

TET.cpp

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C++ program for calling genotypes from tetraploid high-throughput sequencing data

This C++ program is for calling genotypes from nucleotide read quartets (read counts of A, C, G, and T) derived from individual high-throughput sequencing data for multiple tetraploid individuals from a population by a maximum-likelihood (ML) method. Although this program can be used to call genotypes from low-coverage sequencing data, high coverage is needed to call accurate genotypes with this program. At each site, the genotype and sequence-error rate for each individual are estimated by maximizing the likelihood of the observed data. To avoid analyzing false polymorphisms, number of alleles at the site is estimated based on significant genotypes with respect to the genotype homozygous for the most abundant nucleotide in the population sample. We use a likelihood-ratio test, which is expected to be asymptotically chi-square distributed with one degree of freedom, to examine the statistical significance.

Input file. The input file is a tab-delimited text file, consisting of the reference nucleotide and individual nucleotide read quartets at each position. The format is the same as that for our genotype-frequency estimator (Maruki and Lynch 2015), and can be made using our software package GFE (<https://github.com/Takahiro-Maruki/Package-GFE>). The first and second columns are the scaffold and position identifiers. The third column denotes the nucleotide of the reference sequence. Thereafter, the nucleotide read quartet for each individual is presented in each of the columns.

Output file. The output file is also a tab-delimited file. The meanings of the first eight columns are: 1) scaffold (chromosome) identifier; 2) site identifier (coordinate); 3) nucleotide of the reference sequence; 4) number of alleles; 5) depth of coverage in the population sample (sum of the coverage over the individuals); 6) number of called genotypes; 7) mean error-rate estimate among called genotypes; 8) nucleotide of the major allele. Next, the called genotypes are shown for each individual in each of the columns. The last four columns show the maximum values of the likelihood-ratio test statistics of called genotypes containing A, C, G, and T, respectively. These are expected to be asymptotically chi-square distributed with one degree of freedom.

Reference

If you use this program, please cite the following paper:

Maruki, T., and Lynch, M., (in press) Genotype calling from population-genomic sequencing data. *G3: Genes / Genomes / Genetics*.

Instructions

Below are specific procedures for using the program:

1. Make the input file using GFE (<https://github.com/Takahiro-Maruki/Package-GFE>).

2. Compile the program by typing the following command:

```
g++ -o TET TET.cpp -lm
```

3. Run the program using the following command:

```
./TET -in In_TET.txt -out Out_TET.txt
```

- In_TET.txt and Out_TET.txt are default names of the input and output files, respectively. The input and output file names can be specified by adding the '-in' and '-out' options, respectively.

- The chi-square critical value for the polymorphism test can be specified by adding the '-cv' option

- The minimum required coverage and allowed maximum coverage to call a genotype of an individual can be specified by adding the '-min_cov' and '-max_cov' options, respectively. Their default values are 1 and 2,000,000,000, respectively.

- The maximum allowed error rate per read per site can be specified by adding the '-max_e' option.

- Results only at significantly polymorphic sites can be outputted by adding the '-snpc' option.

- A usage help message explaining these options can be shown by typing the following command:

```
./TET -h
```

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Contact

If you have difficulty using this software, please send the following information to Takahiro Maruki (tmaruki@indiana.edu):

1. Brief explanation of the problem.
2. Command entered.
3. Part of the input file.
4. Part of the output file.