

Resource Summary Report

Generated by [dkNET](#) on Aug 21, 2026

GM00097

RRID:CVCL_V747

Type: Cell Line

Proper Citation

Coriell Cat# GM00097, RRID:CVCL_V747

Cell Line Information

URL: https://web.expasy.org/cellosaurus/CVCL_V747

Proper Citation: Coriell Cat# GM00097, RRID:CVCL_V747

Description: Cell line GM00097 is a Finite cell line with a species of origin Homo sapiens (Human)

Sex: Female

Disease: Simpson Golabi Behmel syndrome type 1

Defining Citation: [PMID:1679663](#), [PMID:2498246](#), [PMID:4139001](#), [PMID:10377420](#)

Comments: Karyotypic information: 46,X,t(X;1)(q26;q21) (PubMed=10377420)., Population: Caucasian.

Category: Finite cell line

Organism: Homo sapiens (Human)

Name: GM00097

Synonyms: GM-97, GM0097, GM 0097, GM00097A

Cross References: CLO:CLO_0025819, Coriell:GM00097, Wikidata:Q54836039

ID: CVCL_V747

Vendor: Coriell

Catalog Number: GM00097

Record Creation Time: 20220427T215904+0000

Record Last Update: 20260815T232844+0000

Ratings and Alerts

No rating or validation information has been found for GM00097.

No alerts have been found for GM00097.

Data and Source Information

Source: [CelloSaurus](#)

Usage and Citation Metrics

We have not found any literature mentions for this resource.