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NCBI News, September 2014

NCBI Sequence Viewer version 3.4 available

Tuesday, September 30, 2014

NCBI Sequence Viewer has recently been updated and now has improved visualization of graphs, sequence track and other text, as well as a reworked configuration dialog. A full list of new features, improvements and fixes is included in the release notes.

Sequence Viewer provides a graphical view of sequences and color-coded annotations on regions of sequences stored in the Nucleotide and Protein databases.

HIV-1, human interaction database updated

Monday, September 29, 2014

The HIV-1, human interaction database has been updated and is now on an improved page. The improved interface includes help documentation and supports structured queries against Gene, as well as browsing, filtering and downloading the protein and replication interaction data sets. The most recent data release (June 2014) includes 12,785 HIV-1, human protein-protein interactions for 3,142 human genes and 1,316 replication interactions for 1,250 human genes.

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HIV-1 Human Interaction Database

- About the database
- Help
- Publications
- Releases

Browse and Download Data

Protein and Replication Interactions

Search NCBI Gene Records With HIV-1 Interaction Data

Searching human genes with all selected criteria. For HIV-1 genes, click here.

20 10 10 10 10 10 10 10 10 10 10 10 10 10	THE STATE OF THE S
Gene Ontology (GO)	type to choose GO term
Protein domain name	type to choose protein domain name
Properties	☐ Has phenotype
	Has gene expression data
	☐ Has Homologene Cluster
	□ Has >1 RefSeq transcript
	Has biological pathways
Entrez Gene keywords	enter optional keywords

Figure 1. The HIV-1 interactions database homepage.

The HIV-1, human interactions project collates published reports of two types of interactions: HIV-1, human protein interactions, and human gene knock-downs that affect virus replication which are reported as "replication interactions".

Virus Variation Resource pages for Ebolavirus, MERS coronoavirus give quick and easy access to related sequences and other data

Friday, September 19, 2014

NCBI has created resource pages for Ebolavirus and MERS coronavirus, giving users an easy way to find all sequences related to these pathogens. These pages aggregate links to virus data at NCBI and also provide important links out to other information at the CDC, WHO, and HealthMap.



Figure 1. Ebolavirus Resource page. The main column gives a brief description of the resource and displays the NCBI Graphical Sequence Viewer. In the right column, from top to bottom, are links to: the Ebolavirus database, other NCBI Ebolavirus resources, and links out to HealthMap, CDC, and WHO. The MERS coronavirus resource page has the same layout and links.

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The Virus Variation resource pages all include a description of the resource components: the database, the reference genome graphical display and links to other resources, both external and within NCBI.

Dedicated Virus Variation databases for Ebolavirus and MERS coronavirus have been developed. These databases allow searching for nucleotide and protein sequences by a variety of criteria including host, sequence patterns, region or country of isolation, and collection or release dates. The databases allow you to:

- Quickly find the sequences you need, through an intuitive search interface for all viral sequences using standardized protein/gene names and metadata
- Select the latest sequences based on date criteria or sorting of results
- Download sequences in many formats or find links to sequences in NCBI databases.

Visit the Virus Variation homepage to see resource pages for other viruses.

Simplified FASTA headers included on new NCBI Genomes FTP site

Wednesday, September 17, 2014

Last month, a major revision of the NCBI Genomes FTP site was announced. In response to user feedback, a new format for FASTA headers of genome, protein and transcript records has been implemented. This new format is limited to records in the /all/, /refseq/, and /genbank/ directories on the new Genomes FTP site and does not affect the Nucleotide database web FASTA displays.

Now, instead of ">gi|xx|dbsrc|accession.version|description", the new format is simply ">accession.version description".

For example, the header on the FASTA record for Homo sapiens chromosome 1 was previously:

>gi|568336023|gb|CM000663.2| Homo sapiens chromosome 1, GRCh38 reference primary assembly.

On the new Genomes FTP site, the header is now:

>CM000663.2 Homo sapiens chromosome 1, GRCh38 reference primary assembly.

NCBI has traditionally used a compound FASTA sequence identifier string in which multiple IDs were separated by "|" characters. This format provides more information, but requires that the individual sequence identifiers be parsed out of the compound string. The simpler sequence identifier string is identical to that used in the GFF annotation files on the genomes FTP site. Providing sequence and annotation files with matching sequence identifiers supports their use in commonly used RNA-Seq analysis packages and in other analysis pipelines that rely on simple string comparison to match sequence identifiers.

More information about the revised Genomes FTP site, including the new FASTA header format, is available on the Genomes Download FAQ page.

RefSeq release 67 available on FTP

Thursday, September 11, 2014

The full RefSeq release 67 is now available on the FTP site with over 61 million records describing 45,166,402 proteins, 8,163,775 RNAs, and sequences from 41,913 different NCBI TaxIDs.

More details about the RefSeq release 67 are included in the release statistics and release notes. In addition, reports indicating the accessions included in the release and the files installed are available.

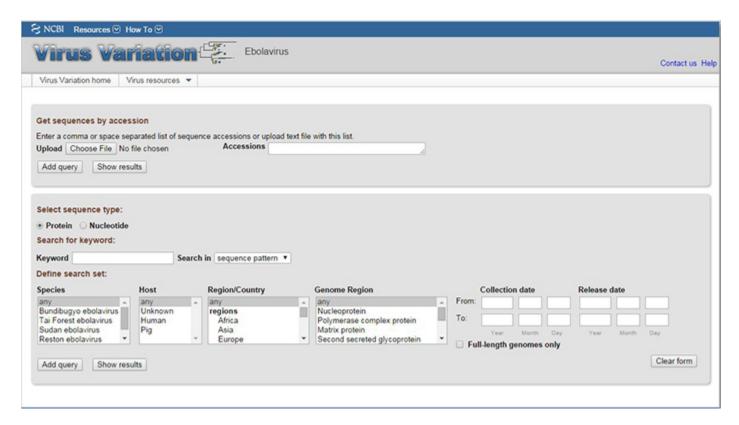


Figure 2. Ebolavirus database. Users can get sequences by accession or browse by searching for a keyword, host, or region/country, among other options.

Identical Protein Report Display option added to Protein database

Tuesday, September 09, 2014

A new display option has been added to the Protein database - the "Identical Protein Report". When viewing an individual record, this display allows you to access a list of all other identical proteins including those submitted as translations to GenBank, as well as RefSeq, UniProtKB/Swiss-Prot, PIR, PDB, and patented protein records.

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60S ribosomal protein L23a [Trypanosoma brucei]

GenBank: AAX79509.1
GenPept FASTA Graphics

RefSeq Selected Product: XP 846140.1, 164 amino acids

Name: 60S ribosomal protein L23a [Trypanosoma brucei brucei strain 927/4 GUTat10.1]

Source	CDS Region in Nucleotide	Protein	Organism	Superkingdom
RefSeq	NC_007280_1:1362946-1363440+	XP_846140.1	Trypanosoma brucei brucei strain 927/4 GUTat10.1	Eukaryota
RefSeq	XM_841047.1:1-495+	XP_846140.1	Trypanosoma brucei brucei strain 927/4 GUTat10.1	Eukaryota
Swiss-Prot	N/A	P41165.1	Trypanosoma brucei brucei	Eukaryota
PDB	N/A	3ZF7_X	Trypanosoma brucei brucei strain 927/4 GUTat10.1	Eukaryota
INSDC	L21172.1:43-539+	AAC37186.1	Trypanosoma brucei	Eukaryota
INSDC	AC105378.10:65131-65625+	AAX79509.1	Trypanosoma brucei	Eukaryota
INSDC	CP000070.1:1362946-1363440+	AAZ12581.1	Trypanosoma brucei brucei strain 927/4 GUTat10.1	Eukaryota
INSDC	FN554970.1:1431278-1431772+	CBH12694.1	Trypanosoma brucei gambiense DAL972	Eukaryota
Patent	N/A	AAQ39714.1	Unknown	Unknown
Patent	N/A	ACC09504.1	Unknown	Unknown

Figure 1. The "Identical Protein Report" display setting in the Protein database, showing identical proteins for 60S ribosomal protein L23 [*Trypanosoma brucei*].

As shown in Figure 1, the page title reflects the protein record from which you started. Beneath that, there is information on the suggested RefSeq preferred protein accession, protein length, and protein name. Identical proteins are presented in a tabular format that includes information on the database source (e.g., RefSeq, INSDC, etc.), the corresponding nucleotide CDS accession and location, the organism name, and the superkingdom. The displayed table can be downloaded for further use, and is also available through Eutils.

The Identical Protein Report display setting provides important functions, such as:

- A mapping table between protein accessions and the nucleotide record(s) on which they are annotated, when relevant:
- For the RefSeq autonomous non-redundant protein dataset, a mapping table to the organisms to which the protein is relevant;
- And identification of highly conserved proteins when the identical protein sequence is found annotated on divergent species.