Hope Adams A15652616

Find a gene project

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: Kinesin-like protein KIF11

Accession: P52732 Species: Homo Sapiens

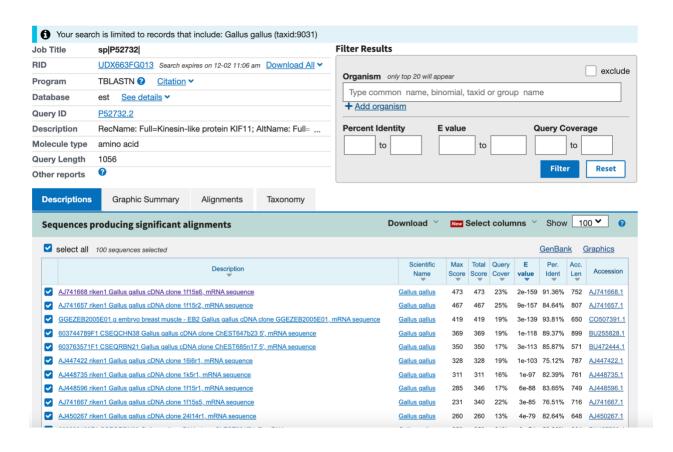
[Q2] Perform a BLAST search against a DNA database, such as a database consisting of genomic DNA or ESTs. The BLAST server can be at NCBI or elsewhere. Include details of the BLAST method used, database searched and any limits applied (e.g. Organism).

Method: tblastn search against Gallus gallus EST's

Database: Expressed Sequence Tags (est)

Organism: Gallus gallus (taxid:9031)

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.



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.

Chosen match: Accession: AJ741668.1, a 752 basepair mRNA from Gallus gallus



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.

AJ741668 riken1 Gallus gallus cDNA clone 1f15s6, mRNA sequence

GenBank: AJ741668.1

FASTA Graphics

```
Go to: 🔽
LOCUS
           AJ741668
                                     752 bp
                                                       linear EST 12-FEB-2011
                                               mRNA
DEFINITION AJ741668 riken1 Gallus gallus cDNA clone 1f15s6, mRNA sequence.
ACCESSION AJ741668
VERSION AJ741668.1
DRIINK BioSample: SAMN00170383
KEYWORDS EST.
           Gallus gallus (chicken)
SOURCE
  ORGANISM Gallus gallus
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Archelosauria; Archosauria; Dinosauria; Saurischia; Theropoda;
            Coelurosauria; Aves; Neognathae; Galloanserae; Galliformes;
            Phasianidae; Phasianinae; Gallus.
REFERENCE
           1 (bases 1 to 752)
  AUTHORS
           Caldwell, R.B., Kierzek, A.M., Arakawa, H., Bezzubov, Y., Zaim, J.,
            Fiedler, P., Kutter, S., Blagodatski, A., Kostovska, D., Koter, M.,
           Plachy, J., Carninci, P., Hayashizaki, Y. and Buerstedde, J.M.
          Full-length cDNAs from chicken bursal lymphocytes to facilitate
  TITLE
           gene function analysis
  JOURNAL Genome Biol. 6 (1), R6 (2005)
  PUBMED <u>15642098</u>
COMMENT
            Contact: Caldwell RB
            GSF - Forschungszentrum, Institut fuer Molekulare Strahlenbiologie
           Ingolstaedter Landstr. 1, D-85764 Neuherberg, GERMANY.
FEATURES
                     Location/Qualifiers
     source
                     1..752
                     /organism="Gallus gallus"
                     /mol_type="mRNA"
                     /db xref="taxon:9031"
                     /-1----"1615-6"
```

[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.

➤ Gallus gallus protein (taken from BLAST result)

Query	72	FGASTKQIDVYRSVVCPILDEVIMGYNCTIFAYGQTGTGKTFTMEGERSPNEEYTWEEDP	131
		F A KQIDVYRSVVCPILDEVIMGYNCT+FAYGQTGTGKTFTMEGERSPNEEYTWEEDP	
Sbjct	731	FRAQAKQIDVYRSVVCPILDEVIMGYNCTVFAYGQTGTGKTFTMEGERSPNEEYTWEEDP	552
Query	132	LAGIIPRTLHQIFEKLTDNGTEFSVKVSLLEIYNEELFDLLNPSSDVSERLQMFDDPRNK	191
		LAGIIPRTLHQIFEKLT+NGTEFSVKVSLLEIYNEELFDLLNP+ DV ERLQMFDDPRNK	
Sbjct	551	LAGIIPRTLHQIFEKLTENGTEFSVKVSLLEIYNEELFDLLNPAPDVGERLQMFDDPRNK	372
Query	192	RGVIIKGLEEITVHNKDEVYQILEKGAAKRTTAATLMNAYSSRSHSVFSVTIHMKETTID	251
		RGVIIKGLEE+TVHNK+EVYQILE+GAAKRTTAAT MNAYSSRSHSVFS+TIHMKETT+D	
Sbjct	371	RGVIIKGLEEVTVHNKNEVYQILERGAAKRTTAATYMNAYSSRSHSVFSITIHMKETTVD	192
Query	252	GEELVKIGKLNLVDLAGSENIGRSGAVDKRAREAGNINQSLLTLGRVITALVERTPHVPY	311
		GEELVKIGKLNLVDLAGSENIGRSGAVDKRAREAGNINQSLLTLGRVI+ALVER P	
Sbjct	191	GEELVKIGKLNLVDLAGSENIGRSGAVDKRAREAGNINQSLLTLGRVISALVERAPAYSI	12
Query	312	RES 314	
		+ES	
Sbjct	11	QES 3	

Gallus gallus protein (taken from BLAST result)

FRAQAKQIDVYRSVVCPILDEVIMGYNCTVFAYGQTGTGKTFTMEGERSPNEEYTWEEDPLAGIIPRTLHQIFEKLT ENGTEFSVKVSLLEIYNEELFDLLNPAPDVGERLQMFDDPRNKRGVIIKGLEEVTVHNKNEVYQILERGAAKRTTAA TYMNAYSSRSHSVFSITIHMKETTVDGEELVKIGKLNLVDLAGSENIGRSGAVDKRAREAGNINQSLLTLGRVISAL VERAPAYSIQES

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.

Name: Kinesin-like protein KIF11

Species: Gallus gallus

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Archelosauria; Archosauria; Dinosauria; Saurischia; Theropoda;

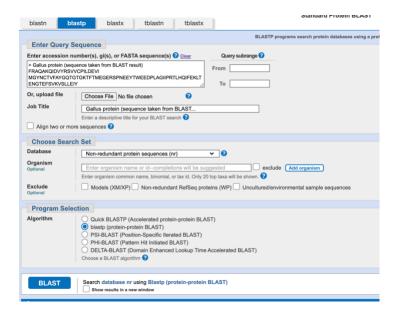
Coelurosauria; Aves; Neognathae; Galloanserae; Galliformes;

Phasianidae; Phasianinae; Gallus.

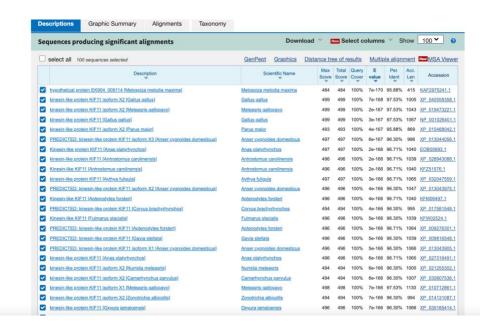
[Q4] Prove that this gene, and its corresponding protein, are novel. For the purposes of this project, "novel" is defined as follows. Take the protein sequence (your answer to [Q3]), and use it as a query in a blastp search of the nr database at NCBI.

- 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.

A BLASTP search against the nr database gave a top hit result to a protein from Gallus Gallus (Red junglefowl Birds):



The highest percent identity was for the species Gallus Gallus but is still under 100%



[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.

>Human KIF11 | CX788535.1

VQENIQQKSKDIVNKMTFHSQKFCADSDGFSQELRNFNQEGTKLVEESVKHSDKLNGNLEKISQETE QRCESLNTRTVYFSEQWVSSLNEREQELHNLLEVVSQCCEASSSDITEKSDGRKAAHEKQHNIFLDQ MTIDEDKLIAQNLELNETIKIGLTKLNCFLEQDLKLDIPTGTTPQRKSYLYPSTLVRTEPREHLLDQ LKRKQPELLMMLNCSENNKEETIPDVDVEEAVLGQYTEEPLSQEPSVDAGVDCSSIGGVPFFQHKKS HGKDKE

>GallusGallus protein (taken from BLAST result)

FRAQAKQIDVYRSVVCPILDEVIMGYNCTVFAYGQTGTGKTFTMEGERSPNEEYTWEEDPLAGIIPRTLHQIFEKLT ENGTEFSVKVSLLEIYNEELFDLLNPAPDVGERLQMFDDPRNKRGVIIKGLEEVTVHNKNEVYQILERGAAKRTTAA

TYMNAYSSRSHSVFSITIHMKETTVDGEELVKIGKLNLVDLAGSENIGRSGAVDKRAREAGNINQSLLTLGRVISAL VERAPAYSIQES

> RedGrouse KIF11 | GW700457.1

QTFRFDYAFDETAPNEMVYRFTARPLVETIFERGMATCFAYGQTGSGKTHTMGGDFSGK---NQDCSKGIYALAARDVFLMLKKPNYKKLELQVYATFFEIYSGKVFDLLNRKT----KLRVLED-GKQQVQVVGLQEREVKCVEDVLKLIEIGNSCRTSGQTSANAHSSRSHAVFQIILRRKGKL-----HGKFSLIDLAGNERGADTSSADROTRLEGAEINKSLLALKECIRALGRNKPHTPFRASKLTQV

>ZebraFinch KIF11 | FE719925.1

KSKSSESVRVVVRCRPMNSKEQTASYEKVVNVDVKLGQVSVKNPRGSSHELPKTFTFDAV YDWNSKQVELYDETFRPLVDSVLQGFNGTIFAYGQTGTGKTYTMEGVRGDPEK------RGVIPNSFDHIFTHISRSQNQQYLVRASYLEIYQEEIRDLL--SKDQSKRLELKERPDT-GVFVKDLTTIVTKSVKEIEHIMNLGNQNRSVGATNMNEHSSRSHAIFQITIECSELGLD

>Mallard KIF11 | DR765078.1

RGEDEKGIPVRVALRCRPLVPKETSEGCQXCLSFVPGEPQVVV------GSDKVFSYDFVFXPTVXQEEVFNTXVAPLVRGIFKGYNATVIAYGQTGSGKTYSMGGAYTASQEH---- DPSVGIIPRVINAAISGEGSGGXDWGI

>SocietyFinch KIF11 | DC282391.1

DGTEFSVKVSLLEIYNEELFDLLNPTPDVGERLQMFDDPRNKRGVIIKGLEEVTVHNKNQVYQILER GAAKRTTAATYMNAYSSRSHSVFSITIHMKETTVDGEELVKIGKLNLVDLAGSENIGRSGAVDKRAR EAGNINQSLLTLGRVITALVERAPHIPYRESKLTRILQDSLGGRTKTSIIATISPASINLEETLSTL EYAHRAKNIMNKPEVNQKLTKKA

>WhiteSparrow KIF11 | XP 005481265.1

KKEEKGKNIQVVVRCRPFNASELKVSSYAVVDCDQARKEVSIRTGGVTDKSSRKTYTFDMVFGAQAK QIDVYRSVVCPILDEVIMGYNCTVFAYGQTGTGKTFTMEGERSPNEEYTWEEDPLAGIIPRTLHQIF EKLTENGTEFSVKVSLLEIYNEELFDLLNPTPDVGERLQMFDDPRNKRGVIIKGLEEVTVHNKNQVY QILERGAAKRTTAATYMNAYSSRSHSVFSITIHMKETTVDGEELVKIGKLNLVDLAGSENIGRSGAV DKRAREAGNINQSLLTLGRVITALVERAPHIPYRESKLTRILQDSLGGRTKTSIIATISPASVNLEE TLSTLEYAHRAKNIMNKPEVNQKLTKKALIKEYTEEIERLKRDLAAAREKNGVYISAENYEALNGKL TVQEEQITEYIDKISVMEEEVKRVTELFRVSKNELEQCKTDLQIKEKELEETQKDLQETKVQLAEEE YVVSVLESTEQELHDTASQLLTTVEETTRDVSGLHAKLERKRAVDQHNAAVQNTFAGQMNASFSKIQ DSITENSLKQQQMLTYYTNCIGDLLSTSSSTADMFASVVSASFACLKELVSTEVSHISEKITQHENL SLDCKAELLRLIEEHQTGLGRAVNS-

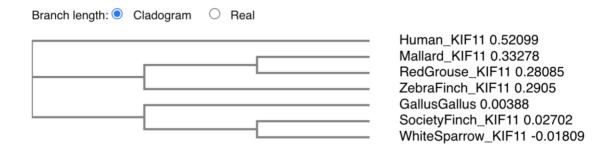
LTPMVEFVLGLNCQFQSNMKKYSAVADQMEDHKKEMDTFFADLSLTLKKIQEQTAGGFAQLQHNCDS LKEEVEMMRLAHRKSAAELMSSLQSQLDLFAQETQKSLTDVLTRNGSLKTTITAMQENVHLKTTDLV SSTNSNHNKFAASLDNFSQELRSINAENKAMLEESNDHCQHLLTNLKNVAQHTNTWGEFTTAQMVNF TNQHLLSFKDEKQQFQYLQKKNEENCDQAIAEIADHIGSQKAAEEKVLNGLLDQIKVDQEILVEQKL ALREQVQHGLTQVNGFLQEDLKVDVPTGTTPQRKDYSYPVTLVRTEPRQLLLEQLRQKQPNLDAMLS SVGKEMEDSAGQDLLEEGVLQEPSESLACDKYSMDTNVYCHTNGGIPFFQHKRSLKKGKENKSAAPL E-NKMEDMTEELLQKSKHPLR

Multiple sequence alignment using MUSCLE at EBI: Human KIF11 Mallard KIF11 RGEDEKGIPVRVALRCRPLVPKETSEGCQXCLSFVPGEPQVVVGSD-----KVFSYD RedGrouse KIF11 -----QTFRFD ZebraFinch_KIF11 --KSKSSESVRVVVRCRPMNSKEQTASYEKVVNVDVKLGQVSVKNPRGSSHELPKTFTFD ______ GallusGallus SocietyFinch KIF11 WhiteSparrow KIF11 -KKEEKGKNIQVVVRCRPFNASELKVSSYAVVDCDQARKEVSIRTGGVTDKSSRKTYTFD QKSKDIVNKMTFHSQKFCADSD----GFSQELRNFNQEGT--KLVEESVKHSDKL----Human KIF11 Mallard_KIF11 FVFXPTVXQEEVFNTXVAPLVRGIFKGYNATVIAYGQTGSGKTYSMGGAYTASQEH---RedGrouse_KIF11 YAFDETAPNEMVYRFTARPLVETIFERGMATCFAYGQTGSGKTHTMGGDFSGKNQ---ZebraFinch_KIF11 AVYDWNSKQVELYDETFRPLVDSVLQGFNGTIFAYGQTGTGKTYTMEGVRGDPEK---GallusGallus --FRAQAKQIDVYRSVVCPILDEVIMGYNCTVFAYGQTGTGKTFTMEGERSPNEEYTWEE SocietyFinch_KIF11 MVFGAQAKQIDVYRSVVCPILDEVIMGYNCTVFAYGQTGTGKTFTMEGERSPNEEYTWEE Human KIF11 Mallard KIF11 RedGrouse KIF11 ZebraFinch_KIF11 VGERLQMFDDPRNKRGVIIKGLEEVTVHNKNEVYQILERGAAKRTTAATYMNAY----VGERLQMFDDPRNKRGVIIKGLEEVTVHNKNQVYQILERGAAKRTTAATYMNAY-----GallusGallus SocietyFinch KIF11 WhiteSparrow_KIF11 VGERLQMFDDPRNKRGVIIKGLEEVTVHNKNQVYQILERGAAKRTTAATYMNAY-----Human KIF11 LDIPTGTTPQRKSYLYPSTLVRTEPREHLLDQLKRKQPELLMMLNCSENNKEETIPDVDV Mallard_KIF11 -----XDWGI-----RedGrouse KIF11 -----SSRSHAVFQIIL----RRKGKLH-----GKFSLIDLAGNERGADTSSADR RedGrouse_KIII ZebraFinch_KIF11 -----SSRSHAIFOITI---ECSELGLD------GallusGallus -----SSRSHSVFSITI---HMKETTVDGEELVKIGKLNLVDLAGSENIGRSGAVDK SocietyFinch_KIF11 -----SSRSHSVFSITI--HMKETTVDGEELVKIGKLNLVDLAGSENIGRSGAVDK WhiteSparrow_KIF11 -----SSRSHSVFSITI--HMKETTVDGEELVKIGKLNLVDLAGSENIGRSGAVDK Human KIF11 EEAVLGQYTEEPLSQEPSVDAGVDCSSIGGVPFFQHKKSHGKDKE--------Mallard KIF11 RedGrouse KIF11 OTRLEGAEINKSLLALKECIRAL----GRNKPHTPFRASKLTOV--------ZebraFinch KIF11 GallusGallus RAREAGN-INQSLLTLGRVISAL----VERAPAYSIQES------SocietyFinch_KIF11 RAREAGN-INQSLLTLGRVITAL----VERAPHIPYRESKLTRILQDSLGGRTKTSIIAT WhiteSparrow_KIF11 RAREAGN-INQSLLTLGRVITAL-----VERAPHIPYRESKLTRILQDSLGGRTKTSIIAT

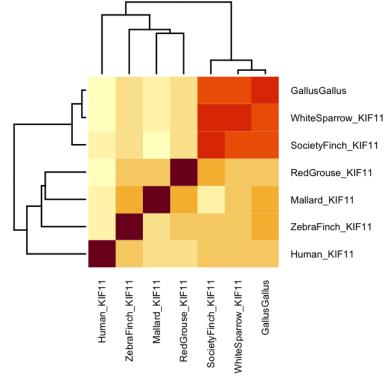
Human_KIF11 Mallard_KIF11 RedGrouse_KIF11 ZebraFinch_KIF11 GallusGallus SocietyFinch_KIF11 WhiteSparrow_KIF11	ISPASINLEETLSTLEYAHRAKNIMNKPEVNQKLTKKAISPASVNLEETLSTLEYAHRAKNIMNKPEVNQKLTKKALIKEYTEEIERLKRDLAAAREK
Human_KIF11 Mallard_KIF11 RedGrouse_KIF11 ZebraFinch_KIF11 GallusGallus SocietyFinch_KIF11 WhiteSparrow_KIF11	NGVYISAENYEALNGKLTVQEEQITEYIDKISVMEEEVKRVTELFRVSKNELEQCKTDLQ
Human_KIF11 Mallard_KIF11 RedGrouse_KIF11 ZebraFinch_KIF11 GallusGallus SocietyFinch_KIF11 WhiteSparrow_KIF11	IKEKELEETQKDLQETKVQLAEEEYVVSVLESTEQELHDTASQLLTTVEETTRDVSGLHA
Human_KIF11 Mallard_KIF11 RedGrouse_KIF11 ZebraFinch_KIF11 GallusGallus SocietyFinch_KIF11 WhiteSparrow_KIF11	KLERKRAVDQHNAAVQNTFAGQMNASFSKIQDSITENSLKQQQMLTYYTNCIGDLLSTSS
Human_KIF11 Mallard_KIF11 RedGrouse_KIF11 ZebraFinch_KIF11 GallusGallus SocietyFinch_KIF11 WhiteSparrow_KIF11	STADMFASVVSASFACLKELVSTEVSHISEKITQHENLSLDCKAELLRLIEEHQTGLGRA
Human_KIF11 Mallard_KIF11 RedGrouse_KIF11 ZebraFinch_KIF11 GallusGallus SocietyFinch_KIF11 WhiteSparrow_KIF11	VNSLTPMVEFVLGLNCQFQSNMKKYSAVADQMEDHKKEMDTFFADLSLTLKKIQEQTAGG
Human_KIF11 Mallard_KIF11 RedGrouse_KIF11 ZebraFinch_KIF11 GallusGallus SocietyFinch_KIF11 WhiteSparrow_KIF11	FAQLQHNCDSLKEEVEMMRLAHRKSAAELMSSLQSQLDLFAQETQKSLTDVLTRNGSLKT

Human_KIF11 Mallard_KIF11 RedGrouse_KIF11 ZebraFinch_KIF11 GallusGallus SocietyFinch_KIF11 WhiteSparrow_KIF11	TITAMQENVHLKTTDLVSSTNSNHNKFAASLDNFSQELRSINAENKAMLEESNDHCQHLL
Human_KIF11 Mallard_KIF11 RedGrouse_KIF11 ZebraFinch_KIF11 GallusGallus SocietyFinch_KIF11 WhiteSparrow_KIF11	TNLKNVAQHTNTWGEFTTAQMVNFTNQHLLSFKDEKQQFQYLQKKNEENCDQAIAEIADH
Human_KIF11 Mallard_KIF11 RedGrouse_KIF11 ZebraFinch_KIF11 GallusGallus SocietyFinch_KIF11 WhiteSparrow_KIF11	IGSQKAAEEKVLNGLLDQIKVDQEILVEQKLALREQVQHGLTQVNGFLQEDLKVDVPTGT
Human_KIF11 Mallard_KIF11 RedGrouse_KIF11 ZebraFinch_KIF11 GallusGallus SocietyFinch_KIF11 WhiteSparrow_KIF11	TPQRKDYSYPVTLVRTEPRQLLLEQLRQKQPNLDAMLSSVGKEMEDSAGQDLLEEGVLQE
Human_KIF11 Mallard_KIF11 RedGrouse_KIF11 ZebraFinch_KIF11 GallusGallus SocietyFinch_KIF11 WhiteSparrow_KIF11	PSESLACDKYSMDTNVYCHTNGGIPFFQHKRSLKKGKENKSAAPLENKMEDMTEELLQKS
Human_KIF11 Mallard_KIF11 RedGrouse_KIF11 ZebraFinch_KIF11 GallusGallus SocietyFinch_KIF11 WhiteSparrow_KIF11	 KHPLR

[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 E-value and sequence identity to your query. Please also add annotation details of these structures. For example, include the annotation terms PDB identifier

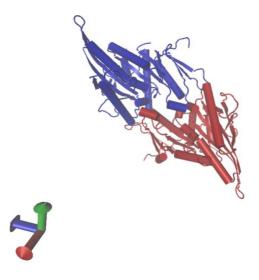
(structure Id), Method used to solve the structure (experimental Technique), resolution (resolution), and source organism (source).

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.

ID	Technique	Resolution	Source	E-value	Identity
4ZCA_A	X-ray	2.3	Homo	0.00e+00	90.529
	Diffraction		sapiens		
1IA0_K	Electron	15.0	Mus	1.67e-66	37.500
	microscope		musculus		
2WBE_C	Electron	9.4	Drosophila	3.68e-140	57.713
	microscope		melanogaster		

[Q9] Generate a molecular figure of one of your identified PDB structures using VMD. 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?



This structure is very likely to be similar in structure to Kinesin-like protein according to the high similarity (90.529%) and the very low e value of 0.00e+00.

[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? CHEMBL details 8 Binding Assay (CHEMBL4581) and 163 Functional Assays. There were multiple points on the ligand efficiency graph shown below.



