

CryostaX Geneknown™

Single Freeze Pooled Cryopreserved Human Hepatocytes

HPCH.OATP1B1.HA Lot No. 1510250

OATP1B1 Genotype: *1/*1B + *5/*5

Pool of 2

Assured Minimum Yield: 4.5 x 10⁶ cells per vial

Viability: >70.0%

Individual Donor CYP Genotype Information:

Donor	CYP1A1	CYP1A2	CYP2A6	CYP2B6	CYP2D6	CYP2C8	CYP2C9	CYP2C19	CYP2E1	CYP3A4	CYP3A5
945	*1/*1	*1/*1	*1/*1	*1/*6	*1/*68x2+*4	*1/*1	*1/*1	*1/*2	*1/*1	*1/*1	*3/*3
1218	*2/*4	*1C/*1	ND	*1/*1	*1/*1	*1/*1	*1/*1	*1/*1	*1/*1	*1/*1	*3/*3

Enzyme	Marker Substrate Reaction	[S] (µM)	Rate (pmol/million cells/min)
CYP1A2	Phenacetin O-dealkylation	100	22.8
CYP2A6	Coumarin 7-hydroxylation	50	9.6
CYP2B6	Bupropion hydroxylation	500	28.7
CYP2C8	Amodiaquine N-dealkylation	20	124
CYP2C9	Diclofenac 4'-hydroxylation	100	249
CYP2C19	S-Mephenytoin 4'-hydroxylation	400	3.54
CYP2D6	Dextromethorphan O-demethylation	80	44.3
CYP2E1	Chlorzoxazone 6-hydroxylation	500	150
CYP3A4/5	Testosterone 6β-hydroxylation	250	99.4
CYP3A4/5	Midazolam 1'-hydroxylation	30	12.9

Values for enzyme activities are averages of the individual donor activities for each specified CYP, therefore the values indicated should be considered theoretical.

ND: Not determined

Donor Information

Gender:Male (1), Female (1)Age:24-68 years of ageRace:Caucasian (2)

Cause of Death: Head trauma (1), Cerebrovascular accident (1)

Cytomegalovirus (CMV): Negative (2)

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.

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Datasheet prepared 23 January 2019

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