

# Resource Summary Report

Generated by NIF on Jul 29, 2026

## GT0001

RRID:CVCL\_8W53

Type: Cell Line

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### Proper Citation

ECACC Cat# 96011125, RRID:CVCL\_8W53

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### Cell Line Information

**URL:** [https://web.expasy.org/cellosaurus/CVCL\\_8W53](https://web.expasy.org/cellosaurus/CVCL_8W53)

**Proper Citation:** ECACC Cat# 96011125, RRID:CVCL\_8W53

**Description:** Cell line GT0001 is a Transformed cell line with a species of origin Homo sapiens (Human)

**Sex:** Male

**Disease:** Holoprosencephaly

**Comments:** Karyotypic information: 46,XY,inv(13)(q31.3;q32.3) (ECACC=96011125)., Population: Asian., Part of: ECACC chromosomal abnormality collection.

**Category:** Transformed cell line

**Organism:** Homo sapiens (Human)

**Name:** GT0001

**Cross References:** ECACC:96011125, Wikidata:Q54871822

**ID:** CVCL\_8W53

**Vendor:** ECACC

**Catalog Number:** 96011125

**Record Creation Time:** 20220427T220220+0000

**Record Last Update:** 20260725T233826+0000

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### Ratings and Alerts

No rating or validation information has been found for GT0001.

No alerts have been found for GT0001.

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## Data and Source Information

**Source:** [Cellosaurus](#)

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## Usage and Citation Metrics

We have not found any literature mentions for this resource.