

MafDb.ExAC.r1.0.nonTCGA.GRCh38

September 27, 2023

MafDb.ExAC.r1.0.nonTCGA.GRCh38-package

Annotation package for minor allele frequency data from the Exome Aggregation Consortium

Description

This annotation package stores minor allele frequency (MAF) data derived from the variant set release 1.0 of the Exome Aggregation Consortium (ExAC), excluding TCGA samples. The data are exposed to the user in the form of a [GScores](#) object, named after the package and loaded into main memory only as different chromosomes and populations are being queried. The class definition and methods to access [GScores](#) objects are found in the [GenomicScores](#) software package. To minimize disk space and memory requirements, MAF values larger or equal than 0.1 are stored using two significant digits, while MAF values smaller than 0.1 are stored using one significant digit.

Please consult the ExAC FAQ page at <http://exac.broadinstitute.org/faq> before you use these data for your own research.

Format

[MafDb.ExAC.r1.0.nonTCGA.GRCh38](#) [GScores](#) object containing MAF values from ExAC downloaded on March 2018 from

Author(s)

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Source

Lek M et al. Analysis of protein-coding genetic variation in 60,706 humans. *Nature*, 536:285-291, 2016.

The Exome Aggregation Consortium (ExAC), Cambridge, MA (URL: <http://exac.broadinstitute.org>) [March, 2018, accessed]

See Also

[GScores-class gscores](#) [GenomicScores](#)

Examples

```
library(SNPlocs.Hsapiens.dbSNP149.GRCh38)
library(MafDb.ExAC.r1.0.nonTCGA.GRCh38)

ls("package:MafDb.ExAC.r1.0.nonTCGA.GRCh38")

mafdb <- MafDb.ExAC.r1.0.nonTCGA.GRCh38
mafdb
citation(mafdb)

populations(mafdb)

## lookup allele frequencies for rs1129038, a SNP associated to blue and brown eye colors
## as reported by Eiberg et al. Blue eye color in humans may be caused by a perfectly associated
## founder mutation in a regulatory element located within the HERC2 gene inhibiting OCA2 expression.
## Human Genetics, 123(2):177-87, 2008 [http://www.ncbi.nlm.nih.gov/pubmed/18172690]

snpdb <- SNPlocs.Hsapiens.dbSNP149.GRCh38
rng <- snpsById(snpdb, ids="rs1129038")
rng
gscores(mafdb, rng)
gscores(mafdb, GRanges("15:28111713"))
```

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