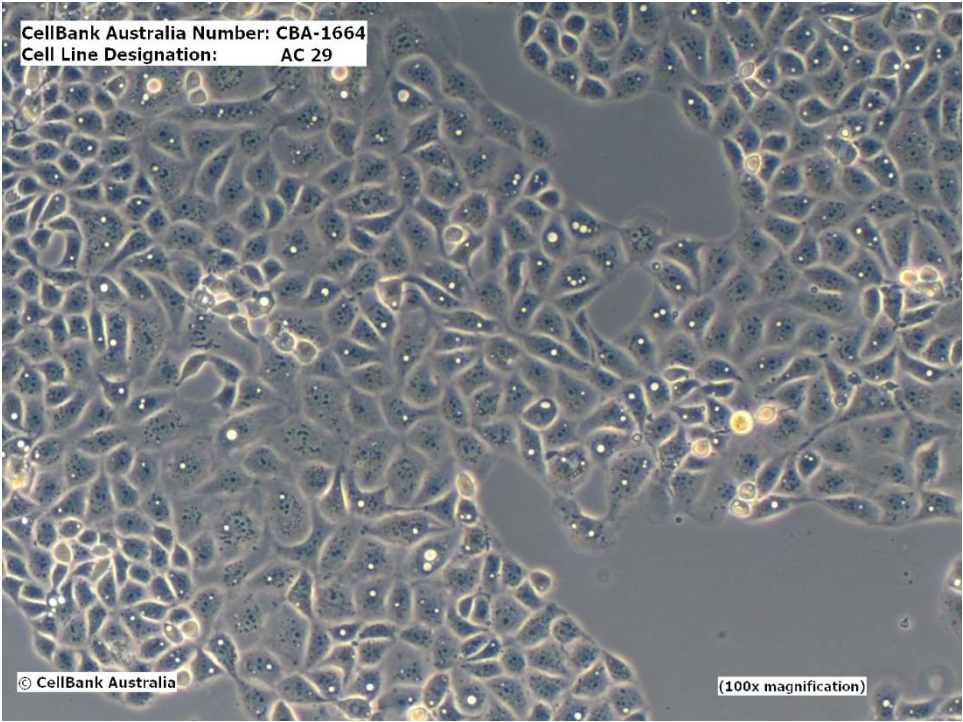


## Cell Line Information Sheet – AC 29

<b>CellBank Catalogue No.:</b>	<b>CBA-1664</b>
<b>Lot Number:</b>	<b>16640919E</b>
<b>Passage Number:</b>	80
<b>Total Cell Number:</b>	2.7 x 10 <sup>6</sup> cells
<b>Expected Cell Viability:</b>	93%
<b>Cell Line Description:</b>	<p>AC29 is a mouse mesothelioma cell line.</p> <p>Mice (CBA strain, female, 6-8 weeks old) were exposed to crocidolite asbestos through intraperitoneal injection, resulting in tumour development (both ascites and solid tumours). Cultures were established from malignant mesothelial cells obtained from ascites fluid.</p>
<b>Organism:</b>	Mouse ( <i>Mus musculus</i> )
<b>Strain:</b>	CBA
<b>Tissue:</b>	Mesothelium, malignant mesothelial cells were obtained from ascites fluid
<b>Growth Properties:</b>	Adherent
<b>Morphology:</b>	Epithelial-like
<b>Image:</b>	

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<b>Growth Medium:</b>	RPMI 1640 (with 2mM L-Glutamine + 25mM HEPES) + 5% Foetal Calf Serum
<b>Resuscitation</b>	<b>Remove protective cryoflex layer around the ampoule prior to thawing.</b> Thaw the ampoule by gently agitating in a 37°C waterbath; thawing should be rapid (around 2 minutes). A centrifugation step to remove the cryoprotectant after thawing is necessary for this cell line. For Lot 16640911E thaw 1 vial into 1xT75
<b>Subculturing Procedure:</b>	<b>Medium Renewal:</b> 2-3 times per week.  <b>Subcultivation ratio:</b> 1:8-1:16, Seeding density 0.6-1.0 x10 <sup>4</sup> cells/cm <sup>2</sup> Split subconfluent cultures (70-80%). Harvest the cells using 0.05% Trypsin/EDTA at 37°C for 5 minutes.  <b>Culture conditions:</b> Incubate the culture at 37°C with 5% CO <sub>2</sub> .  <b>Cryoprotectant Medium:</b> 10% DMSO + 90% FCS
<b>Safety Precaution:</b>	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.
<b>Handling Procedure for Frozen Cells:</b>	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
<b>Biosafety Level:</b>	Cell line of mouse origin.  CellBank Australia recommends that cell lines be handled at category PC-2* containment level.  *AS/NZS 2243.3:2010

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<b>Additional Information:</b>	<p>Cells are tumourigenic in syngeneic immunocompetent mice.</p> <p>CBA cells did not adapt as well to in vitro conditions, with a lower proportion of cell lines relative to BALB/c cells.</p> <p>AC29 cells express WT1 and have been used in the study of Wilms' tumour.</p>
<b>Depositor:</b>	Richard Lake - University of Western Australia
<b>References:</b>	<p><b>Original Reference</b></p> <p>Davis MR, Manning LS, Whitaker D, Garlepp MJ, Robinson BW (1992) Establishment of a murine model of malignant mesothelioma. Int J Cancer 52: 881-886. PubMed: <a href="https://pubmed.ncbi.nlm.nih.gov/1459729/">1459729</a></p>
<b>Use Restrictions:</b>	These cells are distributed for research purposes only - refer to the <a href="#">Sales Terms and Conditions</a> .
<b>CellBank Warranty:</b>	While CellBank Australia uses reasonable efforts to include accurate and up-to date information on this product sheet, CellBank Australia makes no warranties or representations as to its accuracy. Citations from scientific literature and patents are provided for informational purposes only. CellBank Australia does not warrant that such information has been confirmed to be accurate.
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