

## In vitro coculture

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### Organisation

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## SCOPE OF THE METHOD

<b>The Method relates to</b>	Animal health, Human health
<b>The Method is situated in</b>	Basic Research
<b>Type of method</b>	In vitro - Ex vivo
<b>Specify the type of cells/tissues/organs</b>	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 rationale 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\*10<sup>4</sup> 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

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