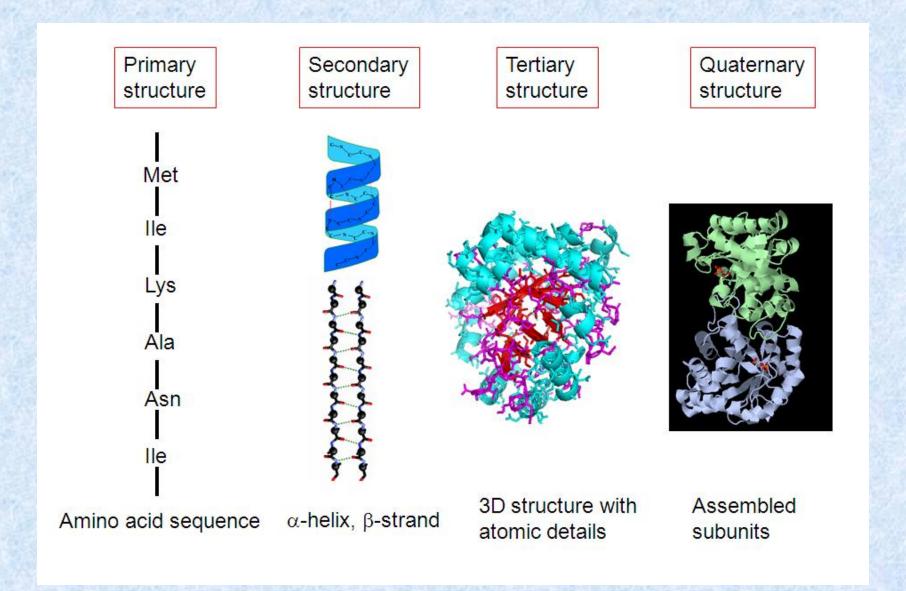
Protein Structure



Primary structure: human hemoglobin

>sp|P68871|HBB_HUMAN Hemoglobin subunit beta OS=Homo sapiens

VHLTPEEKSAVTALWGKVNVDEVGGEALGRLLVVYPWTQRFFESFGDLSTPD AVMGNPKVKAHGKKVLGAFSDGLAHLDNLKGTFATLSELHCDKLHVDPENFR LLGNVLVCVLAHHFG KEFTPPVQAAYQKVVAGVANALAHKYH

Primary structure describes the linear sequence of amino acid residues in a protein.

It includes all covalent bonds between amino acids.

The relative arrangement of the linked amino acids is not specified.

Databases for protein sequences

EXProt

Munich Information Center for Protein Sequences (MIPS)

NCBI Protein database

PIR - Protein Information Resource (Georgetown University)

PIR-NREF

PRF

SWISS-PROT (Swiss Institute of Bioinformatics)

TrEMBL

UniProt - The Universal Protein Knowledgebase

Protein Information Resource

PRO: Protein family classification

iProClass: integrated protein knowledgebase

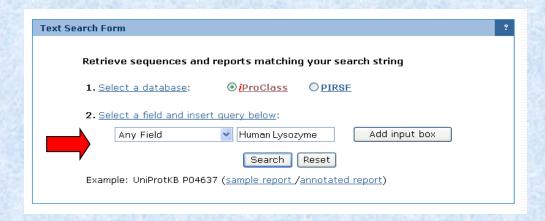
iProLink: literature, information and knowledge



http://pir.georgetown.edu/

Search with iProClass

The iProClass database provides value-added information reports on protein sequences, structures, families, functions, interactions, expressions and modifications.



1 selected (show) • BLAST • FASTA • Pattern Match • Pairwise Alignment • Multiple Alignment • Domain Dis						
☐ Protein AC/ID	Protein Name	Length	Organism Name	PIRSF ID	Related Seq. +	Matched Fields
P79239/LYSC_PONPY /ProClass UniProtKB/Swiss-Prot	Lysozyme C precursor BioThesaurus	148	Pongo pygmaeus (Bornean orangutan)	PIRSF001064	300	Paper Title=>human lysozyme
P61628/LYSC_PANTR /ProClass UniProtKB/Swiss-Prot	Lysozyme C precursor BioThesaurus	148	Pan troglodytes (Chimpanzee)	PIRSF001064	300	Paper Title=>human lysozyme
61627/LYSC_PANPA	Lysozyme C precursor BioThesaurus	148	Pan paniscus (Pygmy chimpanzee) (Bonobo)	PIRSF001064	300	Paper Title=>human lysozyme
P61626/LYSC_HUMAN /ProClass UniProtKB/Swiss-Prot	Lysozyme C precursor BioThesaurus	148	Homo sapiens (Human)	PIRSF001064	300	Paper Title=>human lysozyme; Pape Title=>human
P79179/LYSC_GORGO /ProClass UniProtKB/Swiss-Prot	Lysozyme C precursor BioThesaurus	148	Gorilla gorilla (Lowland gorilla)	PIRSF001064	300	Paper Title=>human lysozyme
P02788/TRFL_HUMAN /ProClass UniProtKB/Swiss-Prot	Lactotransferrin precursor BioThesaurus	710	Homo sapiens (Human)	PIRSF002549; PIRSF500683	300	Paper Title=>human lysozyme
Q6PCD2/Q6PCD2_HUMAN /ProClass UniProtKB/Trembl	GABRE protein BioThesaurus	365	Homo sapiens (Human)		300	Paper Title=>human lysozyme
B2R4C5/B2R4C5_HUMAN /ProClass UniProtKB/Trembl.	Lysozyme (Renal amyloidosis), isoform CRA_a BioThesaurus	148	Homo sapiens (Human)		300	Paper Title=>human lysozyme; Pape Title=>human
Q876Z9/Q876Z9_ASPOR /ProClass UniProtKB/Trembl.	Predicted protein BioThesaurus	600	Aspergillus oryzae	PIRSF037788; PIRSF500676	300	Paper Title=>human lysozyme
Q4R8K7/Q4R8K7_MACFA /ProClass UniProtKB/TremBL	Testis cDNA clone: QtsA-12244, similar to human lysozyme homolog (LOC57151), BioThesaurus	109	Macaca fascicularis (Crab eating macaque) (Cynomolgus monkey)		300	Protein Name=>human lysozyme



1 selected Show						
✓ Protein AC/ID	Protein Name	Length	Organism Name	PIRSF ID	Related Seq. +	Matched Fields
P61626/LYSC_HUMAN /ProClass UniProtKB/Swiss-Prot	Lysozyme C precursor BioThesaurus	148	Homo sapiens (Human)	PIRSF001064	300	UniProtKB AC=>P61626



TLALPN----RKAVADHLLM
LIGCLRNCSAVTAAAKQLAE
VTGFSN----AKTTAQHVKK
.:.*.::
Text Search:

iProClass Summary Report for UniProtKB Entry: P61626

Search/Analysis

		Relate	<u>ed Sequences</u> <u>BioTf</u>	<u>hesaurus</u>	ID Mapping	
GENERAL INFORMATION						
	UniProtKB ID	Protein Name	Protein Name			
8	LYSC HUMAN	P61626; P00695; Q13170; Q9UCF8	Lysozyme C precursor			
Protein Name and ID	PIR-PSD: LZHU RefSeq: NP 000230.1 GenPept: AAA59535.1; AAC63078.1; EAW97222.1; AAA59536.1; CAA32175.1; AAH04147.1; EAW97221.1; ACO37637.1; AAA36188.1 IPI: IP100019038					
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.					
Gene Name	LYZ; LZM					
Keywords	3d-structure; amyloid; amyloidosis; antimicrobial; bacteriolytic enzyme; direct protein sequencing; disease mutation; disulfide bond; glycosidase; hydrolase; polymorphism; polysaccharide degradation; signal					
Function	Lysozymes have primarily a bacteriolytic function; those in tissues and body fluids are associated with the monocyte- macrophage system and enhance the activity of immunoagents.					
Subunit	Monomer.					

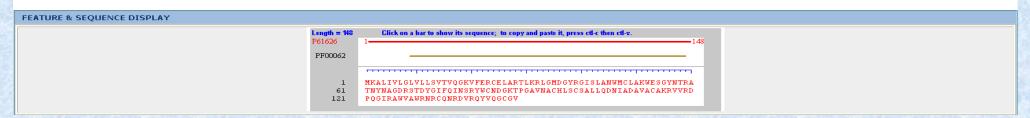
Support

Download

CROSS-REFERENCES	
Bibliography	► <u>View Bibliography Information</u> ► <u>Submit Bibliography</u> Annotated references: PMID: 8105095; 10350481; 10469827; 10561612; 11887182; 11927576; 11986950 [PDB/GeneRIF] More
	Other references: PMID: 11849445; 12675840; 15745733; 8765309; 9659355; 9745729; 18391951; 9359845; 8566845; 17353931; 9883972; 366724; 10534505; 12477932; 10558865; 18591461
DNA Sequence	GenBank/EMBL/DDBJ: M21119; J03801; M19045; X14008; U25677; BC004147

Structure	185U: SCOP CATH FSSP MMDB PDBsum 185V: SCOP CATH FSSP MMDB PDBsum 185W: SCOP CATH FSSP MMDB PDBsum 185W: SCOP CATH FSSP MMDB PDBsum 185X: SCOP CATH FSSP MMDB PDBsum 185X: SCOP CATH FSSP MMDB PDBsum 185X: SCOP CATH FSSP MMDB PDBsum 187L: SCOP CATH FSSP MMDB PDBsum 187M: SCOP CATH FSSP MMDB PDBsum 187M: SCOP CATH FSSP MMDB PDBsum 187N: SCOP CATH FSSP MMDB PDBsum 187P: SCOP CATH FSSP MMDB PDBsum 187S: SCOP CATH FSSP MMDB PDBsum 1884: SCOP CATH FSSP MMDB PDBsum
PIR Feature & Post Translational Modifications	FEAT1; active site: Glu, Asp (53,71) [predicted] FEAT2; binding site: substrate (Asp) (120) [predicted] FEAT3; disulfide bonds: (24-146,48-134,83-99,95-113) [experimental] FEAT3; domain: signal sequence (1-18) [predicted] FEAT5; product: lysozyme (19-148) [experimental] Phosphosite: P61626
	PIR Feature & Post Translational

FAMILY CLASSIFICATION	
UniRef	UniRef100 P61626; UniRef90 P61626; UniRef50 P61626
PIRSF	PIRSF001064 lysozyme c
Pfam Domain	Pfam: PF00062: C-type lysozyme/alpha-lactalbumin family (19-146)
Prosite Motif	Prosite: P500128: PDOC00119: Alpha-lactalbumin / lysozyme C signature. Prosite: P551348: PDOC00119: Alpha-lactalbumin / lysozyme C family profile.
InterPro	InterPro: LYSC HUMAN IPR001916: Glycoside hydrolase, family 22 IPR000974: Glycoside hydrolase, family 22, lysozyme
SCOP Fold	►Class: Alpha and beta proteins (a+b); Fold: Lysozyme-like; Superfamily: Lysozyme-like; Family: C-type lysozyme [133L:A; 134L:A; 185U:A; 185U:A; 185V:A; 185X:A; 185X:A; 185Z:A; 185Z:B; 187L:A; 187M:A; 187N:A; 187O:A; 187P:A; 187P:A; 187S:A; 1883:B; 1884:B; 1884:B; 1885:B; 1885:B; 1043:A; 1045:A; 1045:A; 1046:A; 107P:A; 107P
Other Classification	BLOCKS: IPB000974 Lysozyme signature PRINTS: PR00137 LYSOZYME PRINTS: PR00135 LYZLACT SMART: SM00263 LYZ1



Swiss-prot/Uniprot

Annotated protein sequence database established in 1986 and maintained collaboratively, since 1987, by the Department of Medical Biochemistry of the University of Geneva and the EMBL Data Library.

It 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 domain structure, post-translational modifications and variants), a minimal level of redundancy and a high level of integration with other databases.

TrEMBL is a computer annotated supplement of SWISS-PROT that contains all the translations of EMBL nucleotide sequence entries not yet integrated in SWISS-PROT.

Currently, SWISS-PROT and TrEMBL have 0.55 and 73.7 million sequences, respectively.

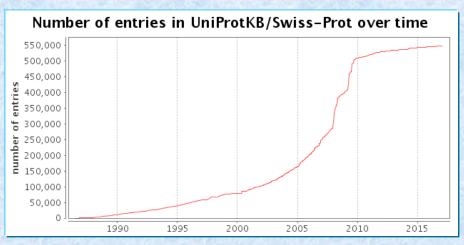
Total: 74.4 million

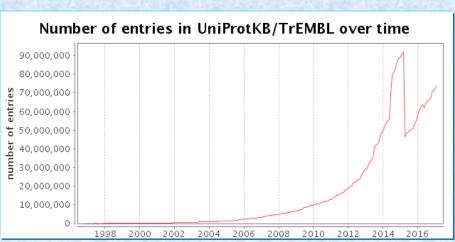
http://www.ebi.ac.uk/swissprot/

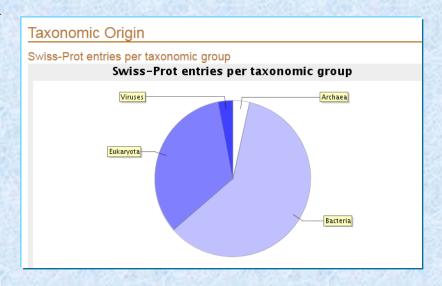
http://www.uniprot.org/uniprot/

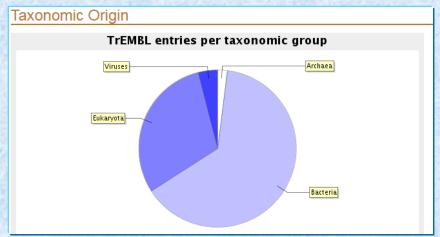
Uniprot: statistics

Number of entries: ~74.4 million



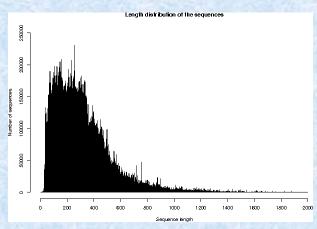


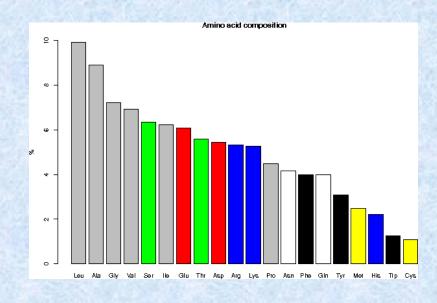


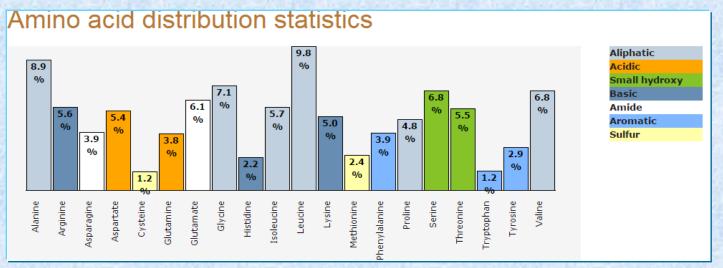


Uniprot: statistics

Average sequence length: 315 amino acids







Name and origin of the protein

protein attributes

general information

Ontologies

sequence annotation

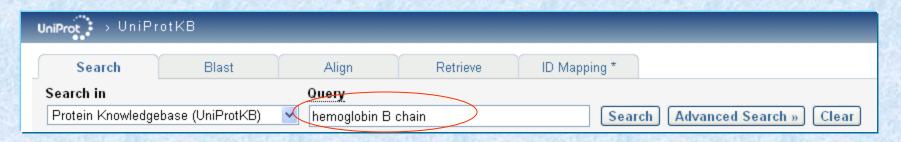
amino acid sequence

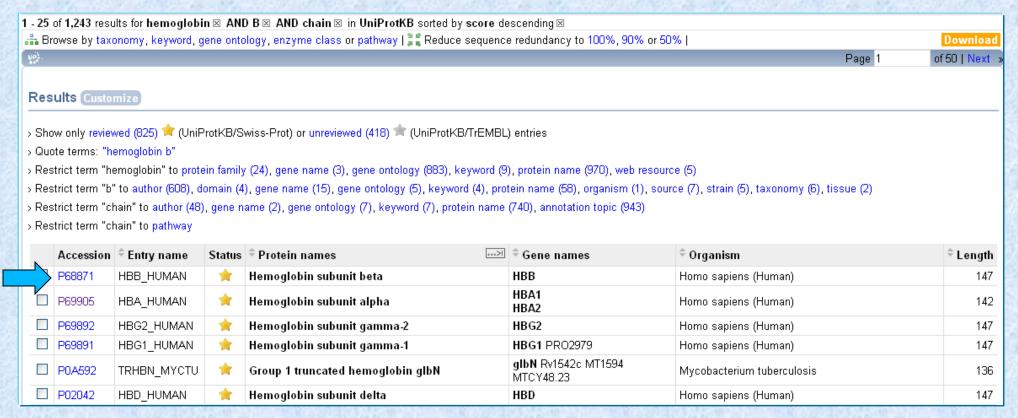
bibliographic references

cross-references with sequence, structure and interaction databases

entry information.

Uniprot: search results





Names and origin			
Protein names Recommended name: Hemoglobin subunit beta Alternative name(s): Beta-globin Hemoglobin beta chain Cleaved into the following chain: 1. LVV-hemorphin-7			
Gene names	Name: HBB		
Organism	Homo sapiens (Human) [Complete proteome]		
Taxonomic identifier	9606 [NCBI]		
Taxonomic lineage	Eukaryota > Metazoa > Chordata > Craniata > Vertebrata > Euteleostomi > Mammalia > Eutheria > Euarchontoglires > Primates > Haplorrhini > Catarrhini > Hominidae > Homo		

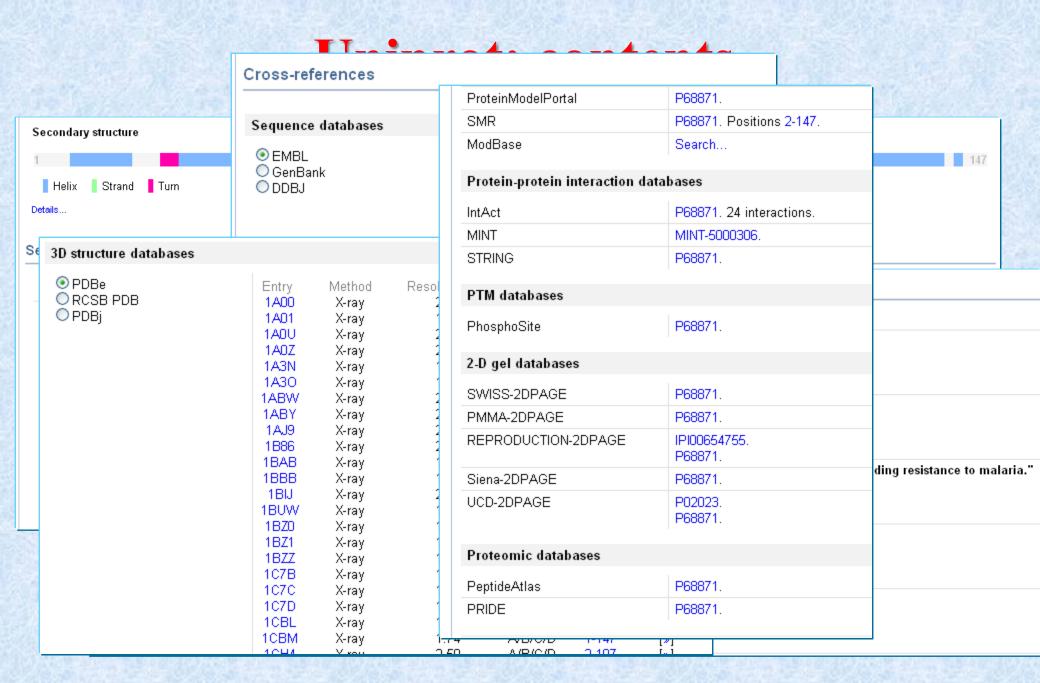
Protein attributes				
Sequence length	147 AA.			
Sequence status	Complete.			
Sequence processing	The displayed sequence is further processed into a mature form.			
Protein existence	Evidence at protein level.			

General annotation (Comm	ents)
Function	Involved in oxygen transport from the lung to the various peripheral tissues. Ref.35 LVV-hemorphin-7 potentiates the activity of bradykinin, causing a decrease in blood pressure. Ref.35
Subunit structure	Heterotetramer of two alpha chains and two beta chains in adult hemoglobin A (HbA).
Tissue specificity	Red blood cells.
Post-translational modification	Glucose reacts non-enzymatically with the N-terminus of the beta chain to form a stable ketoamine linkage. This takes place slowly and continuously throughout the 120-day life span of the red blood cell. The rate of glycation is increased in patients with diabetes mellitus.
	S-nitrosylated; a nitric oxide group is first bound to Fe ²⁺ and then transferred to Cys-94 to allow capture of O ₂ .
	Acetylated on Lys-60, Lys-83 and Lys-145 upon aspirin exposure. Ref.34 reports the identification of HBB acetylated on Lys-145 in the cytosolic fraction of HeLa cells. This may have resulted from contamination of the sample.
Involvement in disease	Defects in HBB may be a cause of Heinz body anemias (HEIBAN) [MIM:140700]. This is a form of non-spherocytic hemolytic anemia of Dacie type 1. After splenectomy, which has little benefit, basophilic inclusions called Heinz bodies are demonstrable in the erythrocytes. Before splenectomy, diffuse or punctate basophilia may be evident. Most of these cases are probably instances of hemoglobinopathy. The hemoglobin demonstrates heat lability. Heinz bodies are observed also with the Ivemark syndrome (asplenia with cardiovascular anomalies) and with glutathione peroxidase deficiency. Ref.125 (Ref.126) (Ref.129)
	Defects in HBB are the cause of beta-thalassemia (B-THAL) [MIM:604131]. A form of thalassemia. Thalassemias are common monogenic diseases occurring mostly in Mediterranean and Southeast Asian populations. The hallmark of beta-thalassemia is an imbalance in globin-chain production in the adult HbA molecule. Absence of beta chain causes beta(0)-thalassemia, while reduced amounts of detectable beta globin causes beta [†] -thalassemia. In the severe forms of beta-thalassemia, the excess alpha globin chains accumulate in the developing erythroid precursors in the marrow. Their deposition leads to a vast increase in erythroid apoptosis that in turn causes ineffective erythropoiesis and severe microcytic hypochromic anemia. Clinically, beta-thalassemia is divided into thalassemia major which is transfusion dependent, thalassemia intermedia (of intermediate severity), and thalassemia minor that is asymptomatic. Ref52
	Defects in HBB are the cause of sickle cell anemia (SKCA) [MIM:603903]; also known as sickle cell disease. Sickle cell anemia is characterized by abnormally shaped red cells resulting in chronic anemia and periodic episodes of pain, serious infections and damage to vital organs. Normal red blood cells are round and flexible and flow easily through blood vessels, but in sickle cell anemia, the abnormal hemoglobin (called Hb S) causes red blood cells to become stiff. They are C-shaped and resembles a sickle. These stiffer red blood cells can led to microvascular occlusion thus cutting off the blood supply to nearby tissues.
	Defects in HBB are the cause of beta-thalassemia dominant inclusion body type (B-THALIB) [MIM:603902]. An autosomal dominant form of beta thalassemia characterized by moderate anemia, lifelong jaundice, cholelithiasis and splenomegaly, marked morphologic changes in the red cells, erythroid hyperplasia of the bone marrow with increased numbers of multinucleate red cell precursors, and the presence of large inclusion bodies in the normoblasts, both in the marrow and in the peripheral blood after splenectomy. (Ref.52)
Miscellaneous	One molecule of 2,3-bisphosphoglycerate can bind to two beta chains per hemoglobin tetramer.

Ontologies						
Keywords						
Biological process	Oxygen transport Transport					
Coding sequence diversity	Polymorphism					
Disease	Congenital dyserythropoietic anemia Disease mutation Hereditary hemolytic anemia					
Ligand	Heme Iron Metal-binding Pyruvate					
Molecular function	Hypotensive agent Vasoactive					
PTM	Acetylation Glycation Glycoprotein Phosphoprotein S-nitrosylation					
Technical term	3D-structure Complete proteome Direct protein sequencing					

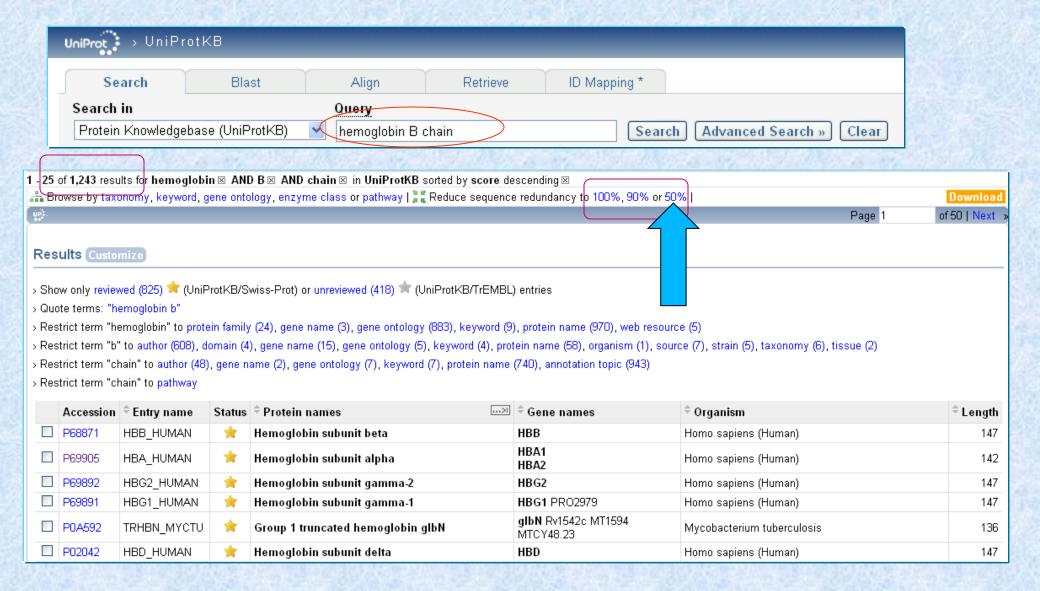
Gene Ontology (GO)		
Biological process	blood coagulation Traceable author statement. Source: Reactome	
	hydrogen peroxide catabolic process Inferred from direct assay. Source: BHF-UCL	
	nitric oxide transport Non-traceable author statement. Source: UniProtKB	
	positive regulation of cell death Inferred from direct assay. Source: BHF-UCL	
	positive regulation of nitric oxide biosynthetic process Non-traceable author statement. Source: UniProtKB	
	protein heterooligomerization Inferred from direct assay. Source: BHF-UCL	
	regulation of blood pressure Inferred from electronic annotation. Source: UniProtKB-KW	
	regulation of blood vessel size Inferred from electronic annotation. Source: UniProtKB-KW	
Cellular component	haptoglobin-hemoglobin complex Inferred from direct assay. Source: BHF-UCL	
	hemoglobin complex Non-traceable author statement (Ref.33)(Ref.71). Source: UniProtKB	
Molecular function	heme binding Inferred from electronic annotation. Source: InterPro	
	hemoglobin binding Inferred from direct assay. Source: UniProtKB	
	oxygen binding Inferred from direct assay. Source: UniProtKB	
	oxygen transporter activity Non-traceable author statement (Ref.71). Source: UniProtKB	

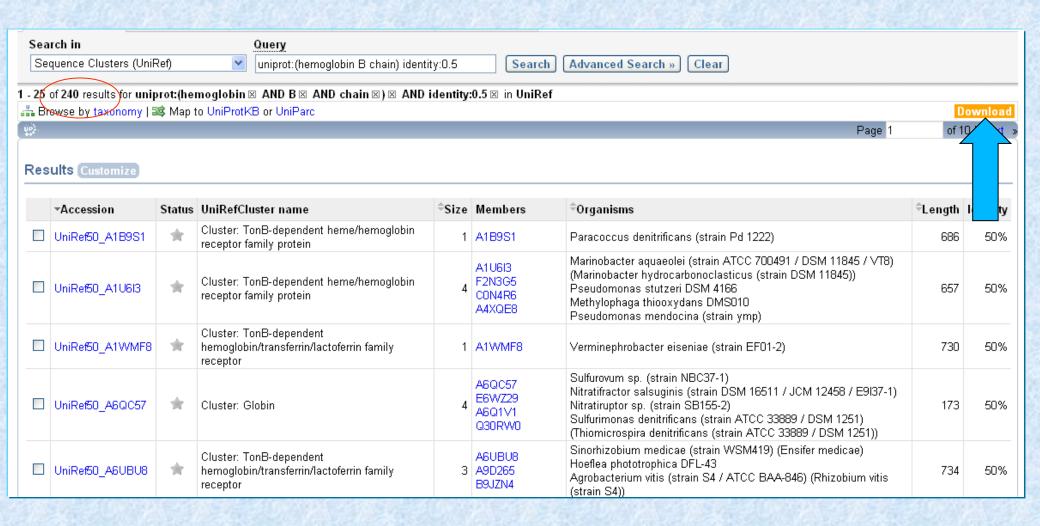
Bina	Binary interactions								
With	With Entry #Exp. IntAct No.			Natu	ıral variations				
HBA	1 P69905 1	EBI-715554,EBI-714680			Natural variant	2	1	V → A in Raleigh; O(2) affinity down. [dbSNP:rs33949930]	
				'		Natural variant	3	1	H → L in Graz. [dbSNP:rs35906307] (Ref.78)
Sequ	ience annotatio	n (Features	s)			Natural variant	3	1	H → Q in Okayama; O(2) affinity up. [dbSNP:rs713040]
				l		Natural variant	3	1	H → R in Deer Lodge; O(2) affinity up. [dbSNP:rs33983205]
	Feature key	Position(s)	Length	Descrip		Natural variant	3	1	H → Y in Fukuoka. [dbSNP:rs35906307]
Mole	Molecule processing				Natural variant	6	1	P → R in Warwickshire. [dbSNP:rs34769005]	
	Initiator methionine	1	1	Remov		Natural variant	7	1	E → A in G-Makassar.
	Chain	2 – 147	146			Natural variant	7	1	E → K in C. Ref.3 Ref.49
	Peptide	33 – 42	10	LVV-he		Natural variant	7	1	E → Q in Machida. [dbSNP:rs33930165]
Sites						Natural variant	7	1	E → V in S; sickle cell anemia. [dbSNP:rs334] Ref.10 Ref.39
			4	l		Natural variant	8	1	E → G in G-San Jose; mildly unstable. [dbSNP:rs34948328]
	Metal binding Metal binding	64 93	1	Iron (he Iron (he		Natural variant	8	1	E → K in G-Siriraj. [dbSNP:rs34948328]
	Binding site	2	1	2,3-bis		Natural variant	9	1	K → E in N-Timone. [dbSNP:rs33932981] Ref.101
	Binding site	3	1	2,3-bis		Natural variant	9	1	K → Q in J-Luhe. [dbSNP:rs33926764]
	Binding site	83	1	2,3-bis		Natural variant	9	1	K → T in Rio Grande. Ref.114
	Binding site	144	1	2,3-bis		Natural variant	10	1	S → C in Porto Alegre; O(2) affinity up. [dbSNP:rs33918131]
						Natural variant	11	1	A → D in Ankara. [dbSNP:rs33947457] Ref.55



Genome annotation databases				Enzyme and pathway databases					
Ensembl		ENST00000335295; ENSP		Reactome		REACT_604. Hemostasis.			
GenelD		3043.		Gene expression databases					
KEGG		hsa:3043.							
UCSC		uc001mae 1 human		ArrayExpress P68871.		P68871.			
Organism-specific	Entry information								
CTD	Entry name HBB			_HUMAN					
GeneCards				Primary (citable) accession number: P68871 Secondary accession number(s): A4GX73 🕶 Q9UCP9					
H-InvDB				ntegrated into UniProtKB/Swiss-Prot: July 21, 1986					
HGNC	Ü		Last	ast sequence update: January 23, 2007					
HPA				st modified: June 28, 2011 is is version 97 of the entry and version 2 of the sequence. [Complete history]					
MIM	Entry status		Reviewed (UniProtKB/Swiss-Prot)						
	Annotation program		Chordata Protein Annotation Program				1 hit.		
	Disclaimer		Any medical or genetic information present in this entry is provided for research, a used as a substitute for professional medical advice, diagnosis, treatment or care						
neXtProt			useu	as a substitute for profession	mai medic	ar advice, diagnosis, treatment of care			
Orphanet	Relevant	documents							
	Human chromosome 11 Human chromosome 11: entries, gene names and cross-references to MIM								
PharmGKB		ries with polymorphisms							
GenAtlas	List of hum	an entries with polymorph	nisms	s or disease mutations					
Phylogenomic databases				DrugBank		DB00893. Iron Dextran.			
, ,				NextBio 12048.					
HOVERGEN		HBG009709.		PMAP-CutDB		P68871.			

Uniprot: search results





240 results for uniprot:(hemoglobin ⋈ AND B ⋈ AND chain ⋈) ⋈ AND identity:0.5 ⋈ in UniRef

> Download data compressed or uncompressed

Tab-Delimited

Summary information from the result view.

[Download | Open | Open first 10]

Excel 🛎

Summary information from the result view for MS Excel™.

[Download | Open | Open first 10]

FASTA

Sequence data in FASTA format.

[Download (200 KB*) | Open | Open first 10]

XML

Complete data in XML for

[Download (400 KB*) | Open | Open first 10]

RDF/XML

Complete data in RDF format.

[Download (500 KB*) | Open | Open first 10]

List

List of accession numbers.

[Download (5 KB*) | Open | Open first 10]

>UniRef50_A1B9S1 TonB-dependent heme/hemoglobin receptor family MPRHSIRGALLAGTACLTALTFTAPLLAQERAGADSAQSTYVLDQIVLRAGKPKVASEVP QSVSVVDSRQLEDIAPIHIGEVLATVPGVAGVGSGSFFGQGFNIRGFGSSGAAASESGIV QLIDGEEKYYESYRQGALFVEPDFLRQVEVLRGPGSSTLYGSGALGGVIAMETIEAGDLI AEGQTFGGRTKLGYASNPDTVLGSVALGWRPAEDFEALAAFAWRKLGDTKDADGNTTVRA NSKTPNLLLKAKKTFGDQYVAFSYQHLEAKGDDQDFNQLEGAQVGLFPGFPGWGVGDITT RDQTARFIWGWNPEDNRYVDLTATLSYTNTLKDVRQGDDPDEPIMDSLLGERDYRLWKFK LSNVADLSGAGYDHHLTTGAEVLKQDRSSSVPSSSHPEAYTRAWAAYALSELTWGDLTIN SGLRYEKQRTEPKSSVTVTDDTYDADSVEPQVAAIYRLNDSLSVFGSVAFVNRMPTVDEL YDSFMGGAPSGDLKDEKGKNIELGLSYHGSGILTASDEAVVKLTLFRNHIDDMIVRTNAP APMPAYVNIDRAYLRGGELEATYSVAAWEFGAAFSVVNGVDQDGADLDTLPNNRVTLQAI WQASDALRLGLRSTLADGRDKPNGTHRAGYGVHDVFATWVPQGGAAAGIEVHVGVDNVTD RDYTPATWLSGPAPGRNFKLSVSRSF

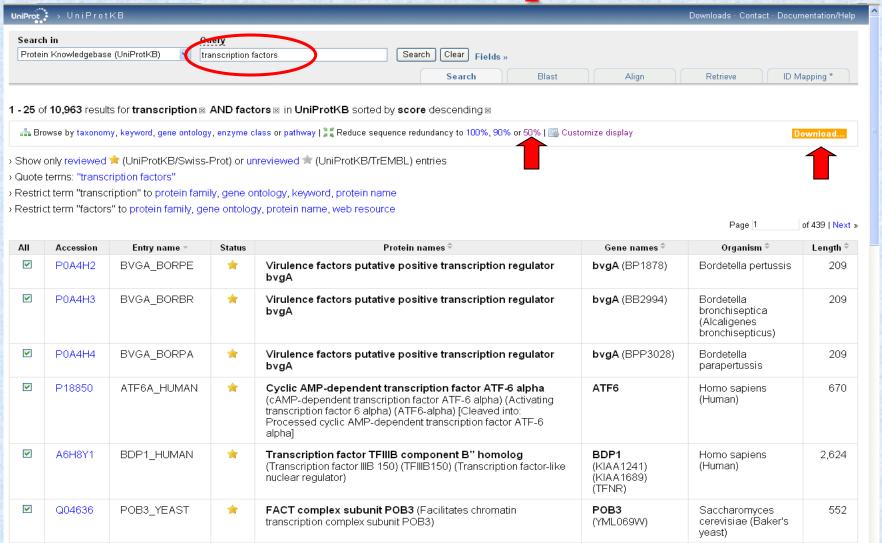
>UniRef50_A1U6I3 TonB-dependent heme/hemoglobin receptor family MANSSPMKQPRFRRNTLWLALMAAPLAHAQPVSLDPIQVTADREADADTVVDAETIERF QADDLEDVFAGQPDVSVGGSNSIAQKVYVRGFDDPLLNVSVDGATQAGALFHHSGRLSVE PELLKQVEVNAGAGRATEGAGALGGSIRFVTKDPDDLLRPGESAGALVKFGSFSNTDGYK ASGTAFGRLSDNWSTLVSVSQSDHEPFKDGSGDRIAGSDARQQLGFAKLVGQLPADQTIK LSHEVRTDEGERPQRPQWVVSSFNRLYELDGRRDTTTLNYGYAPAGNALVDLEATVYHTE SDIEQNVEDRWGRYFGFSRNIGGDLRNTSRFGGHSLTYGVDYREDKVNAGYQEDKRAEQQ TGEVLGVYLQGDLWLTSRLLFSAGARYDDYRLKDNDDQRFSEDEVSPNANLAWEVVDGLT LKAGYAEAFRGPTTQDAFKLEGSENDPDLEGEKARNTEVGFDYRYETFRLSAEVYRSEIK DAIADPLLPFRESIYKNIGDLESDGYLISAGYQWQALSAGLSFHSNDAEVDGQPLTVYEH NLLGNTMGDTWIADLAYRWDRNLEFGWQGRFVEGIDNLDTSVGTIDKPGYGVHDLYLHWL PTGNEDLRLSLTIKNVGDKQYLAHASNADYQHIEDYEGIVGMPEPGRDIRVGLAMRF

Question

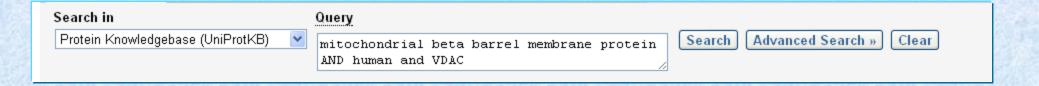
Obtain the sequences of "transcription factors" with less than 50% sequence identity.

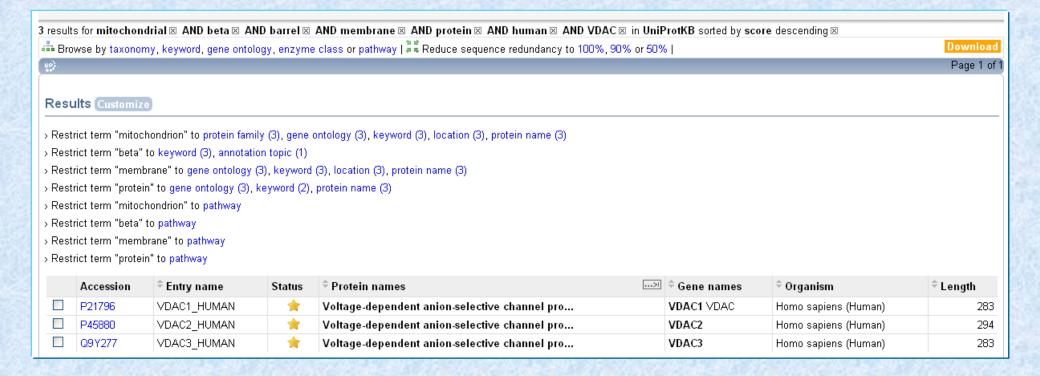
Find the amino acid sequence of human mitochondrial beta barrel membrane protein VDAC

Dataset for transcription factors



```
>sp|P18485|1A12_SOLLC 1-aminocyclopropane-1-carboxylate synthase 2 OS=Solanum lycopersicum GN=ACS2 PE=1 SV=2
MGFEIAKTNSILSKLATNEEHGENSPYFDGWKAYDSDPFHPLKNPNGVIOMGLAENOLCL
DLIEDWIKRNPKGSICSEGIKSFKAIANFQDYHGLPEFRKAIAKFMEKTRGGRVRFDPER
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ARKMSSFGLVSTQTQYFLAAMLSDEKFVDNFLRESAMRLGKRHKHFTNGLEVVGIKCLKN
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SPLVR
>sp|P16375|7UP1 DROME Steroid receptor seven-up, isoforms B/C OS=Drosophila melanogaster GN=svp PE=1 SV=1
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PGTTTGSVATGGGGTTPSSVASOOSAVIKODLSCPSLNOAGSGHHPGIKEDLSSSLPSAN
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PSM
>sp|P16376|7UP2 DROME Steroid receptor seven-up, isoform A OS=Drosophila melanogaster GN=svp PE=2 SV=3
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```





Sequences											
		Sequence			Length Ma	ass (Da)	Tools				
[P21796 [UniPa	irc].		FASTA	283	30,773	Blast	~	go	
		Last modified January 23, 2007. Version 2. Checksum: 89BA3378B04020D5									
		1 <u>0</u> MAVPPTYADL	_	3 <u>0</u> GYGFGLIKLD				6 <u>0</u> ET			
		7 <u>0</u> KYRWTEYGLT	_	9 <u>0</u> LGTEITVEDQ		11 <u>0</u> ssfspntgkk		.2 <u>0</u> TKR			
		13 <u>0</u> EHINLGCDMD	_	15 <u>0</u> ALVLGYEGWL				.8 <u>0</u> .8 <u>0</u>			
		19 <u>0</u> HTNVNDGTEF	_	21 <u>0</u> KLETAVNLAW	22 <u>0</u> TAGNSNTRFG	_		:4 <u>0</u> INS			
		25 <u>0</u> SLIGLGYTQT	_	27 <u>0</u> ALLDGKNVNA		FQA					

>sp|P21796|VDAC1_HUMAN Voltage-dependent anion-selective channel protein 1 OS=Homo sapiens GN=VDAC1 PE=1 SV=2
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KYRWTEYGLTFTEKWNTDNTLGTEITVEDQLARGLKLTFDSSFSPNTGKKNAKIKTGYKR
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SLIGLGYTQTLKPGIKLTLSALLDGKNVNAGGHKLGLGLEFQA