Resource Summary Report

Generated by NIF on Sep 19, 2026

GM01981

RRID:CVCL_F589
Type: Cell Line

Proper Citation

RRID:CVCL_F589

Cell Line Information

URL: https://web.expasy.org/cellosaurus/CVCL_F589

Proper Citation: RRID:CVCL_F589

Description: Cell line GM01981 is a Finite cell line with a species of origin Homo sapiens (Human)

Sex: Male

Disease: Menkes disease

Defining Citation: PMID:7438975, PMID:7977350

Comments: Population: Caucasian.

Category: Finite cell line

Organism: Homo sapiens (Human)

Name: GM01981

Synonyms: GM-1981, GM1981

Cross References: CLO:CLO_0032339, BioSample:SAMN00807346, Coriell:GM01981, Wikidata:Q54837212

ID: CVCL_F589

Record Creation Time: 20220427T215919+0000

Record Last Update: 20260913T003207+0000

Ratings and Alerts

No rating or validation information has been found for GM01981.

No alerts have been found for GM01981.

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 NIF .

Zlatic SA, et al. (2018) Rare Disease Mechanisms Identified by Genealogical Proteomics of Copper Homeostasis Mutant Pedigrees. Cell systems, 6(3), 368.