## HowTo: Creating HMM objects from BeadStudio-processed Illumina arrays

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This vignette describes how to create an instance of an RatioSnpSet from Illumina data.

## 1 Reading in the data

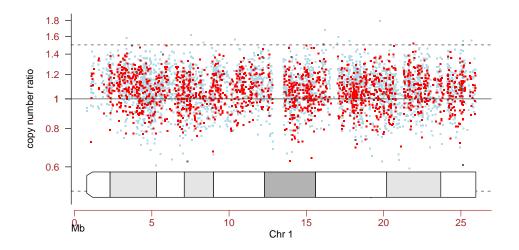
To illustrate, an example of BeadStudio output obtained from the Pevsner website (http://pevsnerlab.kennedykrieger.org/SNPtrio03.htm) is included with this package. The Pevsner laboratory provide the following instructions for saving the data using Illumina's BeadStudio in the appropriate format:

- 1. select the "Full Data Table" tab
- 2. click on the "Column Chooser" icon
- 3. in the "Displayed Columns" area, keep "Name", "Chr" and "Position", hide the rest
- 4. the "Displayed Subcolumns" area, keep "GType" and "Log R Ratio", hide the rest
- 5. click on "Export Displayed Data to File" icon; finally, save the file

A subset of 1000 SNPs is included with this package and can be loaded by

The following code converts this data frame to an object of class oligoSnpSet:

```
> gt <- illuminaEx[, "S1135.GType", drop = FALSE]</pre>
> gt[gt == "AA"] <- 1
> gt[gt == "BB"] <- 3
> gt[gt == "AB"] <- 2
> gt[gt == "NC"] <- 4
> gt <- as.matrix(as.integer(gt[[1]]))</pre>
> ratio <- 2^as.matrix(as.numeric(illuminaEx[, "S1135.Log.R.Ratio"]))</pre>
> colnames(gt) <- colnames(ratio) <- "S1135"
> rownames(ratio) <- rownames(gt) <- illuminaEx[, "Name"]</pre>
> fd <- new("AnnotatedDataFrame", data = data.frame(position = illuminaEx[,
      "Position"], chromosome = illuminaEx[, "Chr"],
      stringsAsFactors = FALSE), varMetadata = data.frame(labelDescription = c("position",
      "chromosome")))
> featureNames(fd) <- illuminaEx[, "Name"]</pre>
> callsConfidence <- ratioConfidence <- matrix(NA,
      nrow = nrow(ratio), ncol = ncol(ratio))
> rownames(callsConfidence) <- rownames(ratioConfidence) <- rownames(ratio)
> colnames(callsConfidence) <- colnames(ratioConfidence) <- colnames(ratio)
> snpset <- new("RatioSnpSet", ratio = ratio, ratioConfidence = ratioConfidence,
      calls = gt, callsConfidence = callsConfidence,
      featureData = fd, phenoData = annotatedDataFrameFrom(ratio,
          byrow = FALSE), annotation = "Illumina550k")
> chrom <- chromosome2numeric(chromosome(snpset))</pre>
> snpset <- snpset[order(chrom, position(snpset)),</pre>
      ]
> stopifnot(validObject(snpset))
   We can now use methods from the R package SNPchip to plot the data:
> gp <- plotSnp(snpset)</pre>
> gp$cex <- 3
> gp$ylab <- "copy number ratio"</pre>
> gp$abline <- TRUE
> gp$abline.h <- c(0.5, 1, 3/2)
> gp$abline.col <- "grey20"</pre>
> gp$abline.lty <- c(2, 1, 2)
> print(gp)
```



## **Session Information**

The version number of R and packages loaded for generating the vignette were:

- R version 2.7.0 alpha (2008-04-06 r45127), powerpc-apple-darwin8.11.0
- Locale: C
- Base packages: base, datasets, grDevices, graphics, methods, stats, tools, utils
- Other packages: Biobase 1.99.7, SNPchip 1.3.24, VanillaICE 1.1.21, oligoClasses 1.1.21