

BLAST

**Slides adapted & edited from a set by
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Kerfeld CA, Scott KM (2011) Using BLAST to Teach “E-value-tionary” Concepts.
PLoS Biology 9(2):e1001014

Starts with a Query Sequence in FASTA Format

Amino acid sequence:

```
>ribosomal protein L7/L12 [Thiomicrospira crunogena XCL-2]
MAITKDDILEAVANMSVMEVELVEAMEEKFGVSAAVAVAGPAGDAGAA
GEEQTEFDVVLTGAGDNKVAIAKAVRGATGLGLKEAKSAVESAPFTLKEG
VSKEEAETLANELKEAGIEVEVK
```

Note the description line
Starts with ">", ends with carriage return
Not read as sequence data

Nucleotide sequence:

```
>gi|118139508:333094-333465 Thiomicrospira crunogena XCL-2
ATGGCAATTACAAAAGACGATATTTAGAAGCAGTTGCTAACATGTCAGTAATGGAAG
TTGTTGAACCTGTTGAAGCAATGGAAGAGAAGTTGGTGTTCCTGCAGCAGCAGTTGC
GGTTGCAGGTCTGCAGGTGATGCTGGCGCTGCTGGTAAGAACAAACAGAGTTGAC
GTTGCTTGACTGGTGTGGTACAACAAAGTTGCAGCAATCAAAGCCGTTCGTGGCG
CAACTGGTCTTGGCTTAAAGAAGCAGAAACTCTTGCAAATGAGCTTAAAGAAGCA
TAAAGAGGGTGTCTAAAGAAGAAGCAGAAACTCTTGCAAATGAGCTTAAAGAAGCA
GGTATTGAAGTCGAAGTTAAATAA
```

NCBI BLAST Interface

(blastp: for protein-protein alignments)

NCBI BLAST Home Recent Results Saved Strategies Help

Basic Local Alignment Search Tool

NCBI BLAST/blastp suite

blast blastp blastx tblastn tblastx

Enter Query Sequence

BLASTP programs search protein databases using a protein query. more...

Reset page Bookmark

Enter accession number, gi, or FASTA sequence

(Paste FASTA format sequence here)

From
To

Or, upload file

Job Title Enter a descriptive title for your BLAST search.

Align two or more sequences

Choose Search Set

Database Non-redundant protein sequences (nr)

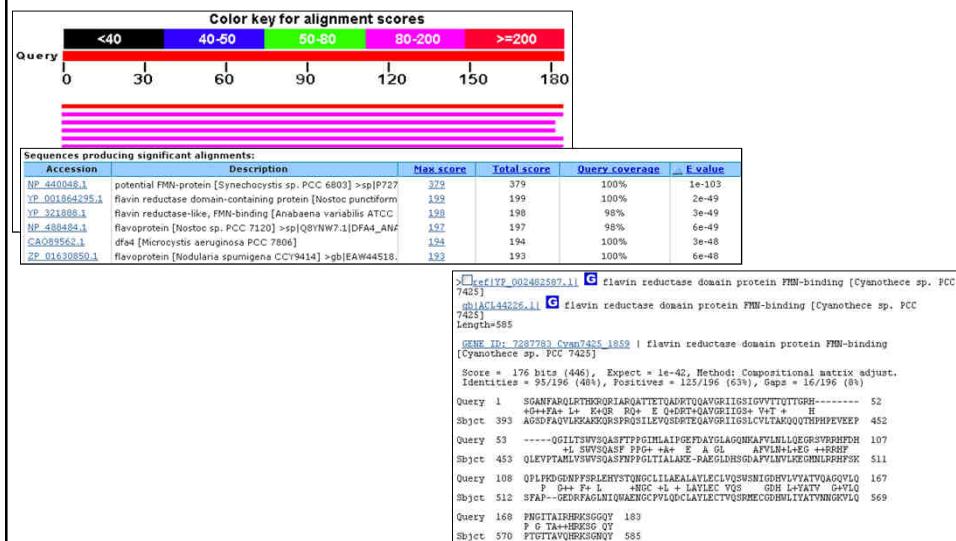
Organism Optional Enter organism name or id-completions will be suggested Exclude

Enter organism common name, binomial, or tax id. Only 20 top taxa will be shown.

Exclude Optional Models (Y/M/X/P) Uncultured/environmental sample sequences

Entrez Query Optional Enter an Entrez query to limit search

NCBI BLAST Results Page: Potential homologs retrieved from database



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Overview of BLAST

1. Segment the query sequence into short “words”
2. Use the query sequence segments to scan the database for matching sequences
3. Extend the matched segments in either direction to find local alignments.
4. Create a list of hits & alignments, with best matches first

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BLAST Phase 1: Segment the query sequence and identify words that could form potential alignments

Query Sequence:

>gi|16329320 (residues 412 to 594)
SGANFARQLRTHKRQRIARQATTETQADRTQQAVGRIIGSIGVVTQTTG
RHQGILTSWVSQASFTEPGIMLAIPGEEFDAYGLAQQNKAFLNLQEGRS
VRRHFDHQPLPKDGDNPFSRLEHYSTQNGCLILAEALAYLECLVQSWSNI
GDHVLVYATVQAGQVLPNGITAIRHRKSGGQY

Fragmentation into words:

SWVSQASFTEPGIM → SWV WVS VSQ SQA QAS ASF SFT ...

Selection of words scoring above threshold (for word SWV):

Substitution Matrix*							
R	G	I	K	F	S	T	W
R	5	-1	-1	-2	1	0	-3
G	6	-4	-2	-3	0	-2	-3
I	4	-3	0	-2	-1	-3	3
K		5	-3	0	-1	-3	-2
F		6	-2	2	1	-1	
S			4	1	-3	-2	
T				5	-2	0	
W					11	-3	
V						4	

*A portion of the BLOSUM 62 matrix

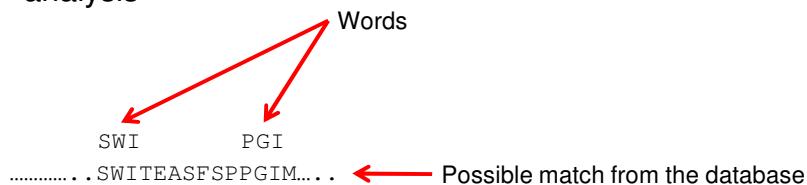
- Segment the query sequence into pieces (“words”)
 - Default word length: 3 amino acids or 11 nucleic acids
- Create a list of synonyms and their scores for comparing query words to target words
 - Uses scoring matrix to calculate scores for synonyms that might be found in the database
- Save the scores (and synonyms) exceeding a given threshold T

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BLAST Phase 2: Using the query sequence word list, scan the database for synonyms (hits)

- Scan the database for matches to the word list with acceptable T values
- Require two matches (“hits”) within the target sequence
- Set aside sequences with matches above T for further analysis

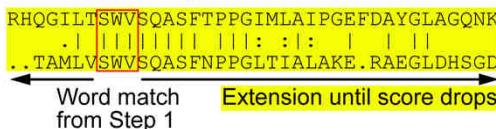


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BLAST Phase 3: Extending the hits

- Search 5' and 3' of the word hit on both the query and target sequence
- Add up the score for sequence identity or similarity until value exceeds S
- Alignment is dropped from subsequent analyses if value never exceeds S



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So, to summarize:

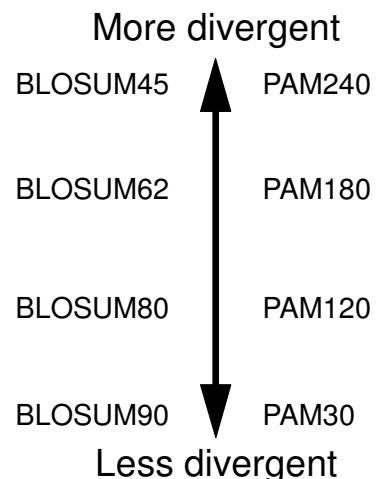
- BLAST segments query sequence into “words” and scores potential word matches
- Scans this list for alignments that meet a threshold score T
 - uses a scoring matrix to calculate this (e.g., **BLOSUM62**)
- Uses this list of ‘synonyms’ to scan the database
- Extends the alignments to see if they meet a cutoff score S
 - uses a scoring matrix to calculate this
- Reports the alignments that exceed S

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PAM and BLOSUM Matrices

- Scoring matrices are calibrated to capture different degrees of sequence similarity
- In practice, this means choosing a matrix appropriate to the suspected degree of sequence identity between the query and its hits
- PAM: empirically derived for close relatives
- BLOSUM: empirically derived for distant relatives



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Raw Scores (S values) from an Alignment

$$S = (\sum M_{ij}) - cO - dG,$$

where

M = score from a similarity matrix

for a particular pair of amino acids (ij)

c = number of gaps

O = penalty for the existence of a gap

d = total length of gaps

G = per-residue penalty for extending
the gap

Limitations of Raw Scores

- S values depend on the substitution matrix, gap penalties
- Impossible to compare S values from hits retrieved from BLAST searches when different matrices and gap penalties are used

Going from Raw Scores to Bit Scores

$$S' = [\lambda S - \ln(K)] / \ln(2)$$

where

S' = bit score

λ and K = normalizing parameters of the specific matrices and search spaces

↑
(as in 0 vs 1)

- Larger raw scores result in larger bit scores
- Allows user to compare scores obtained by using different matrices and search spaces

Limitations of Bit Scores

- How high does a bit score have to be to suggest common ancestry?
 - Hard to evaluate hits as homologs or not, based solely on bit scores

E-value

- Number of distinct alignments with scores greater than or equal to a given value expected to occur in a search against a database of known size, based solely on chance, not homology.
 - Large E-values suggest that the query sequence and retrieved sequence similarities are due to chance
 - Small E-values suggest that the sequence similarities are due to shared ancestry (or potentially convergent evolution)

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Calculating E-values

$$E = (n \times m) / 2^S$$

where

m = effective length of the query sequence
= length of query sequence – average length of alignments
(Controls for fewer alignments occurring at the ends
of the query sequence)

n = effective length of the database sequence
(total number of bases)

The value of E decreases exponentially with increasing S

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BLAST Parameters

- Expect
- Word size
- Matrix
- Gap costs
- Filter
- Mask

The screenshot shows the 'Algorithm parameters' section of the BLAST search interface. It includes fields for 'Max target sequences' (set to 100), 'Expect threshold' (set to 10), 'Word size' (set to 3), 'Matrix' (set to BLOSUM62), 'Gap Costs' (set to Existence: 11 Extension: 1), and 'Compositional adjustments' (set to Conditional compositional score matrix adjustment). Below this is the 'Filters and Masking' section with options for 'Filter' (Low complexity regions) and 'Mask' (Mask for lookup table only, Mask lower case letters). At the bottom is a 'BLAST' button and search options.

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E value Threshold

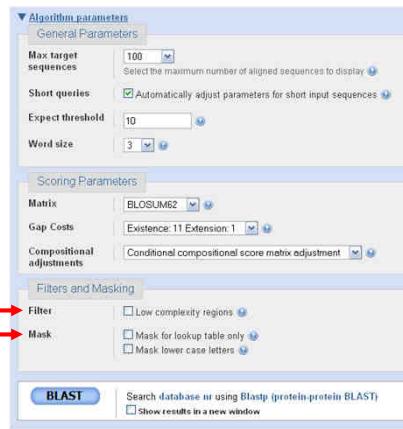
- Alignments will be reported with E-values less than or equal to the expect values threshold
 - Setting a larger E threshold will result in more reported hits
 - Setting a smaller E threshold will result in fewer reported hits

The screenshot shows the 'Algorithm parameters' section of the BLAST search interface. It includes fields for 'Max target sequences' (set to 100), 'Expect threshold' (set to 10), 'Word size' (set to 3), 'Matrix' (set to BLOSUM62), 'Gap Costs' (set to Existence: 11 Extension: 1), and 'Compositional adjustments' (set to Conditional compositional score matrix adjustment). Below this is the 'Filters and Masking' section with options for 'Filter' (Low complexity regions) and 'Mask' (Mask for lookup table only, Mask lower case letters). At the bottom is a 'BLAST' button and search options. A red arrow points to the 'Expect threshold' field.

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Filter and Mask

- Filter: Low complexity
 - Replaces the following with N (nucleotides) or X (amino acids)
 - Dinucleotide repeats
 - Amino acid repeats
 - Leader sequences
 - Stretches of hydrophobic residues
- Mask: Lower case
 - Replaces lowercase letters in sequence with N or X
 - Lowercase letters typically indicate base or amino acid not known with certainty



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Parameter Summary is Found at the Bottom of the Output.....

Search Parameters		
Program:	blastp	
Word size:	3	
Expect value:	10	
Hitlist size:	100	
Gapcosts:	11,1	
Matrix:	BLOSUM62	
Filter string:	F	
Genetic Code:	1	
Window Size:	40	
Threshold:	11	
Composition-based stats:	2	
Database		
Posted date:	Sep 6, 2010 4:42 AM	
Number of letters:	4,014,994,744	
Number of sequences:	11,756,863	
Entrez query:	none	
Karlin-Altschul statistics		
Lambda	0.319424	0.267
K	0.13352	0.041
H	0.397413	0.14
Results Statistics		
Length adjustment:	129	
Effective length of query:	54	
Effective length of database:	2498359417	
Effective search space:	134911408518	
Effective search space used:	134911408518	

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Evaluating BLAST Results

Color key for alignment scores

Query	<40	40-50	50-80	80-200	>=200	
0	30	60	90	120	150	180

Sequences producing significant alignments:

Accession	Description	Max score	Total score	Query coverage	E value
NP_440481.1	potential FMN-protein [Synechocystis sp. PCC 6803] >sp P727	379	379	100%	1e-103
YP_001864285.1	flavin reductase domain-containing protein [Nostoc punctiforme]	193	199	100%	2e-49
YP_321888.1	flavin reductase-like, FMN-binding [Anabaena variabilis ATCC 198	198	198	98%	3e-49
NP_488484.1	flavoprotein [Nostoc sp. PCC 7120] >sp Q8YNW7.1 DFA4_ANA	197	197	98%	6e-49
CA089562.1	dfa4 [Microcystis aeruginosa PCC 7806]	194	194	100%	3e-48
ZP_01630850.1	flavoprotein [Nodularia spumigena CCY9414] >gb EAW44518.	193	193	100%	6e-48

Sequence details:

```

>ref|YP_002482587.1| G flavin reductase domain protein FMN-binding [Cyanothec sp. PCC 7425]
7425)                               G flavin reductase domain protein FMN-binding [Cyanothec sp. PCC 7425)
Length=585
GENE ID: 7287783 Cyan7425_1859 | flavin reductase domain protein FMN-binding
(Cyanothec sp. PCC 7425)

Score = 176 bits (446), Expect = 1e-42, Method: Compositional matrix adjust,
Identities = 95/196 (48%), Positives = 125/196 (63%), Gaps = 16/196 (8%)
Query 1      SGANFARQLRTHKRQRRIARQATTEQQADRTQQA VGRRIIGSIVGVVTTQTGRRH----- 52
+G++FA+ L+ K+QR RQ+ E Q+DRT+QAVGRRIIGS+ V+T + H
Sbjct 393 AGSDFAQVLKKAKKQRSPRQSILEVQSDRTEQAVGRRIIGSCLVLTAKQQTHPHEVEEP 452
-----+QGILTSWVSQASFTPPGIMLAIPGEFDAYGLAGQNKA FVNLLQEGRSWRHF D 107
Query 53      -----+QGILTSWVSQASFTPPGIMLAIPGEFDAYGLAGQNKA FVNLLQEGRSWRHF D----- 52
+SUVSQASF PPG+ +A+ E A GL AFVLM+L+EG ++RRHF
Sbjct 453 QLEVP TAMLVSVVSQASFNPPLGTTIALAKE-RAEGLDHSGDAFVLNVLKEGMNLLRHF SK 511
-----+QPLPKGDGDNFSRSLREHYSTQNGCLILAEALAYLECLVQWSWSNIGDHVLVYATVQAGQV LQ 167
Query 108     QPLPKGDGDNFSRSLREHYSTQNGCLILAEALAYLECLVQWSWSNIGDHVLVYATVQAGQV LQ 167
P G++ F+ L +NGC +L + LAYLEC VQS GDH L+YATV G+V LQ
Sbjct 512 SFAP--GE DRFAGLNQIWAENGCPVLDCLAYLECTVQSRMECGDHWLYATVNNNGKV LQ 569
-----+PNGITAIRHRKSGQY 183
P G TA++HHRKSG QY
Sbjct 570 PTGTTAVQHRKSGNQY 585

```

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Examine the BLAST Alignment

Sequence details:

```

>ref|YP_002482587.1| G flavin reductase domain protein FMN-binding [Cyanothec sp. PCC 7425]
7425)                               G flavin reductase domain protein FMN-binding [Cyanothec sp. PCC 7425)
Length=585
Does it cover the whole length of both the query and subject sequences?
GENE ID: 7287783 Cyan7425_1859 | flavin reductase domain protein FMN-binding
(Cyanothec sp. PCC 7425)

Score = 176 bits (446), Expect = 1e-42, Method: Compositional matrix adjust,
Identities = 95/196 (48%), Positives = 125/196 (63%), Gaps = 16/196 (8%)
Query 1      SGANFARQLRTHKRQRRIARQATTEQQADRTQQA VGRRIIGSIVGVVTTQTGRRH----- 52
+G++FA+ L+ K+QR RQ+ E Q+DRT+QAVGRRIIGS+ V+T + H
Sbjct 393 AGSDFAQVLKKAKKQRSPRQSILEVQSDRTEQAVGRRIIGSCLVLTAKQQTHPHEVEEP 452
-----+QGILTSWVSQASFTPPGIMLAIPGEFDAYGLAGQNKA FVNLLQEGRSWRHF D 107
Query 53      -----+QGILTSWVSQASFTPPGIMLAIPGEFDAYGLAGQNKA FVNLLQEGRSWRHF D----- 52
+SUVSQASF PPG+ +A+ E A GL AFVLM+L+EG ++RRHF
Sbjct 453 QLEVP TAMLVSVVSQASFNPPLGTTIALAKE-RAEGLDHSGDAFVLNVLKEGMNLLRHF SK 511
-----+QPLPKGDGDNFSRSLREHYSTQNGCLILAEALAYLECLVQWSWSNIGDHVLVYATVQAGQV LQ 167
Query 108     QPLPKGDGDNFSRSLREHYSTQNGCLILAEALAYLECLVQWSWSNIGDHVLVYATVQAGQV LQ 167
P G++ F+ L +NGC +L + LAYLEC VQS GDH L+YATV G+V LQ
Sbjct 512 SFAP--GE DRFAGLNQIWAENGCPVLDCLAYLECTVQSRMECGDHWLYATVNNNGKV LQ 569
-----+PNGITAIRHRKSGQY 183
P G TA++HHRKSG QY
Sbjct 570 PTGTTAVQHRKSGNQY 585

```

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High E-value: Discovery of a Distant Homolog or Garbage?

- Take another look at the target (subject) sequence(s) that have high E-values
 - Similar length?
 - Recurring motifs?
 - Similar biological functions?
- Use target sequences as query sequences for another BLAST search
 - Does the original query sequence come up in report?