scanRBP

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What is scanRBP?

scanRBP loads RNA-protein binding motif PWM and computes the log-odds scores for all the loaded RBPs across a given genomic sequence + draws a heatmap of the scores.

The scores can be described as follows (biopython docs):

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Here we can see positive values for symbols more frequent in the motif than in the background and negative for symbols more frequent in the background. 0.0 means that it's equally likely to see a symbol in the background and in the motif.

Using the background distribution and PWM with pseudo-counts added, it's easy to compute the log-odds ratios, telling us what are the log odds of a particular symbol to be coming from a motif against the background.

For more information, see the biopython docs.

Installation

The easiest way to install **scanRBP** is to simply run:

Unset pip install scanRBP

Note that on some systems, **pip** is installing the executable scripts under ~/.local/bin. However this folder is not in the PATH which will result in "command not found" if you try to run "scanRBP" on the command line. To fix this, please execute "export PATH="\$PATH:~/.local/bin" (and add this to your .profile). Another suggestion is to install inside a virtual environment (using virtualenv).

If you would like instead to install the latest developmental version from this repository:

```
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# clone scanRBP GitHub repository
git clone https://github.com/grexor/scanRBP.git
# build and install
./build.sh
```

Quick Start

scanRBP quick start:

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Usage for single sequence: scanRBP sequence output [options]
    * one sequence provided on the command line, generates output.png/pdf +
output.tab
Usage for processing FASTA file: scanRBP filename.fasta [options]
    * one heatmap/matrix will be generated per sequence
    * output name of the files will be sequence ids provided in the fasta file
Options:
    -annotate
                            Annotate each heatmap cell with the number
    -xlabels
                            Display sequence (x-labels), default False
    -only_protein TARDBP Only analyze binding for the specific protein /
search by name
    -all_protein TARDBP
                          Additionally to one motif per protein (for all
proteins), also include all motifs (PWMs) for this specific protein (search by
name)
                            (note that one protein can have several PWMs)
    -figsize "(10,20)"
                            Change matplotlib/seaborn figure size for the
heatmap, example width=10, height=20
    -heatmap title
                      Make heatmap (png+pdf) with title
    -output_folder folder Store all results to the output folder (default:
current folder)
                            All negative vector values are set to 0, not
    -nonzero
enabled by default
```

Examples:

Database

Currently, scanRBP is using the mCross PWM database of 112 RBPs from the paper:

Feng H, Bao S et al.

Modeling RNA-Binding Protein Specificity In Vivo by Precisely Registering Protein-RNA Crosslink Sites Molecular Cell, 2019

```
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# to download PWMs
wget http://zhanglab.c2b2.columbia.edu/data/mCross/eCLIP_mCross_PWM.tgz
--no-check-certificate
tar xfz eCLIP_mCross_PWM.tg
```

Additional PWM datasets

https://genomebiology.biomedcentral.com/articles/10.1186/s13059-023-02913-0 https://static-content.springer.com/esm/art%3A10.1186%2Fs13059-023-02913-0/MediaObjects/13059_20 23_2913_MOESM6_ESM.txt

CLIP datasets

bedGraph files

https://www.encodeproject.org/metadata/?status=released&internal_tags=ENCORE&assay_title=eCLIP&biosample_ontology.term_name=K562&biosample_ontology.term_name=HepG2&type=Experiment&files.analyses.status=released&files.preferred_default=true

Gene Annotation

Gene metadata (names, aliases) donwloaded from: https://www.ncbi.nlm.nih.gov/gene/?term=human[organism]

scanRBP Github: https://github.com/grexor/scanRBP