

Updated on: 24th May 2023

CERTIFICATE OF ANALYSIS

Lot#: HC-03

PRODUCT DESCRIPTION

Reference: HuHECP/4Product: Cryopreserved Human Hepatocytes
Category: Plateable

Isolation date: 28th November 2017
Initial Isolation Viability: 84.00%
Storage conditions: -196°C using LN₂

Spheroid qualified: NO Sterility test: negative for bacteria, yeast, and

(see details below: 3D Spheroid formation section) fungi

DONOR DEMOGRAPHICS

Species	Gender	Race	Age	ВМІ	Smoker	Alcohol Use	Drug Use
Human	Male	Caucasian	71	22.98	No	No	No
P	athology		Serological Data ¹				
	N/A		Tested negative less than 3 months before surgery				

Patient informed consent was obtained. ¹The donor was serologically tested negative for following infectious diseases: HIV, Hepatitis B, and C. Donor medical history was also examined prior to accepting this donor. *For donor's medication information, please contact us.*

CHARACTERIZATION FOR PLATEABLE CELLS

Post Thaw Lot information	Result	SD	n
Number of viable cells (cells/vial):	3.62x10 ⁶	± 0.25x10 ⁶	2
Post-thaw viability (%):	84.44	± 3.12	2
Days in culture after thaw (24w):	3	± 0.00	1
Days in culture after thaw (96w):	3	± 0.00	1

MONOLAYER ASSESSMENT² Plateable: YES Confluence 24h: 80.00% Seeding density in 24 well recommended: 2.37x10⁵ cells/cm² Seeding density in 96 well recommended: 2.81x10⁵ cells/cm²

Cell morphology 24h



Human hepatocytes were thawed and seeded according to Cytes Biotechnologies culture protocol. The yield and viability were determined by a trypan blue exclusion assay after the thawing process. ²Resuspended human hepatocytes from post-thaw assessment were plated in collagen-coated 24-well plates in hepatocyte plating medium. Cells were refreshed with hepatocytes maintenance medium at first medium during the first change of medium on the day of thawing. Maintenance medium was replaced in the culture every day. If images from the 96-well plates are needed, please contact us.

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3D SPHEROID FORMATION

Spheroid morphology

Cytes **does not guarantee** that these primary hepatocytes will be suitable for 3D culture and creation of spheroid structures while using Cytes protocols.

If you need help for an experiment, just contact us, our experts will be pleased to assist you

CERTIFICATION:

The viability and performance of the primary human hepatocytes provided depend primarily on the use of appropriate media and reagents, as well as the use of sterile plastics. Likewise, proper handling protocols must be followed. Please note that if these parameters are not carefully considered, the cellular response obtained in the assays may be lower than expected.

Name	Tittle	Signature	Cytes Biotechnologies, S.L.	Date
Pilar Sainz de la Maza	Quality Manager	Plan Jamber	CYTES BIOTECHHOLOGIES S.L.	24/05/23

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CELL COUNTING

Lot #:			Date	:/_		
MORPHOLOGY						
☐ Clear cytoplasm☐ Clear membranes	☐ Rounded shape ☐ Membrane blebbing	☐ Cell swelling ☐ Lipid drople		☐ Hardly any debris☐ Prevalent debris		
TRYPAN BLUE COUNTING RESULTS						
		NEUBAUER CHAMBER COUNTING				
01	Quadrant	Live cells +	Dead cells		tal cells	
Q1 Q2	Quadrant 1	+	Dead cens	=	rear cens	
	Quadrant 2	+		=		
	Quadrant 3	+		=		
	Quadrant 4	+		=		
Q3	Total	+		=		
(Live cells) (Total cells) $x 100 = 0$ Viability (%) YIELD (Total cells) $x (Dilution factor)$ $x 10^4 * x (Current volume)$ ml (Counted quadrants) *This factor (10 ⁴) is applicable when it is used a Hemocytometer SEEDING DENSITY (Desired number of cells) cells $x (Current volume)$ ml (Total number of cells) cells (Total number of cells) cells (Total number of cells) cells (Total volume per dish or plate to use (Volume needed) and then calculate the needed volume to add: (Total volume well) $ml - (Cells total volume)$ $ml = 0$						
Surface of the most cor	mmon plates for culture	ThermoFisher Corning® Falcon® Eppendorf	24-well plate 1.90 cm ² /well 2.00 cm ² /well 1.90 cm ² /well 2.08 cm ² /well	96-well plate 0.32 cm²/well 0.36 cm²/well 0.32 cm²/well 0.37 cm²/well		
			COUNTED BY:			

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