

# Package ‘manhattanVAAST’

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**Type** Package

**Title** test

**Version** 1.0

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**Description** no

**License** none

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manhattanVAAST-package
<i>manhattanVAAST</i>

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## Description

manhattanVAAST plot VAAST pvalues over range. Similar to GWAS style manhattan plots.

## Details

Package:	manhattanVAAST
Type:	Package
Version:	1.001
Date:	2012-02-08
License:	none

**Author(s)**

Zev N. Kronenberg

Maintainer: <zev.kronenberg@gmail.com> Lab page: <<http://www.yandell-lab.org/software/vaast.html>>

**References**

A probabilistic disease-gene finder for personal genomes Yandell M Huff CD Hu H Singleton M Moore B Xing J Jorde L Reese MG Genome Res. 2011 Jul

Using VAAST to Identify an X-Linked Disorder Resulting in Lethality in Male Infants Due to N-Terminal Acetyltransferase Deficiency Rope AF Wang K Evjenth R Xing J Johnston JJ Swensen JJ Johnson WJ Moore B Huff CD Bird LM Carey JC Opitz JM Stevens CA Jiang T Schank C Fain HD Robison R Dalley B Chin S South ST Pysher TJ Jorde LB Hakonarson H Lillehaug JR Biesecker LG Yandell M Arnesen T Lyon GJ Am J Hum Genet. 2011 Jul 15;89(1):28-43

**See Also**

~~ Optional links to other man pages, e.g. ~~

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manhattanVAAST

*a function to plot VAAST simple reports.*


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**Description**

given some data and few lines parameters the package will generate a plot.

**Usage**

```
manhattanVAAST(n.features=23000, sig.level=0.05, vaast.simple, title, sig.line = TRUE, axis.text
```

**Arguments**

n.features	the number of genes or features VAAST scored
sig.level	The significane level sig.level = 0.05
vaast.simple	The File to do work on
title	main title
sig.line	Do you want the significance line in your plot?
axis.text	Do you want the seqids on the plot?
custom.xlab	Place your x-label here
sig.hjust	move the sig.line text horizontally + right - left (on same scale). use large numbers.
sig.vjust	move the sig.line text vertically + up - down (on same scale)

**Details**

no additional details

**Author(s)**

Zev N. Kronenberg

**Examples**

```
##---- Should be DIRECTLY executable !! ----
##-- ==> Define data, use random,
##--or do help(data=index) for the standard data sets.

## The function is currently defined as
function (n.features, sig.level, vaast.simple, title, sig.line = TRUE,
  axis.text = TRUE, custom.xlab = "index", sig.hjust = 0, sig.vjust = 0)
{
  dat <- read.table(vaast.simple, header = FALSE, skip = 1,
    sep = "\t", fill = TRUE)
  dat <- dat[dat$V4 > 0, ]
  dat <- dat[, 1:6]
  dat <- dat[grepl(dat$V5, pattern = "chr"), ]
  pos.dat <- strsplit(as.character(dat$V5), split = ":",
    perl = TRUE)
  dat$seqid <- sapply(pos.dat, FUN = function(xx) {
    return(xx[[1]][1])
  })
  dat$pos <- as.numeric(as.character(sapply(pos.dat, FUN = function(xx) {
    return(xx[[2]][1])
  })))
  dat <- dat[order(dat$seqid, dat$pos), ]
  dat$index.vec <- cumsum(1:length(dat$pos))
  dat$len.correction1 <- unlist(tapply(dat$pos, INDEX = dat$seqid,
    FUN = function(x) {
      c(max(x), rep(0, length(x) - 1))
    }, simplify = TRUE))
  dat$len.correction3 <- unlist(tapply(dat$pos, INDEX = dat$seqid,
    FUN = function(x) {
      rep(max(x), length(x))
    }, simplify = TRUE))
  dat <- dat[order(-dat$len.correction3, dat$seqid, dat$pos),
    ]
  dat$len.correction2 <- cumsum(dat$len.correction1)
  dat$relative.pos <- as.numeric(as.character(dat$len.correction2)) +
    as.numeric(as.character(dat$pos))
  axis.name <- unique(dat$seqid)
  axis.name.pos.rel <- as.vector(tapply(dat$index, INDEX = dat$seqid,
    FUN = function(x) {
      middle <- (max(x) + min(x))/2
    })))
  axis.name.pos.real <- sort(as.vector(tapply(dat$relative.pos,
    INDEX = dat$seqid, FUN = function(x) {
      middle <- (max(x) + min(x))/2
    })))
  plot(y = -log10(as.numeric(as.character(dat$V3))), x = dat$relative.pos,
    col = as.factor(dat$seqid), xaxt = "n", xlab = custom.xlab,
    pch = 20, ylab = "-log10(p-value)", main = title)
  if (sig.line == TRUE) {
    abline(h = -log10(sig.level/n.features), lty = 2, lwd = 3,
      col = "grey")
  }
}
```

```
      text(x = (0.5 * max(dat$relative.pos)) + sig.hjust, y = 0.5 +
        -log10(sig.level/n.features) + sig.vjust, "genome-wide sig. level",
        cex = 1)
    }
    if (axis.text == TRUE) {
      axis(1, at = axis.name.pos.real, labels = axis.name,
        las = 2)
    }
    if (axis.text == FALSE) {
      axis(1, at = axis.name.pos.real, labels = FALSE, las = 2)
    }
    return(dat)
  }
}
```

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