

An Image based Cervical Cancer Segmentation and Detection Using ANFIS Classifier

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Abstract—An image-based detection of cervical cancer at early stages is used by an Interior point algorithm. Soft tissue sarcomas are hard to spot because they can promote anywhere in our body. Regular Pap smear screening is the successful attempt of medical science and practice for the early detection of cervical cancer. Manual section of the cervical cells is time-consuming, laborious and error-prone. They may also feel firm or resolute and might be fixed to the tissue. By bounding, region method shape of the tissue can be found. Firm or solid level of tissue can be found by comparing consecutive Interior pointing algorithm. Interior pointing algorithm refers to more parameters like pH level of the tissue, deep gradient level, Non- gradient value of tissue, etc. In proposed method deals with these parameters to ensure the possibility of soft sarcoma tissue. This may useful to find cancer at an earlier stage, it may cause to avoid radiation or chemotherapy treatments. As a result, of detection process will display the normal or abnormal tissues, in that abnormal tissue may turn as cancer.

Index Terms— Segmentation, Feature Extraction, Classifier, Sensitivity, Specificity, Accuracy.

I. INTRODUCTION

Cervical cancer in women, is the most average cancer worldwide, next only to breast cancer. In India, cervical cancer is the most regular woman-related cancer, followed by breast cancer. Every year cervical cancer is diagnosed in around 500,000 women globally and is authentic for more than 280,000 deaths annually. There is a bumper variation in the incidence of cervical cancer across the globe.

Cancer detection has always been a major release for the pathologists and medical practitioners for Process planning. The manual identification of cancer from microscopic biopsy images is deference innature and may vary from a droit to adroit depending on their expertise. In other factors which include lack of special and exact quantitative measures to classify the biopsy images as normal or cancerous one.

Cervical cancer could be exceedingly preventable in many nations because of the wide accessibility of the screening test and immunization to a verthumanpapillomainfection (HPV) disease. HPV is the primary cause of cervical cancer and has been recognized in 90 to 100% of cervical cancer growth incidents [WHO, 2000].

In 32% of the Indian population gets cancer at some capeduring their lifetide. Cancer is one of the sweeping diseases in India which has an obligation to maximum mortality with about 0.3 million deaths per year. The possibility of tenor from cancer is increased due to recently United progression in medicine and engineering. The assortment of dealing with cancer totally depends on its level of malignancy. Medical professionals use numerous performances for the detection of cancer. These techniques may include diverse imaging modalities such as X-ray, Computed Tomography (CT) Scan, Positron Emission Tomography (PET), Ultrasound, and Magnetic

Resonance Imaging (MRI) and pathological tests such as urine test and blood test. In histopathology, the cancer detection alleviates normally consist of categorizing the image biopsy into cancerous or non-cancerous. In microscopic biopsy twin investigation doctors and pathologists observe many of the irregularities and classify the sample based on various characteristics of the cell nuclei such as color, icon, Magnitude, and proportion to the cytoplasm. High determination microscopic biopsy affords dependable information for differentiating irregular and standard tissues.

II. DETECTION AND DIAGNOSIS OF CANCER

For the recognition and analysis of cancer from microscopic biopsy images, the histopathologists normally look at the definite features in the cells and tissue shape. These several communal features used for the recognition and analysis of cancer from the microscopic biopsy images include the shape and size of cells, icon, and magnitude of cell nuclei, and distribution of the cell.

III. IDENTIFICATION OF PARAMETER

A. Shape and Size of The Cells

It has been observed that the overall outline and scope of cells in the tissues are mostly normal. The cellular structures of the cancerous cell's strength are either larger or shorter than standard cells. The standard cells have even shapes and functionality. Cancer cells usually do not a role in a useful way and their shapes are often not even.

B. Size and Shape of The Cell's Nucleus

The outline and size of the nucleus of a cancer cell are often not normal. The nucleus is reorganized in the cancer cells. An image of the cell aspects like an omelet, in which the basilic yolk is the nucleus and the surrounding white in the cytoplasm. The nuclei of cancer cells are grander than the normal cells and deviated from the center of the mass. The nucleus of the cancer cell is darker. The segmentation steps mainly concentrations on separation of regions of interests from background tissues as well as the departure of nuclei from the cytoplasm.

C. Distribution of The Cells in Tissue

The function of each tissue is contingent on the delivery and arrangements of the normal cells. These adjectives of microscopic biopsy descriptions have been elaborate an outline and morphology-based landscapes, textured landscapes, etc., which is more organically interpretable and clinically significant.

D. Figures and Tables

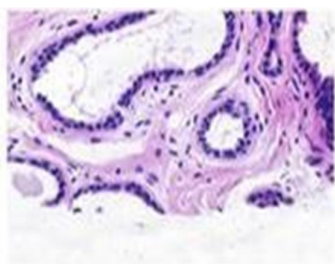


Figure 1. Original

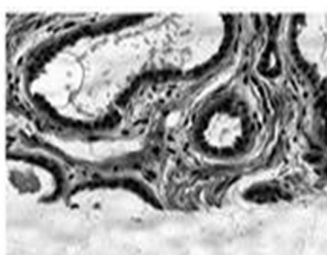
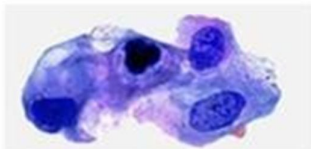
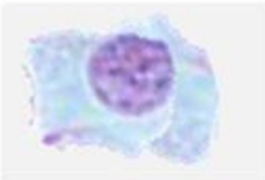


Figure 2. Enhanced Biopsy Image

The main aim of this paper is to design and progress a frame work and a software tool for computerized detection and classification of cancer from microscopic biopsy images. This paper focuses on picking a fitting method for each design stage of the frame work after making a comparative analysis of the various frequently used methods in each category. The various steps elaborate in the estimated methodology include

the development of microscopic images, segmentation of circumstantial cells, landscapes extraction, and lastly the classification.

TABLE I. DIFFERENCE BETWEEN NORMAL AND CANCEROUS CELLS

Normal cells	Cancerous cells	Description of cancerous cells
		Large and variably shaped nuclei
		Many dividing cells and disorganized arrangements
		Variations in size and shape of nuclei
		Loss of normal feature (shape and morphology)

IV. METHODS AND MODELS

For the enhancement of the microscopic biopsy images, the contrast limited adaptive histogram equalization approach is used. In segmentation of background, cells mean segmentation algorithms are used. Finally, the performances of the classifiers are judge using well-known parameters and from results and analysis. It is observed that the fuzzy KNN based classifier is performing better for the elected features set.

V. EXISTING METHOD

Current screening for cervical cancer is highly dependent upon the type of resources available in the population being screened. In high-resource settings, routine screening includes pap smears throughout a lifetime to evaluate for cervical dysplasia. Depending on the age of the patient, an evaluation may or may not include screening for high-risk HPV. If abnormal cytology is detected, then the patient may either have more frequent pap smears. This type of screening allows for close evaluation of the cervix and early excision of high-grade dysplasia in appropriate cases.

The method comprised of clear-cut segments in which information identified with the author, year of publication, category and issues to cervical cancer counteractive action benefit use. After careful perusing of the

article, information identified related cervical cancer issues and was entered into a matrix containing columns reflecting the issue categories. However, the rasping assumption used by the pixel classification schemes that the pixels of the entire image are distributed in two discrete classes, such as nuclei pixels and other pixels. It is not appropriate for Pap smear images, which exhibit great complexity and the section of all the pixels of the image. Later, a two-stage fully automated method for the accurate determination of the nuclei boundaries in Pap smear images, which may contain both isolated cells and cell clusters. More specifically, in the first step, nuclei markers are detected with a procedure based on reconstruction for the extraction of the areas of regional minima in the image, which usually correspond to nucleilocations.

A. Disadvantages

1. Highratio:

False positive (FP): Healthy people incorrectly diagnosed as sick False negative (FN): Sick people incorrectly diagnosed as healthy.

2. It can take large steps.

3. It is effective on some problems with non-smooth constraints.

It's a large-scale algorithm. (requires more data/constraints)

VI. PROPOSED METHODOLOGY

Fuzzy Cluster is a method where data points can belong in more than one group (cluster). Clustering divides data points into groups based on the similarity between items and the similarity between an item in a set. Clusters are identified via similarity measures. Adaptive histogram (AH) equalization is a contrast enhancement method in computer image processing technique. Used to enhance the definitions of edges in each region.

IPA contains the following processing steps to find cancer cells at an earlier stage in an effective manner. Gabor filter, GLCM, Fuzzy Cluster, Adaptive histogram, Neural network classifier are the processing steps involved in IPA.

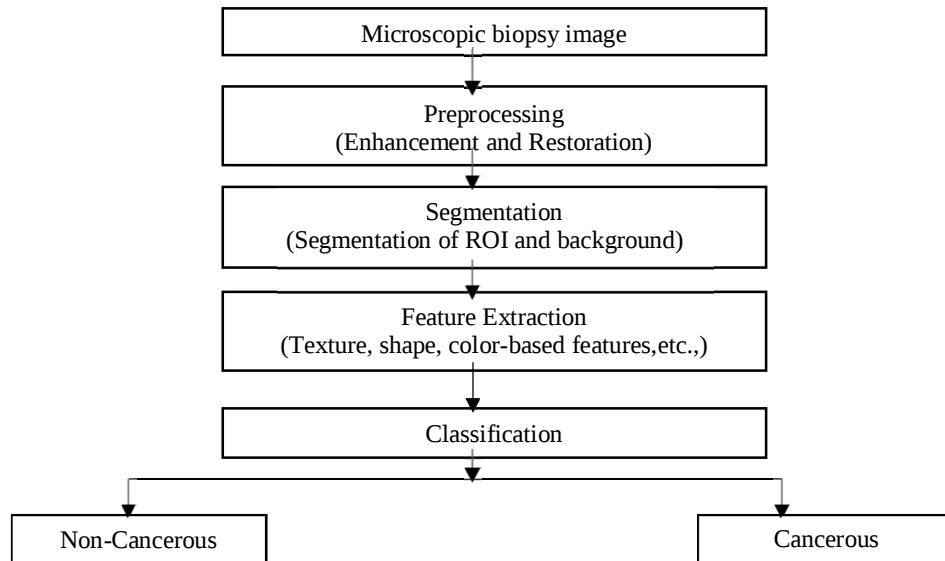


Figure 3. Block Diagram

Finally, the K-nearest neighborhood (KNN), fuzzy KNN, and support vector machine (SVM) based classifiers are examined for classifying the standard and cancerous biopsy images.

A. Enhancement

The main determination of the preprocessing is to remove specific dilapidation such as noise reduction and contrast enhancement of the region of interests. The contrast limited adaptive histogram equalization method is used for the improvement of microscopic biopsy images.

B. Segmentation

Several separation methods have been modified for cytoplasm, cell, and nucleus segmentation from microscopic biopsy images like threshold based, region-based, and clustering-based algorithms. However, the collections of separation methods depend on the type of features to be conserved and extracted. For the segmentation of the region of interest, the ground truth of the images is manually cropped and created from the histology data set. The k-means clustering based segmentation algorithm is used because of the preservation of the desired information. From the obtained results through experimentation it is observed that the clustering-based algorithm specifically k-means based method is the best suited for microscopic biopsy images. For testing and experimentation purpose, twenty microscopic biopsy images available from the histology data set were used. These images were randomly selected for segmentation. The ground truth images are manually created by cropping the region of interest.

Finally, the region of interest segmented image of microscopic biopsy is associated with ground truth images for the measurable evaluation of various segmentation methods for all 20 sample images from the histology data set. The performance of the various segmentation approaches such as K-means, fuzzy C-means, texture-based segmentation, and color-based segmentation is evaluated in terms of various popular parameters of segmentation measures. These parameters include accuracy, sensitivity, specificity, false positive rate, probability random index.

C. Equations

Probability Random Index: Probability random index is the non-parametric measure of goodness of segmentation algorithms. A random index between Stets and ground truths was estimated by summing the number of pixel pairs with same label and number of pixel pairs having various labels in both S and G and then dividing it by a total the number of pixel pairs. Given a set of ground truth segmentation G_K , the PRI is used. In such that C_{ij} is an event that describes a pixel pair (i,j) having the same label in the test image S_{test} .

$$PRI(Stest, Gk) = \frac{1}{(N/2)} \sum_{i,j \in S} [C_{ij}P_{ij} + (1 - C_{ij})(1 - P_{ij})] \quad (1)$$

VII. FEATURE EXTRACTION

After segmentation of image structures are extracted from the regions of interest to perceive and grade potential cancers. The features are mined at the tissue level and cell level of microscopic biopsy images for improved predictions. To improved capture the shape information, we use both region-based and contour-based method to extract anti circularity, area irregularity, and contour irregularity of nuclei. There are three structural features to replicate the irregularity of nuclei in biopsy images. The cellular level feature emphasizes on measuring the properties of individual cells without considering spatial dependency between them. In biopsy images for a single cell, the shape and morphological, textural, a histogram of concerned with gradients and wavelet features are mined. The tissue level features to measure the supply of the cells across the tissue; for that, it primarily makes use of either the spatial dependency of the cells or the gray level dependency of the pixels. Based on these features, some important shape and morphological based features are explained as follows.

A. Nucleus Area (A)

The nucleus area can be signified by a nucleus region containing a total number of pixels; it is shown in

$$A = \sum_{i=1}^n \sum_{j=1}^m B(i, j) \quad (2)$$

Where A is nucleus area and B is a segmented image of i rows and j columns.

B. Brightness of Nucleus

The normal value of an intensity of the pixels belonging to the nucleus region is known as nucleus brightness.

VIII. RESULT

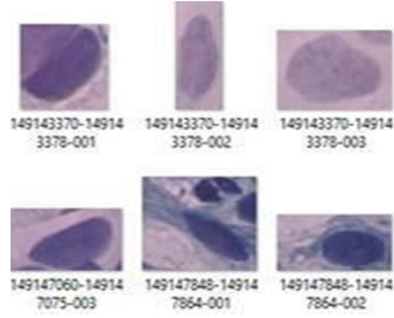


Figure 4. Input Image

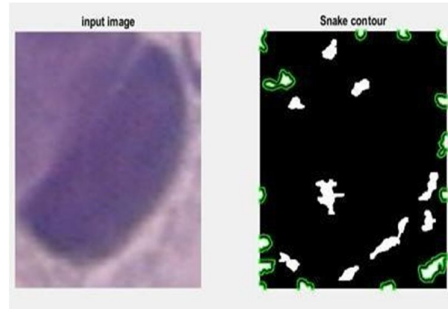


Figure 5. Select an Input Image

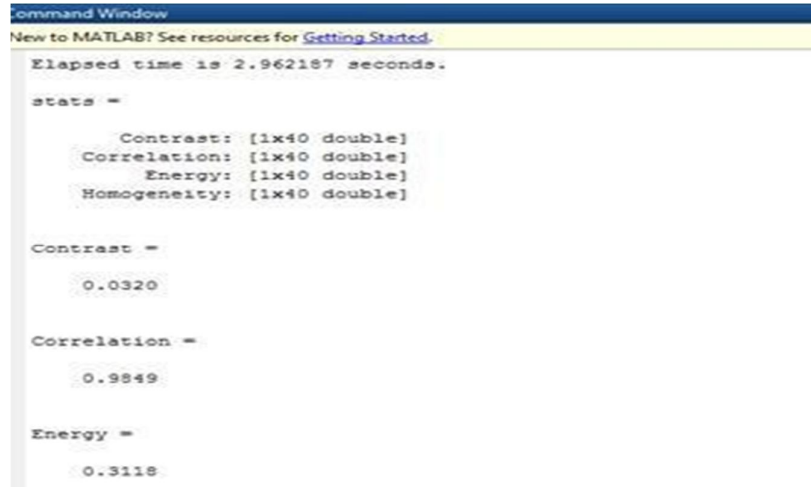


Figure 6. Parameter Calculation

IX. CONCLUSION

An automated detection and classification procedure is a supply for detection of cancer from microscopic biopsy images using a clinically remarkable and living interpretable set of features. The proposed analysis is based on tissues level microscopic observations of cell and nuclei for cancer detection and classification. For the development of microscopic biopsy, images contrast partial adaptive histogram equalization-based method was used. For segmentation of images -means clustering technique is used. After segmentation of images, a total of 115 hybrid sets of landscapes. They are mined for 2828 sample images taken from the database.

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