

Cell Line Information Sheet for LIM2550

Cell Line Designation LIM2550

CellBank Catalogue No. CBA-0169

Lot Number 01690810E

Total Cell Number 2.9x 10⁶ cells

Expected Cell Viability 90%

Passage number P12

Brief Description Colon carcinoma

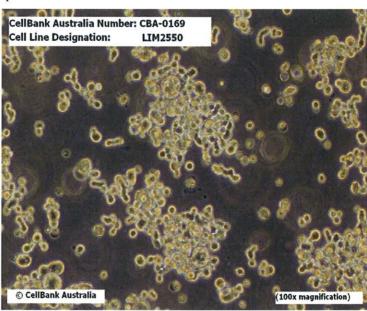
Organism Human (Homo Sapiens)

Tissue Colon

Growth Properties Cell line grows as floating and loose aggregates with some

adherent cells.

Morphology Epithelial



Image

Growth Medium

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

Subcultivation Ratio

Split sub confluent flasks (70-80%). Optimal split ratio 1:4-1:8 using 0.05%Trypsin /EDTA at 37°C for 5 minutes. Seeding density 1-2x10⁴ cells/cm²



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Establishing and Maintaining your Culture

Cells maintained at 37°C and 5% CO₂. LIM2550 requires growth medium to be changed 3 times each week. Feed and passage as for semi-adherent culture (collect medium with floating cells and then trypsinise the adherent population, pool both together before reseeding).

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

Derived from colon carcinoma, adherent and loose aggregates, MSI, A33 negative, truncated APC (stop at aa 1544), mutated p53 (aa G245D), heterozygous mutated K-Ras (aa Q61H), mutated PI3K (aa H1047R p110alpha).

Professor Tony Burgess

Depositor

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Australia

References

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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