

**Designation: PC-3**

CLS order number: Cryovial: 300312
Vital: 330312

Origin and General Characteristics		
Organism:	Homo sapiens (human)	
Ethnicity:	Caucasian	
Age:	62 years old	
Gender:	Male	
Tissue:	Prostate; from metastatic site: bone	
Morphology:	Epithelial	
Cell type:	Prostate Adenocarcinoma, grade IV	
Growth Properties:	Monolayer; the cells form clusters in soft agar and can be adapted to suspension growth	
Description:	The cells exhibit low acid phosphatase and testosterone-5- alpha reductase activities.	
References:	Kaighn ME et al. Prostate carcinoma : tissue culture cell lines. Natl Cancer Inst Monogr 49 : 17-21, 1978.	
Culture Conditions and Handling		
Culture Medium:	DMEM:Ham's F12 medium (1:1 mixture) supplemented with L-glutamine and 5% fetal bovine serum (MG-40, CLS order number 820400).	
Subculturing:	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 TrypleExpress (1-2ml per T25, 2.5ml per T75 cell culture flask), the cell sheet must be covered completely. Incubate at 37°C for 10 minutes. Carefully resuspend the cells, the addition of medium is optional but not necessary, and dispense into new flasks which contain fresh medium.	
Split Ratio:	A ratio of 1:3 to 1:6 is recommended	
Seeding density:	Start culture after freezing at 3 x 10 ⁴ cells/cm ² . After cell recovery, use the seeding density of 1 x 10 ⁴ cells/cm ² for the subsequent splitting steps.	
Fluid Renewal:	2 to 3 times weekly	
Doubling time:	40 ± 10 hrs	
Freeze Medium:	CM-1 (CLS order number: 800125, 25ml, 800150, 50ml), contains serum	
Freezing recovery:	About 100% cells were recovered	
Sterility:	Fluorescence (DAPI) test: negative; Mycoplasma specific PCR: negative; Bacteria 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 nude mice	
Viruses:	SMRV: Negative, as confirmed by Real-Time PCR	
DNA Profile (STR):	Amelogenin: X,X CSF1PO: 11 D13S317: 11	wWA: 17 D3S1358: 16 D21S11: 29,31.2

	D16S539: 11 D5S818: 13 D7S820: 8,11 THO1: 6,7 TPOX: 8,9	D18S51: 14,15 Penta E: 10,17 Penta D: 9 D8S1179: 13 FGA: 24
Antigen Expression:	HLA A1, A9	

Certificate of Analysis:	The Certificate of Analysis for each batch can be requested by e-mail at service@clsgmbh.de .
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Recommendations for handling of adherent cell cultures following delivery

Cryopreserved cells	If immediate culturing is not intended, the cryovial(s) must be stored below -150°C or at least at -80°C after arrival. If immediate culturing is intended, please follow these instructions: 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. From now on, all operations should be carried out under aseptic conditions. 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. Resuspend the cells carefully in 10ml fresh cell culture medium and transfer them into two T25 cell culture flasks. All further steps are described in the Subculture section.
Proliferating Cultures	The cell culture flasks are completely filled with cell culture medium to prevent loss of cells during transit. Remove the entire medium except for a sufficient volume to cover the bottom of the flask. Incubate at 37°C for 24 hrs. Sometimes the cultures are handled roughly during transit, and most of the cells detach and float in the culture medium. If this has occurred remove the entire content of the flask and centrifuge at 300x g for 5 minutes. Take off the supernatant, resuspend the cells in 10 ml of culture medium and transfer the entire cell suspension into cell culture flasks of suitable size (size (do not seed in more than 1T75 flask).

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.