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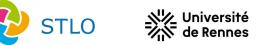
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Characterization of the mucus lining a co-culture model of Caco-2/HT29-MTX cells

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Introduction

- · Co-cultures integrating enterocyte-like cells (Caco-2) and goblet cells (HT29-MTX), are promising models of a functional epithelial
- · It is expected that such cultures are lined by a mucus layer but the spatial characteristics of this layer is poorly described.

Objectives

- To visualize mucins in a co-culture of Caco-2/HT29-MTX
- To set up a method of mucus characterization based on image analysis

Materials and methods

Caco2 + HT29-MTX (9:1) seeding density 2.105 cell/cm2



Incubation at 5% CO2 / 37°C 21 days



Differenciated monolayer fixation with 4% PFA

Open-source software ilastik



classification of pixels into three categories (corresponding to bright staining mucin clusters, diffuse mucin staining, background)



Ten 3-D images (80 stacked images on average) from 2 independent wells



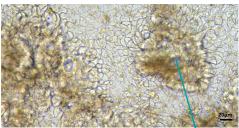
Confocal microscopy Zeiss-LSM 880



Staining: F-actin with phalloidin-rhodamine → cytoskeleton Sialic acid with WGA-Alexa 488 → mucins

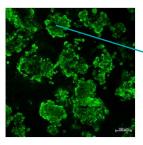
Results

1- Uneven distribution of the mucus



Observation of cultures by optical microscopy (X20)

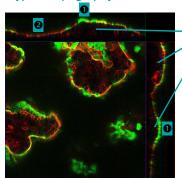
Cells are confluent and they form a monolyer but some cells are grouped and lined with a blurry material.



Mucins are abundantly expressed as clusters

Observation of cultures after staining of sialic acid

2- A typical topography



Observation of one stack of a 3-D reconstituted image

The culture is not planar. Domes (visible on z-views) are present. Height up to 135 µm.

Mucins are present mainly at the apical sides of domes (see bright staining: sites of secretion) but also between domes (see 2 diffuse staining: secreted mucins spreading over the culture, loose mucus)

3- Development of an image analysis workflow to characterize the mucus structure

Machine learning: annotation of pixels representative of bright / diffuse staining on a subset of images, followed by automatic classification on all images.

Extraction of quantitative data

	Dense mucin clusters	Loose mucus
Average thickness	6.9 µm	21.2 µm
Average volume	183000 µm ³	7502000 μm ³

On-going: segmentation and classification of cells

Conclusions

- · Staining of sialic acid and confocal microscopy enable three-dimensional visualization of the mucus layer
- · The mucus on the co-culture model is heterogenous in density
- The protocol of image analysis is suitable to extract quantitative data on the mucus layer (NB: lower thickness than in vivo data)

These methodological developments open the way to studying the impact of digested food constituents or digested foods on the intestinal mucus

(1) Béduneau et al. 2014. A Tunable Caco-2/HT29-MTX Co-Culture Model Mimicking Variable Permeabilities of the Human Intestine Obtained by an Original Seeding Procedure. Eur J Pharm

(2) Berg et al. 2019. A. Ilastik: Interactive Machine Learning for (Bio)Image Analysis. Nat Methods 16 (12), 1226–1232.