Quantification of fiber production by preadipocytes



Adipose Tissues • 2D Cell Models • High Content Screening • Automatic Detection • Dermatology

YOUR NEEDS

OUR SOLUTIONS

- Monitor pre-adipocytes differentiation
- Monitor the fibro-inflammation process (collagen, fibronectin)
- Fully automatic imaging and image analysis
- Robust data, short delay and cost-saving

General Procedure



D | V A Cell isolation, culture and labelling supported by DIVA Expertise

Image acquisition:

- Acquisition done with structured light or confocal microscopy
- Several fields of view to maximize the amount of data
- Image stack for each field of view to get each cellular structure on their focal plane

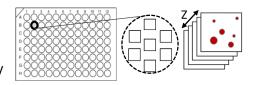
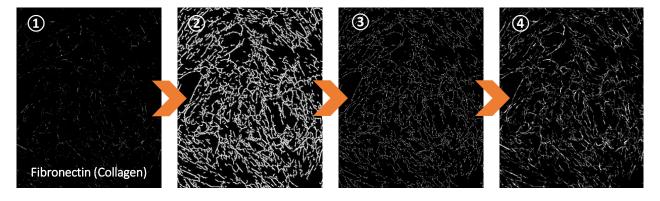


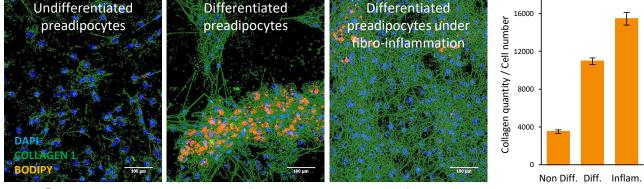
Image processing:

- (1) Detail of an image after acquisition with spinning disk confocal microscopy
- (2) Denoising and segmentation of every fiber
- (3) Refining the segmented object and calculating the criteria: total fiber length, thickness of fibers
- (4) Automatic generation of illustrative images for every field of view



Application example

During weight gain, adipose tissue is characterized by the presence of low-grade inflammation and matrix remodelling. This leads to a greater tissue stiffness and to resistance to weight loss.



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