

Resource Summary Report

Generated by [RRID](#) on Sep 3, 2026

F35T

RRID:CVCL_E2X2

Type: Cell Line

Proper Citation

RRID:CVCL_E2X2

Cell Line Information

URL: https://web.expasy.org/cellosaurus/CVCL_E2X2

Proper Citation: RRID:CVCL_E2X2

Description: Cell line F35T is a Transformed cell line with a species of origin Homo sapiens (Human)

Sex: Female

Disease: Fuchs endothelial dystrophy

Defining Citation: [PMID:33157036](#), [PMID:38024683](#)

Comments: Caution: PubMed=38024683 and Patent=US11512312 erroneously indicate the number of CTG18.1 repeats to be ~1500., Miscellaneous: STR profile and donor ethnicity from personal communication of Zarouchlioti, Christina., Donor information: Established from a patient undergoing corneal transplant surgery (PubMed=33157036)., Population: Caucasian.

Category: Transformed cell line

Organism: Homo sapiens (Human)

Name: F35T

ID: CVCL_E2X2

Record Creation Time: 20240911T070903+0000

Record Last Update: 20260904T020412+0000

Ratings and Alerts

No rating or validation information has been found for F35T.

No alerts have been found for F35T.

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](#) .

Zarouchlioti C, et al. (2024) Tissue-specific TCF4 triplet repeat instability revealed by optical genome mapping. EBioMedicine, 108, 105328.