Resource Summary Report

Generated by RRID on Sep 7, 2026

GM16825

RRID:CVCL_U890
Type: Cell Line

Proper Citation

RRID:CVCL_U890

Cell Line Information

URL: https://web.expasy.org/cellosaurus/CVCL_U890

Proper Citation: RRID:CVCL_U890

Description: Cell line GM16825 is a Finite cell line with a species of origin Homo sapiens (Human)

Sex: Male

Disease: Alexander disease

Comments: Population: Caucasian.

Category: Finite cell line

Organism: Homo sapiens (Human)

Name: GM16825

Synonyms: ALX-2

Cross References: CLO:CLO_0018411, Coriell:GM16825, Wikidata:Q54848761

ID: CVCL_U890

Record Creation Time: 20220427T220109+0000

Record Last Update: 20260905T175805+0000

Ratings and Alerts

No rating or validation information has been found for GM16825.

No alerts have been found for GM16825.

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 .

Li L, et al. (2018) GFAP Mutations in Astrocytes Impair Oligodendrocyte Progenitor Proliferation and Myelination in an hiPSC Model of Alexander Disease. Cell stem cell, 23(2), 239.