Sequence Comparison: Significance of similarity scores

Genome 373

Genomic Informatics

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Review

Global alignment algorithm:

Needleman-Wunsch.

		G	A	A	T	C
	0	-4	-8	-12	-16	-20
C	-4	-5	-9	-13	-12	-6
A	-8	-4	5	1	-3	-7
T	-12	-8	1	0	11	7
A	-16	-12	2	11	7	6
C	-20	-16	-2	7	11	**17**

Local alignment algorithm:

Smith-Waterman.

		A	A	G
	0	0	0	0
G	0	0	0	2
A	0	2	2	0
A	0	2	4	0
G	0	0	0	6
G	0	0	0	2
C	0	0	0	0

Are these proteins related?

The intuitive answer:

```
SEQ 1: RVVNLVPS--FWVLDATYKNYAINYNCDVTYKLY
           L P      L   Y N    Y C     L
SEQ 2: QFFPLMPPAPYFILATDYENLPLVYSCTTFFWLF
```

score = -1 → NO?

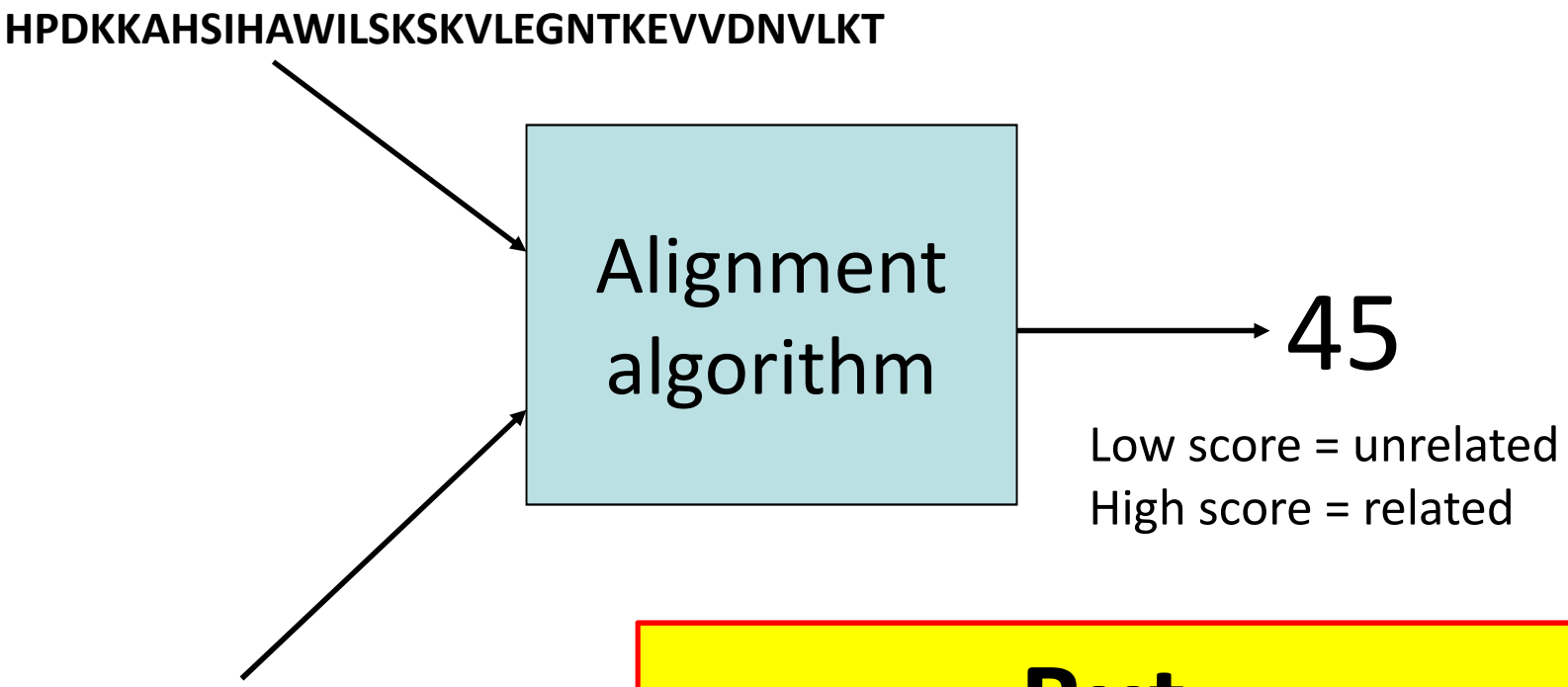
```
SEQ 1: RVVNLVPS--FWVLDATYKNYAINYNCDVTYKLY
           L P    W LDATYKNYA  Y C     L
SEQ 2: QFFPLMPPAPYWILDATYKNYALVYSCTTFFWLF
```

score = 15 → PROBABLY?

```
SEQ 1: RVVNLVPS--FWVLDATYKNYAINYNCDVTYKLY
       RVV L PS   W LDATYKNYA  Y CDVTYKL
SEQ 2: RVVPLMPSAPYWILDATYKNYALVYSCDVTYKLF
```

score = 24 → YES?

Significance of scores



But …
How high is high enough?

The null hypothesis

- We want to know how **surprising** a given score is, …

  assuming that the two sequences are not related.
- This assumption is called the null hypothesis.
- The purpose of most statistical tests is to determine whether the observed result provides a reason to reject the null hypothesis.

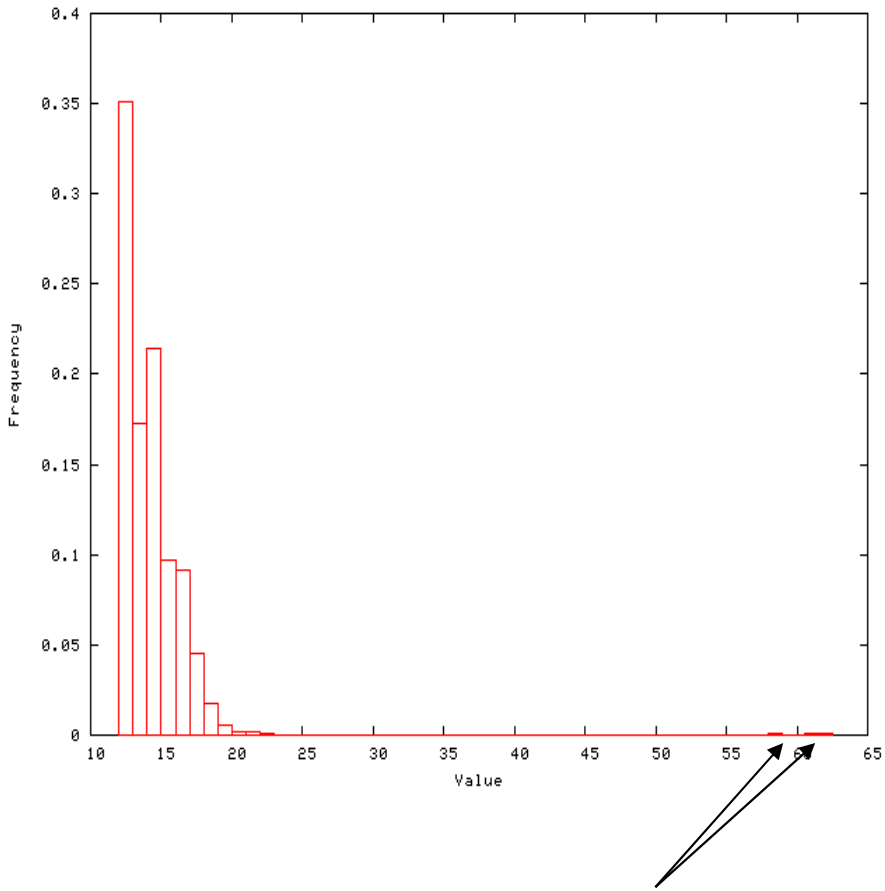
- We want to characterize the distribution of scores from pairwise sequence alignments.

Sequence similarity score distribution

- Search a **randomly generated** database of sequences using a given query sequence.
- What will be the form of the resulting distribution of pairwise alignment scores?

Empirical score distribution

- This shows the distribution of scores from a real database search using BLAST.
- This distribution contains scores from a few related and lots of unrelated pairs.

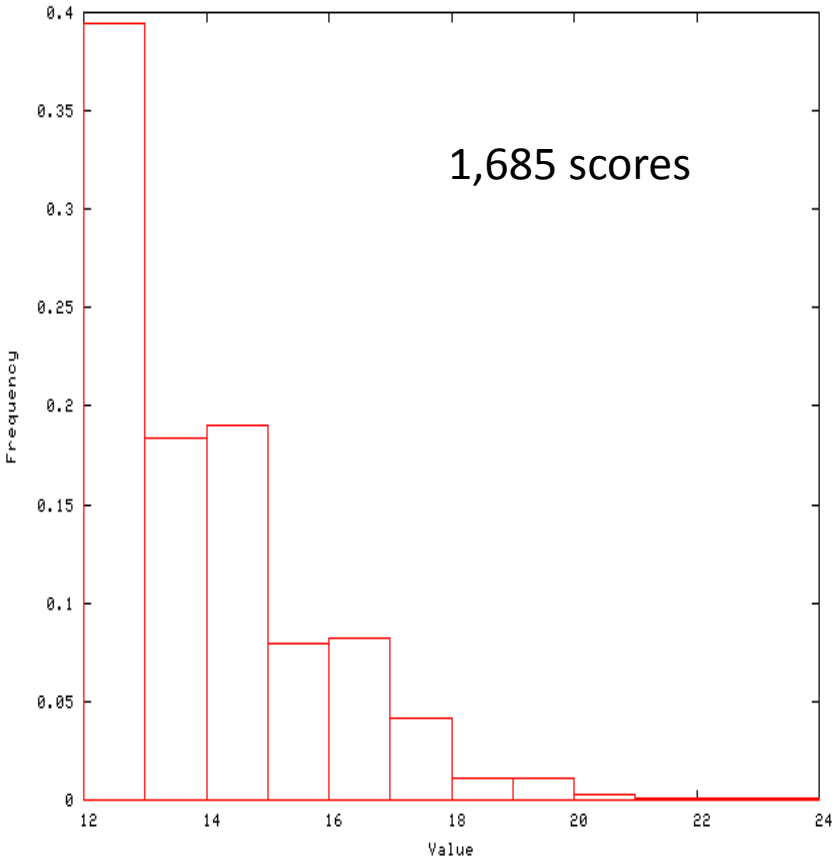


High scores from related sequences

(note - there are lots of lower scoring alignments not reported)

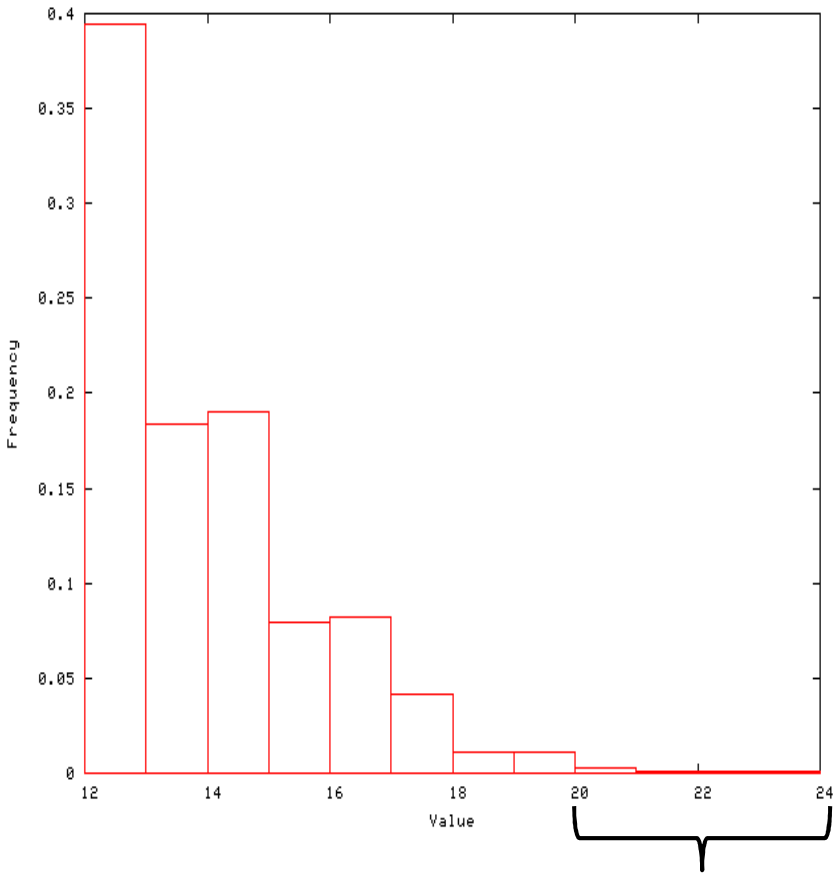
Empirical null score distribution

- The distribution of scores obtained from aligning a given sequence to a database of randomized sequences (e.g., each sequence was shuffled)



(note - there are lots of lower scoring alignments not reported)

Computing an empirical p-value



- The probability of observing a score >=X is the area under the curve to the right of X.
- This probability is called a p-value.
- p-value = Pr(data|null)
  (read as probability of data given a null hypothesis)

e.g. out of 1,685 scores, 28 received a score of 20 or better. Thus, the p-value associated with a score of 20 is ~28/1685 = 0.0166.

Problems with empirical distributions

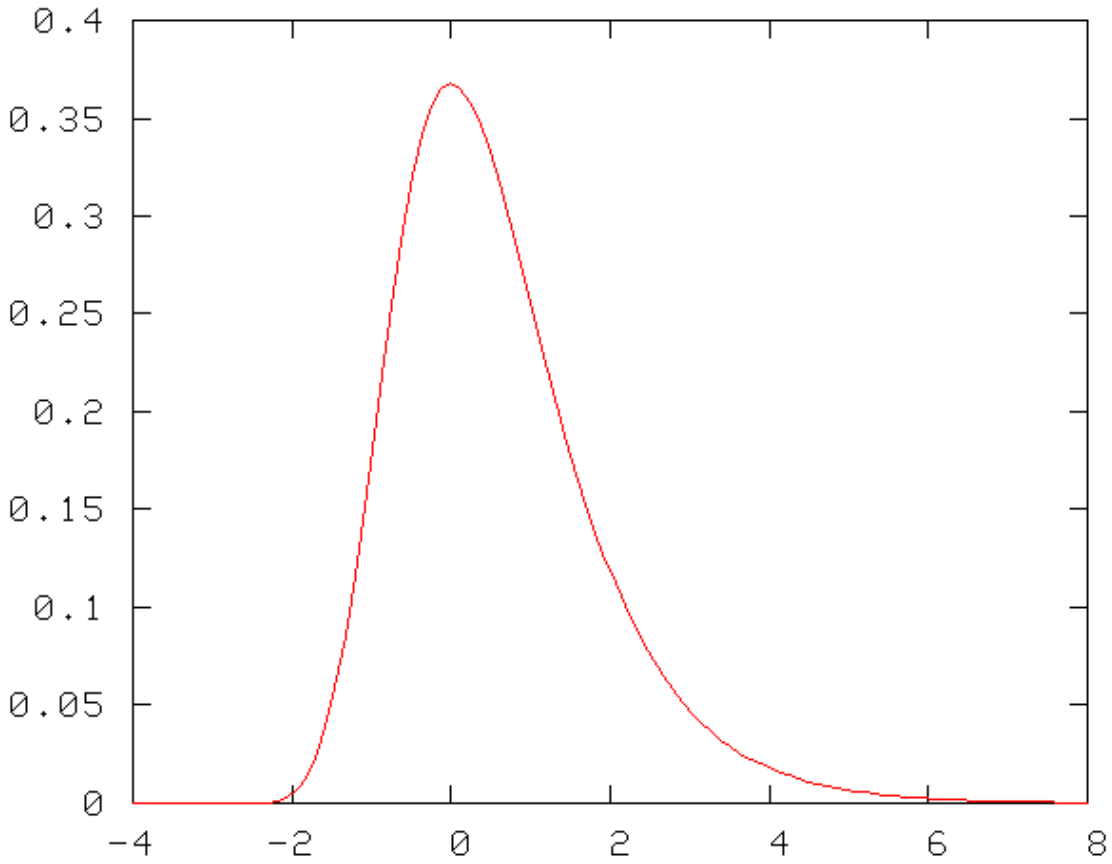
- We are interested in very small probabilities.
- These are computed from the *tail* of the null distribution.
- Estimating a distribution with an accurate tail is feasible but computationally very expensive because we have to make a very large number of alignments.

A solution

- Characterize the form of the score distribution **mathematically**.
- Fit the parameters of the distribution empirically (or compute them analytically).
- Use the resulting distribution to compute accurate p-values.

  (first solved by Karlin and Altschul)

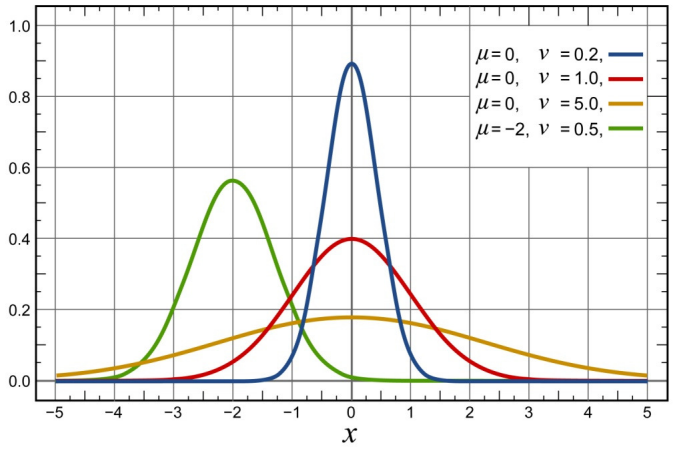
Extreme value distribution



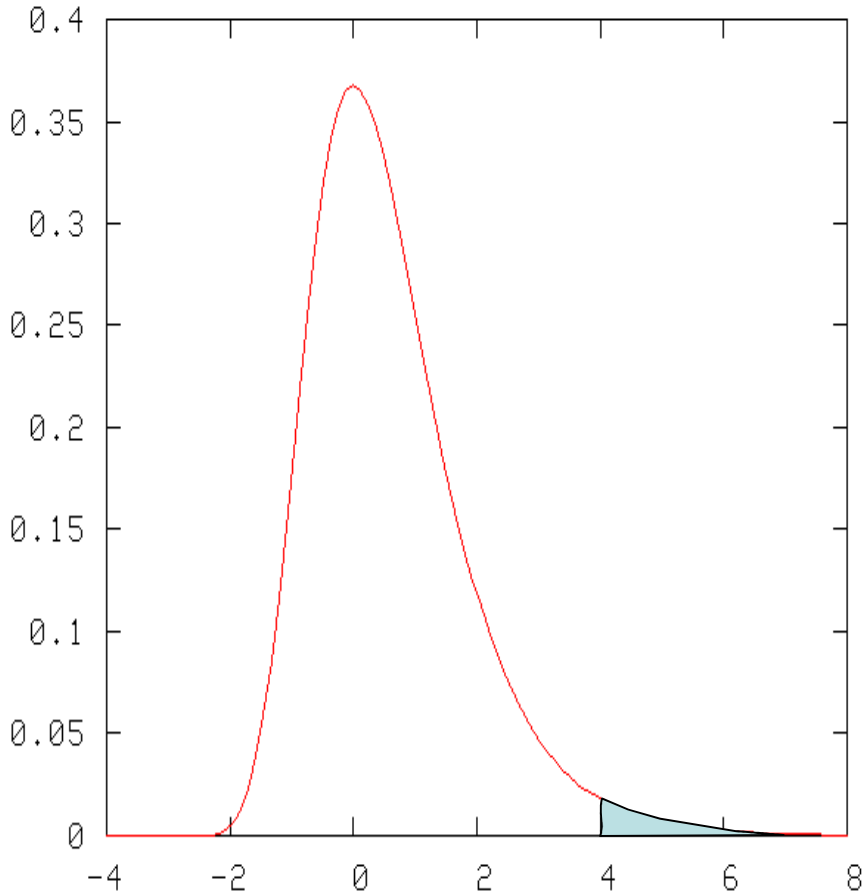
This distribution is roughly normal near the peak, but characterized by a larger tail on the right.

- For an *Unscaled EVD:*

$$P\left(S \geq x\right) = 1 - e^{\left(-e^{-x}\right)}$$



Computing a p-value



- The probability of observing a score >=4 is the area under the curve to the right of 4.
- For an *Unscaled EVD:*

$$P(S \geq x) = 1 - e^{(-e^{-x})}$$

$$P(S \geq 4) = 1 - e^{(-e^{-4})}$$

$$P(S \geq 4) = 0.018149$$

What p-value is significant?

- The most common thresholds are 0.01 and 0.05.
- A threshold of 0.05 means you are 95% sure that the result is significant.
- Is 95% enough? It depends upon the *cost* associated with making a mistake.
- Examples of costs:
  - Doing extensive wet lab validation (expensive)
  - Making clinical treatment decisions (very expensive)
  - Misleading the scientific community (very expensive)
  - Doing further simple computational tests (cheap)
  - Telling your grandmother (very cheap)

Multiple testing

- Say that you perform a statistical test with a 0.05 threshold, but you repeat the test on twenty different observations (e.g. 20 different blast runs)
- Assume that all of the observations are explainable by the null hypothesis.
- What is the chance that at least one of the observations will receive a p-value < 0.05?

$$1 - 0.95^{20} = 0.6415$$

Bonferroni correction

- Assume that individual tests are *independent*.
- Divide the desired p-value threshold by the number of tests performed.

Database searching

- Say that you search the non-redundant protein database at NCBI, containing roughly one million sequences (i.e. you are doing 10^6 pairwise tests).
- and … you want to use a p-value of 0.01.

- Recall that you would observe such a p-value by chance approximately every 100 times in a random database.
- That is, without correcting for multiple testing you will get **~10,000 false positives**!!!
- A Bonferroni correction would suggest using a p-value threshold of $0.01 / 10^6 = 10^{-8}$.

E-values

- A p-value is the probability of making a mistake.
- An E-value is the expected number of times that the given score would appear in a random database of the given size.
- One simple way to compute the E-value is to multiply the p-value times the size of the database.
- Thus, for a p-value of 0.001 and a database of 1,000,000 sequences, the corresponding E-value is 0.001 × 1,000,000 = 1,000.

(BLAST actually calculates E-values in a more complex way, but they mean the same thing)

Sequences producing significant alignments:		Score (bits)	E Value
gi|112670|sp|P15711|104K_THEPA	104 KD MICRONEME-RHOPTRY ANT...	1352	0.0
gi|14268530|gb|AAK56556.1|	104 kDa microneme-rhoptry antige...	243	1e-62
gi|14268528|gb|AAK56555.1|	104 kDa microneme-rhoptry antige...	242	4e-62
gi|14268526|gb|AAK56554.1|	104 kDa microneme-rhoptry antige...	238	7e-61
gi|31210185|ref|XP_314059.1|	ENSANGP00000015608 [Anopheles ...	37	2.1
gi|22971724|ref|ZP_00018655.1|	hypothetical protein [Chloro...	35	9.7
gi|32403566|ref|XP_322396.1|	hypothetical protein [Neurospo...	35	12
gi|24639766|ref|NP_572189.1|	CG2861-PA [Drosophila melanoga...	34	17
gi|30348569|emb|CAC84361.1|	hypothetical protein [Saimiriin...	34	19
gi|6492132|gb|AAF14193.1|	spherical body protein 3 [Babesia...	34	20
gi|9629342|ref|NP_044542.1|	virion protein [Human herpesvir...	34	21
gi|24639768|ref|NP_726958.1|	CG2861-PB [Drosophila melanoga...	34	21
gi|4757118|emb|CAB42096.1|	TashAT2 protein [Theileria annul...	34	22
gi|17534529|ref|NP_495288.1|	putative protein (2G676) [Caen...	34	22
gi|15241089|ref|NP_195809.1|	leucine-rich repeat transmembr...	33	23
gi|43489677|gb|EAD99646.1|	unknown [environmental sequence]	33	23
gi|44419062|gb|EAJ13596.1|	unknown [environmental sequence]	33	25
gi|43969222|gb|EAG41329.1|	unknown [environmental sequence]	33	29
gi|15792145|ref|NP_281968.1|	putative oxidoreductase [Campy...	33	34
gi|43926327|gb|EAG18073.1|	unknown [environmental sequence]	33	37
gi|39595869|emb|CAE67372.1|	Hypothetical protein CBG12848 [...	33	38
gi|30020082|ref|NP_831713.1|	Glycosyltransferase [Bacillus ...	33	40
gi|43723946|gb|EAF16931.1|	unknown [environmental sequence]	33	41
gi|11545212|gb|AAG37800.1|	hypothetical telomeric SfiI frag...	33	44
gi|40788024|emb|CAE47751.1|	ubiquitin specific proteinase 5...	32	51
gi|42656951|ref|XP_052597.6|	ubiquitin specific protease 53...	32	51
gi|32698642|ref|NP_872557.1|	DNA-ligase [Adoxophyes orana g...	32	52
gi|12840300|dbj|BAB24814.1|	unnamed protein product [Mus mu...	32	54
gi|28899333|ref|NP_798938.1|	4-diphosphocytidyl-2C-methyl-D...	32	55
gi|7243081|dbj|BAA92588.1|	KIAA1350 protein [Homo sapiens]	32	62

Summary

- A distribution plots the frequencies of types of observation.
- The area under the distribution curve is 1.
- Most statistical tests compare observed data to the expected result according to a null hypothesis.
- Sequence similarity scores follow an extreme value distribution, which is characterized by a long tail.
- The p-value associated with a score is the area under the curve to the right of that score.
- Selecting a significance threshold requires evaluating the cost of making a mistake.
- Bonferroni correction: Divide the desired p-value threshold by the number of statistical tests performed.
- The E-value is the expected number of times that a given score would appear in a random database of the given size.