

Resource Summary Report

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RBE

RRID:CVCL_4896

Type: Cell Line

Proper Citation

(RRID:CVCL_4896)

Cell Line Information

URL: https://web.expasy.org/cellosaurus/CVCL_4896

Proper Citation: (RRID:CVCL_4896)

Sex: Female

Defining Citation: [PMID:9358283](#), [PMID:15767549](#), [PMID:27231123](#), [PMID:31077409](#),
[PMID:35640676](#), [PMID:39026794](#)

Comments: Omics: Transcriptome analysis by microarray., Omics: Deep exome analysis., Population: Japanese., Part of: JFCR45 cancer cell line panel., Part of: Cancer Dependency Map project (DepMap) (includes Cancer Cell Line Encyclopedia - CCLE)., Part of: Biliary tract cancer cell line atlas.

Category: Cancer cell line

Name: RBE

Cross References: CLO:CLO_0050848, BioSample:SAMN03472125, cancercelllines:CVCL_4896, CCRID:1101HUM-PUMC000675, CCRID:3101HUMTCHu179, Cell_Model_Passport:SIDM01850, CLS:305019, Cosmic:1187341, Cosmic:1571785, Cosmic:2629289, DepMap:ACH-001856, KCB:KCB 2014049YJ, RCB:RCB1292, Wikidata:Q54949330

ID: CVCL_4896

Record Creation Time: 20250131T202425+0000

Record Last Update: 20250131T204316+0000

Ratings and Alerts

No rating or validation information has been found for RBE.

No alerts have been found for RBE.

Data and Source Information

Source: [CelloSaurus](#)

Usage and Citation Metrics

We found 15 mentions in open access literature.

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

Li J, et al. (2024) Babaodan overcomes cisplatin resistance in cholangiocarcinoma via inhibiting YAP1. *Pharmaceutical biology*, 62(1), 314.

Chen T, et al. (2024) S100A6 drives lymphatic metastasis of liver cancer via activation of the RAGE/NF-kB/VEGF-D pathway. *Cancer letters*, 587, 216709.

Myint KZ, et al. (2024) Therapeutic Implications of Ceritinib in Cholangiocarcinoma beyond ALK Expression and Mutation. *Pharmaceuticals (Basel, Switzerland)*, 17(2).

Ratanabunyong S, et al. (2024) Exploring the apoptotic effects of sericin on HCT116 cells through comprehensive nanostring transcriptomics and proteomics analysis. *Scientific reports*, 14(1), 2366.

Zhang C, et al. (2024) Re-evaluation of the relationship between PrP^c expression and patient prognosis in primary esophageal squamous cell carcinoma and primary hepatocellular carcinoma. *Scientific reports*, 14(1), 31122.

Hu J, et al. (2024) Modulating PCGF4/BMI1 Stability Is an Efficient Metastasis-Regulatory Strategy Used by Distinct Subtypes of Cancer-Associated Fibroblasts in Intrahepatic Cholangiocarcinoma. *The American journal of pathology*, 194(7), 1388.

Noronha KJ, et al. (2024) NAPRT Silencing in FH-Deficient Renal Cell Carcinoma Confers Therapeutic Vulnerabilities via NAD⁺ Depletion. *Molecular cancer research : MCR*, 22(10), 973.

Alva-Ruiz R, et al. (2024) YAP-TEAD inhibition is associated with upregulation of an androgen receptor mediated transcription program providing therapeutic escape. *FEBS open bio*, 14(11), 1873.

Mohan A, et al. (2023) Devimistat in Combination with Gemcitabine and Cisplatin in Biliary Tract Cancer: Preclinical Evaluation and Phase Ib Multicenter Clinical Trial (BiT-04). *Clinical*

cancer research : an official journal of the American Association for Cancer Research, 29(13), 2394.

Jamnongsong S, et al. (2022) Comprehensive drug response profiling and pan-omic analysis identified therapeutic candidates and prognostic biomarkers for Asian cholangiocarcinoma. iScience, 25(10), 105182.

Dong L, et al. (2022) Proteogenomic characterization identifies clinically relevant subgroups of intrahepatic cholangiocarcinoma. *Cancer cell*, 40(1), 70.

Liu Z, et al. (2022) Liver kinase B1 in exosomes inhibits immune checkpoint programmed death ligand 1 and metastatic progression of intrahepatic cholangiocarcinoma. *Oncology reports*, 48(3).

Liu W, et al. (2022) Transcriptional repression and apoptosis influence the effect of APOBEC3A/3B functional polymorphisms on biliary tract cancer risk. *International journal of cancer*, 150(11), 1825.

Chen S, et al. (2022) Polymethoxylated flavone sudachitin is a safe anticancer adjuvant that targets glycolysis in cancer-associated fibroblasts. *Oncology letters*, 24(1), 236.

Hao X, et al. (2021) STK39 enhances the progression of Cholangiocarcinoma via PI3K/AKT pathway. iScience, 24(11), 103223.