

# Optical digital biopsy; the physical photo of pigmentary epithelium

## Introduction

In the secuentionation of image by pixelometry of isolated cells, studing the pigmentary epithelium of retina we can appreciate three areas: the nucleus and the cytoplasm corresponding to cells of the pigmentary epithelium, physical, alive, in situ, achieved through the sequencing of optical coherence tomography (OCT) images (Figure 1).<sup>1-16</sup>

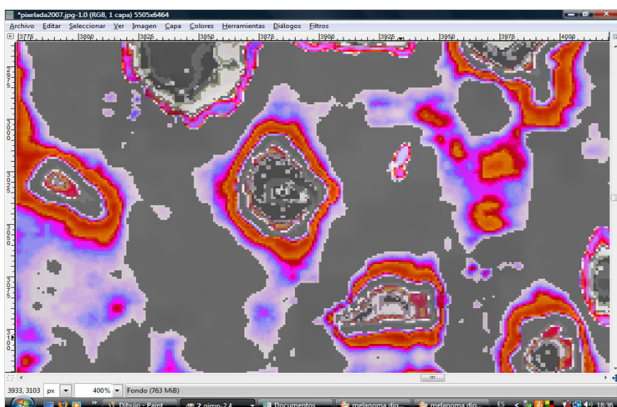
The steps for this determination can be summarized in:

1. Determine the total number of pixels in the image.
2. Determine the number of pixels contained within the cytoplasmic membrane.
3. Determine the number of pixels contained in the kernel.
4. Determine the number of pixels in the cytoplasm.
5. Determine the ratio of the number of pixels in the nucleus to those in the cytoplasm.

This procedure aims to adapt the image for later analysis. Therefore, the following operations have been carried out:

1. Band removal
2. Smoothing the image
3. Rotation
4. Extraction of the region

This procedure has allowed us to: first extract a band to be able to then operate on it, this is convenient as it is necessary within the process convert the image to binary image. Then, soften the image using a median filter; this because the totality of the detail that contains the image, since they behave as noise for the end pursued. And finally the image has been rotated in order to place it in its better angle, in such a way, that the extraction of the region of interest contains pixels that represent it optimally. The adequate selection of the region of interest allows operating only on the pixels that contain relevant information.



**Figure 1** Physical cells of the pigment epithelium of the retina. Photo obtained from an OCT after the sequencing of images, that is, without an invasive component.

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Obtaining the required parameters for work on the image has been made by segmenting the image. This procedure has allowed us to subdivide the images into their constituent objects: nucleus and cytoplasm, taking advantage of the fact that the gray levels of the nucleus and the cytoplasm are differentiable. The segmentation of the images was done by the thresholding method, using information provided by the histograms of the levels of gray of the processed images and the corresponding averages, as

it shows in the following diagram:

## Obtaining results

The determination of the number of pixels represented by the cell as a whole, It was done using as a threshold the average of the total pixels of the image. This it has allowed us to label each pixel as belonging to the background or the cell. To obtain the pixels corresponding to the nucleus, the threshold was used gray level values corresponding to the left inflexion points and right of the curve to the left of the histogram; and, for the pixels corresponding to the cytoplasm, the left and right inflexion points of the curve to the right of the histogram. The methodology is repeatedly pixelometric, pixelográfica and could be added pixeloarquitectural. Each binary image, which taken to the subcellular level can be a protein has a code based on pixels, with so many probabilities that it exceeds the genome.

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## Conflicts of interest

Author declares that there are no conflicts of interest.

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