

# Resource Summary Report

Generated by [FDI Lab - SciCrunch.org](https://fdi-lab.org) on Mar 29, 2025

## Caco-2

RRID:CVCL\_0025

Type: Cell Line

### Proper Citation

(RRID:CVCL\_0025)

### Cell Line Information

**URL:** [https://web.expasy.org/cellosaurus/CVCL\\_0025](https://web.expasy.org/cellosaurus/CVCL_0025)

**Proper Citation:** (RRID:CVCL\_0025)

**Sex:** Male

**Defining Citation:** [PMID:327080](#), [PMID:833871](#), [PMID:2914637](#), [PMID:3349466](#), [PMID:3518877](#), [PMID:6935474](#), [PMID:7459858](#), [PMID:7764660](#), [PMID:8253353](#), [PMID:8508948](#), [PMID:9294210](#), [PMID:10092214](#), [PMID:10612807](#), [PMID:10737795](#), [PMID:11414198](#), [PMID:11416159](#), [PMID:11668190](#), [PMID:12584437](#), [PMID:14599474](#), [PMID:15316659](#), [PMID:15731278](#), [PMID:15868485](#), [PMID:16418264](#), [PMID:16854228](#), [PMID:18258742](#), [PMID:20570890](#), [PMID:20606684](#), [PMID:20831567](#), [PMID:21607810](#), [PMID:23272949](#), [PMID:23932154](#), [PMID:24042735](#), [PMID:24755471](#), [PMID:25485619](#), [PMID:25841592](#), [PMID:25877200](#), [PMID:25926053](#), [PMID:25944804](#), [PMID:25960936](#), [PMID:26537799](#), [PMID:26589293](#), [PMID:26869432](#), [PMID:28196595](#), [PMID:28683746](#), [PMID:29101300](#), [PMID:29787057](#), [PMID:30894373](#), [PMID:31068700](#), [PMID:31981602](#), [PMID:33389257](#), [PMID:34339474](#)

**Comments:** Omics: Transcriptome analysis by serial analysis of gene expression (SAGE)., Omics: Transcriptome analysis by RNAseq., Omics: Transcriptome analysis by microarray., Omics: SNP array analysis., Omics: Protein expression by reverse-phase protein arrays., Omics: N-glycan profiling., Omics: miRNA expression profiling., Omics: Deep proteome analysis., Omics: Deep phosphoproteome analysis., Omics: Deep exome analysis., Omics: Deep antibody staining analysis., Omics: H3K4me3 ChIP-seq epigenome analysis., Omics: H3K36me3 ChIP-seq epigenome analysis., Omics: H3K27me3 ChIP-seq epigenome analysis., Omics: CTCF ChIP-seq epigenome analysis., Virology: Susceptible to infection by SARS coronavirus 2 (SARS-CoV-2) (COVID-19) (PubMed=33389257; PubMed=34339474)., Virology: Highly susceptible to infection by SARS coronavirus (SARS-CoV) (PubMed=15316659; PubMed=15731278)., Population: Caucasian., Part of: MD Anderson Cell Lines Project., Part of: ENCODE project common cell types; tier 3., Part of: Cancer Dependency Map project (DepMap) (includes Cancer Cell Line Encyclopedia - CCLE)., Part

of: AstraZeneca Colorectal cell line (AZCL) panel.

**Category:** Cancer cell line

**Name:** Caco-2

**Synonyms:** CaCo-2, CACO-2, Caco 2, CACO 2, CACO2, CaCo2, CaCO2, Caco2, Caco-2/ATCC, Caco-II

**Cross References:** BTO:BTO\_0000195, CLO:CLO\_0002172, CLO:CLO\_0050627, EFO:EFO\_0001099, MCCL:MCC:0000120, CLDB:cl614, CLDB:cl615, CLDB:cl616, CLDB:cl617, CLDB:cl618, CLDB:cl619, CLDB:cl621, CLDB:cl5174, Abcam:ab275464, AddexBio:C0009009/29, ArrayExpress:E-MTAB-2706, ATCC:HTB-37, BCRJ:0059, BioSample:SAMN03473140, BioSample:SAMN05292461, BioSample:SAMN10989600, cancercellines:CVCL\_0025, CCRID:1101HUM-PUMC000100, CCRID:1102HUM-NIFDC00087, CCRID:3101HUMSCSP5027, CCRID:3101HUMTCHu146, CCRID:4201HUM-CCTCC00153, CCRID:5301HUM-KCB07010YJ, CCTCC:GDC0153, Cell\_Model\_Passport:SIDM00891, ChEMBL-Cells:ChEMBL3307519, ChEMBL-Targets:ChEMBL614058, CLS:300137, ColonAtlas:CACO2, Cosmic:720340, Cosmic:873696, Cosmic:875297, Cosmic:876709, Cosmic:887244, Cosmic:889531, Cosmic:948127, Cosmic:948829, Cosmic:983736, Cosmic:995392, Cosmic:1043805, Cosmic:1122320, Cosmic:1132564, Cosmic:1132694, Cosmic:1184081, Cosmic:1187299, Cosmic:1310931, Cosmic:1466811, Cosmic:1479620, Cosmic:1524334, Cosmic:1552180, Cosmic:1676742, Cosmic:1708415, Cosmic:1803940, Cosmic:1933015, Cosmic:2036652, Cosmic:2145574, Cosmic:2156940, Cosmic:2301543, Cosmic:2301964, Cosmic:2667879, Cosmic:2800568, DepMap:ACH-000003, DSMZ:ACC-169, DSMZCellDive:ACC-169, ECACC:09042001, ECACC:86010202, EGA:EGAS00001000610, EGA:EGAS00001002554, ENCODE:ENCBS311PHE, ENCODE:ENCBS390JMI, ENCODE:ENCBS391ENC, ENCODE:ENCBS530YXL, ENCODE:ENCBS700RKG, ENCODE:ENCBS890JZY, FCS-free:192-2-378-1-3-3, GEO:GSM206450, GEO:GSM274711, GEO:GSM274712, GEO:GSM274725, GEO:GSM383861, GEO:GSM472900, GEO:GSM472933, GEO:GSM513908, GEO:GSM514182, GEO:GSM741244, GEO:GSM749689, GEO:GSM749691, GEO:GSM749748, GEO:GSM784006, GEO:GSM845394, GEO:GSM843481, GEO:GSM843482, GEO:GSM945162, GEO:GSM945203, GEO:GSM945206, GEO:GSM945236, GEO:GSM1006210, GEO:GSM1006211, GEO:GSM1006212, GEO:GSM1346866, GEO:GSM1374426, GEO:GSM1448160, GEO:GSM2549989, IARC\_TP53:21749, IBRC:C10094, ICLC:HTL97023, IZSLER:BS TCL 87, KCB:KCB 200710YJ, KCLB:30037.1, LINCS\_LDP:LCL-1170, Lonza:36, MeSH:D018938, MetaboLights:MTBLS227, MetaboLights:MTBLS328, NCBI\_Iran:C139, PharmacDB:Caco2\_161\_2019, PRIDE:PXD001550, PRIDE:PXD005354, PRIDE:PXD005355, PRIDE:PXD018357, PRIDE:PXD019645, PRIDE:PXD023760, Progenetix:CVCL\_0025, PubChem\_Cell\_line:CVCL\_0025, RCB:RCB0988, TOKU-E:762, Ubigen:YC-D003, Wikidata:Q5016050

**ID:** CVCL\_0025

**Record Creation Time:** 20250131T194636+0000

**Record Last Update:** 20250131T195142+0000

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## Ratings and Alerts

No rating or validation information has been found for Caco-2.

No alerts have been found for Caco-2.

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## Data and Source Information

**Source:** [Cellosaurus](#)

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## Usage and Citation Metrics

We found 9054 mentions in open access literature.

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

Mottawea W, et al. (2025) Multi-level analysis of gut microbiome extracellular vesicles-host interaction reveals a connection to gut-brain axis signaling. *Microbiology spectrum*, 13(2), e0136824.

Slaninová V, et al. (2024) The Hippo pathway terminal effector TAZ/WWTR1 mediates oxaliplatin sensitivity in p53 proficient colon cancer cells. *BMC cancer*, 24(1), 587.

Shin Y, et al. (2024) MMP-9-dependent proteolysis of the histone H3 N-terminal tail: a critical epigenetic step in driving oncogenic transcription and colon tumorigenesis. *Molecular oncology*, 18(8), 2001.

Bai Y, et al. (2024) Shaping immune landscape of colorectal cancer by cholesterol metabolites. *EMBO molecular medicine*, 16(2), 334.

Tian S, et al. (2024) Design, performance, processing, and validation of a pooled CRISPR perturbation screen for bacterial toxins. *Nature protocols*.

Zhang L, et al. (2024) ACE2-independent sarbecovirus cell entry can be supported by TMPRSS2-related enzymes and can reduce sensitivity to antibody-mediated neutralization. *PLoS pathogens*, 20(11), e1012653.

Mitrofanova O, et al. (2024) Bioengineered human colon organoids with in vivo-like cellular complexity and function. *Cell stem cell*, 31(8), 1175.

Dong B, et al. (2024) NK Receptor Signaling Lowers TCR Activation Threshold, Enhancing Selective Recognition of Cancer Cells by TAA-Specific CTLs. *Cancer immunology research*, 12(10), 1421.

Yang Q, et al. (2024) Reduced representative methylome profiling of cell-free DNA for breast cancer detection. *Clinical epigenetics*, 16(1), 33.

Mao L, et al. (2024) Olgotrelvir, a dual inhibitor of SARS-CoV-2 Mpro and cathepsin L, as a standalone antiviral oral intervention candidate for COVID-19. *Med (New York, N.Y.)*, 5(1), 42.

Nanlohy NM, et al. (2024) Exploring host-commensal-pathogen dynamics in cell line and organotypic human intestinal epithelial models. *iScience*, 27(5), 109771.

Meng C, et al. (2024) Chlorogenic acid regulates the expression of NPC1L1 and HMGCR through PXR and SREBP2 signaling pathways and their interactions with HSP90 to maintain cholesterol homeostasis. *Phytomedicine : international journal of phytotherapy and phytopharmacology*, 123, 155271.

Pearson GJ, et al. (2024) ER-export and ARFRP1/AP-1-dependent delivery of SARS-CoV-2 Envelope to lysosomes controls late stages of viral replication. *Science advances*, 10(14), ead15012.

Liu C, et al. (2024) HuR promotes triglyceride synthesis and intestinal fat absorption. *Cell reports*, 43(5), 114238.

Carregari VC, et al. (2024) Diving into the proteomic atlas of SARS-CoV-2 infected cells. *Scientific reports*, 14(1), 7375.

Zhang L, et al. (2024) SARS-CoV-2 BA.2.86 enters lung cells and evades neutralizing antibodies with high efficiency. *Cell*, 187(3), 596.

Huang J, et al. (2024) A Gene-Switch Platform Interfacing with Reactive Oxygen Species Enables Transcription Fine-Tuning by Soluble and Volatile Pharmacologics and Food Additives. *Advanced science (Weinheim, Baden-Wurtemberg, Germany)*, 11(20), e2306333.

Zhu X, et al. (2024) The feedback loop of EFTUD2/c-MYC impedes chemotherapeutic efficacy by enhancing EFTUD2 transcription and stabilizing c-MYC protein in colorectal cancer. *Journal of experimental & clinical cancer research : CR*, 43(1), 7.

Tian D, et al. (2024) Bacterial mucopeptides promote OXPHOS and suppress mitochondrial stress in mammals. *Cell reports*, 43(4), 114067.

Knippler CM, et al. (2024) Bisbiguanide analogs induce mitochondrial stress to inhibit lung cancer cell invasion. *iScience*, 27(4), 109591.