

Resource Summary Report

Generated by [RRID](#) on Jul 30, 2026

WI-L2-729HF2

RRID:CVCL_6741

Type: Cell Line

Proper Citation

RRID:CVCL_6741

Cell Line Information

URL: https://web.expasy.org/cellosaurus/CVCL_6741

Proper Citation: RRID:CVCL_6741

Description: Cell line WI-L2-729HF2 is a Transformed cell line with a species of origin Homo sapiens (Human)

Sex: Male

Disease: Hereditary spherocytosis

Defining Citation: [PMID:3101287](#), [PMID:3365703](#), [PMID:6350451](#), [PMID:6699402](#)

Comments: Anecdotal: Because this cell line was deposited at ATCC on April 2, 1981 and the patent describing it was filed one week earlier (March, 26), the patent was first rejected. This decision was later reversed in a ruling that changed the deposition rules (DOI=10.1038/nbt1185-971)., Population: Caucasian., Group: Patented cell line., Group: Hybridoma fusion partner cell line.

Category: Transformed cell line

Organism: Homo sapiens (Human)

Name: WI-L2-729HF2

Synonyms: WI-L2-729-HF2, WI-L2-729 HF2, 729-HF2, UC729-HF2, UC729HF2, HF2

Cross References: CLO:CLO_0009603, ATCC:CRL-8062, Wikidata:Q54994037

ID: CVCL_6741

Hierarchy: cvcl_d039

Record Creation Time: 20220427T221720+0000

Ratings and Alerts

No rating or validation information has been found for WI-L2-729HF2.

Warning: Discontinued: ATCC; CRL-8062

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Data and Source Information

Source: [Cellosaurus](#)

Usage and Citation Metrics

We have not found any literature mentions for this resource.