

GenomeGraphs: Integrated Genomic Visualization

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- Goals:
 - ▶ High-throughput data sets
 - ▶ Changing annotation
 - ▶ Various data sources/formats
- Other Tools:
 - ▶ [rtracklayer](#)
 - ▶ [Ensembl Genome Browser](#)
 - ▶ [NCBI Entrez Map Viewer](#)
 - ▶ [UCSC's Golden Path Genome Browser](#)
 - ▶ [Statistical Viewer](#), interpretation of linkage and association data by providing a plug-in for data uploaded to the Ensembl Genome Browser.
 - ▶ [X:Map](#) genome annotation database and [exonmap](#) for exon arrays.

```
> require(GenomeGraphs)
```

```
> args(gdPlot)
```

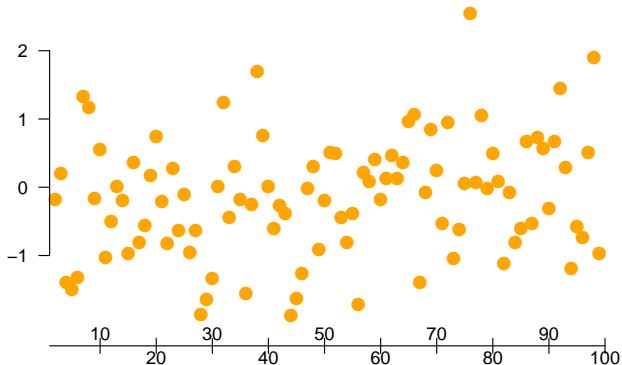
```
function (gdObjects, minBase = NA, maxBase = NA, overlays =
```

```
    labelColor = "black", labelCex = 1, labelRot = 90)
```

```
NULL
```

- gdObject
 - ▶ Gene, GeneRegion, Transcript, TranscriptRegion, Ideogram, AnnotationTrack
 - ▶ Title, Legend, GenomeAxis
 - ▶ GenericArray, ExonArray, BaseTrack
- DisplayPars
- Overlay, RectangleOverlay, TextOverlay
- Segmentable

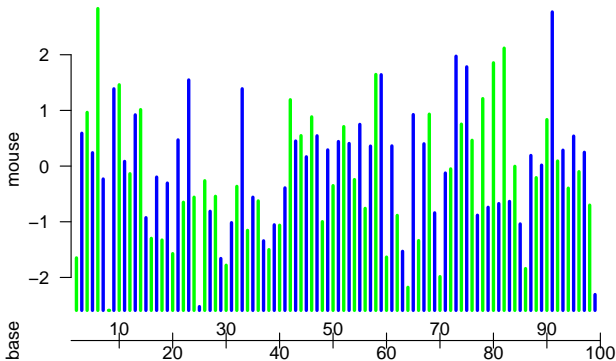
```
> lst <- list(makeBaseTrack(1:100,  
+           rnorm(100)), makeGenomeAxis())  
> gdPlot(lst)
```



- Annotate the y -axis (name the tracks).
- Change the color of the plotting symbols.
- Change the plot type (vertical bars).

Exercise 0: Answers

```
> gdPlot(list(mouse = makeBaseTrack(1:100,  
+   rnorm(100), dp = DisplayPars(type = "h",  
+   lwd = 3, color = c("blue",  
+   "green")))), base = makeGenomeAxis())
```



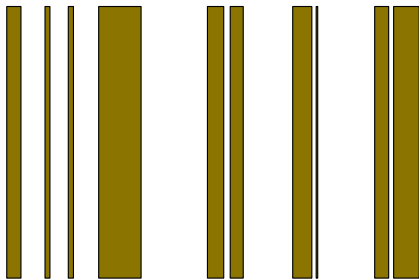
```
> showDisplayOptions("BaseTrack")  
  
color = orange  
lty = solid  
lwd = 1  
size = 5  
type = p  
  
> showDisplayOptions("GenericArray")  
  
color = darkred  
lty = solid  
lwd = 1  
pch = 16  
pointSize = 0.2  
size = 5  
type = point
```



```
> mart <- useMart("ensembl", "scerevisiae_gene_ensembl")
> data("seqDataEx")
> head(seqDataEx$daavid)

  chr location strand  expr
1   4  1300003     -1 -0.20
2   4  1300007      1  0.61
3   4  1300011     -1  0.07
4   4  1300015      1  1.25
5   4  1300019     -1 -0.29
6   4  1300023      1  0.61

> gdPlot(makeGeneRegion(10000, 50000,
+   chr = "IV", strand = "+", biomaRt = mart),
+   10000, 50000)
```



Exercise 1: Tying Annotation with Data

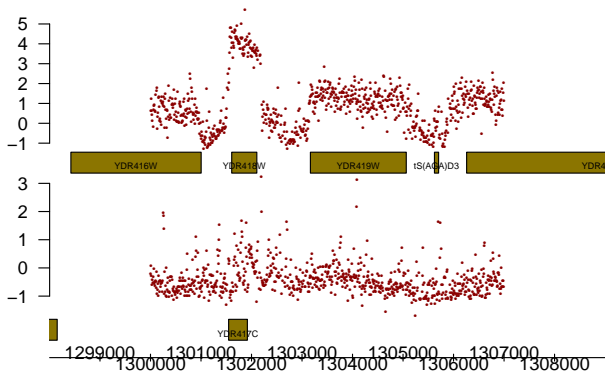
- Add the microarray data of David et al.
- Split things by strand, both annotation and array.
- Add a GenomicAxis.
- Add the gene names.

Exercise 1: Tying Annotation with Data

```
> array <- as.data.frame(seqDataEx$david)
> lst <- lapply(c("+", "-"), function(s) {
+   a <- as.matrix(subset(array,
+     strand == ifelse(s == "+",
+       1, -1)))
+   c(makeGenericArray(a[, "expr",
+     drop = FALSE], a[, "location"]),
+     makeGeneRegion(start = min(array[,
+       "location"]), end = max(array[,
+       "location"]), chr = "IV",
+       strand = s, biomart = mart,
+       dp = DisplayPars(plotId = TRUE,
+         idRotation = 0,
+         cex = 0.5, idColor = "black")))
+ })
```

Exercise 1: Tying Annotation with Data

```
> gdPlot(yeastLst <- c(unlist(lst),  
+   makeGenomeAxis()))
```

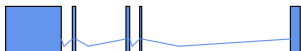
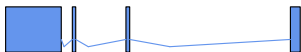


Exercise 2: Human Annotation

- Plot all the transcripts for gene: ENSG00000168309

Exercise 2: Human Annotation

```
> hMart <- useMart("ensembl", "hsapiens_gene_ensembl")  
> gdPlot(makeTranscript("ENSG00000168309",  
+   biomart = hMart))
```



```
> data("unrData", package = "GenomeGraphs")
> class(unrData)

[1] "matrix"

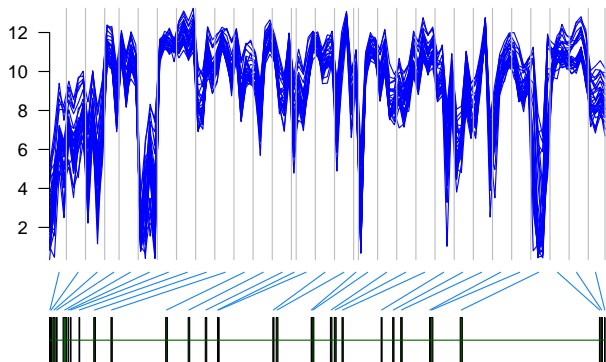
> head(unrPositions, 2)

  probesetId chromosome      start
1    2429278           1 115061081
2    2429279           1 115061152
      stop
1 115061119
2 115061198
```



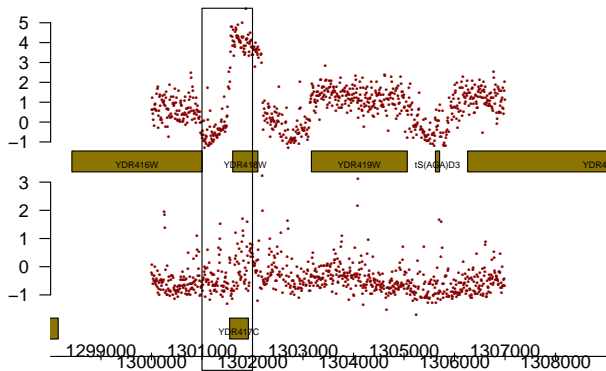
```
> exon <- makeExonArray(intensity = unrData,  
+   probeStart = unrPositions[,  
+     3], probeEnd = unrPositions[,  
+     4], probeId = as.character(unrPositions[,  
+     1]), nProbes = unrNProbes,  
+   dp = DisplayPars(color = "blue",  
+     mapColor = "dodgerblue2"),  
+   displayProbesets = FALSE)  
> geneModel <- makeGeneModel(start = unrPositions[,  
+   3], end = unrPositions[, 4])  
> gdPlot(list(exon, geneModel))
```

Exon Arrays



Overlays provide a way to plot across tracks, either using genomic coordinates or absolute coordinates. `Overlay` is an abstract class, you cannot instantiate it, only its subclasses: `RectangleOverlay`, `TextOverlay`

```
> overlay <- makeRectangleOverlay(1301000,  
+ 1302000)  
> gdPlot(yeastLst, overlays = overlay)
```



Exercise 3: Overlays

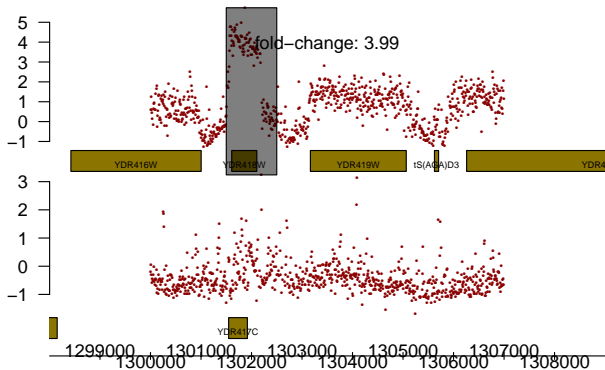
- Fix the overlay to only cover the “+” strand tracks (genes and data).
- Make the overlay semi-transparent.
- Add a text overlay adjacent to the upregulated region with the average expression of the probes in the gene YDR418W.

Exercise 3: Overlays

```
> array <- array[array$strand ==  
+   1, ]  
> anno <- yeastLst[[2]]@ens  
> anno <- anno[anno$ensembl_gene_id ==  
+   "YDR418W", c("exon_chrom_start",  
+   "exon_chrom_end")]  
> foldChange <- paste("fold-change:",  
+   round(mean(split(array$expr,  
+     findInterval(array$location,  
+     as.numeric(anno))))[[2]]),  
+   2))  
> overlay <- makeRectangleOverlay(1301500,  
+   1302500, region = c(1, 2),  
+   dp = DisplayPars(alpha = 0.5))  
> tovlay <- makeTextOverlay(foldChange,
```

Exercise 3: Overlays

```
+ 1303500, 0.75, region = c(1,  
+ 1), dp = DisplayPars(color = "black"))  
> gdPlot(yeastLst, overlays = c(ovlay,  
+ tovlay))
```

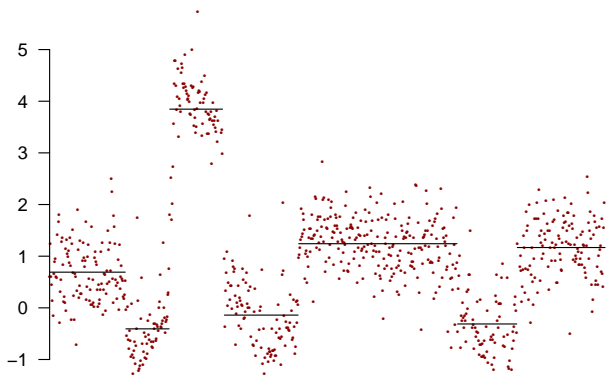


Data classes implement the interface `Segmentable` which allows one to add a piecewise constant function as an overlay. In the future, this will be expanded to more general functions.

```
> require(tilingArray)
> Y <- yeastLst[[1]]
> s <- segment(Y@intensity, maxk = 500,
+             maxseg = 20)
> bins <- findInterval(1:length(Y@intensity),
+                     s@breakpoints[[7]])
> means <- tapply(Y@intensity, bins,
+                 mean)
> ranges <- do.call(rbind, tapply(Y@probeStart,
+                               bins, range))
> Y@segmentation <- makeSegmentation(ranges[,
```

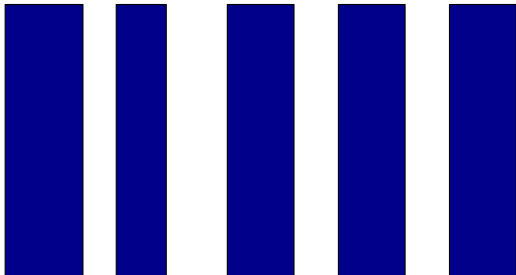


```
+      1], ranges[, 2], means)  
> gdPlot(Y)
```



We can construct custom annotation by using the `makeAnnotationTrack` function.

```
> S <- seq(1, 100, by = 20)
> E <- S + rpois(length(S), 10)
> anno <- makeAnnotationTrack(start = S,
+   end = E, feature = "gene",
+   dp = DisplayPars(gene = "darkblue"))
> gdPlot(anno, minBase = 0, maxBase = 100)
```



Exercise 4: Custom Annotation

Download the annotation file:

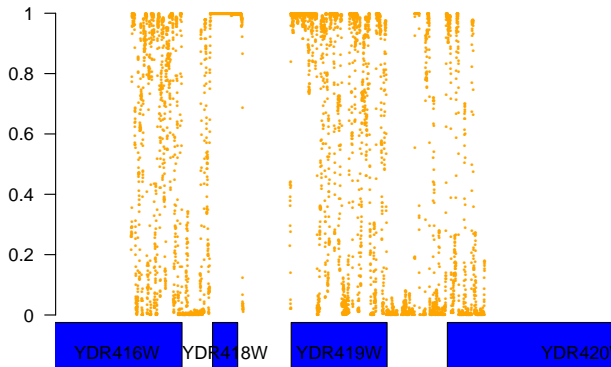
`wiki.biostat.berkeley.edu/~bullard/aleatoria/yeast_anno_4.c`

Construct a annotation track using data from this file and plot along with conservation data from the example dataset `seqDataEx`. Add gene names and determine what the group column does.

Exercise 4: Custom Annotation

```
> anno <- read.csv("http://wiki.biostat.berkeley.edu/~bullard")
> data("seqDataEx")
> aTrack <- makeAnnotationTrack(start = anno$start[anno$feature ==
+   "CDS"], end = anno$end[anno$feature ==
+   "CDS"], feature = "CDS", ID = anno$name[anno$feature ==
+   "CDS"], dp = DisplayPars(CDS = "blue",
+   plotId = TRUE, idColor = "black",
+   idRotation = 0))
> bTrack <- makeBaseTrack(seqDataEx$conservation[,
+   "location"], seqDataEx$conservation[,
+   "score"], dp = DisplayPars(type = "p",
+   lwd = 0.2))
> gdPlot(list(bTrack, aTrack), minBase = 1300000 -
+   1500, maxBase = 1307000 + 2500)
```

Exercise 4: Custom Annotation



- pch option to BaseTrack
- image option for generic array
- Add a suitable minBase and maxBase determiner for Annotation objects