

Resource Summary Report

Generated by [dkNET](#) on Sep 9, 2026

DD0102

RRID:CVCL_8U69

Type: Cell Line

Proper Citation

ECACC Cat# 89021601, RRID:CVCL_8U69

Cell Line Information

URL: https://web.expasy.org/cellosaurus/CVCL_8U69

Proper Citation: ECACC Cat# 89021601, RRID:CVCL_8U69

Description: Cell line DD0102 is a Finite cell line with a species of origin Homo sapiens (Human)

Sex: Male

Disease: Fragile X syndrome

Comments: Part of: ECACC chromosomal abnormality collection.

Category: Finite cell line

Organism: Homo sapiens (Human)

Name: DD0102

Cross References: ECACC:89021601, Wikidata:Q54828755

ID: CVCL_8U69

Vendor: ECACC

Catalog Number: 89021601

Record Creation Time: 20220427T215747+0000

Record Last Update: 20260905T175129+0000

Ratings and Alerts

No rating or validation information has been found for DD0102.

No alerts have been found for DD0102.

Data and Source Information

Source: [CelloSaurus](#)

Usage and Citation Metrics

We have not found any literature mentions for this resource.