Resource Summary Report

Generated by RRID on Sep 6, 2026

GM04026

RRID:CVCL_AX77
Type: Cell Line

Proper Citation

RRID:CVCL_AX77

Cell Line Information

URL: https://web.expasy.org/cellosaurus/CVCL_AX77

Proper Citation: RRID:CVCL_AX77

Description: Cell line GM04026 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: GM04026

Cross References: CLO:CLO_0016230, Coriell:GM04026, Wikidata:Q54838375

ID: CVCL_AX77

Record Creation Time: 20220427T215933+0000

Record Last Update: 20260905T175459+0000

Ratings and Alerts

No rating or validation information has been found for GM04026.

No alerts have been found for GM04026.

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 RRID .

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.