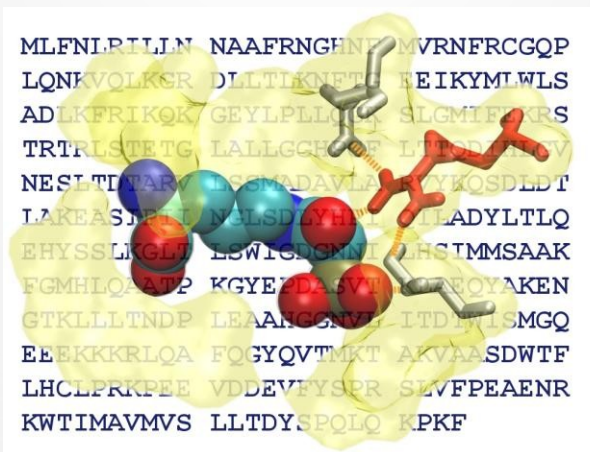
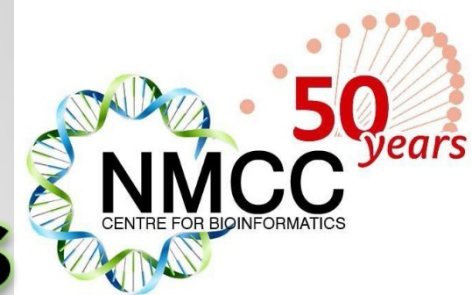




UGC SPONSORED NATIONAL WORKSHOP ON
BIOINFORMATICS AND GENOME ANALYSIS
FOR COLLEGE TEACHERS



LECTURE TOPIC: **PROTEIN DATABASE**

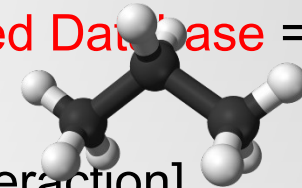
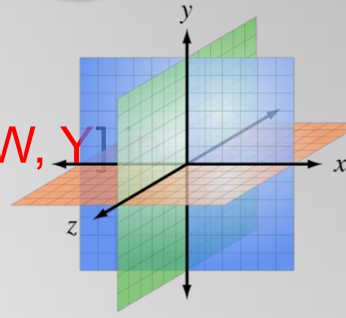
T. ASHOK KUMAR
HEAD, DEPARTMENT OF BIOINFORMATICS
NOORUL ISLAM COLLEGE OF ARTS AND
SCIENCE
KUMARACOIL, THUCKALAY - 629180

TOPICS COVERED

- **Protein Terms & Definitions** – Computational biology aspect of protein
- **ExPASy** – SIB Bioinformatics Resource Portal (<http://www.expasy.org>)
- **UniProt/Swiss-Prot** – A comprehensive, non-redundant, expert manually annotated protein sequence database (<http://www.uniprot.org/>)
- **NBRF/PIR** – A comprehensive, non-redundant, expertly manually annotated, fully classified and extensively cross-referenced protein sequence database (<http://pir.georgetown.edu/>)
- **PDB** – A single worldwide repository of information about the 3D structures of large biological molecules, including proteins and nucleic acids (<http://rcsb.org/pdb>)
- **SCOP** – Knowledge-based expert analysis and classification of proteins that are structurally characterized and deposited in the Protein Data Bank (<http://scop2.mrc-lmb.cam.ac.uk/>)
- **CATH** – A hierarchical domain classification of protein structures in the Protein Data Bank (<http://www.cathdb.info/>)
- **MOTIF** – Finds sequence motifs in a query sequence, also provides functional and genomic information of the found motifs using DBGET and LinkDB as the hyperlinked annotations (<http://www.genome.jp/tools/motif/>)
- **Pfam** – Database of protein HMM profiles that define domain families (<http://pfam.xfam.org/>)
- **PROSITE** – Database of protein motifs expressed as patterns or profiles (<http://prosite.expasy.org/>)

PROTEIN TERMS & DEFINITIONS

- Protein Sequence – 20 a.a. characters [A, C, D, E, F, G, H, I, K, L, M, N, P, Q, R, S, T, V, W, Y, sequence]
- Protein Structure – 3D of atomic co-ordinates [x-axis, y-axis, z-axis]
- Types of Biological Databases – [Raw Database = Plain text, Object-oriented Database = Table (Records), Relational Database = Table of tables]
- 3D Atom Model – [Sphere = Atom, Cylinder = Bond, Dotted Line = Bond Interaction]
- Sequence Alignment – [Match = Similar Character, Mismatch = Dissimilar Character, Gap = No Substitute Character, Word = Sub-string, Sequence = Super-string, Score = Rating, Identity = Similar in function]
- Motif – Short, conserved sequence associated with a distinct function.
- Domain – Evolutionarily conserved sequence region that corresponds to a structurally independent 3D unit associated with a particular functional role. It is usually much larger than a motif.
- Pattern – Sequence with symbol representation for a expression. Example: N{P}[ST]{P}
- Regular Expression – Representation format for a sequence motif, which includes positional information for conserved and partly conserved residues. Similar to Pattern, but applies to MSA.
- Profile – Scoring matrix that represents a multiple sequence alignment. It contains probability or



EXPASY

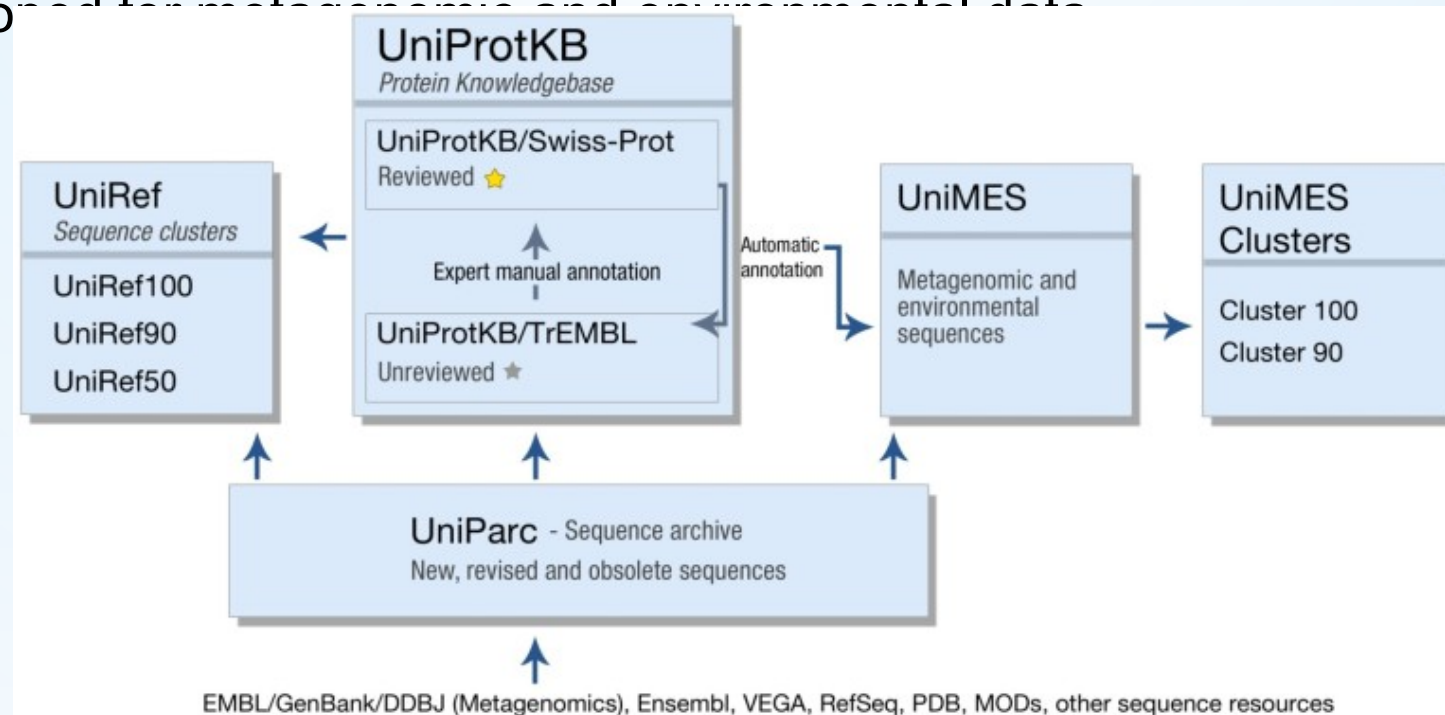
- ExPASy (**Ex**pert **Protein Analysis System**) is a bioinformatics resource portal operated by the Swiss Institute of Bioinformatics (SIB).
- ExPASy was the **first website of the life sciences**.
- Extensible and integrative portal for accessing many **scientific resources, databases** and **software tools**.
- Wide range of resources in many different domains, such as **proteomics, genomics, phylogeny/evolution, systems biology, population genetics, transcriptomics**, etc.
- **Proteomics server** to analyze **protein sequences and structures** and **2D Page gel electrophoresis**.
- **Databases, online and offline software tools** are hosted by different groups of the **SIB and partner institutions**. --- CFSSP
- ExPASy references the **proteome knowledgebase, UniProtKB/Swiss-Prot, and its computer annotated supplementary database, UniProtKB/Trembl**.



Resource Portal

ARCHITECTURE OF UNIPROT/SWISS-PROT

- Universal Protein Resource (UniProt) is a comprehensive resource for **protein sequence** and **annotation data**
- The UniProt databases are the UniProt Knowledgebase (**UniProtKB**), the UniProt Reference Clusters (**UniRef**), and the UniProt Archive (**UniParc**)
- UniProt Metagenomic and Environmental Sequences (UniMES) database is a repository specifically developed for metagenomic and environmental data



BACKGROUND OF UNIPROT/SWISS-PROT

- UniProt is a collaboration between the European Bioinformatics Institute (~~EMBL-EBI~~), the Swiss Institute of Bioinformatics (~~SIB~~) and the Protein Information Resource (~~PIR~~)
- EMBL-EBI and SIB together used to produce **Swiss-Prot** and **TrEMBL**, while PIR produced the Protein Sequence Database (~~PIR-PSD~~)
- Translated EMBL Nucleotide Sequence Data Library (~~TrEMBL~~) was originally created because sequence data was being generated at a pace that exceeded Swiss-Prot's ability to keep up
- PIR maintained the PIR-PSD and related databases, including iProClass, a database of protein sequences and curated families



<http://www.uniprot.org/>

| | | | | |
|-------------------------------------|-------|--------|-------------------|------------|
| Search | Blast | Align | Retrieve | ID Mapping |
| Search in | | Query | | |
| Protein Knowledgebase (UniProtKB) ▾ | | serpin | | |
| | | Search | Advanced Search » | Clear |

WELCOME

The mission of UniProt is to provide the scientific community with a comprehensive, high-quality and freely accessible resource of protein sequence and functional information.

What we provide

| | |
|-----------------|--|
| UniProtKB | Protein knowledgebase, consists of two sections: <ul style="list-style-type: none"> ★ Swiss-Prot, which is manually annotated and reviewed. ★ TrEMBL, which is automatically annotated and is not reviewed. Includes complete and reference proteome sets . |
| UniRef | Sequence clusters, used to speed up sequence similarity searches. |
| UniParc | Sequence archive, used to keep track of sequences and their identifiers. |
| Supporting data | Literature citations , taxonomy , keywords , subcellular locations , cross-referenced databases and more. |

Getting started

- [Text search](#)
- [Sequence similarity searches \(BLAST\)](#)
- [Sequence alignments](#)
- [Batch retrieval](#)
- [Database identifier mapping \(ID Mapping\)](#)



TRY THE NEW UNIPROT WEBSITE



NEWS



UniProt release 2014_07 - Jul 9, 2014

Lark or owl? PER3 is the answer | Cross-references to CCDS and GeneReviews | UniParc cross-references with protein and gene names

- › [Statistics for UniProtKB: Swiss-Prot · TrEMBL](#)
- › [Forthcoming changes](#)
- › [News archives](#)

Follow @uniprot 1,018 followers

SITE TOUR



Learn how to make best use of the tools and data on this site.

PROTEIN SPOTLIGHT

it did it its way July 2014

Temperatures can get cold. And living organisms have to find ways of keeping themselves warm. Humans use clothes. Polar bears grow fur. Whales are lined with blubber. And many animals avoid the cold by migrating to warmer parts of the planet...

Search

Blast

Align

Retrieve

ID Mapping *

Search in

Protein Knowledgebase (UniProtKB) ▾

Query

serpin

Search

Advanced Search »

Clear

1 - 25 of 8,282 results for **serpin** in UniProtKB sorted by **score** descending ☒

Browse by taxonomy, keyword, gene ontology, enzyme class or pathway | Reduce sequence redundancy to 100%, 90% or 50%

Download

Page 1

of 332 | Next »

Results Customize

Select the box to download individual sequence

Click the link to download entire matching sequences

- › Show only reviewed (311) (UniProtKB/Swiss-Prot) or unreviewed (7,971) (UniProtKB/TrEMBL) entries
- › Restrict term "serpin" to protein family (6,881), gene name (93), protein name (2,342), web resource (1)

| Entry | Entry name | Status | Protein names | Gene names | Organism | Length |
|--|--------------|--------|-----------------------------------|--|--|--------|
| <input checked="" type="checkbox"/> Q9V3N1 | Q9V3N1_DROME | ★ | LD24467p | Spn27A serpin-27A Spn27A-RA CG11331 Dmel_CG11331 | Drosophila melanogaster (Fruit fly) | 447 |
| <input type="checkbox"/> P05120 | PAI2_HUMAN | ★ | Plasminogen activator inhibitor 2 | SERPINB2 PAI2 PLANH2 | Homo sapiens (Human) | 415 |
| <input type="checkbox"/> Q5I2A0 | SPA3G_MOUSE | ★ | Serine protease inhibitor A3G | Serpina3g Spi2A | Mus musculus (Mouse) | 440 |
| <input type="checkbox"/> Q8IW75 | SPA12_HUMAN | ★ | Serpin A12 | SERPINA12 | Homo sapiens (Human) | 414 |
| <input type="checkbox"/> Q9UIV8 | SPB13_HUMAN | ★ | Serpin B13 | SERPINB13 PI13 | Homo sapiens (Human) | 391 |
| <input type="checkbox"/> Q9S7T8 | SPZX_ARATH | ★ | Serpin-ZX | At1g47710 F16N3.3 T2E6.22 | Arabidopsis thaliana (Mouse-ear cress) | 391 |
| <input type="checkbox"/> D6WWC4 | D6WWC4_TRICA | ★ | Serpin peptidase inhibitor 23 | serpin23 TcasGA2_TC005771 | Tribolium castaneum (Red flour beetle) | 385 |

1 selected: Q9V3N1

Serpin peptidase inhibitor 31

serpin31 TcasGA2_TC015224

Retrieve

Align

Blast

Clear

Search Blast Align Retrieve ID Mapping

UniProt identifiers or file

Q9V3N1

Choose File No file chosen

Retrieve

Clear

Help
Enter or upload a list of UniProt identifiers to download the corresponding entries, e.g.
P00750
A4_HUMAN
[More...](#)

1 unique item available for download

UniProtKB (1)

Download data [compressed](#) or [uncompressed](#)

FASTA

Sequence data in FASTA format.

[\[Download | Open \]](#)

GFF

Sequence features in GFF

[\[Download | Open \]](#)

Flat Text

Complete data in the original flat text format.

[\[Download | Open \]](#)

XML

Complete data in XML format.

[\[Download | Open \]](#)

RDF/XML

Complete data in RDF format.

[\[Download | Open \]](#)

List

List of identifiers.

[\[Download | Open \]](#)

Click the link to download the sequence to disk in UniProt file format

UNIPROT/SWISS-PROT FILE FORMAT

| Line code | Content | Occurrence in an entry |
|-----------|------------------------------|--|
| ID | Identification | Once; starts the entry |
| AC | Accession number(s) | Once or more |
| DT | Date | Three times |
| DE | Description | Once or more |
| GN | Gene name(s) | Optional |
| OS | Organism species | Once or more |
| OG | Organelle | Optional |
| OC | Organism classification | Once or more |
| OX | Taxonomy cross-reference | Once |
| OH | Organism host | Optional |
| RN | Reference number | Once or more |
| RP | Reference position | Once or more |
| RC | Reference comment(s) | Optional |
| RX | Reference cross-reference(s) | Optional |
| RG | Reference group | Once or more (Optional if RA line) |
| RA | Reference authors | Once or more (Optional if RG line) |
| RT | Reference title | Optional |
| RL | Reference location | Once or more |
| CC | Comments or notes | Optional |
| DR | Database cross-references | Optional |
| PE | Protein existence | Once |
| KW | Keywords | Optional |
| FT | Feature table data | Once or more in Swiss-Prot, optional in TrEMBL |
| SQ | Sequence header | Once |
| (blanks) | Sequence data | Once or more |
| // | Termination line | Once; ends the entry |


NBRF/PIR

- The Protein Information Resource (**PIR**) was established in 1984 by the National Biomedical Research Foundation (**NBRF**) as a resource to assist researchers in the identification and interpretation of protein sequence information.
- In 2002 PIR, along with its international partners, EBI and SIB, were awarded a grant from NIH to create UniProt, by unifying the PIR-PSD, Swiss-Prot, and TrEMBL databases.
- As of 2010, PIR offers a wide variety of resources mainly oriented to assist the propagation and standardization of protein annotation: PRO, iProClass, iProLINK.

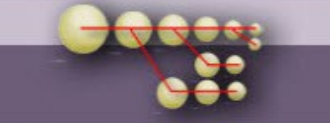


INTEGRATED PROTEIN INFORMATICS RESOURCE FOR
GENOMIC, PROTEOMIC AND SYSTEMS BIOLOGY RESEARCH

Click the
button

 The Universal Protein Resource (UniProt) provides the scientific community with a single, centralized, authoritative resource for protein sequences and functional information.
[UniProtKB](#) | [UniRef](#) | [UniParc](#) Current release: 2014_07


PRO
Protein Ontology



- Representation of protein objects with descriptions and relationships
- [Browse PRO](#)
- Annotate with [RACE-PRO](#)

[*Sample PRO report*](#)

iProClass
Integrated Protein Knowledgebase



- Value-added reports for [UniProtKB](#) and unique [UniParc](#) proteins
- Functional analysis and [protein ID mapping](#)

[*Sample protein report*](#)

iProLINK
Literature Information & Knowledge



- Source for text mining and ontology development
- [RLIMS-P](#) text mining tools
- [Bibliography mapping](#)

[*Sample Biblio. report*](#)

O OTHER RESOURCE

- [Representative Proteomes](#)
- [iProXpress](#)
- [iPTMnet](#)

P PEPTIDE SEARCH ?

DATABASE: UniProtKB

Use single letter amino acid code

T TEXT SEARCH ?

DATABASE: iProClass

NEW Bioinformatics & Computational Biology Graduate Programs:

- [MS program at Georgetown University](#)
- [MS, PSM and Graduate Certificate programs at University of Delaware](#)
 - Now recruiting new students for the year of 2014
 - Scholarships available

Text Search Result

ALL Fields
AND
ALL Fields
AND
ALL Fields
+ add input box
- del input box

Click the link to download the sequence

Display Options Help?

3015 proteins | 302 pages | 10 / page | <<< 1 | 2 | 3 | 4 | 5 >>>

Save Result As: TABLE FASTA

1 selected (show) GO Slim / Pathway... Pairwise Alignment Multiple Alignment Domain Display

| <input type="checkbox"/> Protein AC/ID | Protein Name | Length | Organism Name | PRO ID | PIRSF ID | UniRef50 |
|---|---|--------|--------------------------------------|--|---|---------------------------------|
| <input checked="" type="checkbox"/> F7VJQ1/APRIO_HUMAN <small>/ProClass UniProtKB/Swiss-Prot</small> | Alternative prion protein | 73 | Homo sapiens (Human) | | | UniRef50 F7VJQ1 |
| <input type="checkbox"/> O15394/NCAM2_HUMAN <small>/ProClass UniProtKB/Swiss-Prot</small> | Neural cell adhesion molecule 2 precursor | 837 | Homo sapiens (Human) | PR:000011018; PR:O15394 | PIRSF002507; PIRSF501037 | UniRef50 O15394 |
| <input type="checkbox"/> O43414/ERI3_HUMAN <small>/ProClass UniProtKB/Swiss-Prot</small> | ERI1 exoribonuclease 3 | 337 | Homo sapiens (Human) | PR:000013260; PR:O43414 | | UniRef50 O43414 |
| <input type="checkbox"/> P00441/SODC_HUMAN <small>/ProClass UniProtKB/Swiss-Prot</small> | Superoxide dismutase [Cu-Zn] | 154 | Homo sapiens (Human) | PR:000015399; PR:P00441 | PIRSF000348 | UniRef50 P00441 |
| <input type="checkbox"/> P01031/CO5_HUMAN <small>/ProClass UniProtKB/Swiss-Prot</small> | Complement C5 precursor | 1676 | Homo sapiens (Human) | PR:000004904; PR:P01031 | PIRSF001635 | UniRef50 P01031 |
| <input type="checkbox"/> P01583/IL1A_HUMAN <small>/ProClass UniProtKB/Swiss-Prot</small> | Interleukin-1 alpha precursor | 271 | Homo sapiens (Human) | PR:000001135; PR:P01583 | PIRSF001937 | UniRef50 P01583 |
| <input type="checkbox"/> P01584/IL1B_HUMAN <small>/ProClass UniProtKB/Swiss-Prot</small> | Interleukin-1 beta precursor | 269 | Homo sapiens (Human) | PR:000001136; PR:P01584 | PIRSF001937 | UniRef50 P01584 |
| <input type="checkbox"/> P02745/C1QA_HUMAN <small>/ProClass UniProtKB/Swiss-Prot</small> | Complement C1q subcomponent subunit A precursor | 245 | Homo sapiens (Human) | PR:000004875; PR:P02745 | PIRSF002477 | UniRef50 P02745 |
| <input type="checkbox"/> P02746/C1QB_HUMAN <small>/ProClass UniProtKB/Swiss-Prot</small> | Complement C1q subcomponent subunit B precursor | 253 | Homo sapiens (Human) | PR:000004876; PR:P02746 | PIRSF002477 | UniRef50 P02746 |
| <input type="checkbox"/> P02747/C1QC_HUMAN <small>/ProClass UniProtKB/Swiss-Prot</small> | Complement C1q subcomponent subunit C precursor | 245 | Homo sapiens (Human) | PR:000004878; PR:P02747 | PIRSF002477 | UniRef50 P02747 |

3015 proteins | 302 pages | 10 / page | <<< 1 | 2 | 3 | 4 | 5 >>>



ProClass Summary Report for UniProtKB Entry: F7VJQ1

[BioThesaurus](#) [ID Mapping](#)

GENERAL INFORMATION

| | | | |
|---|--|---------------------|---------------------------|
| Protein Name and ID | UniProtKB ID | UniProtKB Accession | Protein Name |
| | APRIO_HUMAN | F7VJQ1 | Alternative prion protein |
| RefSeq: NP_001258490.1 GenPept: DAA34790.1 IPI: IPI01026282 | | | |
| Taxonomy | Source Organism: Homo sapiens (Human) Taxon Group: Euk/mammal NCBI Taxon: 9606 Lineage: Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Primates; Haplorrhini; Catarrhini; Hominidae; Homo. | | |
| Keywords | complete proteome; membrane; mitochondrion; mitochondrion outer membrane; reference proteome; transmembrane; transmembrane helix | | |
| Function | Involved in the secretion of alkaline protease. | | |
| Tissue Specificity | Detected in brain homogenate, primary neurons, and peripheral blood mononuclear cells (at protein level). | | |

CROSS-REFERENCES

| | |
|-----------------|--|
| Bibliography | <p>View Bibliography Information</p> <p>Annotated references: PMID: 11120925; 11220690; 11244488; 11559357; 11584448; 11593450; 11684342; 11704923; 11756421; 11775001; 11787070; 11833672; 11840201; 11961239; 11986958; 12070046; 12084159; 12161431; 12186633; 12205650 [GeneRIF] More</p> <p>Other references: PMID: 16004966; 2180366; 16294306; 7913755; 10970892; 1683708; 8105771; 1736177; 1357663; 12679034; 2564168; 12514748; 12356908; 17500595; 11438139; 18051367; 1351748; 9786248; 10988071; 1363802; More</p> |
| DNA Sequence | EMBL: AL133396; BK007887 DDBJ: AL133396; BK007887 GenBank: AL133396; BK007887 |
| Genome/Gene | Gene Name: PRNP; prion protein Locus Tag: RP5-1068H6.2; Synonyms: ASCR; AltPrP; CD230; CID; GSS; KURU; PRIP; PrP; PrP27-30; PrP33-35C; PrPc; p27-30; Map Location: 20p13 Entrez Gene: 5621 UniGene: Hs.472010; Hs.610285; Hs.721670 RefSeq: NM_000311.3 NP_000302.1 [Map Viewer]; NM_001080121.1 NP_001073590.1 [Map Viewer]; NM_001080122.1 NP_001073591.1 [Map Viewer]; NM_001080123.1 NP_001073592.1 [Map Viewer]; NM_001271561.1 NP_001258490.1 [Map Viewer]; NM_183079.2 NP_898902.1 [Map Viewer] UCSC: uc002wkw.4 human KEGG: hsa:5621 |
| Gene Expression | GEO |
| Ontology | Gene Ontology: Cellular Component GO:0005739:mitochondrion [GO_REF:0000037; evidence:IEA] GO:0005741:mitochondrial outer membrane [GO_REF:0000037; evidence:IEA][GO_REF:0000039; evidence:IEA][PMID:21478263; evidence:IDA] GO:0016021:integral component of membrane [GO_REF:0000037; evidence:IEA] GO:0016020:membrane [GO_REF:0000037; evidence:IEA] |
| Structure | ProteinModelPortal: F7VJQ1 |
| Other Databases | neXtProt: NX_F7VJQ1 NextBio: 35510517 CTD: 5621 GenomeRNAi: 5621 GeneWiki: PRNP |

FAMILY CLASSIFICATION

| | |
|----------------------|--|
| UniRef | UniRef100_F7VJQ1; UniRef90_F7VJQ1; UniRef50_F7VJQ1 |
| Other Classification | KO: K05634 |

FEATURE & SEQUENCE DISPLAY

Length = 73
 F7VJQ1

Click on a bar to show its sequence; to copy and paste it, press ctrl-c then ctrl-v.

1
 61

MEHNGQPIFGAGQPMRQFLPFTSGRWULGAASUUULGAAPUUULGTASUUULG
 SRRWHFQSVQAE

SEQUENCE RETRIEVED FROM NBRF/PIR IN FASTA FILE FORMAT

```
>F7VJQ1 APRIO_HUMAN Alternative prion protein [Homo  
sapiens]  
MEHWGQPIPGAGQPWRQPLPTSGRWWLGAASWWLGAASWWLGAAPWWLGTASWWL  
G  
SRRWHPQSVEQAE
```

PDB

- The Protein Data Bank (**PDB**) archive is the single worldwide repository of information about the 3D structures of large biological molecules, including proteins and nucleic acids.
- The PDB was established in 1971 at Brookhaven National Laboratory (**BNL**) under the leadership of Walter Hamilton and originally contained 7 structures.
- In 1998, the Research Collaboratory for Structural Bioinformatics (**RCSB**) became responsible for the management of the PDB.
- In 2003, the **wwPDB** was formed to maintain a single PDB archive of macromolecular structural data that is freely and publicly available to the global community.
- The RCSB PDB supports a website where visitors can perform simple and complex queries on the data, analyze, and visualize the results.
- Members of wwPDB are: **RCSB PDB** (USA), **PDB_e** (Europe) and **PDB_j** (Japan), and Biological Magnetic Resonance Data Bank **BMRB** (USA).

<http://rcsb.org/pdb/>



Search
Advanced
Browse

Everything Author Macromolecule Sequence Ligand ?

Chemokine

Search History (1), Previous Results (138)

Customize This Page

Available on the
App Store

ANDROID APP ON
Google play

PDB-101 Show

MyPDB Show

Home Show

Deposition Show

Tools Show

Help Show

Biological Macromolecular Resource

Full Description

Explore Archive Hide

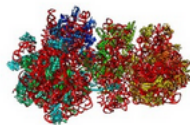
| | |
|-----------------------|-----------------------|
| Organism | Taxonomy |
| Exp. Method | X-ray Resolution |
| Release Date | Polymer Type |
| Enzyme Classification | SCOP Classification |
| Protein Symmetry | Protein Stoichiometry |
| Membrane Proteins | |

Organism

- Homo sapiens (26245)
- Escherichia coli (4855)
- Mus musculus (4419)
- Saccharomyces cerevisiae (2321)
- Bos taurus (2321)
- Rattus norvegicus (215)
- Escherichia coli K-12 (256239)
- Other (56239)

Show all

Latest Structures Hide



4TP8 : Crystal structure of the E. coli ribosome bound to dalfopristin and quinupristin. This file contains the 30S subuni...

Noeske, J., Huang, J., Olivier, N.B., Giacobbe, R.A., Zambrowski, M., Cate, J.H.

Synergy of streptogramin antibiotics occurs independently of their effects on translation.

View in 3D (Jmol)

check out all of the latest structures released.

New user? Try the browser [Compatibility Check](#) and view information on [Getting Started](#).

New Features Hide

Latest release:
April 2014



Browser
Compatibility Check

Website Release Archive: ▾

New Structures Hide

Latest Release
New Structure Papers
Search Unreleased Entries

wwPDB News Hide

2014-08-01
wwPDB Events at IUCr
(August 5-12)

- Improved Representation of Large Structures in the PDB Archive
- Full wwPDB News
- Statement on Retraction of PDB Entries

FTP Archives Hide

Current PDB FTP Archive:
<ftp://wwpdb.org>

Yearly FTP Snapshots:
<ftp://snapshots.wwpdb.org>

Search
 Advanced
 Browse

Everything Author Macromolecule Sequence Ligand ?
 e.g., PDB ID, molecule name, author

Search History (2), Previous Results (138)

↑ PDB-101 Hide

Structural View of Biology
 Understanding PDB Data
 Molecule of the Month
 Educational Resources
 Author Profiles

↑ MyPDB Hide

Login to your Account
 Register a New Account
 MyPDB Help Page
 Query Results (138)
 Query History (2)

↑ Home Hide

News & Publications
 Usage/Reference Policies
 Deposition Policies
 Website FAQ
 Deposition FAQ
 Contact Us
 About Us
 Careers
 External Links
 New Website Features

↑ Deposition Hide

All Deposit Services
 Electron Microscopy
 X-ray | NMR
 Validation Server
 BioSync Beamlines/Facilities
 Related Tools

↑ Tools Hide

Download Files
 Compare Structures
 Drug & Drug Target
 Mapping
 File Formats
 RESTful Web Services
 Widgets

↑ Help Hide

Launch Help System
 Display Settings
 Video Tutorials
 Glossary of Terms
 RCSB PDB Mobile

138 Structure Hits 7 Unreleased Structures 87 Citations 54 Ligand Hits 2 Web Page Hits

Query Parameters:

Text Search for: chemokine

Other search suggestions:

Query Details | Save Query to MyPDB

Query Refinements: Select an item or pie chart ?

Show

Organism
 Taxonomy
 Exp. Method
 X-ray Resolution
 Release Date
 Polymer Type
 Enzyme Classification
 SCOP Classification
 Protein Symmetry
 Protein Stoichiometry
 Membrane Proteins

Refine Query with Advanced Search

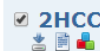
Show only representatives at sequence identity

Showing 1 - 2 of 138 Results

Results : Page: of 69

Filter: Remove Unc View: Detailed

Reports: Select one... Sort: Relevance



SOLUTION STRUCTURE OF THE HUMAN CHEMOKINE HCC-2, NMR, 30 STRUCTURES

Authors: Sticht, H., Escher, S.E., Schweimer, K., Forssmann, W.G., Roesch, P., Adermann, K.

Release: 1999-07-13

Experiment: SOLUTION NMR

Residue Count 66

Compound: 1 Polymer [[Display Full Polymer Details](#) | [Display for All Results](#)]

Citation: **Solution structure of the human CC chemokine 2: A monomeric representative of the CC chemokine subtype.**
 (1999) *Biochemistry* **38**: 5995-6002 [[Display Full Abstract](#) | [Display for All Results](#)]

Search Hit: Classification: CHEMOKINE



PROTON NMR ASSIGNMENTS AND SOLUTION CONFORMATION OF RANTES, A CHEMOKINE OF THE CC TYPE

Authors: Skelton, N.J.

Release: 1995-06-03

Experiment: SOLUTION NMR

Residue Count 136

Compound: 1 Polymer [[Display Full Polymer Details](#) | [Display for All Results](#)]

Citation: **Proton NMR assignments and solution conformation of RANTES, a chemokine of the C-C type.**
 (1995) *Biochemistry* **34**: 5329-5342 [[Display Full Abstract](#) | [Display for All Results](#)]

Search Hit: Classification: CHEMOKINE

1 2 3 4 5 6 7 8 9 10 Next Last

| SECTION | DESCRIPTION | RECORD TYPE |
|---------------------------|---|--|
| Title | Summary descriptive remarks | HEADER, OBSLTE, TITLE, SPLIT, CAVEAT, COMPND, SOURCE, KEYWDS, EXPDTA, NUMMDL, MDLTYP, AUTHOR, REVDAT, SPRSDE, JRNL |
| Remark Annotations | Various comments about entry in more depth than standard records | REMARKs 0-999 |
| Primary structure | Peptide and/or nucleotide sequence and the relationship between the PDB sequence and that found in the sequence database(s) | DBREF, SEQADV, SEQRES MODRES |
| Heterogen | Description of non-standard groups | HET, HETNAM, HETSYN, FORMUL |
| Secondary structure | Description of secondary structure | HELIX, SHEET |
| Connectivity annotation | Chemical connectivity | SSBOND, LINK, CISPEP |
| Miscellaneous features | Features within the macromolecule | SITE |
| Crystallographic | Description of the crystallographic cell | CRYST1 |
| Coordinate transformation | Coordinate transformation operators | ORIGXn, SCALEn, MTRIXn, |
| Coordinate | Atomic coordinate data | MODEL, ATOM, ANISOU, TER, HETATM, ENDMDL |
| Connectivity | Chemical connectivity | CONNECT |
| Bookkeeping | Summary information, end-of-file marker | MASTER, END |

SCOP

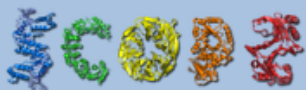
- The Structural Classification of Proteins (**SCOP**) database is a largely manual classification of protein structural domains **based on similarities of their structures and amino acid sequences**.
- A motivation for this classification is to **determine the evolutionary relationship between proteins**.
- Proteins with the **same shapes but having little sequence or functional similarity** are placed in different "superfamilies", and are assumed to have only a very **distant common ancestor**.
- Proteins having the **same shape and some similarity of sequence and/or function** are placed in "families", and are assumed to have a closer common ancestor.
- SCOP has been discontinued and the last official version of SCOP is **1.75**. [SCOP1.75](#) is also known as [SCOP2](#).
- SCOP2 offers two different ways for accessing data: **SCOP2-browser**, and **SCOP2-graph**.
- SCOP2-browser allows navigation in a traditional way by **browsing pages displaying the node information**.
- SCOP2-graph is a **graph-based web tool** for display and navigation.
- The source of protein structures is the Protein Data Bank.

HIERARCHICAL STRUCTURE OF SCOP

- The unit of classification of structure in SCOP is the **protein domain**.
- The levels of SCOP are as follows.
 1. **Class**: Types of folds, e.g., all α , all β , α/β , $\alpha+\beta$, $\alpha&\beta$, etc.
 2. **Fold**: The **different shapes of domains** within a class, e.g., 2 helices; antiparallel hairpin, left-handed twist, etc.
 3. **Superfamily**: The domains in a fold are grouped into superfamilies, which have **at least a distant common ancestor**.
 4. **Family**: The domains in a superfamily are grouped into families, which have **more recent common ancestor**.
 5. **Protein domain**: The domains in families are grouped into protein domains, which are **essentially the same protein**.
 6. **Species**: The domains in "protein domains" are **grouped according to species**.
 7. **Domain**: It is **part of a protein**. For simple proteins, it can be the entire protein.



<http://scop2.mrc-lmb.cam.ac.uk/>



News

November, 2013

During the development of SCOP2, we have identified a new, previously unrecognised type of alpha-alpha superhelix. Unlike other alpha-alpha superhelices..
[More...](#)

January, 2014

SCOP2 article in NAR is published
[More...](#)

January, 2014

The structure of the month
[More...](#)

Welcome to SCOP2!

Citation

Antonina Andreeva, Dave Howorth, Cyrus Chothia, Eugene Kulesha, Alexey Murzin, SCOP2 prototype: a new approach to protein structure mining (2014) Nucl. Acid Res., 42 (D1): D310-D314. [[PDF](#)]

Description of the SCOP2 database

SCOP2 is a successor of Structural classification of proteins ([SCOP](#)). Similarly to SCOP, the main focus of SCOP2 is on proteins that are structurally characterized and deposited in the PDB. Proteins are organized according to their structural and evolutionary relationships, but, in contrast to SCOP, instead of a simple tree-like hierarchy these relationships form a complex network of nodes. Each node represents a relationship of a particular type and is exemplified by a region of protein structure and sequence.

In SCOP2, we try to put in use the knowledge we acquired over the past years and the lessons we have learned during the classification of protein structures. We believe that there are many peculiarities of proteins and their structures that have been missed due to the constraints of the original SCOP hierarchical schema. We hope that our users will find the new resource useful and that it could open new avenues for protein analysis and research.

Quick introduction on how to browse, search and download

SCOP2 offers two different ways for accessing data: [SCOP2-browser](#), that allows navigation through the SCOP2 classification in a traditional way by browsing pages displaying the node information, and [SCOP2-graph](#), which is a graph-based web tool for display and navigation through the SCOP2 classification. Both tools provide search of SCOP2 data by free text, node names, IDs, tags and keywords, as well as external identifiers associated with them, e.g. PDB and UniProt. SCOP2 data can also be retrieved via [REST interface](#) or downloaded from the [SCOP2 Download page](#). For more information visit the [About](#) page.

Web browser compatibility check

To test whether your web browser and its settings are suitable to view SCOP2-graph and to visualize protein structures using Jmol applet [click here](#).

Information about the release

This is a beta release and contains a fraction of protein structure data. We wish to introduce the new database to our users and seek for a comprehensive feedback that would guide us for the future database development and expansion. We plan to regularly update the site with new classification data.

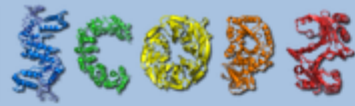
Search Browser

Add an asterisk to search free text (e.g. serine*)

Search Graph

Add an asterisk to search free text (e.g. protein*domain)

OUTPUT OF SCOP



Structural Classification of Proteins 2

[About](#) | [Browser](#) | [Graph](#) | [Download](#) | [Support](#)

MRC

Laboratory of
Molecular Biology

Invertase inhibitor (5001532)

Add an asterisk to search free text (e.g. serine*)

[Protein Relationships](#) | [Structural Classes](#) | [Protein Types](#) | [Evolutionary Events](#) | [Keywords](#)

Parents

- family Plant invertase/pectin methylesterase inhibitor (4000644)
[Pfam 04043](#)
Derived from [SCOP 101149](#)

Protein

[Go to Graph](#)

protein Invertase inhibitor (5001532)
Derived from [SCOP 101150](#)

Children

- species Common tobacco (*Nicotiana tabacum*) (6002192)
(4097)
Derived from [SCOP 101151](#)



Family: Plant invertase/pectin methylesterase inhibitor

[Pfam 04043](#)

Lineage:

1. Root: [scop](#)
2. Class: [All alpha proteins](#) [46456]
3. Fold: [Bromodomain-like](#) [47363]
4 helices; bundle; minor mirror variant of up-and-down topology
4. Superfamily: [Plant invertase/pectin methylesterase inhibitor](#) [101148]
contains a short alpha-hairpin at the N-terminal extension
[Superfamily](#)
5. Family: [Plant invertase/pectin methylesterase inhibitor](#) [101149]
[Pfam 04043](#)

Protein Domains:

1. Invertase inhibitor [101150]
 1. [Common tobacco \(Nicotiana tabacum\) \[TaxId: 4097\]](#) [101151] (9)
2. Pectin methylesterase inhibitor 1, PME1 [116887]
forms dimers and trimers by swapping the extra N-terminal hairpins
 1. [Thale cress \(Arabidopsis thaliana\) \[TaxId: 3702\]](#) [116888] (3)
SQ [Q9LNF2](#) 28-176

Enter [search](#) key:



CATH

- The CATH (**C**lass, **A**rchitecture, **T**opology, and **H**omologous superfamily) is a semi-automatic, hierarchical classification of protein domains.
- CATH shares many broad features with its principal rival, SCOP.
- The four main levels of the CATH hierarchy are as follows:
 - **Class:** the overall secondary-structure content of the domain. e.g., all α , all β , α/β , $\alpha+\beta$, $\alpha\&\beta$, etc.
 - **Architecture:** high **structural similarity** but **no evidence of homology**. Equivalent to a fold in SCOP.
 - **Topology:** a large-scale grouping of topologies which **share particular structural features**
 - **Homologous superfamily:** indicative of a demonstrable evolutionary relationship. Equivalent to the superfamily level of SCOP.

<http://www.cathdb.info/>

CATH

CATH / Gene3D

26 million protein domains classified into 2,738 superfamilies

[Browse »](#)

[Search »](#)

[Download »](#)

[Take the Tour »](#)

What is CATH?

CATH is a classification of protein structures downloaded from the Protein Data Bank. We group protein domains into superfamilies when there is sufficient evidence they have diverged from a common ancestor.

- [Search CATH by text, ID or keyword](#)
- [Search CATH by protein sequence \(FASTA\)](#)
- [Search CATH by PDB structure](#)
- [Browse CATH Hierarchy](#)
- [CATH Release Notes](#)
- [CATH Tutorials](#)

Example pages

- [PDB "2bop"](#)
- [Domain "1cukA01"](#)
- [Relatives of "1cukA01"](#)
- [Superfamily "HUPS"](#)
- [Functional Family](#)
- [FunFam Alignment](#)
- [Search for "enolase"](#)
- [Superfamily Comparison](#)

Citing CATH

If you find this resource useful, please consider citing the reference that describes this work:

New functional families (FunFams) in CATH to improve the mapping of conserved functional sites to 3D structures.

Sillitoe I, Cuff AL, Dessailly BH, Dawson NL, Furnham N, Lee D, Lees JG, Lewis TE, Studer RA, Rentzsch R, Yeats C, Thornton JM, Orengo CA

Nucleic Acids Res. 2013 Jan PubMed: 23203873

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Latest Release Statistics

CATH v4.0 based on PDB dated March 26, 2013

| | |
|---------|------------------------------------|
| 277,687 | CATH Domains |
| 2,738 | CATH Superfamilies |
| 69,058 | Annotated PDBs |

Gene3D v12 released March 18, 2012

| | |
|------------|---|
| 6,131 | Cellular Genomes |
| 21,662,155 | Protein Sequences |
| 25,615,754 | CATH Domain Predictions |

Search CATH

SEARCH LINKS

[Search by Keyword / ID](#)
[Search By FASTA Sequence](#)
[Search By PDB Structure](#)
[Scan History](#)

Matching Results in CATH

> CATH Superfamilies 83



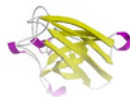
2.40.128.20

CATH superfamily 2.40.128.20

Rhipicephalus appendiculatus, positive regulation of cell proliferation, prostaglandin-D synthase activity, negative regulation of cyto

[View all entries](#)

> CATH Domains 3826



3cqpA00

PDB code 3cqp, chain A, domain 00

Superfamily: 2.60.40.200

hydrogen peroxide biosynthetic process, platelet activation,

[View all entries](#)

Current Search Filters

Remove search filters by clicking on the 'X'

cytokine" X

Filter by Keyword / CATH ID

Start typing and press 'enter' to add a new filter

Top Keywords

Click a keyword to filter results

atp + a protein = adp + a phosphoprotein.

cytokine homo sapiens homo **homo**

sapiens mus musculus myd88-dependent

toll-like receptor signaling pathway ras protein

signal transduction sulfate ion transferase trif-

dependent toll-like receptor signaling pathway

toll signaling pathway zinc ion activation of mapk

activity cellular component movement

membrane platelet activation platelet

degranulation **polymer**

polypeptide(I) positive regulation

of i-kappab kinase/nf-kappab signaling positive

regulation of nf-kappab transcription factor

activity positive regulation of cell proliferation

positive regulation of transcription from rna

polymerase ii promoter stress-activated mapk

cascade tetrameric toll-like receptor 1

signaling pathway toll-like receptor 2 signaling

pathway toll-like receptor 3 signaling pathway

toll-like receptor 4 signaling pathway trimeric

vascular endothelial growth factor receptor signaling

pathway **water** zinc ion binding

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MOTIF

- Motif is a search service provided by GenomeNet to search with a protein query sequence against Motif Libraries.
- Supports several motif databases such as Prosite, BLOCKS, ProDom, Pfam, and PRINTS.
- Allows you to search protein sequence libraries with your patterns.
 - Each residue must be separated with - (minus sign).
 - x represents any amino acids.
 - [DE] means either D or E.
 - {FWY} means any amino acids except for F, W and Y
 - A(2,3) means that A appears 2 to 3 times consecutively.
 - The pattern string must be terminated with . (period).

For example, C-x-{C}-[DN]-x(2)-C-x(5)-C-C.

- Generates a profile from a set of multiple aligned sequences using PFMake or HMMBuild, respectively.

<http://www.genome.jp/tools/motif/>



**Enter your query sequence:**

(sequence only or in Fasta format)

```
>tr|Q9V3N1|Q9V3N1_DROME LD24467p OS=Drosophila melanogaster
GN=Spn27A PE=2 SV=1
MTKMGGNLAVMLLSLFLSALATGNGNSIPTTTTPQGVFETRDKLPGGAASVPSGAGIYD
DIDTFVPRSDSHDPFSWHLLKTVLQNETADKRVIIISPFVSVLVLALLAEAGAGTQTQV
ELANTQTDIRSQNNVREFYRKTILNSFKKENQLHETLSVTRKLFDSFIETQQKFTATLKH
FYDSEVEALDFTNPEAAADAINAWAANITQGRLLQVLVAPDNVRSSVMLLTNLIYFNLWR
RQFATTFQGSFFRSKDDQSRAEFMEQTDYFYITSEKLKAQILRLPYKGNLSLFLVLLPYA
LNGIHDLVKNLENDLKSQAQWAMEEVKVKVTLPKFHFYDQQLKETLRSLGVREIFEDSA
SLPGLTRGADVAGKVKVSNILQKAGINVNEKGEAYAATVVEIENKFGGSTAIEEFNVNR
PFVFFIEEESTGNILFAGKVHSPETTQN
```

Or give the file name containing your sequence:

(plain sequence or in Fasta format)

 No file chosen**Select motif libraries for protein sequences:** ([Help](#))

| Databases | Cut-off score (Click each database to get help for cut-off score) |
|---|---|
| <input checked="" type="checkbox"/> PROSITE Pattern | <input checked="" type="checkbox"/> Skip entries with SKIP-FLAG |
| <input type="checkbox"/> PROSITE Profile | <input checked="" type="checkbox"/> Skip frequently matching (unspecific) profiles |
| <input type="checkbox"/> Pfam | 1.0 * E-value |
| <input type="checkbox"/> User-defined Profile Library (may contain multiple profiles) | <input type="text"/> |

Profile file name: No file chosen

- PROSITE format**
 HMMER format

(Notice) DNA MOTIF search using TRANSFAC (release of December 1999) has been terminated due to the request from Biobase.

MATCHING MOTIF HITS

Your Query Sequence:

MTKMGGNLA VM LLSLFLSALATGNGNSIPTTTTPQGVFETRTDKLPGGAASVPSGAGIYD
DIDTFVPRSDSHDPFSWLLKTVLQNETADKNVIISPFSVKLV LALLAEAAGAGTQTQV
ELANTQTDIRSQNNVREFYRKTLNSFKKENQLHETLSVRTKLFTDSFIETQQKFTATLKH
FYDSEVEALDFTNPEAAADAINAWAANITQGRLQQLVAPDNVRSSVM LLTNLIYFNGLWR
RQFATTFQGSFFRSKDDQSRAEFMEQTDYFYTTSEKLKAQILRLPYKGKNSLFLVLLPYA
LNGIHDLVKNLENDELKSAQWAMEEVKVKVTLPKFHFDYQQNLKETLRSLGVREIFEDSA
SLPGLTRGADVAGKVKVSNILQKAGINVNEKGTEAYAATVVEIENKFGGSTAIEEFNVNR
PFVFFIEEESTGNILFAGKVHSPTTQN

Number of found motifs: 1

PROSITE PATTERN

| Found Motif | Position | PROSITE | Description | Related Sequences | Related Structures |
|-------------|---------------------------------|-------------------------|--------------------|---------------------|---------------------|
| SERPIN | 416..426 Detail | PS00284 | Serpins signature. | 232 | 127 |

PATTERN OF MATCHING MOTIF HITS

Motif SERPIN in your sequence

Prosite ID:

SERPIN (PS00284)

Description:

Serpins signature.

Pattern:

[LIVMFY]-{G}-[LIVMFYAC]-[DNQ]-[RKHQS]-[PST]-F-[LIVMFY]-[LIVMFYC]-x-
[LIVMFAH].

Appearance:

| Position | Found Motif |
|----------|-------------|
| 416..426 | FNVNRPVFFI |

Sequence:

MTKMGGNLAVMLLSLFLSALATGNGNSIPTTTTTPQGVFETRDKLPGGAASVPSGAGIYD
DIDTFVPFRSDSHDPFSWLLKTVLQNETADKNVIISPFVSVKLVALLAEAAGAGTQTQV
ELANTQTDIRSQNNVREFYRKTLSFKKENQLHETLSVRTKLFTDSFIETQQKFTATLKH
FYDSEVEALDFTNPEAAADAINAWAANITQGRLQQLVAPDNVRSSVMLLTNLIYFNGLWR
RQFATTFQGSFFRSKDDQSRAEFMEQTDYFYTTSEKLKAQILRLPYKGKNSLFLVLLPYA
LNGIHDLVKNLENDELKSAQWAMEEVKVKVTLPKFHFQYQQLKETLRLSLGVREIFEDSA
SLPGLTRGADVAGKVKVSNILQKAGINVNEKGTEAYAATVVEIENKFGGSTAIEE**FNVNR**
PFVFFIEEEESTGNILFAGKVHSPTTQN

PFAM

- The Pfam database is a large collection of **protein families**, each represented by multiple sequence alignments and hidden Markov models (HMMs).
- Pfam version 27.0 was produced at the European Bioinformatics Institute using a sequence database called **Pfamseq**, which is based on UniProt.
- The descriptions of Pfam families are managed by the general public using **Wikipedia**.
- The Pfam database contains information about **protein domains and families**.
- Pfam-A is the **manually curated portion of the database** that contains over 10,000 entries.
- Pfam-B contains a **large number of small families derived from clusters** produced by an algorithm called **ADDA** (for automatic generation).

<http://pfam.xfam.org/> Pfam-B families can be useful when no Pfam-A families are found (**but low quality**).



Pfam 27.0 (March 2013, 14831 families)

The Pfam database is a large collection of protein families, each represented by **multiple sequence alignments** and **hidden Markov models (HMMs)**. [More...](#)

QUICK LINKS

[SEQUENCE SEARCH](#)

[VIEW A PFAM FAMILY](#)

[VIEW A CLAN](#)

[VIEW A SEQUENCE](#)

[VIEW A STRUCTURE](#)

[KEYWORD SEARCH](#)

JUMP TO

YOU CAN FIND DATA IN PFAM IN VARIOUS WAYS...

Analyze your protein sequence for Pfam matches

View Pfam family annotation and alignments

See groups of related families

Look at the domain organisation of a protein sequence

Find the domains on a PDB structure

Query Pfam by keywords

Enter any type of accession or ID to jump to the page for a Pfam family or clan, UniProt sequence, PDB structure, etc.

Or view the [help](#) pages for more information






Citing Pfam

If you find Pfam useful, please consider [citing](#) the reference that describes this work:


[The Pfam protein families database](#): R.D. Finn, A. Bateman, J. Clements, P. Coghill, R.Y. Eberhardt, S.R. Eddy, A. Heger, K. Hetherington, L. Holm, J. Mistry, E.L.L. Sonnhammer, J. Tate, M. Punta
Nucleic Acids Research (2014) Database Issue 42:D222-D230

You have hidden the blog posts section. You can restore it [here](#).

Family: *Insulin* (PF00049)

6 architectures 1447 sequences 3 interactions 340 species 923 structures     

Summary

[Domain organisation](#)[Clan](#)[Alignments](#)[HMM logo](#)[Trees](#)[Curation & model](#)[Species](#)[Interactions](#)[Structures](#)**Jump to...** 

Summary: Insulin/IGF/Relaxin family

Pfam includes annotations and additional family information from a range of different sources. These sources can be accessed via the tabs below.

[Wikipedia: Insulin/IGF/Relaxin family](#)[Pfam](#)[InterPro](#)

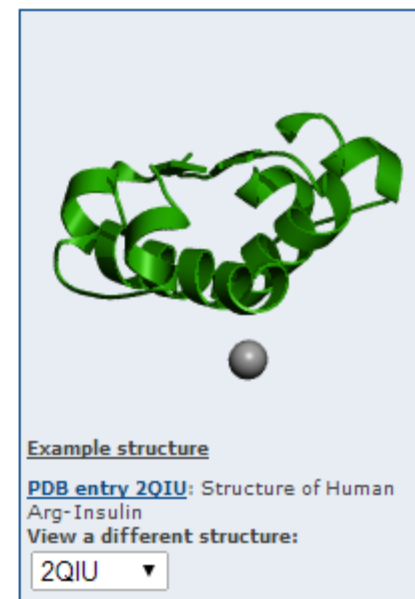
This tab holds the annotation information that is stored in the Pfam database. As we move to using Wikipedia as our main source of annotation, the contents of this tab will be gradually replaced by the Wikipedia tab.

Insulin/IGF/Relaxin family [Provide feedback](#)

Superfamily includes insulins; relaxins; insulin-like growth factor; and bombyxin. All are secreted regulatory hormones. Disulfide rich, all-alpha fold. Alignment includes B chain, linker (which is processed out of the final product), and A chain.

External database links

| | |
|-------------------|---------------------------|
| HOMSTRAD: | ins |
| PANDIT: | PF00049 |
| PRINTS: | PR00276 |
| PROSITE: | PDOC00235 |
| Pseudofam: | PF00049 |
| SCOP: | 1cph |
| SYSTERS: | Insulin |



PROSITE

- PROSITE, a protein domain database for **functional characterization and annotation**.
- PROSITE consists of entries describing the **protein families, domains and functional sites** as well as **amino acid patterns and profiles** in them.
- PROSITE is manually curated by a team of the **Swiss Institute of Bioinformatics** and tightly **integrated into Swiss-Prot protein annotation**.
- PROSITE is complemented by **ProRule**, a collection of rules based on profiles and patterns.
- The rules contain information about biologically meaningful residues, like **active sites, substrate- or co-factor-binding sites, posttranslational modification sites** or **disulfide bonds**, to help function determination.

<http://prosite.expasy.org/>





Database of protein domains, families and functional sites

PROSITE consists of documentation entries describing protein domains, families and functional sites as well as associated patterns and profiles to identify them [[More...](#) / [References](#) / [Commercial users](#)].

PROSITE is complemented by [ProRule](#), a collection of rules based on profiles and patterns, which increases the discriminatory power of profiles and patterns by providing additional information about functionally and/or structurally critical amino acids [[More...](#)].

[Forthcoming changes to the profile format](#)

Release 20.104 of 06-Jul-2014 contains 1703 documentation entries, 1308 patterns, 1087 profiles and 1076 ProRule.

Search

e.g. PDOC00022, PS50089, SH3, zinc finger

Browse

- by documentation entry
- by ProRule description
- by taxonomic scope
- by number of positive hits

Quick Scan mode of ScanProsite

Quickly find matches of your protein sequences to PROSITE signatures (max. 10 sequences). [[?](#)] [[Examples](#)]

```
FVKDVFSTTNRMTVLLITNDVLRGGFKANFTTGYHLGIPEPCADHFQCKNGECVPLVN  
LCDGHLHCEDEGSDAECVRFNNGTINNGLVRFRIQSIWHTACAENWTTQISNDVCQLLG  
LGSGNSSKPIFPTDGGPFVKLNTAPDGHLLITPSQQCLQDSLIRLQCNHKSCGKKLAAQD  
ITPKIVGGSSNAKEGAWPWWVGLYGGRLCGASLVSSDWLVSAACHCVYGRNLEPSKWTAI  
LGLHMKSNLITSPQTVPRLLIDEIVINPHYNRRRKDNDIAMMHLEFKVNYTDYIQPICLPEE  
NQVFPGRNCSIAGWGTVVYQGTANILQEADVPLLSNERCQQMPEYNITENMICAGYE  
EGGIDSCQGDSEGGPLMCQENNRWFLAGVTSFGYKCALPNRPGVYARVSRFTEWIQSFLH
```

Exclude motifs with a high probability of occurrence from the scan

For more scanning options go to [ScanProsite](#)

Other tools

- **PRATT** - allows to interactively generate conserved patterns from a series of unaligned proteins.
- **MyDomains - Image Creator** - allows to generate custom domain figures.



found: 13 hits in 1 sequence

sp-P98073-ENTK_HUMAN (1019 aa)

```
MGSKRGISSRHHSLSSEYIMFAALFAILVVLCAGLIAVSCLTIKESQKGAALGQSHEARATFKITS
GVTYNPNLQDKLSVDFKVLAFDLQQMIDEIFLSSNLKNEYKNSRVLQFENGSIIVVFDLFFAQWVS
DENVKEELIQGLEANKSSQLVTFHIDLNSVDILDKLTITSHLATPGNVSIECLPGSSPCTDALTCI
KADLFCDGEVNCPDGSDENKMCATVCDGRFLLTGSSGSFQATHYKPKPSETSVVCQWIIRVNGLS
IKLSFDDFNTYYTDILDIYEGVGSKILRASIWETNPGTIRIFSNQVTATFLIESDESIVYGFNAT
YTAFNSSSELNNYEKINCNFEDGFCFWVQDLNDDNEWERIQQSTIFSPTGPNFDHTFGNASGFYIST
PTGGGRQERVGLLPLDPTLEPACLSFWYHMYGENVHKLSINISNDQNMKTVFQKEGNYGDNW
NYGQVTLNETVKFKVAFNAFKNKILSDIALDDISLTYGICNGSLYPEPTLVPTPPPELPTDCGGPF
ELWEFNTIFSSNTFNPSYPNLAFVCVWILNAQKGNLQLHFQEFDLENINDVVEIRDGEEADSLLLA
VYTGPGPKVDVFTTNRMTVLLITNDVLRARGGFKANFTTGYHLGPEPCADHFQCKNGECVPLVN
LCDGHLHCEDEADCVRFNNGTINNGLVRFRIQSIWHTACAENWTTQISNDVCQLLGLGSGNS
SKPIFPDGGPFVKLNTAPDGHILITPSQQCLQDSLIRLQCNHKSCGKKLAAQDITPKIVGGSNAK
EGAWPWWVGLYGGRLCGASLVSSDWLVSAACHVYGRNLEPSKWTAILGLHMKSNLITSPQIVPRL
IDEIVINPHYNRRRKNDIAMMHLEFKVNYTDYIQPICLPEENQVFPGRNCSIAGWGTVVYQGTI
ANILQEADVPLLSNERCQQQMPENITENMICAGYEEGGIDSCQGDSSGGLMCCQENNRWFLAGVTS
FGYKCALPNRPGVYARVSRFTEWISFLH
```

Legend:

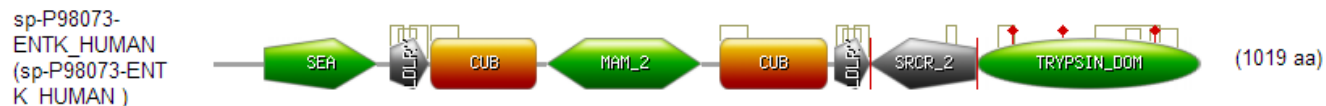
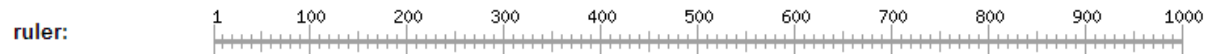


Please note that the graphical representations of domains displayed hereafter are for illustrative purposes only, and that their colors and shapes are not intended to indicate homology or shared function.

For more information about how these graphical representations are constructed, go to <http://prosite.expasy.org/mydomains/>.

hits by profiles: [8 hits (by 6 distinct profiles) on 1 sequence]

Upper case represents match positions, lower case insert positions, and the '-' symbol represents deletions relative to the matching profile.



PS50024 SEA SEA domain profile :

54 - 169: score = 32.979

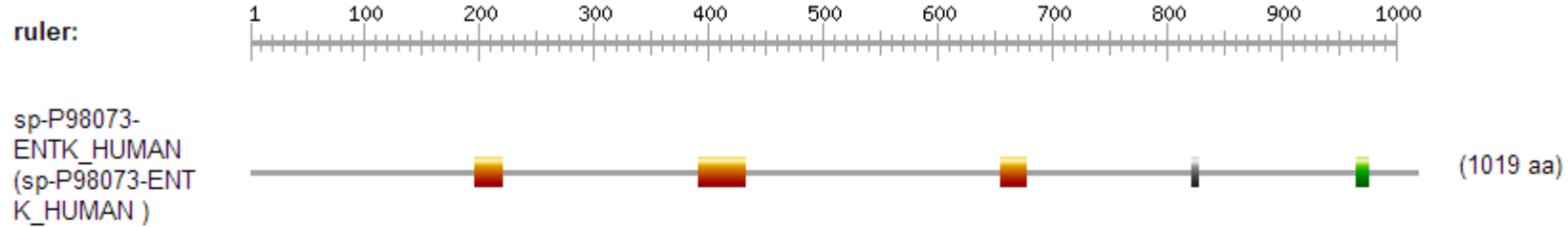
```
QSHEARATFKITsGVTYNPNLQDKLSVDFKVLAFDLQQMIDEIFLSSNLKNEYKNSRVLQ
FEN--GSIIVVFDLFFAQWVSD--ENVKEELIQGLEANKssQLVTFHIDLNSVDILDKLT
```

Predicted feature:

DOMAIN 54 169 SEA

[condition: none]

hits by patterns: [5 hits (by 4 distinct patterns) on 1 sequence]



PS01209 LDLRA_1 *LDL-receptor class A (LDLRA) domain signature :*

197 - 221: [confidence level: (0)] CIkadlf.CDgevNCpdgsDEDnkm..C

655 - 677: [confidence level: (0)] CVplvn1.CDghlHCedg.SDEad...C

PS00740 MAM_1 *MAM domain signature :*

391 - 431: [confidence level: (0)] GfYIstpIgpgrqervg.Lls1pLdptlepaCLsFwYhmyG

PS00134 TRYPsin_HIS *Serine proteases, trypsin family, histidine active site :*

821 - 826: [confidence level: (0)] VSAAHC

Predicted feature:

ACT_SITE 825 Charge relay system (By similarity) [condition: none]

PS00135 TRYPsin_SER *Serine proteases, trypsin family, serine active site :*

965 - 976: [confidence level: (0)] DSsqGDSGGPLM

Predicted feature:

ACT_SITE 971 Charge relay system (By similarity) [condition: none]



THANK YOU