# Resource Summary Report

Generated by RRID on Sep 7, 2026

## GM16756

RRID:CVCL_G041
Type: Cell Line

### Proper Citation

RRID:CVCL_G041

### Cell Line Information

**URL:** https://web.expasy.org/cellosaurus/CVCL_G041

**Proper Citation:** RRID:CVCL_G041

**Description:** Cell line GM16756 is a Transformed cell line with a species of origin Homo sapiens (Human)

**Sex:** Male

**Disease:** Fanconi anemia, complementation group D2

**Defining Citation:** PMID:12361951, PMID:16166284

**Category:** Transformed cell line

**Organism:** Homo sapiens (Human)

**Name:** GM16756

**Synonyms:** PD20.L, PD.20i, PD20

**Cross References:** CLO:CLO_0018426, Coriell:GM16756, Wikidata:Q54848727

**ID:** CVCL_G041

**Record Creation Time:** 20220427T220108+0000

**Record Last Update:** 20260905T175804+0000

### Ratings and Alerts

No rating or validation information has been found for GM16756.

No alerts have been found for GM16756.

## 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 .

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.

Fouquet B, et al. (2017) A homozygous FANCM mutation underlies a familial case of non-syndromic primary ovarian insufficiency. eLife, 6.