

# Resource Summary Report

Generated by [NIF](#) on Sep 20, 2026

## GM06902

RRID:CVCL\_Y703

Type: Cell Line

---

### Proper Citation

RRID:CVCL\_Y703

---

### Cell Line Information

**URL:** [https://web.expasy.org/cellosaurus/CVCL\\_Y703](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.