

Practical

1. Sequence information resource
2. Understanding and use of various web resources: EMBL, Genbank, Entrez, Unigene, Protein information resource (PIR).
3. Understanding and using: PDB, Swissprot, TREMBL
4. Using various BLAST and interpretation of results.
5. Retrieval of information from nucleotide databases.
6. Sequence alignment using BLAST.
7. Multiple sequence alignment using Clustal W.

References

1. Ghosh, Z., & Bibekanand M. (2008). *Bioinformatics: Principles and Applications*. Oxford University Press.
2. Pevsner, J. (2009). *Bioinformatics and Functional Genomics* (2nd ed.). Wiley-Blackwell.
3. Campbell ,A. M., & Heyer, L.J. (2006). *Discovering Genomics, Proteomics and Bioinformatics* (2nd ed.). Benjamin Cummings.

KARPAGAM ACADEMY OF HIGHER EDUCATION, COIMBATORE
DEPARTMENT OF BIOTECHNOLOGY

LIST OF PRACTICAL

Name of the Faculty: Dr. S. Selvakumar	Department: Biotechnology
Course name: Bioinformatics Practical	Course code: 17BTU512A
Academic Year: 2019-2020	Semester: Fifth
Class: III year B.Sc.,	Section: A

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Exp No: 1 Sequence information resources

AIM: To explore unique features and applications of various sequence resources.

Procedure: The National Center for Biotechnology Information (NCBI) is part of the United States National Library of Medicine (NLM), a branch of the National Institutes of Health (NIH). The NCBI houses a series of databases relevant to biotechnology and biomedicine and is an important resource for bioinformatics tools and services.

Gene database has been implemented at NCBI to characterize and organize the information about genes. It serves as a major node in the nexus of genomic map, expression, sequence, protein function, structure and homology data. A unique GeneID is assigned to each gene record that can be followed through revision cycles. Protein database is an important protein resource at NCBI. It maintains the text record for individual protein sequences, derived from many different resources such as NCBI Reference Sequence (RefSeq) project, GenBank, PDB and UniProtKB/Swiss-Prot. Protein records are present in different formats including FASTA and XML and are linked to other NCBI resources. Protein provides the relevant data to the users such as genes, DNA/RNA sequences, biological pathways, expression and variation data and literature.

- The amino acid sequence of proteins, nucleic acids and SNP would be downloaded in the FASTA format from the website <https://www.ncbi.nlm.nih.gov/>.
- Once the FASTA file is downloaded, it can be opened in a word pad file and stored in txt format.

UniProt is a freely accessible database of protein sequence and functional information, many entries being derived from genome sequencing projects. It contains a large amount of information about the biological function of proteins derived from the research literature. UniProt Archive (UniParc) is a comprehensive and non-redundant database, which contains all the protein sequences from the main, publicly available protein sequence databases.

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There are many publicly available protein sequence databases.

FlyBase: the primary repository of genetic and molecular data for the insect family Drosophilidae (FlyBase)

H-Invitational Database (H-Inv)

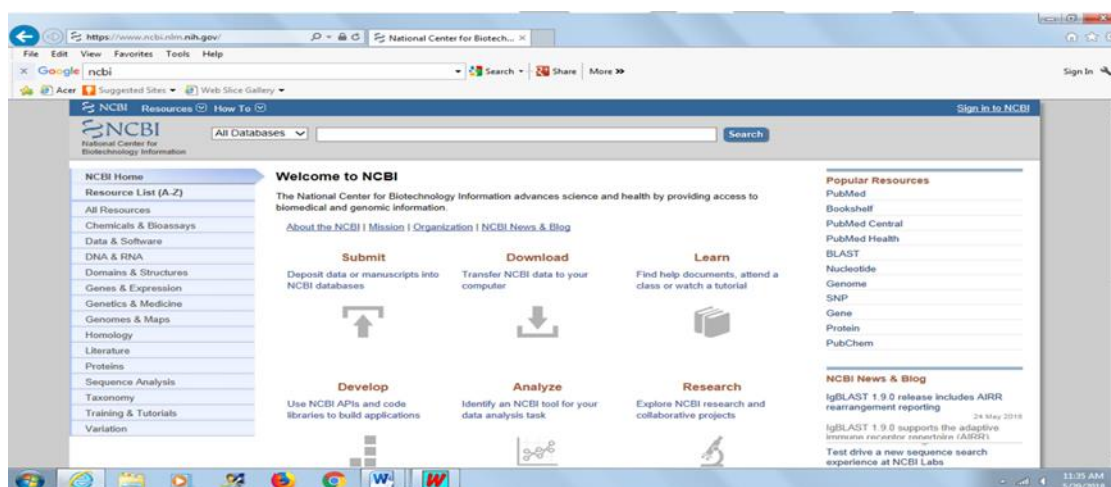
International Protein Index (IPI)

Saccharomyces Genome Database (SGD)

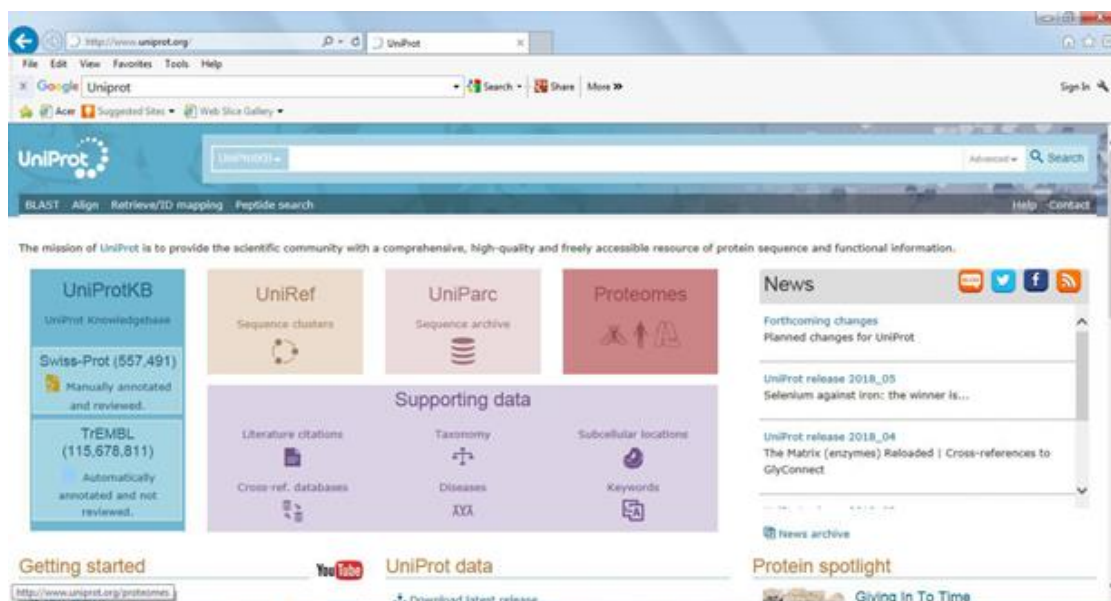
The Arabidopsis Information Resource (TAIR)

Vertebrate and Genome Annotation Database (VEGA)

WormBase



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Information pertaining to primary, secondary and tertiary structures of proteins/nucleic acids can be inferred from the annotation webpages of the corresponding sequences. For instance, annotation data of myoglobin are herein listed out for the quick understanding.

Protein : Myoglobin
Gene : MB
Organism : *Homo sapiens* (Human)
Status : Reviewed
Annotation score : Function

Serves as a reserve supply of oxygen and facilitates the movement of oxygen within muscles.

Sites

Feature key	Position(s)	DescriptionActions	Graphical view	Length
-------------	-------------	--------------------	----------------	--------

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Metal binding ⁱ	65	Iron (heme distal ligand)		1
Metal binding ⁱ	94	Iron (heme proximal ligand)		1

GO - Molecular function

View the complete GO annotation on QuickGO

GO - Biological process

Keywords

Molecular function	Muscle protein
Biological process	Oxygen transport, Transport
Ligand	Heme, Iron, Metal-binding

Enzyme and pathway databases

Reactome ⁱ	R-HSA-8981607 Intracellular oxygen transport
-----------------------	--

Protein family/group databases

TCDB ⁱ	1.A.107.1.3 the pore-forming globin (globin) family
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Names & Taxonomy

Protein names	Recommended name: Myoglobin
Gene names	Name: MB
Organism	Homo sapiens (Human)
Taxonomic identifier	9606 [NCBI]
Taxonomic lineage	Eukaryota › Metazoa › Chordata › Craniata › Vertebrata › Euteleostomi › Mammalia › Eutheria › Euarchontoglires › Primates › Haplorrhini › Catarrhini › Hominidae › Homo
Proteomes	UP000005640 Component: Chromosome 22

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Organism-specific databases

EuPathDB	HostDB:ENSG00000198125.12
HGNC	HGNC:6915 MB
MIM	160000 gene
neXtProt	NX_P02144

Subcellular locationⁱ

Extracellular region or secreted Cytosol Plasma membrane Cytoskeleton Lysosome Endosome Peroxisome ER Golgi apparatus Nucleus Mitochondrion Manual annotation Automatic computational assertion Graphics by Christian Stolte; Source: COMPARTMENTS

- GO - Cellular component
- Cytosol
 - Cytosol Source: Reactome
- Extracellular region or secreted
 - Extracellular exosome Source: UniProtKB

View the complete GO annotation on QuickGO

Pathology & Biotech

Organism-specific databases

DisGeNET	4151
OpenTargets	ENSG00000198125
PharmGKB	PA30658

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Chemistry databases

ChEMBL	CHEMBL2406892
DrugBank	DB02671 1-Methylimidazole
	DB03385 4-Methylimidazole
	DB02379 Beta-D-Glucose
	DB02073 Biliverdine Ix Alpha
	DB03399 Ethyl Isocyanide
	DB04337 Methyl Isocyanide
	DB02396 Methylethylamine
	DB01826 N-Butyl Isocyanide
	DB04050 N-Propyl Isocyanide
	DB02646 Nitrosoethane
	DB01710 Porphyrin Fe(III)
	DB02528 Tetrazolyl Histidine

Polymorphism and mutation databases

BioMuta	MB
DMDM	127661

Results:

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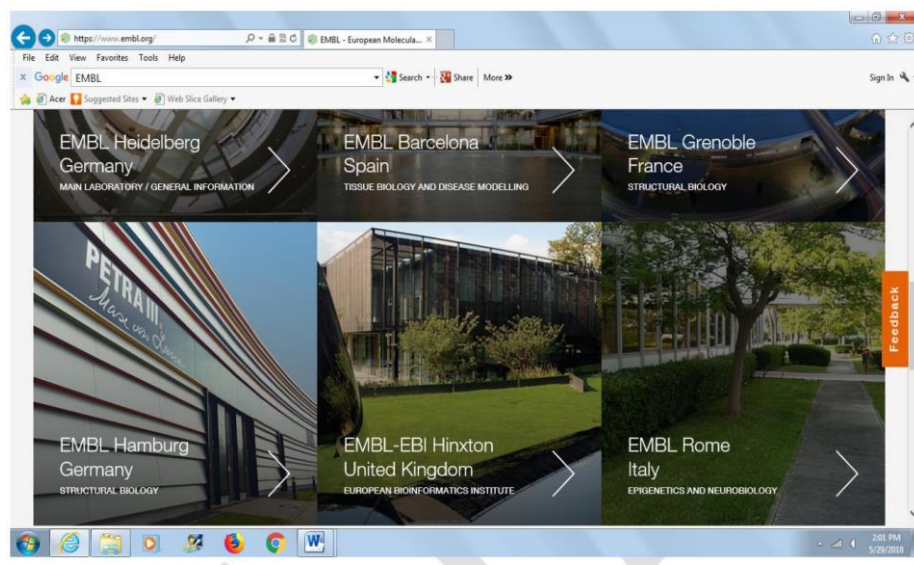
Exp No: 2. Understanding and use of various web resources: EMBL, GenBank, Entrez, Unigene and Protein information resources

Aim: To understand and exploit the biological data available in various web resources.

Procedure: The European Molecular Biology Laboratory (EMBL) is a molecular biology research institution supported by 22 member states, four prospect and two associate member states. EMBL was created in 1974 and is an intergovernmental organisation funded by public research money from its member states. Research at EMBL is conducted by approximately 85 independent groups covering the spectrum of molecular biology. The Laboratory operates from five sites: the main laboratory in Heidelberg, and outstations in Hinxton (the European Bioinformatics Institute (EBI), in England), Grenoble (France), Hamburg (Germany), Monterotondo (near Rome) and Barcelona (Spain). EMBL groups and laboratories perform basic research in molecular biology and molecular medicine as well as training for scientists, students and visitors. The organization aids in the development of services, new instruments and methods, and technology in its member states. Israel is the only Asian state that has full membership. In March 2010, the EMBL Advanced Training Centre (ATC) was inaugurated on the main campus in Heidelberg. Shaped in the form of a double helix, it hosts conferences and provides training. EMBL also runs an active Science and Society Programme which offers activities and events on current questions in life science research for the general public and the scientific community.

GenBank is the NIH genetic sequence database, an annotated collection of all publicly available DNA sequences. GenBank is part of the International Nucleotide Sequence Database Collaboration, which comprises the DNA DataBank of Japan (DDBJ), the European Nucleotide Archive (ENA), and GenBank at NCBI. These three organizations exchange data on a daily basis.

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Access to GenBank: There are several ways to search and retrieve data from GenBank.

Search GenBank for sequence identifiers and annotations with Entrez Nucleotide, which is divided into three divisions: CoreNucleotide (the main collection), dbEST (Expressed Sequence Tags), and dbGSS (Genome Survey Sequences).

Search and align GenBank sequences to a query sequence using BLAST (Basic Local Alignment Search Tool). BLAST searches CoreNucleotide, dbEST, and dbGSS independently; see BLAST info for more information about the numerous BLAST databases.

Search, link, and download sequences programatically using NCBI e-utilities.

The ASN.1 and flatfile formats are available at NCBI's anonymous FTP server: <ftp://ftp.ncbi.nlm.nih.gov/ncbi-asn1> and <ftp://ftp.ncbi.nlm.nih.gov/genbank>.

Entrez is a molecular biology database system that provides integrated access to nucleotide and protein sequence data, gene-centered and genomic mapping information, 3D structure data, PubMed MEDLINE, and more. The system is produced by the National Center for Biotechnology Information (NCBI) and is available via the Internet

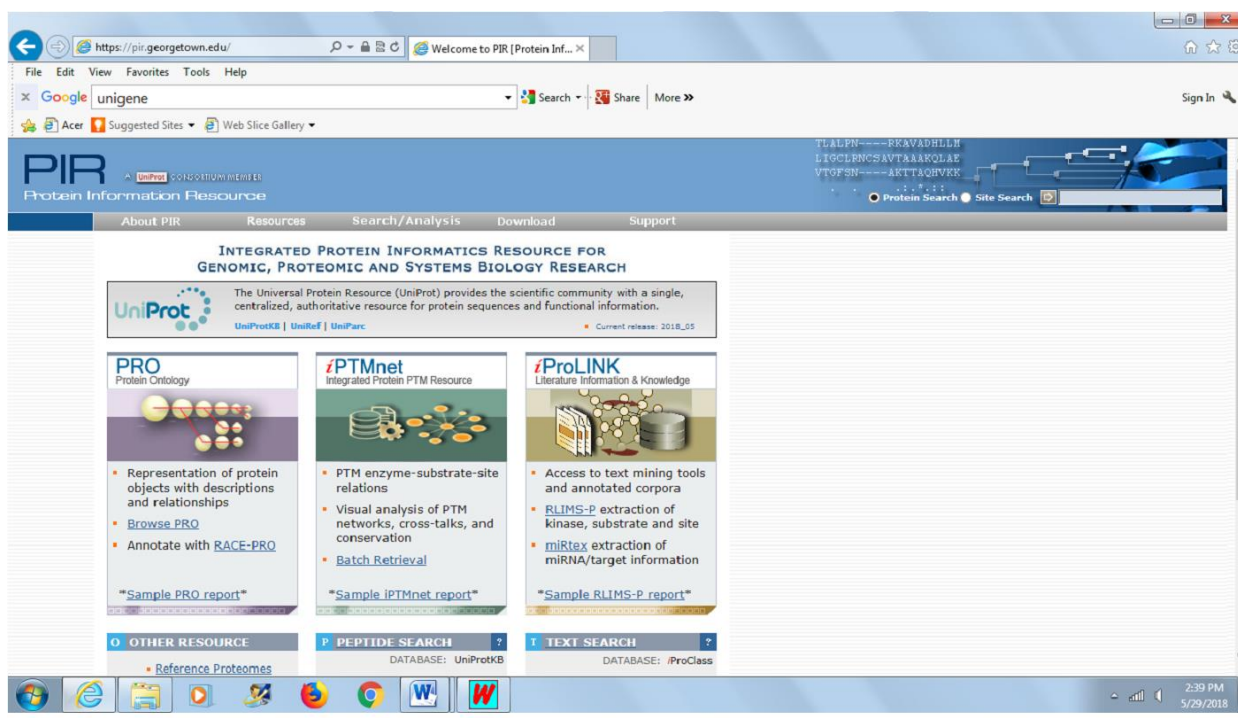
UniGene is an NCBI database of the transcriptome and thus, despite the name, not primarily a

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database for genes. Each entry is a set of transcripts that appear to stem from the same transcription locus (i.e. gene or expressed pseudogene). Information on protein similarities, gene expression, cDNA clones, and genomic location is included with each entry.

The Protein Information Resource (PIR) is an integrated public bioinformatics resource to support genomic, proteomic and systems biology research and scientific studies. The PIR continues to offer world leading resources to assist with proteomic and genomic data integration and the propagation and standardization of protein annotation.



Result:

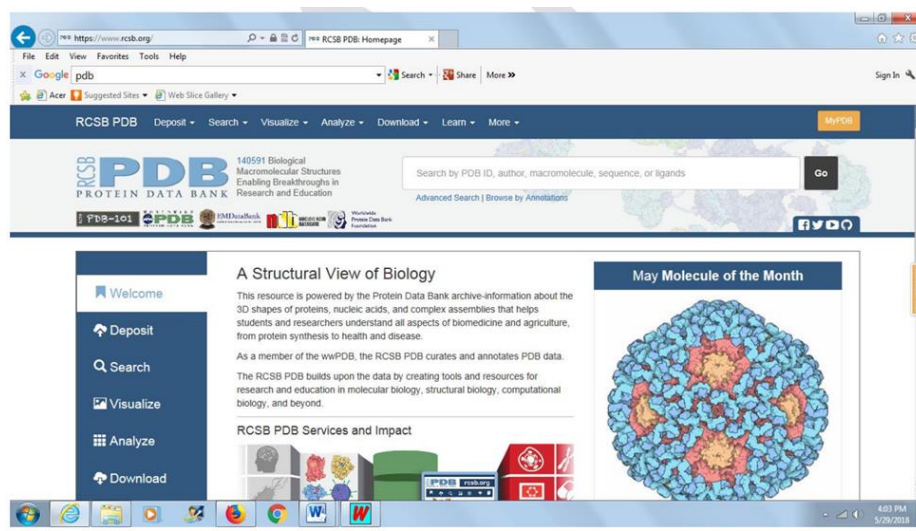
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Exp No: 3. Understanding and use of PDB, Swissprot and TrEMBL

Aim: To analyze various sequence and structural information provided in the PDB, Swiss-Prot and TREMBL databases.

Procedure: SWISS-PROT is a curated protein sequence database which strives to provide a high level of annotation (such as the description of the function of a protein, its domains structure, post-translational modifications, variants, etc.), a minimal level of redundancy and high level of integration with other databases. Recent developments of the database include format and content enhancements, cross-references to additional databases, new documentation files and improvements to TrEMBL, a computer-annotated supplement to SWISS-PROT. TrEMBL consists of entries in SWISS-PROT-like format derived from the translation of all coding sequences (CDSs) in the EMBL Nucleotide Sequence Database, except the CDSs already included in SWISS-PROT. SWISS-PROT is available at: <http://www.expasy.ch/sprot/> and <http://www.ebi.ac.uk/swissprot/>

The Protein Data Bank (PDB) is a crystallographic database for the three-dimensional structural data of large biological molecules, such as proteins and nucleic acids. The data, typically obtained by X-ray crystallography, NMR spectroscopy, or, increasingly, cryo-electron microscopy, and submitted by biologists and biochemists from around the world, are freely accessible on the Internet via the websites of its member organisations (PDBe, PDBj, and RCSB). The PDB is overseen by an organization called the Worldwide Protein Data Bank, wwPDB. The PDB is a key resource in areas of structural biology, such as structural genomics.



TITLE 3 SHORT NEUROTOXINS

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COMPND MOL_ID: 1;

COMPND 2 MOLECULE: CARDIOTOXIN III;

COMPND 3 CHAIN: A;

COMPND 4 ENGINEERED: YES

SOURCE MOL_ID: 1;

SOURCE 2 ORGANISM_SCIENTIFIC: NAJA ATRA;

SOURCE 3 ORGANISM_COMMON: CHINESE COBRA;

SOURCE 4 ORGANISM_TAXID: 8656

KEYWDS CARDIOTOXIN

EXPDTA SOLUTION NMR

AUTHOR R.BHASKARAN,C.C.HUANG,K.D.CHANG,C.YU

REVDAT 3 24-FEB-09 2CRT 1 VERSN

REVDAT 2 01-APR-03 2CRT 1 JRNL

REVDAT 1 01-NOV-94 2CRT 0

JRNL AUTH R.BHASKARAN,C.C.HUANG,D.K.CHANG,C.YU

JRNL TITL CARDIOTOXIN III FROM THE TAIWAN COBRA (NAJA NAJA

JRNL TITL 2 ATRA). DETERMINATION OF STRUCTURE IN SOLUTION AND

JRNL TITL 3 COMPARISON WITH SHORT NEUROTOXINS.

JRNL REF J.MOL.BIOL. V. 235 1291 1994

JRNL REFN ISSN 0022-2836

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JRNL PMID 8308891

JRNL DOI 10.1006/JMBI.1994.1082

REMARK 1

REMARK 2

REMARK 2 RESOLUTION. NOT APPLICABLE.

REMARK 3

REMARK 3 REFINEMENT.

REMARK 3 PROGRAM : X-PLOR

REMARK 3 AUTHORS : BRUNGER

REMARK 3

REMARK 3 OTHER REFINEMENT REMARKS: NULL

REMARK 4

REMARK 4 2CRT COMPLIES WITH FORMAT V. 3.15, 01-DEC-08

REMARK 100

REMARK 100 THIS ENTRY HAS BEEN PROCESSED BY BNL.

REMARK 210

REMARK 210 EXPERIMENTAL DETAILS

REMARK 210 EXPERIMENT TYPE : NMR

REMARK 210 TEMPERATURE (KELVIN) : NULL

The following is atomic coordinates of the protein. It provides information pertaining to atom

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types, atom nomenclature, atom number, atom position and atom occupancy.

ATOM	1	N	LEU	A	1	12.825	6.867	0.006	1.00	1.76
N										
ATOM	2	CA	LEU	A	1	12.032	6.006	0.919	1.00	0.90
C										
ATOM	3	C	LEU	A	1	11.849	4.631	0.268	1.00	0.73
C										
ATOM	4	O	LEU	A	1	12.036	4.471	-0.924	1.00	0.90
O										
ATOM	5	CB	LEU	A	1	10.658	6.639	1.165	1.00	1.16
C										
ATOM	6	CG	LEU	A	1	10.813	8.094	1.633	1.00	1.61
C										
ATOM	7	CD1	LEU	A	1	9.429	8.662	1.957	1.00	2.23
C										
ATOM	8	CD2	LEU	A	1	11.676	8.150	2.900	1.00	2.57
C										
ATOM	9	H1	LEU	A	1	13.292	6.273	-0.707	1.00	2.15
H										
ATOM	10	H2	LEU	A	1	12.194	7.547	-0.470	1.00	2.36
H										
ATOM	11	H3	LEU	A	1	13.547	7.380	0.551	1.00	2.21
H										

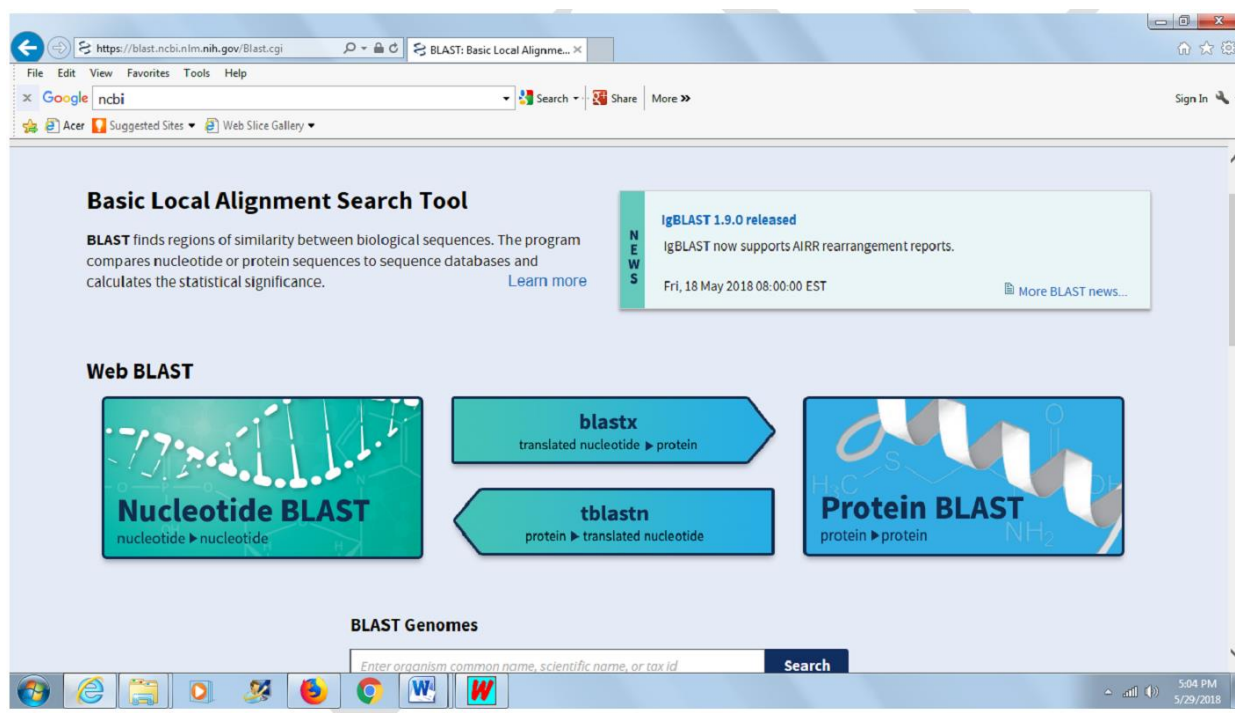
Results:

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Exp No: 4. Using various BLAST and interpretation of results

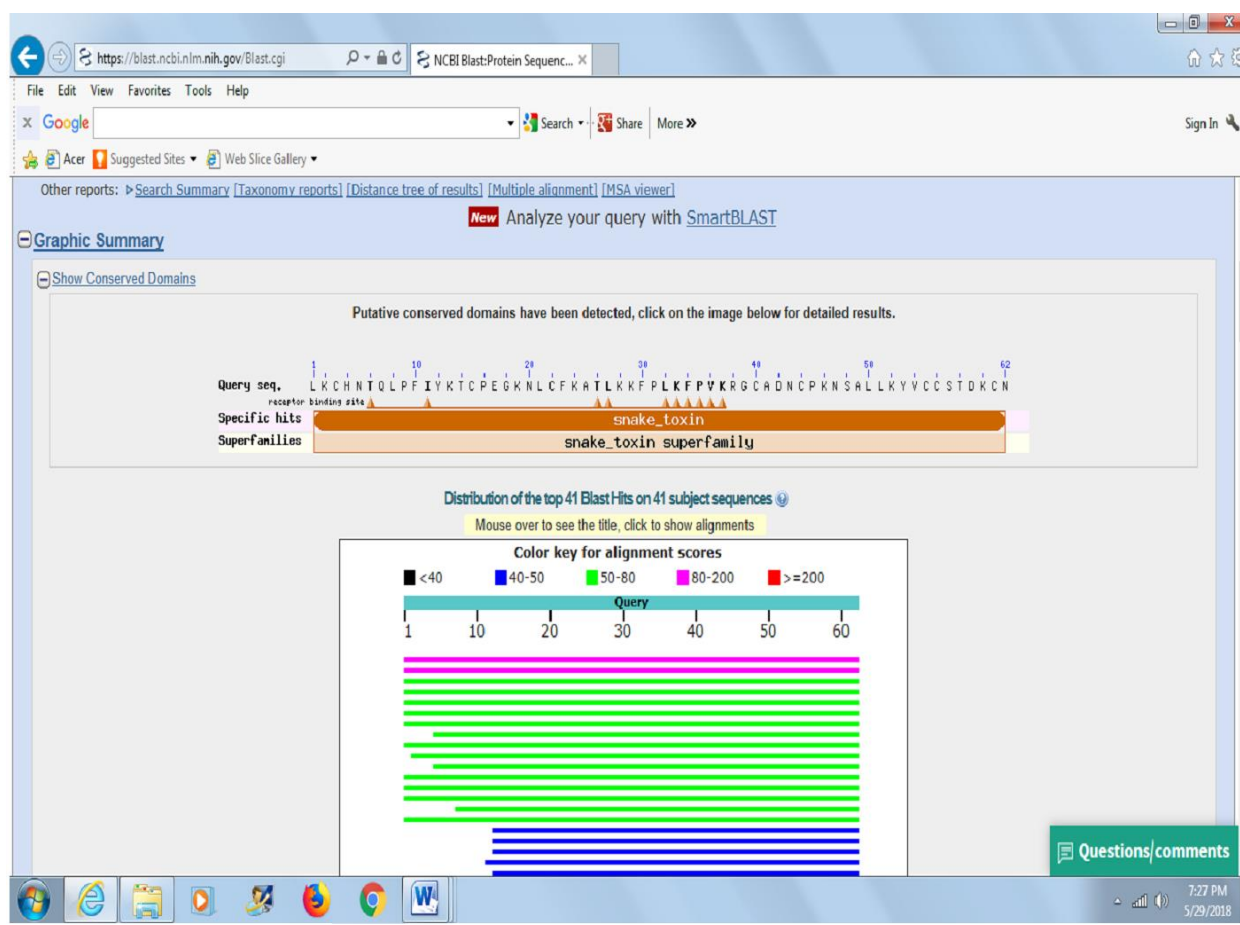
Aim: To retrieve similar sequences for a given protein/nucleotide primary structure using BLAST and rationalizing the outcomes

Procedure: Retrieve a few numbers of protein/nucleotide sequences from NCBI/UniProt databases (<https://www.ncbi.nlm.nih.gov/>) and save the retrieved sequences in FASTA format and accession IDs for all the sequences as well. The ‘sequence similarity search’ can be executed using various BLAST algorithms.



Using protein blast, similar sequences can be obtained either from primary or from structural databases. In general, for distant related and close related sequences, PSI-BLAST, BLASTP algorithms are preferred, respectively. For instance, a cardiotoxin (1CVO) was subjected to BLASTP similarity search and the outcomes were as shown herein below.

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The sequences would be further analyzed on the basis of query coverage, total score, percentage of identities and E-values. The score values and other statistical parameters for a few hits of the query sequences are depicted in the following illustration.

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Sequences producing significant alignments:

Select: All None Selected: 0

Alignments Download GenPept Graphics Distance tree of results Multiple alignment

Description	Max score	Total score	Query cover	E value	Ident	Accession
☐ Chain A. Structure Of Cytotoxin Homolog Precursor	124	124	100%	1e-39	100%	1KXI_A
☐ Chain A. Crystal Structure Of A Three Finger Toxin From Snake Venom	86.3	86.3	100%	2e-24	68%	3VTS_A
☐ Chain A. CARDIOTOXIN V4/II FROM NAJA MOSSAMBICA MOSSAMBICA: THE REFINED CRYSTAL STRUCTURE	78.2	78.2	100%	3e-21	66%	1CDT_A
☐ Chain A. NMR structure with tightly bound water molecule of cytotoxin I from Naja oxiata in aqueous solution (major form)	76.3	76.3	100%	2e-20	65%	1RL5_A
☐ Chain A. Minor form of the recombinant cytotoxin-1 from N. oxiata	76.3	76.3	100%	2e-20	65%	5LUE_A
☐ Chain A. X-RAY STRUCTURE AT 1.55 Å OF TOXIN GAMMA. A CARDIOTOXIN FROM NAJA NIGRICOLLIS VENOM. CRYSTAL PACKING REVEALS A	75.5	75.5	100%	4e-20	61%	1TGX_A
☐ Chain A. CARDIOTOXIN II FROM TAIWAN COBRA VENOM. NAJA NAJA ATRA: STRUCTURE IN SOLUTION AND COMPARISON AMONG HOMOLOG	75.1	75.1	100%	6e-20	65%	1CRE_A
☐ Chain A. SOLUTION STRUCTURE OF CARDIOTOXIN IV. NMR. 1 STRUCTURE	74.3	74.3	93%	1e-19	64%	1KBS_A
☐ Chain A. Nmr Structure Of Ctx A3 At Neutral Ph (20 Structures)	73.6	73.6	100%	2e-19	63%	1IG2_A
☐ Chain A. Crystal Structure Of Beta-cardiotoxin. A Novel Three-finger Cardiotoxin From The Venom Of Ophiophagus Hannah	73.6	73.6	98%	3e-19	57%	3PLC_A
☐ Chain A. DETERMINATION OF THE NUCLEAR MAGNETIC RESONANCE SOLUTION STRUCTURE OF CARDIOTOXIN CTX IIB FROM NAJA MOSSA	72.8	72.8	93%	5e-19	59%	2CCX_A
☐ Chain A. STRUCTURE OF COBRA CARDIOTOXIN CTX IAS DERIVED FROM NUCLEAR MAGNETIC RESONANCE SPECTROSCOPY AND DISTANCE	71.2	71.2	100%	2e-18	61%	2CDX_A
☐ Chain S. Elucidation Of The Solution Structure Of Cardiotoxin Analogue V From The Taiwan Cobra (Naja Naja Atra) Venom	70.1	70.1	100%	5e-18	60%	1CHV_S
☐ Chain A. Crystal Structure Of Cardiotoxin Vi From Taiwan Cobra (Naja Atra) Venom	70.1	70.1	100%	6e-18		

Questions/comments

Similar types of data analyzes can be carried out for nucleotide sequences and as well for other BLAST algorithms.

Results:

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Exp No: 5. Retrieval of information from nucleotide databases

Aim: To retrieve nucleotide sequences and their annotations from various nucleotide databases

Procedure: The nucleotide sequences under interest can be retrieved from any one of the suitable databases such as Gene, UniGene, EST, Nucleotide and Species-specific Genome repository. Using name of the protein, a general search can be carried out at the beginning and then species-specific sequences can be obtained in a systematic search options. Following are the data of myoglobin (*Homo sapiens*) as taken from the 'Nucleotide (NCBI database)'.

Homo sapiens myoglobin (MB) gene, complete cds

GenBank: AH002877.2

LOCUS AH002877 6889 bp DNA linear PRI 01-AUG-2016
DEFINITION Homo sapiens myoglobin (MB) gene, complete cds.
ACCESSION AH002877 M10090 M14602 M14603
VERSION AH002877.2
KEYWORDS Fok family repetitive sequence; direct repeat; myoglobin; repeat region; tandem repeat.
SOURCE Homo sapiens (human)
ORGANISM [Homo sapiens](#)
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Primates; Haplorrhini; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 6889)
AUTHORS Akaboshi, E.
TITLE Cloning of the human myoglobin gene
JOURNAL Gene 33 (3), 241-249 (1985)
PUBMED [2989088](#)
COMMENT On or before Aug 1, 2016 this sequence version replaced [M10090.1](#), [M14602.1](#), [M14603.1](#), [AH002877.1](#).
FEATURES
Location/Qualifiers
source 1..6889
/organism="Homo sapiens"
/mol_type="genomic DNA"
/db_xref="taxon:[9606](#)"
/map="22q11.2-qter"
[repeat region](#) 1100..1750
/note="Fok family repetitive element"
[prim transcript](#) 1895..>2552
/note="myoglobin mRNA and intron"
[gene](#) 1965..5924
/gene="MB"

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```
CDS          join(1965..2059,4067..4289,5778..5924)
               /gene="MB"
               /note="myoglobin"
               /codon_start=1
               /protein_id="AAA59595.1"

/translation="MGLSDGEWQLVLNVWGKVEADIPGHGQEV LIRLFKGH PETLEKF
DKFKHLKSEDEMKASEDLKKHGATVLTALGGILKKKGHHEAEIKPLAQSHATKHKIPV
               KYLEFISECIIQVLQSKHPGDFGADAEGAMNKALELFRKDMASNYKELGFQG"
exon          <1965..2059
               /gene="MB"
               /note="myoglobin; G00-119-378"
               /number=1
intron        2060..>2552
               /gene="MB"
               /note="myoglobin intron A"
gap           2553..2652
               /estimated_length=unknown
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               /note="myoglobin mRNA"
intron        <2653..4066
               /gene="MB"
               /note="myoglobin intron A"
repeat region <2653..2873
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               /gene="MB"
               /note="9 bp direct repeat copy 1"
repeat region 3255..3386
               /gene="MB"
               /note="33 bp tandem repeat (4 copies)"
repeat region 3387..3395
               /gene="MB"
               /note="9 bp direct repeat copy 2"
exon          4067..4289
               /gene="MB"
               /note="G00-119-378"
               /number=2
intron        4290..>5323
               /gene="MB"
               /note="myoglobin intron B"
gap           5324..5423
               /estimated_length=unknown
prim transcript <5424..6336
               /note="myoglobin mRNA"
intron        <5424..5777
               /gene="MB"
               /note="myoglobin intron B"
exon          5778..>5924
               /gene="MB"
```

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```

                                /note="myoglobin; G00-119-378"
                                /number=3
ORIGIN      2 bp upstream of RsaI site.
1  gtactgtatt ttcatctctc ttagttatct ccctaaaaag actctgagtt ccttgaacac
61 aggaaggtgt tttatttgat tttgttatcc tcagcatgta gcagtgtctg acacacagta
121 ggtgctctat cactgtgaga gggatggatg gatgggtgga gttacagatg gatagaagga
181 tagatggagg gatgggtgga ttagtgatgg atagatggat ggagggggga tgatgaatgg
241 agggataatg agtggatgaa tgagggaaat ggtggatgga tggatggagg gatggaggaa
301 cagatagata gatggaggga tgggtgggtg atggatggat agatggatgg agggagggat
361 gatgaatgga gggataatga atggatgaat gaggggatgg gtggatggat gaatggaggg
421 atgatgggtg gatgaatgaa ttgagggatg gatggatgaa cacatggatg gatggataga
481 tggatagatg gaggaactgg tggatttttg atggatgggt ggatggatag atgaatgaat
541 gcctggatag acaaagagat gatggataga tgaatagatg aattaaggga tgtcggatag
601 atggagggat tgatagatgt tggatggatg ggtgggtggat ggatagatga gtgaatgcat
661 ggatagacaa agagatgatg gatggatgaa ttaagggatg acagatggat ggatggatga
721 gtaactggat ggacaagtgg ataaatggat agatgggtga atacctgaat ggattgaagg
781 aggatgcatg gatgtaagat aaggctaata atcctccact ctctttcttt gcaaaacat
841 ccaccattt actcaataaa catttattca gttcaaactt ggcacaaagc accatgtgag
901 gcccaagaga tacgtgggtt aataaaacag agctcctgcc ctctgaaaa ctgcaaagaa
961 aggggctgtg ctctctgagt tcaaattcca actctgccag cgactagctg tacatcagtg
1021 atgtttccct actttctctc aattaaatag ggataatgtc agtacctatc acattgggag
1081 gtcttgccgg gattaaatga gttaccaaat gccagtgtt tgggacaggg cctggcacc
1141 agcaaagtct ctgtgagtg ctggctgcta ttatcctaag ggagaagatg gcatgaaaac
1201 caggaaatag gatgccctt gggaagcaat gcaacaggaa cttacacaaa gaaaggaaag
1261 gaggaagcaa ttagtggtgt tcaaaaggag tatgtcaaga aaaacttttc agagggaaac
1321 ctttgagcag ggtcatgaaa acaggagttc tctaagagat tgtggacttg cttgggacca
1381 cctggctata agcacaaaac catccggttc ctttctgtca cttctggcgg gtgaggggtc
1441 tctggcaaag gggcagaagg tgcgtgagag gttgcgaatg gccaggactg tcctggggcc
1501 agccggggca cctgggtggcc aagcttagaa acatgacagg tcctcttggg agggctgacc
1561 gcagggagcg ttgggtttca ggctgctggc gtcggcttct gtggtgccct ttctgtcggc
1621 tatgagagtc cagacagtgc ccaacctcct ccccttcttt ccacacgcac aaccacccca
1681 cccctgtgtg cctgagctgt cctgcctcgc cacaatggca cctgccctaa aatagcttcc
1741 catgtgaggg ctagagaaag gaaaagatta gaccctccct ggatgagaga gagaaagtga
1801 aggagggcag gggaggggga cagcgagcca ttgagcgatc tttgtcaagc atcccagaag
1861 gtataaaaac gcccttgga ccaggcagcc tcaaacccca gctgttgggg ccaggacacc
1921 cagtgaagcc atacttgctc tttttgtctt cttcagactg cgccatgggg ctcagcgacg
1981 gggaatggca gttggtgctg aacgtctggg ggaaggtgga ggctgacatc ccaggccatg
2041 ggcaggaagt cctcatcagg taaaagggaag agattccatt gccctgcca cccacacct
2101 aagatcaagg gtgttcagct gcaaggtgga aagtttgcac gtggggtagg tcagttggct
2161 gcattagtta aggggtttag aacggtcact tgctttttct ttgcttttaa gtgtcaggga
2221 ttggactcag gagaggaaa ggagccattt caggctgatg tcagcagatt gaggaagcat
2281 gagaatcaaa cctaggatgc tcagagtcca ccaggaagaa ttttagaatt atagacagtc
2341 agagttaaca agggtcctga gagattttgt acagccacct ctcttacagg atgaggacaa
2401 aaagcgactg agaaggggag gacatttcca gagtcacagc tcattaaatg ctcttaaagt
2461 gtcaaggtta agacatgctc ttcaagggga gacagatctg gttctagact tggctctgcc
2521 actgagccac tgggtgacct ttgggaaggt ac
[ gap 100 bp] Expand Ns
2653 gaattctg agaattgctc aaaccagga ggtggagggt gcggtgagca
2701 gagattgcac cactgcactc cagcctgggc aacagagcca gattccatct caaaaaagaa
2761 aaacaaaaaa caaaaaagcc atgaactcat ttccaggttg aggagctcag catcctgggt
2821 gtgaaatacc ctctcataa aaccctggga tggagactac ggggatcagg tgcttccttg
2881 tgacaacttc tgggcatggt ggctcagggc gcaaactgga gtgtggccac aatacatact
2941 gtgtactttt acaaggatgt cacagagcct gggtatcata aaagaggagc ttttcaagga
```

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```

3001 actgaaacca ttagacagga gagagagccc tgggcagaca ggggtgcccg tgccaaacat
3061 ttcagctgtg gcacaaggga aagggtggga gttatgaaac tgttccattt tgggttttagg
3121 tctgggctct gccgctagct agccaagtga ccttgccac ttatctctgt ggtcttccat
3181 gagtaaaagg cggaaactca ctccctacca gagggcaggt ctgactccct ttaaccagca
3241 cccacctgct cacagcagga aggactgagg tctaaagctg gaggtgggca ggaaggactg
3301 aggtctaaag ctggaggtgg gcaggaagga ccgaggtcta aagctggagg tgggcaggaa
3361 ggaccgaggt ctaaagctgg agtggtgctgc tcagagtccc agcagaggcc tctggggcac
3421 ctcactgagt gcctggcagg agtgggtgcc tgtctcaggg ctgggttgag ttgctcccac
3481 caggaccctt cgtcatctgc acagtgagg gactgggagg ttcagagagt cacagcttgg
3541 gctcaaaaca agcaagaggt ttctgagtgt gaggattgct ctggagtggga atggccctca
3601 caggtaggag tgagcctcct gtagctagag gtattttaagc agctgaagga caatccctgg
3661 gcaggaagct gcagagatgg tcgcagcgtg gactagaact gctgttttgg tcaactcagac
3721 ctcatctcag cctggcttct ctggacagca cccctgcaat agtgagctgg tgacttttacg
3781 cctcagaacc tcggtttcta catctgtaaa atgggaatta tatgacactc actatgtgcc
3841 agacaccctg ttggtacata gcacacacta tctcacttaa tccttcaagt agggacaagt
3901 tatccccatc ctttatatga ggaagctgag gcacagagag gtgaagtga tggcccaagg
3961 tcacacagct gggaagacag ggagctaaac ttgaactcta gtctggctgc cccagacct
4021 cacaccgcac ctcccatgcc gactccagcc ttccctgtgc ccacaggctc tttaagggtc
4081 acccagagac tctggagaag tttgacaagt tcaagcacct gaagtccagag gacgagatga
4141 aggcattctga ggacttaaa aagcatggtg ccactgtgct caccgccctg ggtggcatcc
4201 ttaagaagaa ggggcatcat gaggcagaga ttaagccctt ggcacagtcg catgccacca
4261 agcacaagat ccccgtaag tacctggagg taggaggcag agcctgggca ggtgggagga
4321 tgcggggaag gcctcgggtg gggcaatggg atctgggttc gagtccaagc tcagccacta
4381 acttggtgga tgacctatgc cactcttctc tgtgccccag gtttctcatt tgtaaagggg
4441 actgccaccc actttgcctt cctcctggga ttgttgagaa tgaacacatt tagcattttt
4501 aatttagtat gccaaattca catcttatta ccaaagagga aaggagaggg ggaattggg
4561 tgcaaaattt gcatcctctc catgggtagg taccattatc atatccactt gatagatggg
4621 gaaactgagg ctcacagagg ttaagcagct tgtccacggt cacaggaggt ggataatggc
4681 agagccaaga ttcaaacgca ggtctctatt actacagaac cccagccctt aactgctgtg
4741 ccactgggag tctggtacat gcaggactta tgtggcagga gctcagcaag tggggctcaa
4801 tttgggggtg gggtgaccag caggctggct ctattgggtc cagcatcttc acagatgaag
4861 agacaggacc tcggtttcca gcacaagcaa ttggtttgaa cctcctgaga tgggttgga
4921 agttgggtg atcagggttg ggggcaggag cctgggcttc aggttggtg tctataactg
4981 gtgggaggag gcgatttggg gagaggagg agctggggat gaaggaccac agggacaggt
5041 gcatcccccg agggtagaaa cagcaggaag tctggtgcag ccatgaggat taggatgtgg
5101 tgatagctac ccgctgggat gggccacagt gagcatttgc tgccatgcct agcacatgca
5161 tccatcctca aagttgcctc atggccaaaa tgactgcaag agctccagcc agctcttcta
5221 tattccaac tggaagcagg agaaagagag gaatgtcttc ttttgaggag tttaaggagt
5281 cccagaaatc tcatccaaca attttattta catctcattg gcc
[ gap 100 bp] Expand Ns
5424 gagctct cacctgggtt cagtggggtc tacatcctga
5461 tggagtggag ggggctgtga gtaagagcgt gggctccgga gccggccctc ctgggtccaa
5521 atgtcccttc cattcaacct cccctgcct cagtttctgc atctgtaa atcggggcag
5581 tgtagtatct atctcacagt ggttggtggg atcaaagggg ttcattccgt gagatcacac
5641 agactctcac ctggtgccta gcaagtgtc aatacacggt cctggaataa agagaaggta
5701 ggaggacaac tgactcccat ctggcccttg gcttgctcca cctgggtgac cattttctct
5761 cctcaccctc cctgcagttc atctcggaat gcatcatcca ggttctgcag agcaagcatc
5821 ccggggactt tgggtctgat gccgagggg ccatgaacaa ggccctggag ctgttccgga
5881 aggacatggc ctccaactac aaggagctgg gcttccaggg ctaggccctt gccgctccca
5941 ccccccacca tctgggcccc gggttcaaga gagagcgggg tctgatctcg ttagccata
6001 tagagtttgc ttctgagtgt ctgctttgtt tagtagaggt gggcaggagg agctgagggg
6061 ctggggctgg ggtgttgaag ttggctttgc atgccagcg atgcgcctcc ctgtgggatg
6121 tcatcaccct gggaaccggg agtggccctt ggctcactgt gttctgcatg gtttggatct

```

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```
6181 gaattaattg tcctttcttc taaatcccaa ccgaacttct tccaacctcc aaactggctg
6241 taaccccaaa tccaagccat taactacacc tgacagtagc aattgtctga ttaatcactg
6301 gccccttgaa gacagcagaa tgtccctttg caatgaggag gagatctggg ctgggcgggc
6361 cagctgggga agcatttgac tatctggaac ttgtgtgtgc ctctcaggt atggcagtga
6421 ctcacctggg tttaataaaa caacctgcaa catctcagtt tctgcctggc atttttcatc
6481 tcctagagta aatgatgcc ccaccagcac cagcatcaag gaagaaatgg gaggaaggca
6541 gaccctgggc ttgtgtgtgc agagagcctc aggaaagagg agaaggggag gaggaaggc
6601 aggaggggta gagggacagg agcccaccct ccctgggcca ccgctcagag gcaggcccag
6661 tgcagggcat ggggaaatgg aagggacagg cttggcccca gccttgggag caccttctct
6721 tcgggggagg tgggaggcag cgaacagacc tctgcaatac gaggagagag tgacagggtgc
6781 gccaggctgt gggaaccag aggagagggg aagccatcat catcatggct gcaatacctt
6841 cagtaacgtg ggaaggtcac cctgctagta agtggcagag ctgggactc
```

//

Similarly, the nucleotide sequences from other databases can also be analyzed in a forthright manner.

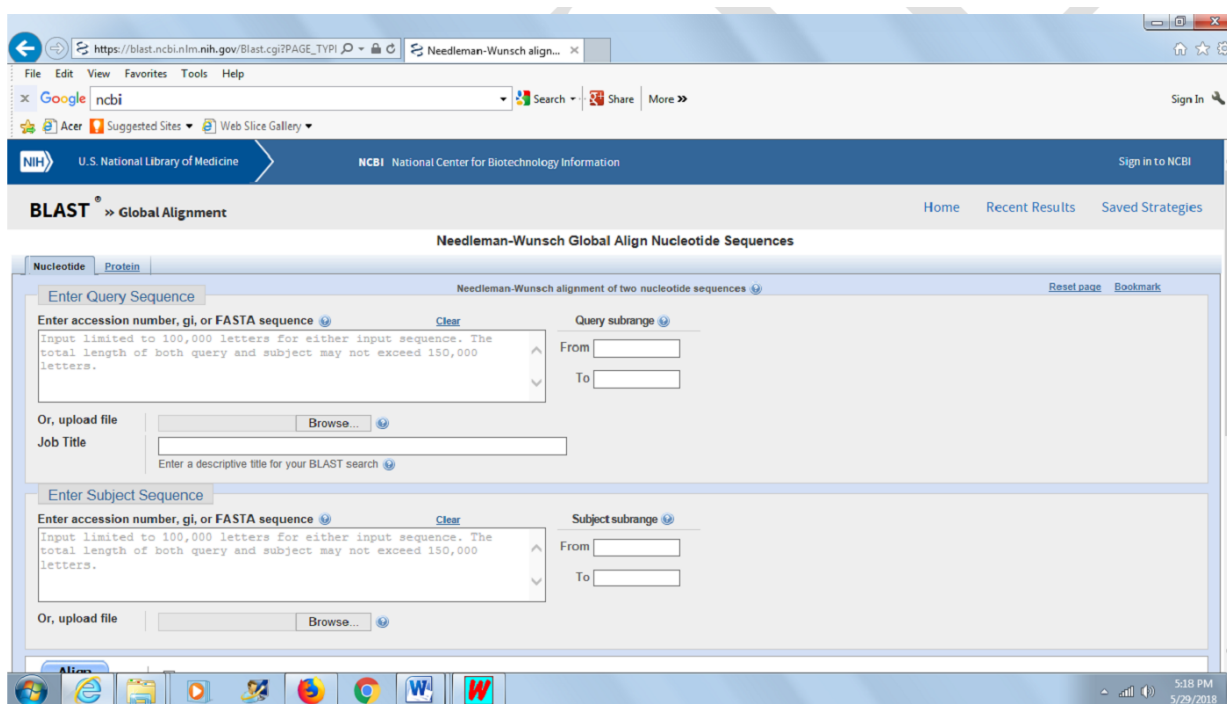
Results:

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Exp No: 6. Sequence alignment using BLAST

Aim: To examine pair-wise alignments of protein sequences using BLAST

Procedure: Retrieve the two protein sequences under interest either from primary or structural databases and save them in 'Fasta' format. Either the sequences or accession IDs of the sequences can be pasted in the space provided under categories of 'Query sequence' and 'Subject sequence' of the algorithm. Upon selecting appropriate alignment algorithm and gap penalty options, the program can be run for aligning the sequences.



For example, two cardiotoxins bearing PDB IDs of 1CRF and 1CVO were aligned and the results are shown below herein.

Seq2

Sequence ID: Query_206921Length: 62Number of Matches: 1

Related Information

Prepared by: Dr. Selvakumar, S. Assistant Professor, Department of Biotechnology, KAHE.

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Range 1: 1 to 62 [Graphics](#) [Next Match](#) [Previous Match](#) [First Match](#)

Alignment statistics for match #1

NW Score	Identities	Positives	Gaps	Frame
213	40/62(65%)	46/62(74%)	2/62(3%)	

Features:

Query 1 LKC-NKLVPLFYKTCFAGKNLCYKMFMS-NLTVPVKRGCIDVCPKNSALVKYVCCNTDR 58

LKC N +P YKTCF GKNLC+K + L PVKRG C D CPKNSAL+KYVCC+TD+

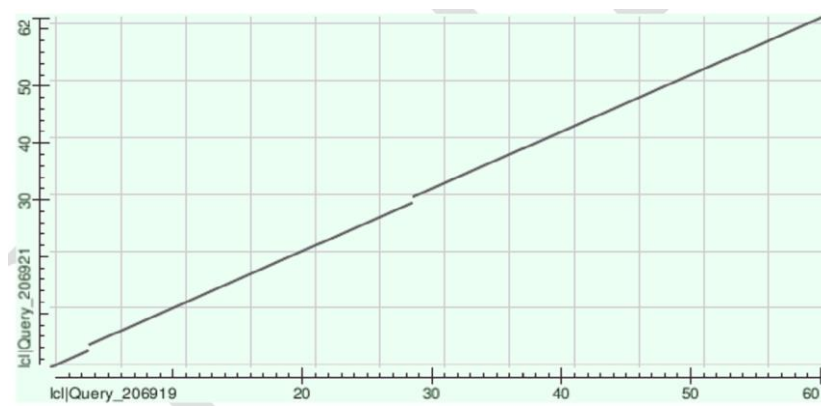
Sbjct 1 LKCHNTQLPFIYKTCFEGKNLCFKATLKKFPLKFPVKRGCADNCPKNSALLKYVCCSTDK 60

Query 59 CN 60

CN

Sbjct 61 CN 62

The dot plot for the sequences is depicted in the figure shown herein.



The sequence alignment can also be performed by using SmartBLAST and the results are as

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shown below herein.

Query: Seq1

Query length: 60 aa Identical to: 1CRE_A

DOMAIN: Snake toxin domain, present in short and long neurotoxins, cytot...

RecName: Full=Cytotoxin 1; AltName: Full=Cytotoxin I; Short=CTI
cytotoxin 13

RecName: Full=Cytotoxin 1; Short=NK-CT1

RecName: Full=Cytotoxin 2; AltName: Full=Cardiotoxin 1A; AltName: Full=Cardi...

Your query: Chain A, CARDIOTOXIN II FROM TAIWAN COBRA VENOM, NA...

RecName: Full=Cytotoxin 5b; AltName: Full=Cardiotoxin 5b; Short=CTX-5b; Sho...

Select: All None Selected:0

Description	Max score	Total score	Query cover	E value	Ident	Accession
RecName: Full=Cytotoxin 2; AltName: Full=Cardiotoxin 1A; AltName: Full=Cardiotoxin 2; Short=CTX-2; AltName: Full=Cardiotoxin A2; Short=CTX A2; AltName: Full=Card...	122	122	100%	9e-42	100%	P01442.2
RecName: Full=Cytotoxin 5b; AltName: Full=Cardiotoxin 5b; Short=CTX-5b; Short=Cto5b; Flags: Precursor	121	121	100%	9e-42	100%	P60310.1
RecName: Full=Cytotoxin 1; Short=NK-CT1	113	113	100%	1e-38	93%	P0CH80.1
cytotoxin 13 [Naia naia]	113	113	100%	2e-38	93%	BAU24670.1
RecName: Full=Cytotoxin 1; AltName: Full=Cytotoxin I; Short=CTI	110	110	100%	2e-37	88%	P01451.1

Results:

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Exp No: 7. Multiple sequence alignment using ClustalW/Clustal Omega

Aim: To examine multiple sequences alignment of protein sequences using Clustal Omega

Procedure: Clustal Omega is a new multiple sequence alignment program that uses seeded guide trees and HMM profile-profile techniques to generate alignments between three or more sequences. This tool can align up to 4000 sequences or a maximum file size of 4 MB.

The following 11 sequences were first retrieved from PDB database and the data were stored in 'Fasta' format.

```
>1CRF Naja Naja Atra
LKCENKLVPLFYKTCFAGKNLCYKMFVSNLTVPVKRGCIDVCPKNSALVKYVCCNTDRCN

>1CRE Naja Naja Atra
LKCENKLVPLFYKTCFAGKNLCYKMFVSNLTVPVKRGCIDVCPKNSALVKYVCCNTDRCN

>1CHV Naja Naja Atra
LKCENKLVPLFYKTCFAGKNLCYKMFVSNKMPVKRGCIDVCPKSSLLVKYVCCNTDRCN

>P60301.1
MKTLLLTLVVVTIVCLDLGYTLKCNKLVPLFYKTCFAGKNLCYKMFVATPKVPVKRGCIDVCPKSSLLV
KYVCCNTDRCN

>P80245.2
MKTLLLTLVVVTIVCLDLGYTLKCNQLIPPFYKTCAGKNLCYKMFVAAAPKVPVKRGCIDVCPKSSLLV
KYVCCNTDRCN

>P60304.1
MKTLLLTLVVVTIVCLDLGYTLKCNKLIPIASKTCFAGKNLCYKMFMSDLTIPVKRGCIDVCPKNSLLV
KYVCCNTDRCN

>1CXO
LKCENQLIPPFWKTCFAGKNLCYKMTMRAAPMPVKRGCIDVCPKSSLLIKYMCCNTDKCN

>1CXN
LKCENQLIPPFWKTCFAGKNLCYKMTMRAAPMPVKRGCIDVCPKSSLLIKYMCCNTDKCN

>P01442.2
MKTLLLTLVVVTIVCLDLGYTLKCNKLVPLFYKTCFAGKNLCYKMFVSNLTVPVKRGCIDVCPKNSALV
KYVCCNTDRCN

>1CVO
LKCHNTQLPFIYKTCPEGKNLCFKATLKKFPLKFPVKRGCADNCPKNSALLKYVCCSTDCKN

>P62375.1
MKTLLLTMTVVVTIVCLDLGYTLKCHNTQLPFIYKTCPEGKNLCFKATLKKFPLKFPVKRGCADNCPKNSA
LLKYVCCSTDCKN
```

All the 11 sequences were pasted one-by-one in the 'sequence space' provided in the Clustal

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Omega program. After setting appropriate alignment algorithm, matrix and output format, the program was executed and the outcomes are shown below herein'

CLUSTAL O(1.2.4) multiple sequence alignment

```

1CVO      -----
LKCHNTQLPFIYKTCPEGKNLCFKATLKKFPLKF
PVKRG 39 P62375.1
MKTLLLTMLVVVTIVCLDLGYTLKCHNTQLPFIYKTCPEGKNLCFKATLKKFPLKFPVKRG 60
1CXO      -----LKCNO-LIPPFWKTCPKGKNLCYKMTMRA-
APMPVKRG 37
1CXN      -----LKCNO-LIPPFWKTCPKGKNLCYKMTMRA-
APMPVKRG 37
P60301.1  MKTLLLTMLVVVTIVCLDLGYTLKCNK-LVPLFYKTCPAGKNLCYKMFMA-
TPKVPVKRG 58
P80245.2  MKTLLLTMLVVVTIVCLDLGYTLKCNQ-LIPPFYKTCAGKNLCYKMFMA-
APKVPVKRG 58
P60304.1  MKTLLLTMLVVVTIVCLDLGYTLKCNK-LIPIASKTCPAGKNLCYKMFMS-
DLTIPVKRG 58
1CRF      -----LKCNO-LVPLFYKTCPAGKNLCYKMFMS-
NLTVPVKRG 37
1CRE      -----LKCNO-LVPLFYKTCPAGKNLCYKMFMS-
NLTVPVKRG 37
P01442.2  MKTLLLTMLVVVTIVCLDLGYTLKCNK-LVPLFYKTCPAGKNLCYKMFMS-
NLTVPVKRG 58
1CHV      -----LKCNO-LVPLFYKTCPAGKNLCYKMFMS-
NKMVPVKRG 37

          ***::: * *** *****:* :
. *****

1CVO      CADNCPKNSALLKYVCCSTDKN 62
P62375.1  CADNCPKNSALLKYVCCSTDKN 83
1CXO      CIDVCPKSSLLIKYMCCNTDKN 60
1CXN      CIDVCPKSSLLIKYMCCNTDKN 60
P60301.1  CIDVCPKSSLLVKYVCCNTDRCN 81
P80245.2  CIDVCPKSSLLVKYVCCNTDRCN 81
P60304.1  CIDVCPKNSLLVKYVCCNTDRCN 81
1CRF      CIDVCPKNSALVKYVCCNTDRCN 60
1CER      CIDVCPKNSALVKYVCCNTDRCN 60
P01442.2  CIDVCPKNSALVKYVCCNTDRCN 81
1CHV      CIDVCPKSSLLVKYVCCNTDRCN 60
          * * ***. * *:***:***.***:***

```

Results: