

# Package ‘FlowSorted.Blood.450k’

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**Version** 1.46.0

**Title** Illumina HumanMethylation data on sorted blood cell populations

**Description** Raw data objects for the Illumina 450k DNA methylation microarrays, and an object depicting which CpGs on the array are associated with cell type.

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**License** Artistic-2.0

**Depends** R (>= 2.13.0), minfi (>= 1.21.2)

**LazyData** yes

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FlowSorted.Blood.450k *Illumina Human Methylation data from 450k on sorted blood cell populations*

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## Description

This RGset contains Illumina 450k DNA methylation measurements on 60 samples from Reinius et al. 2012, which can be used by the [minfi](#) package to estimate cellular composition from whole blood samples. This data may also be useful to individuals as example Illumina 450k data for trying preprocessing methods across a variety of Bioconductor packages.

## Usage

```
data(FlowSorted.Blood.450k)
```

## Format

An object of class `RGChannelSet`.

## Details

The `FlowSortedBlood.450k` objects is based on samples assayed as part of Reinius et al (2012). Please cite this paper, if the data is used. If you're using this data together with the [minfi](#) package, please see the package vignette for details on how to cite that package.

A script for obtaining this dataset is available in the `scripts` directory of this package.

## References

Reinius, L. E. et al. *Differential DNA Methylation in Purified Human Blood Cells: Implications for Cell Lineage and Studies on Disease Susceptibility*. PLoS ONE (2012), 7:e41361. doi:[10.1371/journal.pone.0041361](#)

Houseman E.A. et al *DNA methylation arrays as surrogate measures of cell mixture distribution*. BMC Bioinformatics (2012), 13:p86. doi:[10.1186/147121051386](#)

Jaffe A.E. and Irizarry R.A. *Accounting for cellular heterogeneity is critical in epigenome-wide association studies* Genome Biology (2013), 15:R31. doi:[10.1186/gb2014152r31](#)

## See Also

[FlowSorted.Blood.450k.JaffeModelPars](#) and [FlowSorted.Blood.450k.compTable](#) for additional datasets derived from this one. See the `minfi` package for tools for estimating cell type composition in blood using these data..

## Examples

```
data(FlowSorted.Blood.450k)
```

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FlowSorted.Blood.450k.compTable

*Cell Composition Association Table*


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## Description

Association of each probe on the Illumina 450k with blood cell composition Note that probes on the sex chromosomes and those that contain annotated single nucleotide polymorphisms (SNPs) have been filtered (see Methods section of Jaffe and Irizarry 2013, below)

## Usage

```
data(FlowSorted.Blood.450k.compTable)
```

## Format

Name: CpG identifier from the Illumina 450k Fstat: f-statistic for composition from the ANOVA containing 6 samples/biological replicates per cell type across 6 cell types p.value: corresponding p-value for f-statistic CD8T\_mean: mean DNA methylation (DNAm) across the 6 CD8+ T-cell replicates, on the beta/proportion methylation scale CD4T\_mean: mean DNA methylation (DNAm) across the 6 CD4+ T-cell replicates, on the beta/proportion methylation scale NK\_mean: mean DNA methylation (DNAm) across the 6 Natural Killer cell replicates, on the beta/proportion methylation scale Bcell\_mean: mean DNA methylation (DNAm) across the 6 B-cell replicates, on the beta/proportion methylation scale Mono\_mean: mean DNA methylation (DNAm) across the 6 monocyte replicates, on the beta/proportion methylation scale Gran\_mean: mean DNA methylation (DNAm) across the 6 granulocyte replicates, on the beta/proportion methylation scale DNAm\_min: minimum beta values across the 36 samples DNAm\_max: maximum beta values DNAm\_range: range of beta values

## Details

We recommend using the CpG identifiers to match each probe from a user's differential methylation analysis in their whole blood data to obtain the corresponding composition p-value - if there are many small p-values for significant differentially methylated sites for the exposure/outcome/trait of interest, this may be a sign of confounding via composition differences, in which case we recommend estimating cellular components using the minfi Bioconductor package, and formally exploring this potential correlation between the trait, composition, and DNA methylation.

## References

- Reinius, L. E. et al. *Differential DNA Methylation in Purified Human Blood Cells: Implications for Cell Lineage and Studies on Disease Susceptibility*. PLoS ONE 7, e41361 (2012). <http://dx.doi.org/10.1371/journal.pone.0041361>
- Houseman E.A. et al *DNA methylation arrays as surrogate measures of cell mixture distribution*. BMC Bioinformatics 13, p86 (2012). <http://dx.doi.org/10.1186/1471-2105-13-86>
- Jaffe A.E. and Irizarry R.A. *Accounting for cellular heterogeneity is critical in epigenome-wide association studies* Under Review (2013).

## See Also

[FlowSorted.Blood.450k](#) for the original data, [FlowSorted.Blood.450k.JaffeModelPars](#) for an additional dataset derived from this one. See the `minfi` package for tools for estimating cell type composition in blood using these data.

## Examples

```
data(FlowSorted.Blood.450k.compTable)
```

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```
FlowSorted.Blood.450k.JaffeModelPars
```

*Model Parameters for Blood Cell Type Estimation*

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## Description

This object can be used by [minfi](#) to estimate the relative proportion of each cell type, given user's whole blood Illumina 450k data.

## Usage

```
data(FlowSorted.Blood.450k.JaffeModelPars)
```

## Format

A matrix where rows are selected CpGs that are differentially methylated by cell type, and columns are particular cell types.

## Details

For details on how this coefficient object was created, see Methods section of Jaffe and Irizarry, 2013, below. For statistical details on how the cell estimation procedure is performed, refer to Houseman et al 2012.

## References

Reinius, L. E. et al. *Differential DNA Methylation in Purified Human Blood Cells: Implications for Cell Lineage and Studies on Disease Susceptibility*. PLoS ONE 7, e41361 (2012). <http://dx.doi.org/10.1371/journal.pone.0041361>

Houseman E.A. et al *DNA methylation arrays as surrogate measures of cell mixture distribution*. BMC Bioinformatics 13, p86 (2012). <http://dx.doi.org/10.1186/1471-2105-13-86>

Jaffe A.E. and Irizarry R.A. *Accounting for cellular heterogeneity is critical in epigenome-wide association studies* Under Review (2013).

## See Also

[FlowSorted.Blood.450k](#) for the original data, [FlowSorted.Blood.450k.compTable](#) for an additional dataset derived from this one. See the `minfi` package for tools for estimating cell type composition in blood using these data.

**Examples**

```
data(FlowSorted.Blood.450k.JaffeModelPars)
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