

CERTIFICATE OF ANALYSIS

CRYOPLATEABLE HUMAN HEPATOCYTES

FOR RESEARCH USE ONLY. CAUTION: Not intended for human diagnostic or therapeutic uses. Users should treat all human cells as potential pathogens. Wear protective clothing and eyewear. Practice appropriate disposal techniques for potentially pathogenic or bio-hazardous materials.

PRODUCT INFORMA	ATION:		
Product name:		Lot identifier:	
Post-thaw viability of ≥%		Post-thaw confluency ≥ % on day 5	
Contains a minimum of	x 10 ⁶ viable cells/mL		
DONOR DEMOGRAP	PHICS:		
Sex: Age:	BMI: Ethnicity:	COD:	
Donor negative for: HIV, HCV, HBV, RPR		Culture negative for: Gram+, Gram-, Mycoplasm, Fungi	
PLATING:			
Recommended seeding density (million cells/mL):		Recommended thaw medium:	
Lot	hours post-plating	Lot	hours post-plating

3D spheroid formation of Lot _____

days post-thaw.

CDFDA Stain of Lot ______ after _____hours post-plating

ENZYME INDUCTION ACTIVITY

ENZYME	TREATMENT	mRNA FOLD-CHANGE
CYP1A2	Omeprazole (25 µM)	
CYP2B6	Phenobarbital (2 mM)	
CYP3A4	Rifampin (10 µM)	

Hepatocytes were seeded on a 24-well plate pre-coated with collagen I, and cultured for 48 hours at 37° C in 5% CO₂ and 90% humidity. The hepatocytes were then incubated at 37° C in 5% CO₂ and 90% humidity with omeprazole (25μ M, CYP1A2), phenobarbital (2μ M, CYP2B6), rifampin (10μ M, CYP3A4) and vehicle control for 48 hours, respectively. The media were replaced every 24 hours with fresh media containing the positive inducers and vehicle. The mRNA levels of CYP1A2, CYP2B6 and CYP3A4 were determined using RT-PCR.

METABOLIC ACTIVITY

ENZYME	SUBSTRATE	CONCENTRATION (μΜ)	ENZYME ACTIVITY (pmole/min/million cells)
CYP1A2	Phenacetin	100	
CYP2A6	Coumarin	50	
CYP2B6	Bupropion	500	
CYP2C8	Amodiaquine	20	
CYP2C9	Diclofenac	25	
CYP2C19	S-mephenytoin	250	
CYP2D6	Dextromethorphan	15	
CYP2E1	Chlorzoxazone	250	
CYP3A4	Testosterone	200	
CYP3A4	Midazolam	20	
Phase I CYP	7-ethoxycoumarin	100	
SULT	7-hydroxycoumarin sulfation	100	
UGT	7-hydroxycoumarin glucuronidation	100	

Hepatocytes in suspension (0.5 million/mL) were incubated with substrate at 37°C in 5% CO₂ and 90% humidity for 30 minutes, respectively. The concentrations of the metabolites were determined using LC-MS/MS methods.

by these institutions from the donor or the donor's legal next of kin, for use of the tissue and its derivatives for research purposes.

Quality Assurance

Date

The cells from this lot were derived from tissue obtained from accredited institutions. Consent was obtained



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