

Resource Summary Report

Generated by [RRID](#) on Sep 7, 2026

HSC 99

RRID:CVCL_G050

Type: Cell Line

Proper Citation

RRID:CVCL_G050

Cell Line Information

URL: https://web.expasy.org/cellosaurus/CVCL_G050

Proper Citation: RRID:CVCL_G050

Description: Cell line HSC 99 is a Transformed cell line with a species of origin Homo sapiens (Human)

Sex: Female

Disease: Fanconi anemia, complementation group A

Defining Citation: [PMID:2805228](#), [PMID:6809308](#), [PMID:7987813](#), [PMID:7994019](#), [PMID:8340107](#)

Category: Transformed cell line

Organism: Homo sapiens (Human)

Name: HSC 99

Synonyms: HSC-99, HSC99

Cross References: Wikidata:Q54896378

ID: CVCL_G050

Record Creation Time: 20220427T220615+0000

Record Last Update: 20260905T180806+0000

Ratings and Alerts

No rating or validation information has been found for HSC 99.

No alerts have been found for HSC 99.

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](#) .

Oppezzo A, et al. (2020) Microphthalmia transcription factor expression contributes to bone marrow failure in Fanconi anemia. The Journal of clinical investigation, 130(3), 1377.