# Package 'ohtadstats'

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Title Tomoka Ohta D Statistics
<b>Description</b> Calculate's Tomoka Ohta's partitioning of linkage disequilibrium, deemed D-statistics, for pairs of loci. Beissinger et al. (2016) <doi:10.1038 hdy.2015.81="">.</doi:10.1038>
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beissinger\_data

Chicken Genotype Data

## Description

This file is a matrix of genotypes from 96 chickens encompassing 5 breeds, genotyped as part of the Synbreed Project. Individuals are in rows. Marker genotypes are in columns, coded as 0, 1, and 2. Row names are a breed index so all rows named "1" are from breed 1, all rows named "2" are from breed 2, and so on. Column names are marker names. These data are a subset of the data used by Beissinger et al. (2016). The full dataset is hosted on Figshare at the which can be at the link below.

## Usage

```
data(beissinger_data)
```

#### **Format**

A matrix with 1417 rows and 100 columns.

### **Source**

(https://figshare.com/articles/Synbreed\_Biodiversity\_Panel\_Genotypes/1497961)

### References

Beissinger et al. (2016) Heredity. (https://www.nature.com/articles/hdy201581)

dheatmap

Heatmap Plot

### **Description**

Plots a matrix of D statistics, output from dwrapper, as a heatmap.

## Usage

```
dheatmap(d_matrix, colors = c("white", "lightblue", "blue", "darkblue",
   "black"), mode = "linear", tick.labels = TRUE, nbins = 5)
```

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## Arguments

d\_matrix A matrix of D statistics or a matrix of D statistic ratios.

colors An optional color vector. Optionally modify the color scheme of the heatmap.

If mode = 'binned', must be of length 5.

mode A string indicating desired coloring scheme. The option "linear" scales colors

linearly, "truncated" truncates values greater than 1, and "binned" returns a dis-

cretedistribution of colors.

tick.labels A logical indicating whether or not marker labels should be drawn.

nbins An integer specifying the number of bins to be used. Only relevent if mode is

"binned".

#### **Details**

The d\_matrix input should be one of the matrices output by dwrapper. Options are d2it\_mat, d2is\_mat, d2st\_mat, dp2is\_mat, npops\_mat, ratio1, and ratio2. More customized plots can be developed using the "levelplot" package.

## Value

A color plot

## **Examples**

```
data(miyashita_langley_data)
miyashita_langley_subset <- miyashita_langley_data[,1:15]
ml_results <- dwrapper(miyashita_langley_subset)
dheatmap(ml_results[["d2it_mat"]], mode = 'linear')

## Not run:
data(miyashita_langley_data)
ml_results <- dwrapper(miyashita_langley_data)
dheatmap(ml_results[["d2it_mat"]], mode = 'linear')

## End(Not run)</pre>
```

dparallel

Compute Ohta's D Statistics in a manner optimized for parallelization

## Description

Infers the comparisons that this instance of the function is supposed to perform given job\_id and comparisons\_per\_job. Returns the results of those comparisons to an SQL database.

#### Usage

```
dparallel(data_set, tot_maf = 0.1, pop_maf = 0.05, comparisons_per_job,
   job_id, outfile = "Ohta")
```

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## **Arguments**

data\_set The data set that is to be analysed.

tot\_maf Minimum minor allele frequency across the total population for a marker to be included in the analysis.

pop\_maf Minimum minor allele frequency across a subpopulation for that subpopulation to be included in analysis.

comparisons\_per\_job

The number of comparisons that each instance of dparallel will compute.

A number indicating that this is the nth instance of this function.

outfile Prefix for the file name that results will be written to. May be a path. Do not

include extension.

## **Examples**

job\_id

dstat

Tomoka Ohta's D Statistics

## **Description**

Implements Ohta's D statistics for a pair of loci. Statistics are returned in a vector in the following order: Number of populations, D2it, D2is, D2st, D'2st, D'2is.

## Usage

```
dstat(index, data_set, tot_maf = 0.1, pop_maf = 0.05)
```

## **Arguments**

index	A two-element vector of column names or numbers for which Ohta's D Statistics will be computed.
data_set	Matrix containing genotype data with individuals as rows and loci as columns. Genotypes should be coded as 0 (homozygous), 1 (heterozygous), or 2 (homozygous). Rownames must be subpopulation names and column names should be marker names.
tot_maf	Minimum minor allele frequency across the total population for a marker to be included in the analysis.
pop_maf	Minimum minor allele frequency across a subpopulation for that subpopulation to be included in analysis.

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### **Details**

When the loci being evaluated fail to pass the filtering thresholds determined by tot\_maf and pop\_maf, NAs are returned.

#### Value

nPops Number of subpopulations used for computation, after filtering.

D2it A measure of the correlation of alleles at two loci on the same gametes in a subpopulation relative to their expectation according to allele frequencies in the total population.

D2is Expected variance of LD for subpopulations.

D2st Expected correlation of alleles in a subpopulation relative to their expected correlation in the total population.

Dp2st Variance of LD for the total population computed over alleles only.

Dp2is Correlation of alleles at two loci on the same gamete in subpopulations relative to their expected correlation in the total population.

#### References

Beissinger et al. (2016) Heredity. (https://www.nature.com/articles/hdy201581) & Ohta. (1982) Proc. Natl. Acad. Science. (http://www.pnas.org/content/79/6/1940)

## **Examples**

```
data(beissinger_data)
dstat(index = c(5,6), data_set = beissinger_data)
```

dwrapper

Ohta D Statistic Wrapper

### **Description**

Pairwise computation of Ohta's D Statistics for each pair of polymorphisms in a given dataset.

## Usage

```
dwrapper(data_set, tot_maf = 0.1, pop_maf = 0.05)
```

### **Arguments**

data\_set Matrix containing genotype data with individuals as rows and loci as columns.

Genotypes should be coded as 0 (homozygous), 1 (heterozygous), or 2 (homozygous). Rownames must be subpopulation names and column names should be

marker names.

tot\_maf Minimum minor allele frequency across the total population for a marker to be

included in the analysis.

pop\_maf

Minimum minor allele frequency across a subpopulation for that subpopulation to be included in analysis.

#### **Details**

This wrapper implements the dstat function for all pairs of loci in a genotype matrix. If the input matrix includes n loci, choose(n,2) pairs are evaluated. Therefore, the computation time scales quadratically, and is not feasible for large datasets. We suggest manual parallelization across computational nodes for a large-scale (ie thousands of markers) implementation.

#### Value

A list of matrices containing the pairwise comparisons for each D statistic. Also included is the number of subpopulations evaluated in each comparison and the ratio of d2is\_mat to d2st\_mat (ratio1) and dp2st\_mat to dp2is\_mat (ratio2). The result of a comparison between marker M and marker N will be found in the Mth row at the Nth column.

## **Examples**

```
data(beissinger_data)
beissinger_subset <- beissinger_data[,1:15]
dwrapper(beissinger_subset, tot_maf = 0.05, pop_maf = 0.01)
## Not run:
data(beissinger_data)
dwrapper(beissinger_data, tot_maf = 0.05, pop_maf = 0.01)
## End(Not run)</pre>
```

miyashita\_langley\_data

Drosophila melanogaster genotypes

### **Description**

Genotype data obtained from Miyashita & Langley (1988). A matrix representing 85 loci in 64 individuals. Individuals are in rows. Rownames "RL", "TX", or "FK", indicate the subpopulation from which the sample was taken.

## Usage

```
data(miyashita_langley_data)
```

#### Format

A matrix with 64 rows and 85 columns.

# References

Miyashita & Langley (1988) Genetics 120:199-212 (https://www.ncbi.nlm.nih.gov/pmc/articles/PMC1203490/)

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