

The SIPAKMED database is publicly available and it can be used for experimental purposes with the request to cite the following paper:

M. E. Plissiti, P. Dimitrakopoulos, G. Sfikas, C. Nikou, O. Krikoni, A. Charchanti, SIPAKMED: A new dataset for feature and image based classification of normal and pathological cervical cells in Pap smear images, IEEE International Conference on Image Processing (ICIP) 2018, Athens, Greece, 7-10 October 2018.

Description of Cell Features

In each image of our database, the boundaries of the regions of interest, i.e. the area of the cytoplasm and the nucleus of the cells, were manually defined by expert observers. The coordinates of the contour of each area are provided for both the images of cell clusters (in *im_(class)* directories) and the isolated cell images (in the CROPPED subdirectory of each class).

In every region of interest, we calculate 26 features concerning the intensity (average intensity, average contrast), the texture (smoothness, uniformity, third moment, entropy in all three color channels), and the shape (area, major and minor axis length, eccentricity, orientation, equivalent diameter, solidity and extent). These features were calculated for both the region of the nucleus and the cytoplasm of each cell and they are stored in 10 tables (cytoplasm and nuclei features for each class) of 28 columns (the two added fields denote the number of the image and the cell correspondingly). The feature tables are stored in Feature_CELL directory and they have the following structure:

| Column | Feature | Column | Feature |
|--------|--------------------------|--------|----------------------------|
| 1 | Index of the Image | 15 | Uniformity in RED |
| 2 | Index of the Cell | 16 | Entropy in RED |
| 3 | Area | 17 | Average intensity in GREEN |
| 4 | MajorAxisLength | 18 | Average contrast in GREEN |
| 5 | MinorAxisLength | 19 | Smoothness in GREEN |
| 6 | Eccentricity | 20 | Third moment in GREEN |
| 7 | Orientation | 21 | Uniformity in GREEN |
| 8 | EquivDiameter | 22 | Entropy in GREEN |
| 9 | Solidity | 23 | Average intensity in BLUE |
| 10 | Extent | 24 | Average contrast in BLUE |
| 11 | Average intensity in RED | 15 | Smoothness in BLUE |
| 12 | Average contrast in RED | 26 | Third moment in BLUE |
| 13 | Smoothness in RED | 27 | Uniformity in BLUE |
| 14 | Third moment in RED | 28 | Entropy in BLUE |

Description of Image and Deep Features

The trained VGG-19 Convolution neural network was used as a feature extractor by feeding it an input cell image and using the intermediate layer pre-activations to construct a feature vector.

File ConvFeatures.mat :

In this file there are features extracted by using the pre-activations of the last convolutional layer (layer conv5).

These features are then aggregated by sum-pooling to form a feature vector of size 512 to describe each cell image. For every cell there are 3 types of features: raw features 512 size, compressed features 256 size, compressed features 32 size

File FcFeatures.mat:

In this file there are features extracted by using the pre-activations of the first fully-connected layer (layer fc6).

These features are of size equal to the number of neurons in the corresponding layer, i.e. 4096. For every cell there are 3 types of features: raw features 4096 size, compressed features 256 size, compressed features 32 size.

Both files shared the following structure:

- Folds 1-5 (5-fold cross-validation)
- Cell Classes (Superficial-Intermediate, Parabasal, Koilocytotic, Dyskeratotic, Metaplastic)
- Size of cell features (Raw values, PCA 256, PCA 32)
- Feature for each cell (Cell cluster image id, cell id, [features])