Resource Summary Report

Generated by dkNET on Sep 16, 2026

GM05131

RRID:CVCL_AX78
Type: Cell Line

Proper Citation

Coriell Cat# GM05131, RRID:CVCL_AX78

Cell Line Information

URL: https://web.expasy.org/cellosaurus/CVCL_AX78

Proper Citation: Coriell Cat# GM05131, RRID:CVCL_AX78

Description: Cell line GM05131 is a Finite cell line with a species of origin Homo sapiens (Human)

Sex: Male

Disease: Fragile X syndrome

Comments: Population: Caucasian.

Category: Finite cell line

Organism: Homo sapiens (Human)

Name: GM05131

Cross References: CLO:CLO_0025352, Coriell:GM05131, Wikidata:Q54838877

ID: CVCL_AX78

Vendor: Coriell

Catalog Number: GM05131

Record Creation Time: 20220427T215939+0000

Record Last Update: 20260913T003248+0000

Ratings and Alerts

No rating or validation information has been found for GM05131.

No alerts have been found for GM05131.

Data and Source Information

Source: Cellosaurus

Usage and Citation Metrics

We found 1 mentions in open access literature.

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

Susco SG, et al. (2022) Molecular convergence between Down syndrome and fragile X syndrome identified using human pluripotent stem cell models. Cell reports, 40(10), 111312.