Resource Summary Report

Generated by NIF on Sep 21, 2026

GM01057

RRID:CVCL_H964
Type: Cell Line

Proper Citation

Coriell Cat# GM01057, RRID:CVCL_H964

Cell Line Information

URL: https://web.expasy.org/cellosaurus/CVCL_H964

Proper Citation: Coriell Cat# GM01057, RRID:CVCL_H964

Description: Cell line GM01057 is a Finite cell line with a species of origin Homo sapiens (Human)

Sex: Male

Disease: Menkes disease

Defining Citation: PMID:7438975

Comments: Population: Caucasian.

Category: Finite cell line

Organism: Homo sapiens (Human)

Name: GM01057

Synonyms: GM-1057

Cross References: CLO:CLO_0030350, BioSample:SAMN00803596, Coriell:GM01057, Wikidata:Q54836624

ID: CVCL_H964

Vendor: Coriell

Catalog Number: GM01057

Record Creation Time: 20220427T215911+0000

Record Last Update: 20260920T002659+0000

Ratings and Alerts

No rating or validation information has been found for GM01057.

No alerts have been found for GM01057.

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.