Sequence analysis

Top-Down Crawl: a method for the ultra-rapid and motif-free alignment of sequences with associated binding metrics

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Abstract

Summary: Several high-throughput protein–DNA binding methods currently available produce highly reproducible measurements of binding affinity at the level of the k-mer. However, understanding where a k-mer is positioned along a binding site sequence depends on alignment. Here, we present Top-Down Crawl (TDC), an ultra-rapid tool designed for the alignment of k-mer level data in a rank-dependent and position weight matrix (PWM)-independent manner. As the framework only depends on the rank of the input, the method can accept input from many types of experiments (protein binding microarray, SELEX-seq, SMiLE-seq, etc.) without the need for specialized parameterization. Measuring the performance of the alignment using multiple linear regression with 5-fold cross-validation, we find TDC to perform as well as or better than computationally expensive PWM-based methods.

Availability and implementation: TDC can be run online at https://topdowncrawl.usc.edu or locally as a python package available through pip at https://pypi.org/project/TopDownCrawl.

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Supplementary information: Supplementary data are available at Bioinformatics online.

1 Introduction

High-throughput in vitro binding methods, such as protein binding microarrays (Berger et al., 2006), SELEX-seq (Riley et al., 2014; Slattery et al., 2011) and SMiLE-seq (Ishikawa et al., 2017), have given researchers the ability to precisely quantify transcription factor (TF) binding in a controlled environment using unbiased pools of DNA. For each of these methods, the enrichment of each individual probe is not as informative as the enrichment of k-mers. While each full-length probe may occur a few times within a sample, shorter k-mers will occur more frequently, providing highly reproducible measures of binding affinity. Since k-mer enrichment is inherently context-free, it is common to see a high level of enrichment for k-mers that only covers a portion of the binding site. Determining which part of the binding site a k-mer covers depends on alignment. Alignment allows researchers to pinpoint TF–DNA interactions along the binding site and is a necessary step in the application of conventional machine learning approaches such as multiple linear regression (MLR). Previously described approaches, such as MEME (Bailey & Elkan, 1994), BEESEM (Ruan et al., 2017) and SelectGLM (Zhang et al., 2018) are designed to generate position weight matrices (PWMs) which can subsequently be used to align k-mer level data, but they were not developed for this purpose. Furthermore, using a PWM to summarize binding preferences for a TF is an unnecessary abstraction from the original k-mer level data and results in the loss of information regarding interdependencies between positions of the binding site. It is already known that DNA shape is dependent on local interactions across several base pairs (bp) and plays a significant role in protein–DNA binding for many TFs (Yang et al., 2017). Here, we describe a new approach called Top-Down Crawl (TDC), which can rapidly align large sets of k-mer level quantitative binding data in a rank-dependent manner that does not depend on experiment-specific parameterization.

2 TDC implementation

TDC was developed with one goal in mind: the usage of high-affinity sequences to describe the binding of similar, but lower-affinity sequences. Then, use those sequences to align other similar sequences. More specifically, the algorithm starts by assigning the k-mer with the largest binding metric a shift of 0 bp and is set as the first reference. All unaligned k-mers that are one single bp mutation away from the reference are then added to the alignment and
assigned a shift equal to that of the reference (0 in this case). All unaligned k-mers overlapping the reference by up to k–2 bp are then added to the alignment with a shift equal to that of the reference ± 1 or 2 bp depending on if that sequence is overlapping on the 5’ or 3’ end of the reference (Fig. 1A). For example, the sequence AGTAAAC would overlap with the 5’ end of GTAACA in a shift of –1 bp. After this round, the reference sequence is marked as ‘complete’ and the next reference is determined as the most enriched k-mer amongst those which have been aligned, excluding sequences already marked as complete. As before, the new reference is used as a starting point for the addition of more k-mers to the alignment, so long as they have not been added previously. This process is terminated when all sequences added to the alignment have been marked as ‘complete’.

TDC is provided through a freely accessible webserver and requires only a list of sequences with their corresponding binding measurements as input. After processing the upload, the user is given a summary of the alignment as well as a representative PWM, weighting each sequence by its associated binding metric. This is most appropriate for SELEX-seq data, for which the relative enrichment is expected to approximate the relative binding affinity (Riley et al., 2014; Slattery et al., 2011). The alignment itself is provided as a tab-delimited file where gaps are represented by the ‘_’ character. The output is stored for 48 h and can be accessed using a unique link generated for the submission. TDC can also be run locally as a python package available through pip.

3 Results and comparisons
Given a PWM from a motif-generating method such as MEME, BEESEM or SelexGLM, k-mers can be assigned to their most likely ‘shift’ relative to the reference, similar to TDC. This is done by padding a given PWM with neutral positions on the 5’ and 3’ ends, then sliding each k-mer along every window along the PWM to see which shift results in the highest score. The alignments generated by these methods can then be directly compared with those provided by TDC. A generalizable workflow for PWM-based k-mer alignment and evaluation is provided at https://github.com/bhcooper/TDC_evaluation.

MEME is a well-established method for the alignment of sequences, but it does not take quantitative data as input. Therefore, every sequence is weighted equivalently in the construction of the PWM. For this reason, we run MEME using only k-mers with a log enrichment two standard deviations above the mean. The resulting PWM can then be used for the alignment of all k-mers as described above. Alternatively, BEESEM was made specifically for creating PWMs from SELEX-seq data but is computationally limited to producing motifs no longer than 10 bp and relies on subsampling for particularly large datasets (Ruan et al., 2017). Although SelexGLM is able to generate much longer PWMs (Zhang et al., 2018), the currently available implementation has considerable memory requirements (Supplementary Table S2, Supplementary Fig. S1), and the output depends on the specification of several hyperparameters.

To compare TDC with alternate alignment methods, we include the analysis of 12 SELEX-seq datasets that have previously been published (Abe et al., 2015; Dantas Machado et al., 2020; Zhang et al., 2018). We use a k-mer length of 10 bp, which covers the known binding site for most of the TFs considered and contains more information about suboptimal binding sites compared to longer k-mers. Although the goal of TDC is alignment rather than PWM generation, we can generate a logo from each alignment, weighting each sequence by its relative enrichment (Fig. 1C, Supplementary Fig. S2). We found the resulting PWMs to be most similar to those generated by BEESEM, with additional information outside the center of the binding site, covering about 15 informative positions for the androgen and glucocorticoid receptor binding sites (Supplementary Fig. S2).

For a more in-depth comparison, we determine what percent of significantly enriched 10-mers are assigned to the same shift, using TDC as the reference. TDC showed a high level of agreement with BEESEM, followed by MEME and SelexGLM (Supplementary Table S3, Supplementary Fig. S1). To evaluate the quality of each alignment, an MLR model was trained to predict the log enrichment of aligned 10-mers which were significantly enriched. Base pairs were one-hot encoded for each position, and the predicted minor groove width and electrostatic potential were included to account for interdependencies between positions (Chiu et al., 2016). For MLR to perform well, sequences need to be aligned at such that position-specific permutations along the binding site predictably modulate binding affinity. We found TDC to exhibit the best average performance as measured by the median $R^2$ using 5-fold cross validation (Fig. 1B, Supplementary Table S4). Comparing the wall-clock times, BEESEM was the slowest, requiring hours to complete, whereas TDC only takes seconds (Supplementary Table S5 and Supplementary Fig. S1). SelexGLM was faster than BEESEM but required a large amount of memory in the tests performed (Supplementary Table S2 and Supplementary Fig. S1). The primary reason these methods are more computationally demanding is because they work at the level of the full-length read rather than at the level of the k-mer. Since MEME was only used to align k-mers with a log enrichment two standard deviations above the mean, its wall-clock time was dependent on the number of sequences passing this threshold. While the quickest batch, including 653 sequences, was aligned in about 5 s, the slowest batch, including just 4889 sequences took 18 min, demonstrating a 216-fold increase in wall-clock time for a 7.5-fold increase in the number of sequences aligned (Supplementary Table S5). Finally, we tested the MLR performance of various length k-mers aligned with TDC and found the optimal length to be 10 bp (Supplementary Fig. S3).

4 Conclusions
Although we demonstrate TDC’s speed and performance using SELEX-seq data, the alignment framework is highly flexible as it...
only depends on the rank of the sequences provided. This allows for the alignment of binding data from a variety of experimental approaches used today and those that are produced in the future.

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References
Abe, N. et al. (2015) Deconvolving the recognition of DNA shape from sequence. Cell, 161, 307–318.
Bailey, T.L. and Elkan, C. (1994) Fitting a mixture model by expectation maximization to discover motifs in biopolymers. Proc. Int. Conf. Intell. Syst. Mol. Biol., 2, 28–36.
Berger, M.F. et al. (2006) Compact, universal DNA microarrays to comprehensively determine transcription-factor binding site specificities. Nat. Biotechnol., 24, 1429–1435.
Chiu, T.P. et al. (2016) DNAshapeR: an R/bioconductor package for DNA shape prediction and feature encoding. Bioinformatics, 32, 1211–1213.
Dantas Machado, A.C. et al. (2020) Landscape of DNA binding signatures of myocyte enhancer factor-2B reveals a unique interplay of base and shape readout. Nucleic Acids Res., 48, 8529–8544.
Isakov, A. et al. (2017) SMiLE-seq identifies binding motifs of single and dimeric transcription factors. Nat. Methods, 14, 316–322.
Riley, T.R. et al. (2014) SELEX-seq: a method for characterizing the complete repertoire of binding site preferences for transcription factor complexes. In: Graba, Y., Rezsohazy, R. (eds) Hox Genes. Methods in Molecular Biology, Vol. 1196, Humana Press, New York, NY, pp. 255–278.
Ruan, S. et al. (2017) REESEM: estimation of binding energy models using HT-SELEX data. Bioinformatics, 33, 2288–2295.
Slattery, M. et al. (2011) Cofactor binding evokes latent differences in DNA binding specificity between Hox proteins. Cell, 147, 1270–1282.
Yang, L. et al. (2017) Transcription factor family-specific DNA shape readout revealed by quantitative specificity models. Mol. Syst. Biol., 13, 910.
Zhang, L. et al. (2018) SelexGLM differentiates androgen and glucocorticoid receptor DNA-binding preference over an extended binding site. Genome Res., 28, 111–121.