

```

> options(width = 75, continue = " ")
> library("Gviz")
> library("trioClasses")
> data("cnv", package = "trioClasses")
> data("pedigrees", package = "CleftCNVAssoc")
> se <- SummarizedExperiment(assays = SimpleList(cnv = t(cnv.obj$cnv.mat)),
  colData = DataFrame(id = rownames(cnv.obj$cnv.mat), row.names = rownames(cnv.obj$cnv.ma
  rowData = cnv.obj$cmp.gr)
> beaty.trios <- MinimumDistance::trios(beaty.pedigree)
> beaty.ped <- DataFrame(famid = do.call("rbind", strsplit(beaty.trios$0,
  "_"))[, 1], id = beaty.trios$0, fid = beaty.trios$F,
  mid = beaty.trios$M, sex = NA, dx = NA)
> ped <- PedClass(beaty.ped)
> fe <- FamilyExperiment(se, pedigree = ped)

> fe.parents <- fe[, colnames(fe) %in% parents(fe)]

> freq.vec <- colSums(cnv(fe.parents))/nrow(cnv(fe.parents))

> trioAssay <- trioClasses::TrioAssay(fe, type = "cnv")
> trioStates <- with(trioAssay, matrix(paste0(F, M, 0), nrow = nrow(0),
  ncol = ncol(0)))
> dimnames(trioStates) <- dimnames(trioAssay$0)

> table.list <- apply(trioStates, 2, "table")

> trans.vec <- as(lapply(table.list, trioClasses::trans.tab),
  "numeric")
> head(table.list[which(trans.vec <= 0.05/length(trans.vec))])

$comp1994

000 001 010 011 100 101 110 111
149  7  26  57  28  71  5 102

$comp1995

000 001 010 011 100 101 110 111
116  5  30  63  38  75  9 109

$comp1996

000 001 010 011 100 101 110 111
113  5  30  64  37  74  11 111

$comp1997

```

000 001 010 011 100 101 110 111
109 5 30 64 39 71 12 115

\$comp1998

000 001 010 011 100 101 110 111
109 5 30 64 38 72 12 115

\$comp1999

000 001 010 011 100 101 110 111
108 5 30 63 38 73 12 116


```

> chr <- 8
> gr.cnp <- cnv.obj$cmp.gr
> gr.cnp.chr <- gr.cnp[as.logical(seqnames(gr.cnp) == paste0("chr",
  chr))]
> p.vec <- trans.vec
> p.vec.chr <- p.vec[as.logical(seqnames(gr.cnp) == paste0("chr",
  chr))]
> freq.vec.chr <- freq.vec[as.logical(seqnames(gr.cnp) == paste0("chr",
  chr))]
> library("TxDb.Hsapiens.UCSC.hg18.knownGene")
> TranscriptDb <- TxDb.Hsapiens.UCSC.hg18.knownGene
> atrack <- AnnotationTrack(reduce(gr.cnp.chr), name = "CNP comp.",
  fill = "darkgreen")
> gtrack <- GenomeAxisTrack()
> dtrack <- DataTrack(range = gr.cnp.chr, data = -log10(p.vec.chr),
  type = "p", cex = 1, name = "-log10(p)", ylim = c(0,
  10), col = "red")
> dtrack2 <- DataTrack(range = gr.cnp.chr, data = ifelse(freq.vec.chr >
  0.01, freq.vec.chr, NA), type = "s", col = "blue", cex = 1,
  name = "Frequency", ylim = c(0, 1), lwd = 2)
> itrack <- IdeogramTrack(genome = "hg18", chromosome = paste0("chr",
  chr), lty = 1, lwd = 1)
> grtrack <- GeneRegionTrack(TranscriptDb, genome = "hg18",
  chromosome = chr, name = "UCSC Known Transcripts", showId = TRUE)
> fud <- 5e+05/2
> plotTracks(list(dtrack, dtrack2, grtrack, gtrack, itrack),
  background.panel = "#FFFEDB", background.title = "darkblue",
  from = 39356825 - fud, to = 39497557 + fud)

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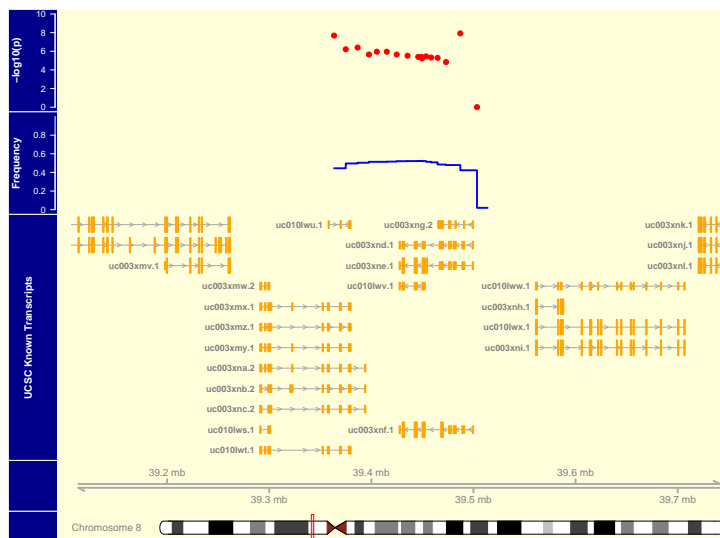


Figure 1: Foo!