



# The New BLAST® Results Page

Enhanced graphical presentation and added functionality

<http://blast.ncbi.nlm.nih.gov/>

National Center for Biotechnology Information • National Library of Medicine • National Institutes of Health • Department of Health and Human Services

## Scope

NCBI has introduced an enhanced report for search results generated by the BLAST® web service. This will eventually supersede the current display first introduced in April 2006. The new report provides access to displays of the alignment results in the NCBI Sequence Viewer and adds a function for downloading sequences for the aligned regions. It also provides easily accessible links to related information for matched sequences and a Description table with customizable columns. Additionally, a benefit of this update is the optimization of BLAST result delivery, which allows the browser to stay responsive to user input while the alignment data is rendered. This is especially beneficial for displaying results containing large numbers of matches.



## Access to BLAST and the new BLAST result page

Access to NCBI BLAST web services through its homepage ([blast.ncbi.nlm.nih.gov](http://blast.ncbi.nlm.nih.gov/)) remains the same as before. The change is in the layout of the results, which are displayed after a search is run.

The enhanced report format is the default display. The YouTube link (A) below points to a video tutorial explaining this new report format. The link to this handout is given to the right. Two ways are available to set the result display back to the old format, particularly with other formatting adjustments are also necessary:

- Using the “Formatting options” link (B) by checking the “Old view” checkbox in the form and then clicking the “Reformat” button.
- Checking the “Old view” checkbox in the “Form request” form, which can be reached after entering a valid RID in the “Recent Results” tab and clicking the “Go” button (C)

BLAST® Basic Local Alignment Search Tool

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NCBI/ BLAST/ blastn suite/ Formatting Results - 6639UARX016

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ref|NM\_000249| (2662 letters)

RID 6639UARX016 (Expires on 10-21 06:52 am)

Query ID gi|263191547|ref|NM\_000249.3| Database Name Human G+T (2 databases)

Description Homo sapiens mutL homolog 1 (MLH1), transcript variant 1, mRNA Description [See details](#)

Molecule type nucleic acid Program BLASTN 2.2.28+ [Citation](#)

Query Length 2662

Other reports: [Search Summary](#) [Taxonomy reports](#) [Distance tree of results](#) [Genome view](#)

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Formatting options [Reformat](#)

Show Alignment as HTML  Old View [Reset form to defaults](#)

Alignment View Pairwise [Reset form to defaults](#)

BLAST® Basic Local Alignment Search Tool

Home Recent Results Saved Strategies Help

NCBI/ BLAST/ Recent Results

Links to your unexpired BLAST jobs appear below. [more...](#)

Lookup BLAST Job

Request ID:  [Go](#)

NCBI/ BLAST/ Format Request

Query ref|NM\_000249| (2662 letters)

Database GPIPE/9606/current/all\_top\_level GPIPE/9606/current/mna

Job title ref|NM\_000249| (2662 letters)

Request ID  [View report](#)  Show results in a new window

Format

Show Alignment as HTML  Old View [Reset form to defaults](#)

Alignment View Pairwise [Reset form to defaults](#)

Display  Graphical Overview  NCBI-gi  CDS feature [Reset form to defaults](#)

Masking Character: Lower Case Color: Grey [Reset form to defaults](#)

## The new BLAST result pages

The general structure of the BLAST result page has not changed. This consists of the following sections: Summary, Graphical Overview, Descriptions table, and Alignments section. The changes are predominantly in the Descriptions table and the Alignments sections.

### The Descriptions table

The Descriptions table (shown below) provides a summary of the database sequences identified by BLAST to be similar to the input query. Two selection controls at the top of the table, "All" and "None" (A), allow for the quick selection and de-selection of matched database sequences. Individual sequences in the table can be selected/de-selected using the checkboxes to the left (B). Selecting database sequences activates links (C) at the top (see details on p.3). From left to right, the Descriptions table columns provide the following information:

- the description/title of matched database sequence
- the highest alignment score (Max score) from that database sequence
- the total alignment scores (Total score) from all alignment segments
- the percentage of query covered by alignment to the database sequence
- the best (lowest) Expect value (E value) of all alignments from that database sequence
- the highest percent identity (Max ident) of all query-subject alignments, and
- the Accession of the matched database sequence

Clicking a column header (D) changes the default column used for sorting the table. The default sorting is by E-value.

For example, when aligning an mRNA to an assembled genome, sorting by "Query coverage" column could help bring the true alignments with lower scores to the top (E). These true alignments might be of low scores due to breakage along the intron/exon boundaries. Columns shown in the table can be customized by using the column selecting palette activated through clicking on the gear icon (F).

The screenshot shows the BLAST Descriptions table with several annotations:

- A**: Select controls (All, None, Selected:7)
- B**: Checkboxes for selecting individual sequences
- C**: Action links (Download, GenPept, Graphics, Distance tree of results, Multiple alignment)
- D**: Column headers (Max score, Total score, Query coverage, E value, Max ident, Accession)
- E**: A sequence title (Rhesus Macaque BAC CH250-38L3) highlighted in red
- F**: Gear icon for column selection

The "Available columns" palette shows the following options:

- Description
- Max Score
- Total Score
- Coverage
- E-value
- Ident
- Accession

Two alternative sorting options are shown:

**Default: sort by E-value**

Description	Max score	Total score	Query coverage	E value	Max ident
creatine kinase B-type [Homo sapiens] >qil332843151[refl]XP_510185.3[PREDICTED: creatine kinase	639	639	100%	0.0	100%
creatine kinase B-type [Macaca mulatta]	637	637	100%	0.0	99%
creatine kinase B-type [Bos taurus]	630	630	100%	0.0	97%
creatine kinase, brain [Sus scrofa]	629	629	100%	0.0	97%
creatine kinase B-type [Oryctolagus cuniculus]	626	626	100%	0.0	97%
creatine kinase B-type [Mus musculus]	625	625	100%	0.0	97%
creatine kinase B-type [Rattus norvegicus]	625	625	100%	0.0	97%
putative creatine kinase B variant 1 [Taeniopygia guttata]	625	625	100%	0.0	97%

**One alternative: sort by Query coverage**

Description	Max score	Total score	Query coverage	E value	Max ident
Rhesus Macaque BAC CH250-38L3 () complete sequence	433	3077	96%	3e-117	100%
Homo sapiens cell division cycle 20 homolog (S. cerevisiae) pseudogene 1 (CDC20f	2582	2582	94%	0.0	96%
Human DNA sequence from clone RP11-276H19 on chromosome 9 Contains the 3'	2582	2582	94%	0.0	96%
Homo sapiens CDC20 cell division cycle 20 homolog (CDC20) gene, complete cds	411	2786	92%	1e-110	100%
Homo sapiens chromosome 1 clone RP11-282K6, complete sequence	411	2792	92%	1e-110	100%
Human DNA sequence from clone RP1-92014 on chromosome 1p33-34.2 Contains	411	2786	92%	1e-110	100%

In the table body, clicking a sequence title (G) quickly scrolls the page display to the alignment section for that sequence where details of the alignment can be examined. Clicking an accession (H) retrieves that record from the corresponding sequence database.

## The Descriptions table: functions provided by the links at the top

At the top of the Descriptions table, clicking the “Alignments” link (A) scrolls the display to the Alignments section. The remaining links work on the selected database sequence(s) for which checkboxes have been checked:

- “Download” activates a menu to select download format (B). The first three options select full sequence records or their aligned regions. The remaining options select the alignment data for these selected sequences.
- “GenBank” or “GenPept” (C) retrieves the selected sequences from the source database.
- “Graphics” (D) spawns a new browser window (or tab depending on the browser setting) to display the summary of query-anchored alignments in the NCBI Sequence Viewer. It is recommended that an accession or gi (with subsequence range if necessary) be used to help make this display more informative.
- “Distance tree of results” (E) opens a new page depicting the relationship among the selected database sequences and the query in a dendrograph. The distances for the tree view are derived from the pairwise local alignment between query and that of selected database sequences. Functions are available on this page to manipulate the display.
- For results from blastp searches, a “Multiple alignment” link (not shown) will be available within this group. Clicking this link performs multiple sequence alignment for the query protein and selected database sequences using the Constraint-based alignment tool (COBALT).

The screenshot illustrates the NCBI BLAST Results page. At the top, the "Descriptions" table shows "Sequences producing significant alignments:" with 13 selected. A yellow arrow labeled 'A' points to the "Alignments" link. A yellow arrow labeled 'B' points to the "Download" dropdown menu, which is open to show options like "FASTA (complete sequence)", "FASTA (aligned sequences)", "GenBank (complete sequence)", "Hit Table (text)", "Hit Table (CSV)", "Text", "XML", and "ASN.1". A yellow arrow labeled 'C' points to the "GenBank" link. A yellow arrow labeled 'D' points to the "Graphics" link. A yellow arrow labeled 'E' points to the "Distance tree of results" link, which opens a "Tree view for RID: 4MDFVJTE014, query ID: NM\_148898.3, database: refseq\_rna". The tree view shows a dendrogram with various species and their forkhead box P2 (FOXP2) mRNA sequences. A yellow arrow labeled 'F' points to the "Tools" and "Configure" menus in the Sequence Viewer. The Sequence Viewer displays the query sequence (NM\_148898.3: 1.6.4K (6.4Kbp)) and its alignment with several database sequences (e.g., NG\_007491.2, AP005269.1, AC199014.3, AC020606.7, AC146090.2). A red box highlights the "Cleaned Alignments" section, which shows a summary view of the alignment coverage from each of the selected database sequences, one subject sequence per row. A red box also highlights the "Tools" and "Configure" menus, with a text box explaining that using an accession or gi as input query enables the SV display to retrieve and display the feature annotation to make the graphical presentation more informative.

The graphical display in Sequence Viewer can be customized using the “Tools” and “Configure” (F) menu and controls. More information on Sequence Viewer is available through the handout and help document accessible online:

- Sequence Viewer help document [www.ncbi.nlm.nih.gov/projects/sviewer/help.html](http://www.ncbi.nlm.nih.gov/projects/sviewer/help.html)
- Sequence Viewer handout [ftp.ncbi.nih.gov/pub/factsheets/Factsheet\\_Graphical\\_SV.pdf](http://ftp.ncbi.nih.gov/pub/factsheets/Factsheet_Graphical_SV.pdf)

## The Alignments section

The Alignments section (A, below) contains the detailed pairwise alignments between query and database sequences. Segments of alignments, also known as high scoring pairs (HSPs), from the same database sequence are grouped under the same separator (B) and are sorted by ascending E-value. Links within the separator, from left to right, provide the following functions:

- “Download” enables downloading the database sequence or its aligned region (C).
- “GenBank” retrieves the database sequence record from the source database.
- “Graphics” presents the alignment in Sequence Viewer (D) for interactive examination under the context of the database sequence and its feature annotation.
- “Sort by” (E) pull-down menu enables sorting of the HSPs from the same database sequence by specific orders, such as by the “Query start position” (F) to place mRNAs in a natural order to genomic alignments.
- “Next” and “Previous” allows for quick navigation in the Alignments section.

On the right side of individual alignments, the new layout also prominently displays additional information available for each database sequence from NCBI databases such as Gene, UniGene, Map Viewer, GeoProfiles, and Structure (G). The detailed alignment display remains essentially the same as the old display, with alignment statistics now summarized in tables (H) given at the top of each HSP. The “Next Match” and “Previous Match” links (I) provide quick navigation among different HSPs from the same database sequence.

The screenshot displays the BLAST Alignments section with several key components highlighted by yellow callouts:

- A:** The main alignment list header, including options for Download, GenBank, Graphics, and Sort by (E value).
- B:** The separator between different High Scoring Pairs (HSPs) from the same database sequence.
- C:** The Download menu, which allows users to download the database sequence or its aligned region.
- D:** The Sequence Viewer (Graphics) showing a genomic map with the query sequence aligned to the database sequence.
- E:** The Sort by pull-down menu, which allows users to sort HSPs by various criteria.
- F:** The Query start position pull-down menu, which allows users to sort HSPs by their query start position.
- G:** The Related Information section, which provides additional information about the database sequence, such as Gene, UniGene, Map Viewer, GeoProfiles, and Structure.
- H:** The HSP summary table, which provides a summary of the alignment statistics for each HSP.
- I:** The Next Match and Previous Match links, which allow users to navigate between different HSPs from the same database sequence.

## Feedback

We welcome feedback on the new BLAST report and have provided a feedback link in the BLAST result page. Comments, bug reports and feedback can be sent to [blast-help@ncbi.nlm.nih.gov](mailto:blast-help@ncbi.nlm.nih.gov), while questions on other NCBI resources should be addressed to [info@ncbi.nlm.nih.gov](mailto:info@ncbi.nlm.nih.gov).