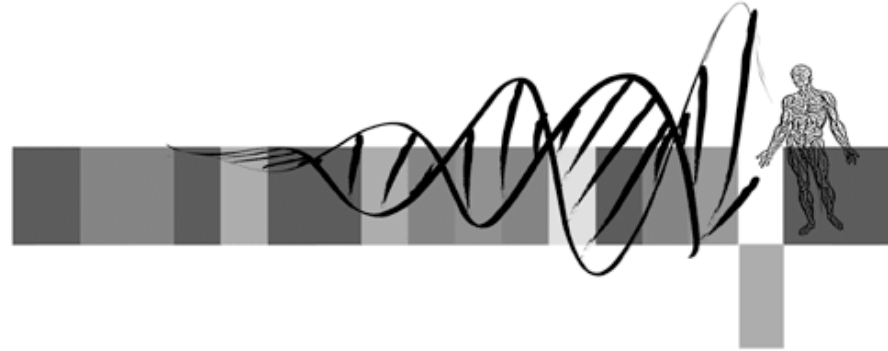




Bioinformatics for Biologists

Sequence Analysis: Part I. Pairwise alignment and database searching

Fran Lewitter, Ph.D.
Head, Biocomputing
Whitehead Institute

Bioinformatics Definitions

“The use of computational methods to make biological discoveries.”

Fran Lewitter

“An interdisciplinary field involving biology, computer science, mathematics, and statistics to analyze biological sequence data, genome content, and arrangement, and to predict the function and structure of macromolecules.”

David Mount

Topics to Cover

- Introduction
- Scoring alignments
- Alignment methods
- Significance of alignments
- Database searching methods

Topics to Cover

- Introduction
 - Why do alignments?
 - A bit of history
 - Definitions
- Scoring alignments
- Alignment methods
- Significance of alignments
- Database searching methods

Simian sarcoma virus onc gene, v-sis, is derived from the gene (or genes) encoding a platelet-derived growth factor.

Doolittle RF, Hunkapiller MW, Hood LE, Devare SG, Robbins KC, Aaronson SA, Antoniades HN. *Science* 221:275-277, 1983.

Cancer Gene Found

Cell, Vol. 75, 1027–1038, December 3, 1993, Copyright © 1993 by Cell Press

The Human Mutator Gene Homolog *MSH2* and Its Association with Hereditary Nonpolyposis Colon Cancer

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can give rise to mismatched bases (Friedberg, 1985). For example, the deamination of 5-methylcytosine creates a thymine and, therefore, a G·T mispair (Duncan and Miller, 1980). Second, misincorporation of nucleotides during DNA replication can yield mismatched base pairs and nucleotide insertions and deletions (Modrich, 1991). Finally,

Homology to bacterial and yeast genes shed new light on human disease process

Evolutionary Basis of Sequence Alignment

- *Similarity* - observable quantity, such as per cent identity
- *Homology* - conclusion drawn from data that two genes share a common evolutionary history; no metric is associated with this

Some Definitions

- An *alignment* is a mutual arrangement of two sequences, which exhibits where the two sequences are similar, and where they differ.
- An *optimal alignment* is one that exhibits the most correspondences and the least differences. It is the alignment with the highest score. May or may not be biologically meaningful.

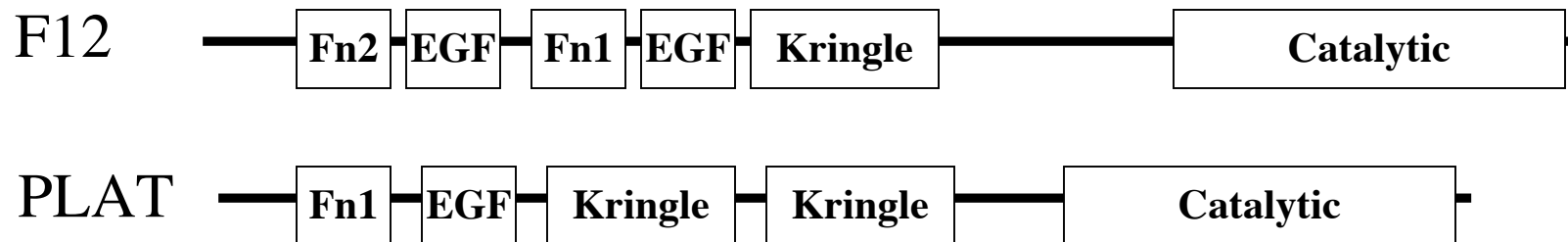
Alignment Methods

- *Global alignment* - Needleman-Wunsch (1970) maximizes the number of matches between the sequences along the entire length of the sequences.
- *Local alignment* - Smith-Waterman (1981) is a modification of the dynamic programming algorithm gives the highest scoring local match between two sequences.

Alignment Methods

Global vs Local

Modular proteins



Possible Alignments

A: T C A G A C G A G T G

B: T C G G A G C T G

I. T C A G A C G A G T G

T C G G A - - G C T G

II. T C A G A C G A G T G

T C G G A - G C - T G

III. T C A G A C G A G T G

T C G G A - G - C T G

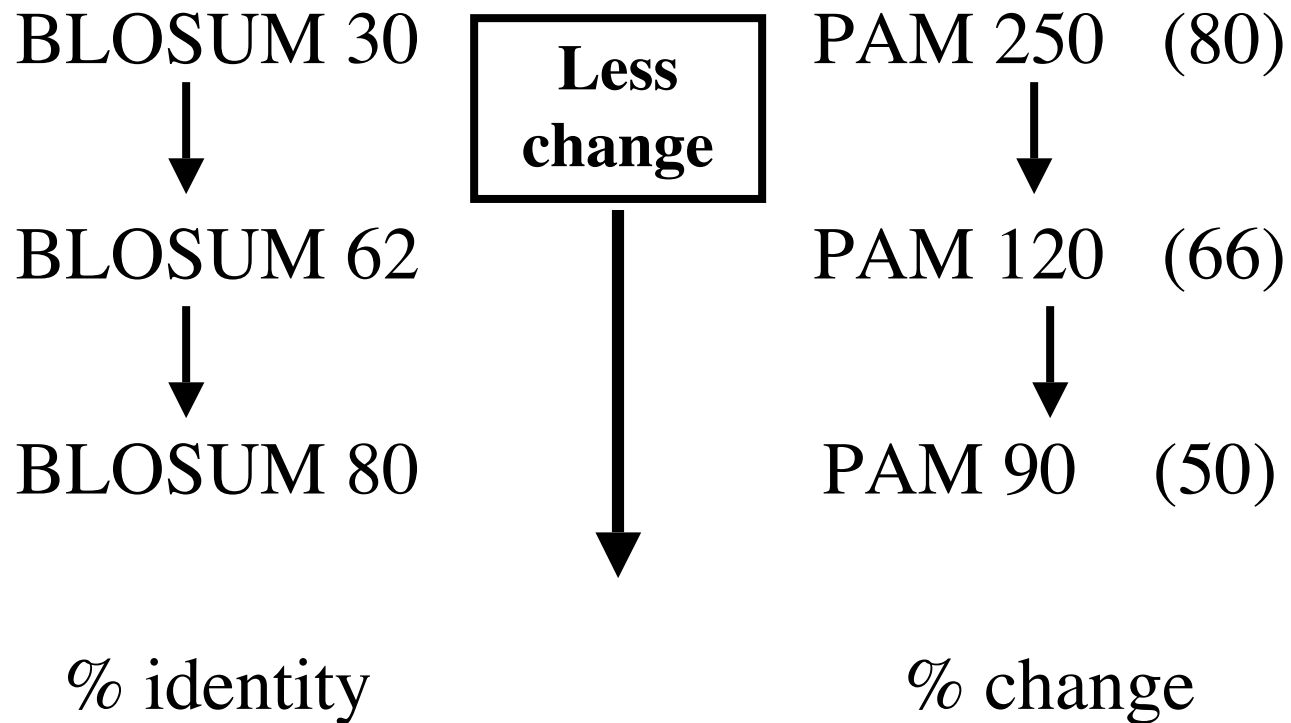
Topics to Cover

- Introduction
- Scoring alignments
 - Nucleotide vs Proteins
- Alignment methods
- Significance of alignments
- Database searching methods

Amino Acid Substitution Matrices

- **PAM** - point accepted mutation based on *global* alignment [evolutionary model]
- **BLOSUM** - block substitutions based on *local* alignments [similarity among conserved sequences]

Substitution Matrices



Part of BLOSUM 62 Matrix

	C	S	T	P	A	G	N
C	9						
S	-1	4					
T	-1	1	5				
P	-3	-1	-1	7			
A	0	1	0	-1	4		
G	-3	0	-2	-2	0	6	
N	-3	1	0	-2	-2	0	

Log-odds = $\frac{\text{obs freq of aa substitutions}}{\text{freq expected by chance}}$

Part of PAM 250 Matrix

	C	S	T	P	A	G	N
C	1	2					
S	0	2					
T	-2	1	3				
P	-3	1	0	6			
A	-2	1	1	1	2		
G	-3	1	0	-1	1	5	
N	-4	1	0	-1	0	0	

Log-odds = $\frac{\text{pair in homologous proteins}}{\text{pair in unrelated proteins by chance}}$

Gap Penalties

- *Insertion and Deletions* (indels)
- *Affine gap costs* - a scoring system for gaps within alignments that charges a penalty for the existence of a gap and an additional per-residue penalty proportional to the gap's length

Example of simple scoring system for nucleic acids

- Match = +1 (ex. A-A, T-T, C-C, G-G)
- Mismatch = -1 (ex. A-T, A-C, etc)
- Gap opening = - 2
- Gap extension = -1

T	C	A	G	A	C	G	A	G	T	G	
T	C	G	G	A	-	-	G	C	T	G	
+1	+1	-1	+1	+1	-2	-1	-1	-1	+1	+1	= 0

Scoring for BLAST 2 Sequences

Score = 94.0 bits (230), Expect = 6e-19

Identities = 45/101 (44%), Positives = 54/101 (52%), Gaps = 7/101 (6%)

Query: 204 YTGPFCDV----DTKASCYDGRGLSYRGLARTTSLGAPCQPWASEATYRNVTAEQ---AR 256

Y+ FC + + CY G G +YRG T SGA C PW S V Q A+

Sbjct: 198 YSSEFCSTPACSEGNSDCYFGNGSAYRGTHSLTESGASCLPWNSMILIGKVYTAQNPSAQ 257

Query: 257 NWGLGGHAFCRNPDNDIRPWCFVLNRDRLSWEYCDLAQCQT 297

GLG H +CRNPD D +PWC VL RL+WEYCD+ C T

Sbjct: 258 ALGLGKHNYCRNPDGDAKPWCHVLKNRRLTWEYCDVPSCST 298

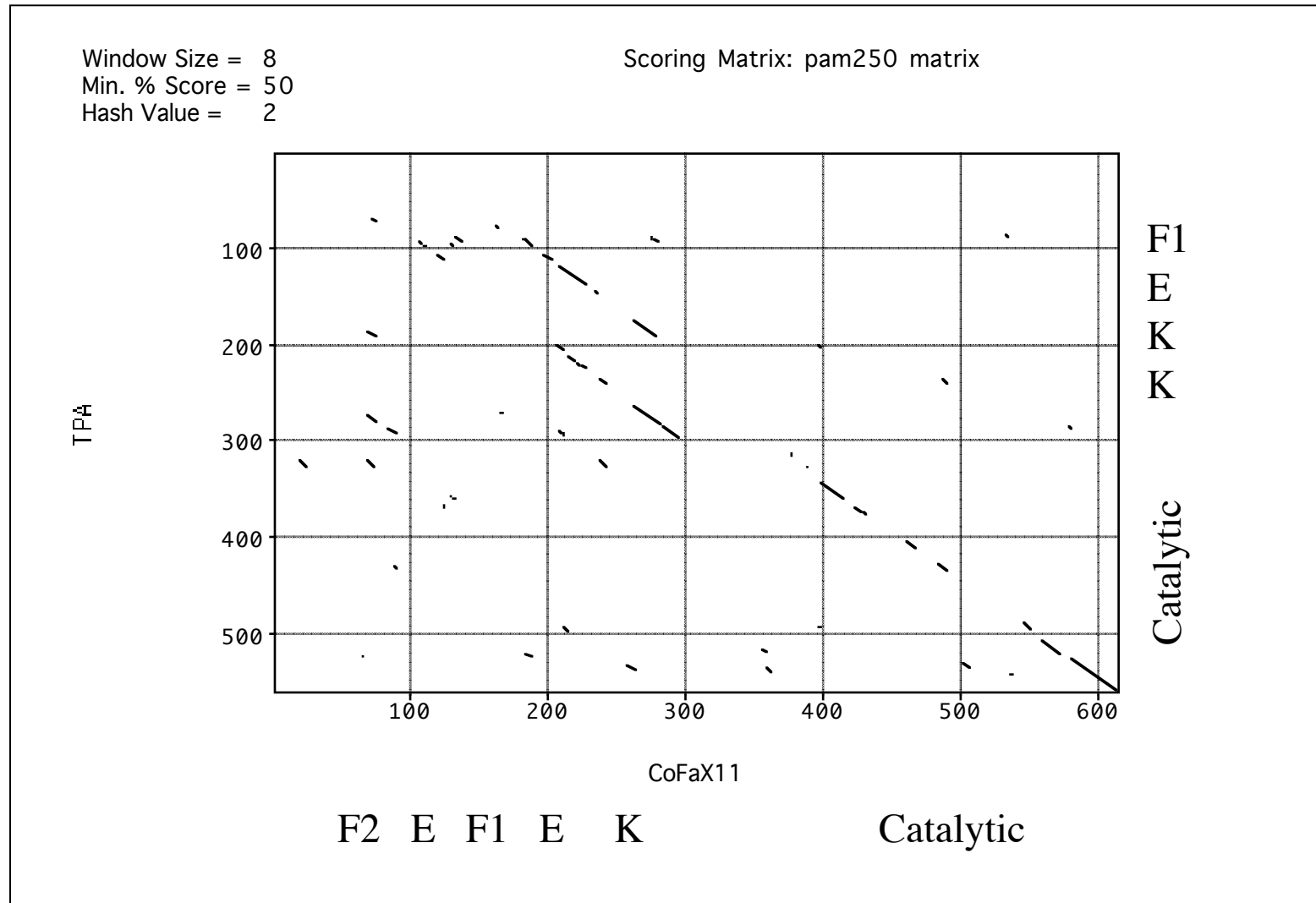
Based on
BLOSUM62

Position	1:	Y	-	Y	=	7
Position	2:	T	-	S	=	1
Position	3:	G	-	S	=	0
Position	4:	P	-	E	=	-1
		.	.	.		
Position	9:	-	-	P	=	-11
Position	10:	-	-	A	=	-1
		.	.	.		
					Sum	230

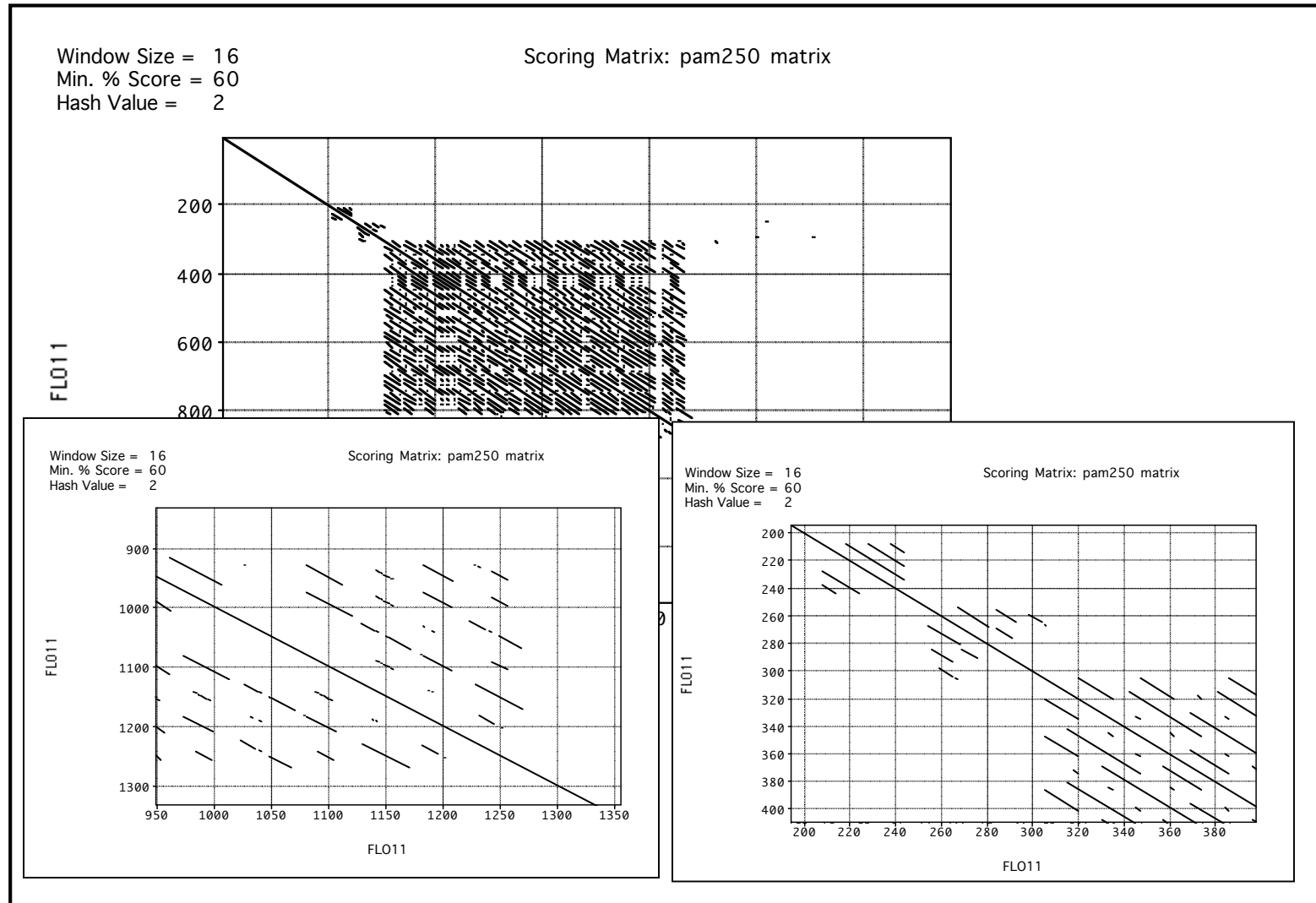
Topics to Cover

- Introduction
- Scoring alignments
- Alignment methods
 - Dot matrix analysis
 - Exhaustive methods; Dynamic programming algorithm (Smith-Waterman (Local), Needleman-Wunsch (Global))
 - Heuristic methods; Approximate methods; word or k-tuple (FASTA, BLAST, BLAT)
- Significance of alignments
- Database searching methods

Dot Matrix Comparison



Dot Matrix Comparison



Dynamic Programming

- Provides very best or optimal alignment
- Compares every pair of characters (e.g. bases or amino acids) in the two sequences
- Puts in gaps and mismatches
- Maximum number of matches between identical or related characters
- Generates a score and statistical assessment
- Nice example of global alignment using N-W:
<http://www.sbc.su.se/~per/molbioinfo2001/dynprog/dynamic.html>

Global vs Local Alignment

(example from Mount 2001)

	GAP	M	N	A	L	S	D	R	T
GAP	0	-12	-16	-20	-24	-28	-32	-36	-40
M	-12	6 ⁽⁶⁾	-6 ⁽⁻²⁾	-10	-14	-18	-22	-26	-30
G	-16	-6 ⁽⁻³⁾	6 ⁽⁰⁾	-5	-10	-13	-17	-22	-26
S	-20	-10	-5	7	-5	-8	-13	-17	-21
D	-24	-14	-8	-5	3	-5	-4	-14	-17
R	-28	-18	-14	-10	-8	3	-6	2	-10
T	-32	-22	-18	-13	-12	-7	3	-7	5
T	-36	-26	-22	-17	-15	-11	-7	2	-4
E	-40	-30	-25	-22	-20	-15	-8	-8	0
T	-44	-34	-30	-24	-24	-21	-15	-9	-5

	GAP	M	N	A	L	S	D	R	T
GAP	0	0	0	0	0	0	0	0	0
M	0	6	0	0	4	0	0	0	0
G	0	0	6	1	0	5	1	0	0
S	0	0	1	7	0	2	5	1	1
D	0	0	2	1	3	0	6	4	1
R	0	0	0	0	0	3	0	12	3
T	0	0	0	1	0	1	3	0	15
T	0	0	0	1	0	1	1	2	3
E	0	0	1	0	0	0	4	0	2
T	0	0	0	2	0	1	0	3	3

sequence 1	M	-	N	A	L	S	D	R	T	
sequence 2	M	G	S	D	R	T	T	E	T	
score	6	-12	1	0	-3	1	0	-1	3	= -5

sequence 1	S	D	R	T	
sequence 2	S	D	R	T	
score	2	4	6	3	= 15

Original "Ungapped" BLAST Algorithm

- To improve speed, use a word based hashing scheme to index database
- Limit search for similarities to only the region near matching words
- Use **Threshold** parameter to rate neighbor words
- Extend match left and right to search for high scoring alignments

Original BLAST Algorithm (1990)

Query word (W=3)

Query: GSVEDTTGSQSLAALLNKCKT**PQG**QRLVNQWIKQPLM

Neighborhood
words

PQG	18	PHG	13
PEG	15	PMG	13
PNG	13	PTG	12
PDG	13	Etc.	

Neighborhood
Score threshold
(T=13)

Query: 325

SLAALLNKCKT**PQG**QRLVNQWIKQPLMDKNRIEERLNLVEA

+LA++L+ TP G R++ +W+ P+ D + ER I A

Sbjct: 290

TLASVLDCTVT**PMG**SRMLKRWLHMPVRDTRVLLERQQTIGA

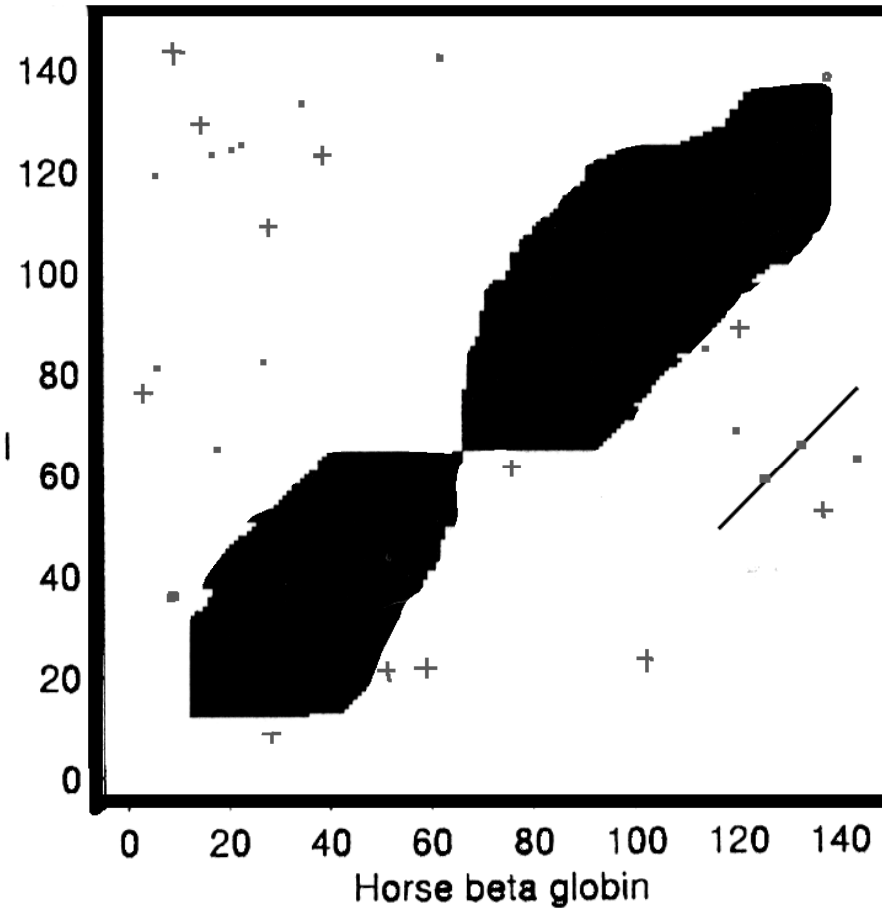
BLAST Refinements (1997)

- “two-hit” method for extending word pairs
- Gapped alignments
- Iterate with position-specific matrix (PSI-BLAST)
- Pattern-hit initiated BLAST (PHI-BLAST)

Gapped BLAST

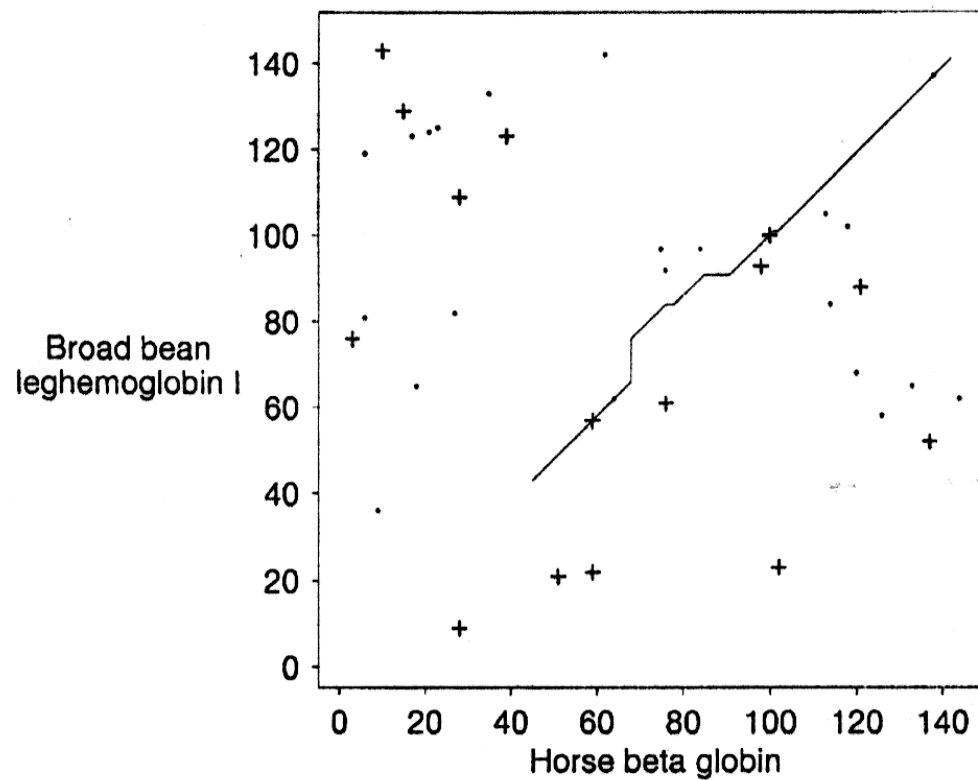
15(+) > 13
22(•) > 11

Broad bean
leghemoglobin I



(Altschul et al 1997)

Gapped BLAST



(Altschul et al 1997)

Programs to Compare two sequences - Unix or Web

NCBI

BLAST 2 Sequences

EMBOSS

water - Smith-Waterman

needle - Needleman - Wunsch

dotmatch (dot plot)

einverted or palindrome (inverted repeats)

equicktandem or etandem (tandem repeats)

Other

lalign (multiple matching subsegments in two sequences)

Topics to Cover

- Introduction
- Scoring alignments
- Alignment methods
- Significance of alignments
- Database searching methods
- Demo

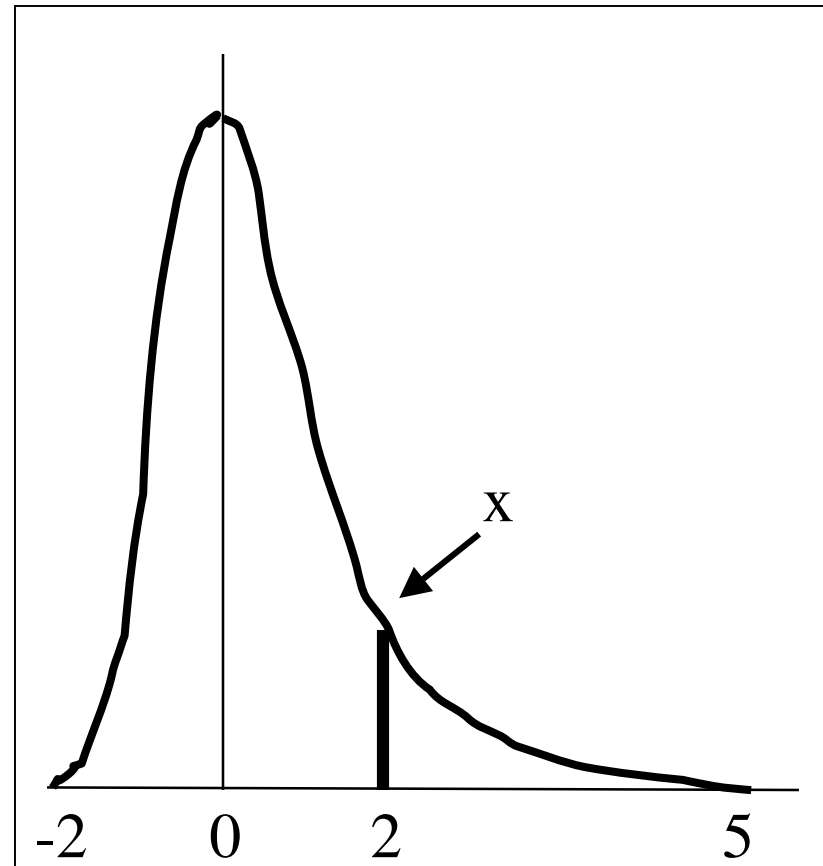
Significance of Alignment

How strong can an alignment be expected by chance alone?

- Real but non-homologous sequences
- Real sequences that are shuffled to preserve compositional properties
- Sequences that are generated randomly based upon a DNA or protein sequence model

Extreme Value Distribution

- When 2 sequences have been aligned optimally, the significance of a local alignment score can be tested on the basis of the distribution of scores expected by aligning two random sequences of the same length and composition as the two test sequences.



Statistical Significance

- **Raw Scores** - score of an alignment equal to the sum of substitution and gap scores.
- **Bit scores** - scaled version of an alignment's raw score that accounts for the statistical properties of the scoring system used.
- **E-value** - expected number of distinct alignments that would achieve a given score by chance. Lower E-value => more significant.

Some formulas

$$E = Kmn e^{-\lambda S}$$

This is the Expected number of high-scoring segment pairs (HSPs) with score at least S for sequences of length m and n .

This is the E value for the score S .

Topics to Cover

- Introduction
- Scoring alignments
- Alignment methods
- Significance of alignments
- Database searching methods
 - BLAST - ungapped and gapped
 - BLAST vs. FASTA
 - BLAT

Questions

- Why do a database search?
- What database should be searched?
- What alignment algorithm to use?
- What do the results mean?

Issues affecting DB Search

- Substitution matrices
- Statistical significance
- Filtering
- Database choices

BLASTP Results

Sequences producing significant alignments:			Score	E	
			(bits)	Value	
gi 34862150 ref XP_345634.1 	similar to mismatch repair pro...		<u>209</u>	5e-54	L
gi 36949366 ref NP_002431.2 	mutS homolog 4; mutS (E. coli)...		<u>162</u>	9e-40	L
gi 34481396 emb CAC79990.1 	sperm protein [Homo sapiens]		<u>152</u>	1e-36	L
gi 34861090 ref XP_227831.2 	similar to MutS homolog 4 [Rat...		<u>147</u>	3e-35	L
gi 34872785 ref XP_213395.2 	similar to hypothetical protei...		<u>33</u>	0.62	L
gi 34853116 ref XP_345138.1 	similar to hypothetical protei...		<u>32</u>	1.3	L
gi 34783109 qb AAH01726.2 	Unknown (protein for IMAGE:35345...		<u>32</u>	1.6	
gi 16307283 qb AAH09731.1 AAH09731	Similar to hypothetical ...		<u>31</u>	3.1	L
gi 34868124 ref XP_221530.2 	similar to mKIAA0719 protein [...		<u>31</u>	3.4	L
gi 34853816 ref XP_344817.1 	similar to FGFR1 oncogene part...		<u>30</u>	7.8	L

Alignments

☐ >[gi|34862150|ref|XP_345634.1|](#) **L** similar to mismatch repair protein MSH6 [Rattus norvegicus]
 Length = 1541

Score = 209 bits (533), Expect = 5e-54
 Identities = 174/617 (28%), Positives = 283/617 (45%), Gaps = 78/617 (12%)

Low Complexity Regions

- **Local regions of biased composition**
- **Common in real sequences**
- **Generate false positives on BLAST search**
- **DUST for BLASTN (n's in sequence)**
- **SEG for other programs (x's in sequence)**

*Filtering is only applied to the query sequence
(or its translation products), not to database
sequences.*

Filtered Sequence

>HUMAN MSH2

NEEYTKNKTEYEE

QGMPEKPTTTVRLFDRGDFYTAHGEDALLAAR
VLSKMNFESEFVKDLLVRQYRVEVYKNRAGNK
ILFGNNDMSASIGVVGKMSAVDQQRQVGVGY
ALLIQIGPKECVLPGGETAGDMGKLRQIIQRG

GILITERKPADFSTKDIYQDLNRLLLKGGKGEQMNSAVLPEMENQVAVSSLSAVIK
FLELLSDDSNFGQFELTTFDLSQYMKLDIAAVRALNLFQGSVEDTTGSQSLAALL
NKCKTPQGQRIVNQWIKQPLMDKNRIEE
DLNRLAKKFQRQAANLQDCYRLYQGINQ
TDLRSDFSKFQEMLETTLDMDQVENHEE
LISAARDLGLDPGRQIKLDSSAQFGYYE
FTNSKLTSLNEEYTKNKTEYEEEAQDAIVKEIVNISSGYVEPMOTLNDVLAQLDAV
VSFAHVSNGAPVPYVRPAILEKGQGRIILKASRHACVEVQDEIAFIPNDVYFEKD
KQMFHIITGPNMGGKSTYIRQTGVIVLMAQIGCFVPCESAEVSIVDCILARVGAG
DSQLKGVSTFMAEMLETASILRSATKDSLIIIDELGRQESTYDGFGLAWAISEYI
ATKIGAFCMFATHFHELTALANQIPTVNNLHVTALTTEETLTMLYQVKKGVCDQS
FGIHVAELANFPKHVIECAKQKALELEEFQYIGESQGYDIMEPAAKKCYLEREQG
EKIIQEFLSKVKQMPFTEMSEENITIKLKQLKAEVIAKNNSFVNEIISRIKVTT

TALTTEETLT

Example Alignment w/o filtering

Score = 29.6 bits (65), Expect = 1.8

Identities = 22/70 (31%), Positives = 32/70 (45%), Gaps = 12/70 (17%)

```
Query: 31   PPPTTQGAPRTSSFTPTTLT-----NGTSHSPTALNGAPSPPNGFS 71
           PPP+ Q   R   S +   T T               NG+S S ++ + + S   +   S
Sbjct: 1221 PPSVQNQQRWGSSSVITTTTCQQRQQSVSPHSNGSSSSSSSSSSSSSSSSSSSTS 1273

Query: 72   NGPSSSSSSSLANQQLP 88
           +   SSSS+SS      Q P
Sbjct: 1274 SNCSSSSASSCQYFQSP 1290
```

Example BLAST w/ filtering

Score = 36.6 bits (83), Expect = 0.67

Identities = 21/58 (36%), Positives = 25/58 (42%), Gaps = 1/58 (1%)

```
Query: 471 AEDALAVINQQEDSSDESCWNCGRKASETCSGCNTARYCGSFCQHKDWE-KHHHICGQT 527
          A D  V  Q + + C  CG  A  TCS C  A YC      Q  DW+  H   C Q+
Sbjct: 61  ASDTECVCLQLKSGAHLCRVCGCLAPMTCSRCKQAHYCSKEHQTLDWQLGHKQACTQS 118
```

Score = 37.0 bits (84), Expect = 0.55

Identities = 18/55 (32%), Positives = 22/55 (39%)

```
Query: 483 DSDESCWNCGRKASETCSGCNTARYCGSFCQHKDWEKHHHICGQTLQAQQQGDTP 537
          D    C  CG  A++ C+ C  ARYC      Q  DW  H   C +          D P
Sbjct: 75  DGPGLCRICGCSAAKKCAKCQVARYCSQAHQVIDWPAHKLECAKAATDGSITDEP 129
```

WU-BLAST vs NCBI BLAST

- WU-BLAST first for gapped alignments
- Use different scoring system for gaps
- Report different statistics
- WU-BLAST does not filter low-complexity by default
- WU-BLAST looks for and reports multiple regions of similarity
- Results will be different

BLAT

- *B*last-*L*ike *A*lignment *T*ool
- Developed by Jim Kent at UCSC
- For DNA it is designed to quickly find sequences of $\geq 95\%$ similarity of length 40 bases or more.
- For proteins it finds sequences of $\geq 80\%$ similarity of length 20 amino acids or more.
- DNA BLAT works by keeping an index of the entire genome in memory - non-overlapping 11-mers (< 1 GB of RAM)
- Protein BLAT uses 4-mers (~ 2 GB)

FASTA

- Index "words" and locate identities
- Rescore best 10 regions
- Find optimal subset of initial regions that can be joined to form single alignment
- Align highest scoring sequences using Smith-Waterman

NCBI Programs for nt vs nt

- Discontiguous megablast
- Megablast
- Nucleotide-nucleotide BLAST (blastn)
- Search for short, nearly exact matches

NCBI Programs for proteins

- Protein-protein BLAST (blastp)
- PHI- and PSI-BLAST
- Search for short, nearly exact matches
- Search the conserved domain database (rpsblast)
- Search by domain architecture (cdart)

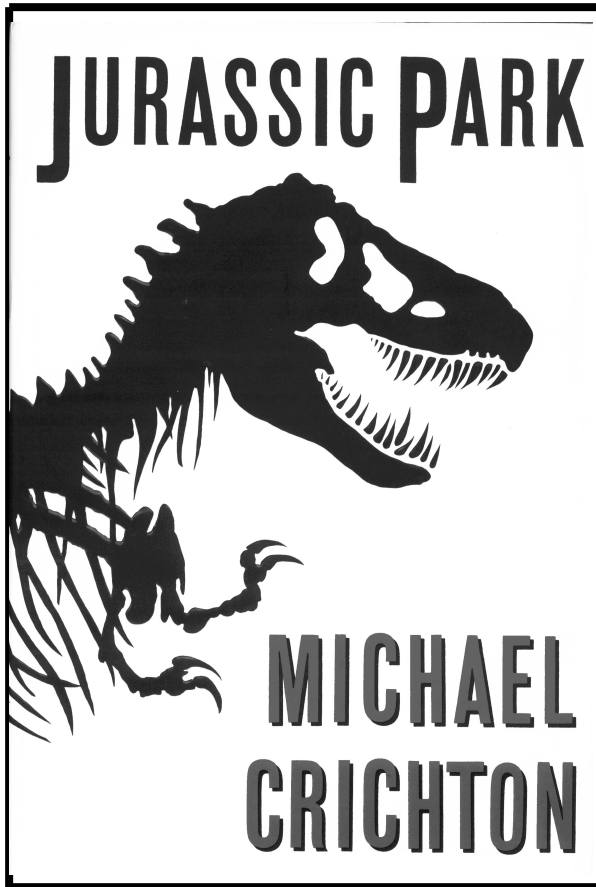
NCBI Programs w/ translations

- Translated query vs. protein database
(blastx)
- Protein query vs. translated database
(tblastn)
- Translated query vs. translated database
(tblastx)

Basic Searching Strategies

- Search early and often
- Use specialized databases
- Use multiple matrices
- Use filters
- Consider Biology

Sequence of Note



```
1 GCGTTGCTGG CGTTTTTCCA TAGGCTCCGC
31 CCCCCTGACG AGCATCACAA AAATCGACGC
61 GGTGGCGAAA CCCGACAGGA CTATAAAGAT
```

.....

```
1371 GTAAAGTCTG GAAACGCGGA AGTCAGCGCC
```

“Here you see the actual structure of a small fragment of dinosaur DNA,” Wu said. “Notice the sequence is made up of four compounds - adenine, guanine, thymine and cytosine. This amount of DNA probably contains instructions to make a single protein - say, a hormone or an enzyme. The full DNA molecule contains *three billion* of these bases. If we looked at a screen like this once a second, for eight hours a day, it’d still take more than two years to look at the entire DNA strand. It’s that big.” (page 103)

DinoDNA "Dinosaur DNA" from Crichton's THE LOST WORLD p. 135

GAATTCCGGAAGCGAGCAAGAGATAAGTCCTGGCATCAGA
TACAGTTGGAGATAAGGACGACGTGTGGCAGCTCCCGCAG
AGGATTCACTGGAAGTGCATTACCTATCCCATGGGAGCCA
TGGAGTTCGTGGCGCTGGGGGGGGCCGGATGCGGGGCTCCCC
CACTCCGTTCCCTGATGAAGCCGGAGCCTTCCTGGGGGCTG
GGGGGGGGGCGAGAGGACGGAGGCGGGGGGGGCTGCTGGCCT
CCTACCCCCCCTCAGGCCGCGTGTCCCTGGTGCCGTGGCA
GACACGGGTACTTTGGGGACCCCCCAGTGGGTGCCGCCCG
CCACCCAAATGGAGCCCCCCCCACTACCTGGAGCTGCTGCA
ACCCCCCGGGGCAGCCCCCCCCCATCCCTCCTCCGGGGCCC
CTACTGCCACTCAGCAGCGCCTGCGGCCTCTACTACAAAC

>Erythroid transcription factor (NF-E1 DNA-binding protein)

```
Query: 121  MEFVALGGPDAGSPTFPDEAGAFGLGGGERTEAGGLLASYPPSGRVSLVPWADTGTLG 300
           MEFVALGGPDAGSPTFPDEAGAFGLGGGERTEAGGLLASYPPSGRVSLVPWADTGTLG
Sbjct: 1    MEFVALGGPDAGSPTFPDEAGAFGLGGGERTEAGGLLASYPPSGRVSLVPWADTGTLG 60

Query: 301  TPQWVPPATQMEPPHYLELLQPPRGSPPHPSSGPLLPLSSGPPPCEARECVMARKNCGAT 480
           TPQWVPPATQMEPPHYLELLQPPRGSPPHPSSGPLLPLSSGPPPCEARECV      NCGAT
Sbjct: 61   TPQWVPPATQMEPPHYLELLQPPRGSPPHPSSGPLLPLSSGPPPCEARECV----NCGAT 116

Query: 481  ATPLWRRDGTGHYLCNWSACGLYHRLNGQNRPLIRPKKRLLVSKRAGTVCSHERENCQT 660
           ATPLWRRDGTGHYLCN      ACGLYHRLNGQNRPLIRPKKRLLVSKRAGTVCS      NCQT
Sbjct: 117  ATPLWRRDGTGHYLCN---ACGLYHRLNGQNRPLIRPKKRLLVSKRAGTVCS---NCQT 169

Query: 661  STTTLWRRSPMGDPVCNNIHACGLYYKLHQVNRPLTMRKDG IQTRNRKVSSKGKKRRPPG 840
           STTTLWRRSPMGDPVCN      ACGLYYKLHQVNRPLTMRKDG IQTRNRKVSSKGKKRRPPG
Sbjct: 170  STTTLWRRSPMGDPVCN---ACGLYYKLHQVNRPLTMRKDG IQTRNRKVSSKGKKRRPPG 226

Query: 841  GGNPSATAGGGAPMGGGGDPSMPPPPPPPPAAAPPQSDALYALGPVVLSGHFLPFGNSGGF 1020
           GGNPSATAGGGAPMGGGGDPSMPPPPPPPPAAAPPQSDALYALGPVVLSGHFLPFGNSGGF
Sbjct: 227  GGNPSATAGGGAPMGGGGDPSMPPPPPPPPAAAPPQSDALYALGPVVLSGHFLPFGNSGGF 286

Query: 1021 FGGGAGGYTAPPGLSPQI 1074
           FGGGAGGYTAPPGLSPQI
Sbjct: 287  FGGGAGGYTAPPGLSPQI 304
```

>Erythroid transcription factor (NF-E1 DNA-binding protein)

```

Query: 121  MEFVALGGPDAGSPTFPDEAGAFGLGGGERTEAGGLLASYPPSGRVSLVPWADTGTLG 300
            MEFVALGGPDAGSPTFPDEAGAFGLGGGERTEAGGLLASYPPSGRVSLVPWADTGTLG
Sbjct: 1     MEFVALGGPDAGSPTFPDEAGAFGLGGGERTEAGGLLASYPPSGRVSLVPWADTGTLG 60

Query: 301  TPQWVPPATQMEPPHYLELLQPPRGSPPHPSSGPLLPLSSGPPPCEARECYMARKNCGAT 480
            TPQWVPPATQMEPPHYLELLQPPRGSPPHPSSGPLLPLSSGPPPCEARECY      NCGAT
Sbjct: 61   TPQWVPPATQMEPPHYLELLQPPRGSPPHPSSGPLLPLSSGPPPCEARECY----NCGAT 116

Query: 481  ATPLWRRDGTGHYLC[WASACGLYHRLNGQNRPLIRPKKRLLVSKRAGTVCS]HERE[NCQT 660
            ATPLWRRDGTGHYLC[WASACGLYHRLNGQNRPLIRPKKRLLVSKRAGTVCS]      NCQT
Sbjct: 117  ATPLWRRDGTGHYLC[----ACGLYHRLNGQNRPLIRPKKRLLVSKRAGTVCS]----NCQT 169

Query: 661  STTTLWRRSPMGDPVCFNIHACGLYYKLHQVNRPLTMRKDG[QTRNRKVSSKGKKRRPPG 840
            STTTLWRRSPMGDPVCFNIHACGLYYKLHQVNRPLTMRKDG[QTRNRKVSSKGKKRRPPG
Sbjct: 170  STTTLWRRSPMGDPVCFNIHACGLYYKLHQVNRPLTMRKDG[QTRNRKVSSKGKKRRPPG 226

Query: 841  GGNPSATAGGGAPMGGGGDPSMPPPPPPPPAAAPPQSDALYALGPVVLSGHFLPFGNSGGF 1020
            GGNPSATAGGGAPMGGGGDPSMPPPPPPPPAAAPPQSDALYALGPVVLSGHFLPFGNSGGF
Sbjct: 227  GGNPSATAGGGAPMGGGGDPSMPPPPPPPPAAAPPQSDALYALGPVVLSGHFLPFGNSGGF 286

Query: 1021 FGGGAGGYTAPPGLSPQI 1074
            FGGGAGGYTAPPGLSPQI
Sbjct: 287  FGGGAGGYTAPPGLSPQI 304

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Useful Web Links

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

<http://www.ebi.ac.uk/blast2/>

<http://www2.ebi.ac.uk/fasta33/>

http://www2.ebi.ac.uk/bic_sw/

<http://genome-test.cse.ucsc.edu/cgi-bin/hgBlat>