

# Package ‘HaploGeno’

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**Title** Haplotype-Based Genomic Prediction

**Version** 0.3.0

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**Description** Implements a high-performance framework for haplotype-based genomic prediction and QTL discovery. Leverages file-backed matrices via 'bigstatsr' to handle large-scale genomic datasets efficiently. Key features include LD-based haplotype block construction, kernel ridge regression (KRR) for genomic prediction, and local genomic estimated breeding value (GEBV) analysis for identifying significant genomic regions.

**License** MIT

**Encoding** UTF-8

**LazyData** true

**Imports** R6, bigstatsr, data.table, Rcpp, methods, future, future.apply, bigsnpr, progressr, ggplot2

**Suggests** knitr, rmarkdown, testthat (>= 3.0.0)

**VignetteBuilder** knitr

**LinkingTo** Rcpp, RcppArmadillo

**RoxygenNote** 7.3.3

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encode_block_fast	<i>Encode Haplotypes (Optimized)</i>
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### Description

Maps unique rows of a genotype matrix to integers without string conversion.

### Usage

```
encode_block_fast(mat)
```

### Arguments

mat	A numeric matrix of genotypes (0, 1, 2).
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### Value

Integer vector of haplotype IDs.

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HaploObject	<i>HaploObject Class</i>
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### Description

HaploObject Class

HaploObject Class

### Public fields

geno Filebacked Big Matrix of genotypes.

map Data.table containing marker map information.

pheno Vector of phenotypes.

blocks Data.table defining haplotype blocks.

haplo\_mat Matrix of encoded haplotypes.

hrm Haplotype Relationship Matrix.

marker\_effects Vector of estimated marker effects.

local\_gebv List containing local GEBV matrix and variances.

significance Data.table of significance test results.

active\_markers Integer vector of indices of markers with non-zero variance.

## Methods

### Public methods:

- `HaploObject$new()`
- `HaploObject$print()`
- `HaploObject$import_genotypes()`
- `HaploObject$load_map()`
- `HaploObject$load_pheno()`
- `HaploObject$get_subset()`
- `HaploObject$define_blocks_fixed()`
- `HaploObject$define_blocks_ld()`
- `HaploObject$estimate_marker_effects()`
- `HaploObject$encode_haplotypes()`
- `HaploObject$calculate_local_gebv()`
- `HaploObject$test_significance()`
- `HaploObject$compute_hrm()`
- `HaploObject$fit_krr()`
- `HaploObject$cross_validate()`
- `HaploObject$plot_manhattan()`
- `HaploObject$plot_pca()`
- `HaploObject$plot_gebv_image()`
- `HaploObject$identify_superior_haplotypes()`
- `HaploObject$score_stacking()`
- `HaploObject$plot_stacking_trend()`
- `HaploObject$impute_genotypes()`
- `HaploObject$filter_monomorphic()`
- `HaploObject$save_project()`
- `HaploObject$clone()`

**Method** `new()`: Initialize a new HaploObject.

*Usage:*

```
HaploObject$new(backing_file_path)
```

*Arguments:*

`backing_file_path` Path to the backing file for the FBM.

**Method** `print()`: Print a summary of the HaploObject.

*Usage:*

```
HaploObject$print()
```

**Method** `import_genotypes()`: Import genotypes from a file or matrix.

*Usage:*

```
HaploObject$import_genotypes(matrix_or_path)
```

*Arguments:*

`matrix_or_path` A matrix or path to a file (bed/vcf/rds).

**Method** `load_map()`: Load marker map.

*Usage:*

```
HaploObject$load_map(map_data)
```

*Arguments:*

map\_data A data.frame or path to a file.

**Method** load\_pheno(): Load phenotypes.

*Usage:*

```
HaploObject$load_pheno(pheno_data)
```

*Arguments:*

pheno\_data A vector, data.frame, or path to a file.

**Method** get\_subset(): Get a subset of the genotype matrix.

*Usage:*

```
HaploObject$get_subset(row_ind, col_ind)
```

*Arguments:*

row\_ind Row indices.

col\_ind Column indices.

**Method** define\_blocks\_fixed(): Define haplotype blocks using fixed window size.

*Usage:*

```
HaploObject$define_blocks_fixed(window_size)
```

*Arguments:*

window\_size Number of markers per block.

**Method** define\_blocks\_ld(): Define haplotype blocks using LD scan.

*Usage:*

```
HaploObject$define_blocks_ld(  
  r2_threshold = 0.1,  
  tolerance = 3,  
  window_size = 2000  
)
```

*Arguments:*

r2\_threshold r2 threshold for block definition.

tolerance Number of failures allowed before ending a block.

window\_size Maximum window size to scan.

**Method** estimate\_marker\_effects(): Estimate marker effects using Ridge Regression.

*Usage:*

```
HaploObject$estimate_marker_effects(lambda = 1)
```

*Arguments:*

lambda Regularization parameter.

**Method** encode\_haplotypes(): Encode haplotypes into integer IDs.

*Usage:*

```
HaploObject$encode_haplotypes(n_cores = 1)
```

*Arguments:*

n\_cores Number of cores to use.

**Method** `calculate_local_gebv()`: Calculate local GEBVs.

*Usage:*

```
HaploObject$calculate_local_gebv(n_cores = 1)
```

*Arguments:*

`n_cores` Number of cores to use.

**Method** `test_significance()`: Test significance of local GEBVs.

*Usage:*

```
HaploObject$test_significance()
```

*Returns:* Data.table of p-values.

**Method** `compute_hrm()`: Compute Haplotype Relationship Matrix.

*Usage:*

```
HaploObject$compute_hrm(n_cores = 1)
```

*Arguments:*

`n_cores` Number of cores to use.

**Method** `fit_krr()`: Fit Kernel Ridge Regression model.

*Usage:*

```
HaploObject$fit_krr(lambda = 0.1, use_cg = NULL, tol = 1e-05, max_iter = 1000)
```

*Arguments:*

`lambda` Regularization parameter.

`use_cg` Whether to use Conjugate Gradient solver.

`tol` Tolerance for CG.

`max_iter` Maximum iterations for CG.

**Method** `cross_validate()`: Cross-validate KRR model.

*Usage:*

```
HaploObject$cross_validate(k = 5, lambdas = NULL, n_cores = 1, folds = NULL)
```

*Arguments:*

`k` Number of folds.

`lambdas` Vector of lambdas to test.

`n_cores` Number of cores to use.

`folds` Optional vector of fold assignments.

**Method** `plot_manhattan()`: Plot Manhattan plot of local GEBV significance.

*Usage:*

```
HaploObject$plot_manhattan(threshold = 0.05)
```

*Arguments:*

`threshold` Significance threshold (p-value).

**Method** `plot_pca()`: Plot PCA of HRM.

*Usage:*

```
HaploObject$plot_pca(groups = NULL)
```

*Arguments:*

`groups` Optional vector of groups for coloring.

**Method** `plot_gebv_image()`: Plot heatmap of local GEBVs.

*Usage:*

```
HaploObject$plot_gebv_image(block_range = NULL)
```

*Arguments:*

`block_range` Optional range of blocks to plot.

**Method** `identify_superior_haplotypes()`: Identify superior haplotypes.

*Usage:*

```
HaploObject$identify_superior_haplotypes(top_n = 50)
```

*Arguments:*

`top_n` Number of top blocks to consider.

**Method** `score_stacking()`: Calculate stacking scores.

*Usage:*

```
HaploObject$score_stacking(superior_haplos)
```

*Arguments:*

`superior_haplos` Data.table of superior haplotypes.

**Method** `plot_stacking_trend()`: Plot stacking trend.

*Usage:*

```
HaploObject$plot_stacking_trend(scores = NULL, superior_haplos = NULL)
```

*Arguments:*

`scores` Vector of stacking scores.

`superior_haplos` Optional table of superior haplotypes (if scores not provided).

**Method** `impute_genotypes()`: Impute missing genotypes.

*Usage:*

```
HaploObject$impute_genotypes(method = "mean")
```

*Arguments:*

`method` Imputation method. Currently only "mean" is supported.

**Method** `filter_monomorphic()`: Filter monomorphic markers (zero variance).

*Usage:*

```
HaploObject$filter_monomorphic()
```

**Method** `save_project()`: Save the project to an RDS file.

*Usage:*

```
HaploObject$save_project(path)
```

*Arguments:*

`path` Path to the output .rds file.

**Method** `clone()`: The objects of this class are cloneable with this method.

*Usage:*

```
HaploObject$clone(deep = FALSE)
```

*Arguments:*

`deep` Whether to make a deep clone.

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load_demo_data	<i>Load Demo Dataset</i>
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**Description**

Loads a pre-processed demo dataset for testing and visualization. The dataset contains 50 samples and 500 markers, with blocks defined, haplotypes encoded, and local GEBVs calculated.

**Usage**

```
load_demo_data()
```

**Value**

A HaploObject instance.

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load_haplo_project	<i>Load HaploGeno Project</i>
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**Description**

Load HaploGeno Project

**Usage**

```
load_haplo_project(path)
```

**Arguments**

path	Path to the .rds file.
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**Value**

A HaploObject instance.

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print.HaploObject	<i>Print method for HaploObject</i>
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**Description**

Print method for HaploObject

**Usage**

```
## S3 method for class 'HaploObject'  
print(x, ...)
```

**Arguments**

x	A HaploObject
...	Additional arguments

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summary.HaploObject	<i>Summary method for HaploObject</i>
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**Description**

Summary method for HaploObject

**Usage**

```
## S3 method for class 'HaploObject'  
summary(object, ...)
```

**Arguments**

object	A HaploObject
...	Additional arguments