

Lentiviral reprogramming of human umbilical cord-derived mesenchymal stromal cells towards hepatocyte-like cells

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Organisation

Name of the organisation Vrije Universiteit Brussel (VUB)

Department Pharmaceutical and Pharmacological Sciences

Specific Research Group or Service In Vitro Toxicology and Dermato-Cosmetology

Country Belgium

Geographical Area Brussels Region

SCOPE OF THE METHOD

The Method relates to	Human health
The Method is situated in	Basic Research
Type of method	In vitro - Ex vivo
Specify the type of cells/tissues/organs	umbilical cord

DESCRIPTION

Method keywords

human adult stem cells
hepatocyte-like cells
lentiviral reprogramming
umbilical cord

Scientific area keywords

in vitro liver model
Drug-induced liver injury (DILI)
human hepatocyte-like cells
liver enriched transcription factor
human adult stem cells

Method description

Human umbilical cord-derived mesenchymal stromal cells (hUC-MSCs) express several key liver-specific transcription factors as well as hepatic progenitor markers. However, they still lack the hepatocyte nuclear factors 1-alpha (HNF1a) and 4-alpha (HNF4a), indispensable for their reprogramming towards hepatocyte-like cells. This method

comprises the reprogramming of hUC-MSCs towards hepatocyte-like cells through HNF1a lentiviral over-expression. Whole genome microarray analysis revealed that the expression of the nuclear receptor retinoid X receptor (RXR) gamma and the nuclear transcription factor HNF4a, in HNF1a-transduced hUC-MSCs, was significantly upregulated compared to the control conditions. This expression was even higher than found in human hepatocytes. The same was observed for the uridine 5'-diphosphoglucuronosyltransferase (UGT) 1A family. Further, a significant upregulation was observed for alpha-foetoprotein (AFP), alpha1-antitrypsin (A1-AT), the phase I biotransformation enzymes cytochrome P450 (CYP) 1A2 and CYP2A6 and the drug transporter multidrug resistance protein (MDR) 1.

Lab equipment

Incubator ($37 \pm 1^\circ\text{C}$, $90 \pm 5\%$ humidity, $5.0 \pm 1\%$ CO_2/air);
Type 2 laminar airflow HEK293T cells;
Ultracentrifuge Water bath ($37 \pm 1^\circ\text{C}$);
Inverse-phase contrast microscope;
Pipettes;
Pipettors;
Colorimetric reverse transcriptase assay;
Human hepatocytes;
Affymetrix microarray technologies;
Partek Genomics Suite Software.

Method status

Still in development

PROS, CONS & FUTURE POTENTIAL

Advantages

Persistent expression of HNF1a transcription factor in hUC-MSCs;
Endogenous induction of HNF4a expression.

Challenges

Genomic integration of the lentiviral vector.

Modifications

Usage of a non-integrative reprogramming method e.g. mRNA transfection.

Future & Other applications

Generation of hepatocyte-like cells for the development of functional human liver-based *in vitro* models for pharmaco-toxicological purposes.

REFERENCES, ASSOCIATED DOCUMENTS AND OTHER INFORMATION

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