**BIMM-143: INTRODUCTION TO BIOINFORMATICS**

The find-a-gene project assignment <https://bioboot.github.io/bimm143_S20/>

Dr. Barry Grant

**Overview:**

The find-a-gene project is a required assignment for BIMM-143. You should prepare a written report in **PDF** format that has responses to each question labeled **[Q1] - [Q10]** below. You may wish to consult the scoring rubric at the end of this document and the example report provided online.

The objective with this assignment is for you to demonstrate your grasp of database searching, sequence analysis, structure analysis and the R environment that we have covered in class.

**Due Date:**

Your responses to questions Q1-Q4 are due at the beginning of class **Tuesday May 5th** (05/05/20) at 12pm San Diego time. Note that these answers can be obtained very quickly (at best within 10 or 15 minutes), so if you don’t succeed at first, just keep trying.

The complete assignment, including responses to all questions, is due **Friday June 5th** (06/05/20) at 12pm San Diego time.

**Submission instructions:**

Your report formatted as a **PDF document** should be uploaded to ***GradeScope****.*  Please make sure to include your UCSD email and PID number on the first page.

**Be sure to include your UCSD email and PID number on the first page of your report.**

Submit your preliminary report with answers to Q1-Q4 as soon as you can so we can determine if you have found a novel gene. Submit this preliminary report as one document with screen shots of the results inserted appropriately.

See the demonstration report linked to on the course website for an example of format. I will email you my decision; proceed with subsequent questions only after we are sure you have found a novel gene.

For the final report add your results for Q5-Q10 to the preliminary report and submit the final document containing your results for all questions - **Please do not send only Q5-Q10 answers as the final report**.

**Questions:**

[**Q1**] Tell me the name of a protein you are interested in. Include the species and the accession number. This can be a human protein or a protein from any other species as long as it's function is known.

If you do not have a favorite protein, select human RBP4 or KIF11. Do not use beta globin as this is in the worked example report that I provide you with online.

# Name: neurexin-1 isoform alpha4 precursor

Accession: NP\_001317007

Organism: Homo sapiens

[**Q2**] Perform a BLAST search against a DNA database, such as a database consisting of genomic DNA or ESTs. The BLAST server can be at NCBI or elsewhere. Include details of the BLAST method used, database searched and any limits applied (e.g. Organism).

Method: TBLASTN (BLOSUM62) against crustaceans ESTs (expect threshold = 10)

Database: Expressed sequence tags (est)

Organism: Crustaceans (Taxid: 6657)

Also include the output of that BLAST search in your document. If appropriate, change the font to Courier size 10 so that the results are displayed neatly. You can also screen capture a BLAST output (e.g. alt print screen on a PC or on a MAC press ⌘-shift-4. The pointer becomes a bulls eye. Select the area you wish to capture and release. The image is saved as a file called Screen Shot [].png in your Desktop directory). It is **not** necessary to print out all of the blast results if there are many pages.

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On the BLAST results, clearly indicate a match that represents a protein sequence, encoded from some DNA sequence, that is homologous to your query protein. I need to be able to inspect the pairwise alignment you have selected, including the E value and score. It should be labeled a "genomic clone" or "mRNA sequence", etc. - but include no functional annotation.

Accession number:  [**CI998089.1**](https://www.ncbi.nlm.nih.gov/nucleotide/CI998089.1?report=genbank&log$=nuclalign&blast_rank=1&RID=KTGY0UDN013)

cDNA clone from immature Penaeus japonicus, 635 bp, mRNA sequence

Query: neurexin-1 isoform alpha4 precursor [Homo sapiens] Query ID: NP\_001317007.1 Length: 1507

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>CI998089 Marsupenaeus japonicus eyestalk immature Penaeus japonicus cDNA clone YAQ01A02NGRM0003\_C10 5', mRNA sequence

Sequence ID: CI998089.1 Length: 635

Range 1: 28 to 633

Score:188 bits(477), Expect:6e-53,

Method:Compositional matrix adjust.,

Identities:96/233(41%), Positives:138/233(59%), Gaps:32/233(13%)

Query 1056 VDLNGRLPDLI-SDALFCNGQIERGCEGPSTTCQEDSCSNQGVCLQQWDGFSCDCSMTSF 1114

+DLNG PD D + + GC+GPST C ++C+N G C+QQW+ +SC+C MTSF

Sbjct 28 LDLNGEAPDPAGKDVPLISSPVFPGCDGPSTKCHRNACANGGTCVQQWNSYSCNCDMTSF 207

Query 1115 SGPLCNDPGTTYIFSKGGGQITYKWPPNDRPSTRADRLAIGFSTVQKEAVLVRVDSSSGL 1174

+GP C+D T Y F G G +T+++P P TR D LA+GF T Q++AV++R+DS++

Sbjct 208 TGPTCSDESTAYEFGGGSGLMTFQYPEGRWPDTRRDLLALGFMTSQEDAVMLRLDSANS- 384

Query 1175 GDYLELHIHQGKIGVKFNVGTDDIAIEESNAIINDGKYHVVRFTRSGGNATLQVDSWPVI 1234

DY+EL I G I + +N+GT+D + E A +NDG YHVVRF RSG NAT+Q+D + V

Sbjct 385 NDYMELEIVDGNIFMVYNMGTEDHPVGEVMAKVNDGIYHVVRFVRSGSNATVQIDDYEVR 564

Query 1235 ERYPAGNNDNERLAIARQRIPYRLGRVVDEWLLDKGRQLTIFNSQATIIIGGK 1287

+ P KG QL++FN+Q+ + IGG+

Sbjct 565 SKNP------------------------------KGHQLSVFNAQSRLQIGGR 633

In general, [Q2] is the most difficult for students because it requires you to have a “feel” for how to interpret BLAST results. You need to distinguish between a perfect match to your query (i.e. a sequence that is not “novel”), a near match (something that might be “novel”, depending on the results of [Q4]), and a non-homologous result.

If you are having trouble finding a novel gene try restricting your search to an organism that is poorly annotated.

[**Q3**] Gather information about this “novel” **protein**. At a minimum, show me the protein sequence of the “novel” protein as displayed in your BLAST results from [Q2] as FASTA format (you can copy and paste the aligned sequence subject lines from your BLAST result page if necessary) or translate your novel DNA sequence using a tool called EMBOSS Transeq at the EBI. Don’t forget to translate all six reading frames; the ORF (open reading frame) is likely to be the longest sequence without a stop codon. It may not start with a methionine if you don’t have the complete coding region. Make sure the sequence you provide includes a header/subject line and is in traditional FASTA format.

>CI998089.1 CI998089 Marsupenaeus japonicus eyestalk immature Penaeus japonicus cDNA clone YAQ01A02NGRM0003\_C10 5', mRNA sequence

GAATTCCCGGGTCGACCCACGCGTCCGCTCGACTTGAACGGCGAAGCTCCCGACCCAGCAGGCAAAGACG

TTCCTCTGATCTCGAGCCCCGTCTTCCCAGGCTGTGACGGACCAAGTACAAAATGCCATCGGAATGCATG

TGCAAATGGTGGGACGTGCGTGCAACAGTGGAATTCCTATTCATGCAACTGTGACATGACCTCCTTTACC

GGACCTACGTGTTCTGATGAGTCCACGGCGTATGAGTTCGGCGGCGGCTCTGGCCTCATGACCTTCCAGT

ACCCTGAGGGCAGATGGCCTGATACCCGCCGTGACCTGTTGGCACTCGGTTTTATGACCTCGCAGGAGGA

TGCGGTCATGCTGAGGCTGGATTCGGCCAATTCGAATGATTACATGGAATTGGAAATCGTCGACGGCAAT

ATCTTCATGGTCTACAACATGGGCACGGAGGACCACCCCGTAGGCGAGGTCATGGCAAAGGTCAACGACG

GCATATATCACGTGGTTCGCTTCGTTAGGTCAGGTTCCAACGCCACGGTACAGATTGACGACTATGAAGT

TCGTTCAAAGAATCCAAAAGGACACCAGCTCTCCGTCTTCAACGCACAGTCACGCCTCCAGATCGGTGGA

CGAAG

>CI998089.1\_1 Marsupenaeus japonicus eyestalk immature Penaeus japonicus cDNA clone YAQ01A02NGRM0003\_C10 5', mRNA sequence

EFPGRPTRPLDLNGEAPDPAGKDVPLISSPVFPGCDGPSTKCHRNACANGGTCVQQWNSY

SCNCDMTSFTGPTCSDESTAYEFGGGSGLMTFQYPEGRWPDTRRDLLALGFMTSQEDAVM

LRLDSANSNDYMELEIVDGNIFMVYNMGTEDHPVGEVMAKVNDGIYHVVRFVRSGSNATV

QIDDYEVRSKNPKGHQLSVFNAQSRLQIGGRX

>CI998089.1\_2 Marsupenaeus japonicus eyestalk immature Penaeus japonicus cDNA clone YAQ01A02NGRM0003\_C10 5', mRNA sequence

NSRVDPRVRST\*TAKLPTQQAKTFL\*SRAPSSQAVTDQVQNAIGMHVQMVGRACNSGIPI

HATVT\*PPLPDLRVLMSPRRMSSAAALAS\*PSSTLRADGLIPAVTCWHSVL\*PRRRMRSC

\*GWIRPIRMITWNWKSSTAISSWSTTWARRTTP\*ARSWQRSTTAYITWFASLGQVPTPRY

RLTTMKFVQRIQKDTSSPSSTHSHASRSVDEX

>CI998089.1\_3 Marsupenaeus japonicus eyestalk immature Penaeus japonicus cDNA clone YAQ01A02NGRM0003\_C10 5', mRNA sequence

IPGSTHASARLERRSSRPSRQRRSSDLEPRLPRL\*RTKYKMPSECMCKWWDVRATVEFLF

MQL\*HDLLYRTYVF\*\*VHGV\*VRRRLWPHDLPVP\*GQMA\*YPP\*PVGTRFYDLAGGCGHA

EAGFGQFE\*LHGIGNRRRQYLHGLQHGHGGPPRRRGHGKGQRRHISRGSLR\*VRFQRHGT

D\*RL\*SSFKESKRTPALRLQRTVTPPDRWTK

>CI998089.1\_4 Marsupenaeus japonicus eyestalk immature Penaeus japonicus cDNA clone YAQ01A02NGRM0003\_C10 5', mRNA sequence

SSTDLEA\*LCVEDGELVSFWIL\*TNFIVVNLYRGVGT\*PNEANHVIYAVVDLCHDLAYGV

VLRAHVVDHEDIAVDDFQFHVIIRIGRIQPQHDRILLRGHKTECQQVTAGIRPSALRVLE

GHEARAAAELIRRGLIRTRRSGKGGHVTVA\*IGIPLLHARPTICTCIPMAFCTWSVTAWE

DGARDQRNVFACWVGSFAVQVERTRGSTREF

>CI998089.1\_5 Marsupenaeus japonicus eyestalk immature Penaeus japonicus cDNA clone YAQ01A02NGRM0003\_C10 5', mRNA sequence

FVHRSGGVTVR\*RRRAGVLLDSLNELHSRQSVPWRWNLT\*RSEPRDICRR\*PLP\*PRLRG

GPPCPCCRP\*RYCRRRFPIPCNHSNWPNPASA\*PHPPARS\*NRVPTGHGGYQAICPQGTG

RS\*GQSRRRTHTPWTHQNT\*VR\*RRSCHSCMNRNSTVARTSHHLHMHSDGILYLVRHSLG

RRGSRSEERLCLLGRELRRSSRADAWVDPGIX

>CI998089.1\_6 Marsupenaeus japonicus eyestalk immature Penaeus japonicus cDNA clone YAQ01A02NGRM0003\_C10 5', mRNA sequence

LRPPIWRRDCALKTESWCPFGFFERTS\*SSICTVALEPDLTKRTT\*YMPSLTFAMTSPTG

WSSVPML\*TMKILPSTISNSM\*SFELAESSLSMTASSCEVIKPSANRSRRVSGHLPSGYW

KVMRPEPPPNSYAVDSSEHVGPVKEVMSQLHE\*EFHCCTHVPPFAHAFRWHFVLGPSQPG

KTGLEIRGTSLPAGSGASPFKSSGRVGRPGNS

Name: Penaeus neurexin

CI998089

Species: [Penaeus japonicus](https://www.ncbi.nlm.nih.gov/Taxonomy/Browser/wwwtax.cgi?id=27405)

Eukaryota; Metazoa; Ecdysozoa; Arthropoda; Crustacea;

Multicrustacea; Malacostraca; Eumalacostraca; Eucarida; Decapoda;

Dendrobranchiata; Penaeoidea; Penaeidae; Penaeus.

Here, tell me the name of the novel protein, and the species from which it derives. It is very unlikely (but still definitely possible) that you will find a novel gene from an organism such as *S. cerevisiae*, human or mouse, because those genomes have already been thoroughly annotated. It is more likely that you will discover a new gene in a genome that is currently being sequenced, such as bacteria or plants or protozoa.

[**Q4**] Prove that this gene, and its corresponding protein, are novel. For the purposes of this project, “novel” is defined as follows. Take the protein sequence (your answer to [Q3]), and use it as a query in a blastp search of the nr database at NCBI.

* If there is a match with 100% amino acid identity to a protein in the database, from the same species, then your protein is NOT novel (even if the match is to a protein with a name such as “unknown”). Someone has already found and annotated this sequence, and assigned it an accession number.
* If the top match reported has less than 100% identity, then it is likely that your protein is novel, and you have succeeded.
* If there is a match with 100% identity, but to a different species than the one you started with, then you have likely succeeded in finding a novel gene.
* If there are no database matches to the original query from [Q1], this indicates that you have partially succeeded: yes, you may have found a new gene, but no, it is not actually homologous to the original query. You should probably start over.

Graphical user interface, text, application, email

Description automatically generatedThe top result is to a protein from Penaeus Japonicus (caridean shrimp), see second screen shot below for alignment details: Graphical user interface, text

Description automatically generated

[**Q5**] Generate a multiple sequence alignment with your novel protein, your original query protein, and a group of other members of this family from different species. A typical number of proteins to use in a multiple sequence alignment for this assignment purpose is a minimum of 5 and a maximum of 20 - although the exact number is up to you. Include the multiple sequence alignment in your report. Use Courier font with a size appropriate to fit page width.

Side-note: Indicate your sequence in the alignment by choosing an appropriate name for each sequence in the input unaligned sequence file (i.e. edit the sequence file so that the species, or short common, names (rather than accession numbers) display in the output alignment and in the subsequent answers below). The goal in this step is to create an interesting an alignment for building a phylogenetic tree that illustrates species divergence.

[**Q6**] Create a phylogenetic tree, using either a parsimony or distance-based approach. Bootstrapping and tree rooting are optional. Use “simple phylogeny” online from the EBI or any respected phylogeny program (such as MEGA, PAUP, or Phylip). Paste an image of your Cladogram or tree output in your report.

[**Q7**] Generate a sequence identity based **heatmap** of your aligned sequences using R.

If necessary convert your sequence alignment to the ubiquitous FASTA format (Seaview can read in clustal format and “Save as” FASTA format for example). Read this FASTA format alignment into R with the help of functions in the **Bio3D package**. Calculate a sequence identity matrix (again using a function within the Bio3D package). Then generate a heatmap plot and add to your report. Do make sure your labels are visible and not cut at the figure margins.

[**Q8**] Using R/Bio3D (or an online blast server if you prefer), search the main protein structure database for the most similar atomic resolution structures to your aligned sequences.

List the top 3 *unique* hits (i.e. not hits representing different chains from the same structure) along with their Evalue and sequence identity to your query. Please also add annotation details of these structures. For example include the annotation terms PDB identifier (structureId), Method used to solve the structure (experimentalTechnique), resolution (resolution), and source organism (source).

HINT: You can use a single sequence from your alignment or generate a consensus sequence from your alignment using the Bio3D function consensus(). The Bio3D functions blast.pdb(), plot.blast() and pdb.annotate() are likely to be of most relevance for completing this task. Note that the results of blast.pdb() contain the hits PDB identifier (or pdb.id) as well as Evalue and identity. The results of pdb.annotate() contain the other annotation terms noted above.

Note that if your consensus sequence has lots of gap positions then it will be better to use an original sequence from the alignment for your search of the PDB. In this case you could chose the sequence with the highest identity to all others in your alignment by calculating the row-wise maximum from your sequence identity matrix.

[**Q9**] Generate a molecular figure of one of your identified PDB structures using the **NGL viewer** online (or **VMD/PyMol)**. You can optionally highlight conserved residues that are likely to be functional. Please use a white or transparent background for your figure (i.e. not the default black).

Based on sequence similarity. How likely is this structure to be similar to your “novel” protein?

[**Q10**] Perform a “Target” search of ChEMBEL ( <https://www.ebi.ac.uk/chembl/> ) with your novel sequence. Are there any **Target Associated Assays** and **ligand efficiency data** reported that may be useful starting points for exploring potential inhibition of your novel protein?

**Scoring Rubric**:

[45 total points available]

**Q1** (4 points)

Protein name 1

Species 1

Accession number 1

Function known 1

**Q2** (6 points)

Blast method 1

Database searched 1

Limits applied 1

Search output list (top hits) 1

Alignment of choice 1

Evalue and other alignment stats 1

**Q3** (3 points)

Protein sequence of choice matches Subject above 1

Name in header 1

Species 1

**Q4** (3 point)

Blastp output list with identities & Evalue 1

Top alignment shown with alignment statistics 1

Results indicates a “novel” gene found 1

**Q5** (3 points)

MSA labeled with useful names 1

MSA trimmed appropriately (i.e. no gap overhangs) 1

Pasted MSA fits report page width (i.e. font, format) 1

**Q6** (1 point)

Figure illustrates sequence clustering pattern 1

**Q7** (10 points)

Heatmap figure included in report 5

Heatmap is legible (i.e. no labels obscured) 5

**Q8** (10 points)

PDB identifiers from multiple species reported 5

Annotation of PDB source, resolution and technique 4

Annotation of Evalue and Sequence Identity 1

**Q9** (4 points)

Structure figure provided 2

Uses white background for molecular figure 1

Figure of high resolution (i.e. not just snapshot) 1

**Q10** (1 point)

Evidence of ChEMBEL searches 1