Automated Segmentation and Shape Tracking of Fluorescent Cancerous Cells by Wavelet Otsu model

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Abstract

In recent days due to the urbanisation and socialism among people the diseases such as cancers are very common and they need to be treated as early as possible. Reports suggest that 60% of the people are suffering with various traits of cancer. Cancer is caused because of the splitting and merging of infected cells So in order to detect the vulnerability of the disease we have to study the entire nature of the cells. The main goal of the project is to demonstrate that the proposed tracking scheme is more accurate and significantly faster than the other state-of-the-art tracking by model evolution approaches. The crucial tasks are, in particular, segmenting, tracking, and evaluating movement tracks and morphological changes of cells, sub-cellular components, and other particles. In recent days the evolving shape of the fluorescent cells, the mechanisms of cell motility and their regulation is an important challenge in biomedical research such as the immune response, embryonic development, or tumoro-genesis. The system proposed here not only tracks and segments the cells, it also detects the percentage or vulnerability of the disease by using classes of segmentators. This can be obtained by wavelet otsu model in which the classified levels of cells are found out and they are used for isolating the infected cells. This approach is carried out in rat adipose mesenchymal stem cells, or the carcinogenic cells.

Keywords: Fluorescent cells, tumorogenesis, wavelet, histogram, vulnerability

1. Introduction

In computer vision, image segmentation is the process of partitioning a digital image into multiple segments (sets of pixels, also known as super pixels). The applications of any computer vision system require cell tracking to be fast, affordable and, most importantly, precise and robust. Segmentation of the cells is important for
allowing different regions of the body to develop differentially for different uses. The process of detecting and tracking biological features such as cancerous cell growth and nuclei is complicated by the fact that they constantly change their shape. Shape changes happen both continuously as the biological features grow and discontinuously as they divide or die. This can be done effectively if they are fluorescent. So fluorescence microscopy technique has to used for segmenting and tracking of cells. The extraction of fluorescence time course data is a major bottleneck in high-throughput live-cell microscopy. An extendible framework is based on the open-source image analysis, which aims in particular at analyzing the expression of fluorescent reporters through cell divisions. The ability to track individual cell lineages is essential for the analysis of gene regulatory factors involved in the control of cell fate and identity decisions.

2. Existing Methodologies

The idea raised for the development of this system for replacing the chanvese model [1] by using morphological operations. With the invention of fast cell segmenting and shape tracking methodologies and recent increase of computer power and decrease of man power in the field of medical science especially for tumorogenesis, it has become very common to see a cell tacking and segmenting software that will analyse the disease nature in detail with more ease and accuracy.

2.1 Chan–Vese model

The Segmentation And Shape Tracking of Fluorescent cells is initiated by using the Chan-Vese model [1]. The methodologies such as Cell tracking, level set and graph cut Optimisation are used efficiently along with the Chan vese model in which the system is used so as to track the multiple cells simultaneously if the number of frames is also maximum. It is a fast and robust approach to tracking the evolving shape of whole fluorescent cells in time-lapse series. It has 2 steps. First, coherence-enhancing diffusion filtering is applied on each frame to reduce the amount of noise and enhance flow-like structures. Second, the cell boundaries are detected by minimizing the Chan-Vese model in the fast level set-like and graph cut frameworks. The major advantage of this method is Ellapsing of frames is reduced, and also multiple tracking is enabled. There are several limitations that we intend to address in future work to improve the overall performance of the proposed tracking scheme. First, a manual separation of cells clustered in the first frame is required to track each of them correctly over time. This complicates the use of the proposed tracking scheme in experiments with high density of tightly packed cells. Furthermore, coherence-enhancing diffusion filtering takes up to about 85% of the total execution time. Therefore, a different choice of the filtering technique would make the proposed tracking scheme significantly faster and more suitable for high-throughput applications.

2.2 Coupled Active Segmentation

Segmenting and tracking fluorescent cells in dynamic 3-D microscopy with coupled active surfaces a semi automatic segmentation and tracking method designed to
enable quantitative analyses of cellular shape and motion from dynamic three-dimensional microscopy data. The method uses multiple active surfaces with or without edges, coupled by a penalty for overlaps, and a volume conservation constraint that improves outlining of cell/cell boundaries. Cell migrations and deformations play essential roles in biological processes, such as parasite invasion, immune response, embryonic development, and cancer. Its main advantages are robustness to low signal-to-noise ratios and the ability to handle multiple cells that may touch, divide, enter, or leave the observation volume.

3. Proposed Scheme
The proposed system which uses a morphological technique is applied on the fluorescent cells so as to get a clear cut segmented image. The main advantage of the system is it is fully automatic. The frames that are extracted from the video should undergo various preliminary screening techniques. This system eliminates the existing traditional methodologies on segmenting whereas it utilizes some of the techniques used for segmenting traditionally. In this system the cells are segmented and shape tracked based on morphological operations. The morphological operations include the morphological mathematical functions. For this the wavelet otsu paradigm is used in where the image or frame is filtered, segmented and finally they are made to undergo so many stages such as the classes whereas in every classes they should be transformed by means of iterative scannings. Otsu suggested minimizing the weighted sum of within-class variances of the foreground and background pixels to establish an optimum threshold. The Otsu method gives satisfactory results when the numbers of pixels in each class are close to each other. The Otsu method still remains one of the most referenced thresholding methods. This methodology makes use of Thresholding in cells which is entirely new in the domain of medical imaging whereas the cells are fragmented only by means of frequencies and not on thresholding criterians. The class means which will be detected is also used for finding effective splitting and spreading stage of infected and uninfected cells.

4. Experimental Analysis
An object can be easily detected in an image if the object has sufficient contrast from the background. We use edge detection and basic morphology tools to detect a prostate cancer cell. Fluorescent microscopy allows the direct visualization of molecules at the subcellular level, in both live and fixed cells. Molecules of interest are marked with either green fluorescent protein (GFP), another fluorescent protein, or a fluorescently-labeled antibody. The shape tracking methods can be decomposed in two tracking types: the line tracking and the junction tracking. Next the tracking process depends on used tracking elements. These ones are of two types: pixel element and surface element. The experiment is carried out by these methods. The video that got splitted into frames are allowed to pass through various stages by taking the frames.

4.1 Preprocessing
The preprocessing of the images or frames obtained by the video are performed with
the AHE algorithm. Adaptive histogram equalization (AHE) is a computer image processing technique used to improve contrast in images. It differs from ordinary histogram equalization in the respect that the adaptive method computes several histograms, each corresponding to a distinct section of the image, and uses them to redistribute the lightness values of the image.

4.2 Anisotropic diffusion filtering
Anisotropic diffusion resembles the process generates a parameterized family of successively more and more blurred images based on a diffusion process. Each of the resulting images in this family are given as a convolution where the width of the filter increases with the parameter. As a consequence, anisotropic diffusion is nonlinear and space-variant transformation of the original image which mainly focuses on removing noise. This diffusion process is a linear and space-invariant transformation of the original image. Anisotropic diffusion is a generalization of this diffusion process: it produces a family of parameterized images, but each resulting image is a combination between the original image and a filter that depends on the local content of the original image.

4.3 K-Means Clustering
Image after running k-means with $k = 16$. Note that a common technique to improve performance for large images is to downsample the image, compute the clusters, and then reassign the values to the larger image if necessary. The K-means algorithm is an iterative technique that is used to partition an image into $K$ clusters. The basic algorithm is: Pick $K$ cluster centers, either randomly or based on some heuristic. Assign each pixel in the image to the cluster that minimizes the distance between the pixel and the cluster center. Re-compute the cluster centers by averaging all of the pixels in the cluster. Repeat steps 2 and 3 until convergence is attained (i.e. no pixels change clusters).

4.4 Morphological Analysis
Morphological analysis is important to study the cellular organization and the physiological state of the cells, and thus it can be commonly used as a qualitative and quantitative measure of various biological assays. Analysis of cell morphology remains increasingly important, as the image analysis aids in the detailed examination of microscopic cells; study of cell behavior, and also provides quantitative measure of its curvature, area, perimeter, eccentricity and additional metrics of nuclear morphology for large populations of cells. Analysis of the cells based on their morphological differences is applied to study the differentiation of stem cells, cancer cells, and in hematology. A wide variety of image analysis software packages have been developed that helps to convert to the microscopic images into more relatively quantitative measurements. In order to control the growth of the infectious cancerous cells during, a fully-automated sampling and analysing system has been developed.
Figure 1: Input image when segmented through various phases of our project

Figure 2: Initial and current frames

Figure 3: Overlapped frames with segmented regions showing variations
5. Conclusion
In this paper, we have presented a fast and robust approach to tracking whole fluorescent cells in time-lapse series. The proposed tracking scheme combines CED filtering with diffusion. It allows simultaneous tracking of multiple cells over time by applying a topological prior that exploits the object indication function. The experimental evaluation was performed on 2-D and 3-D time-lapse series of rat adipose-derived mesenchymal stem cells and human squamous cell carcinoma cells, respectively. It clearly verified the improved accuracy up to about 9%. Thus, they could be preferred in studies focused on local morphological changes in the cell shape, in which as-accurate-as-possible cell boundaries are the most crucial task. This complicates the use of the proposed tracking scheme in experiments with high density of tightly packed cells.

6. References