## **Resource Summary Report**

Generated by dkNET on Apr 25, 2025

# **WTCHG Genome Scan Viewer**

RRID:SCR\_001635 Type: Tool

## **Proper Citation**

WTCHG Genome Scan Viewer (RRID:SCR\_001635)

### **Resource Information**

URL: http://mus.well.ox.ac.uk/gscandb/

Proper Citation: WTCHG Genome Scan Viewer (RRID:SCR\_001635)

**Description:** THIS RESOURCE IS NO LONGER IN SERVICE. Documented on September 23,2022. Database / display tool of genome scans, with a web interface that lets the user view the data. It does not perform any analyses - these must be done by other software, and the results uploaded into it. The basic features of GSCANDB are: \* Parallel viewing of scans for multiple phenotypes. \* Parallel analyses of the same scan data. \* Genome-wide views of genome scans \* Chromosomal region views, with zooming \* Gene and SNP Annotation is shown at high zoom levels \* Haplotype block structure viewing \* The positions of known Trait Loci can be overlayed and queried. \* Links to Ensembl, MGI, NCBI, UCSC and other genome data browsers. In GSCANDB, a genome scan has a wide definition, including not only the usual statistical genetic measures of association between genetic variation at a series of loci and variation in a phenotype, but any quantitative measure that varies along the genome. This includes for example competitive genome hybridization data and some kinds of gene expression measurements.

#### Abbreviations: GSCANDB

**Synonyms:** Wellcome Trust Centre for Human Genetics Genome Scan Viewer, Genome Scan Viewer, Genome Scan Database

Resource Type: database, service resource, data or information resource

**Keywords:** genome, gene, snp, trait, genotype, phenotype, visualization, region, chromosome, quantitative trait locus, hybridization, gene expression

Funding: NIAAA U01AA014425;

NCRR R24RR015116; NIGMS R01GM072863; NINDS R01NS049445; NIMH P20-MH 62009; NIAAA U24AA13513

Availability: THIS RESOURCE IS NO LONGER IN SERVICE

Resource Name: WTCHG Genome Scan Viewer

Resource ID: SCR\_001635

Alternate IDs: nlx\_153902

Record Creation Time: 20220129T080208+0000

Record Last Update: 20250425T055209+0000

## **Ratings and Alerts**

No rating or validation information has been found for WTCHG Genome Scan Viewer.

No alerts have been found for WTCHG Genome Scan Viewer.

## Data and Source Information

Source: SciCrunch Registry

## **Usage and Citation Metrics**

We found 3 mentions in open access literature.

Listed below are recent publications. The full list is available at <u>dkNET</u>.

Nicod J, et al. (2016) Genome-wide association of multiple complex traits in outbred mice by ultra-low-coverage sequencing. Nature genetics, 48(8), 912.

Vered K, et al. (2014) Susceptibility to Klebsiella pneumonaie infection in collaborative cross mice is a complex trait controlled by at least three loci acting at different time points. BMC genomics, 15(1), 865.

Krohn J, et al. (2014) Genetic interactions with sex make a relatively small contribution to the heritability of complex traits in mice. PloS one, 9(5), e96450.