

Resource Summary Report

Generated by [RRID](#) on Sep 6, 2026

[AG06922](#)

RRID:CVCL_X793

Type: Cell Line

Proper Citation

Coriell Cat# AG06922, RRID:CVCL_X793

Cell Line Information

URL: https://web.expasy.org/cellosaurus/CVCL_X793

Proper Citation: Coriell Cat# AG06922, RRID:CVCL_X793

Description: Cell line AG06922 is a Finite cell line with a species of origin Homo sapiens (Human)

Sex: Male

Disease: Down syndrome

Defining Citation: [PMID:17668376](#)

Comments: Population: Caucasian.

Category: Finite cell line

Organism: Homo sapiens (Human)

Name: AG06922

Synonyms: AG06922A

Cross References: CLO:CLO_0035478, Coriell:AG06922, Wikidata:Q54740453

ID: CVCL_X793

Vendor: Coriell

Catalog Number: AG06922

Record Creation Time: 20220427T215136+0000

Record Last Update: 20260905T184457+0000

Ratings and Alerts

No rating or validation information has been found for AG06922.

No alerts have been found for AG06922.

Data and Source Information

Source: [CelloSaurus](#)

Usage and Citation Metrics

We found 2 mentions in open access literature.

Listed below are recent publications. The full list is available at [RRID](#) .

Hwang S, et al. (2019) Suppressing Aneuploidy-Associated Phenotypes Improves the Fitness of Trisomy 21 Cells. Cell reports, 29(8), 2473.

Galati DF, et al. (2018) Trisomy 21 Represses Cilia Formation and Function. Developmental cell, 46(5), 641.