

Resource Summary Report

Generated by [NIF](#) on Sep 23, 2026

GM06902

RRID:CVCL_Y703

Type: Cell Line

Proper Citation

RRID:CVCL_Y703

Cell Line Information

URL: https://web.expasy.org/cellosaurus/CVCL_Y703

Proper Citation: RRID:CVCL_Y703

Description: Cell line GM06902 is a Finite cell line with a species of origin Homo sapiens (Human)

Sex: Male

Disease: Ornithine carbamoyltransferase deficiency disease

Comments: Population: Caucasian.

Category: Finite cell line

Organism: Homo sapiens (Human)

Name: GM06902

Synonyms: GM6902

Cross References: CLO:CLO_0036484, Coriell:GM06902, Wikidata:Q54842378

ID: CVCL_Y703

Record Creation Time: 20220427T215950+0000

Record Last Update: 20260920T002813+0000

Ratings and Alerts

No rating or validation information has been found for GM06902.

No alerts have been found for GM06902.

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](#) .

Lee JS, et al. (2018) Urea Cycle Dysregulation Generates Clinically Relevant Genomic and Biochemical Signatures. Cell, 174(6), 1559.