Introduction to the plotAlongChrom function

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1 Introduction to the example data

The purpose of this vignette is to demonstrate some of the functionalities of the plotAlongChrom in the tilingArray package. We use a small subset data from an expression profiling paper [1]; The data only include the region from 35000bp to 50000bp in yeast chromosome one. Expression profiling is done in YPE and YPD conditions, 3 replicates each. Further information about the experimental design can be found at the paper website http://steinmetzlab.embl.de/NFRsharing/.

```
> library("grid")
> library("RColorBrewer")
> library("tilingArray")
> data("segnf")
> class(segnf)
[1] "environment"
> ls(segnf)
[1] "1.+" "1.-"
> segnf$"1.+"
Object of class 'segmentation':
Data matrix: 1775 x 6
Change point estimates for number of segments S = 1:17
Confidence intervals for 1 fits from S = 17 to 17
Selected S = 17
> head(segnf$"1.+"@y)
      YPE1 YPE2 YPE3 YPD1 YPD2 YPD3
[1,] -1.55 -2.70 -2.52 -4.78 -6.66 -4.62
[2,] -1.95 -4.20 -3.55 -3.93 -4.75 -4.74
[3,] -2.24 -2.24 -1.86 -6.15 -4.74 -3.82
[4,] -2.59 -2.52 -2.39 -4.31 -4.44 -4.40
[5,] -2.62 -3.91 -4.41 -4.94 -5.19 -4.65
[6,] -4.30 -4.82 -4.61 -5.62 -5.14 -5.60
> dim(segnf$"1.+"@y)
```

[1] 1775

```
> head(segnf$"1.+"@x)
[1] 35001 35009 35017 35025 35033 35041
> length(segnf$"1.+"@x)
[1] 1775
> segnf$"1.+"@logLik
 [1] -Inf -5136 -4772 -4627 -4562 -4498 -4428 -4372 -4314 -4280 -4235 -4198
[13] -4148 -4106 -4082 -4059 -4045
> segnf$"1.+"@nrSegments
[1] 17
> head(segnf$"1.+"@breakpoints[[segnf$"1.+"@nrSegments]])
  lower estimate upper
              13
1
     13
                    13
    158
             158
2
                    158
3
    177
             177
                    177
4
    242
             242
                    242
5
    258
             259
                   260
6
    273
             273
                   273
```

The segnf object is an environment which contains two objects of class segmentation. segnf is the output of the segChrom in the tilingArray package. The segmentation object in segnf stores the probe expression information in the slot y. As can be seen, it contains 1775 probes and 6 array hybes in two conditions. The genomic coordinates where the probes aligned to is stored in the slot x. The order of the slot x is the same as the probe row order in slot y. The segment boundary information is stored in the slot breakpoints which is a list that contains all the optimal placement of 1 segment to the designate number(here in this data set is 17) of segments for this data. A log likelihood score for each placement is stored in slot logLik from which the best one is choosen and stored in the slot nrSegments. Further information about how the segmentation algorithm works, please read the vignette segmentation demo.

```
> data(gffSub)
```

> head(gffSub))
----------------	---

	id	chr	start	end	strand	source	feature	Name	orf_classification
41	41	1	35156	36304	+	SGD	gene	YAL060W	Verified
42	42	1	35156	36304	+	SGD	CDS	YAL060W	Verified
43	43	1	36497	36919	_	SGD	gene	YALO59C-A	Dubious
44	44	1	36497	36919	_	SGD	${\tt CDS_dubious}$	YALO59C-A	Dubious
45	45	1	36510	37148	+	SGD	gene	YALO59W	Verified
46	46	1	36510	37148	+	SGD	CDS	YALO59W	Verified
	gen	e							
41	BDH	[1							

42 BDH1

43 <NA>

4.4 437.4.5

44 <NA>

45 ECM1

46 ECM1

The gffSub object is a data frame that contains the SGD annotated features of the region 35000bp-50000bp for yeast chromosome one.

2 Visualizing the expression profiling with the plotAlongChrom function

The function plotAlongChrom accepts an environment as its first argument, which is expected to contain objects of class *segmentation* with names given by paste(chr, c("+", "-"), sep="."), where chr is the chromosome identifier.

The following code generates Figure 1, a dot plot that averaged across all hybes.

```
> grid.newpage()
> plotAlongChrom(segnf,chr=1, coord=c(35000,50000),what="dots", gff=gffSub)
```

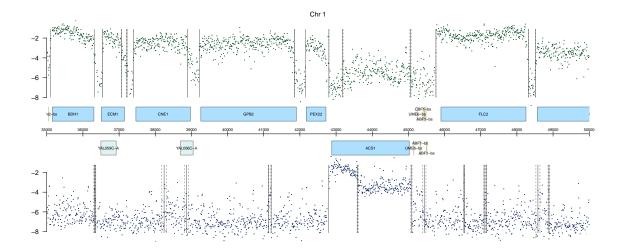


Figure 1: Along-chromosome dot plot of the averaged value across all hybes.

We could also make separate dot plot for different hybes by setting the parameter sepPlot as TRUE. The following code generates Figure 2 that plots the expression separately for the two conditions.

However, with the number of hybes increases, it is very hard to see the difference in dot plots in a normal screen. Thus, if the number of hybes is more than 4, the function will force to take the average. A better alternative of displaying multiple hybes is to use the heatmap. The following code generates Figure 3 that makes the heatmap plot.

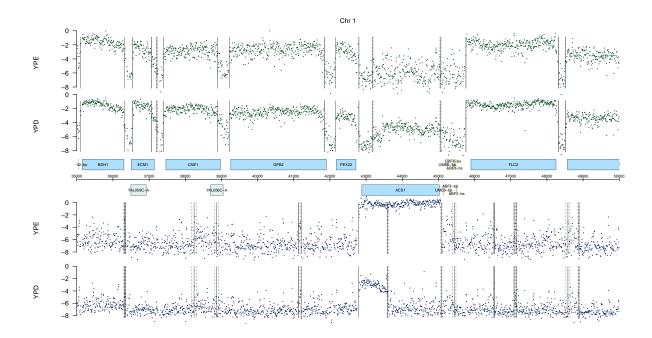


Figure 2: Along-chromosome dot plot of the averaged value among different replicates for YPD and YPE condition.

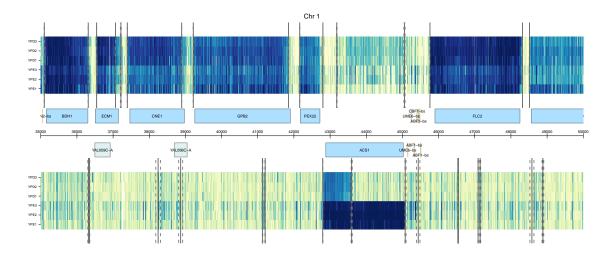


Figure 3: Along-chromosome heatmap plot of all the replicates in YPD and YPE condition.

Start with R 2.11.0, the *grid* package introduced the raster array image function <code>grid.raster</code> which is a faster and efficient way of generating heatmap images. From R 2.11.0, The *tilingArray* package will use the <code>grid.raster</code> function as default to make heatmap images replacing the previous <code>grid.rect</code> function. The choice between the two drawing functions can be changed by the parameter <code>makeRasterImage</code>. The following code generates Figure 4 that makes the raster heatmap plot.

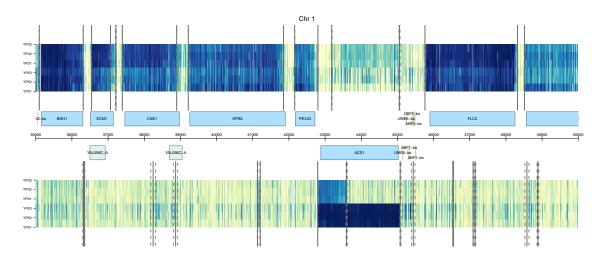


Figure 4: Along-chromosome raster heatmap plot of all the replicates in YPD and YPE condition.

The color gradient of the heatmap could be changed by the parameter *colHeatmap*. The following code generates Figure 5 that makes the raster heatmap plot using a blue color gradient.

```
> grid.newpage()
> plotAlongChrom(segnf,chr=1, coord=c(35000,50000),what="heatmap", gff=gffSub,
+ rowNamesHeatmap=nmLabel,makeRasterImage=TRUE,
+ colHeatmap = colorRamp(brewer.pal(9, "Blues")))
```

References

[1] Zhenyu Xu, Wu Wei, Julien Gagneur, Fabiana Perocchi, Sandra Clauder-Munster, Jurgi Camblong, Elisa Guffanti, Francoise Stutz, Wolfgang Huber and Lars M. Steinmetz Bidirectional promoters generate pervasive transcription in yeast. *Nature*, 2009. 1

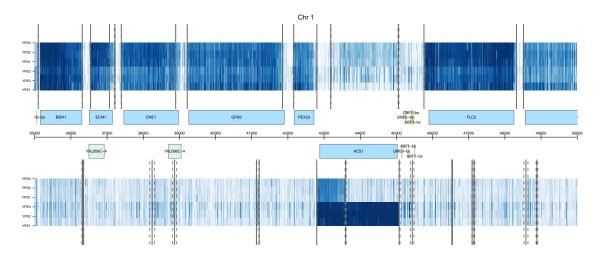


Figure 5: Along-chromosome raster heatmap plot of all the replicates in YPD and YPE condition with a blue color gradient.