

TECHNICAL NOTE

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New finite-size correction for local alignment score distributions

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Abstract

Background: Local alignment programs often calculate the probability that a match occurred by chance. The calculation of this probability may require a “finite-size” correction to the lengths of the sequences, as an alignment that starts near the end of either sequence may run out of sequence before achieving a significant score.

Findings: We present an improved finite-size correction that considers the distribution of sequence lengths rather than simply the corresponding means. This approach improves sensitivity and avoids substituting an *ad hoc* length for short sequences that can underestimate the significance of a match. We use a test set derived from ASTRAL to show improved ROC scores, especially for shorter sequences.

Conclusions: The new finite-size correction improves the calculation of probabilities for a local alignment. It is now used in the BLAST+ package and at the NCBI BLAST web site (<http://blast.ncbi.nlm.nih.gov>).

Background

Local alignments are an essential tool for biologists and often provide the first information about the function of an unknown nucleotide or protein sequence. An important question concerns the relationship of the score of a local alignment with the probability that the alignment occurred by chance. Karlin and Altschul [1] developed an asymptotic theory for local alignments, assuming that no gaps are permitted. For two random sequences **I** and **J** of lengths m and n , respectively, the resulting distribution of the optimal alignment score \hat{M} approximates a Gumbel distribution [2]

$$\mathbb{P}\{\hat{M} > y\} \approx 1 - \exp(-kmne^{-\lambda y}). \quad (1)$$

The two statistical parameters in Equation (1) are λ , the scale parameter, and k , the pre-factor.

Several authors [3-12] extended this framework to local alignments with gaps and showed that the Gumbel distribution from Equation (1) is still valid, though different values for λ and k are required. Altschul [13] discussed the need for a “finite-size correction” to the lengths m and n to improve the accuracy of Equation

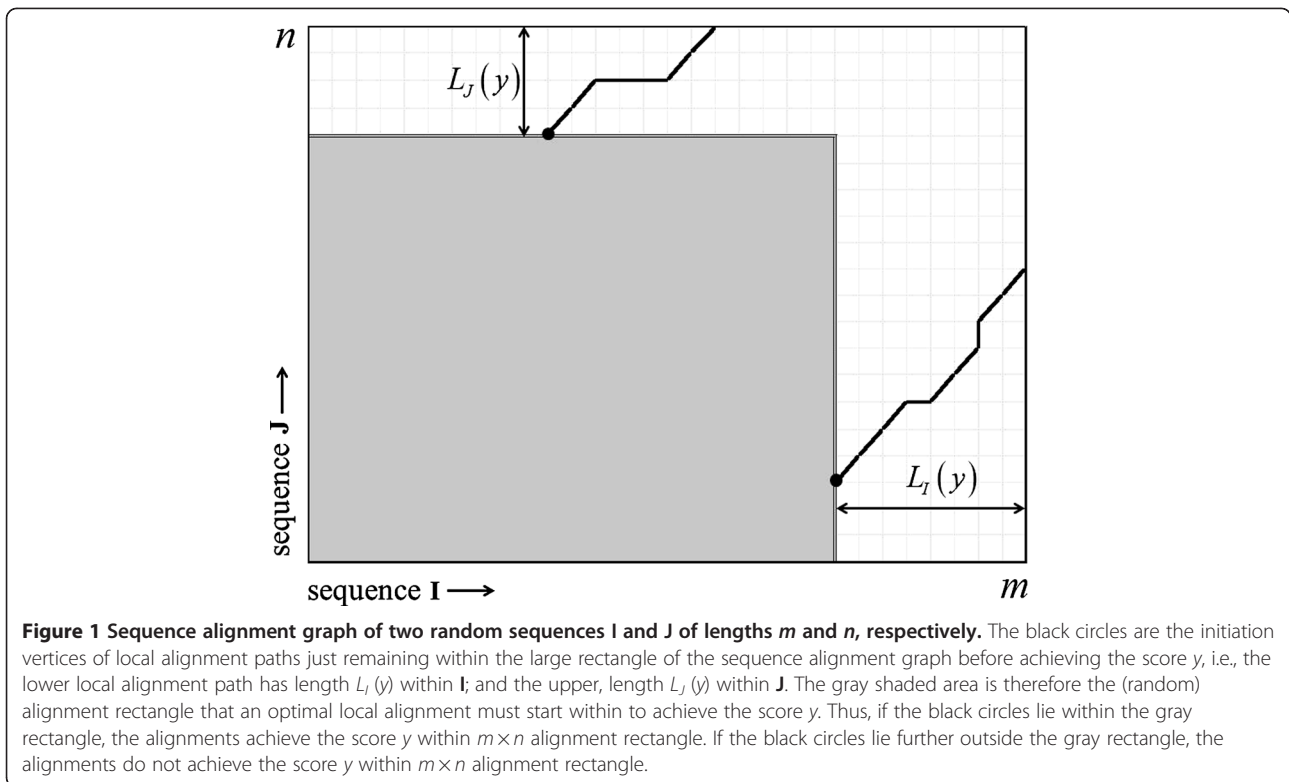
(1). The resulting statistics are an integral part of the Basic Local Alignment Search Tool (BLAST) [14].

The following presentation emphasizes intuition over mathematical formality, to explain how the finite-size correction can account for the finite sequence lengths m and n to improve the accuracy of Equation (1). Let us begin with an optimal local alignment, which starts from score 0 and requires a non-zero sequence length within both **I** and **J**, before it achieves score y . Let $L_I(y)$ ($L_J(y)$) be the required random lengths within both **I** (**J**), and let $l_I(y) = \mathbb{E}\{L_I(y)\}$ ($l_J(y) = \mathbb{E}\{L_J(y)\}$) be the corresponding means. The main idea is that the optimal local alignment cannot start anywhere along the full length m (n) of sequence **I** (**J**), because there might be insufficient sequence to permit it to achieve the score y (Figure 1). The finite-size correction described in [13] and used in BLAST therefore replaced the area mn of the alignment matrix for Equation (1) by

$$[m - l_I(y)][n - l_J(y)]. \quad (2)$$

Equation (2) approximates the area within the alignment matrix where the optimal local alignment can start and on average still have enough space to exceed the score y . If $m < l_I(y)$ or $n < l_J(y)$, however, the resulting value in Equation (2) might become negative. The BLAST code for the old finite-size correction therefore set the corrected sequence length to an *ad hoc* value

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(typically 1). For very short query or database sequences, the *ad hoc* correction could underestimate the significance of an alignment by many orders of magnitude.

The purpose of this note is to present a new finite-size correction formula for the BLAST statistics. It avoids the *ad hoc* correction and improves on them by considering the (approximately normal) distributions of the random lengths $L_I(y)$ and $L_J(y)$ explicitly, and not just the corresponding means $l_I(y)$ and $l_J(y)$. We demonstrate below that the new finite-size correction is better than the older one, both in theory and in practice. All BLAST+ protein-protein applications (i.e., BLASTP, BLASTX) use the new finite-size correction by default, starting with version 2.2.26.

Findings

New finite-size correction

As with the old finite-size correction, the expectation $l_I(y) = \mathbb{E}L_I(y)$ is approximated linearly:

$$l_I(y) = a_I y + b_I. \quad (3)$$

Most practical scoring systems are symmetric, with $s(A, B) = s(B, A)$ for any two letters A and B , and for a symmetric scoring matrix and symmetric sequence compositions, expectations corresponding to I and J are the same, e.g., $l_I(y) = l_J(y) = l(y)$. For asymmetric scoring systems or asymmetric sequence compositions, however,

the variates $L_I(y)$ and $L_J(y)$ can have different distributions, so the following retains the subscripts I and J.

The new finite-size correction replaces mn in Equation (1) by

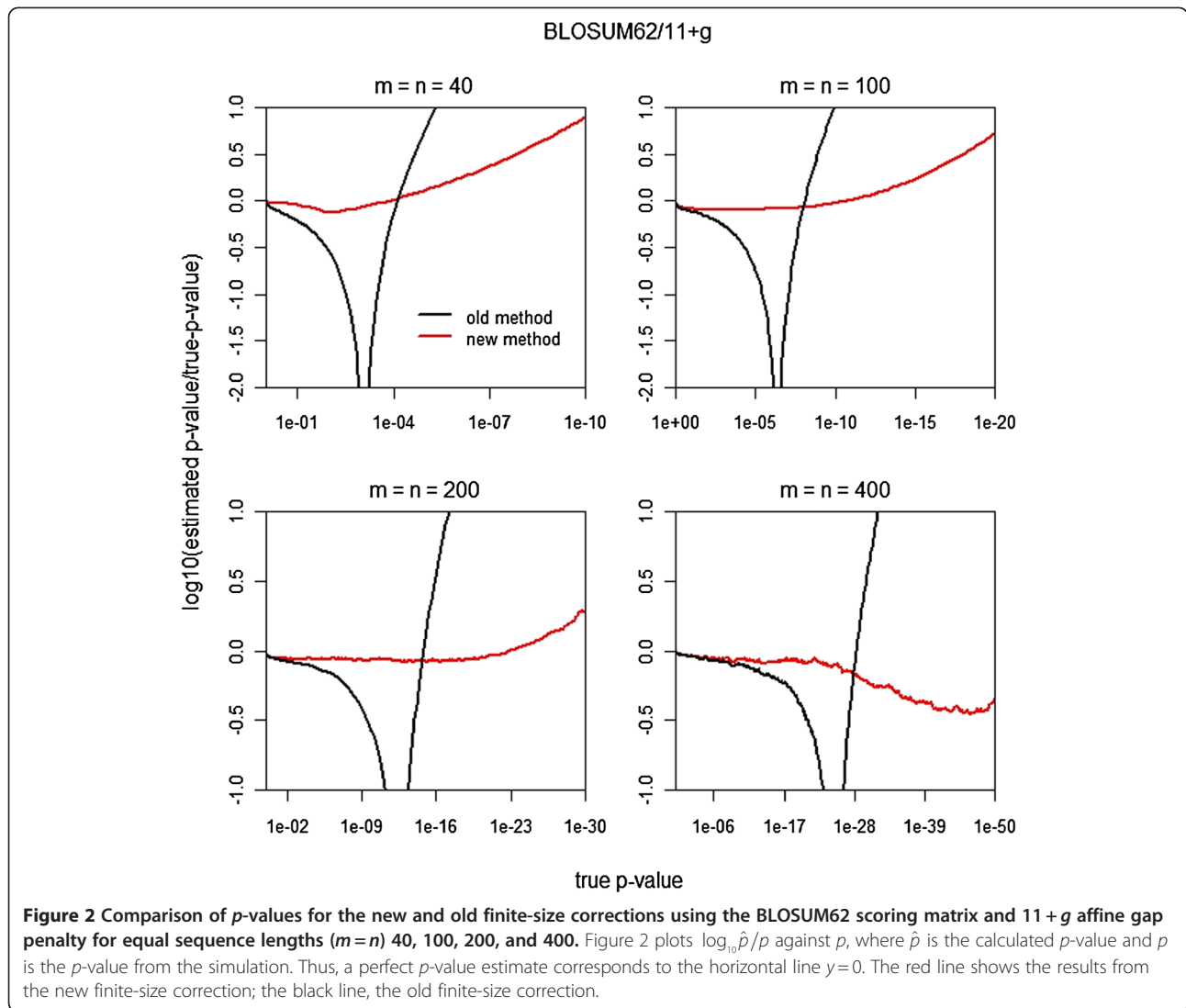
$$\mathbb{E}\{[m - L_I(y)]^+ [n - L_J(y)]^+\}, \quad (4)$$

where $x^+ = \max\{x, 0\}$. Rather than taking the expectation of $L_I(y)$ and $L_J(y)$ as in Equation (2), Equation (4) is the expected area within the alignment rectangle where an optimal local alignment can start and have enough random sequence length to reach the score y (Figure 1).

The practical computation of Equation (4) approximates the distribution of $(L_I(y), L_J(y))$ with a bivariate normal distribution, with means $l_I(y) = \mathbb{E}L_I(y)$ and $l_J(y) = \mathbb{E}L_J(y)$, variances $\text{var } L_I(y) = v_I(y)$ and $\text{var } L_J(y) = v_J(y)$, and covariance $\text{cov}(L_I(y), L_J(y)) = c(y)$, all assumed to be linear in the score y , i.e.,

$$\begin{aligned} l_I(y) &= a_I y + b_I, l_J(y) = a_J y + b_J, \\ v_I(y) &= \alpha_I y + \beta_I, v_J(y) = \alpha_J y + \beta_J, \\ c(y) &= \sigma y + \tau. \end{aligned} \quad (5)$$

The estimation of the parameters $a_I, a_J, \alpha_I, \alpha_J$ and σ has mathematical depth and involves many unproved speculations, but involves a heuristic modeling of a random sequence alignment with Markov additive processes [15], ultimately with use of the renewal-reward theorem.



The Appendix presents formulas for computing a_I , a_J , α_I , α_J and σ .

BLAST p -values are relatively insensitive to the values of the intercepts b_I , b_J , β_I , β_J , and τ , so the practical computation approximates them, as follows. Let a_u (α_u) be

the value of a_I (α_I) for ungapped alignment. The mathematical theories for random walks and for renewals yield analytic formulas for a_u and α_u [16]. For an ungapped optimal alignment, the alignment length required to exceed the score y is the same within the sequences I and J, because it lacks gaps. Thus, a_u and α_u do not depend on the sequence (I or J) under consideration, so they contain no subscripts I or J. In a gapped alignment, let a gap of length 1 incur a penalty G . The following uncontrolled approximations hold [17]:

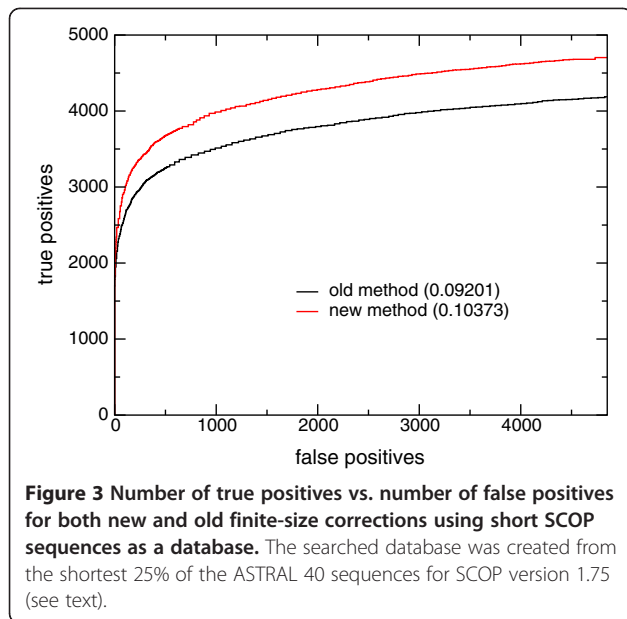
$$\begin{aligned} b_I &= 2G(a_u - a_I), b_J = 2G(a_u - a_J) \\ \beta_I &= 2G(\alpha_u - \alpha_I), \beta_J = 2G(\alpha_u - \alpha_J) \\ \tau &= 2G(\alpha_u - \sigma). \end{aligned} \tag{6}$$

Under the normal approximation, routine computation shows that Equation (4) is approximately

Table 1 Retrieval accuracy for different subsets of SCOP database with the new and old finite-size correction

Method	25 th percentile	50 th percentile	Full database
New correction	0.10373 ± 0.00022	0.10073 ± 0.00019	0.08535 ± 0.00013
Old correction	0.09201 ± 0.00020	0.09282 ± 0.00017	0.08358 ± 0.00014

The three subsets contain proteins shorter than 91 residues (25th percentile by length), shorter than 137 residues (50th percentile by length), and the full database. ROC-4852 scores are presented with an error (one standard deviation). The 25th percentile database contains 2533 sequences, the 50th percentile database contains 5008 sequences, and the full database contains 10,569 sequences. There are 4852 queries.



$$\mathbb{E}([m - l_I(y)]^+ [n - l_J(y)]^+) \approx \frac{[(m - l_I(y))\mathbb{P}(X \leq m_\phi) - \sqrt{v_I(y)}\mathbb{E}(X; X \leq m_\phi)] \times [(n - l_J(y))\mathbb{P}(X \leq n_\phi) - \sqrt{v_J(y)}\mathbb{E}(X; X \leq n_\phi)] + c(y)\mathbb{P}(X \leq m_\phi)\mathbb{P}(X \leq n_\phi)}{c(y)}$$
(7)

where $m_\phi := [m - l_I(y)] / \sqrt{v_I(y)}$, $n_\phi := [n - l_J(y)] / \sqrt{v_J(y)}$, and X is a standard normal variate. The final product $\mathbb{P}(X \leq m_\phi)\mathbb{P}(X \leq n_\phi)$ is an uncontrolled independence approximation for the bivariate normal distribution.

Comparison of p -values for the new and old finite-size corrections

We compared p -values for the new finite-size correction with those for the old finite-size correction using the BLOSUM62 scoring matrix and affine gap penalty $11 + g$. Hartmann used a rare-event simulation method to compute the local alignment score distribution for ranges that included small p -values like $p = 10^{-50}$ [18], thereby producing a theoretical standard for small p -values.

Figure 2 plots relative errors in logarithmic scale against true p -values for equal sequence lengths $m = n = 40, 100, 200,$ and 400 . Using Hartmann's theoretical standard, the new finite-size correction outperforms as the p -value decreases, sometimes by orders of magnitude.

Evaluation of accuracy

We evaluated the performance of the new finite-size correction using the ASTRAL SCOP 40 subset [19] of release 1.75 of the Structural Classification of Proteins

(SCOP) [20] database. We sorted the SCOP domains by lexicographic order and used the even numbered sequences as our query set, but removed any query that was the sole member of the superfamily in ASTRAL 40. For a given query sequence, we considered any database sequence belonging to the same SCOP superfamily as a true positive, and any database sequence belonging to a different SCOP fold as a false positive. Following [21], in the retrieval list for each query, we censored all sequences belonging to the same fold but different superfamily, so those sequences contributed neither true or false positives to the retrieval.

We report the performance in terms of the Receiver Operator Characteristics (ROC). Specifically, we report the ROC_n score, which is obtained by pooling the results of all queries, ordering them by expect value, but only keeping results up the n -th false positive [21]. The expect value for the database search was obtained from the pairwise p -values using a length-proportional correction that takes the ratio of the database length to the target sequence length into account [13].

As discussed above, the new finite-size correction should show the greatest improvement for short sequences. Therefore, we also produced ROC_n scores for different subsets of the SCOP database. One database subset has sequences shorter than the 25th percentile length (95 residues), and another has sequences shorter than the 50th percentile length (137 residues).

Table 1 presents ROC_n scores for the full database as well as the two subsets described above. These scores have an average of one false positive per query (4852), a threshold found useful in other studies (Altschul SE, private communication). The ROC -4852 scores for the full database demonstrate a small improvement of the new finite-size correction over the older one. The subsets show a more impressive improvement. For the 50th percentile subset, the ROC -4852 score improves by 9%. For the 25th percentile subset, the ROC -4852 score shows a 13% improvement. In the 25th percentile subset, the new finite-size correction produces roughly 12% more true positives overall at 4852 false positives than the old finite-size correction (Figure 3). These results confirm our expectation that the new finite-size correction will display greatest improvement in retrieval for short sequences.

To assess the significance of this improvement on BLAST searches, one may look to the length distribution of sequences in a heavily used protein BLAST database. The non-redundant ("nr") database is the default protein database at the NCBI BLAST web site. Of the sequences in the nr database, 11% are 95 residues or shorter; and 21%, 137 residues or shorter. The new finite-size correction improves the retrieval accuracy for a noticeable fraction of the proteins in the nr database.

Conclusion

We have described a new finite-size correction. The new correction has a more rigorous derivation than the current finite-size correction and avoids the use of an *ad hoc* value for short sequences. We have tested the retrieval accuracy of the new finite-size correction on the gold standard SCOP set, and have shown that the improvement is most important for short sequences. This correction has been made part of the BLAST+ protein-protein applications (e.g., BLASTP, BLASTX) as well as at the NCBI BLAST web site. In the future, we plan to implement this correction for nucleotide-nucleotide comparisons.

Availability and requirements

Project Name: BLAST Statistical Parameters

Project home page: http://www.ncbi.nlm.nih.gov/CBBresearch/Spouge/html_ncbi/html/blast/

Operating systems: Windows, MacOSX, LINUX, UNIX
 Programming language: C++

License: Public Domain (see <http://www.ncbi.nlm.nih.gov/books/NBK22952/>)

Any restrictions to use by non-academics: None

Appendix

Let $\mathbb{Z}_+ = \{0, 1, \dots\}$. Consider two semi-infinite random sequences, yielding alignment scores $S_{i,j}$ at each vertex $(i, j) \in \mathbb{Z}_+^2$ within their alignment graph. Define the edge maximum score $E_n = \max\{\max_{0 \leq i \leq n} S_{i,n}, \max_{0 \leq j \leq n} S_{n,j}\}$. Let $\kappa_0 = E_{\kappa_0} = 0$ and $\kappa_i = \inf\{n : n > \kappa_{i-1}, E_n > E_{\kappa_{i-1}}\}$ for $i \geq 1$. We call κ_i the i^{th} SALE (strict ascending ladder epoch) and E_{κ_i} the i^{th} SALE score. Let $\Delta E_i := E_{\kappa_i} - E_{\kappa_{i-1}}$, the increment between the $(i-1)^{\text{th}}$ and i^{th} SALE scores.

Let $L_I(y) = \inf\{i : S_{i,j} \geq y, (i, j) \in \mathbb{Z}_+^2\}$ and $L_J(y) = \inf\{j : S_{i,j} \geq y, (i, j) \in \mathbb{Z}_+^2\}$. We also define $I_n = \inf\{i : S_{i,j} = E_{\kappa_n}, (i, j) \in \mathbb{Z}_+^2\}$ and $J_n = \inf\{j : S_{i,j} = E_{\kappa_n}, (i, j) \in \mathbb{Z}_+^2\}$. Let $\Delta I_i := I_i - I_{i-1}$, the incremental sequence length between $(i-1)^{\text{th}}$ and i^{th} SALEs in sequence **I**, and $\Delta J_j := J_j - J_{j-1}$, the incremental sequence length between $(j-1)^{\text{th}}$ and j^{th} SALEs in sequence **J**. Last, we define $\mathbb{E}^*[\Delta I_i] := \mathbb{E}[\Delta I_i e^{\lambda E_{\kappa_i}}, \kappa_i < \infty]$, $\mathbb{E}^*[\Delta J_j] := \mathbb{E}[\Delta J_j e^{\lambda E_{\kappa_j}}, \kappa_j < \infty]$, and $\mathbb{E}^*[\Delta E_i] := \mathbb{E}[\Delta E_i e^{\lambda E_{\kappa_i}}, \kappa_i < \infty]$.

The formulas for computing $a_I, a_J, \alpha_I, \alpha_J$ and σ are:

$$a_I = \lim_{i \rightarrow \infty} \frac{\mathbb{E}^*[\Delta I_i]}{\mathbb{E}^*[\Delta E_i]}, a_J = \lim_{j \rightarrow \infty} \frac{\mathbb{E}^*[\Delta J_j]}{\mathbb{E}^*[\Delta E_j]},$$

$$\alpha_I = \lim_{i \rightarrow \infty} \frac{\text{var}^*[\Delta I_i]}{\mathbb{E}^*[\Delta E_i]}, \alpha_J = \lim_{j \rightarrow \infty} \frac{\text{var}^*[\Delta J_j]}{\mathbb{E}^*[\Delta E_j]},$$

$$\sigma = \lim_{i \rightarrow \infty} \frac{\text{cov}^*[\Delta I_i, \Delta J_i]}{\mathbb{E}^*[\Delta E_i]},$$

where var^* and cov^* represent the variance and covariance associated with the probability measure underlying

the expectation \mathbb{E}^* . In practice, for computational efficiency, we use importance sampling to estimate the parameters above [15]. The parameters are estimated separately at each SALE. We then apply asymptotic regression to estimate the values of $a_I, a_J, \alpha_I, \alpha_J$ and σ as $i \rightarrow \infty$ in the equation above [22].

Competing interests

The authors declare that they have no competing interests.

Authors' contributions

YP, TM and JS drafted the manuscript. YP designed the p -value evaluation method. SS implemented the new finite-size correction. NM integrated the correction into the BLAST+ code, ran tests, and calculated the ROC scores. JS devised the new finite-size correction. YP and SS are equal contribution first authors for this article. TLM and JLS are equal contribution last authors for this article. All authors read and approved the final manuscript.

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References

- Karlin S, Altschul SF: **Methods for assessing the statistical significance of molecular sequence features by using general scoring schemes.** *Proc Natl Acad Sci U S A* 1990, **87**(6):2264–2268.
- Galambos J: *The asymptotic theory of extreme order statistics.* New York: Wiley; 1978.
- Mott R: **Maximum-likelihood-estimation of the statistical distribution of Smith-Waterman local sequence similarity scores.** *Bull Math Biol* 1992, **54**(1):59–75.
- Waterman MS, Vingron M: **Rapid and accurate estimates of statistical significance for sequence data base searches.** *Proc Natl Acad Sci U S A* 1994, **91**(11):4625–4628.
- Altschul SF, Gish W: **Local alignment statistics.** *Methods Enzymol* 1996, **266**:460–480.
- Bundschuh R: **Rapid significance estimation in local sequence alignment with gaps.** *J Comput Biol* 2002, **9**(2):243–260.
- Chia N, Bundschuh R: **A practical approach to significance assessment in alignment with gaps.** *J Comput Biol* 2006, **13**(2):429–441.
- Newberg LA: **Significance of gapped sequence alignments.** *J Comput Biol* 2008, **15**(9):1187–1194.
- Agrawal A, Brendel VP, Huang X: **Pairwise statistical significance and empirical determination of effective gap opening penalties for protein local sequence alignment.** *Int J Computat Biol Drug Des* 2008, **1**(4):347–367.
- Poleksic A: **Island method for estimating the statistical significance of profile-profile alignment scores.** *BMC Bioinformatics* 2009, **10**:112.
- Ortet P, Bastien O: **Where does the alignment score distribution shape come from?** *Evol Bioinformatics* 2010, **6**:159–187.
- Agrawal A, Huang X: **Pairwise statistical significance of local sequence alignment using sequence-specific and position-specific substitution matrices.** *IEEE/ACM Trans Comput Biol Bioinformatics* 2011, **8**(1):194–205.
- Altschul SF: **Evaluating the statistical significance of multiple distinct local alignments.** In *Theoretical and computational methods in genome research.* Edited by Suhai S. New York: Plenum Press; 1997:1–14.
- Altschul SF, Madden TL, Schaffer AA, Zhang J, Zhang Z, Miller W, Lipman DJ: **Gapped BLAST and PSI-BLAST: a new generation of protein database search programs.** *Nucleic Acids Res* 1997, **25**(17):3389–3402.
- Park Y, Sheetlin S, Spouge JL: **Estimating the gumbel scale parameter for local alignment of random sequences by importance sampling with stopping times.** *Ann Stat* 2009, **37**(6A):3697–3714.
- Asmussen S: *Applied probability and queues.* New York: Springer; 2003.
- Altschul SF, Bundschuh R, Olsen R, Hwa T: **The estimation of statistical parameters for local alignment score distributions.** *Nucleic Acids Res* 2001, **29**(2):351–361.

18. Hartmann AK: **Sampling rare events: statistics of local sequence alignments.** *Phys Rev E* 2002, **65**(5). doi:10.1103/PhysRevE.65.056102.
19. Chandonia JM, Hon G, Walker NS, Lo Conte L, Koehl P, Levitt M, Brenner SE: **The ASTRAL Compendium in 2004.** *Nucleic Acids Res* 2004, **32**:D189–D192.
20. Murzin AG, Brenner SE, Hubbard T, Chothia C: **SCOP - a structural classification of proteins database for the investigation of sequences and structures.** *J Mol Biol* 1995, **247**(4):536–540.
21. Schaffer AA, Aravind L, Madden TL, Shavirin S, Spouge JL, Wolf YI, Koonin EV, Altschul SF: **Improving the accuracy of PSI-BLAST protein database searches with composition-based statistics and other refinements.** *Nucleic Acids Res* 2001, **29**(14):2994–3005.
22. Sheetlin S, Park Y, Spouge JL: **Objective method for estimating asymptotic parameters, with an application to sequence alignment.** *Phys Rev E* 2011, **84**(3). doi:10.1103/PhysRevE.84.031914.

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