

## PSI-Search: iterative HOE-reduced profile SSEARCH searching

Weizhong Li<sup>1</sup>, Hamish McWilliam<sup>1</sup>, Mickael Goujon<sup>1</sup>, Andrew Cowley<sup>1</sup>, Rodrigo Lopez<sup>1,\*</sup> and William R. Pearson<sup>2,\*</sup>

<sup>1</sup>EMBL – European Bioinformatics Institute, Wellcome Trust Genome Campus, Hinxton, Cambridge CB10 1SD, UK and <sup>2</sup>Department of Biochemistry and Molecular Genetics, Charlottesville, VA 22908, USA

Associate Editor: Alfonso Valencia

### ABSTRACT

**Summary:** Iterative similarity searches with PSI-BLAST position-specific score matrices (PSSMs) find many more homologs than single searches, but PSSMs can be contaminated when homologous alignments are extended into unrelated protein domains—homologous over-extension (HOE). PSI-Search combines an optimal Smith–Waterman local alignment sequence search, using SSEARCH, with the PSI-BLAST profile construction strategy. An optional sequence boundary-masking procedure, which prevents alignments from being extended after they are initially included, can reduce HOE errors in the PSSM profile. Preventing HOE improves selectivity for both PSI-BLAST and PSI-Search, but PSI-Search has ~4-fold better selectivity than PSI-BLAST and similar sensitivity at 50% and 60% family coverage. PSI-Search is also produces 2- for 4-fold fewer false-positives than JackHMMER, but is ~5% less sensitive.

**Availability and implementation:** PSI-Search is available from the authors as a standalone implementation written in Perl for Linux-compatible platforms. It is also available through a web interface ([www.ebi.ac.uk/Tools/sss/psisearch](http://www.ebi.ac.uk/Tools/sss/psisearch)) and SOAP and REST Web Services ([www.ebi.ac.uk/Tools/webservices](http://www.ebi.ac.uk/Tools/webservices)).

**Contact:** [pearson@virginia.edu](mailto:pearson@virginia.edu); [rodrigo.lopez@ebi.ac.uk](mailto:rodrigo.lopez@ebi.ac.uk)

Received on March 29, 2012; revised on March 29, 2012; accepted on April 17, 2012

### 1 INTRODUCTION

PSI-BLAST (Altschul *et al.*, 1997) uses an iterative strategy to construct a protein profile, in the form of a position-specific score matrix (PSSM), which dramatically improves homology detection in diverse protein families. Improved versions of PSI-BLAST have more accurate statistics and more sensitive consensus profiles (Agrawal *et al.*, 2009; Altschul *et al.*, 2005, 2009; Bhadra *et al.*, 2006; Li *et al.*, 2011; Przybylski and Rost, 2008; Stojmirović *et al.*, 2008), but the most common cause of PSI-BLAST errors is contamination of the PSSM by extension of an homologous domain into a non-homologous region (homologous over-extension, HOE) (Gonzalez and Pearson, 2010a). Even searches with a single well-defined domain do not guarantee uncontaminated profiles (Kim *et al.*, 2010). Some HOE errors can be reduced by ‘profile cleaning’; HangOut (Kim *et al.*, 2010) focuses on long insertions, but requires insertion boundaries to be specified by the user, thus assuming *a priori* knowledge of the domain structure of the query protein.

Here we present PSI-Search, an iterated profile search application for identifying distantly related protein sequences. PSI-Search is similar to PSI-BLAST, but substitutes a rigorous Smith–Waterman local alignment (Smith and Waterman, 1981) search strategy (SSEARCH, Pearson, 1991) to produce optimal local alignment scores from the profile PSSM. PSI-Search includes an optional alignment boundary-masking procedure that reduces HOE errors in the PSSM profile. SCANPS (Walsh *et al.*, 2008) implements a similar iterative search strategy using Smith–Waterman alignments; however, it does not currently scale to large protein databases and does not include boundary masking.

### 2 METHODS

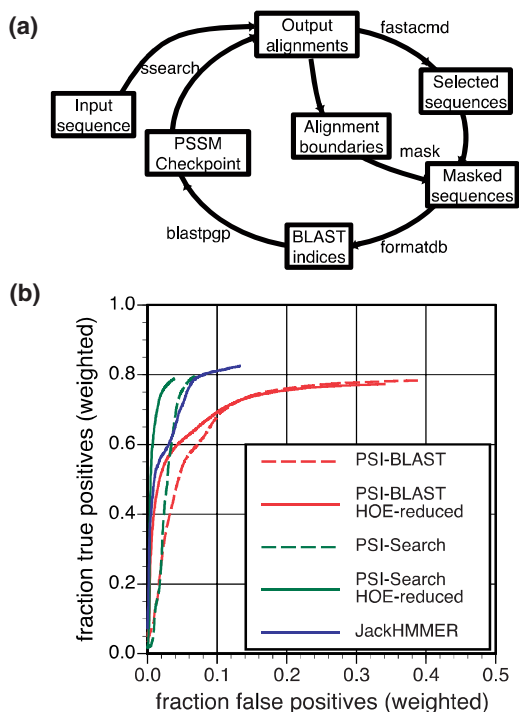
In PSI-Search, library searches are performed with *ssearch*, selected hit sequences from the result are processed with an automated sequence boundary-masking procedure, and PSSM profiles are built using *blastpgp*. The PSI-Search iteration workflow (Fig. 1a) iterates through search and alignment/PSSM construction steps:

- (1) The initial iteration is a normal *ssearch* run with a sequence input.
- (2) During the second iteration, aligned sequences with statistically significant scores from the previous search are retrieved using *fastacmd*; details of the alignment boundaries are stored; sequence regions outside the boundaries are masked with ‘X’s to remove potential HOE regions; masked sequences are formatted into BLAST indexes using *formatdb* with an additional 10 000 random protein sequences created by *makeprotseq* (Rice *et al.*, 2000); and a PSSM checkpoint constructed with a *blastpgp* search; finally *ssearch* is run with the input sequence, using the generated PSSM, to complete the second iteration and output alignments.
- (3) Further iterations repeat Step (2). To avoid HOEs, PSI-Search always uses the alignment boundary information from the first significant alignment in which a library sequence appears. Thus, if the first significant alignment with a library sequence aligns residues 25–125 at iteration *i*, later alignment boundaries at iteration *i*+1 and beyond are ignored; only the initially aligned region (25–125) is used to form the PSSM.

### 3 RESULTS

Five iterative search strategies—PSI-BLAST (standard and HOE-reduced), PSI-Search (standard and HOE-reduced) and JackHMMER (Eddy, 2011)—were evaluated on the RefProtDom (Gonzalez and Pearson, 2010b) benchmark queries (500 sampled domain-embedded sequences) against the RefProtDom benchmark database using an *E*-value threshold of 0.001. JackHMMER is another iterative search tool that uses Hidden Markov Models

\*To whom correspondence should be addressed.



**Fig. 1.** (a) HOE-reduced PSI-Search iteration workflow. (b) Fraction of true-positives versus false-positives found by PSI-BLAST, PSI-BLAST HOE-reduced, PSI-Search, PSI-Search HOE-reduced, and JackHMMER. Weighted true-positives and false-positives are calculated as  $1/500 \sum_1^{500} tp_f$  (or  $fp_f$ )/ $total_f$  where  $tp_f$  (or  $fp_f$ ) is the number of true positives (or false positives) at iteration 5 and  $total_f$  is the total number of homologs for query  $f$  in the RefProtDom benchmark database. Alignments containing HOEs with >50% of the alignment outside the homologous boundary are counted as both true and false positives

(HMMs) (Johnson *et al.*, 2010) rather than a PSSM. The output alignments from the fifth iteration were classified into true positives (TPs) and false positives (FPs, Fig. 1b). At 50% family coverage, PSI-Search reduces the weighted fraction of errors from 4.5% (PSI-BLAST) to 2.9% (PSI-Search). Reducing HOE improves sensitivity even more, to 1.7% for HOE-reduced PSI-BLAST and 0.5% for HOE-reduced PSI-Search. At 50% coverage, JackHMMER performs very well using its statistical alignment envelope, producing only 1% weighted FPs, but its selectivity is worse than PSI-Search or HOE-reduced PSI-Search at 60% and 75% coverage. Overall, HOE-reduced PSI-Search is 9-fold more selective than PSI-BLAST. At the end of iteration 5, 78.3, 79.5, 77.3, 78.8 and 82.5% of weighted homologs are found by PSI-BLAST, PSI-Search, HOE-reduced PSI-BLAST, HOE-reduced PSI-Search

and JackHMMER respectively. Thus, (i) HOE-reduction greatly improves search selectivity with a small cost in sensitivity in both PSI-BLAST and PSI-Search; (ii) Both PSI-Search and JackHMMER are more sensitive and selective than PSI-BLAST; (iii) HOE-reduced PSI-Search is more selective, but slightly less sensitive, than JackHMMER. JackHMMER is the most sensitive tool, but HOE-reduced PSI-Search is the most selective iterative tool.

## ACKNOWLEDGEMENTS

**Funding:** This research was supported by the National Library of Medicine (NIH grant LM04969 to W.R.P.); European Molecular Biology Laboratory; and European Commission Research Infrastructures of the FP7 [grant agreement number 226073 SLING (Integrating Activity)].

**Conflict of Interest:** none declared.

## REFERENCES

- Agrawal,A. and Huang,X. (2009) PSIBLAST\_PairwiseStatSig: reordering PSI-BLAST hits using pairwise statistical significance. *Bioinformatics*, **25**, 1082–1083.
- Altschul,S.F. *et al.* (1997) Gapped BLAST and PSI-BLAST: a new generation of protein database search programs. *Nucleic Acids Res.*, **25**, 3389–3402.
- Altschul,S.F. *et al.* (2005) Protein database searches using compositionally adjusted substitution matrices. *FEBS J.*, **272**, 5101–5109.
- Altschul,S.F. *et al.* (2009) PSI-BLAST pseudocounts and the minimum description length principle. *Nucleic Acids Res.*, **37**, 815–824.
- Bhadra,R. *et al.* (2006) Cascade PSI-BLAST web server: a remote homology search tool for relating protein domains. *Nucleic Acids Res.*, **34**, W143–W146.
- Eddy,S.R. (2011) Accelerated profile HMM searches. *PLoS Comput. Biol.*, **7**, e1002195.
- Gonzalez,M.W. and Pearson,W.R. (2010a) Homologous over-extension: a challenge for iterative similarity searches. *Nucleic Acids Res.*, **38**, 2177–2189.
- Gonzalez,M.W. and Pearson,W.R. (2010b) RefProtDom: a protein database with improved domain boundaries and homology relationships. *Bioinformatics*, **26**, 2361–2362.
- Johnson,L.S. *et al.* (2010) Hidden Markov model speed heuristic and iterative HMM search procedure. *BMC Bioinformatics*, **11**, 431.
- Kim,B.H. *et al.* (2010) HangOut: generating clean PSI-BLAST profiles for domains with long insertions. *Bioinformatics*, **26**, 1564–1565.
- Li,Y. *et al.* (2011) A performance enhanced PSI-BLAST based on hybrid alignment. *Bioinformatics*, **27**, 31–37.
- Pearson,W.R. (1991) Searching protein sequence libraries: comparison of the sensitivity and selectivity of the Smith–Waterman and FASTA algorithms. *Genomics*, **11**, 635–650.
- Przybylski,D. and Rost,B. (2008) Powerful fusion: PSI-BLAST and consensus sequences. *Bioinformatics*, **24**, 1987–1993.
- Rice,P. *et al.* (2000) EMBOSS: the European molecular biology open software suite. *Trends Genet.*, **16**, 276–277.
- Smith,T.F. and Waterman,M.S. (1981). Identification of common molecular subsequences. *J. Mol. Biol.* **147**, 195–197.
- Stojmirović,A. *et al.* (2008) The effectiveness of position- and composition-specific gap costs for protein similarity searches. *Bioinformatics*, **24**, i15–i23.
- Walsh,T.P. *et al.* (2008) SCANPS: a web server for iterative protein sequence database searching by dynamic programming, with display in a hierarchical SCOP browser. *Nucleic Acids Res.*, **36**, W25–W29.