

**Designation: AR42J**

CLS order number: Cryovial: 500478
Vital: 550478

Origin and General Characteristics	
Organism:	<i>Rattus norvegicus</i> (rat)
Strain:	Wistar
Tissue:	Pancreas tumor, exocrine
Morphology:	Epitheloid
Growth Properties:	Cells grow slowly, in clusters and appear as hollow spheroid colonies. They can pile up and attach loosely.
Description:	Stimulation by glucocorticoids can enhance the secretory activity which is followed by extensive reorganization of the endoplasmic reticulum.
References:	Longnecker DS et al. Effect of age on nodule induction by azaserine and DNA synthesis in rat pancreas. J Natl Cancer Inst 58: 1769-75, 1977.
Culture Conditions and Handling	
Culture Medium:	DMEM supplemented with 4.5 g/L glucose, 2mM L-glutamine and 10% fetal bovine serum (MG-30, CLS order number 820300)
Subculturing:	It is recommended to cover the tissue culture flasks with gelatine prior to cell cultivation. Gelatine is added to the flask, incubated for 30min at 37°C and washed once with PBS. Remove medium and rinse the adherent cells using PBS without calcium and magnesium (3-5 ml PBS for T25, 5-10ml for T75 cell culture flasks). Add Accutase (1-2ml per T25, 2.5ml per T75 cell culture flask), the cell sheet must be covered completely. Incubate at ambient temperature for 8-10 minutes. Carefully resuspend the cells with medium (10 ml), centrifuge for 3 min at 300xg, resuspend cells in fresh medium and dispense into new flasks which contain fresh medium.
Split Ratio:	A split ratio of 1:2 every 48 hours. Is recommended.
Seeding density:	1x10 ⁴ cells/cm ²
Fluid Renewal:	2 to 3 times weekly
Freeze Medium:	CM-1 (CLS order number: 800125, 25ml, 800150, 50ml)
Freezing recovery:	After thawing, plate the cells at 5x10 ⁴ cells/cm ² and allow the cells to recover from the freezing process and to adhere for at least 48 hrs.
Sterility:	Fluorescence (DAPI) test: negative; Mycoplasma specific PCR: negative
Biosafety Level:	1
Safety precautions:	If the cryovial is planned to be stored in liquid nitrogen and to be thawed in the future, special safety precautions should be followed: Protective gloves and clothing should be used and a facemask or safety goggles must be worn when transferring frozen samples into or removing from the liquid nitrogen tank. The removal of a cryovial from liquid nitrogen may result in the explosion of the frozen vial creating flying fragments. Caputo, J.L. Biosafety procedures in cell culture. J. Tissue Cult. Methods 11:223-227, 1988. ATCC Quality Control Methods for Cell Lines, 2nd edition, 1992.
Special Features of the Cell Line	
Tumorigenic:	yes, in athymic mice
Viruses:	SMRV: Negative, as confirmed by Real-Time PCR

Authentication :	The rat origin was verified by Real-Time PCR.
Receptors Expressed:	Insulin; glucocorticoid
Products:	Amylase and other exocrine enzymes

Certificate of Analysis:	The Certificate of Analysis for each batch can be requested by e-mail at service@clsghmbh.de .
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Recommendations for handling of cells growing in suspension following delivery

Cryopreserved cells	<p>The cells come deep-frozen shipped on dry ice. Please make sure that the vial is still frozen.</p> <p>If immediate culturing is not intended, the cryovial(s) must be stored below -150°C after arrival.</p> <p>If immediate culturing is intended, please follow these instructions:</p> <p>Quickly thaw by rapid agitation in a 37°C water bath within 40-60 seconds. The water bath should have clean water containing an antimicrobial agent. As soon as the sample has thawed, remove the cryovial from the water bath. Note: A small ice clump should still remain and the vial should still be cold.</p> <p>From now on, all operations should be carried out under aseptic conditions.</p> <p>Transfer the cryovial to a sterile flow cabinet and wipe with 70% alcohol. Carefully open the vial and transfer the cell suspension into a 15 ml centrifuge tube containing 8 ml of culture medium (room temperature). Resuspend the cells carefully. Centrifuge at 300xg for 3 min and discard the supernatant. The centrifugation step may be omitted, but in this case the remains of the freeze medium have to be removed 24 hours later.</p> <p>Resuspend the cells carefully in 10ml fresh cell culture medium and transfer them into one T25 cell culture flask. All further steps are described in the Subculture section.</p>
Proliferating Cultures	<p>The cell culture flask, 1xT25, comes filled with cell culture medium.</p> <p>Incubate at 37°C for a minimum of 24 hrs.</p> <p>Count the cells, spin down the cell suspension at 300x g for 3 minutes to collect the cells. Resuspend the cells in an appropriate amount of fresh cell culture medium and transfer to new cell culture flasks.</p> <p>Incubate at 37°C for a minimum of 24 hrs.</p>

Warranty:	CLS warrants for a high cell viability and culture performance only if the product(s) is (are) stored and cultured according to the information described above. Using cell culture media and supplements other than the ones recommended in this product information may result in satisfactory proliferation and viabilities. CLS, however, does not warrant for cell recovery, proliferation and function if differing formulations are employed.
Disclaimer:	The customer shall not be entitled to employ this product for purposes other than research. Commercial utilization shall not be permitted; in particular, the cell line, its components or materials made therefrom shall not be sold or transferred to any third party. In addition, the term 'Commercial use' shall mean any activity by a party for consideration and may include, but is not limited to, use of the product or its components in manufacturing, for providing services, e.g. fee for service testing, in quality control or assurance processes within the manufacturing of products for sale, for therapeutic, diagnostic or prophylactic purposes, or for resale.