SAPPHIRE: a neural network based classifier for σ70 promoter prediction in Pseudomonas

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Abstract

Background: In silico promoter prediction represents an important challenge in bioinformatics as it provides a first-line approach to identifying regulatory elements to support wet-lab experiments. Historically, available promoter prediction software have focused on sigma factor-associated promoters in the model organism E. coli. As a consequence, traditional promoter predictors yield suboptimal predictions when applied to other prokaryotic genera, such as Pseudomonas, a Gram-negative bacterium of crucial medical and biotechnological importance.

Results: We developed SAPPHIRE, a promoter predictor for σ70 promoters in Pseudomonas. This promoter prediction relies on an artificial neural network that evaluates sequences on their similarity to the –35 and –10 boxes of σ70 promoters found experimentally in P. aeruginosa and P. putida. SAPPHIRE currently outperforms established predictive software when classifying Pseudomonas σ70 promoters and was built to allow further expansion in the future.

Conclusions: SAPPHIRE is the first predictive tool for bacterial σ70 promoters in Pseudomonas. SAPPHIRE is free, publicly available and can be accessed online at www.biosapphire.com. Alternatively, users can download the tool as a Python 3 script for local application from this site.

Background

Promoter prediction in prokaryotes has received a lot of attention over the past two decades, to enhance the understanding and construction of gene regulatory networks [14]. Several tools implementing diverse algorithms, ranging from simple motif searches to complex machine learning techniques such as neural networks and support vector machines, have been developed and made available to the scientific community [7, 10, 11]. A unique approach was proposed in a paper studying thermodynamic stability of DNA as a feature for promoter prediction, rather than DNA motifs [9]. A key limitation to many of these bacterial promoter prediction tools is their bias towards σ factors from Escherichia coli. Consequently, application of these tools to other bacterial species yields suboptimal promoter predictions.
We here describe the Sequence Analyser for the Prediction of Prokaryotic Homology Inferred Regulatory Elements (SAPPHIRE), a tool developed to predict σ70 promoters in *Pseudomonas aeruginosa* and *Pseudomonas putida*. *P. aeruginosa* strains are opportunistic, multidrug-resistant pathogens of the highest global priority [8, 15]. *Pseudomonas putida* is a promising bacterial chassis for synthetic biology applications with an industrial scope [5, 6]. Currently, 4660 and 127 sequenced genomes of *P. aeruginosa* and *P. putida* are available on the Pseudomonas Genome Database, respectively [16]. Yet only a small fraction of these genomes contains annotated promoters, illustrating the shortage of available tools for promoter annotation.

The underlying model of SAPPHIRE combines the strong predictive power of a fully-connected artificial neural network with the traditional approach of relying on –35 and –10 boxes of σ70 promoter sequences, which are the distinguishing features of σ70 promoters and have been thoroughly analysed using information theory [12].

**Implementation**

**Data**

SAPPHIRE was trained using a dataset of 170 unique Pseudomonas σ70 promoters (Additional file 1). Ninety four of these sequences were taken from experimentally validated *P. aeruginosa* and *P. putida* σ70 promoters [3]. The 76 remaining σ70 promoters were retrieved from the NCBI Nucleotide database (database query for annotated *P. aeruginosa* and *P. putida* sequences containing keywords “minus_35_signal” and “minus_10_signal”).

Sixteen thousand background sequences were randomly extracted from intergenic genomic regions that were not annotated as promoters in the *P. aeruginosa* PAO1 genome. Extraction of background sequences from these Pseudomonas sources provided a biologically meaningful set of negative examples for training as these intergenic sequences compete with σ70 promoters for RNAP binding in the cytoplasm. The strong imbalance between positive and negative examples in the dataset is justified by a similar ratio of promoter to non-promoter sequences in bacterial genomes. Prior to training, the complete dataset of positive and negative examples was randomly divided in a training set and test set using a 9:1 ratio.

To ensure the quality of the data, the negative sequences were positively verified to contain no obvious sequence motifs that could unwillingly be learned by the neural network, using the MEME suite [1]. Furthermore, the degree of conservation of the 12 nucleotides in the –10 and –35 boxes of between any two of the positive and any two of the negative sequences were found to be 5.6/12 and 3.2/12 on average. For the positive sequences, this degree of conservation, averaging below 50%, confirms satisfactory independence of the training sequences. This is also the case for the negative examples, with a degree of conservation that is not significantly higher than what would be expected if they were randomly generated sequences (3/12, 25%). The positive and negative examples contained average GC-contents of 47 and 61%, which is a notable difference and a feature that could possibly be unwillingly incorporated in the neural network. However, this low GC-content was not compensated for, as it provides a potentially valuable distinguishing characteristic of promoter regions for a neural network, as illustrated by the consensus sequence TTGACA-TATAAT (17% GC).
Features
From the sequences that were collected to train the neural network, only the six nucleotides in both the −35 and −10 boxes were used as features for training. For each sequence, these twelve nucleotides were one-hot encoded, providing data features that can be passed to a neural network.

Neural network architecture
The core architecture of the neural network (Fig. 1) consists of two consecutive fully connected layers, feeding into a single-node third layer. The first layer contains $30 \times 4$ nodes, the depth of 4 being a remnant of the one-hot encoding of the input sequences. The second layer (16 nodes) flattens the two dimensions into a single dimension. A single node in the third layer presents the output of the network. The activation function of the nodes in the first/second layer is the rectified linear unit (ReLU). A sigmoid activation function was chosen for the final node, as it is well-suited for binary classifiers due to its output value between 0 and 1. The choices of all neural network design parameters are the result of optimization by manual tuning of the network, using the performance of fivefold cross-validation on the training set as a measure.
Significance estimation

We introduced a framework to estimate the $p$-value of positive hits, providing a measure for the significance of sequences classified as promoters. We defined the $p$-value of a hit as the probability of a randomly generated sequence to produce a value at the output node of the neural network equal to or greater than the output value produced by that positive hit. In order to estimate this probability, an estimation of the posterior probability distribution of the output value of the neural network was generated using Monte Carlo sampling (10 million random sequences). An approximation of the $p$-value is then correspondingly calculated as the fraction of random output values in this distribution equal to or greater than the output value produced by a new positive hit.

Results

Benchmarking

Due to the imbalanced nature of the dataset, accuracy was expected to accommodate a bias towards specificity and mask the minority-class performance (true positive rate). Therefore, accuracy would not provide a suitable metric in this case. To avoid this bias, sensitivity (true positive rate) and specificity (true negative rate) were chosen as metrics for evaluation. After optimization of the design parameters of the network, the model yielded values of 76.6 and 88.1% for mean sensitivity and mean specificity respectively in fivefold cross-validation on the training set. These results indicate a good overall performance of the model. Additionally, the model for *Pseudomonas* σ70 promoters specifically was compared to two established online tools for bacterial promoter prediction, BPROM [13] and CNNPromoter_b [14]. Both the complete and test dataset were analyzed using SAPPHIRE, BPROM and CNNPromoter_b (Table 1).

The results show that SAPPHIRE outperforms both BPROM and CNNPromoter_b in terms of sensitivity and shows similar performance in terms of specificity, scoring slightly better than BPROM and slightly worse than CNNPromoter_b. The large difference in sensitivity implies that SAPPHIRE effectively better distinguishes σ70 promoters from background sequences in *Pseudomonas*. Furthermore, the notably poor score of 18.8% for sensitivity on the complete dataset (32 out of 170 promoter sequences detected) from both BPROM and CNNPromoter_b corroborates the need for *Pseudomonas* specific promoter prediction tools.

Case study: uncovering σ70 promoters in *Pseudomonas* phages

To further verify the quality of SAPPHIRE as a predictive tool, we applied it to scan the intergenic regions on the genome of *Pseudomonas aeruginosa* bacteriophage LUZ19. The family of the *Autographivirinae*, of which LUZ19 is a member, is known to rely on the

| Tool             | Test set | Complete dataset |
|------------------|----------|------------------|
|                  | Sensitivity | Specificity | Sensitivity | Specificity |
| BPROM            | 23.5%    | 78.8%           | 18.8%      | 79.9%      |
| CNNPromoter_b    | 23.5%    | 86.9%           | 18.8%      | 87.2%      |
| SAPPHIRE         | 88.2%    | 82.9%           | 87.1%      | 82.0%      |

Table 1 Benchmarking of SAPPHIRE against online available promoter prediction tools

Sensitivity and specificity for each tool and both the test set and complete dataset.
host’s σ70 transcriptional apparatus during early infection [4]. Correspondingly, three σ70 promoters at genomic locations 913, 982 and 1147 have already been identified and annotated early on the LUZ19 genome. Using SAPPHIRE with a p-value cutoff of $2 \times 10^{-4}$, four additional promoters could be identified in LUZ19’s intergenic regions. Figure 2 shows these promoters at their specific locations on the LUZ19 genome, along with the transcriptomic landscape of LUZ19 during early infection, as determined by RNA-seq [2]. Each of the predicted sequences correlates to the start of a transcribed genomic region, substantiating the ability of SAPPHIRE as a tool to identify new σ70 promoters. Remarkably, one of the identified promoters on LUZ19 seems to drive the early expression of a cluster of genes including the head-tail connector, scaffolding protein and major capsid protein, structural phage genes which are generally thought to be expressed under a phage RNAP-specific promoter during late infection in members of the Autographivirinae.

Furthermore, the intergenic regions of the genomes of four representative members of the Autographivirinae, infecting a variety of Pseudomonas species were subjected to the SAPPHIRE software (Fig. 3). In addition to identifying the promoters that had previously been annotated on these genomes, SAPPHIRE predicts additional promoters on these genomes. These newly predicted promoter sequences deviate from the −35 and −10 consensus sequences, offering an explanation for why they had not yet been annotated. However, all of the newly discovered promoter sequences are consistent with the genome organization architecture of this clade of viruses. Indeed, Autographivirinae, are known to encode σ70 promoters at the left end of their genomes, driving expression of a phage-encoded RNA polymerase that subsequently transcribes the remainder of phage genes. These biologically consistent findings on phage genomes suggest that SAPPHIRE provides suitable predictions for multiple members across the Pseudomonas genus.

**Conclusion**

SAPPHIRE is the first online predictive program that specifically targets σ70 promoters in Pseudomonas and its viruses. This new tool combines the traditional approach to promoter prediction of searching −35 and −10 boxes with the strong predictive capabilities offered by a neural network architecture. Our stringently selected dataset and a focused number of nucleotide features used by the neural network ensured a model
with high sensitivity and specificity. In line with its initial objective, SAPPHIRE was shown to outperform other σ70 promoter prediction tools for Pseudomonas. In future, this tool can be expanded towards other sigma factors and species, depending on the availability of experimental datasets.

Availability and requirements
Project name: SAPPHIRE
Project home page: www.biosapphire.com
Operating system(s): Platform independent
Programming language: Python 3.
Other requirements: Python3 and python packages: numpy, keras, biopython.
License: Creative Commons Attribution 4.0 International License (http://creativecommons.org/licenses/by/4.0/).
Any restrictions to use by non-academics: None.

Supplementary information
Supplementary information accompanies this paper at https://doi.org/10.1186/s12859-020-03730-z.

Additional file 1.

Abbreviations
RNAP: Ribonucleic acid polymerase; RNA-seq: RNA-sequencing; SAPPHIRE: Sequence Analyser for the Prediction of Prokaryotic Homology Inferred Regulatory Elements; ReLU: Rectified Linear Unit

Acknowledgements
The authors thank Maarten Boon for his MSc supervision of L.C.

Authors’ contributions
L.C. and R.L. conceived, designed and tested the SAPPHIRE software, and wrote the manuscript. The authors have read and approved the manuscript.

Funding
This work was supported by a European Research council consolidator grant ‘BIONICBACTERIA’ (819800) awarded to R.L. (personnel & publication costs).

Availability of data and materials
The dataset used during the study is available in Additional file 1.
Ethics approval and consent to participate
Not applicable.

Consent for publication
Not applicable.

Competing interests
The authors declare that they have no competing interests.

Received: 11 February 2020 Accepted: 31 August 2020
Published online: 22 September 2020

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