

The hepatic stem cell line

PRODUCT SPECIFICATIONS

DIFFERENTIATED HEPARG[®] CRYOPRESERVED

CATALOG NUMBER: HPR116

For in vitro and research use only

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www.HepaRG.com

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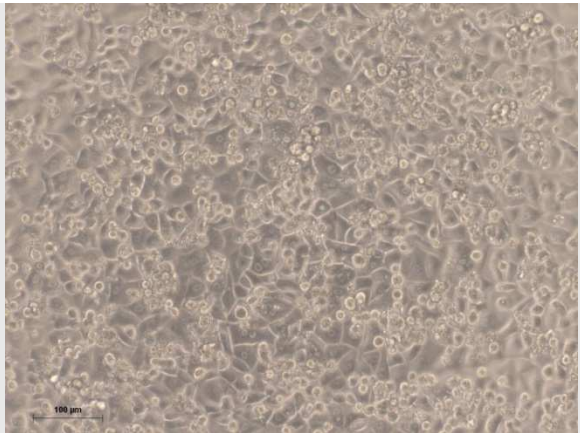
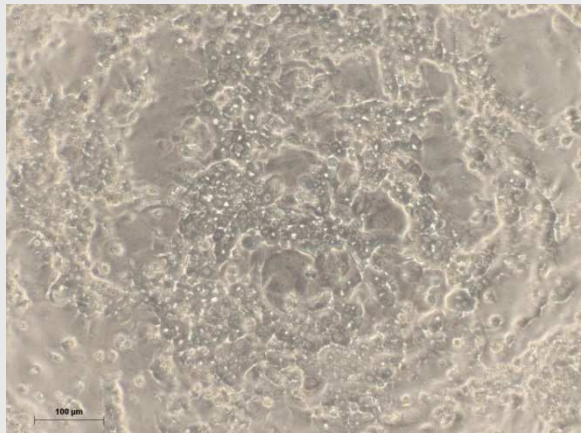
HepaRG[®] cells are patented and their use is strictly limited; consider the cells as a single-use, disposable product that must be destroyed upon conclusion of a study or experiment. Propagating, reproducing, cloning, subcloning or any other use of the cells following the conclusion of a study is prohibited. Use of the cells to produce or manufacture commercial products for general sale or for use in the manufacture of products intended for general sale is prohibited. Transfer of the cells to anyone not employed within the same organization, whether for financial benefit or not, is prohibited. **If you are unwilling to accept the terms of this LIMITED USE LICENSE, do not ORDER or use them, and immediately return the cells for credit. Violators of this Limited Use License will be prosecuted to the fullest extent of the law.**

1. PRODUCT DESCRIPTION

Species	Human	
Origin	Liver tumor of a patient suffering from hepatocarcinoma and hepatitis C infection	
Safety data	Absence of hepatitis B, hepatitis C and HIV1 viruses checked by PCR on the cell suspension. <i>Caution: although controls are performed, human material has to be considered as potentially dangerous. Take maximum care in order to protect yourself and your colleagues.</i>	
Biosafety level	Level 2, as recommended in the CDC-NIH Manual, Biosafety in Microbiological and Biomedical Laboratories, 5th edition, N°21-1112 Revised December 2009	
Description	HepaRG [®] cells are human hepatic progenitor cells able to give rise to adult fully differentiated hepatocytes in appropriate culture conditions. HepaRG [®] cells, catalog number HPR116, are cryopreserved after differentiation using a proprietary process developed by Biopredic International. A few days after thaw and culture, the cells will form a coculture of hepatocytes and of biliary epithelial-like cells. Gripon P. <i>et al. Proc. Natl. Acad. Sci. 99, 15655-15660, 2002.</i>	
Packaging	HPR116-TA08: 1 mL vial with $\geq 8 \times 10^6$ viable cells HPR116-TA12: 1 mL vial with $\geq 12 \times 10^6$ viable cells	
Cell seeding	One vial of HPR116-TA08 will fully seed one 96-well plate (72 000 cells/well) One vial of HPR116-TA12 will fully seed one 24-well plate (480 000 cells/well)	
CYP genotyping data	Obtained in collaboration with N. Picard at Limoges hospital (France) using a validated TaqMan allelic discrimination assay (ABI PRISM 7000 Sequence Detection System, Applied-Biosystems).	
SNPs description SNPs: Single Nucleotide Polymorphisms wt: wild- type mut: mutation **: wt=G T=minor allele in Caucasians	CYP2D6	
	*2 (2850C>T)	*2/wt
	*3 (2549delA)	wt/wt
	*4 (1846G>A)	wt/wt
	*7 (2935A>C)	wt/wt
	*10 (100C>T)	wt/wt
	CYP3A4	
	*1B (392G>A)	wt/wt
	OATP1B1 (=OATP2)	
	*5 (521T>C)	wt/*5
	OATP1B3 (=OATP8)	
	334T>G	wt/wt**
	MRP2	
	-24C>T	wt/wt
1249G>A	wt/mut	
3972C>T	wt/mut	

2. PRODUCT CHARACTERIZATION

Cell controls

Criteria	Method	Specification
Number of viable cells per vial	Microscopic observation	$\geq 8 \times 10^6$
Post-thaw viability (Day 0) ^a	Trypan blue exclusion test	$\geq 80 \%$
Cell density ^b	Microscopic observation 6 hours after thawing and seeding cells	$\geq 85 \%$
		 <p>X 100 – After thawing of cells and 6 hours of culture in the HepaRG[®] Thawing/Plating/General Purpose medium using ADD670</p>
Cell morphology	Microscopic observation 4 days after thawing and seeding cells, 50 % of typical hepatocyte- like cells are organized in well-delineated clusters with bright canaliculi-like structures.	 <p>X 100 – After thawing of cells in the HepaRG[®] Thawing/Plating/General Purpose medium using ADD670, and 3 days in the HepaRG[®] Maintenance/Metabolism Medium using ADD620</p>
Mycoplasma detection	Biochemical test	Negative
Microbial sterility	Under standard use conditions	No microbial growth detectable

^a: After thawing, differentiated HepaRG[®] viability is determined by Trypan blue (0.05% in PBS) exclusion test at 10 min post-thaw.

^b: Differentiated HepaRG[®] are seeded on collagen-coated, multi-well culture plates in the HepaRG[®] Thawing/Plating/General Purpose medium using ADD670 according to Biopredic's description and use guide for thawing, culture and use of cryopreserved differentiated HepaRG[®] (fduHPR116).

Vmax value of Phase I dependent activities (nmole/h/million cells)

Activity	Enzyme	Method	Specification
Phenacetin O-deethylase activity	CYP1A2	After thawing of cells in the HepaRG [®] Thawing/Plating/General Purpose medium using ADD670, cells were incubated in suspension for 1 hour at 37°C with the following test substrates: - phenacetin (200µM), - midazolam (50µM), - bupropion (100µM), - dextrometorphan (100µM). Metabolites formed were measured by LC-MS/MS.	≥ 0.2
Midazolam 1' hydroxylase activity	CYP3A4/5		≥ 1
Bupropion hydroxylase activity	CYP2B6		≥ 0.05
Dextrometorphan O-demethylase activity	CYP2D6		≥ 0.05

Activities are expressed as nanomole of metabolite formed/h/million cells. To convert to pmole of metabolite/min/million cells multiply the results by 16.66.

Induction of metabolic activity

Inducer	Enzyme	Assay	Method	Fold induction* Activity of treated cells/activity of control cells
				Specification
Omeprazole, 50µM	CYP1A2	Phenacetin O-deethylation	After thawing and 3 days of culture in the HepaRG [®] Thawing/Plating/General Purpose medium using ADD670, cells were incubated for 48h in the HepaRG [®] Induction medium using ADD640 with the test inducers. Cells were then incubated with the test substrates:	≥ 4
Phenobarbital, 1mM	CYP2B6	Bupropion-hydroxylation	- for 2 hours with midazolam (50µM) - for 4 hours with bupropion (100µM), - for 5 hours with phenacetin (200µM).	≥ 3
Rifampicin, 10µM	CYP3A4/5	Midazolam 1'-hydroxylation	The formation of metabolites was measured by LC-MS/MS.	≥ 4

* The fold induction in CYP1A2, CYP2B6 and CYP3A4/5 activity was determined as a measure of specific metabolite formation relative to vehicle control cells (un-induced).

Vmax value of uptake transporter activities (pmole/min/million cells)

Substrate	Transporter	Method	Specification
[³ H] MPP+, 250µM	OCT	After thawing, cells were incubated in suspension at 37°C in uptake medium with either [³ H]MPP+, or [³ H]taurocholate for 3 minutes. The uptake transporter activity (Vmax) is measured as cell-associated radioactivity with a liquid scintillation counter and expressed in pmol/min/million cells. *Taurocholate is also a substrate of OATP.	≥ 50
[³ H] Taurocholate*, 25µM	NTCP		≥ 6

3. CONDITIONS OF STORAGE AND DELIVERY

STABILITY, STORAGE, DELIVERY	
Stability and Storage	5 years in liquid nitrogen
Delivery	In dry ice or in liquid nitrogen If in dry ice, ensure shipment time did not exceed 48 hours
Use	Follow description and use guide for thawing, culture and use of cryopreserved differentiated HepaRG [®] cells (fduHPR116).

4. COMPANION PRODUCTS

CULTURE MEDIA*	
Denomination	Catalog number
HepaRG [®] Thawing/Plating/General Purpose Medium Supplement with antibiotics, to be combined with 100 mL of Basal medium	ADD670
HepaRG [®] Maintenance/Metabolism Medium Supplement with antibiotics, to be combined with 100 mL of Basal medium	ADD620
HepaRG [®] Induction Medium Supplement with antibiotics, to be combined with 100 mL of Basal medium	ADD640
HepaRG [®] Serum-free Induction Medium Supplement with antibiotics, to be combined with 100 mL of Basal medium	ADD650

**For details on use of supplements, consult the Description and Use Guide for thawing, culture and use of cryopreserved differentiated HepaRG[®] cells (fduHPR116).*

CULTURE SUPPORT	
Denomination	Catalog number
25 cm ² flask coated with collagen I*	FLA125
6-well plate coated with collagen I*	PLA135
12-well plate coated with collagen I*	PLA138
24-well plate coated with collagen I*	PLA137
48-well plate coated with collagen I*	PLA139
96-well plate coated with collagen I*	PLA136

**BPI proprietary coating process to ensure proper seeding and culture of the HPR116*

5. OTHER OFFERING OF THE HEPARG[®] CELLS

Denomination	Catalog number
Undifferentiated, cryopreserved HepaRG [®] cells for on-site propagation and differentiation. Please see on our HepaRG [®] web site www.heparg.com for the conditions to obtain the cell line.	HPR101
HepaRG [®] differentiated cells, cryopreserved, using a proprietary freezing process developed by Biopredic International that allows direct thawing and seeding of the differentiated cells without the need for post-thaw washing, centrifugation and counting steps.	HPR116NS
HepaRG [®] undifferentiated cells, cryopreserved with a sufficient quantity of culture medium for production of 50 plates within a 6 month period of time.	KIT901