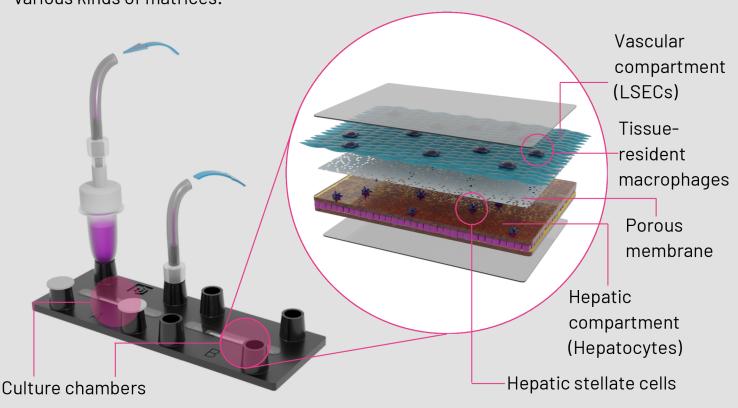


Our human immunocompetent liver-on-chip model is designed for investigating central drug metabolism and first pass toxicological studies. Further, recycling of biologicals and the infiltration of immune cells upon inflammatory triggers can be studied. The liver model comprises a vascular and a hepatic compartment. The vasculature is formed by liver sinusoidal endothelial cells and tissue-resident macrophages serving as Kupffer cell surrogates. The hepatic compartment comprises hepatocytes and optionally hepatic stellate cells embedded in various kinds of matrices.

# **LIVER** ON **CHIP**



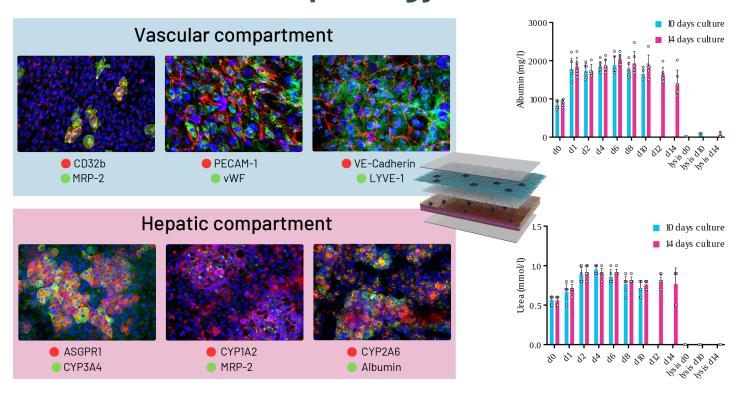
### **Applications**

- / Compound uptake
- / Compound metabolism
- / Toxicological studies
- / Immune cell infiltration
- / Disease modelling (e.g. Inflammation, fibrosis)
- / Infection modelling

### **Key features**

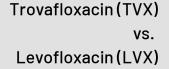
- / Up to 14 days stable model operation
- / Expression & functionality of major CYP enzymes
- / Immunocompetent (Integration of resident/ circulating immune cells)
- / Secretion of clinical parameters (Albumin, urea, LDH, ALT, AST)
- / Various configurations and different cell sources possible (primary cells or cell lines)

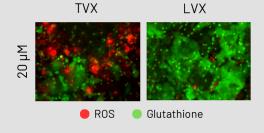
## Liver model - morphology

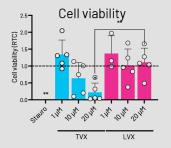


# **Application highlights**

#### Drug-induced liver injury (DILI)

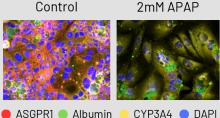


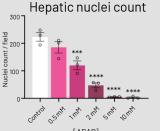






Acetaminophen (APAP)



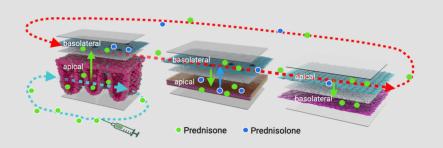




● ASGPR1 ● Albumin ● CYP3A4 ● DAPI (APAP)

#### Multi-organ models

Gut-liverplacenta model for ADME & PBPK





Paper







