

Cell Line Information Sheet for LIM2405

Cell Line Designation LIM2405

CellBank Catalogue No. CBA-0165

> Lot Number 016511111S

25 **Passage Number**

 4.0×10^6 cells **Total Cell Number**

94% **Expected Cell Viability**

> Adherent cell line derived from adenocarcinoma of the caecum of **Brief Description**

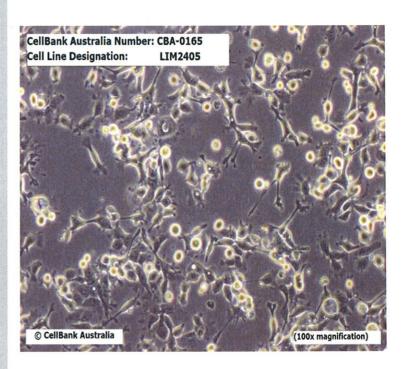
male patient.

Human (Homo Sapiens) Organism

> Tissue Colon

Growth Properties Adherent

> **Epithelial** Morphology



Image

Growth Medium

RPMI1640 (with 2mM L-Glutamine + 25mMHepes)+10%FCS, Insulin 0.6μg/ml, Hydrocortisone 1μg/ml, 1-Thioglycerol 10μM

Subcultivation Ratio

sub-confluent flasks (70-80% confluent) 0.05%Trypsin/EDTA at 37°C for 5 minutes. The optimal split ratio is 1:4-1:6 .Seeding density 1.6x10⁴ cells/cm²

Cells may take 2-3 days to seed after thawing or trypsinisation.



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Establishing and Maintaining your Culture

Cells are maintained at 37°C and 5% CO₂. LIM2405 requires growth medium to be changed 3 times each week. Passage every 4-5 days. Refer to Technical & Customer Service Information pamphlet for further information

Cryoprotectant Medium

10% DMSO + 90% FCS.

Biosafety Level

Cell line of human origin. Cellbank Australia recommends that cell lines be handled at category PC-2* containment level.

*AS/NZS 2243.3:2010

Use Restrictions

These cells are distributed for research purposes only - refer to the Material Transfer Agreement (MTA).

Safety Precaution

Where cell lines are shipped as frozen ampoules there is a small risk that the ampoule may be pressurised, due to the expansion of trapped liquid nitrogen and could explode on warming. It is recommended that persons handling ampoules of frozen cells wear appropriate personal protective equipment including laboratory coat, insulated gloves and a full protective face shield.

Handling Procedure for Frozen Cells

Upon receipt, frozen ampoules should be transferred directly to liquid nitrogen storage without delay, if not to be used immediately. Storage at -80°C may result in loss of viability. Remove protective cryoflex layer around the ampoule prior to thawing. A precentrifugation step to remove the cryoprotectant after thawing is necessary for this cell line.

Additional Information

Cells are adherent, spindly, grow as xenografts, heterozygous APC mutation (stop at aa 2198), B-Raf mutation (V600E), MSI, A33 negative.

Depositor

Professor Tony Burgess Ludwig Institute for Cancer Research, Australia

Whitehead R.H *et al* .Retention of tissue specific phenotype in a panel of colon carcinoma cell lines: Relationship to clinical correlates. Immunol Cell Biol. 1992; 70: 227-36

Reference

Zhang H. *et al.* Selective inhibition of proliferation in colorectal carcinoma cell lines expressing mutant APC or activated B-Raf Int.J.Cancer 2009 July15; 125(2):297-307



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