

# Package ‘GeuvadisTranscriptExpr’

October 24, 2025

**Type** Package

**Title** Data package with transcript expression and bi-allelic genotypes  
from the GEUVADIS project

**Version** 1.37.1

**Date** 2025-06-27

**Description** Provides transcript expression and bi-allelic genotypes  
corresponding to the chromosome 19 for CEU individuals from the  
GEUVADIS project, Lappalainen et al.

**Depends** R (>= 3.3.0)

**License** GPL (>= 3)

**LazyData** true

**biocViews** Homo\_sapiens\_Data, SNPData, Genome, RNASeqData,  
SequencingData, ExpressionData

**VignetteBuilder** knitr

**Suggests** limma, rtracklayer, GenomicRanges, Rsamtools,  
VariantAnnotation, tools, BiocStyle, knitr, testthat

**NeedsCompilation** no

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**git\_url** <https://git.bioconductor.org/packages/GeuvadisTranscriptExpr>

**git\_branch** devel

**git\_last\_commit** bd75e6a

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**Repository** Bioconductor 3.23

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|        |                                      |
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| counts | <i>Sample data for sQTL analysis</i> |
|--------|--------------------------------------|

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**Description**

Subsets of raw data available in this package and saved as Rdata objects for faster loading.

**Usage**

```
counts  
  
gene_ranges  
  
genotypes  
  
snp_ranges
```

**Format**

counts is a data frame with subset of counts from TrQuantCount\_CEU\_chr19.tsv  
gene\_ranges is a GRanges object containing subset of gene coordinates from genes\_chr19.bed  
genotypes is a data frame with subset of genotypes from genotypes\_CEU\_chr19.tsv  
snp\_ranges is a Granges object containing subset of SNP coordinates from genotypes\_CEU\_chr19.tsv  
For all the details on how these data sets were produced, see examples.

**Value**

counts, gene\_ranges, genotypes, snp\_ranges

**Source**

Lappalainen T, Sammeth M, Friedlander MR, et al. Transcriptome and genome sequencing uncovers functional variation in humans. Nature. 2013;501(7468):506-11

**Examples**

```
library(rtracklayer)  
data_dir <- system.file("extdata", package = "GeuvadisTranscriptExpr")  
  
gene_id_subset <- readLines(file.path(data_dir, "gene_id_subset.txt"))  
snp_id_subset <- readLines(file.path(data_dir, "snp_id_subset.txt"))  
  
# Load gene ranges with names!  
gene_ranges <- import(file.path(data_dir, "genes_chr19.bed"))  
names(gene_ranges) <- mcols(gene_ranges)$name  
  
gene_ranges <- gene_ranges[gene_id_subset, ]
```

```
# Load transcript counts
counts <- read.table(file.path(data_dir, "TrQuantCount_CEU_chr19.tsv"),
                     header = TRUE, sep = "\t", as.is = TRUE)

counts <- counts[counts$Gene_Symbol %in% gene_id_subset, ]

# Load genotypes
genotypes <- read.table(file.path(data_dir, "genotypes_CEU_chr19.tsv"),
                        header = TRUE, sep = "\t", as.is = TRUE)

genotypes <- genotypes[genotypes$snpId %in% snp_id_subset, ]

# Create SNP ranges with names!
snp_ranges <- GRanges(Rle(genotypes$chr), IRanges(genotypes$start,
                                                  genotypes$end))
names(snp_ranges) <- genotypes$snpId
```

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