

# In vitro coculture

Created on: 20-09-2022 - Last modified on: 09-02-2023

# **Contact person**

Amar van Laar

# Organisation

Name of the organisation Ghent University (UGent)

Department Food Technology, Safety and Health

Country Belgium

# **SCOPE OF THE METHOD**

The Method relates to	Animal health, Human health
The Method is situated in	Basic Research
Type of method	In vitro - Ex vivo
Specify the type of cells/tissues/organs	Caco-2 and HepG2 cells

# **DESCRIPTION**

# **Method keywords**

cell culture

Cell interactions

in vitro digestion

Metabolism

# Scientific area keywords

in vitro cell culture

liver metabolism

Diabetes research
Sugar metabolism
Fat metabolism

# Method description

This method is used to let cells interact for a better simulation of processes occurring in the body. The rational of the method is that monoculture experiments do not capture the complexity of *in vivo* interactions between different organs. In the most simple setup (published work Nutrients), Caco-2 cells were seeded on the (in our setup 24-well) transwell insert and HepG2 cells on the (in our setup 5\*104 cells in a 24-well) plate below. Caco-2 cells were seeded first. HepG2 cells were seeded in a separate plate upon differentiation of the Caco-2 cells and were put together when HepG2 cells formed a confluent layer. The Caco-2 cells were exposed to sugars and fatty acids, allowing digestion and transport to the HepG2 cells.

Acheter Dapoxetine 60mg sans Ordonnance en France Comprar Cialis Genérico Sin Receta en España - Tadalafilo 20mg Köpa Original Cialis Tadalafil 20mg utan recept

# Lab equipment

- Coculture plates with inserts
- Cell culture flow cabinet
- TEER machine or Lucifer Yellow

#### Method status

Published in peer reviewed journal

# PROS, CONS & FUTURE POTENTIAL

# Advantages

It increases the relevance of the cell culture model compared to the monoculture variants. Furthermore, it allows to study the effects of nutrients and medicinal compounds on organs that are only in contact with metabolites of these compounds *in vivo* (the method allows to incorporate digestion, absorption and metabolism).

# Challenges

The upper layer has to be confluent during the entire exposure (has to be checked before and after coculture and after the exposure). Since the cells should also be sufficiently fresh, timing is important. This timing depends on the cell type.

#### **Modifications**

This method could basically be used with any cell type and could even be used with more than 2 types of cells interacting (which requires some adjustment).

# **Future & Other applications**

The method can be applied for all type of cell research, not just in the field of diabetes.

# REFERENCES, ASSOCIATED DOCUMENTS AND OTHER INFORMATION

#### References

van Laar, A., Grootaert, C., Van Nieuwerburgh, F., Deforce, D., Desmet, T., Beerens, K., & Van Camp, J. (2022). Metabolism and Health Effects of Rare Sugars in a CACO-2/HepG2 Coculture Model. Nutrients, 14(3), 611.

Sadeghi Ekbatan, S., Iskandar, M. M., Sleno, L., Sabally, K., Khairallah, J., Prakash, S., & Kubow, S. (2018). Absorption and metabolism of phenolics from digests of polyphenol-rich potato extracts using the Caco-2/HepG2 co-culture system. Foods, 7(1), 8.

Lammi, C., Zanoni, C., Ferruzza, S., Ranaldi, G., Sambuy, Y., & Arnoldi, A. (2016). Hypocholesterolaemic activity of lupin peptides: Investigation on the crosstalk between human enterocytes and hepatocytes using a co-culture system including Caco-2 and HepG2 cells. Nutrients, 8(7), 437.

Scheers, N. M., Almgren, A. B., & Sandberg, A. S. (2014). Proposing a Caco-2/HepG2 cell model for in vitro iron absorption studies. The Journal of nutritional biochemistry, 25(7), 710-715.

Coordinated by





Financed by



