Resource Summary Report

Generated by PRECISE-TBI on Sep 19, 2026

CGH Explorer

RRID:SCR_010920
Type: Tool

Proper Citation

CGH Explorer (RRID:SCR_010920)

Resource Information

URL: http://www.softgenetics.com/CGHExplorer.html

Proper Citation: CGH Explorer (RRID:SCR_010920)

Description: An easy-to-use software tool for analyzing two color copy number alteration arrays from multiple platforms, including Agilent Technologies, Illumina, AffyMetrix, NimbleGen and others.

Abbreviations: CGH Explorer

Resource Type: software resource

Availability: Commercially available

Resource Name: CGH Explorer

Resource ID: SCR_010920

Alternate IDs: OMICS_00708

Record Creation Time: 20220129T080301+0000

Record Last Update: 20260919T195202+0000

Ratings and Alerts

No rating or validation information has been found for CGH Explorer.

No alerts have been found for CGH Explorer.

Data and Source Information

Source: SciCrunch Registry

Usage and Citation Metrics

We found 2 mentions in open access literature.

Listed below are recent publications. The full list is available at PRECISE-TBI .

, et al. (2018) Assessment of established techniques to determine developmental and malignant potential of human pluripotent stem cells. Nature communications, 9(1), 1925.

Davidsson J, et al. (2008) Deletion of the SCN gene cluster on 2q24.4 is associated with severe epilepsy: an array-based genotype-phenotype correlation and a comprehensive review of previously published cases. Epilepsy research, 81(1), 69.