

HCP100.H15 Lot No. 2210212

Pool of 100 (50 Males and 50 Females)

Assured Minimum Yield: 5.0 x 10⁶ cells per vial

Viability: 81%

This product was pooled from individual human hepatocytes that have been frozen and thawed. The yield and viability given above are based on experiments performed at XenoTech using our thawing protocol that includes a density gradient fractionation and the K8500 OptiThaw Kit. Details of XenoTech's hepatocyte thawing protocol can be found at www.xenotech.com. It is recommended to use XenoTech's thawing protocol, which includes a density gradient step, to maximize the viability of the recovered cells.

Enzyme	Marker Substrate Reaction	[S] (µM)	Rate (pmol/million cells/min)
CYP1A2	Phenacetin O-dealkylation	100	50.6 ± 0.6
CYP2A6	Coumarin 7-hydroxylation	50	45.8 ± 3.9
CYP2B6	Bupropion hydroxylation	500	38.7 ± 3.5
CYP2C8	Amodiaquine N-dealkylation	20	313 ± 10
CYP2C9	Diclofenac 4'-hydroxylation	100	205 ± 22
CYP2C19	S-Mephenytoin 4'-hydroxylation	400	12.3 ± 0.6
CYP2D6	Dextromethorphan O-demethylation	80	42.5 ± 0.9
CYP2E1	Chlorzoxazone 6-hydroxylation	500	99.2 ± 4.1
CYP3A4/5	Testosterone 6β-hydroxylation	250	167 ± 6
CYP3A4/5	Midazolam 1'-hydroxylation	30	48.9 ± 0.8
UGT	7-Hydroxycoumarin glucuronidation	100	319 ± 32
SULT	7-Hydroxycoumarin sulfonation	100	12.8 ± 1.3

Values for enzyme activities were determined at a single substrate concentration run with triplicate determinations.

To measure cytochrome P450 (CYP), UDP-glucuronosyl transferase (UGT) and sulfotransferase (SULT) activities, hepatocytes (1 x 10^6 cells/mL) in suspension were incubated in triplicate at $37 \pm 2^{\circ}$ C for 30 minutes in Optilncubate and marker substrate, at the final concentrations indicated. Metabolite formation was determined by validated LC-MS/MS methods with deuterated metabolites as internal standards.

Uptake Activity Data

Uptake Transporter	Marker Substrate	[S] (µM)	Rate (pmol/million cells/min)
OATP1B1	Estrone sulfate	1	13.5
OATP1B3	CCK-8	1	1.6
OCT1	MPP+	1	5.2
NTCP	TCA	1	4.6

To measure uptake activities, hepatocytes (1.0 x 10⁶ cells/mL) in suspension were incubated in triplicate at 4°C and 37°C for 1 minute in Krebs-Henseleit buffer and marker substrate, at the final concentrations indicated. Uptake of substrate was measured by scintillation counter.

Donor Information

Gender: Males (50), Females (50)

Age: 8-74 years of age

Race: Caucasian (80), African American (9), Asian (3), Hispanic (8)
Cause of Death: Anoxia (39), Head trauma (25), Cerebrovascular accident (36)

Antibody to Cytomegalovirus (CMV): Positive (55), Negative (45)

All donors tested negative for Human Immunodeficiency Virus (HIV), Hepatitis B Surface Antigen (HBsAg), Hepatitis C Virus, and Rapid Plasma Reagin.



Store in liquid nitrogen, vapor phase

CAUTION: This sample should be considered as a potential biohazard and universal precautions should be followed. Intended for in vitro use only.

These data were generated by and are the property of XenoTech. These data are not to be reproduced, published or distributed without the express written consent of XenoTech.

This data sheet serves as a Certificate of Analysis and has been approved by Stephanie Helmstetter, Senior Manager.

Signature and Date: Stephanic Helmstetter 25 October 2022