Expanded primary human liver sinusoidal endothelial cells (upcyte® LSECs) as a tool to complement hepatotoxicity studies

Timo Johannsen, Nikolett Nagy, Torge Evenburg, Astrid Noerenberg
upcyte technologies GmbH, Osterfeldstr. 12-14, D-22529 Hamburg, Germany

INTRODUCTION

Isolated liver cells, such as hepatocytes, liver sinusoidal endothelial cells (LSECs), Kupffer Cells and hepatic stellate cells, are frequently used to study hepatic metabolism, toxicity and diseases. The current in vitro culture models however, have several disadvantages, e.g. short culture longevity and artificial culture conditions that focus mainly on a single cell type in 2D culture. The use of primary cells in vitro is compromised by the limited quantity of cells that can be isolated from one donor, a lack of or very restricted proliferation capacity (hepatocytes and LSECs) and/or the change from a quiescent to an activated state (hepatic stellate cells).

Therefore, we investigated whether the transduction of proliferation-inducing genes could extend the lifespan of primary LSECs without losing their primary characteristics (so-called “upcyte® technology”) as previously demonstrated with hepatocytes (upcyte® hepatocytes).

RESULTS

Why LSECs?

LSECs constitute the sinusoidal wall and can be regarded as unique capillaries, which differ from other capillaries in the body, as they possess open pores or fenestrae lacking a diaphragm and a basal lamina underneath the endothelium. Fenestrae, arranged in so-called selectives “wim plates”, filter fluids, solutes and particles that are exchanged between the sinusoidal lumen and the space of Disse. Among the various substances that are known to be endocytosed by LSECs are proteins, glycoproteins, lipoproteins, and glycosaminoglycans.

Foreign soluble macromolecules and colloids are eliminated from the circulation mainly by receptor-mediated pinocytosis. There are only three different receptors, which have been functionally observed in LSECs and are responsible for uptake of a large number of different ligands:

(1) The Mannose receptor (MR) eliminates collagen waste molecules, preventing accumulation of intravascular clotting and recruits lysosomal enzymes for degradation.

(2) The hyaluronan/scavenger receptor (HA/S-R) plays an important role in the physiological mechanism contributing to maintaining homeostasis and preventing atherosclerosis by binding to type I and III procollagen and atherogenic molecules such as oxidized LDL and advanced glycation end products.

(3) The Fcγ-receptor (FcγR2BII, CD32B) takes up IgG coated particles and soluble IgG immune complexes (taken up especially in the liver). Other functions include interaction with viruses (e.g. HIV-1) [Smedsrod et al., 2004].

Generation of upcyte® LSECs

Transduction of primary cells with a defined cocktail of lentiviral vectors

Previously, a novel technique was reported which enabled the generation of non-transformed proliferating liver cells from primary human hepatocytes with maintained adult phenotype, so-called “upcyte® cells.” upcyte® cells start to grow from primary cells after transduction with a defined cocktail of lentiviral vectors carrying proliferation inducing genes. upcyte® cells have the ability to proliferate for additional cell doublings, depending on the cell type, without losing functional and phenotypic characteristics of mature cells.

Characterization of upcyte® LSECs: the cells express...

...primary endothelial cell markers

- vWF
- CD31
- Lamina reticulum

The morphology was comparable to primary cell LSECs. CD31 was present in all cells investigated and showed a typical membrane localization. vWF (von-Willebrand-factor) was evident as punctate structures that were more intense around the nucleus, indicating its presence in the endoplasmic reticulum.

- Tube formation
- Calcein AM staining
- LDL uptake
- UEA-1

The ability to form tubes in Matrigel™ was analyzed in upcyte® LSECs at different densities. A seeding density of 13,500 cells/ml was found to be optimal, since those upcyte® LSECs formed tubes from single cells. Uptake of lipid Ac-LDL [Alexa483-AcLDL], indicating the presence of the SR-A (Scavenger receptor A), a common endothelial cell receptor, was evident. Lectin (Ulex Europaeus LectinI) was strongly expressed.

...LSECs specific markers

- CD31
- vWF
- MR – FITC Ovalbumin
- HA/S-R – FITC-FSA
- FcγR – FITC-AGG

Immunoﬂuorescence staining of LSEC-speciﬁc receptors

The cells express the three major uptake receptors: there was a high expression of MR and FcγR in upcyte® LSECs. The staining of HA/S-R-receptor was visible, but weak and not present in all cells. A possible approach for better marker expression is elongation of culture time and medium optimization.

Co-culture with hepatocytes

Different studies demonstrate that LSECs are sensitive direct targets for early toxicity to APAP. 30 min after the administration LSECs become swollen and begin to lose their ability to endocytose FITC-FSA. After 2 h, prior to any effect on hepatocytes, fenestrae are lost [McCoy, 2006/2008].

LSEC alone were considerably more sensitive to APAP than when they were co-cultured with upcyte® hepatocytes (either in 2D or in Matrigel). ARPA is detoxified to glucuronide and sulfate metabolites by hepatocytes [Hewatt et al., 2007]. Therefore, this detoxification mechanism may have a protective effect for both hepatocytes and LSECs.

SUMMARY & CONCLUSION

In conclusion, we developed a novel technique which causes primary human LSECs to proliferate additional population doublings whilst still retaining an adult phenotype when cultured at confluence. The resulting cells called “upcyte® LSECs” retained important endothelial cell markers, such as CD31 and vWF, and exhibited functional uptake of LDL, as well as the ability to form tubes in Matrigel™, LSEC-specific uptake of ligands or the expression of the corresponding uptake receptors (MR, FcγR2BII and HA/S-R) could be detected. These data support that upcyte® LSECs are very uniquely and applicable to cell based assays as co-culture (e.g. with hepatocytes) and toxicity studies. Moreover, this technology allows for the generation of large batches of upcyte® LSECs enabling a reproducible and standardized experimental setting.