

Resource Summary Report

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HEK293T/17

RRID:CVCL_1926

Type: Cell Line

Proper Citation

(CLS Cat# 305117, RRID:CVCL_1926)

Cell Line Information

URL: https://web.expasy.org/cellosaurus/CVCL_1926

Proper Citation: (CLS Cat# 305117, RRID:CVCL_1926)

Sex: Female

Defining Citation: [PMID:7690960](https://pubmed.ncbi.nlm.nih.gov/7690960/)

Comments: Group: Patented cell line.

Category: Transformed cell line

Name: HEK293T/17

Synonyms: HEK-293T/17, HEK 293T/17, 293T/17

Cross References: BTO:BTO_0006328, CLO:CLO_0001235, ATCC:CRL-11268, BioSample:SAMN03471332, CCRID:1101HUM-PUMC000212, CCRID:3101HUMGNHu44, CLS:305117, TOKU-E:245, Wikidata:Q28178058

ID: CVCL_1926

Vendor: CLS

Catalog Number: 305117

Record Creation Time: 20250131T200355+0000

Record Last Update: 20250131T201610+0000

Ratings and Alerts

No rating or validation information has been found for HEK293T/17.

No alerts have been found for HEK293T/17.

Data and Source Information

Source: [Cellosaurus](#)

Usage and Citation Metrics

We found 1460 mentions in open access literature.

Listed below are recent publications. The full list is available at [FDI Lab - SciCrunch.org](#).

Alexander KA, et al. (2025) Nuclear speckles regulate functional programs in cancer. *Nature cell biology*, 27(2), 322.

Chakrabarty Y, et al. (2024) The HRI branch of the integrated stress response selectively triggers mitophagy. *Molecular cell*, 84(6), 1090.

Su C, et al. (2024) Vascular injury activates the ELK1/SND1/SRF pathway to promote vascular smooth muscle cell proliferative phenotype and neointimal hyperplasia. *Cellular and molecular life sciences : CMLS*, 81(1), 59.

Cai SW, et al. (2024) POT1 recruits and regulates CST-Pol β /primase at human telomeres. *Cell*, 187(14), 3638.

Wang JY, et al. (2024) PolyQ-expanded ataxin-2 aggregation impairs cellular processing-body homeostasis via sequestering the RNA helicase DDX6. *The Journal of biological chemistry*, 300(7), 107413.

Momenilandi M, et al. (2024) FLT3L governs the development of partially overlapping hematopoietic lineages in humans and mice. *Cell*, 187(11), 2817.

Hirsch T, et al. (2024) IRF4 impedes human CD8 T cell function and promotes cell proliferation and PD-1 expression. *Cell reports*, 43(7), 114401.

Zhang QE, et al. (2024) SARS-CoV-2 Omicron XBB lineage spike structures, conformations, antigenicity, and receptor recognition. *Molecular cell*, 84(14), 2747.

Li Y, et al. (2024) Zinc transporter 1 functions in copper uptake and cuproptosis. *Cell metabolism*, 36(9), 2118.

Muik A, et al. (2024) Immunity against conserved epitopes dominates after two consecutive exposures to SARS-CoV-2 Omicron BA.1. *Cell reports*, 43(8), 114567.

Saxena S, et al. (2024) Unprocessed genomic uracil as a source of DNA replication stress in cancer cells. *Molecular cell*, 84(11), 2036.

Liu Z, et al. (2024) Neutralization of SARS-CoV-2 BA.2.86 and JN.1 by CF501 adjuvant-enhanced immune responses targeting the conserved epitopes in ancestral RBD. *Cell reports. Medicine*, 5(3), 101445.

Ma X, et al. (2024) A programmable targeted protein-degradation platform for versatile applications in mammalian cells and mice. *Molecular cell*.

Ke YD, et al. (2024) Targeting 14-3-3 σ -mediated TDP-43 pathology in amyotrophic lateral sclerosis and frontotemporal dementia mice. *Neuron*.

Wu M, et al. (2024) Bi-directional regulation of type I interferon signaling by heme oxygenase-1. *iScience*, 27(3), 109185.

Lu Y, et al. (2024) HDAC5 enhances IRF3 activation and is targeted for degradation by protein C6 from orthopoxviruses including Monkeypox virus and Variola virus. *Cell reports*, 43(3), 113788.

Yu T, et al. (2024) NLRP3 Cys126 palmitoylation by ZDHHC7 promotes inflammasome activation. *Cell reports*, 43(4), 114070.

Yang L, et al. (2024) SARS-CoV-2 infection causes dopaminergic neuron senescence. *Cell stem cell*, 31(2), 196.

Zhang T, et al. (2024) Structure-guided development of selective caseinolytic protease P agonists as antistaphylococcal agents. *Cell reports. Medicine*, 5(12), 101837.

Johnson K, et al. (2024) Small molecule telomerase inhibitors are also potent inhibitors of telomeric C-strand synthesis. *RNA (New York, N.Y.)*, 30(9), 1213.