CYTES BIOTECHNOLOGIES S.L.

Updated on: 24th march 2023

CERTIFICATE OF ANALYSIS

Lot#: BHum15074

PRODUCT DESCRIPTION

Reference: HuHECP/6+
Product: Cryopreserved Human Hepatocytes
Category: Plateable
Spheroid qualified: NO
(see details below: 3D Spheroid formation section)

Isolation date: 23rd December 2015
Initial Isolation Viability: 81%
Storage conditions: -196°C using LN_2
Sterility test: negative for bacteria, yeast, and fungi

DONOR DEMOGRAPHICS

Species	Gender	Race	Age	BMI	Smoker	Alcohol Use	Drug Use
Human	Male	Caucasian	66	25.28	No	No	No

Pathology	Serological Data[1]
Colorectal cancer	Tested negative less than 3 months before surgery

Patient informed consent was obtained. [1]The 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):	7.70 x10^6	± 0.00	1
Post-thaw viability (%):	93.00	± 0.00	1
Days in culture after thaw (24w):	5	± 0.00	1
MONOLAYER ASSESSMENT[2] **Plateable: YES**			

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. [2]Resuspended 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.

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.

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

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			24/03/23

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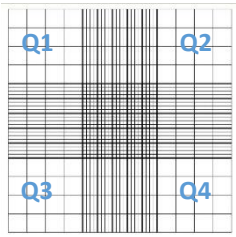
CELL COUNTING

Lot #: _____ **Date:** ____/_____/_____

MORPHOLOGY

❑ Clear cytoplasm	❑ Rounded shape	❑ Cell swelling	❑ Hardly any debris
❑ Clear membranes	❑ Membrane blebbing	❑ Lipid droplets	❑ Prevalent debris

TRYPAN BLUE COUNTING RESULTS



NEUBAUER CHAMBER COUNTING				
Quadrant	Live cells	+	Dead cells	= Total cells
Quadrant 1		+		=
Quadrant 2		+		=
Quadrant 3		+		=
Quadrant 4		+		=
Total		+		=

VIABILITY

$$\frac{(Live\ cells)}{(Total\ cells)} x100 = \quad Viability\ (\%)$$

YIELD

$$\frac{(Total\ cells) \quad x\ (Dilution\ factor) \quad x\ 10^4\ ^*x\ (Current\ volume) \quad ml}{(Counted\ quadrants)} = \quad cells\ (Total\ number\ of\ cells)$$

**This factor (10^4) is applicable when it is used a Hemocytometer*

SEEDING DENSITY

$$\frac{(Desired\ number\ of\ cells) \quad cells\ x\ (\ Current\ volume) \quad ml}{(Total\ number\ of\ cells) \quad cells} = \quad ml\ (Volume\ needed\ for\ your\ cells)$$

Keep in mind the final volume per dish or plate to use (Volume needed) and then calculate the needed volume to add: $(Total\ volume\ well) \quad ml\ -(Cells\ total\ volume) \quad ml = \quad ml\ (Volume\ to\ add)$

Surface of the most common plates for culture:

Brand	24-well plate	96-well plate
ThermoFisher	1.90 cm^2/well	0.32 cm^2/well
Corning®	2.00 cm^2/well	0.36 cm^2/well
Falcon®	1.90 cm^2/well	0.32 cm^2/well
Eppendorf	2.08 cm^2/well	0.37 cm^2/well

COMMENTS

COUNTED BY: