

Resource Summary Report

Generated by [dkNET](#) on Aug 7, 2026

GM10742

RRID:CVCL_Y652

Type: Cell Line

Proper Citation

RRID:CVCL_Y652

Cell Line Information

URL: https://web.expasy.org/cellosaurus/CVCL_Y652

Proper Citation: RRID:CVCL_Y652

Description: Cell line GM10742 is a Transformed cell line with a species of origin Homo sapiens (Human)

Sex: Male

Disease: Leber hereditary optic atrophy

Defining Citation: [PMID:16120335](#)

Category: Transformed cell line

Organism: Homo sapiens (Human)

Name: GM10742

Synonyms: GM10742A

Cross References: CLO:CLO_0023698, BioSample:SAMN00800184, Coriell:GM10742, Wikidata:Q54844630

ID: CVCL_Y652

Record Creation Time: 20220427T220018+0000

Record Last Update: 20260802T000609+0000

Ratings and Alerts

No rating or validation information has been found for GM10742.

No alerts have been found for GM10742.

Data and Source Information

Source: [CelloSaurus](#)

Usage and Citation Metrics

We have not found any literature mentions for this resource.