# **Resource Summary Report**

Generated by dkNET on Apr 16, 2025

# **BEAGLECALL**

RRID:SCR\_013301

Type: Tool

### **Proper Citation**

BEAGLECALL (RRID:SCR\_013301)

#### **Resource Information**

**URL:** http://faculty.washington.edu/browning/beaglecall/beaglecall.html

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**Description:** Software package for simultaneous genotype calling and haplotype phasing for unrelated individuals. BEAGLECALL produces output posterior genotype probabilities and output phased haplotypes. BEAGLECALL generates extremely accurate genotype calls because it uses both allele signal intensity data and inter-marker correlation to call genotypes. BEAGLECALL is designed for use with high-density SNP arrays, and it uses the BEAGLE haplotype frequency model to model inter-marker correlation. (entry from Genetic Analysis Software)

**Abbreviations:** BEAGLECALL

**Resource Type:** software resource, software application

Keywords: gene, genetic, genomic, java, ms-windows, unix, linux, macos

**Funding:** 

Resource Name: BEAGLECALL

Resource ID: SCR\_013301

Alternate IDs: nlx 154239

**Record Creation Time:** 20220129T080315+0000

Record Last Update: 20250416T063639+0000

## **Ratings and Alerts**

No rating or validation information has been found for BEAGLECALL.

No alerts have been found for BEAGLECALL.

### **Data and Source Information**

Source: SciCrunch Registry

## **Usage and Citation Metrics**

We found 1 mentions in open access literature.

**Listed below are recent publications.** The full list is available at dkNET.

Guo T, et al. (2017) Genome-Wide Association Study to Find Modifiers for Tetralogy of Fallot in the 22q11.2 Deletion Syndrome Identifies Variants in the GPR98 Locus on 5q14.3. Circulation. Cardiovascular genetics, 10(5), e001690.