Designation:	ΗK	
Synonym(s):	FDC/	
CLS order number:	Cryov Vital:	

FDC/HK Cryovial: 300204 Vital: 330204 DNA: 300204GD

Origin and General Characteristics		
Organism:	Homo sapiens (human)	
Ethnicity:	Caucasian	
Age:	Unknown, child	
Gender:	Sex unspecified. Female, acc. to CLS authentication by STR analysis	
Disease:	Tonsillitis	
Tissue:	Tonsil	
Morphology:	Fibroidal	
Cell type:	Follicular dendritic reticulum cells (FDCs)	
Growth Properties:	Adherent	
Description:	The cell line was established from tonsillar tissue after routine tonsillectomy of a child in 1994. In coculture with activated (anti-CD40/anti- μ) human lymphocytes soluble factors are released by HK which support growth of normal and malignant lymphocytes.	
References:	Kim H-S, Zhang X, Choi YS. Activation and proliferation of follicular dendritic cell-like cells by activated T lymphocytes. J Immunol 153:2951-2961 (1994).	
	Kim H-S, Zhang X, Klyushnenkova E, Choi YS. Stimulation of germinal center B lymphocyte proliferation by an FDC-like cell line, HK. J Immunol 155:11101-1109 (1995).	
	Kagami Y, Jung J, Choi YS, Osumi K, Nakamura S, Morishima Y, Seto M. Establishment of a follicular lymphoma cell line (FLK-1) dependent on follicular dendritic cell-like cell line HK. Leukemia 15:148-156 (2001).	
	Chihara D, Kagami Y, Kato H, Yoshida N, Kiyono T, Okada Y, Kinoshita T, Seto M. IL- 2/IL-4, Ox40L and FDC-like cell line support the in vitro tumor cell growth of adult T-cell leukemia/lymphoma. Leukemia Res 38:608-612 (2014).	
Cellosaurus citation:	FDC/HK (RRID:CVCL_IY38)	
Culture Conditions and I	Handling	
Culture Medium:	EMEM supplemented with NEAA, L-glutamine and 20% fetal bovine serum.	
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 Accutase (1-2ml per T25, 2.5ml per T75 cell culture flask), the cell sheet must be covered completely. Incubate at ambient temperature for 5-8 minutes. Carefully resuspend the cells with medium (10 ml), centrifuge for 5 min at 300xg, resuspend cells in fresh medium and dispense into new flasks which contain fresh medium.	
Seeding density:	Start new culture in plating the freshly thawed cells into $2xT25$ cell culture flasks. Actively proliferating cells can be plated at $1x10^4$ cells/cm ² .	
Fluid Renewal:	1 to 2 times weekly	
Doubling time:	n.d.	
Freeze Medium:	CM-1 (CLS order number 800150, 50ml)	
Freezing recovery:	Within 24 hours	
Sterility:	Mycoplasma specific PCR: negative Mycoplasma specific PlasmoTest: negative	
Biosafety Level:	2	

cell lines service

	HK was tested positive for EBV. According to the test of test	ording to the German Law for the Protection against G), this cell line falls under risk group L2, and can ing a valid permit of the respective authority (IfSG
Safety precautions:	If the cryovial is planned to be stored special safety precautions should be Protective gloves and clothing should worn when transferring frozen sample The removal of a cryovial from liquid vial creating flying fragments. Caputo, J.L. Biosafety procedures in cell Quality Control Methods for Cell Lines, 2r	in liquid nitrogen and to be thawed in the future, followed: I be used and a facemask or safety goggles must be es into or removing from the liquid nitrogen tank. nitrogen may result in the explosion of the frozen culture. J. Tissue Cult. Methods 11:223-227, 1988. ATCC id edition, 1992.
Special Features of the Cell Line		
Viruses:	Contains EBV.	
Surface antigens:	CD14+, CD40+, ICAM-1+, VCAM-1+	
HLA-typing:	n.d.	
DNA Profile (STR):	Amelogenin: X,X CSF1PO: 10,11 D13S317: 10,13 D16S539: 9,12 D5S818: 12 D7S820: 9,11 TH01: 8,9 TPOX: 10,11	D3S1358: 14,16 D21S11: 28,30 D18S51: 12,19 D8S1179: 10,14 FGA: 22,22 vWA: 16,17 Penta E: 7,11 Penta D: 9,12
Applications:	Feeder cell for growth of normal B lymphocytes and lymphomas/leukemias. Studies on B cell development in germinal centers of lymph nodes. Possibly research on virus infection of FDCs	

Certificate of Analysis:	The Certificate of Analysis for each batch can be requested by e-mail at
	service@clsgmbh.de.

Recommendations for handling of cells growing in suspension following delivery		
Cryopreserved cells	The cells come deep-frozen shipped on dry ice. Please make sure that the vial is still frozen. If immediate culturing is not intended, the cryovial(s) must be stored below -150°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 one T25 cell culture flask. All further steps are described in the Subculture section.	
Vital, proliferating cells:	EBV-containing cells are not shipped out as vital cells.	

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.
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