

PICS: Probabilistic Inference for ChIP-Seq

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A step-by-step guide in the analysis of ChIP-Seq data using the PICS package in R

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1 Licensing

Under the Artistic license 2.0, you are free to use and redistribute this software. However, if you use this package for your own publication, we would ask you to cite the following paper:

X. Zhang, G. Robertson, M. Krzywinski, K. Ning, A. Droit, S. Jones and R. Gottardo. (2009). PICS: Probabilistic Inference for Chip-seq. *Biometrics* 67, 151-163.

2 Introduction

ChIP-Seq, which combines chromatin immunoprecipitation with massively parallel short-read sequencing, can profile in vivo genome-wide transcription factor-DNA association with higher sensitivity, specificity and spatial resolution than ChIP-chip. While it presents new opportunities for research, ChIP-Seq poses new challenges for statistical analysis that derive from the complexity of the biological systems characterized and the variability and biases in its digital sequence data. We propose a method called PICS (Probabilistic Inference for ChIP-Seq) for extracting information from ChIP-Seq aligned-read data in order to identify regions bound by transcription factors. PICS identifies enriched regions by modeling local concentrations of directional reads, and uses DNA fragment length prior information to discriminate closely adjacent binding events via a Bayesian hierarchical t-mixture model. Its per-event fragment length estimates also allow it to remove from analysis regions that have atypical fragment lengths. PICS uses pre-calculated, whole-genome read mappability profile and a truncated t-distribution to adjust binding event models to compensate for reads that are missing due to local genome repetitiveness. PICS estimates uncertainties in model parameters, and these can be used to define confidence regions on binding event locations and to filter estimates.

3 PICS pipeline

A typical PICS analysis consists of the following steps:

1. Convert data to a 'GRanges' for efficient processing
2. Genome segmentation via 'segmentPICS'
3. Estimation of binding site positions and other PICS parameters via 'PICS'
4. Repeat 1-2 by swapping the IP and Control samples
5. Use 1-3 to estimate the FDR via 'picsFDR'
6. Output enriched regions and 'PICS' profiles with bed and wig files

As with any package, you first need to load it with the following command

```
library(PICS)

## Loading required package: BiocGenerics
## Loading required package: parallel
##
## Attaching package: 'BiocGenerics'
##
## The following objects are masked from 'package:parallel':
##
##   clusterApply, clusterApplyLB, clusterCall, clusterEvalQ,
##   clusterExport, clusterMap, parApply, parCapply, parLapply,
##   parLapplyLB, parRapply, parSapply, parSapplyLB
##
## The following object is masked from 'package:stats':
##
```

```
##      xtabs
##
## The following objects are masked from 'package:base':
##
##      anyDuplicated, append, as.data.frame, as.vector, cbind,
##      colnames, do.call, duplicated, eval, evalq, Filter, Find, get,
##      intersect, is.unsorted, lapply, Map, mapply, match, mget,
##      order, paste, pmax, pmax.int, pmin, pmin.int, Position, rank,
##      rbind, Reduce, rep.int, rownames, sapply, setdiff, sort,
##      table, tapply, union, unique, unlist, unsplit
##
## Loading required package: stats4
```

4 Data Input and Formating

The first step of the PICS pipeline consists of converting the aligned reads (from both IP and control samples) into a format that allows efficient segmentation of the genome into a set of candidate regions that have enough Forward and Reverse reads. The data formatting could be a bed type dataframes as well as 'AlignedReads' objects as returned by the function 'readAligned' which can read various file formats including Eland, MAQ, Bowtie, etc. Please refer to the 'ShortRead' vignette for more details on how to read data into R, other than from a BED file. In the example listed below, we use bed type files, which consists of tab delimited files with the following columns: space, start, end, strand, where each line represent a single read.

In addition to the IP and Control datasets, we use another argument consisting of a mappability profile for the genome interrogated. For each chromosome, a mappability profile for a specific read length consists of a vector of zeros and ones that gives an estimated read mappability 'score' for each base pair in the chromosome. A score of one at a position means that we should be able to align a read of that length uniquely at that position, while a score of zero indicates that no read of that length should be uniquely alignable at that position. As noted, reads that cannot be mapped to unique genomic locations are typically discarded. For convenience, and because transitions between mappable and non-mappable regions are typically much shorter than these regions, we compactly summarize each chromosome's mappability profile as a disjoint union of non-mappable intervals that specify only zero-valued profile. We store this information as a bed file where each line correspond a non-mappable intervals. PICS makes use of such mappability profiles to estimate the reads that are missing because they could not be aligned. Mappability profiles for a range of reads lengths and species are available from:

http://wiki.rglab.org/index.php?title=Public:Mappability_Profile

To make your own mappability profiles, use the script available at:

<https://github.com/SRenan/proMap/>

Once the data have been read, we can create the 'GRanges' object, which stores the sequences of genome locations and associatedd annotations. Each element of the list contains the IP and control reads as well as the mappability profiles for the corresponding chromosome. When reading the data, we can also sort and remove duplicated reads, which is recommended for a PICS analysis (sort is actually required).

In this documentation, the path for the data is: .../PICS/inst/extdata folder.

```
datadir <- system.file("extdata",package="PICS")
dataIP<-read.table(file.path(datadir, "Treatment_tags_chr21_sort.bed"), header=TRUE)
dataIP<-as(dataIP, "GRanges")

dataCont<-read.table(file.path(datadir, "Input_tags_chr21_sort.bed"), header=TRUE)
dataCont<-as(dataCont, "GRanges")
```

```
map <- read.table(file.path(datadir, "mapProfileShort"), header=TRUE)
map <- as(map, "GRanges")
```

5 PICS analysis

5.1 Genome segmentation

We segment the genome into candidate regions by pre-processing bidirectional aligned reads data from a single ChIP-Seq experiment to detect candidate regions with a minimum number of forward and reverse reads. These candidate regions will then be processed by PICS. The ‘minReads’ parameter will heavily influence the number of candidate regions returned. In principle, it is better to use a small value for ‘minReads’ in order not to miss any true binding sites. However, a value too small will likely return many false positive regions, which can result a longer computing time for the PICS function. If a control sample is available you may also chose a value of ‘minReads’ that would give you a ratio of the number of candidate regions in the control over the number of candidate regions in the IP of about 25%. A ratio larger than 25% would most likely result in a large FDR when using `picsFDR`, and therefore it is not necessary to use too low of a threshold. See below for an illustration.

```
seg<-segmentPICS(dataIP, dataC=dataCont, map=map, minReads=1)
```

5.2 Parameter estimation

After having segmented the genome into candidate regions, we use the PICS function to detect binding regions in a probabilistic way, and returns binding site estimates, confidence intervals, etc. In the code below, we assume that you are processing transcription factor data, as specified with the ‘dataType’ option set to ‘TF’ for transcription factor. We also plan to support histone modification data in a future release of PICS, which will be turned on by specifying ‘dataType=‘H’.

In order to improve the computational efficiency of the PICS package, we recommend the utilisation of the `parallel` package, which allows for easy parallel computations. In what follows, we assume that `parallel` is installed on your machine and you have at least two cores. If not, you could omit the first line and the argument ‘nCores’, and calculations will occur on a single CPU. By default the function will use only one core.

```
library(parallel)
```

In our case, assuming that we have already segmented the genome using `segmentPICS`, we can proceed with the following command:

```
pics <- PICS(seg, nCores = 2)

## Using the default paraEM
## Using the default paraPrior
## Using the parallel version of PICS with 2 cpus or cores
```

5.3 FDR estimation

In order to estimate the FDR, we need to rerun our analysis after swapping the IP and control samples. Note that this requires the presence of a control sample. We proceed with the following commands to compute the FDR after removing binding site estimates with noisy parameters as specify by the filter.

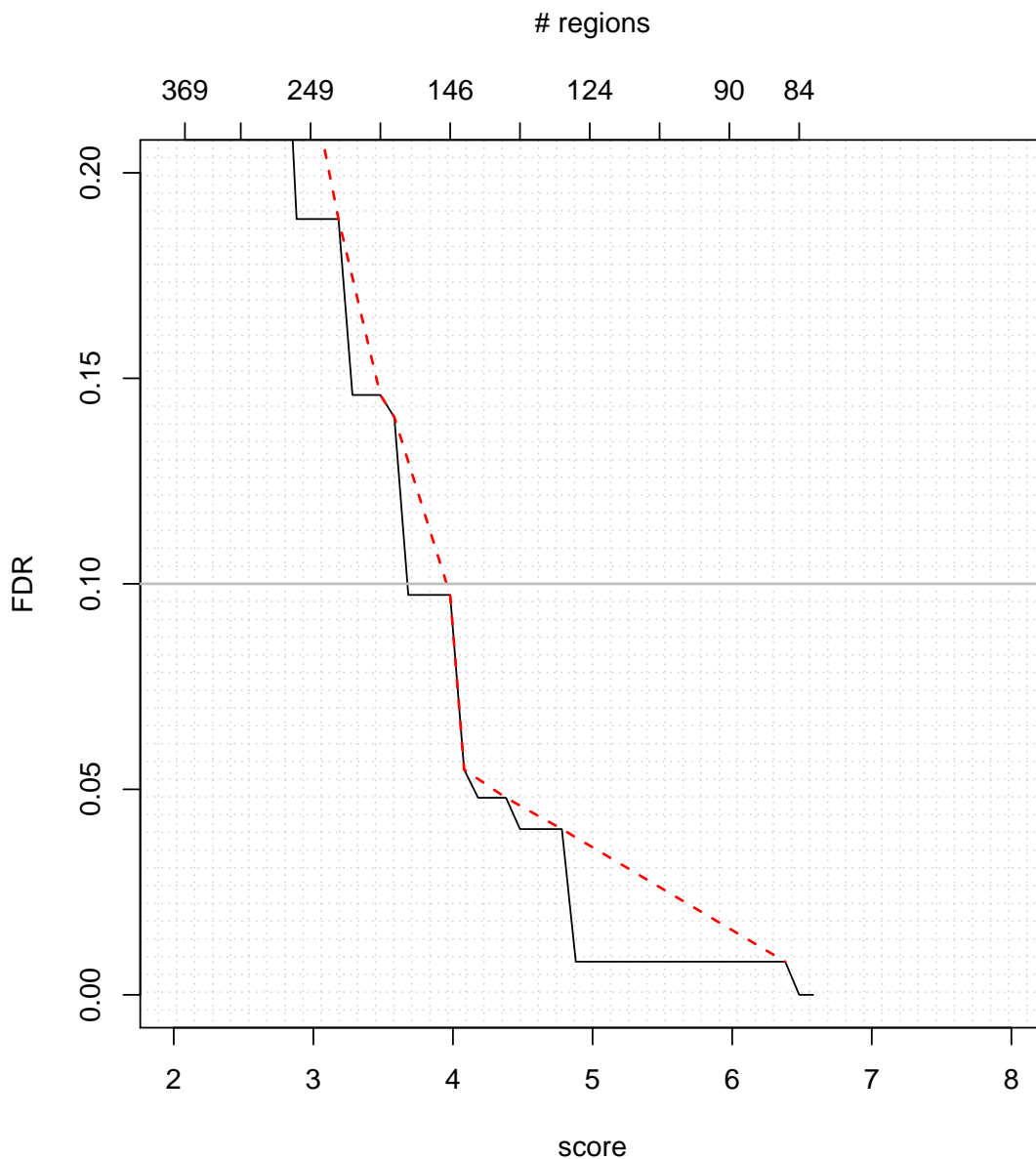
```
segC <- segmentPICS(dataCont, dataC = dataIP, map = map, minReads = 1)
picsC<-PICS(segC)

## Using the default paraEM
## Using the default paraPrior
## Using the serial version of PICS

fdr<-picsFDR(pics,picsC,filter=list(delta=c(50,Inf),se=c(0,50),
  sigmaSqF=c(0,22500),sigmaSqR=c(0,22500)))
```

Then once the FDR has been estimated, one can visualize the FDR as a function of the score to pick a threshold that would lead to the desired FDR.

```
plot(pics, picsC, xlim=c(2,8), ylim = c(0,.2),
  filter = list(delta = c(50,300), se = c(0,50), sigmaSqF = c(0,22500),
  sigmaSqR = c(0,22500)), type = "l")
```



You can also visualize the FDR as a function of the number of regions.

```
plot(fdr[,c(3,1)])
```

