

Cellartis® Enhanced hiPS-HEP

Frozen hepatocytes as effective tools for *in vitro* evaluation of hepatotoxicity and drug metabolism studies







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Cellartis® Enhanced hiPS-HEP is a highly homogenous population of human iPS-derived heptocytes. These cells are ideal for use in drug metabolism and toxicology related applications that demand a highly reproducible platform,

supply of material and substantial expression of drug metabolism enzymes.

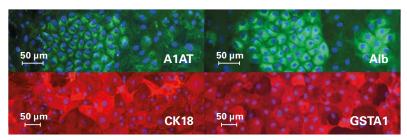


Figure 1: Cryopreserved Cellartis® Enhanced hiPS-HEP express α 1-antitrypsin (A1AT), Albumin (Alb), Cyto-keratin 18 (CK18) and Glutathione S-transferase A1 (GSTA1).

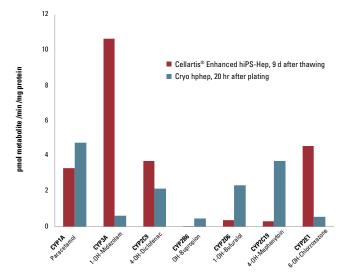


Figure 2: CYP activities of cryopreserved Cellartis® Enhanced hiPS-Hep compared to 4 donors of cryopreserved human primary hepatocytes (hphep)

ADVANTAGES

- Relevant & predictive data
- Human iPS cell derived hepatocytes with substantial expression of drug metabolism enzymes relevant for hepatotoxicity
- Reproducible data
- Low lot-to-lot variation
- Robust differentiation protocols resulting in highly homogenous population (more than 90 % pure)
- Convenient & flexible formats
- Frozen format
- 2D-cultures

APPLICATIONS

- Toxicity testing
- Drug discovery and metabolism studies
- Vaccine development
- High content analysis

CHARACTERISTIC	ASSAY	ANALYTE	
CYP activity	LC/MS	CYP1A (paracetamol), CYP2B6 (OH-bupropion), CYP2C9 (OH-diclofenac), CYP2C19 (OH-mephenytoin), CYP2D6 (OH-bufuralol), CYP2E1 (OH-chlorzoxazone), CYP3A (OH-midazolam)	
Protein markers	ICC	α 1-antitrypsin, Albumin, HNF4 α , CK18, GSTA1	
Gene expression	qPCR	Phase I, e.g., CYP1A1, 1A2, 2B6, 2C9, 2C19, 2D6, 3A4, 3A5, low 3A7	
Gene expression	qPCR	Phase II, e.g., UTGT1A1, 2B7, GSTA1.1	
Gene expression	qPCR	Transporters, e.g., BSEP, BCRP, MDR-1 (P-gp), MRP2, NTCP, OATP1B1, OCT-1	
Gene expression	qPCR	Albumin, α 1-antitrypsin, low AFP	
Capability to store	PAS staining	Glycogen	

PRODUCT	CATALOGUE#	SOURCE	FORMAT
Frozen Cellartis® Enhanced hiPS-HEP	HEP-104-VIAL-KIT	hiPS cell line 18	1 vial suitable for one 96-well plate

Selected references

1. Uhestad M *et al*, Drug metabolizing enzyme and transporter protein profiles of hepatocytes derived from human embryonic and induced pluripotent stem cells. Biochem Pharmacol. 20

2. Yildiniman R *et al*, Human embryonic stem cell derived hepatocyte-like cells as a tool for *in vitro* hazard assessment of chemical carcinogenicity, Toxicol Sci. 2011 Dec. 12/4(2): 278-90

3. Manten N *et al*, Fotential markers of attenuation of YF virus after infection of stem cell-derived human hepatocytes with wild-type Asibi or live-attenuated YF 17D virus, Supplement to the American Journal of tropical Medicine and Hygiene, Volume 83, November 2010, Number 5, abstacts 12

4. Heins *et al*. Stem Cells 2004; 22: 387-376 United States National Stem Cell Bank; http://www.nationalstemcellbank.org col. 2013 Sep 1;86(5):691-702