start)))
}

## read the query sequence
fasta_file = readLines("732A51_BioinformaticsHT2018_Lab01Ex03.fasta")
query_sequence = paste(fasta_file[-1], collapse = "")

## Starting and end indexes of exons.
exon1_start = 6936
exon1_end = 7110

exon2_start = 7158
exon2_end = 7393

exon3_start = 7433
exon3_end = 7609
```



```

exon4_start = 7651
exon4_end = 7818

## get exons with our split function
exon1 = get_exon(query_sequence, exon1_start, exon1_end)
exon2 = get_exon(query_sequence, exon2_start, exon2_end)
exon3 = get_exon(query_sequence, exon3_start, exon3_end)
exon4 = get_exon(query_sequence, exon4_start, exon4_end)

## save exons for protein analysis
writeLines(exon1, file("q3.5_exon1.txt"))
writeLines(exon2, file("q3.5_exon2.txt"))
writeLines(exon3, file("q3.5_exon3.txt"))
writeLines(exon4, file("q3.5_exon4.txt"))
cat(substr(exon1,1,50), "...")
cat(substr(exon2,1,50), "...")
cat(substr(exon3,1,50), "...")
cat(substr(exon4,1,50), "...")

```