

Designation	NCH612
Synonyms	n.a.
Catalog number - cryovial	300121
Catalog number - vital cells	330121

Please read first

Biosafety level	1
Citation	NCH612 (CLS catalog number 300121)

Origin and General Characteristics

Organism	Human
Ethnicity	Caucasian
Age	39
Gender	Male
Tissue	Brain
Disease	Anaplastic oligodendroglioma, WHO °III, IDH1 mutant (R132H)
Growth properties	Spheroid culture
Depositor	C. Herold-Mende
Cellosaurus accession ID	CVCL_X913
Wikidata	Q54907711
NCBI_TaxID	9606
References	References are available in the product page at www.cls.shop

Culture Conditions and Handling

Culture medium	GBM-MG, 0% FBS
Subculturing	Mechanical dissociation by pipetting (5-10 times) with an Eppendorf
	pipette and 1000 μ l filter tips after centrifugation of the cells (5min,
	300g, RT).
Split ratio	A ratio of 1:2 to 1:5 is recommended
Seeding density	1 x 10^ cells/ml
Fluid renewal	Fresh medium must be added every 2-3 days (2-5 ml depending on
	the size of the cell culture flask).
Freeze medium	CM-ACF (CLS catalog number 800650)
Freeze medium	Slow. After thawing allow the cells to recover from the freezing
	process for at least 48 hours.
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Sterility

Plasmotest: negative; Mycoplasma specific PCR: negative

Special Features of the Cell Line

DNA profile (STR)	Amelogenin	X,X	D18S51	13
	CSF1PO	11,12	Penta E	11,14
	D13S317	10	Penta D	9,12
	D16S539	11,13	D8S1179	13
	D5S818	11,13	FGA	21
	D7S820	10,11	D1S1656	n.a.
	TH01	6,7	D6S1043	n.a.
	ТРОХ	8,12	D2S1338	n.a.
	vWA	17	D12S391	n.a.
	D3S1358	14,18	D19S433	n.a.
	D21S1	28,31		
HLA-typing	A*	n.a., n.a.	DQB1*	n.a., n.a.
	B*	n.a., n.a.	DPA1*	n.a., n.a.
	C*	n.a., n.a.	DPB1*	n.a., n.a.
	DRB1*	n.a., n.a.	E	n.a., n.a.
	DRB3*	n.a., n.a.	G	n.a., n.a.
	DRB4*	n.a., n.a.	F	n.a., n.a.
	DRB5*	n.a., n.a.	MR-1	n.a., n.a.
	DQA1*	n.a., n.a.		



Recommendations for handling of cells 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 must be stored below -150°C after arrival. If
	immediate culturing is intended, please follow the below
	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. 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 300 x g 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 10 ml 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, 2xT25, come filled with cell culture medium.
	Collect the entire medium in 2 x 50 ml centrifuge tubes. Carefully
	add 5 ml of cell culture medium to each of the two T25 cell culture
	flasks. Control the cell morphology and confluency under the
	microscope. Incubate at 37 °C for a minimum of 24 hours. Spin
	down the collected medium at 300 x g for 3 minutes to collect the
	cells which may have detached during transit. If a cell pellet is
	visible, resuspend the cells in 5 ml of cell culture medium and
	transfer to 1 x T25 cell culture. Incubate at 37 $^\circ C$ for a minimum of
	24 hours.



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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.
Warranty	CLS warrants for a high cell viability and culture performance only
	if the products are stored and cultured according to the
	information described in this product sheet.
Certificate of Analysis	The certificate of analysis can be requested by email at
	info@cls.shop for each batch. Please indicate the lot number of
	your product in the email.
Disclaimer	This product is intended for laboratory research use only. It is
	strictly prohibited in any case to use the cells, their progeny,
	unmodified derivatives and modifications in human or animal
	subjects for therapeutic, diagnostic or prophylactic purposes. All
	regulations as described in the Material Transfer Agreement or
	Supply Agreement concluded with the order apply.