

UNIVERSITY OF KWAZULU-NATAL
COLLEGE OF AGRICULTURE, ENGINEERING AND SCIENCE



**A Fluorescence Image Segmentation Model Based
on Improved Discrete Active Contours**

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*A thesis submitted in fulfillment of the requirements
for the degree of Master of Science in Engineering*

in the
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COLLEGE OF AGRICULTURE, ENGINEERING AND SCIENCE

Abstract

Master of Science in Engineering

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For multiple fields, from biosciences and pharmaceutical drug development to semiconductor and geology research and beyond, fluorescence microscopy has been established as an essential tool which allows for the visualisation of diminutive organisms and objects that escape visibility from the naked eye. The consequent insight into this world has given us a view from an indispensably unique perspective which has since played in vital role in enriching the livelihood of mankind.

Fluorescence microscopy is not the sole player in this venture. It shares the limelight with another important and just as critical field, and that is image segmentation. However, the course between fluorescence images and extracting a meaningful segmentation is laden with barriers and hurdles such as noise, extreme low contrast, non-uniform illumination, etc. The aim of this research is to better understand fluorescence image properties and to leverage this knowledge to enhance and develop domain-specific segmentation methods, using discrete combinatorial techniques, for extracting meaningful results.

We design a pre-processing scheme which attempts to “undo” the adverse effects inherent in the image acquisition stage. Specifically, we tackle noise removal, boundary completion and enhancement and contrast enhancement. We also modify the Chan-Vese segmentation scheme by defining a weighting process and explicit parameter relations. We then use the information from an initial unsupervised segmentation to estimate the correct parameter settings for the proposed segmentation method, specific to the image of interest. We compare this scheme with other well-known parameter estimation schemes and parameter settings. We also determine the optimal parameter setting for another two energy functions and compare them to the novel energy function for fluorescent image segmentation.

The results of the proposed methods exhibit a significant boost in segmentation correctness and consistency. It also supersedes previous schemes in terms of being applicable over a wider range of images. Furthermore, it boasts a greater sense of generalisation since the parameters can be tuned for other images properties inherent in other domains.

The methods and techniques presented herein will greatly improve the results and stability of higher level decision making as well as removing the need to manually fine-tune segmentation results.

Preface

The research work described in this dissertation was carried out in the School of Engineering, University of KwaZulu-Natal, Durban, from July 2013 to December 2016, under the supervision of Professor Jules-Raymond Tapamo, PhD.

This study is the original work of the author and has not been submitted in any form for any degree or diploma to any tertiary institution. Where use has been made of the work of others, it is duly acknowledged in the text.

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DECLARATION - SUPERVISOR

As the candidates supervisor, I agree to the submission of this dissertation.

Prof. Jules-Raymond Tapamo

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DECLARATION 1 - PLAGIARISM

I, Ryan NAIDOO, declare that

1. The research reported in this thesis, except where otherwise indicated, is my original research.
2. This thesis has not been submitted for any degree or examination at any other university.
3. This thesis does not contain other persons' data, pictures, graphs or other information, unless specifically acknowledged as being sourced from other persons.
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DECLARATION 2 - PUBLICATIONS

Details of contribution to publications that form part and/or include research presented in this dissertation (include publications in preparation, submitted, in press and published and give details of the contributions of each author to the experimental work and writing of each publication)

A Preprocessing Scheme for Fluorescence Microscopy Image Segmentation. Ryan Naidoo and Jules-Raymond Tapamo. In *International Journal of Imaging and Robotics*, pages 1-23, 09 August 2016.

Signed:

"If any of you is deficient in wisdom, let him ask of the giving God [Who gives] to everyone liberally and ungrudgingly, without reproaching or faultfinding, and it will be given him."

James 1.5 AMPC

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List of Abbreviations

ACM	Active Contour Model
ACWE	Active Contours Without Edges
AHE	Adaptive Histogram Equalisation
AOD	Average Optical Density
AIDS	Acquired Immune Deficiency Syndrome
ARV	Antiretroviral
BCC	Boundary Chain Code
BFS	Breadth First Search
BP	Belief Propagation
CCD	Charge-Coupled Device
CED	Coherence Enhancing Diffusion
CLSM	Confocal Laser Scanning Microscopy
CML	Chronic Myelogenous Leukaemia
CRF	Conditional Random Field
DCC	Differential Chain Code
DFS	Depth First Search
DNA	Deoxyribonucleic Acid
DP	Dynamic Programming
DT	Delaunay Triangulation
EGFP	Enhanced Green Fluorescent Protein
EM	Expectation Maximisation
EMGMM	Expectation Maximisation Gaussian Mixture Modelling
FCS	Fluorescence Correlation Spectroscopy
FDA	Food and Drug Administration
FIFO	First-In First-Out
FISH	Fluorescence in-situ Hybridisation
FLIM	Fluorescence Lifetime Imaging Microscopy
FM	Fluorescence Microscopy
FN	False Negatives
FP	False Positives
FRAP	Fluorescence Recovery After Photobleaching
FRET	Fluorescence Resonance Energy Transfer
GA	Genetic Algorithm
GC	Graph Cut
GCBLS	Graph Cut Based Level Set
GFP	Green Fluorescent Protein

GLCM	Gray Level Co-occurrence Matrix
GMM	Gaussian Mixture Modelling
GRF	Gibbs Random Field
GVF	Gradient Vector Flow
HGP	Human Genome Project
HIV	Human Immune Virus
HLF	Highest Level First
ICC	Immunocytochemistry
ICF	Immunocytofluorescence
ICM	Iterated Conditional Modes
IHC	Immunohistochemistry
IHF	Immunohistofluorescence
IOD	Integrated Optical Density
Laser	Light Amplification by Stimulated Emission of Radiation
LBP	Loopy Belief Propagation
LED	Light Emitting Diode
LoG	Laplacian of Gaussian
MAP	Maximum A Posteriori
MCC	Matthews Correlation Coefficient
MIS	Medical Image Segmentation
MLP	Multi-Layered Perceptron
MRF	Markov Random Field
MST	Minimum Spanning Tree
NA	Numerical Aperture
ORI	Optimised Rotational Invariance
OTF	Optical Transfer Function
PSF	Point Spread Function
RF	Random Field
ROC	Receiver Operating Characteristics
RNA	Ribonucleic Acid
SNR	Signal-to-Noise Ratio
SVM	Support Vector Machine
TN	True Negatives
TP	True Positives
TV	Total Variation
UV	Ultraviolet

To God be the glory

Chapter 1

Introduction

The fluorescence microscope is an extremely versatile tool and is used in many fields from petrology and semiconductor research to medical and biological science. In this dissertation we focus on the segmentation of biological images obtained from a fluorescence microscope.

1.1 Motivation

How often is it that a newspaper or any periodical does not contain information on the latest pharmaceutical drugs, the recent advancements in studies of terminal diseases, like cancer, or even the latest ethical-challenging movements in genetic engineering, such as cloning, genetic mutation, etc. In this modern age, this is as rare as not finding a computer in the average household. The insurmountable information in these fields is due to the ability to observe organisms and objects at the micro and even nano level. The consequent insight into this world has allowed us to view the world from an indispensably unique perspective which has since played a vital role in enriching the livelihood of mankind.

Consider just a few highlights of the research and development in drug development, immunology and immunotherapy and genetics from around the turn of the millennium with the corresponding requirements from image analysis.

1. (1996) Antiretroviral (ARV) to treat Human Immune Viruses (HIV) [1]. In 1983/1984 HIV was found to be the causative agent of Acquired Immune Deficiency Syndrome (AIDS). Since then, successful and astonishing research has lead to the development of combative and resistive drugs, known as ARVs. Research in this area is still strong and ongoing. At the time of writing, history has been made with the first large-scale clinical HIV vaccine trials which are taking place in South Africa to prevent a strain of HIV prevalent in Southern Africa.

In these studies, cell count, cell colony growth rates, time-lapse segmentation of cell-to-cell transfer and infection, structural integrity of cellular components, drug reaction and virus life-cycle is off significant concern [2–5].

2. (2001) New targeted therapy transforms treatment for rare leukaemia [6]. Imatinib (Gleevec) was approved after three months of review by the FDA. The fastest in FDA history. It is the first drug proven to act against a molecular defect on the *Philadelphia chromosome*. The drug demonstrated its ability to halt growth of chronic myelogenous leukaemia (CML) in the majority of patients.

In these studies, cell count, cell pluripotency stability, identification and tracking of leukaemic blood cells are off significance [7–9].

3. (2003) Scientists decode the human genome [10]. A thirteen-year collaborative effort, involving scientists from 7 countries and funded primarily by the US, comes to a halt after mapping three billion DNA letters in the human genome. This marked the completion of the Human Genome Project (HGP).

In these studies, RNA, DNA and genome annotation is off importance [11, 12].

4. (1998-2006) First targeted anti-breast cancer drug, trastuzumab (Herceptin), has major impact on care [13]. About 25%-30% of women suffering with advanced breast cancer are diagnosed as *HER2+*. This is where there is an over-production of the protein HER2 which increases the aggressiveness of the tumour. In 2001, the FDA approved a revolutionary drug, trastuzumab (Herceptin), which drastically increases the survival rate of treated women.

In these studies, segmentation, tracking and morphology and cell colony growth rates are off significance [14–16].

5. (2012) Record number of Americans surviving cancer - nearly 14 million [17]. A report published by The National Cancer Institute and the American Cancer Society claims that cancer surviving patients in the US have increased to 13.7 million. At the time, it was the highest recorded cancer survival rate in US history. A drastic over four-fold increase than 1971 which was recorded at three million survivors.

In these studies, segmentation, tracking and morphology and cell colony growth rates are off significance [18, 19].

The common denominator in all research of this nature is the "tool", the fluorescence microscope. It can be easily inferred that fluorescence microscopy is an essential and indespensable field of knowledge that is crucial to the advancement of human-kind. Coupled with the rapid advancement in optical engineering, there is now an ever-increasing amount of image data which needs to be analysed, the sheer volume, of which, presents too much of a burden on manual analysis. This is where we can harness the unmatched computational power of modern computers. A fundamental task in nearly every image analysis endeavour is that of accurate segmentation of the objects of interest. This has placed image segmentation as a crucial tool in the study of life sciences and related fields. However, the course between fluorescence images and extracting a meaningful segmentation is laden with barriers and hurdles such as noise, extreme low contrast, non-uniform illumination, etc.

Despite its importance, fluorescence image segmentation methods are typically comprised of archaic techniques, such as density-based or gradient-based segmentation, histogram segmentation and watershed segmentation variations [20, 21]. Other popular segmentation methods, especially in blood cell segmentation, are K-means and Fuzzy segmentation variations. This is due primarily because fluorescent image properties have not been incorporated well enough into more sophisticated mathematical segmentation models. Although, Active Contour Models have seen rapid and sustained growth in biological image segmentation due to it's robustness and ability to segment closed shapes [19, 22, 23].

1.2 Main Goals and Objectives

The aim of this research is to better understand cellular fluorescence image properties and use this information to enhance and develop domain-specific segmentation methods, using discrete combinatorial techniques, for extracting meaningful results. We focus specifically on incorporating fluorescent image properties into graph cut segmentation models. This is done by tailoring the energy function, which is to be minimised, to suit the need, given the properties of the data.

Graph cuts is an extremely powerful and general mathematical tool. It has also been used in 3D reconstruction, image restoration, image stitching, etc. All that is required, is that the problem be characterised discretely and that the resulting function meet the submodularity constraint, for global minimisation.

We look at the whole fluorescence image segmentation process including image preparation, automatic and interactive segmentation.

The range of effectiveness of the segmentation methods are done by comparing the segmentation results against the ground truth by calculating the appropriate binary classification measures. The two most important measures are accuracy and the Matthew's Correlation Coefficient.

1.3 Contributions and Publications

In this dissertation the main contributions are the following:

1. **Novel pre-processing scheme for fluorescence images as a preparation for segmentation [24].** The fluorescent image acquisition process is far from perfect. The sub-processes involve accumulatively degrading the image. This scheme is designed to, as much as possible, reverse the negative effects imposed on the image and amplify its segmentation characteristics. The output image is the image that will be segmented. The segmented mask is then used to extract the object from the original image.
2. **Novel cell intensity enhancement function based on Bezier curves [24].** Fluorescence imaging is low light, low contrast technique. This poses a major problem for accurate object segmentation. However, there are strong characteristic features that allow for the object data to be enhanced and simultaneously suppress data that isn't likely to belong to the object. This sub-process in the pre-processing scheme designed for fluorescent image enhancement.
3. **Novel parameter relations and modified graph weighting for graph cut active contours without edges segmentation.** A modified weighting technique is proposed that allows for easier mathematical analysis which reveals the implicit relationships between parameters. From these relationships, we devise a general estimation method which is encoded by other variables which can be easily and intuitively tuned. Then we use these "proxy relationship encoding variables" and optimise them for fluorescence image segmentation. Results show a significant boost in classification accuracy as well as consistent results over a large range of fluorescent image types.

4. **Optimal parameter settings for the energy function proposed Boykov and Jolly [25] for interactive fluorescent image segmentation.** The energy function, which is used to weight the graph, that was proposed by Boykov and Jolly, is one of the most commonly used weighting schemes for general graph cut problems. We alter this general form, into one that can be used for segmentation, as proposed in [25], and find the optimal parameters for fluorescent image segmentation based on seeds marked by the user.
5. **Optimal parameter settings for the energy function proposed by Eriksson *et al.* [26] for interactive fluorescent image segmentation.** Similar to the previous point, the energy function, which is used to weight the graph, that was proposed by Eriksson *et al.*, is also a very commonly used weighting system for graph cut image segmentation. We find the optimal parameters for fluorescent image segmentation based on seeds marked by the user.
6. **Novel energy function and the corresponding optimal parameters for interactive fluorescent image segmentation.** We alter the energy function to encode the dominant intensity characteristics of objects in fluorescent images and find the optimal parameters for segmentation. This weighting scheme has two variants: seeding only, and using the seeds as hard constraints. Results show a significant boost in classification accuracy as well as greater stability over varying fluorescent image types in comparison to the two previously mentioned interactive segmentation energy functions.

1.4 Outline

The rest of the dissertation is organised as follows: Chapters 2 and 3 discuss the background material. In Chapter 2 we introduce the fluorescence process. The main points are briefly discussed and focus is put on the problems related to the imaging process. In Chapter 3 we introduce the graph cut technique and its application specifically for 2D image segmentation. The literature review is covered in Chapter 4.

Chapters 5 to 7 cover the contributions and work undertaken. In Chapter 5 we present the novel pre-processing scheme and data enhancement function. In Chapter 6 we present the modified graph weighting and parameter estimation method for ACWE graph cut segmentation. In Chapter 7 we present the fluorescence image specific optimal parameters derived for the energy functions by Boykov and Jolly, and Eriksson *et al.* We also present the novel energy function, derive the optimal parameters for fluorescence image segmentation and compare it to the previous energy functions.

In Chapter 8 we summarise our contributions and point out limitations and possible directions for future work and extensions.

Chapter 2

Fluorescence Microscopy

Fluorescence microscopy has become an essential tool in diverse fields, such as petrology, semiconductors, etc., and it has especially been established as an imaging technique of choice in cellular and molecular biology for visualisation of cells and tissues [27–31]. In this dissertation, we focus our attention on its use in cellular biology.

Certain substances emit radiation when irradiated with a higher intensity light, such as ultraviolet (UV), blue or green, which is of a longer wavelength than that of the exciting light, this is known as *Stokes' Law*. This phenomenon is known as *photoluminescence* [32–34]. There are two types of photoluminescence. If emission persists at an appreciable level after the exciting light is turned off, then we call this *phosphorescence*. If emission persists only so long as the exciting light is on, then we call this *fluorescence* [35, 36].

The first observance and publishing of fluorescence is credited to Sir John Frederick William Herschel circa 1852. Not too later on, in 1852, Sir John George Stokes published a 100-page treatise about this luminescent phenomenon and coined the term *fluorescence*, over Herschel's *dispersive reflection*, when he observed that the mineral *fluorite* emitted red light when irradiated by ultraviolet (UV) light [28, 37].

In the remainder of this chapter we present the underlying principles of fluorescence; how specimens are fluorescently marked; the optical principles of microscope design; image acquisition; image processing and common analysis in cellular biology. This chapter will only go so far as to present a rudimentary understanding of fluorescence microscopy that is necessary for the comprehension of this dissertation.

2.1 Physics of Fluorescence

Fluorescence microscopy is a cross-disciplinary field. It is encouraged, although not necessary, for biologists and computer scientists to have a photo-physical understanding of the principles of fluorescence. The knowledge of the "under the hood" mechanics of fluorescence empower computer scientists to make informed and directed research in terms of image processing. The aim here is to present an elementary introduction into the physics of fluorescence.

The overall principle of fluorescence can be explained in three steps [38] as illustrated in Figure 2.1(a): (1) Energy is absorbed by an atom upon collision with a photon. (2) The atom becomes excited and an electron jumps to a higher energy level. (3) Shortly after jumping to the higher energy level the electron returns to the ground state and emits a photon.

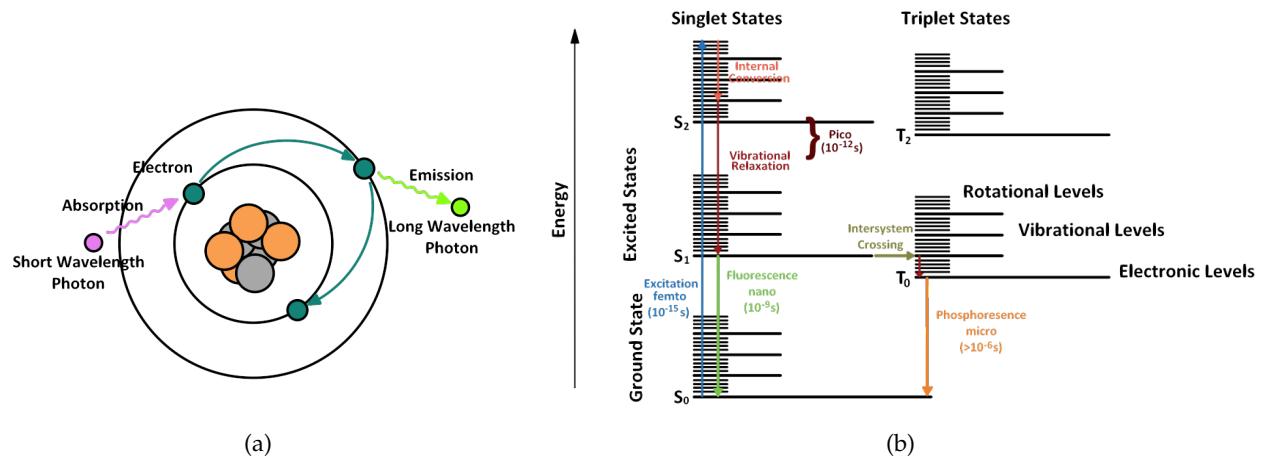


FIGURE 2.1: (a) Simplified fluorescence process. (b) The Jabłoński diagram depicting the electronic states from photon absorption to photon emission.

Excitation and emission One of the best ways to visualise the fluorescing process is using a Jabłoński diagram. A detailed Jabłoński diagram is shown in Figure 2.1(b). When a photon collides with an atom, all its energy is transferred to the atom. The energy of the photon is inversely related to its wavelength, $E = h\frac{c}{\lambda}$, where h is Planck's constant and c and λ are the speed and wavelength of light in a vacuum respectively. This increase in energy moves an electron from the ground state, S_0 , to a higher energy level state. Depending on the energy of the photon and the number of photons absorbed by the electron, the electron can move to different energy levels e.g. S_1 , S_2 , etc. This process happens almost immediately in the order of femtoseconds(10^{-15} s). Before moving to the next lowest energy level, S_1 , some energy is lost due to internal conversion and vibrational relaxation. When the electron spontaneously decays from S_1 to S_0 it emits a photon of a longer wavelength than the absorption photon. This happens within a few nanoseconds (10^{-9} s) after excitation.

The difference in wavelength of the emission photon and excitation photon is known as the *Stokes' Shift* or *Stokes' Law*. The larger the Stokes' Shift the easier it is to separate the emission light from the excitation light [27]. The emission curve is often a mirror image and shifted to longer wavelengths than the excitation curve as illustrated in Figure 2.2.

Fluorophores Substances that exhibit the fluorescent property are called fluorophores, also known as *fluorochromes* or *fluorescent dyes*. Early investigations showed that many substances naturally possess fluorescent properties, such as minerals, crystals, resins, crude drugs, butter, chlorophyll, vitamins, etc. [27]. This is known as *autofluorescence*. Substances that do not fluoresce must be *stained* such that it can be observed through a fluorescent microscope; more on this later.

Intersystem crossing If the electron spin as the electron transfer between energy states is preserved then the energy states are known as *singlet states*. It is also possible for the electron to reverse its spin. However, this is very unlikely and is known as *forbidden transition* in Quantum mechanics. When this happens, the electron is said to be in a *triplet state*, see Figure 2.1(b). The only way for the electron to reach the ground state is to undergo another forbidden transition, which is even less likely. When, eventually, this does happen, the electron may emit a photon; this phenomenon is known as *phosphorescence*. This process takes much longer, in the order of microseconds (10^{-6} s).

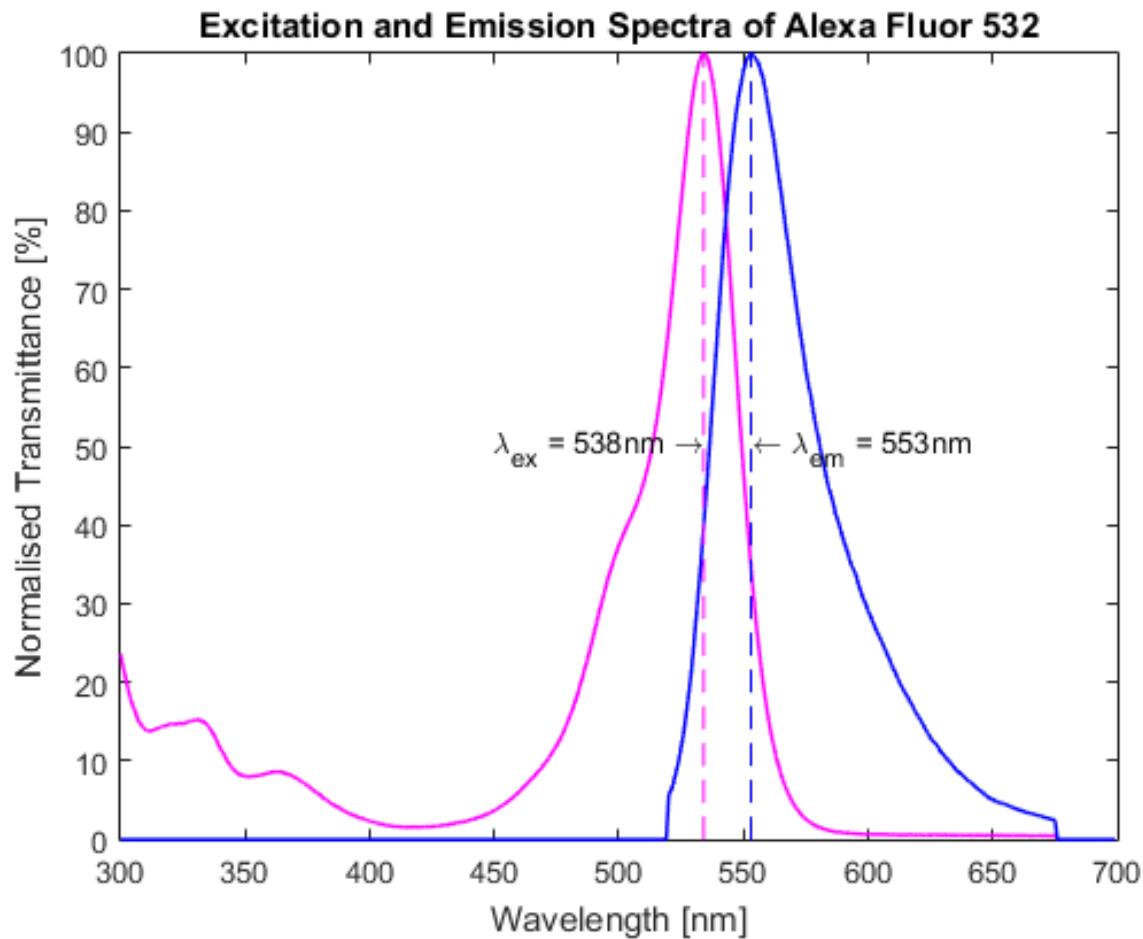


FIGURE 2.2: Normalised Excitation and Emission Spectra of the Alexa Fluor 532 fluorophore. The emission maximum is at 553nm which is a more yellow-green than the excitation maximum at 538nm . Data obtained from ThermoFisher Scientific [39].

2.2 Specimen Labelling

Many of the components of interest, such as cell nuclei, cytoplasm, genes, chromosomes, proteins, do not possess a high degree of, if any, autofluorescence. In this scenario, these components can be marked with a fluorescent dye [40], also known as a fluorophore or fluorochrome, a substance that can bind to a specific target whose excitation and emission spectra are well known. This is known as staining [28, 29, 37]. Once the specimen is stained it can be indirectly observed using a fluorescence microscope.

The most prevalent staining techniques are fluorescence in-situ hybridisation (FISH) and immunostaining [28, 30, 41, 42].

FISH staining FISH is a molecular cytogenetic technique that uses fluorophores that bind to selected regions in nucleic acids [28, 30]. FISH is the most frequently used staining technique which is used primarily for the visualisation and localisation of nucleic acid sequences, chromosomes, cytoplasm or organelles which contain those acids [29]. This makes FISH highly attractive for finding specific

features in DNA and RNA used in genetic diagnosis and research, medicine and species identification [30, 43]. Figure 2.3(a) is a microscope image of mouse chromosomes using the FISH staining technique.

Immunostaining Immunofluorescence is the detection method where an antibody is used to detect an antigen in a tissue or a cell using fluorescence. Fluorophores are usually conjugated onto antibodies, which are proteins that are designed bind to specific antigens, the target proteins, on a cell [44]. The two types of immunofluorescent detection are immunocytofluorescence (ICF) and immunohystofluorescence (IHF). These must not be confused with immunocytochemistry (ICC) and immunohistochemistry (IHC). *Immuno* refers to the immunological technique, i.e. the binding of antibodies to antigens. *Cyto* refers to cells, i.e. cells without the extracellular membrane. *Histo* refers to tissue i.e. cells with the extracellular membrane. *Chemistry* refers to the chemical method of detection, e.g. a change in colour. *Fluorescence* refers to detection by emission of light [45]. Figure 2.3(b) shows the detection of the p53 Binding Protein 1 in perfusion fixed frozen sections of rat kidney.

Live-cell staining The staining techniques discussed previously are not suitable to observe living cells. The fluorescent dyes that are used are phototoxic and cause cells to die. To circumvent this problem an elegant solution has been devised. Instead of staining, the cells are modified to produce a fluorescent substance in the target structures. Derivatives of the *green fluorescent protein* (GFP), isolated from the *Aequorea victoria* jellyfish [30, 40, 46], are used as it generates a strong photon emission and is non-toxic to living cells [28, 29, 37].

2.3 The Epifluorescence Microscope and Image Acquisition

A fluorescence microscope is an optical microscope that is designed specifically to exploit the principle of fluorescence to allow for the observation of fluorescently labelled specimens [30, 47]. There are many types of fluorescent microscopes available but the preferred type among many biologists and geneticists is the epifluorescent microscope [34, 48]. A schematic of the epifluorescent microscopes is illustrated in Figure 2.4.

Light source The light source is typically a high-luminance light source e.g. Mercury or Xenon arc lamps, LEDs, lasers, etc. [48, 49]. The primary criterion for choosing a light source is that its characteristic peaks must coincide with the excitation spectrum of the fluorophores being used [27, 30, 46]. Wavelength coverage spans from near infra-red to UV. Mercury and Xenon arc lamps are expensive. An inexpensive and lightweight alternative are bright LEDs [35, 50].

Excitation filter Typically, the incoming light from the light source is multispectral [36]. The excitation filter is a wavelength selection filter which is placed in the path of the incoming light and filters through only those wavelengths in the absorption spectrum of the fluorescent dye [28, 37].

Dichroic mirror Also known as a *dichroic beam splitter*. This is placed at a 45° angle and reflects the short-wavelength light filtered through the excitation filter at a 90° angle towards the specimen [27, 44] and allows the long-wavelength light from the fluorescing specimen to pass through to the detector [35, 46], thus serving as a separation filter between the absorption and emission light [30, 37].

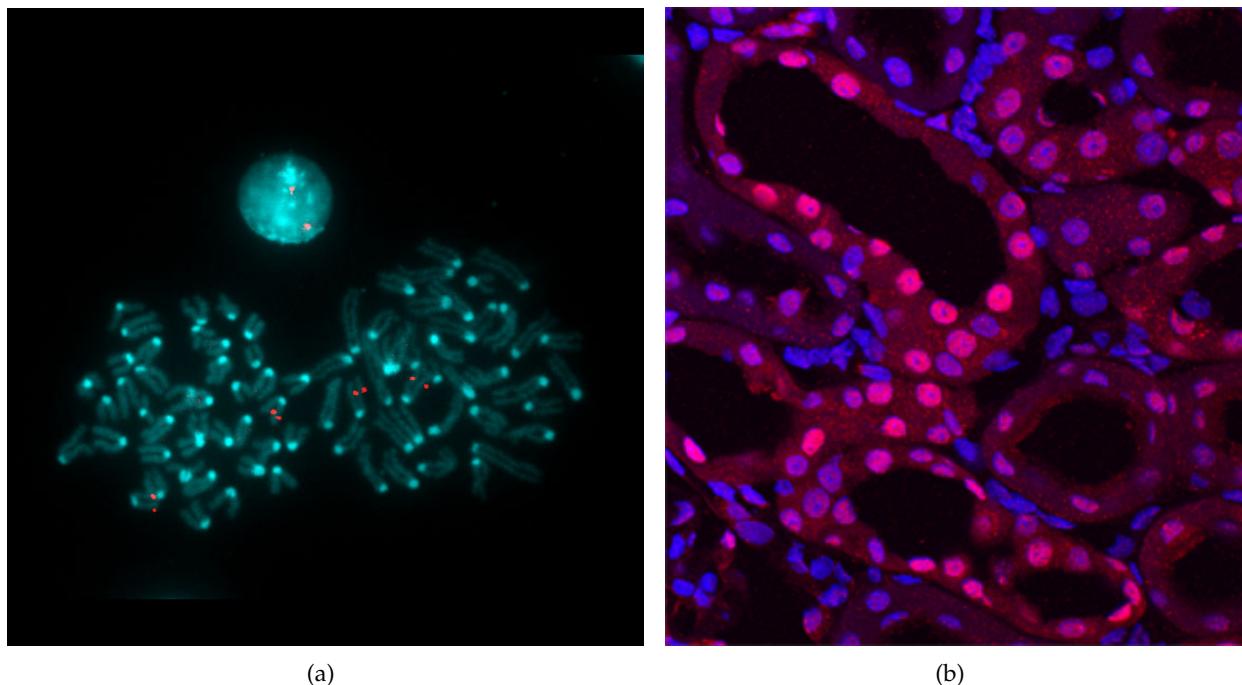


FIGURE 2.3: (a) FISH (Fluorescent 'in-situ' Hybridization) in mouse chromosomes using a BAC clone labeled with Spectrum Orange. The picture shows two metaphases and one interphase with two signals in each exemplifying a homozygous mouse for a transgenic clone. Image Source: "All About the Human Genome Project" Genetic and Genomic Image and Illustration Database. (b) p53 Binding Protein 1 (53BP1) was detected in perfusion fixed frozen sections of rat kidney using Goat Anti-Human 53BP1 Antigen Affinity-purified Polyclonal Antibody (Catalog # AF1877) at 15 µg/mL overnight at 4°C. Tissue was stained using the NorthernLights™557-conjugated Anti-Goat IgG Secondary Antibody (red; Catalog # NL001) and counter-stained with DAPI (blue). Specific staining was localized to nuclei of epithelial cells in convoluted tubules.

Image Source: R&D Systems' IHC image database.

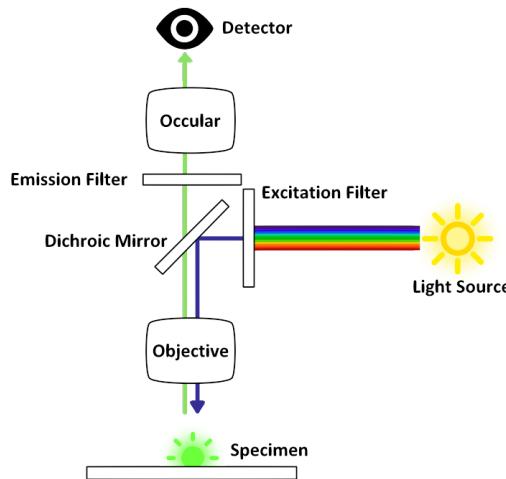


FIGURE 2.4: The schematic of the epifluorescence microscope.

Objective The incoming light reflected off the dichroic mirror passes through the objective lens before reaching the specimen [27, 46]. Emission light from the fluorescing specimen is gathered in the objective lens and passed through to the dichroic mirror.

Specimen The specimen is irradiated by the incoming light from the objective and emits long-wavelength light in all directions. The specimen is stained with a fluorophore whose absorption and emission curves are well known. This is important since the light source and the interference filters are chosen using the peaks of these curves.

Emission filter Also known as a *barrier filter* [27, 35, 46]. The light coming from the specimen contains multiple wavelengths and the dichroic mirror is used to filter out the shorter wavelength light. The emission filter is further used to filter out the wavelengths that correspond to the emission wavelengths of the fluorophore [36, 49].

Detector A detector is used to capture the emission light and can further digital form the image. The detector is usually a photodetector such as a charge-coupled device (CCD) camera or a photomultiplier tube [27, 51]. It is vital that an appropriate detector is chosen as this has a direct impact on the image quality [30].

The type of fluorescence image data that needs to be captured is application dependant and this impacts the decision on which type of microscope is to be used. The *widefield*, or conventional, microscope produces 2D image data. 3D image data cannot be captured directly. Instead, a series of 2D images are captured to form a 2D stack. The 3D image is then constructed in software. In this scenario, the most common choice of microscope is the *confocal laser scanning microscope* (CLSM). This microscope system is expensive and acquisition is slower. An economical alternative is the *confocal spinning disk microscope*. To detect single molecules, the preferred technique is *total internal reflection fluorescence* (TIRF) which can be achieved by a modification of the epifluorescence microscope.

2.4 Image Processing

Due to the physical nature of the fluorescence and image acquisition process, there are many factors that can degrade image quality. There are measures that can be taken to largely mitigate some problems but they can never be completely eliminated. The presence of these factors directly affect segmentation accuracy and subsequently higher level analysis. Therefore, images are processed prior to segmentation to suppress the artefacts and reconstruct the original data [28] or better yet to enhance it. Chapter 5 takes a step in this vein. Here we present some of the commonly occurring factors that reduce image quality and the methods used to mitigate them, and some of the techniques employed for segmentation.

2.4.1 Pre-processing

Non-uniform illumination There are many factors that could contribute to non-uniform illumination. The specimen layer will not be uniformly lit if the arc lamp is not properly focussed on the black aperture. To prevent this from happening a liquid light guide-based light source, which provides even illumination, may be used [46]. If the light brightness diminished towards the edges of the image then this is known as the *Vignette effect*. Common techniques to suppress this distortion is *background correction*, also known as *flat-field correction*, *background flattening* or *shading correction* [28, 30,

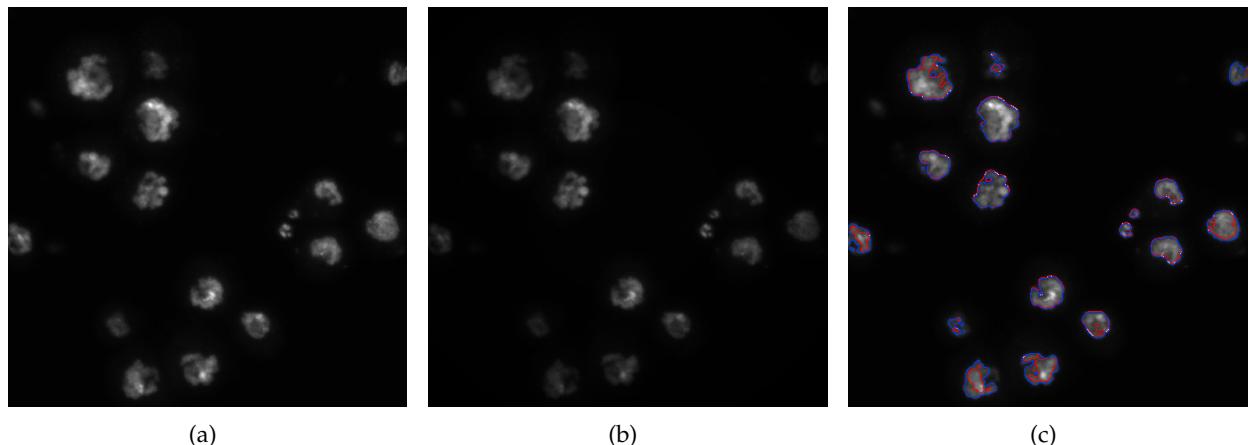


FIGURE 2.5: The effect of non-uniform illumination on segmentation. **(a)** Original image. Image source: The Cell Image Library [56]. **(b)** Non-uniformly illuminated image. The Vignette effect. **(c)** K-means clustering using the Euclidean distance metric and 2 clusters. The blue curve is from the original, the red curve is from the non-uniformly illuminated image. Notice that less of the object is recognised towards the edges.

[51]. Other causes of non-uniform illumination are inhomogeneous detector sensitivity, autofluorescence, dirt particles in the optical system or non-specific sample staining. An example of the Vignette effect is illustrated in Figure 2.5(b). Computational schemes to eliminate non-uniform illumination have been well researched, although recently it has not received too much attention [52–55].

Fading The reduction of emission intensity is called *fading*, of which there are two types: *quenching* and *bleaching*.

Quenching is a reversible loss of fluorescence owing to a variety of non-radiative energy-loss mechanisms such as collisions with nearby acceptor molecules, a phenomenon known as *resonance energy transfer*. This phenomenon is useful in studying molecular interactions below the lateral resolution of the light microscope through a technique called *fluorescence resonance energy transfer* (FRET) [27, 28, 46].

Bleaching refers to all processes that cause a fluorescent signal to fade permanently [46]. From the fluorescence process presented in Section 2.1, one may assume that, under the proper conditions, a fluorochrome has the ability to fluoresce indefinitely. However, this is not the case. There is a limited number of cycles before permanent bleaching [46]. Figure 2.6 illustrates the effect of photobleaching. *Photobleaching* is the most prominent form of bleaching. It is due to the interaction of the fluorophore with an oxygen molecule. This can move the oxygen molecule to an excited singlet state, which then becomes a reactive molecule. When in this state, the oxygen molecule can participate in many chemically destructive reactions with organic molecules causing *phototoxicity* [28]. Photobleaching is used to study diffusion and motion through a technique called *fluorescence recovery after photobleaching* (FRAP) [34, 46].

Photobleaching cannot be avoided but it can be pushed back. The aim in the measures to avoid photobleaching is to take a longer time to reach *reciprocity failure* [34]. This is done by shortening exposure times and using less intensive excitation light. This, however, yields low contrast images. These images with low signal-to-noise [51] ratio (SNR) are more difficult to segment hence contrast enhancement must be performed on the images prior to segmentation [57].

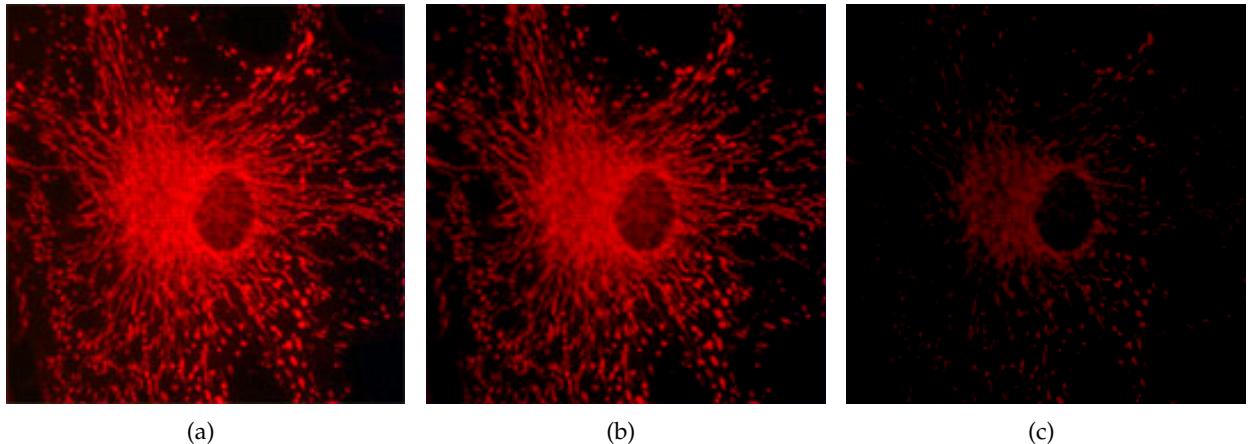


FIGURE 2.6: The effect of photobleaching. Image Source: Molecular Expressions [58]. **(a)** Original image at $t = 0\text{s}$. **(b)** Image after $t = 10\text{s}$. **(c)** Image after $t = 20\text{s}$.

Image distortion The major contributing factors to image distortion are: (a) blurring due to the *point spread function* (PSF) and (b) noise [33]. Image formation can be approximated by

$$O = n(s \otimes h), \quad (2.1)$$

where O is the formed image, n is the noise function, s is the exact image, \otimes is the convolution operator and h is the specific PSF of the optical system. We discuss the PSF first. The observed image is not an exact representation of the real object. Optical systems have an inherent property called ‘the point spread function’ which is the systems optical response to a point light source [28]. The final image is dependent on the spatial position of a point, numerical aperture (NA) and furthermore differs for various emission wavelengths [29, 59]. The final image is a superposition of all points in the illuminated volume where the contribution of each is described by the PSF.

Theoretically, the exact image can be obtained by the deconvolution of the observed image with the PSF. However, there are secondary factors that prevent this. Deconvolution seeks to reconstruct the original image given the PSF and certain assumptions about noise [59]. This is an *ill-posed problem* as little is known about the specific PSF or the noise model. Additionally, if the image is corrupted by too much of noise, deconvolution might still produce unsatisfactory results. The PSF is generally experimentally determined or theoretically modelled, and so the true PSF is never attained. Hence, the original image can never be attained by deconvolving with an approximated PSF. The image formation and restoration process is illustrated in Figure 2.7.

Deconvolution has received a lot of attention in the past with very creative approaches [61–65]. Fluorescence microscopy deconvolution is still a very active field of research. Bayesian methods have become popular in fluorescence microscopy image deconvolution. However, noise in low SNR imaging conditions still poses a challenge [66]. Recent trends in 3D deconvolution in widefield microscopy use blind depth-invariant of the PSF [67]. Most PSF deconvolution systems naïvely assume depth-invariance; however, the PSF changes significantly along the optical axis. There are also deconvolution methods that can preserve detail and possibly enhance image quality in diffraction-limited/super-resolution imaging modalities [68].

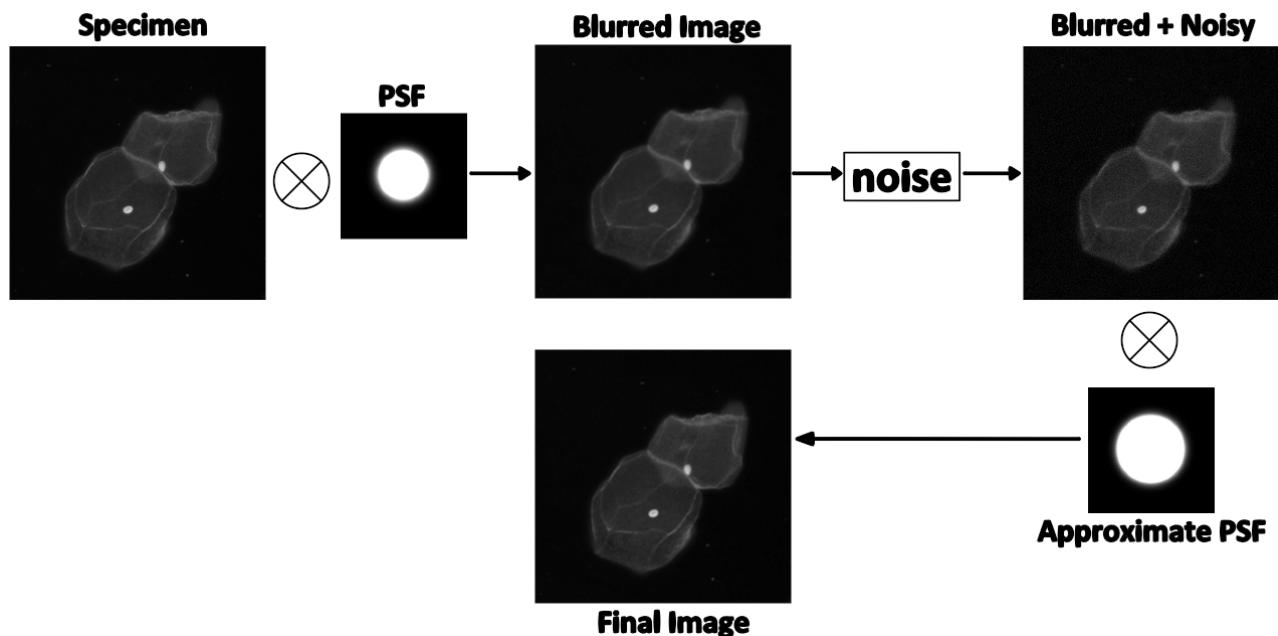


FIGURE 2.7: The image formation process. Image source: [60]

Noise As previously stated, fluorescence imaging is a low light, low contrast and low SNR imaging technique to counter the effects of bleaching. For these reasons, noise becomes prominent. The three types of noise recorded by a camera is *dark noise*, *read noise* and *photon noise* [28, 29, 31, 37].

The electrons in a CCD film are always in motion due to thermal energy. Dark current is due to the extraneous electrons which are excited into the signal. This signal carries a statistical fluctuation known as dark noise. Dark current effects can be reduced by ground image subtraction or cooling the CCD. If there is a significant amount of dark noise, then the background will not be as black as is necessary. For this reason, it is common to mistake dark noise for low-level autofluorescence [24, 37].

Read noise is a result of the conversion process from charge build-up to a voltage and then digitisation.

Photon noise, also known as *shot noise*, is the signal-dependent statistical variation of the counting of photons incident on the CCD or film. This is a naturally occurring phenomenon and cannot be reduced by camera design or system optimisation [24].

In low-light imaging techniques, such as is common in fluorescence imaging, the dominant form of noise is photon noise. Photon noise and dark noise are Poisson distributed [24, 28, 69]. Noise in low-light images was previously modelled using the Gaussian distribution but was found to be a poor description of the noise. The Poisson distribution provides a more physically accurate model especially in photon-limited recording [33]. We study denoising in Section 5.1.1. The effect of noise on segmentation accuracy is illustrated in Figure 2.8.

2.4.2 Segmentation

The primary aim of using fluorescence microscopy imaging is to make diagnoses and to study molecular behaviour and interaction. This means that biologists, geneticists and other professionals alike, not only have a superabundance of microscopic imaging techniques at their disposal, but also

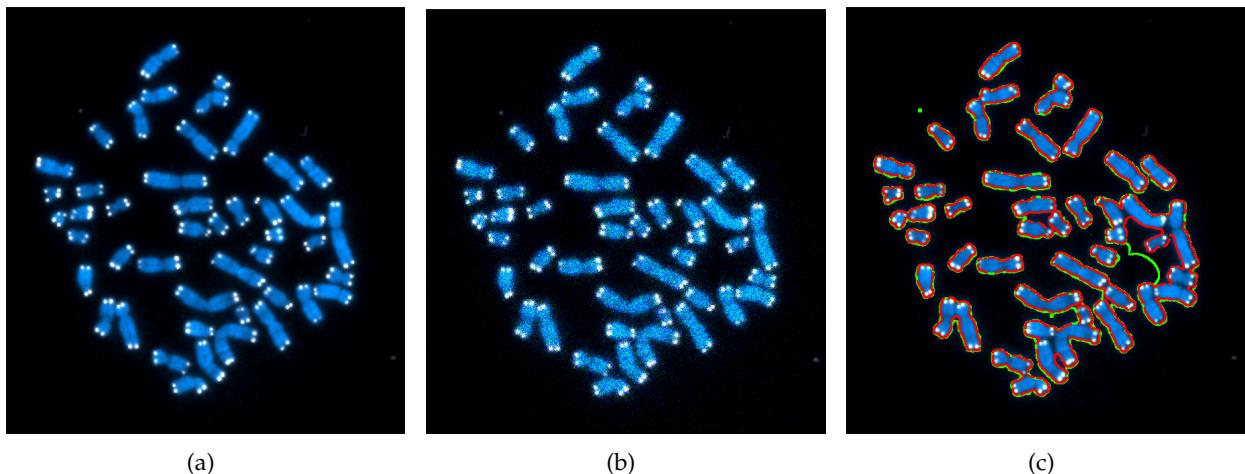


FIGURE 2.8: The effect of noise on segmentation. **(a)** Original image. **(b)** Poisson noise corrupted image. **(c)** ACWE Chan-Vese segmented output. The red curve is from the original, the green curve is from the Poisson noise corrupted image. Notice that there are more artefacts and the segmented output is less accurate especially towards the bottom left from the green curve.

have an immense amount of image data to analyse since the outburst of image acquisition technology. The abundance, diversity, dimensionality and complexity of fluorescence image data obviates manual image processing as this is not tractable, by any means, in terms of time or quality [24, 70]. Consequently, the task has fallen to computers to perform these tasks and has now become essential in advancing these fields [71, 72]. The crux of image analysis lies on the accuracy of image segmentation and has become the principle focus in many studies [73].

Anton van Leeuwenhoek, a Dutch draper and scientist, is credited with the invention of the first real compound microscope in the late 17th century. However, it was not until the mid-1950s that computers became involved when systems were developed to automate the classification of smears of exfoliated cells. These systems used simple thresholding rules on 1D scanning of microscopic lines [74]. In the 1960s, automated systems were developed to count leukocytes on 2D image data based on their colour and morphological measurements [75]. In the 1980s the invention of the confocal microscope made it possible to study cells in 3D but it was not until the 1990s that the computer became powerful enough to process 3D image data or complex 2D images [76]. The trends of increased computing power has made possible the use of more sophisticated cell analysis techniques as well as the ability to use more computationally demanding segmentation methods. The literature on cell segmentation and analysis has experienced exponential growth in the last couple of decades. Many studies comparing segmentation algorithms for cells have been published [77–79]. We review some of the common techniques used in fluorescence microscopy image segmentation.

Intensity thresholding Intensity thresholding assumes a non-overlapping intensity levels between the objects and the background. It is still one of the most common thresholding methods for cell segmentation [80–82]. Locally adaptive thresholding techniques are used when illumination varies across the image. Automated threshold segmentation techniques are usually based on global or local intensity using the histogram [73]. Thresholding produces suboptimal results due to the naïve assumption of mutually exclusive intensity levels.

Morphological segmentation This method uses non-linear mathematical morphological operators like erosion, dilation, opening, closing, etc. with geometrical and topological properties to segment the image [81, 83–86]. Generally, this method is used as a post-processing step to refine a coarse segmentation; or as a pre-processing step to suppress certain image structures [73].

Region accumulation This method starts with selected points, called seeds. The idea is to iteratively add points neighbouring previously labelled pixels based on some conformity measure, usually intensity. The most common implementation is called *region growing*. Most cases assume an image model similar to that of thresholding and suffers the same segmented results problems. Another approach is called the *watershed method* which converts the image into an open 3D shape and "fills the shape with water". The different regions are separated by those "filled with water" and those that are not. A common problem with this method is over-segmentation and usually requires post-processing methods, like *region merging*, to achieve a meaningful result [73, 87, 88].

Edge-based segmentation There are basically two types of edge-based segmentation: *gradient based methods* and *Laplacian based methods*.

Gradient methods are based on the assumption that there is a rapid intensity gradient between the object and the background. Edges are detected by searching for the maximum and minimum in the first derivative, e.g. Prewitt, Roberts and Sobel operators.

Laplacian methods search for zero-crossings in the second derivative, e.g. Marr-Hildreth, Laplacian of Gaussian (LoG), Canny edge detection.

These algorithms can be computed very quickly but the drawback is when closed curves are desired [73], which is a common criterion in biological and molecular segmentation. A solution to this is to use snakes or active shape models.

Energy minimisation The most recent segmentation methods are based on energy minimisation. Most current state-of-the-art techniques fall under this category. This is due to their flexibility and robustness [28]. This group encompasses two main subgroups: *deformable models* and *discrete combinatorial optimisation*.

The aim of deformable model techniques is to fit a deformable model, either a curve or a surface, to the image data. They may be formulated either explicitly, as a parametric contour (2D) or a surface, e.g. snakes [89] or active contours [90–92], or implicitly as a zero-level of a function with one dimension higher than that of the image data, e.g. as a level set [93]. This technique is widely used in fluorescence image segmentation [94–99].

The aim of discrete combinatorial optimisation is to search a finite countable solution space for the optimal solution. Optimality is defined with respect to some energy function, which embeds one or more criteria, which is to be minimised. The most common implementation exploits the graph cut framework. Graph cut segmentation has recently gained a lot of popularity and momentum in medical image segmentation (MIS) [100–107].

Other miscellaneous techniques The techniques presented are just a select few in a plethora segmentation techniques that have been used in microscopy image segmentation. A few other common techniques are unsupervised clustering segmentation using the k-means [108–110], Otsu binarization [18], dynamic programming [111, 112], Voronoi diagrams [113, 114], gradient flow tracking [115], etc.

These are just a few more. Instead of cell segmentation converging to a robust, flexible and unified solution, the number of available options is steadily increasing [70]. There probably exists as many individual unique solutions for cell segmentation as there are problems.

2.5 Object Measurement and Analysis

The aim in fluorescence image analysis is to measure specific properties of interest which enable higher level decision making. Typically, these properties are quantitative measures. In this section, we review some of the important quantitative measurements in digital image analysis. It is important to note that for some of the properties of interest, the accuracy of the measurements depend heavily on the accuracy on the segmentation. The properties of interest are application dependent, one might require solely the object morphology or structure and hence properties like perimeter, area, shape, intensity, colour, etc., are of significance. Alternatively, if one requires the colocalisation of cells, then distance discriminants, such as Euclidean distance, Manhattan distance, Chessboard distance, etc., are of significance [28, 116].

Object measures can be loosely classified into four categories: geometric measures, histogram-based measures, intensity based measures and temporal measures. One can also argue a fifth category, statistical classifiers, although this is generally used in higher level analysis.

Size measures Perimeter, area and volume are common measures to describe the size of objects. Area and volume are suitable measures to describe the general size of an object. The perimeter of an object is distinctly useful in discriminating its shape complexity. Complex and irregular shapes need a larger perimeter to enclose its area.

Pose measures This measure defines an object's location and orientation. The centroid is used as an object's locale and its orientation is the measure of the angle subtended by its major axis.

Shape measures Shape features are used to distinguish objects from one another. These measures are generally translationally, rotationally and scale invariant and can be used independent of, or in conjunction with, the size measures. Commonly assessed shape parameters are thinness ratio to describe the regularity of an object, rectangularity, circularity, Euler number, moments, central moments, object dispersion, rotationally invariant moments, Zernike moments and elongation.

Shape descriptors Shapes descriptors provide a more wholesome way of describing an object's shape than compared to the single parameter shape measures. The differential chain code (DCC) and boundary chain code (BCC) are the two most commonly used shape descriptors that are used to represent the distance around an object. Fourier descriptors is another object distance measure that exploits the periodicity of BCC. There are also graph representations of which the two most common are minimum spanning tree (MST) [117, 118] and Delaunay triangulation (DT) [119, 120].

Distance measures There are many ways to compute the separation between objects. The most commonly assessed distance measures are Euclidean distance, Manhattan distance (also known as the City-block distance or absolute value metric), which is a more computationally efficient approximation of Euclidean distance, and the Chessboard distance (also known as the maximum value metric) [121, 122].

Intensity measures Images are segmented generally into region with low intra-region intensity distribution and high inter-region intensity distribution [81, 123]. Common intensity measures are integrated optical density (IOD) [124, 125], which is simply the sum of all the gray levels that compose the object, it is a reflection of the object's "mass" or "weight"; and average optical density (AOD), which is the IOD divided by the objects area.

Histogram measures These measures provide a measure of an object's intensity distribution. Common histogram-based measures are mean, standard deviation, skew, entropy and energy [126, 127].

Texture measures In image analysis texture refers to the spatial arrangement of gray level values [128] and hence a texture feature quantifies some characteristic of the intensity variation within an object. Common texture measures are statistical texture measures, gray-level co-occurrence matrix (GLCM) [129, 130] and power spectrum features [131, 132].

Ratiometric measures Some fluorescent dyes respond to the changes in Calcium and Hydrogen ion concentration by changing its spectral properties of the fluorescent emission bands. In this case, a ratio of the intensity can be used to calculate concentration of calcium or pH value [37].

Temporal measures Considering the time domain, many interesting properties can be observed. Commonly computed properties of interest are motility [133–135], like velocity and acceleration, rate of growth, rate of change of colour, etc.

These measures are used in higher decision making processes such as the evaluation of a hypothesis to detect the presence of a certain disease. They are also used to aid in the understanding of biological mechanisms, events and interactions [28].

Chapter 3

Mathematical Background

Image segmentation falls under the mathematical classification as being an *ill-posed inverse problem* [136, 137]. It is an inverse problem since we require a model from observation, this simply means: given the results, what are the causes? In image segmentation, this translates to, given a 2D matrix of intensity values, which pixels belong to the object and which belong to the background. Image segmentation is an ill-posed problem since there is a lack of uniqueness or stability of a solution [138], which are two of the three requirements for a solution to be *well-posed*; the other being existence. The ill-posedness exists because an immense amount of information is suppressed in the acquisition processed [139–141]. Many tasks in vision are inherently or can be reformulated as ill-posed inverse problems e.g. scene reconstruction, stereo matching, image restoration, image deconvolution, etc. Computer vision is used heavily in industry, medicine and many life science fields, hence there is a need for a robust, environmentally resistant approach. The *optimisation approach* is an elegant way to obtain a solution. In computer vision, a problem can be posed as an optimisation problem as follows: We are given a coarse, discrete and noisy, approximation of the visual data, d , we aim to infer some hidden quantities x , labels, depth, probable pixel intensity, etc., based on it. We then have to design an *objective function*, also known as an *energy function* or *cost function*,

$$E : (x, d) \rightarrow \mathbb{R},$$

which has to be optimised such that the optimisation of the function provides a solution to the problem. $E(x, d)$ assigns an energy or a cost to each combination (x, d) of the input and hidden quantities. E provides a measure of goodness to how well the candidate solution x fits the expectation given the data d . In the optimisation of this function we seek a minimum energy,

$$x^* = \arg \min_x E(x, d), \quad (3.1)$$

which has roots in Statistical Physics where lower energies correspond to more stable solutions. This gives us a general idea of how we should assign energies to solutions; the better a solution, the lower an energy we should assign to it. In this case, a huge number of inference problems in vision can be solved by minimising the associated energy. A solution is only as good as the energy model and the optimisation technique. Once a precise energy and a minimising algorithm is found, the problem is essentially solved [142].

Early attempts in computer vision would solve problems like these using iteration or relaxation methods [143, 144]. In these attempts the problems are solved in a Calculus of variations framework. This is still a popular approach to optimisation in vision since Poggio *et al.* [136] proposed

an integrated framework to regularisation theory for vision [145]. Many important advancements in computer vision are proposals for a better energy model, a better algorithm or both [142, 146–149]. In this dissertation, we focus on discrete energy optimisation using graph cuts for image segmentation.

Image segmentation falls under a broader category of problems known as *labelling problems*. The aim is to find the best label, foreground/object or background, for each pixel. In Section 3.1 we briefly discuss labelling problems and its formulation as an energy minimisation problem.

3.1 Labelling Problems

Among the many computer vision problems, image segmentation is the easiest labelling problem to understand. A labelling problem is simply assigning to an observation, a label that most accurately explains it. An observation can be anything that we wish to classify e.g. pixels, features, salient points, depth measurements, etc. A label is a description of that observation. There are two types of labels: *semantic labelling* (person, car, tree, sky, face, eye, etc.) or *pixel-wise labelling* (texture, shape, colour, background/object, etc.) [142, 150].

To formulate a labelling problem, we need a set of *sites*, intuitively known as observations, and a set of *labels*, which is a set of explanations. The goal is to find the best explanation given the observations. In computer vision, the observations can be features, image segments, etc. However, they will typically represent pixels in an image with some natural structure or ordering. Let

$$\mathcal{P} = \{1, 2, \dots, n\}$$

be the set of n sites and

$$\mathcal{L} = \{l_1, l_2, \dots, l_k\}$$

be the set of k labels. A discrete labelling is a map $f : \mathcal{P} \rightarrow \mathcal{L}$ that assigns each discrete variable f_p one value from \mathcal{L} and $f = \{f_p\}_{p \in \mathcal{P}}$ which is known as a *configuration*. We are interested in binary segmentation, also known as *binarization*, which implies we have two explanations in our label set, $k = 2$. The labels of interest are the *background* and the *object*. Although the solution space is finite, it is very large and grows exponentially as the image size increases or as the number of labels increases. The number of possible configurations is given by $|\mathcal{L}|^{|\mathcal{P}|}$. Table 3.1 shows the largeness of the solution space even for very small images and a few labels. In practice, the image sizes used in Table 3.1 is too small, hence finding a solution is not easy. Most often, settling for an approximate solution is "good enough".

TABLE 3.1: The impact of the number of sites and labels on the solution space

Image (\mathcal{P})	Number of sites ($ \mathcal{P} $)	Number of labels ($ \mathcal{L} $)	Number of configurations $ \mathcal{L} ^{ \mathcal{P} }$
64×64	$2^{12} = 4096$	2	$2^{2^{12}} = 2^{4096} = n$
128×128	$2^{14} = 16384$	2	$2^{2^{14}} = 2^{16384} = n^4$
64×64	$2^{12} = 4096$	3	$3^{2^{12}} = 3^{4096} \approx n^{1.585}$

3.2 Maximum A Posteriori Estimation for Discrete Models

As previously mentioned, image segmentation can be viewed as a labelling problem. The problem is the huge search space in which the solution, or possibly more than one, exists. We need a metric that is able to appropriately weight a configuration f . *Random Fields* are able to provide a structured and yet flexible probabilistic framework for labelling problems of which *Markov Random Fields* (MRFs) and *Conditional Random Fields* (CRFs) are mostly used in vision tasks. We focus on the discrete image representation provided by MRFs in which we can embed the properties of a desired segmentation solution. MRFs are pivotal in designing, weighting and structuring graphs, so we give a brief introduction into the concepts needed to understand the probabilistic make-up for graph cut image segmentation.

3.2.1 Markov Random Fields

A *random field* (RF) is a stochastic process where each random variable is indexed by a spatial variable [151, 152]. A random field model can be intuitively represented as an undirected graph $\mathcal{G}(\mathcal{V}_{RF}, \mathcal{E}_{RF})$ where $\mathcal{V}_{RF} = \{1, \dots, n\}$ is the set of sites which correspond to a random variable for each pixel in \mathcal{P} , \mathcal{E}_{RF} is the set of undirected edges which links the random variables. In vision, a random variable, which typically corresponds to a pixel, is linked to neighbouring random variables. These links model interdependency in images, nearby pixels exhibit a high degree of spatial correlation (similarity) [153]. Common connectivity arrangements in 2D images are 4 and 8-connectivity. Similarly, higher dimensional data can be represented using graphs. For 3D images, common connectivity arrangements are 6- and 26-connectivity. Connectedness is illustrated in Figure 3.1 for 4-connectivity for 2D image data and 6-connectivity for 3D image data. In this dissertation, we are concerned with 2D images only. Two sites, p and q , are neighbours if edge $(p, q) \cup (q, p) \in \mathcal{E}_{RF}$. The set of neighbours of p are denoted \mathcal{N}_p . The RF associated with \mathcal{P} is denoted as $\mathbf{Y} = \{Y_p : p \in \mathcal{P}\}$, where each Y_p can be assigned one of k labels from \mathcal{L} . A 4-connected RF is illustrated in Figure 3.2. A *clique* c is a fully connected subgraph; it is defined as $\forall p, q \in c, p \in \mathcal{N}_q$ and $q \in \mathcal{N}_p$. In a clique, each site is connected to all other sites.

The joint event $\{Y_1 = y_1, \dots, Y_n = y_n\}$ where $y_p \in \mathcal{L}$ is called a *realisation* or *configuration* for the random field \mathbf{Y} . For readability we simplify the joint event notation to $\mathbf{Y} = \mathbf{y}$ where $\mathbf{y} = \{y_p : p \in \mathcal{P}\}$. The image segmentation problem is now in the form of an inference problem where the image \mathbf{x} is the observation of a hidden random field \mathbf{Y} and the solution is given by the *maximum a posteriori* (MAP), i.e. the solution is given by

$$\mathbf{y}^* = \arg \max_{\mathbf{y} \in \mathcal{Y}} \Pr(\mathbf{y} | \mathbf{x}), \quad (3.2)$$

where \mathcal{Y} denotes the set of all possible labellings. \mathbf{Y} is said to be a *Markov random field* (MRF) if:

$$\Pr(\mathbf{Y} = \mathbf{y}) > 0 \quad \forall \mathbf{y} \in \mathbf{Y}, \quad (\textit{Positivity}), \quad (3.3)$$

$$\Pr(Y_p = y_p | x_{\mathcal{P} \setminus \{p\}}) = \Pr(Y_p = y_p | y_{\mathcal{N}_p}) \quad \forall p \in \mathcal{P}, \quad (\textit{Markovianity}), \quad (3.4)$$

The positivity property constrains all configurations to have non-zero probability and is needed to ensure that the joint probability can be uniquely determined by the local conditional probabilities [154]. The Markovianity property states that a site is conditionally independent of all other sites given its neighbours.

MRFs are one of the most popular probabilistic modelling tools and was introduced to the computer vision community by Geman and Geman [155]; and Besag [156]. MRFs allow us to model local contextual constraints, such as spatial interactions between pixels. According to Baye's rule, the posterior probability relation is:

$$\Pr(\mathbf{y}|\mathbf{x}) \propto \Pr(\mathbf{x}|\mathbf{y}) \Pr(\mathbf{y}), \quad (3.5)$$

$\Pr(\mathbf{x}|\mathbf{y})$ encapsulates the dependency of the labels on the observation, it is the likelihood of observing \mathbf{x} given \mathbf{y} . $\Pr(\mathbf{y})$ is the probability of that specific labelling among all labellings \mathcal{Y} . The joint distribution can be specified as a *Gibbs Random Field* (GRF) [157, 158]:

$$\Pr(x|y) = \frac{1}{Z} \prod_{c \in \mathcal{C}} \exp(-\Psi_c(\mathbf{x}_c)), \quad (3.6)$$

where \mathcal{C} is the set of cliques. Figure 3.3(a) shows a simple MRF construction. This MRF contains cliques of order one and two, i.e.

$$\mathcal{C} = \{1, 2, \dots, 9, \{1, 2\}, \{1, 4\}, \dots, \{8, 9\}\}.$$

Figure 3.3(b) shows a more densely connected MRF where there are cliques of order one, two and three, i.e.

$$\mathcal{C} = \{1, 2, \dots, 9, \{1, 2\}, \{1, 4\}, \{1, 5\}, \dots, \{5, 9\}, \{8, 9\}, \{1, 2, 5\}, \{2, 3, 5\}, \dots, \{5, 6, 9\}\}.$$

However, cliques of orders higher than two are generally ignored due to the computational demands. In Equation (3.6) the term $\Psi_c(\mathbf{x}_c)$ is known as the *potential function* for the clique c , where $\mathbf{x}_c = \{x_i, i \in c\}$. The constant Z is called the *partition function* and ensures that the sum of all probabilities is one. After expanding Equation (3.6) for a maximum clique order of two, the conditional distribution of a pairwise MRF is:

$$\Pr(x|y) = \frac{1}{Z} \prod_{i \in \mathcal{V}} \exp(-\Psi_i(x_i)) \prod_{(i,j) \in \mathcal{E}} \exp(-\Psi_{ij}(x_i, x_j)), \quad (3.7)$$

where $\mathcal{V} = \{1, 2, \dots, n\}$ and \mathcal{E} is the set of pairwise edges, Ψ_i is the unary potential function, for first order cliques, and Ψ_{ij} is the pairwise potential function, for second order cliques.

3.2.2 MAP-MRF Estimation as Energy Minimisation

The equivalence of MRFs and GRFs, proven by the Hammersley-Clifford theorem, means that maximising Equation (3.2) is equivalent to minimising Equation (3.1) [159]:

$$\mathbf{y}^* = \arg \max_{\mathbf{y} \in \mathcal{Y}} \Pr(\mathbf{y}|\mathbf{x}) = \arg \min_{\mathbf{y} \in \mathcal{Y}} E(\mathbf{y}, \mathbf{x}); \quad (3.8)$$

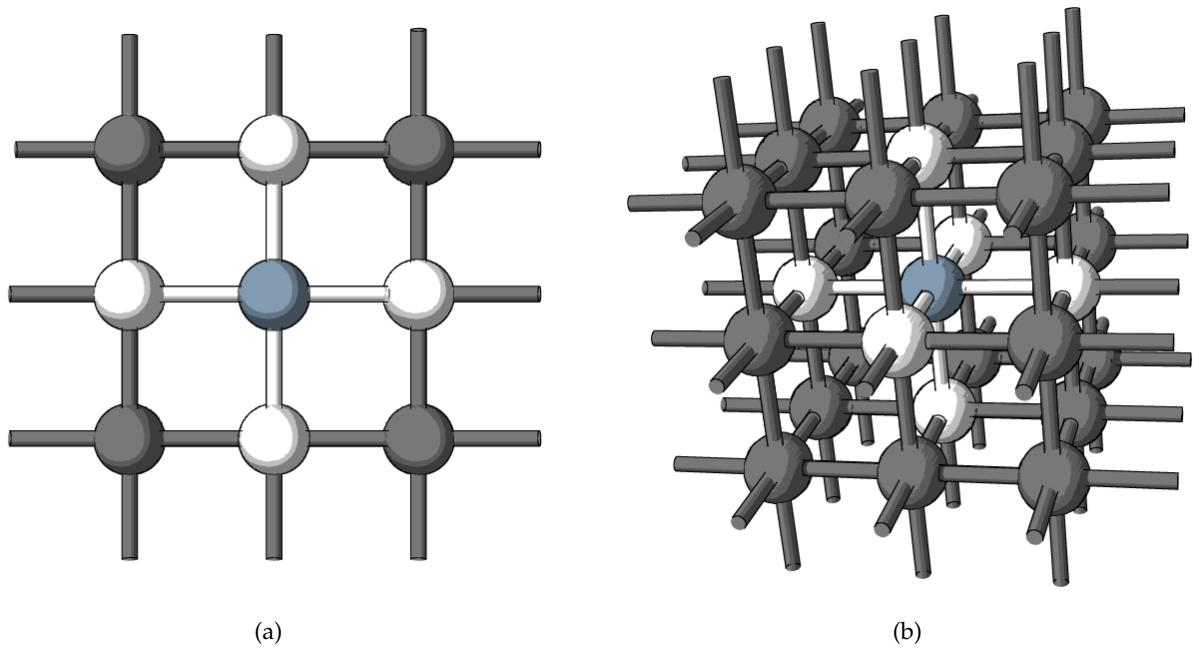


FIGURE 3.1: Common lattice structure for 2D and 3D image data. **(a)** Simplest connection of neighbouring pixels for 2D images. Each non-edge pixel is connected to 4 pixels. This is 4-connectedness. **(b)** Simplest connection of neighbouring voxels for 3D images. Each non-edge voxel is connected to 6 voxels. This is 6-connectedness.

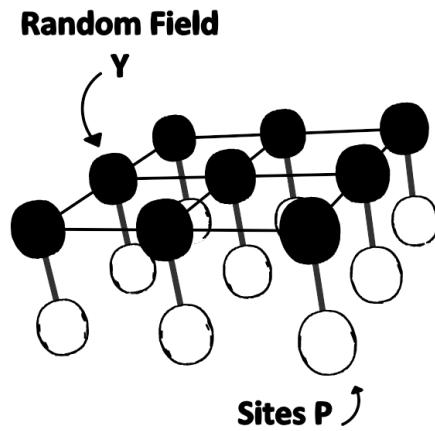


FIGURE 3.2: 4-connected random field \mathbf{Y} over the sites \mathcal{P} .

the most probable labelling yields the lowest energy. Obtaining the optimal labelling from Equation (3.8) does not guarantee that the segmented output will be good. The design of a good energy function that captures all constraints and priors is not easy. However, optimisation is harder still.

From Equation (3.6), if we take the negative log we get:

$$-\log(\Pr(x|y)) = \log(Z) + \sum_{i \in \mathcal{V}} \Psi(x_i) + \sum_{(i,j) \in \mathcal{E}} \Psi_{i,j}(x_i, x_j), \quad (3.9)$$

the constant Z is not needed as it does not affect the final labelling. In this form, the equation is a sum of potentials, i.e. a sum of energies. The first term encodes the data constraints, E_{data} , and

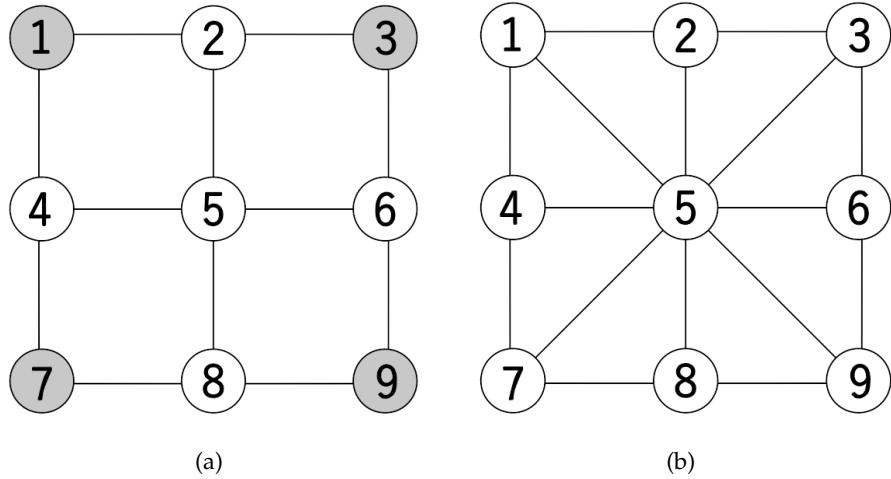


FIGURE 3.3: 2D Neighbourhood structure and node ordering. **(a)** 4-Connected. **(b)** 8-Connected.

the second term encodes the prior constraints, E_{prior} . In addition, there is a factor that controls the relative importance between the data and the prior, λ . The general form of the energy equation is then:

$$E(f) = E_{data}(f) + \lambda E_{prior}(f), \quad (3.10)$$

where f is a particular labelling. The factor λ encodes our belief in the prior i.e. the larger λ is, the more we believe in the prior information. The data energy takes on the following form:

$$E_{data}(f) = \sum_{i \in \mathcal{V}} \Psi(x_i) = \sum_{p \in \mathcal{P}} D_p(f_p). \quad (3.11)$$

D_p measures the level of agreement between the label f_p and the pixel p . A common approximation is to assume independency observations, and this makes designing D_p relatively straightforward. The only restriction is $D_p(f_p) \in \mathbb{R}^+$. The prior energy takes on the following form:

$$E_{prior}(f) = \sum_{(i,j) \in \mathcal{E}} \Psi_{i,j}(x_i, x_j) = \sum_{\{p,q\} \in \mathcal{N}} V_{\{p,q\}}(f_p, f_q). \quad (3.12)$$

$V_{\{p,q\}}(f_p, f_q)$ is known as the *neighbourhood interaction function*. The aim of this function is to encourage neighbouring random variables to take on the same label, i.e. it penalises neighbouring pixels p and q if they have different labels. The form of $V_{\{p,q\}}(f_p, f_q)$ is application dependant and is trickier to design. In image segmentation, the most common prior is that of smoothness, i.e. intra-object pixel intensities are assumed to be the same or vary gradually within some range, it is at edges or boundaries where this assumption is violated. The general form in Equation (3.10) can be rewritten as:

$$E(f) = \sum_{p \in \mathcal{P}} D_p(f_p) + \lambda \sum_{\{p,q\} \in \mathcal{N}} V_{\{p,q\}}(f_p, f_q). \quad (3.13)$$

When we say MAP-MRF estimation we generally mean energy minimisation. Energy minimisation is a non-trivial task given the intractability of the search in the solution space. Energy minimisation can be categorised into *global energy minimisation* and *local energy minimisation*. We briefly discuss some of the energy minimisation techniques.

Iterated Conditional Modes (ICM) This is a deterministic method that converges to a local minimum [160]. It is a greedy technique that was introduced into vision by Besag [156, 161]. The algorithm iteratively chooses the label that results in the largest decrease in energy at each site until convergence. It is extremely sensitive to initialisation as the dimensionality of the space increases with non-convex energies.

Simulated Annealing This is a stochastic optimisation method that simulates the annealing of a material. It is one of the only general-purpose energy minimisation methods. It was developed and published independantly by Černý [162] and Kirkpatrick *et al.* [163] and was introduced into computer vision by Geman and Geman [155]. The algorithm is initialised with a random labelling. Each pixel is then visited and a local random change is made. If the change results in a lower energy, then it is accepted, else the change is accepted based on a probability parameter, i.e. the temperature. With certain cooling schedules the global minimum can be obtained. However, this is horrendously slow in practice, so sub-optimal schedules are used instead [155].

Genetic Algorithms (GAs) GAs have been successfully employed in energy minimisation for image segmentation [164–168]. GAs work by performing simultaneous local searches that optimise the energy function via a random walk in the search space. The algorithm terminates by choosing the search that found the lowest energy for the energy functional. Their drawback is their inability to guarantee a global optimum [169].

Gradient Descent Explicit differentiation under the Euler-Lagrange equations can be used to obtain a solution [169]. Each modified energy functional must be accompanied by derivation of obtaining a minimum [89, 90, 170, 171]. With an artificial time-step, this algorithm deforms a shape, using the gradient descent process, which is equated to the set of Euler-Lagrange equations. When the deformable models come to rest the equations are satisfied. There are two common drawbacks with this method: Firstly, image noise can severely hinder the gradient descent process and this could lead to instability of the deformation process. Secondly, increasing the number of dependant variables increases the complexity of the search space and time to converge to an optimal solution as there are more derivatives to calculate [169].

Loopy Belief Propagation (LBP) The belief propagation algorithm was initially designed to be used on acyclic graphs where it is able to obtain a global minimum [172]. The same algorithm has been successfully applied on cyclic graphs firstly for error-correcting code problems [173] and then later on in vision [174]. Convergence is not guaranteed as the algorithm might get stuck alternating between two labels [172].

Graph Cuts Graph cuts have become an indispensable tool in computer vision. For a restrictive class of energy functions, *submodular functions*, it is able to obtain a global minimum [25, 175–177].

For non-submodular energy functions, it is able to find approximate solutions with strong local optimality [146, 178–180]. Greig *et al.* was the first to use graph cuts in vision to find an exact solution to a certain energy function for the binary image restoration problem [181]. However, it did not receive much attention and remained buried for almost ten years primarily because of the disinterest in binary image restoration and, at the time, its optimisation was notoriously slow which made it an unappealing technique when compared to stochastic optimisation methods, like simulated annealing, which was popular at the time. In the last two decades, graph cut optimisation has been a major focus as a key tool in optimisation since Roy and Cox [182] used it to solve more interesting problems in multi-camera stereo. Shortly after, Boykov *et al.* generalised the method for determining the MAP estimate of MRFs [183]. Graph cut optimisation is the technique of focus in this dissertation.

3.3 Introduction to Graph Cuts

Graph cuts is a combinatorial optimisation method which can be used to minimise energies of the form presented in Equation (3.13). The aim of graph cuts is to partition a graph into mutual exclusive subgraphs by removing the edges whose sum of capacities is a minimum. We are interested in cutting the graph into two subgraphs. Graph cut algorithms existed long before their use was employed in vision, this is primarily due to the lack of computational power available at the time. Fortunately, computational power is no more as rare a resource as it was previously and this has paved a way into exploiting the power of graph data structures. As a result, this has also lead to vision-specific, graph-cut algorithms. There are primarily three types of graph-cut algorithms: *Augmenting Path Algorithms*, such as Ford-Fulkerson algorithm [184], Dinic algorithm [185], Edmond-Karps algorithm [186] etc, *Preflow-Push Algorithms*, also known as Push-Relabel[187], and *Move-Making Algorithms* such as the α - β Swap [146], α -Expansion [146], etc.

3.3.1 Network Theory and the Min-cut Problem

In this section, we briefly cover the foundation aspects to understanding graph cuts. We cover *Flow networks*, a branch of Graph Theory also known as *Transportation networks*, and introduce the *Min-cut problem*. For a solid understanding in Graph Theory and Flow Networks see [188–191]. A brief introduction is given in Appendix A.

Network A network $\mathbf{N} = (V, E)$ is a directed graph with a source node s , a sink node t and a strictly positive capacity on every edge. That is, for each edge $e \in E$, the capacity, $c(\cdot)$, obeys $c(e) \in \mathbb{R}^+$. The **source node** only has out-going edges, $d_{in}(s) = 0$ and $d_{out}(s) \geq 0$. The **sink node** only has incoming edges, $d_{in} \geq 0$ and $d_{out} = 0$. An example of a network is illustrated in Figure 3.4.

Flow A flow $f : V^2 \rightarrow \mathbb{R}^+$ is associated with each edge $e = (u, v)$ such that:

1. for each edge $e \in E$ we have $0 \leq f(e) \leq c(e)$. That is, the flow is positive and cannot exceed the capacity of the edge.
2. for each intermediate node $v \in V \setminus \{s, t\}$ the in- and out-flow of that node $\sum_{u \in V^-(v)} f(u, v) = \sum_{u \in V^+(v)} f(v, u)$.

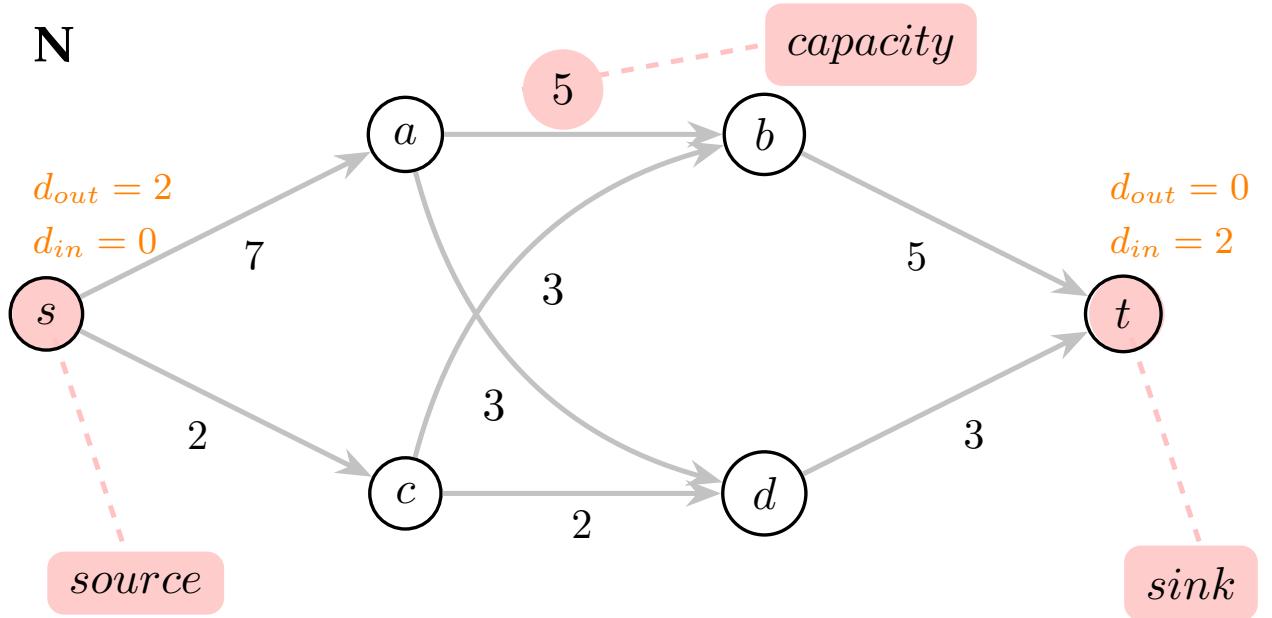


FIGURE 3.4: Network \mathbf{N} with no flow. The in-degree and out-degree for the source, s , and the sink, t , are shown next to the corresponding node.

The **total flow** F of a network is then what leaves the source s or reaches the sink t :

$$F(\mathbf{N}) := \sum_{u \in V} f(s, u) - \sum_{u \in V} f(u, s) = \sum_{u \in V} f(u, t) - \sum_{u \in V} f(t, u) \quad (3.14)$$

An example of a network with non-zero flow is illustrated in Figure 3.5.

Cut A cut of a network $\mathbf{N} = (V, E)$ is a partitioning of the vertex set $V = P \cup \bar{P}$ into two disjoint sets P containing the source node s and \bar{P} containing the sink node t . $P \cap \bar{P} = \emptyset$. The **cost** of a cut is the sum of the capacity of the edges $(u, v) \in V$ where $u \in P$ and $v \in \bar{P}$:

$$\kappa(P, \bar{P}) = \sum_{u \in P; v \in \bar{P}} c(u, v) \quad (3.15)$$

A network with a valid cut is illustrated in Figure 3.6. Invalid cuts are shown for the same network in Figure 3.7 and Figure 3.8.

Maximal Flow The largest amount of flow that is able to reach the sink from the source is known as the maximal flow. A network with a maximal flow, also known as a *max-flow*, is illustrated in Figure 3.9.

Minimal Cut A cut C on a network $\mathbf{N} = (V, E)$ is a minimal cut if there exists no other cut C' where $\kappa(C') < \kappa(C)$. A network with a minimal cut, also known as a *min-cut*, is illustrated in Figure 3.10.

In Figure 3.9 and Figure 3.10 the maximum flow and the minimum cut yield the same answer. This is not a coincidence. In fact, the two problems are duals of each other, known as the *max-flow*

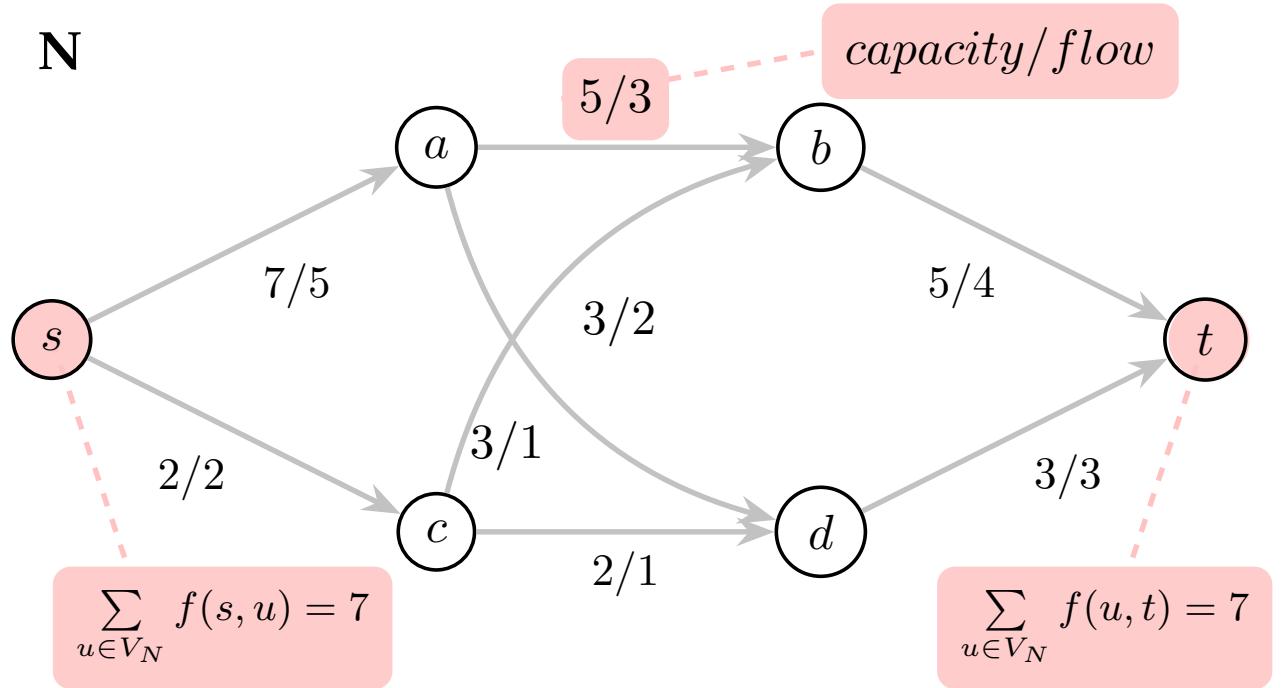


FIGURE 3.5: Network N with flow. The flow out of the source node, s , is equal to the flow into the sink node, t . For all other nodes, the flow-in is equal to the flow-out. This is the conservation of flow principle. This is only part of the network. The remaining part is the residual graph which shows the amount of reverse flow is available on an edge.

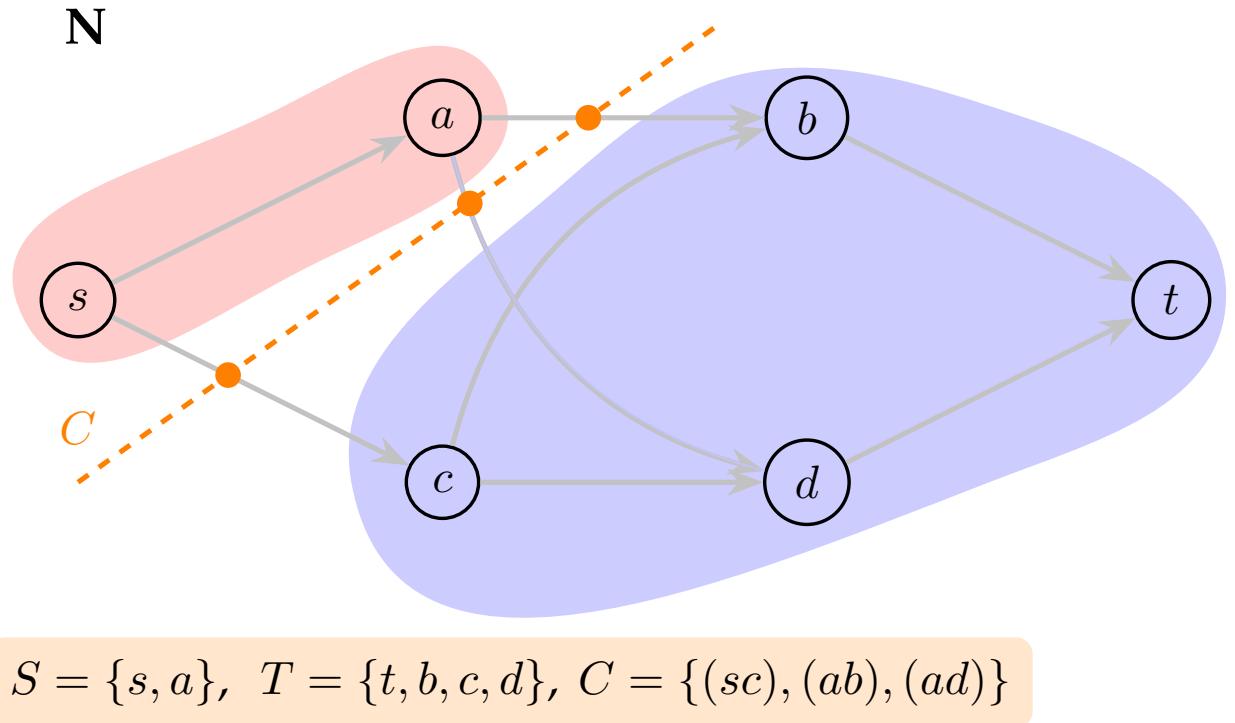


FIGURE 3.6: Network N with a valid cut C . The nodes within the red region are reachable from the source and the nodes within the blue region are able to reach the sink. The cut set, C , is show in the orange-filled block.

min-cut duality. This was proven by P. Elias, A. Feinstein, and C.E. Shannon [192] and by Ford and Fulkerson [184] independently in 1956. This duality is immensely helpful in developing machine

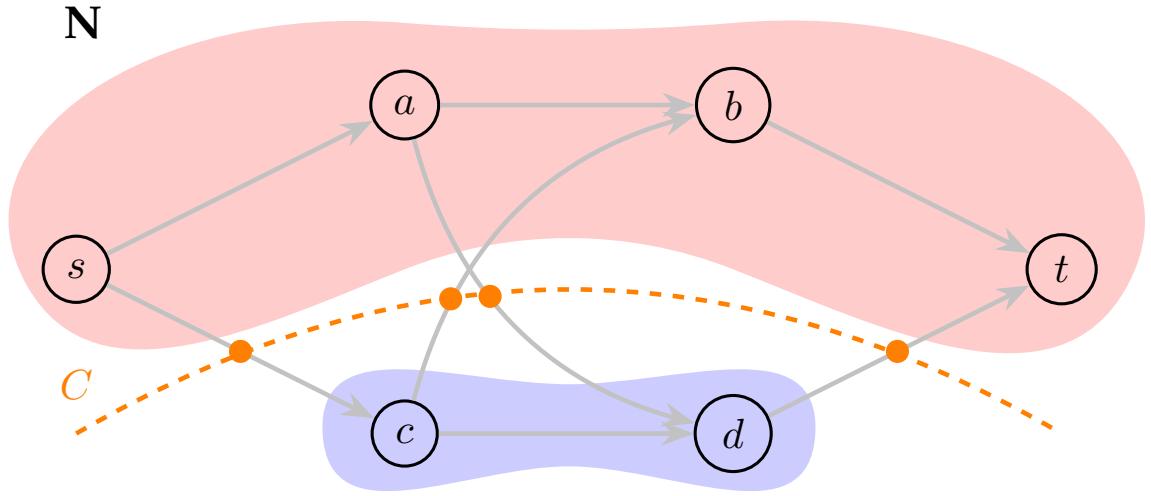


FIGURE 3.7: Network \mathbf{N} with an invalid cut \mathbf{C} . The cut does not partition source node s and sink node t into distinct sets.

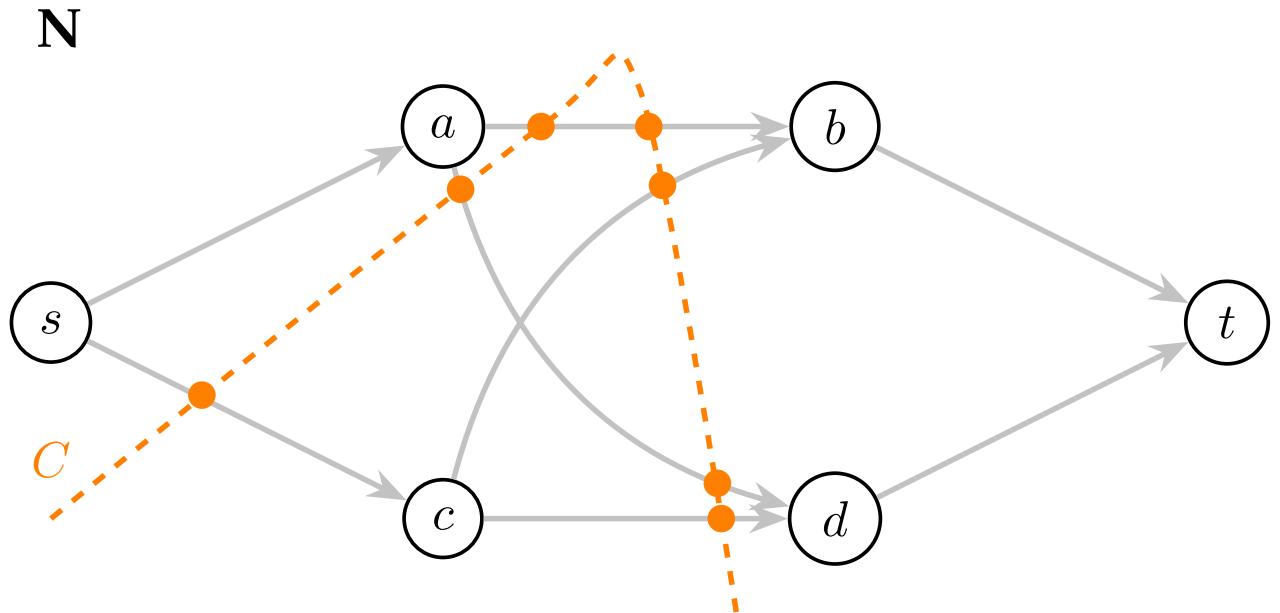


FIGURE 3.8: Network \mathbf{N} with an invalid cut \mathbf{C} . The cut partitions the graph into more than two sets and the cut intersects the edges ab and ad twice.

algorithms to compute the minimum, since it is easier to find a maximum flow. It is important to realise that there may be many cuts that are minimum cuts; and many flow configurations that can yield the maximum flow, hence the solution is not guaranteed to be unique, only an optimum.

3.3.2 Image Segmentation Graph Structure

In graph cut image segmentation the energy function, defined by the image, has to be represented as a graph, specifically a network. We now look at how 2D binary segmentation energy is constructed as a graph. The graph that is constructed uses the MRF, Section 3.2.1, model of the image as its base. Each pixel is a node in the graph. The connections between "pixel nodes" are bi-directional, which

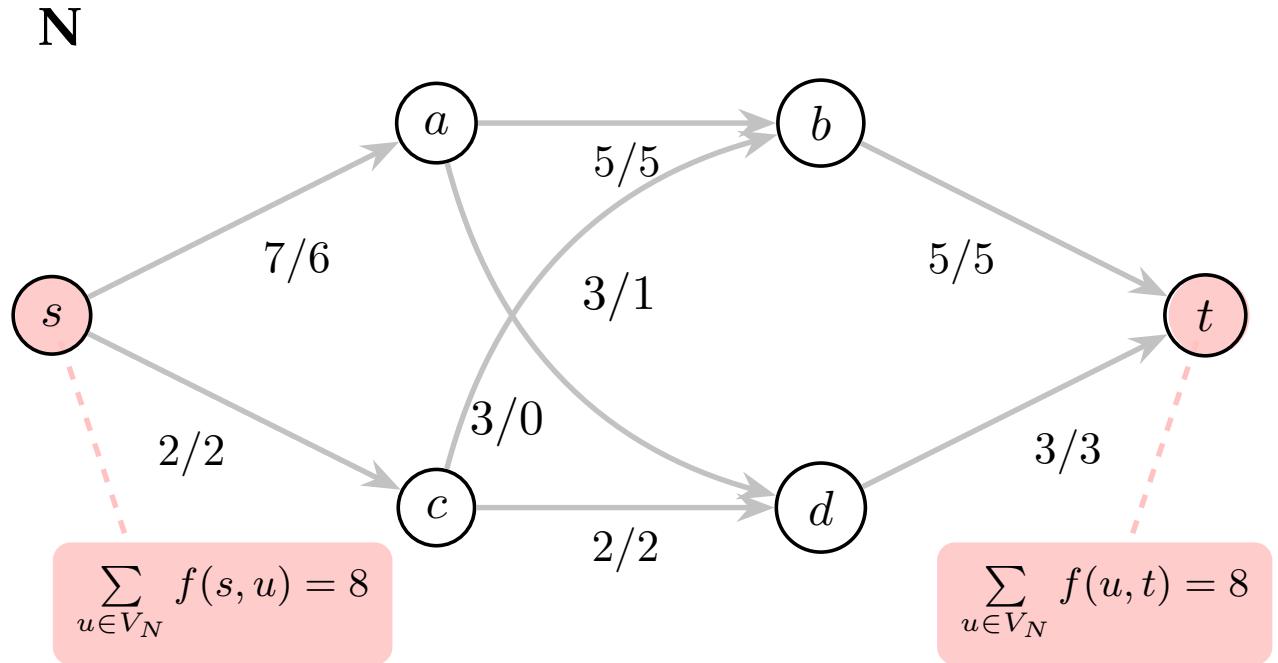


FIGURE 3.9: Network **N** with maximum flow. There is no way to push more flow out of the source into the sink without breaking the rules for the conservation of flow.

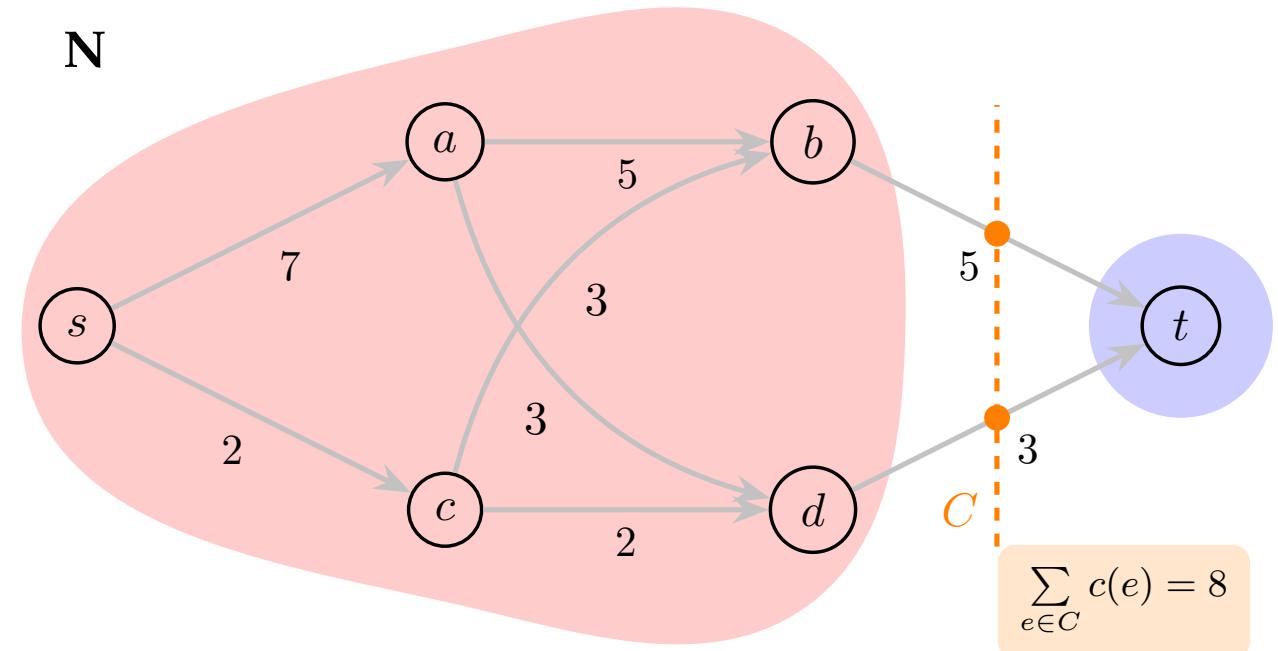


FIGURE 3.10: Network **N** with minimal cut **C**. The sum of the capacity of all the edges in the cut set is the minimum of all possible valid cuts on the network **N**.

is generally decomposed into two uni-directional edges, and are known as *n-links* which is synonymous with *neighbour-links*. Additionally, each label is also represented as a node. Each "pixel node" is attached to all "label nodes". In binary segmentation, there are two labels i.e. object and background. The edges that connect "label nodes" to "pixel nodes" are called *t-links* which is synonymous with

terminal-links. In keeping with network construction, one label will be the source and the other will be the sink. The t-link weights are generally learned from user input seeds or automatically generated. Seeds mark which type of pixels belong to the object and which belong to the background, an illustration is shown in Figure 3.11(a). With the seed data and the neighbourhood interaction, we can construct a graph representation of the energy to be minimised over the image as illustrated in Figure 3.11(b). Once a graph has been constructed, we then call upon a max-flow/min-cut method which will minimise the energy function by partitioning the graph into two subgraphs, as shown in Figure 3.11(c). All pixels are classified according to which "label node" they're still attached to after the max-flow/min-cut algorithm has run, this is illustrated in Figure 3.11(d).

3.3.3 Submodular Functions

In the previous section, we talked about representing the energy function as a graph. However, not all energy functions can be represented as a graph. Moreover, minimising an arbitrary energy function, even if the energy is binary, is NP-hard [193]. There does exist a class of functions which is graph representable and is able to be minimised in polynomial time, i.e. an exact global minimum can be obtained in a single graph cut. These energy functions are known as *submodular functions*. They are sometimes referred to as "discrete analogue of convex functions" since they're the easiest to minimise, much like convex functions. For an energy to be submodular, it must satisfy the submodularity constraint:

$$f^p(a, b) + f^p(a + 1, b + 1) \leq f^p(a, b + 1) + f^p(a + 1, b), \quad \forall a, b \in \mathcal{L}, \quad (3.16)$$

where \mathcal{L} is an ordered label set. The type of energies we are concerned with are second order binary energies, and enforcing the submodularity constraint means that the energy function must satisfy:

$$E_{ij}(0, 0) + E_{ij}(1, 1) \leq E_{ij}(0, 1) + E_{ij}(1, 0), \quad \forall \{i, j\} \in \mathcal{N}, \quad (3.17)$$

where $\mathcal{L} = \{0, 1\}$. It is necessary and sufficient for an energy function to satisfy Equation (3.17) to compute the exact global minimum in polynomial time in a single graph cut. This was first characterised by Kolomogorov and Zabih [175].

3.4 Graph Cut Algorithms for Energy Minimisation

Image segmentation via graph cuts can be seen as three main stages. Firstly, the problem is modelled and a suitable energy function is designed. Generally, this is a trade-off between accuracy and constraining to the submodularity constraint. Secondly, a graph is constructed which represents the energy function. Thirdly, and finally, the energy function is minimised by using a max-flow/min-cut algorithm. In this section we discuss the final stage. The minimisation algorithms are either general or specific to the energy functions. We briefly discuss some of the most common max-flow algorithms.

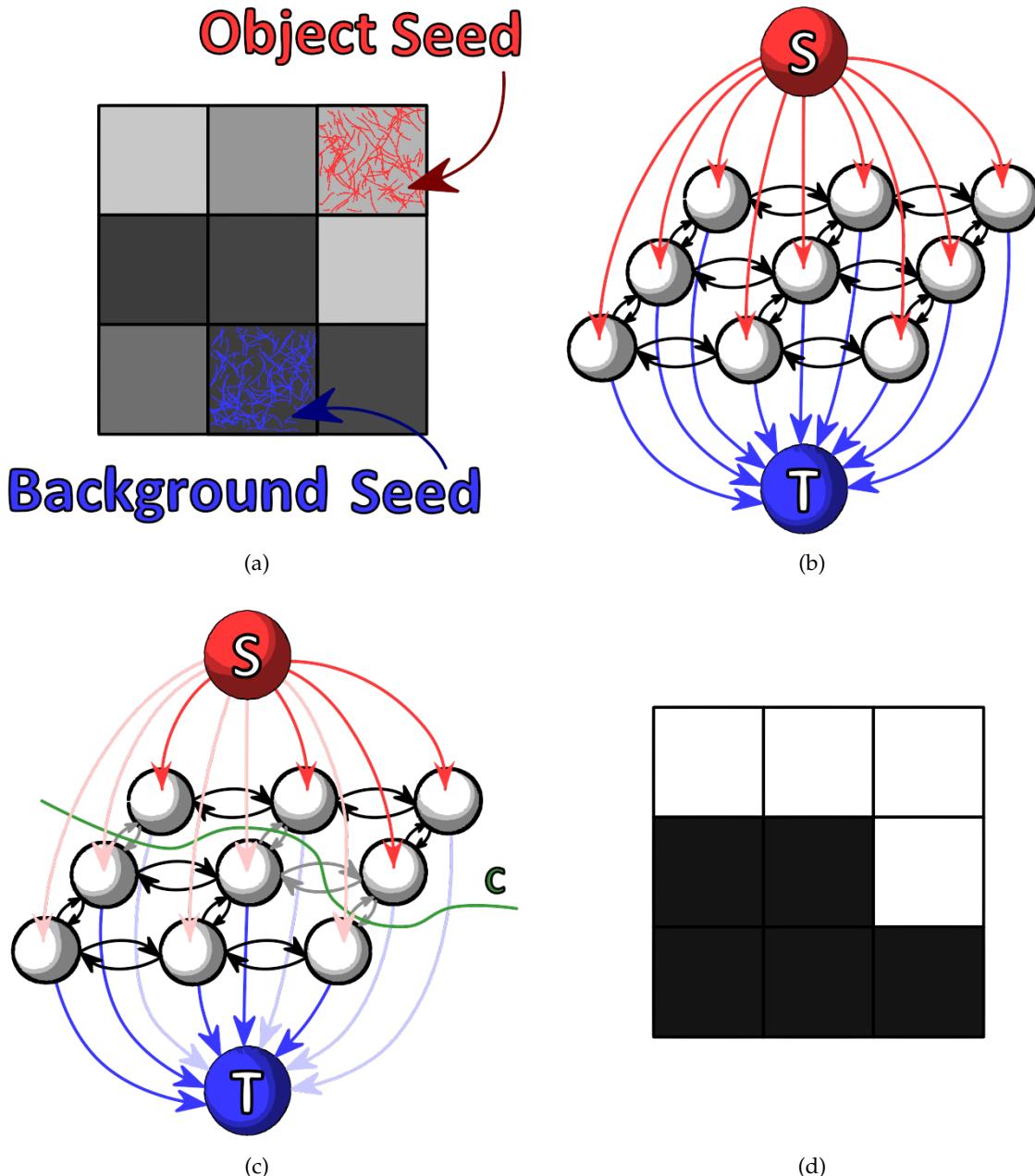


FIGURE 3.11: Overall process of the graph cut image segmentation process. **(a)** Image with object and background seeds. **(b)** Graph representation of the energy function to be minimised over the image. The n-links are represented by black arrows, the t-links from the source are shown in red, and the t-links to the sink are shown in blue. **(c)** Minimised energy function by cut C. The light red, light blue and grey edges are those that belong to the cut set C. **(d)** Segmentation mask after label assignment. In this case, for all nodes that are still attached to the source, their corresponding pixel label is shown in white, similarly, nodes that are connected to the sink have their corresponding pixel labels shown in black.

3.4.1 Ford-Fulkerson

The Ford-Fulkerson max-flow algorithm [184] was designed to work on an arbitrary network with one source and one sink. Ford and Fulkerson proved that the max-flow and min-cut problems are duals of each other, hence solving one means that you have obtained the solution to the other. Augmenting path algorithms iteratively search for open path from the source to the sink. When a path is

Algorithm 1 Ford-Fulkerson Max-flow

```

1: procedure MAXFLOW( $G$ )                                 $\triangleright$  The maximum flow on graph  $G$ 
2:   while  $p = \text{findPath}(s, t)$  do                       $\triangleright$  Find an open path  $p$  between  $s$  and  $t$ 
3:     if  $p = \emptyset$  then                                 $\triangleright$  See if a solution has been found
4:       break
5:     end if

6:      $\text{maxflow}_p = 0$ 
7:     for each edge  $e \in p$  do                           $\triangleright$  Find max-flow on path  $p$ 
8:        $\text{maxflow}_p \leftarrow \min(\text{maxflow}_p, c(e) - f(e))$      $\triangleright$  Residual capacity on edge  $e$ 
9:     end for

10:    for each  $e \in p$  do                             $\triangleright$  Push max-flow on path  $p$ 
11:       $f(e) \leftarrow f(e) + \text{maxflow}_p$             $\triangleright$  Augment each path on path  $p$  with flow  $\text{maxflow}_p$ 
12:    end for
13:  end while
14: end procedure

```

Algorithm 2 Min-cut from Max-flow

- 1: Calculate max-flow on \mathcal{G}
- 2: Partition \mathcal{V} into \mathcal{V}_S and \mathcal{V}_T
- 3: $\mathcal{C} = \{(u, v) \in \mathcal{E} | u \in \mathcal{V}_S \wedge v \in \mathcal{V}_T\}$

found, the maximum flow that can be pushed in that path is pushed through that path by incrementing the flow on each edge of that path. The path is said to be *augmented*. If there are no available paths which can be augmented, the solution has been found and the algorithm terminates. The maximum flow is then obtained by the flow leaving the source or the flow entering the sink. A Ford-Fulkerson max-flow algorithm is shown in Algorithm 1. The min-cut, and segmentation, problem can be solved using Algorithm 2. In image segmentation we only require the sets \mathcal{V}_S and \mathcal{V}_T , so the last step is not necessary. The Ford-Fulkerson algorithm exhibits very poor performance. The primary cause is that each new augmenting path has to be found from scratch. In [176], they countered this by reusing as much information as possible from the existing path.

3.4.2 Dinic/Edmonds-Karp

The Ford-Fulkerson max-flow algorithm is a very inefficient method to finding the max-flow. In practice the runtime is too high making it unfavourable. An improvement on this augmenting path based algorithm was designed by Dinic[185] in 1970 and independently by Edmonds and Karp[186] in 1972. The improvement is a change in searching for a path which can be augmented. The Edmonds-Karp algorithm finds the shortest path from the source to the sink by defining the length of all edges to be one and using a breadth first search from the source. The amount of flow to be augmented into each path is the minimal residual capacity of the edges in the path. This ensures that on each augmentation at least one path is *saturated*. The edges that become saturated are said to be *critical*. The Edmonds-Karp max-flow algorithm is shown in Algorithm 3. The worst-case complexity is $O(|\mathcal{V}|, |\mathcal{E}|^2)$. Depending on the graph structure other searching strategies may be more efficient, e.g. depth first search (DFS) [194], etc.

Algorithm 3 Edmonds-Karp Max-flow

```

1: procedure MAXFLOW( $G$ )                                 $\triangleright$  The maximum flow on graph  $G$ 
2:   while  $p = BFS(s, t)$  do                          $\triangleright$  Find the shortest and open path  $p$  between  $s$  and  $t$ 
3:     if  $length(p) = 0$  then                            $\triangleright$  See if a solution has been found
4:       break
5:     end if

6:      $minrescap_p = \infty$ 
7:     for each edge  $e \in p$  do                       $\triangleright$  Find augmenting flow to saturate an edge
8:        $minrescap_p \leftarrow \min(minrescap_p, c(e) - f(e))$      $\triangleright$  Residual capacity on edge  $e$ 
9:     end for

10:    for each  $e \in p$  do                          $\triangleright$  Push flow on path  $p$ 
11:       $f(e) \leftarrow f(e) + minrescap_p$             $\triangleright$  Augment each edge with flow  $minrescap_p$ 
12:    end for
13:  end while
14: end procedure

```

3.4.3 Push-Relabel

The Push-Relabel was developed by Andrew V. Goldberg and Robert E. Tarjan [187]. Previous algorithms, such as Ford-Fulkerson, use the concept of residual networks and augmenting paths to determine max-flow. Push-Relabel uses the concept of preflow to determine max-flow instead of augmenting paths. Sometimes referred as the *Preflow-Push Algorithm*. Preflow is a concept originally developed by A.V. Karzanov.

The algorithm works at converting a preflow, f , into a normal flow and then terminates. This flow also turns out to be the maximum flow. Goldberg and Tarjan defined a generic Push-Relabel algorithm which solves the maximum flow problem.

Preflow A preflow is a real-valued function, f , on vertex pairs. The total flow into a vertex can exceed the flow out of a vertex but not vice versa.

A preflow where all $v \in V - \{s, t\}$ has a flow excess of zero, $e_f(v) = 0$, is a normal flow. The preflow function is also referred to as the **s-t preflow**.

Preflow must satisfy:

1. Capacity Constraint

$$\forall u, v \in V, f(u, v) \leq c(u, v)$$

2. Anti-symmetry/Skew Symmetry

$$\forall u, v \in V, f(u, v) = -f(v, u)$$

3. Non-negative Constraint

The flow into $v \in V - \{s\}$ must be greater than or equal to the flow out of v . $\forall u \in V, v \in V - \{s\}, \sum f(u, v) > 0$

Flow Excess Flow excess, $e_f(v)$, is the net flow into v where $v \in V$ for some preflow f .

$$e_f(v) = \begin{cases} \infty & \text{if } v = s \\ \sum_{u \in V} f(u, v) & \text{if } v \in V - \{s\} \end{cases}$$

Active Vertex An active vertex/node is a vertex v which satisfies all of the properties:

1. Not a source or sink, $v \in V - \{s, t\}$
2. Positive flow excess, $e_f(v) > 0$
3. Has a valid label, $d(v) < \infty$

Push-Relabel also uses the concept of a residual graph, $G_f^* = (V, E_f)$.

Residual Capacity The residual capacity of a preflow is defined as $r_f(v, w) = c(v, w) - f(v, w)$.

Residual Edges The residual edges for a preflow f is defined as the set of edges with positive residual capacity. $E_f = \{(v, w)\} | r_f(v, w) > 0$.

Labelling Push-Relabel also use a valid labelling function, d , to determine which vertex pairs should be selected for the push operation.

A valid labelling, d , is a non-negative integer function applied to all vertices to denote a label. The labelling is often referred as the height or distance from the sink node, t . This function is sometimes compared to the physical intuition that liquids naturally flow downhill.

A valid labelling for a preflow consists of:

1. For $v \in V, 0 \leq d(v) \leq \infty$
2. $d(s) = |V|$ (source condition)
3. $d(t) = 0$ (sink condition)
4. $d(v) = d(w) + 1$ for every residual edge $(v, u) \in E_f$

A labelling d and a preflow f are said to be compatible if d adheres to the properties above.

The algorithm pushes flow excess starting at the source, s , along all vertices towards the sink, t . The algorithm maintains a compatible vertex labelling function, d , to the preflow, f . The labelling is used to determine where to push the excess flow. The algorithm repeatedly performs either a push or a relabel operation so long as there is an active vertex in G_f^* .

Push Operation The push operation is used to move flow from one vertex to another. The transfer of excess can be performed across the vertex pair $(v, w) \in E_f$ if:

1. v is an active vertex
2. the edge has positive residual capacity, $r_f(v, w) > 0$
3. the label distance $d(v) = d(w) + 1$

Algorithm 4 Push Operation

Input: Preflow f , labels d , and (v, w) where $v, w \in V$ **Output:** Preflow f **Applicable:** if $v \in V - \{s, t\}$, $d(v) < \infty$, $e_f(v) > 0$, $r_f(v, w) > 0$ and $d(v) = d(w) + 1$

```

1: procedure PUSH( $G, G^*, v, w$ )
2:    $\delta \leftarrow \min(e_f(v), r_f(v, w))$                                  $\triangleright$  Find the maximum flow can be pushed from  $v$ 
3:    $f_G(v, w) \leftarrow f(v, w) + \delta$                                  $\triangleright$  Push flow and update residual graph
4:    $f_{G^*}(w, v) \leftarrow f(w, v) - \delta$                              $\triangleright$  Update excess on  $G$  and its residual graph  $G^*$ 
5:    $e_G(v) \leftarrow e_f(v) - \delta$ 
6:    $e_{G^*}(w) \leftarrow e_f(w) + \delta$ 
7:   return  $\delta$ 
8: end procedure

```

Algorithm 5 Relabel Operation

Input: Preflow f , labels d , and $v \in V - \{s, t\}$ **Output:** Labels d **Applicable:** if $v \in V - \{s, t\}$, $d(v) < \infty$, $e_f(v) > 0$, and $\forall w \in V, r_f(v, w) > 0$ which implies $d(v) \leq d(w)$

```

1: procedure RELABEL( $v$ )
2:    $d(v) \leftarrow \infty$ 
3:   for each vertex  $w \in \mathcal{N}_v$  do                                 $\triangleright$  Consider all the neighbours of  $v$ 
4:     if  $\{(v, w) \in E_f\} \neq 0$  then                                 $\triangleright$  Is  $w$  reachable from  $v$ 
5:        $d(v) \leftarrow \min(d(v), d(w) + 1)$ 
6:     end if
7:   end for
8:   return  $d$ 
9: end procedure

```

This allows the algorithm to move δ excess flow: $\delta = \min(e_f(v), r_f(v, w))$ from v to w . A push is considered *saturating* if no more flow can be sent over the edge, $\delta = r_f(v, w)$. A push is considered to be *non-saturating* if all the excess from v the push over the edge and the edge still has some capacity, $\delta = e_f(v)$. The push operation is shown in Algorithm 4.

Relabel Operation The relabel operation is used to increase the label value of a single active vertex so that excess flow can be pushed out of the active vertex. The relabel operation is performed when all the residual edges of the active vertex have positive residual capacity, $r_f(v, w) > 0$. This implies that v 's label is less than or equal to all vertices, $d(v) \leq d(w)$, meaning that no push operation across the edges is possible given the push condition $d(v) = d(w) + 1$.

The relabel operation for some vertex v selects the smallest label for the vertices with positive residual edges, $r_f(v, w) > 0$. The active vertex is then assigned the smallest label value +1 such that $d(v) := \min\{d(v) + 1 | (v, w) \in E_f\}$. This will allow the vertex v to potentially push its excess flow to at least one of the other vertices during the algorithm's next iteration. The relabel operation is shown in Algorithm 5.

Discharge Operation The coordination of pushing excess flow from and relabelling a vertex is handled in a discharge operation. The idea is to push as much excess flow, from the currently picked

Algorithm 6 Discharge Operation

Input: $v \in V - \{s, t\}$ **Output:** State of node**Applicable:** if $v \in V - \{s, t\}$, $d(v) < \infty$, $e_f(v) > 0$

```

1: procedure DISCHARGE( $v$ )
2:    $i \leftarrow v.current\_neighbour$             $\triangleright$  index of the current neighbour under consideration
3:   while ( $e(v) > 0$ )  $\wedge$  ( $i < \text{size}(v.neighbour)$ ) do            $\triangleright$   $neighbour$  is the list of neighbours
4:     if ( $d(v) == d(v.neighbour[i]) + 1$ )  $\wedge$  ( $r_f(v, v.neighbour[i]) > 0$ ) then
5:       push( $v, v.neighbour[i]$ )
6:        $i \leftarrow (i + 1)$ 
7:     end if
8:     if  $e(v) > 0$  then
9:       relabel( $v$ )
10:    end if
11:     $v.current\_neighbour \leftarrow i$             $\triangleright$  Pick up from this neighbour on the next discharge
12:    return  $e(v) > 0$                        $\triangleright$  Is node still active
13:   end while
14: end procedure

```

active vertex, to its neighbours. If no more flow can be pushed but the node is still active, then relabel it. The discharge operation is shown in Algorithm 6.

The algorithm also maintains a list of active nodes to discharge. The complete Push-Relabel algorithm is shown in Algorithm 7. The algorithm repeatedly pushes flow between nodes until there are no more active nodes.

There are many variations of the push-relabel algorithm [187, 195]. Most of these are heuristic implementations to reduce execution time. One such heuristic is the *First-in First-out* (FIFO) [187]. With this implementation, the theoretical run-time is shown to be $O(|V|^3)$. Cubic complexities are unacceptable in practice, however in practice this heuristic displays a great boost in speed optimisation. Empirical runtime over theoretical runtime is accepted in practice.

Instead of discharging nodes in the order that they entered the *Active_node* list, a better heuristic is to discharge nodes in descending order of node level/height. This heuristic is known as *Highest level first* (HLF). Flow is pushed from higher level nodes to lower level nodes first. This allows a greater dissipation of flow.

Another popular heuristic is *Global relabel*. In this heuristic, all nodes are periodically labelled from the distance of the node to the sink. This is can be done by running a BFS from the sink node t in the residual graph G^* . When used in conjunction with the FIFO heuristic, there's a huge performance speedup. The Global relabel heuristic is expensive to perform so it is only run after every $|V|$ relabel operations.

The final heuristic we discuss is known as the *Gap relabel*. This heuristic is based on an important observation, nodes can only push to other nodes that are one level lower. Hence, if there are no nodes of a certain level, d , then it is not possible for higher nodes, at level $d + 1$, to send their flow to the nodes at the next lowest level, $d - 1$, hence this gap in the levels allows us to relabel all nodes that are higher than level d to $|V|$. These relabelled nodes will no more be considered in further push or relabel operations. When the max-flow is determined, graph partitioning can be easily obtained. All

Algorithm 7 Push-Relabel Maxflow Algorithm

Input: $\mathcal{G} = (\mathcal{V}, \mathcal{E})$ **Output:** Maximum flow, f

```

1: procedure MAXFLOW( $\mathcal{G}$ )
2:    $d(s) \leftarrow |\mathcal{V}|$                                  $\triangleright$  Initialise the height of all nodes
3:    $d(t) \leftarrow 0$ 
4:   for  $\forall v \in \mathcal{V} - \{s, t\}$  do
5:      $d(v) \leftarrow 0$ 
6:   end for
7:   for  $\forall v \in \mathcal{N}_s$  do                                 $\triangleright$  Saturate all outgoing edges from the source
8:      $d(v) \leftarrow 1$ 
9:      $f_G(s, v) \leftarrow c(\text{edge}(s, v))$ 
10:     $f_{G^*}(v, s) \leftarrow -c(\text{edge}(s, v))$ 
11:     $e_G(s) \leftarrow -c(\text{edge}(s, v))$ 
12:     $e_G(v) \leftarrow c(\text{edge}(s, v))$ 
13:    Append  $v$  to Active_node                                 $\triangleright$  Build the active node list
14:  end for
15:  while  $\text{size}(\text{Active\_node}) > 0$  do                 $\triangleright$  While there are active nodes, discharge them
16:     $\text{current\_node} \leftarrow \text{pop\_front}(\text{Active\_node})$        $\triangleright$  Get the first active node
17:     $\text{state} \leftarrow \text{discharge}(\text{current\_node})$ 
18:    if  $\text{state} == \text{ACTIVE}$  then           $\triangleright$  If node is still active then put it at the end of the list
19:       $\text{push}(\text{Active\_node}, \text{current\_node})$ 
20:    end if
21:  end while
22:  return  $e(t)$                                  $\triangleright$  Final flow is equal to the flow entering the sink node
23: end procedure

```

nodes that are equal to or higher than the source are part of the source set, i.e $\mathcal{V}_S = \{v \in \mathcal{V} | d(v) \geq |\mathcal{V}|\}$ and $\mathcal{V}_T = \mathcal{V} - \mathcal{V}_S$.

3.4.4 Move-Making Algorithms

Modern move-making algorithms based on combinatorial graph cuts outperform previous move-making methods, such as simulated annealing and ICM, as well as message passing algorithms because of their increased accuracy and efficiency [196]. Specifically, the α -expansion and $\alpha\beta$ -swap [146] has become very popular and gained a large acceptance in the vision community [197].

In the α -expansion, the algorithm iteratively makes the move to expand the α -label set. The label that provides the largest decrease in energy for all labels is kept. When no further label changes can be made, then the algorithm has reached convergence and terminates. In the $\alpha\beta$ -swap, the algorithm iteratively chooses two pixel-sets, with labels α and β , and swaps the labels of the pixels within these sets. The swap between the pair of labels that results in the largest decrease in energy is chosen. Similarly, when no more swap moves can be made, the algorithm has reached convergence and terminates. For binary segmentation, both of these algorithms require just one iteration to reach convergence and the optimal move is determined by using graph cuts.

Chapter 4

Literature Review

It has long been a desire of mankind to know what occurs at the cellular level. Even to get the smallest glimpse of this world would have been extremely rewarding. This was made possible in the 1670s by Antoni van Leeuwenhoek when he took the first steps to build a microscope [70]. The field of cellular biology has rapidly expanded ever since. In more recent times, the fluorescent microscope has allowed biologists and other professionals alike to not only restrict their studies to cellular and subcellular organisms in great detail, but also to store the data in images and videos. This stored data allows scientists to critically analyse and reanalyse the same data to get the most information out of it. As a result, there is a titanic-size volume of image data to be analysed. The image data is so abundantly variant in heterogeneity, dimensionality, complexity and quality such that manual image processing and analysis cannot keep up in terms of time and quality [70]. Most of the useful cell analysis starts at the cell segmentation level and this is a major bottleneck, in time and quality, that needs to be avoided.

What has fluorescence image segmentation been through? Machine assistance in fluorescence image segmentation is not new and dates as far back as the mid-1950s where thresholding on serialised (1D) data was done to enable mass screening for cervical cancer [74]. Since then, biologists, and other professionals alike, began to rely heavily on computerised image segmentation of cells. A timeline of the major advancements in segmentation and cell segmentation is shown in Figure 4.1.

The shift from 1D image processing to 2D image processing happened in the early 1960s [75] and was still threshold-based. It wasn't until the mid-1960s where feature segmentation and mathematical morphology was tested on fluorescence image data [75, 198]. Thresholding still dominated up until the mid-1970s; when research focus shifted to morphological segmentation [199]. It was also the time when researchers in image processing started to get more interested in fluorescence image processing and started to try out other techniques such as those based on Random Fields [200]. None of the emerging techniques worked well enough with the problems posed in fluorescence imaging until the late 1970s when the Watershed method was used [201]. Watershed segmentation began to gain a lot of momentum in comparison to the other new techniques but thresholding was still the technique of preference due to its, by then, long history. At the same time, variants of thresholding became the new focus of cell segmentation. This started with the histogram thresholding method [202] in 1984 and then Otsu segmentation [203] in 1988.

Much of the segmentation techniques developed were very algorithmically simple with little computational demand. This was because computational power was a rare resource. This ended in 1990 when computational power rapidly increased and continued to increase. This made it possible

for more sophisticated and computationally demanding techniques to be implemented in fluorescence image segmentation; such as Active Contour Models (ACM) [204] in 1992. It was also a time when more morphologically accurate segmentations where required and the previous methods really began to fall short of the required expectations. Cellular biologist, and other professional alike, demanded segmentation of specific objects of interest such as cell nuclei, chromosomes, other specific proteins, etc. It was not until 2005 that Graph Cuts (GC), our technique of focus in this dissertation, began to appear on the cell segmentation scene [205].

Although the challenges in fluorescence image segmentation are very unique to the field, there has not been any segmentation techniques developed to exclusively handle fluorescence images [70]. The techniques used in FM segmentation have always been ported from techniques that were designed to segment other types of data. Segmentation techniques in fluorescence imaging seem to follow other segmentation schemes rather than lead or plot its own course. This can be seen in the timeline in Figure 4.1 where, in 1966, mathematical morphology was used in cell segmentation although it had been discovered two years prior. Similarly, the case for the application of Otsu binarization in cell segmentation is even worse with its first application to the field in 1988 [203], although it had been designed in 1979 [206].

On the other hand, rather than embracing newer techniques, the field of fluorescence image analysis is reluctant to accept the modern and sophisticated techniques. Consequently, most of the fluorescence image segmentation done today uses thresholding methods, watershed techniques, feature detection like edge detection, and region accumulation [70]. However, this has not hindered research and regular progress of applying the more sophisticated segmentation models to fluorescence images. Fortunately, modern segmentation methods are increasingly gaining momentum and becoming the method of preference as of late.

Where is fluorescence image segmentation now? We now focus on the relevant research in the last decade. Much of the research involves combining many segmentation methods or designing a variant of an existing method to eliminate a certain problem [70, 207]. These solutions also tend to be very isolated to the specific problem and resultantly suffer from diminished diversity of application.

In 2009, Peng and Hsu [208] designed a variant of adaptive local thresholding for fluorescence cell micrographs. This was to combat the varying brightness and contrast throughout the image. They compared their technique against Otsu binarization [206], Iterative thresholding method [209], Niblack's thresholding method [210] and the then recent adaptive thresholding based on the variational minmax algorithm [211]. Most of these are very old techniques whose weaknesses are well-known. Also, although their experiments did involve images with significant brightness and contrast issues, all the images are very similar. Their segmentation was not automatic, it did require manual tuning and no process or discussion was given on how they had arrived at their parameters. In their efforts to replace manual segmentation with automatic segmentation, they've instead diverted to a parameter tuning problem; which is just as time consuming.

A similar situation is seen when Du *et al.* [212], in 2010, published their performance evaluation studies by comparing K-means, Otsu, EM and Global Minimisation of the Active Contour Model (GMAC) on fluorescence microscopy cell images. They had used a combination of synthetic and

real data and concluded that GMAC performed the best overall. However, no mention was given as to how some of the parameter values were chosen for GMAC. Considering that the performance improvement by GMAC was slightly better than K-means and Otsu, it would be better to use any of the latter techniques, instead of playing around with parameters for a slight improvement when using GMAC.

In 2011, Dima *et al.* [77] compared several commonly used algorithms in fluorescence image segmentation using their novel bivariate similarity index. They had concluded that K-means and Canny edge detection is superior to Otsu, maximum entropy and isodata. However, they had used very simple images.

In 2012, Alexander *et al.* [213] designed an image segmentation scheme which was designed around global thresholding. They ended up with a hard-coded number to be used in their global thresholding method. If history is any indication, then this has extremely limited use and will fall away in the "not-so-long" term. Research into fluorescence image segmentation has almost guaranteed that there is no "magic number". Instead we seek a method that is flexible enough to accommodate the variation of fluorescence images and still deliver a consistent and expected output.

In 2013, there was an explosion of published material of fluorescence image segmentation. This was accompanied by significant application of graph cut segmentation in the domain. Qi [214] designed a novel segmentation scheme combining graph cuts and the convex shape assumption. The proposed initialisation was based on the Poisson model of intensity for FM images. Upon initialisation, a standard graph cut algorithm would perform the segmentation. He used the Boykov and Funka-Lea energy functions [215]. Afterwards, he performs convexity analysis to split overlapping or touching cell nuclei. This was a successful attempt and proved the ability of graph cuts to handle fluorescence images. His scheme outperformed the Watershed method, both direct and gradient variants, Otsu, Mean threshold, Active masks, Merging algorithm, RC threshold and AS threshold. He also tested on a large dataset of 78 images each of size 1940×1940 . The performance was based on the Hausdorff distance. However, he was unable to get rid of the parameter tuning required for that energy function, λ , which controls the relative importance of regional and boundary terms.

In the same year, graph mining was pitted against Supervised machine learning segmentation, Support Vector Machines (SVM) [216]. Graph mining ended with an 85% accuracy beating SVMs which had a 72% accuracy. The process involves a pre-processing scheme of Gaussian noise removal and background removal by thresholding. Their experiments were done over a small set of 10 images. Moreover, no images nor image segmentation mask or contours were published. One would have to assume that there is limited variation of fluorescent embryonic stem cell images for it to be left out.

In 2014, Preethi *et al.* [217] proposed an automatic segmentation and tracking scheme of fluorescent cancerous cells by Wavelet Otsu model. The scheme involved a pre-process of adaptive histogram equalisation (AHE) and anisotropic diffusion filtering. They used morphological techniques to track cell shape. However, they have not published any object results and no comparison to state-of-the-art or other leading techniques.

In 2015, Raza *et al.* [218] proposed a novel automatic segmentation technique by combining textural information in the wavelet domain to counteract the variable intensity in FM images. They tested their technique against active contours [219] and supervised template matching [220]. They

compared the algorithm on various performance measures. On most measures the proposed system outperformed the competitor methods.

Where is fluorescent image segmentation heading? From the recent research and publications, it seems as if there is a new technique designed to work on each sub-field in which fluorescence images are attained. Instead of converging to a unified solution, there seems to be a divergence that is getting increasingly stronger. Many of the novelties are schemes involving a pre-process and following segmentation technique.

Many solutions also involve manual parameter tuning. There is also a lack of published methodologies for parameter tuning in these schemes.

One can conclude that there is a diverging number of solutions for fluorescence image segmentation.

What is needed, is a segmentation method that performs consistently and delivers expected results on a large diversity of fluorescence images that minimises user-interaction. Ideally, it would eliminate user-interaction.

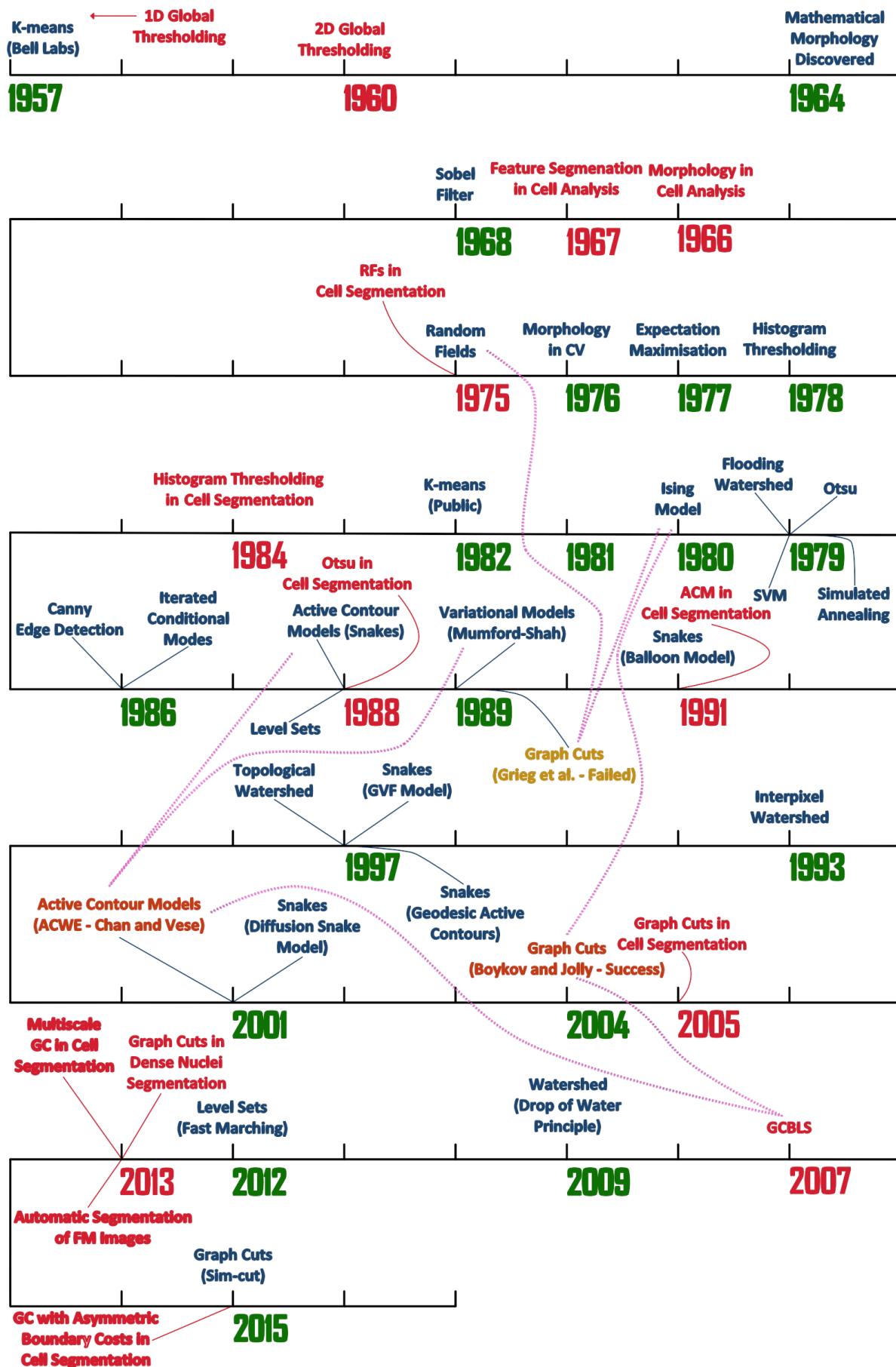


FIGURE 4.1: Progression of cell segmentation techniques. Text in red highlight momentous advancements in fluorescence image segmentation of cells.

Chapter 5

Pre-processing Scheme for Fluorescence Microscopy Images

Often enough, the poor results of a segmentation are not because the segmentation method is not tuned properly, or flawed in some way, it is because the image quality is extremely poor which renders accurate object extraction virtually impossible. The fluorescence image acquisition process is ridden with image degradation factors at almost every step. Yet still, the ever-increasing demand on image segmentation is to extract the objects of interest with greater accuracy in a shorter time. Some of these issues are poor contrast, photo-bleaching, not black enough background, non-fluorescing samples, improper excitation, etc.

Literature is rich with techniques that are able to, a high degree, negate the effect of these factors [221–223]. Given the frequency of occurrence of these problems in fluorescence images, it is surprising that there is lack of definition of a scheme prior to segmentation that prepares an image such that accurate segmentation can be achieved.

The task of defining a set of criteria an image has to meet for reliable segmentation results is not a trivial one. However, segmentation algorithms are designed to work with certain image characteristics and we can design pre-processing schemes that enhance these characteristics such that we would have a "better" image to segment. Enhancement of the original image can facilitate higher level analysis e.g. contrast enhancement. In this chapter, we aim to design a hybrid algorithm that builds on highly efficient algorithms to produce better results. A typical fluorescence image segmentation scheme is shown in Figure 5.1. In this scheme, a proxy image is generated. This image has the enhanced characteristics required for accurate segmentation. In the next step, segmentation is performed on the proxy image which yields a segmentation mask. The segmentation mask often contains speckle and other faux segmentation areas which are then removed in the clean-up stage. Thereafter, depending on the next level of image analysis, just the final segmentation mask may be all that is required, otherwise, the mask is overlayed on the original image and sent through for high level analysis.

The step that we are concerned with, in this chapter, is the Proxy Image Generation stage. Most segmentation schemes in fluorescence image segmentation would segment on intensity, therefore, a typical pre-processing scheme would emphasise the following main steps:

1. Noise reduction
2. Object data enhancement
3. Edge completion and enhancement

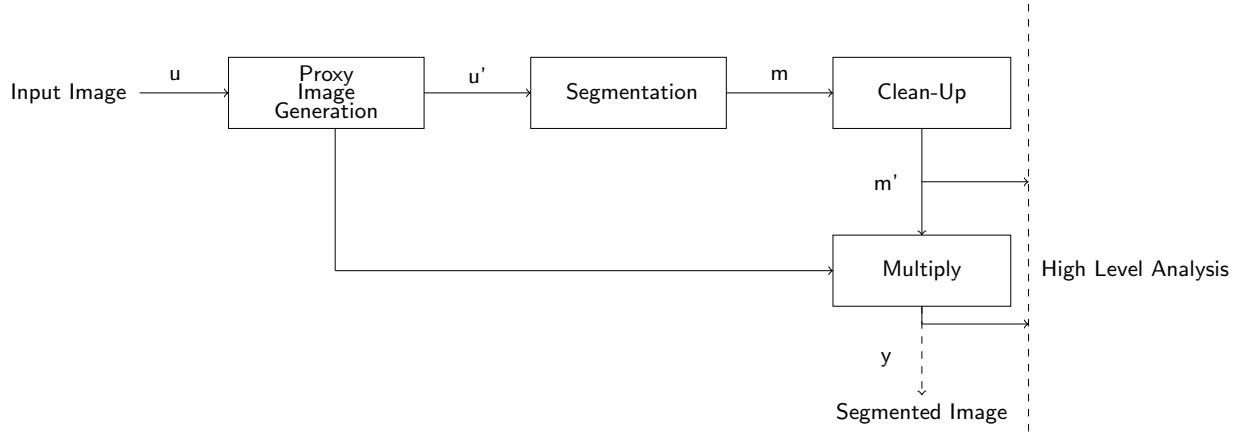


FIGURE 5.1: Segmentation Scheme.

4. Reduction of intra-region variance

We base our scheme on this framework. The step through process of the proposed scheme is illustrated in Figure 5.2. For multi-channel images, we first split the image into its channel components and process each on its own. It is at this stage that extraneous channels are discarded. Then noise removal is performed on the remaining channels. The next step involves object data enhancement by suppressing non-object data and amplifying object data. We then combine the channels into a single image and perform intra-region smoothing to reduce intensity variation.

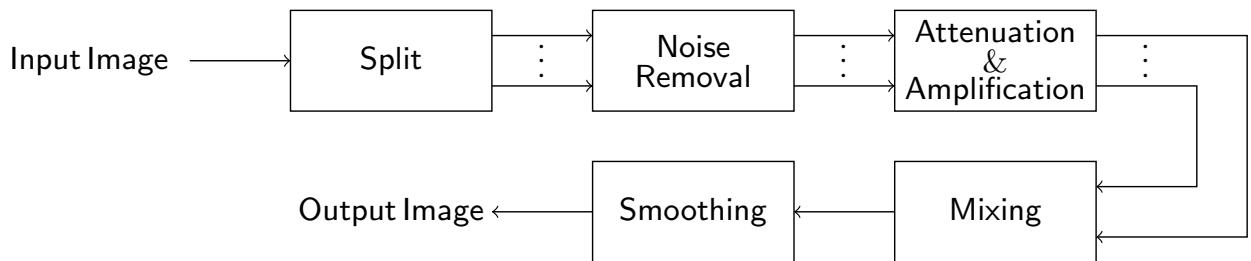


FIGURE 5.2: Proxy image generation.

5.1 Pre-processing Scheme

5.1.1 Denoising

Removing useless channels Many segmentation algorithms are designed to work on gray scale images. If we have a colour image, it is first converted to gray scale. Previously, each fluorescing sample was captured in its own grayscale image. The final colour image was composed on a computer. Through the advancements in optical engineering it is now possible for fluorescence images to be obtained in colour, however not all channels add object data to the image. Often enough, these redundant channels will just be "data-less and noisy", so eliminating this channel will often yield a higher quality image.

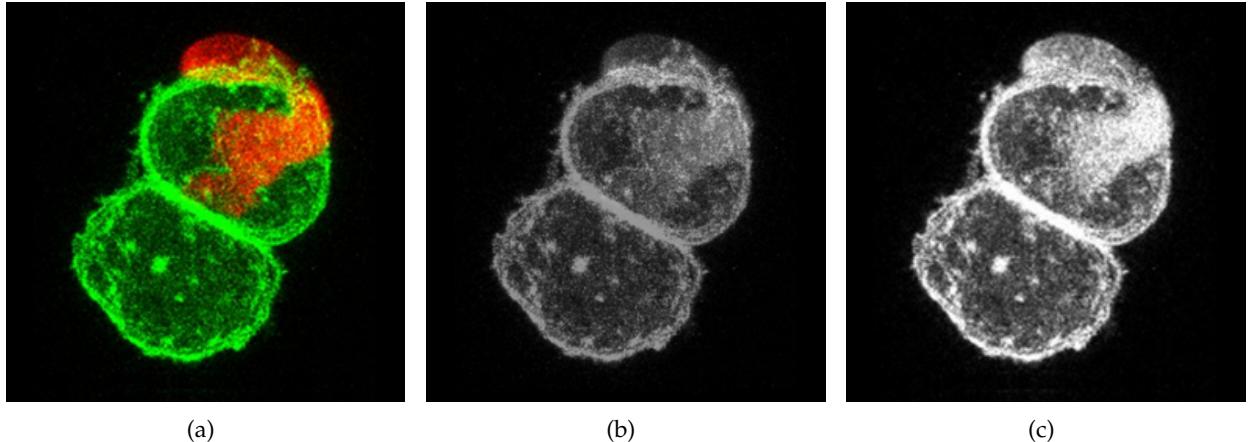


FIGURE 5.3: Comparison of grayscale conversions. **(a)** Original image. **(b)** Direct grayscale conversion. **(c)** Grayscale conversion discarding the blue channel.

For example, a direct conversion of the colour image in 5.3(a) to grayscale, shown in 5.3(b), has lesser brightness, lower contrast and more noise than the grayscale image in 5.3(c), which discarded the data-less blue channel. This is because in a direct grayscale conversion, all channels are averaged to produce the final single-channel image. Hence, the redundant blue channel is suppressing important data.

Poisson Noise Reduction In Section 2.4 we mentioned that the primary form of noise in fluorescent images is Poisson noise. It is important to remove as much noise as possible without dampening the boundary information. To this end, we have used the Total-Variation anisotropic denoising (Bregman Split). Although it was developed to remove Gaussian noise, it was shown by Rodriguez *et al.*[224] that it supersedes other state-of-the-art Poisson noise removal methods, e.g. wavelets [225], platelets[226], minimum description length [227], while maintaining signal integrity.

The Poisson distribution, which has equal mean and standard deviation i.e. $\mu = \sigma$, is defined by

$$P(n, \mu) = \frac{e^{-\mu} \mu^n}{n!} \quad (5.1)$$

Let $y = \{y_i : i = 1, \dots, N\}$ and $x = \{x_i : i = 1, \dots, N\}$ be the observed and the true image, respectively. The sample y_i is a Poisson contaminated form of x_i . We desire to recover the signal x from the observed signal y . From Bayes' Law, we get

$$P(x|y) = \frac{P(y|x)P(x)}{P(y)} \quad (5.2)$$

Therefore, we wish to find the maximum of $P(y|x)P(x)$. If all samples are affected by Poisson noise we have

$$P(y|x) = P(y, x) = \frac{e^{-x_i} x_i^{y_i}}{y_i!} \quad (5.3)$$

Thus the likelihood of observing y given the true image x is given by

$$P(y|x) = \prod_{i=1}^N \frac{e^{-x_i} x_i^{y_i}}{y_i!} \quad (5.4)$$

In anisotropic TV denoising we wish to recover the original image given the noisy image by minimising the constrained problem

$$\min_u \left\| \frac{du}{dx} \right\|_1 + \left\| \frac{du}{dy} \right\|_1 + \frac{\gamma}{2} \|u - f\|_2^2 \quad (5.5)$$

where $\gamma > 0$ is the regularisation parameter which affects the balance between noise removal and signal preservation [228], u is the true image and f is the noisy image. For computational efficiency issues, we actually solve the unconstrained problem

$$\min_{u,dx,dy} \|dx\|_1 + \|dy\|_1 + \frac{\gamma}{2} \|u - f\|_2^2 + \frac{\lambda}{2} \|dx - u_x\|_2^2 + \frac{\lambda}{2} \|dy - u_y\|_2^2 \quad (5.6)$$

This can be solved using the Bregman Split algorithm [229]. The algorithm runs iteratively until the error, $e = \frac{|u' - u|}{\|u\|}$, is less than a user-defined tolerance factor, ϵ , where u' is the image obtained after denoising the input image u .

5.1.2 Object Data Enhancement

The fundamental task of binarization of an image is to split up an image into homogenous regions of two gray levels, l_0 and l_1 . All pixels labelled l_0 share similar values with regard to the feature under consideration. This also means that there is high dissimilarity with pixels that are labelled l_1 . For grayscale images, such as the type we are concerned with, this feature is generally intensity. For segmentation regarding a specific feature, an ideal image is one where there is no overlap in the feature space between the background and the object, as illustrated in Figure 5.4. In real images (non-synthetic) this is almost never the case as there is no known feature space or it does not exist.

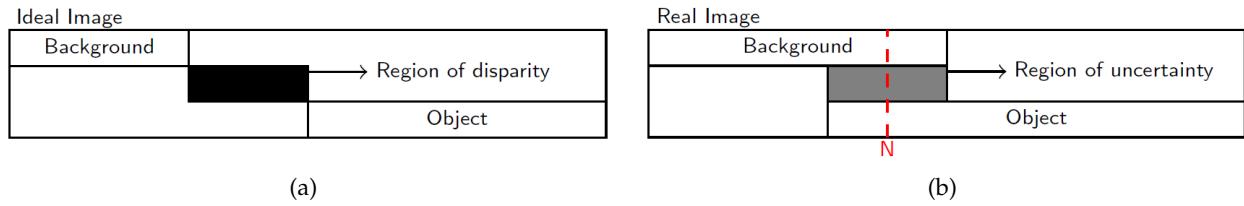


FIGURE 5.4: Feature Distribution for ideal and real images. **(a)** Ideal image. The background and the object occupy distinct non-overlapping partitions of the feature space. The non-overlapping region is called the "Region of Disparity". **(b)** Real Image. The background and the object partially overlap in the feature space. The overlapping section is called the "Region of Uncertainty".

Grayscale medical image segmentation (MIS) typically uses intensity as the dominant feature on which the application of the segmentation algorithms is biased. Hence, it is common to precede segmentation by contrast enhancement [230, 231]. The problem therein lies with the fact that the aim of contrast enhancement is to bring out the details more clearly that are otherwise obscured due to limited dynamic range, non-uniform illumination, etc. In fluorescence image this does not necessarily mean that the contrast enhanced image possesses better "segmentation qualities", in terms of intensity.

In this section, we present a novel mapping function whose aim is to shrink the "region of uncertainty" by non-linearly widening the gap between background pixels and object pixels. This function

is composed of two piece-wise sub-functions; one for data attenuation and one for data amplification. We first design the properties the mapping function should have.

Remapping Function Properties We denote the remap function as $\tilde{x}_i = R(x_i)$, where \tilde{x}_i is the new value which remaps the input pixel, x_i , using the function R . The range on which R works is $[0, L]$. For 8-bit gray scale images the highest value is usually $L = 255$. Let N be the value that contains the greatest classification uncertainty, as illustrated in Figure 5.4(b). R must have the following properties:

1. *R must be non-decreasing in the interval $[0, L]$*

This is a trivial criterion arising from the context in which our problem is defined. We specifically focus black background fluorescence images. Consequently, it is not possible for a lower intensity pixel to have a higher probability of belonging to the object compared to a pixel of a higher gray level intensity.

2. $\tilde{x}_i = R(x_i = N) = N$

This value has no bias as to whether it tends more to the background or the object. It is best left unaltered. This is marked in Figure 5.5 as "2".

3. *Attenuation: $\tilde{x}_i < x_i, \forall x_i < N$*

R must remap gray-level intensities below N , according to $0 \leq \tilde{x}_i \leq x_i$. This is marked in Figure 5.5 as "3".

4. *Amplification: $\tilde{x}_i > x_i, \forall x_i > N$*

R must remap gray-level intensities above N , according to $x_i \leq \tilde{x}_i \leq L$. This is marked in Figure 5.5 as "4".

5. *R' must be non-decreasing in the interval $[0, N]$*

Given two pixels, p_1 and p_2 , with values x_1 and x_2 respectively where $x_1 < x_2$. It is more probable for p_2 to belong to the object since it has a higher value. Also, pixels with gray levels intensities closer to 0 do not need to be attenuated as much.

6. *R' must be non-increasing in the interval $[N, L]$*

As the pixels values approach L , less amplification is needed since the pixels are already more likely to be classified as belonging to the object.

Function Design Given the criteria presented, many functions can be designed. We have decided that one of the better solutions is to make the mapping function a piece-wise quadratic Bezier curve. This is to maintain intuitive tuning of the sub-functions. The two functions are R_{att} , for the attenuation section, and R_{amp} , for the amplification section as shown in Figure 5.6.

There are three anchor points the function must pass through. These are $p_0(0, 0)$, $p_2(N, N)$ and $p_4(L, L)$. If the curve between p_0 and p_4 is a straight line, then there is no change between the output and input image since the gradient of the line is equal to 1. Additional control points are needed to bend the curve. There is a particular class of curves whose shapes are useful for our purpose. This places constraints on the position of the control points. Relative to the straight line, $y = x$, $x \in [0, L]$,

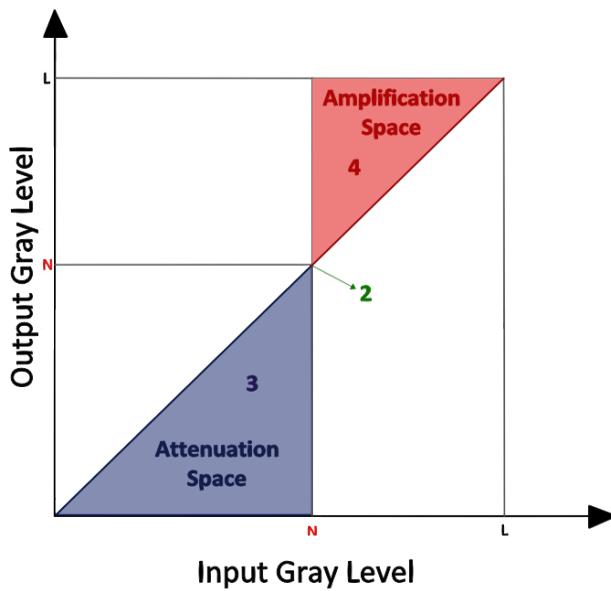


FIGURE 5.5: Remapping function space.

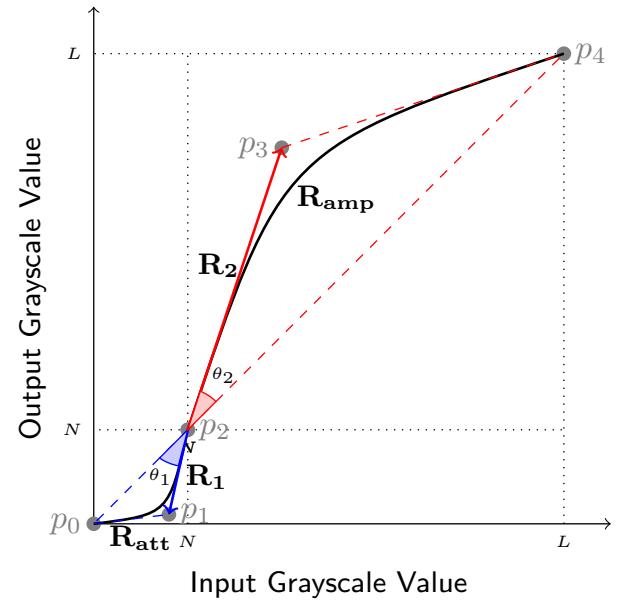


FIGURE 5.6: Plot of remapping function.

the amplification curve would be above the relaxed line, $p_{3y} > p_{3x}$, and similarly the attenuation curve would be below the line, $p_{1y} < p_{1x}$. Also, given that the function is to be a one-to-one function, the control point for the attenuation function $p_{1x} < N$; similarly, the control point for the amplification function $p_{3x} > N$.

It is more geometrically intuitive to represent the control points in polar coordinates with the centre at $p_2(N, N)$. Therefore, control point c is represented as $p_c(R_c, \theta_c)$; where $\theta_c \in [0, \frac{\pi}{4}]$ and $R_c \in \mathbb{R}^+$. The angular deviation is the measure off the straight line which is counter-clockwise for the amplification function and clockwise for the attenuation function. The deviation, θ_c , where $c \in 1, 2$, away from the straight line is further implicitly represented as a range $\kappa_c \in [0, 1]$ where $\kappa_c = 0$ implies $\theta_c = 0$ would mean no deviation, and $\kappa_c = 1$ implies $\theta_c = \frac{\pi}{4}$ would mean maximum deviation. As the function approaches the ends of its domain, less attenuation or amplification occurs, which meets criteria 5 and 6.

The position of the attenuation control point is calculated as

$$p_1 = \left(p_{2x} - R_1 \cos\left[\frac{\pi}{4}(1 + \kappa_1)\right], p_{2y} - R_1 \sin\left[\frac{\pi}{4}(1 + \kappa_1)\right] \right)$$

Similarly, the position of the amplification control point, as shown in Figure 5.7, can be calculated as

$$p_3 = \left(p_{2x} + R_2 \cos\left[\frac{\pi}{4}(1 + \kappa_2)\right], p_{2y} + R_2 \sin\left[\frac{\pi}{4}(1 + \kappa_2)\right] \right)$$

The piecewise functions that define the curve are given by

$$R_{att}(t) = \sum_{i=0}^2 p_i J_i^n \quad (5.7)$$

$$R_{amp}(t) = \sum_{i=2}^4 p_i J_i^n \quad (5.8)$$

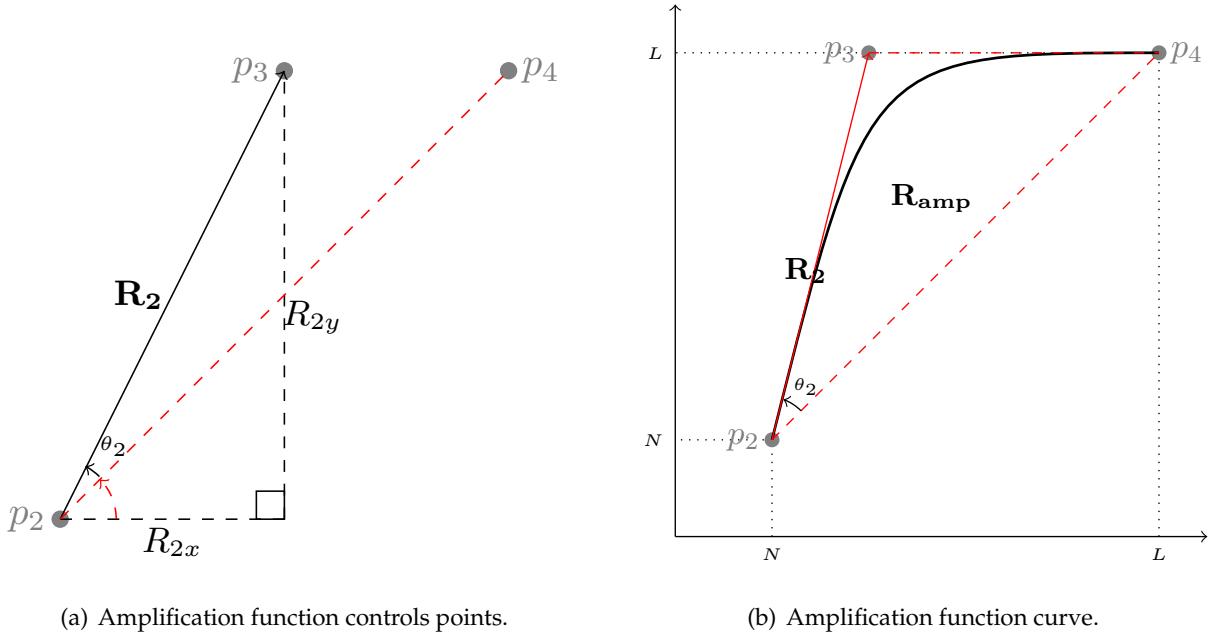


FIGURE 5.7: Amplification function control points and curve.

Where J_i^n is the Bernstein basis function, defined as

$$J_i^n = \binom{n}{i} t^i (1-t)^{n-i}, \quad t \in [0, 1] \quad (5.9)$$

For each input gray level intensity $x \in [0, L]$ we require corresponding output gray level intensity $y \in [0, L]$. The curve is parameterised with respect to t for each of its x and y parameters. For a curve defined by three control points q_0, q_1 and q_2 where $q_0 < q_1 < q_2$, the equation for the curve determined by these points is the quadratic Bezier given by

$$R = \sum_{i=0}^2 q_i J_i^n \quad (5.10)$$

For a quadratic Bezier curve, this simplifies to

$$\begin{aligned} R &= q_0(1-t)^2 + 2q_1(1-t)t + q_2t^2 \\ 0 &= (q_0 - 2q_1 + q_2)t^2 + (-2q_0 + 2q_1)t + (q_0 - R) \end{aligned}$$

Therefore for the parametric function R_x , we require the set of values of t for which $R_x \in [q_{0x}, q_{2x}]$. The value of t for a given R_x is determined by the solution to the quadratic equation.

Let $a = q_{0x} - 2q_{1x} + q_{2x}$, $b = -2q_{0x} + 2q_{1x}$, and $c = q_{0x} - R_x$.

It is seen that $b > 0$, since $q_{1x} > q_{0x}$, and $c \leq 0$, since $R_x \geq q_{0x}$.

Rewrite a as $a = (q_{0x} - q_{1x}) + (q_{2x} - q_{1x})$. For the case of a , there are two cases.

1. For $q_{1x} \in [q_{0x}, \frac{q_{2x}-q_{0x}}{2}]$, $a > 0$.

In this case $-4ac > 0$ hence $\sqrt{b^2 - 4ac} > b$.

\therefore the only value of t which is positive and must be the solution is given by

$$t = \frac{(q_{1x} - q_{0x}) + \sqrt{R_x(q_{0x} - 2q_{1x} + q_{2x}) + (q_{1x}^2 - q_{0x}q_{2x})}}{q_{0x} - 2q_{1x} + q_{2x}} \quad (5.11)$$

2. For $q_{1x} > \frac{q_{2x} - q_{0x}}{2}$, $a < 0$.

It is known that a real solution exists. This means that

$$\begin{aligned} b^2 - 4ac &\geq 0 \\ b^2 &\geq 4ac \\ \implies \sqrt{b^2 - 4ac} &< b \end{aligned}$$

\therefore the only value of t which is positive and must be the solution is given by

$$t = \frac{(q_{1x} - q_{0x}) + \sqrt{R_x(q_{0x} - 2q_{1x} + q_{2x}) + (q_{1x}^2 - q_{0x}q_{2x})}}{q_{0x} - 2q_{1x} + q_{2x}} \quad (5.12)$$

In both cases the solution to t can be calculated using the same formula. It is possible for the remapping function to exceed the range, in this case any value that maps to a value higher than the maximum value will be assigned the maximum value i.e. $\tilde{x}_i = \min(R(x_i), L)$, $x_i \in (N, L]$; similarly any value that maps to a value lower than the minimum value will be assigned the minimum value i.e. $\tilde{x}_i = \max(R(x_i), 0)$, $x_i \in [0, N)$.

Function Properties and Constraints This function presents several properties within the constraints defined as follows:

1. The curves obey the convex hull property. The sub-functions will always be contained within the control polygon determined by the control points [232, 233].
2. No attenuation when $\kappa_1 = 0$ and no amplification when $\kappa_2 = 0$.
3. R is continuous on $[0, L]$.
4. R is weakly monotonically increasing on $[0, L]$.
5. R'_{att} is monotonically increasing on $[0, N]$.
6. R'_{amp} is monotonically decreasing on $[N, L]$.
7. The ends of the curve are coincident with the first and last control points of the control polygon.
8. The direction of the tangent vectors from the end points of the curve are the same as the direction of the vector anchored at the control point and along the line that joins the end point and the centre control point of the control polygon [232, 233].

5.1.3 Channel Mixing

To reconstitute an image from the updated channels we perform channel mixing. In an equi-weighted mixing system, each channel contributes equally to the final image. This simplistic method of channel mixing does not always produce the best image. A consequence of equi-weighted mixing is that some channels might become very suppressed and may be disregarded by the segmentation algorithm. Hence, it is necessary to assign weights which are channel dependant. These weightings must sum to one, $\sum_{i \in C} w_i = 1$, where C is the set of channels to be mixed-down. The mixed-down image is then calculated as $y = \sum_{i \in C} w_i C_i$, where C_i is channel i . Channels with very low gray level values are given a greater weight.

5.1.4 Intra-Region Smoothing and Edge Completion and Enhancement

One of the criteria which is used in identifying an image is that objects tend to have little intra-region variance. It is common for fluorescence images to be plagued with various degrees of lighting and contrast even within objects. This is primarily due to improper excitation, non-fluorescence of particles and ubiquitous measurement errors during acquisition. We used a coherence enhancing diffusion filter with optimised rotational invariance (CED-ORI) presented in [234–236], which very successfully reduces intra-region variance and joins closely-disconnected edges.

The diffusion works by evolving the image, u , over a time using n discrete time steps, t , called the diffusion time. The evolution equation is defined as:

$$\frac{\partial u}{\partial t} = \nabla \cdot (D \nabla u) \quad (5.13)$$

where $D = \begin{pmatrix} a & b \\ b & c \end{pmatrix}$ is the diffusion tensor which can be adapted to the local image structure measure known as the structure tensor. The structure tensor is given by:

$$J_\rho(\nabla u_\sigma) = G_\rho * (\nabla u_\sigma \nabla u_\sigma^T) \quad (5.14)$$

Where G_ρ is the Gaussian kernel with standard deviation ρ , and $u_\sigma := G_\sigma * u$ where G_σ is the Gaussian kernel with standard deviation σ . The eigenvalues of $J_\rho = \begin{pmatrix} J_{11} & J_{12} \\ J_{12} & J_{22} \end{pmatrix}$ are

$$\mu_1 = \frac{1}{2} \left(J_{11} + J_{22} + \sqrt{(J_{11} - J_{22})^2 + 4J_{12}^2} \right) \quad (5.15)$$

$$\mu_2 = \frac{1}{2} \left(J_{11} + J_{22} - \sqrt{(J_{11} - J_{22})^2 + 4J_{12}^2} \right) \quad (5.16)$$

Where the normalised first eigenvector satisfies

$$\begin{pmatrix} \cos\alpha \\ \sin\alpha \end{pmatrix} \parallel \begin{pmatrix} 2J_{12} \\ J_{22} - J_{11} + \sqrt{(J_{11} - J_{22})^2 + 4J_{12}^2} \end{pmatrix} \quad (5.17)$$

The diffusion tensor's, D , eigenvectors are obtained from the structure tensor eigenvectors using:

$$\lambda_1 = c_1 \quad (5.18)$$

$$\lambda_2 = \begin{cases} c_1 & \text{if } \mu_1 = \mu_2 \\ c_1 + (1 - c_1)e^{\frac{c_2}{(\mu_1 - \mu_2)^2}} & \text{otherwise} \end{cases} \quad (5.19)$$

where $c_1 \in (0, 1)$, $c_2 > 0$. The elements of D are then calculated as:

$$a = \lambda_1 \cos^2 \alpha + \lambda_2 \sin^2 \alpha \quad (5.20)$$

$$b = (\lambda_1 - \lambda_2) \sin \alpha \cos \alpha \quad (5.21)$$

$$c = \lambda_1 \sin^2 \alpha + \lambda_2 \cos^2 \alpha \quad (5.22)$$

Further details on the coherence enhancing diffusion filter with optimised rotational invariance is found in [235]. In Masaka *et al.* [237] and Kroon *et al.* [238] this filter was used primarily for noise removal while preserving edge detail, here we use it for its edge completion and smoothing properties.

5.2 Experimental Results

In this section we present the results of the proposed pre-processing scheme compared against other very commonly used fluorescence image enhancement methods. We have used the graph cut version of the Chan-Vese segmentation model, and kept the parameters the same over the same images after different pre-processing methods. The parameter settings used in each scheme is given under the image.

We have compiled a label for label comparison on the segmentation mask and the ground truth in Table 5.1 along with the efficiency measures *precision*, *recall*, *accuracy* and *Matthews Correlation Coefficient (MCC)*. The overview results are shown in Table 5.2. For each image, we have highlighted the method which performs the best in blue, and the worst in red.

We differentiate between methods on the same image as follows:

[imageno]-[method],

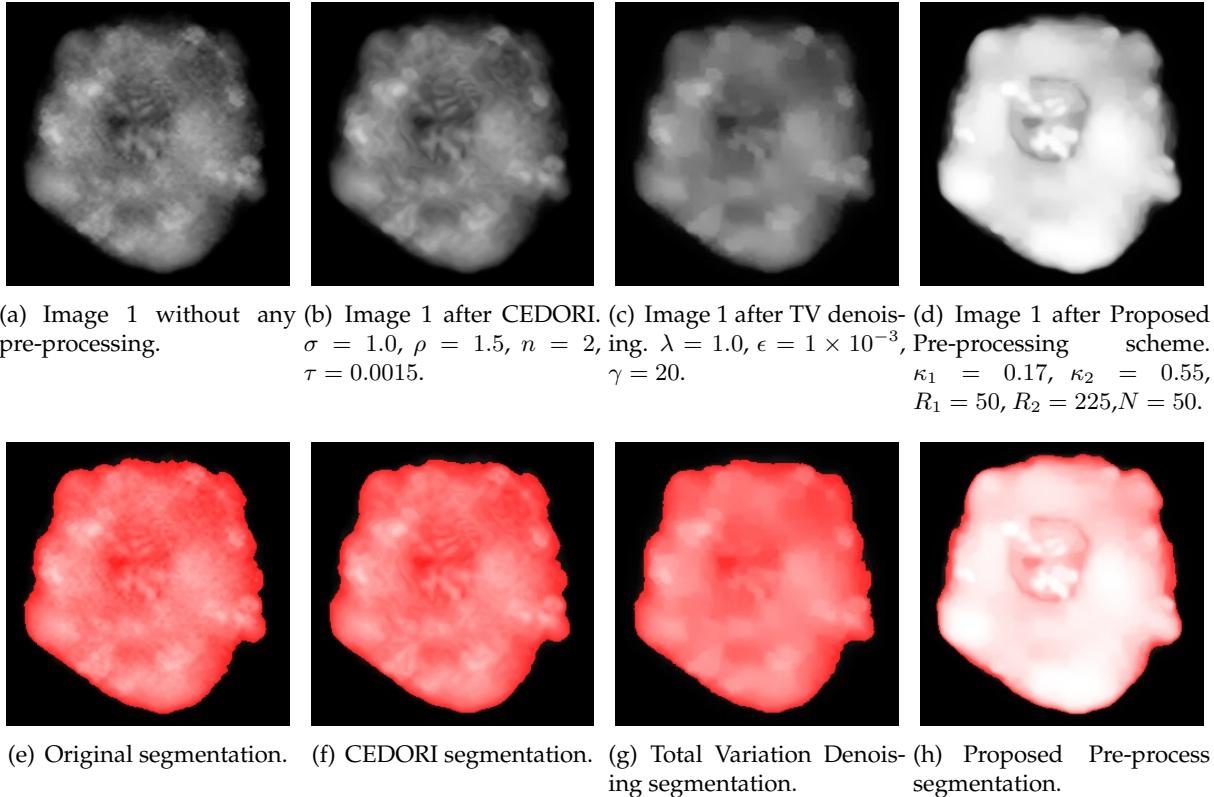
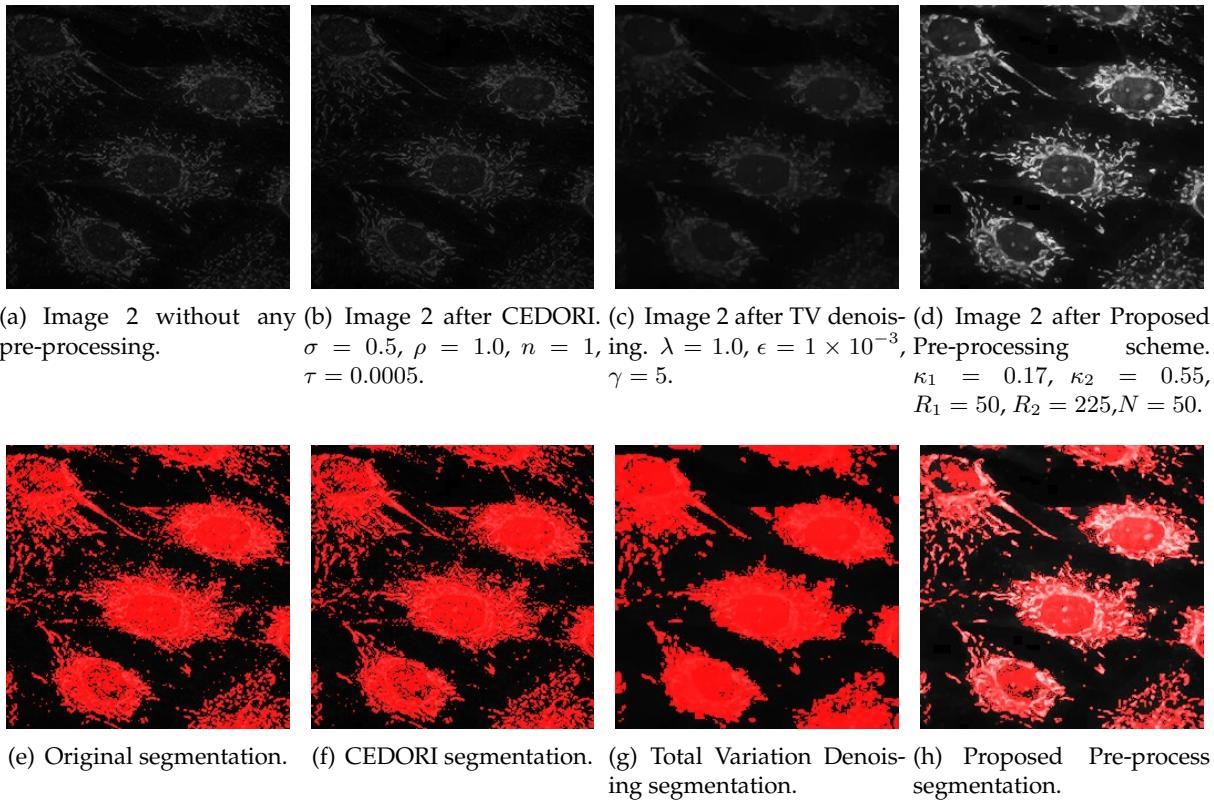
where *imageno* goes from 1 to 25 and *method* is defined as follows:

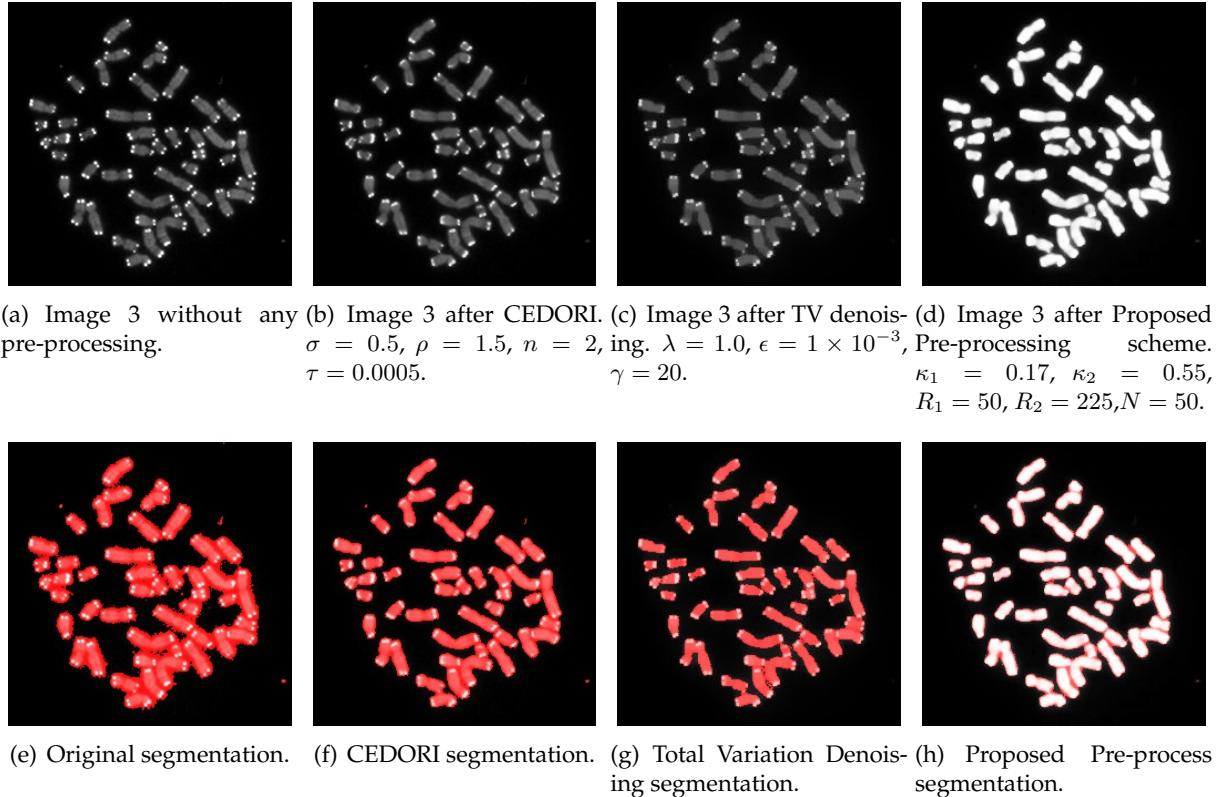
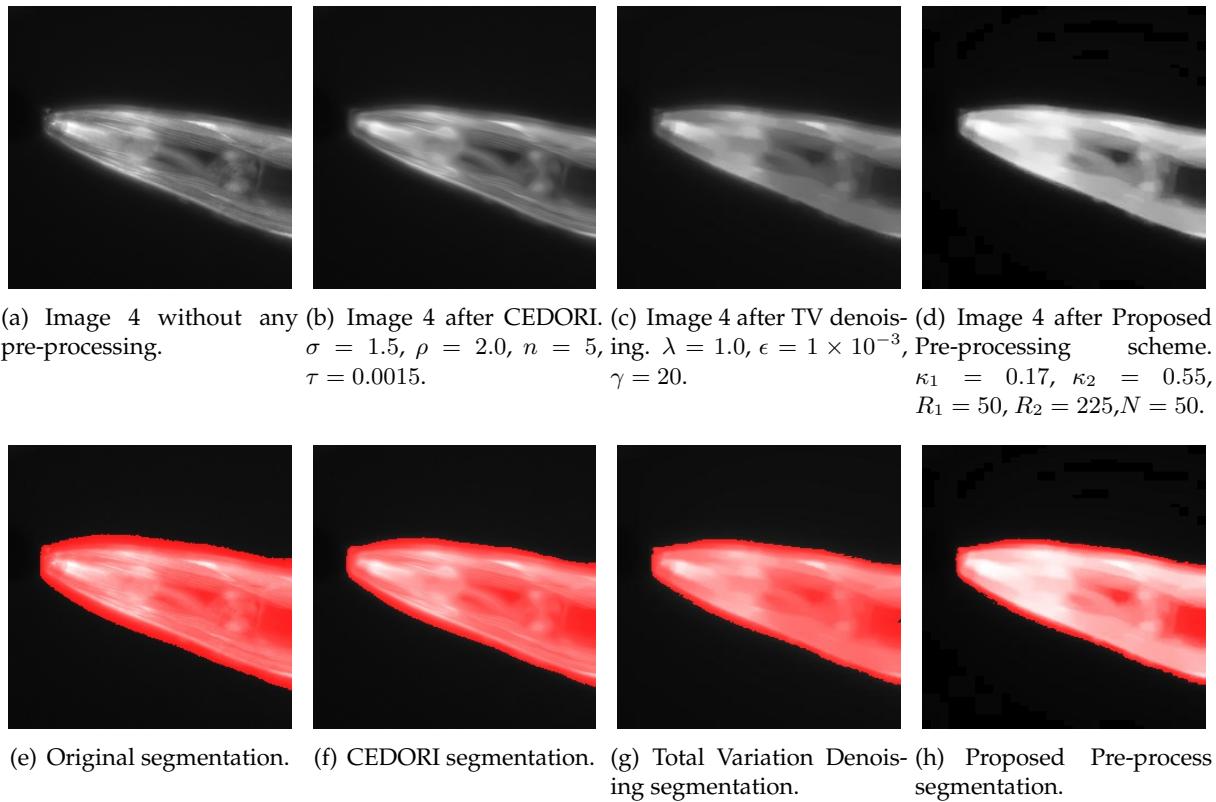
o - Original image without any pre-processing.

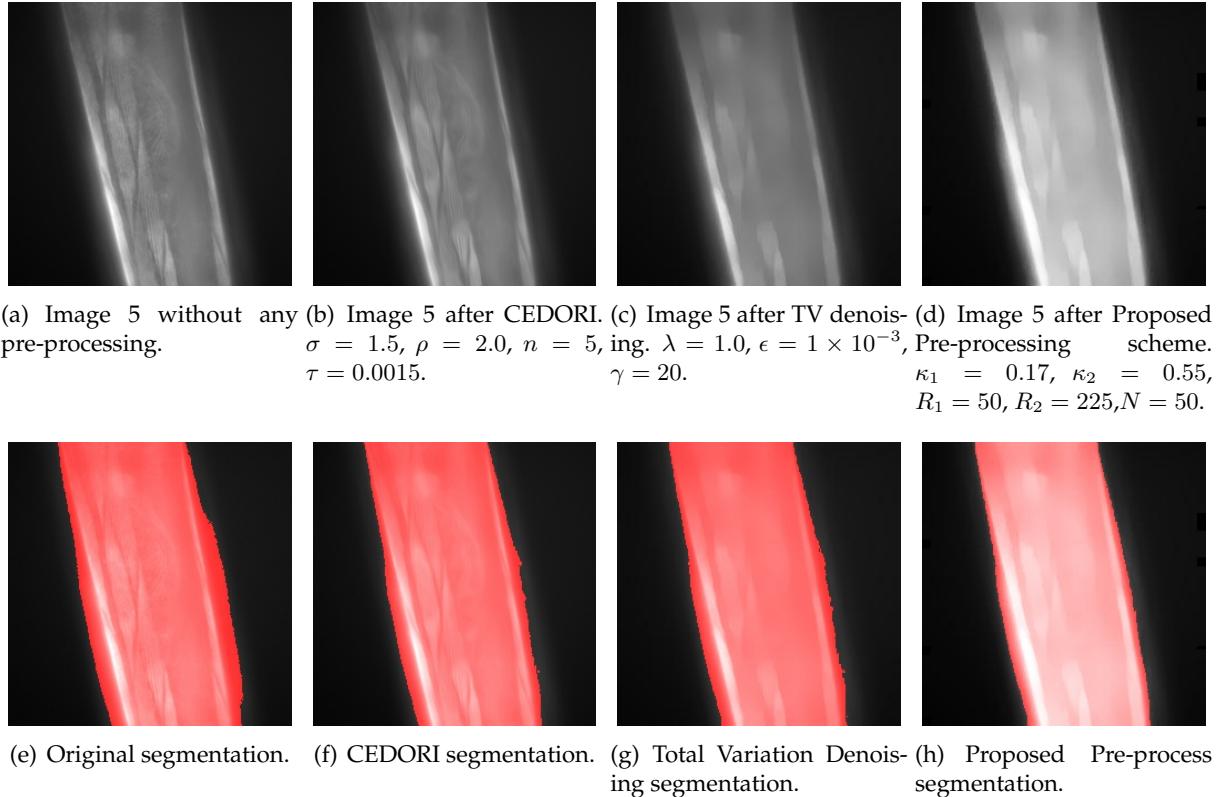
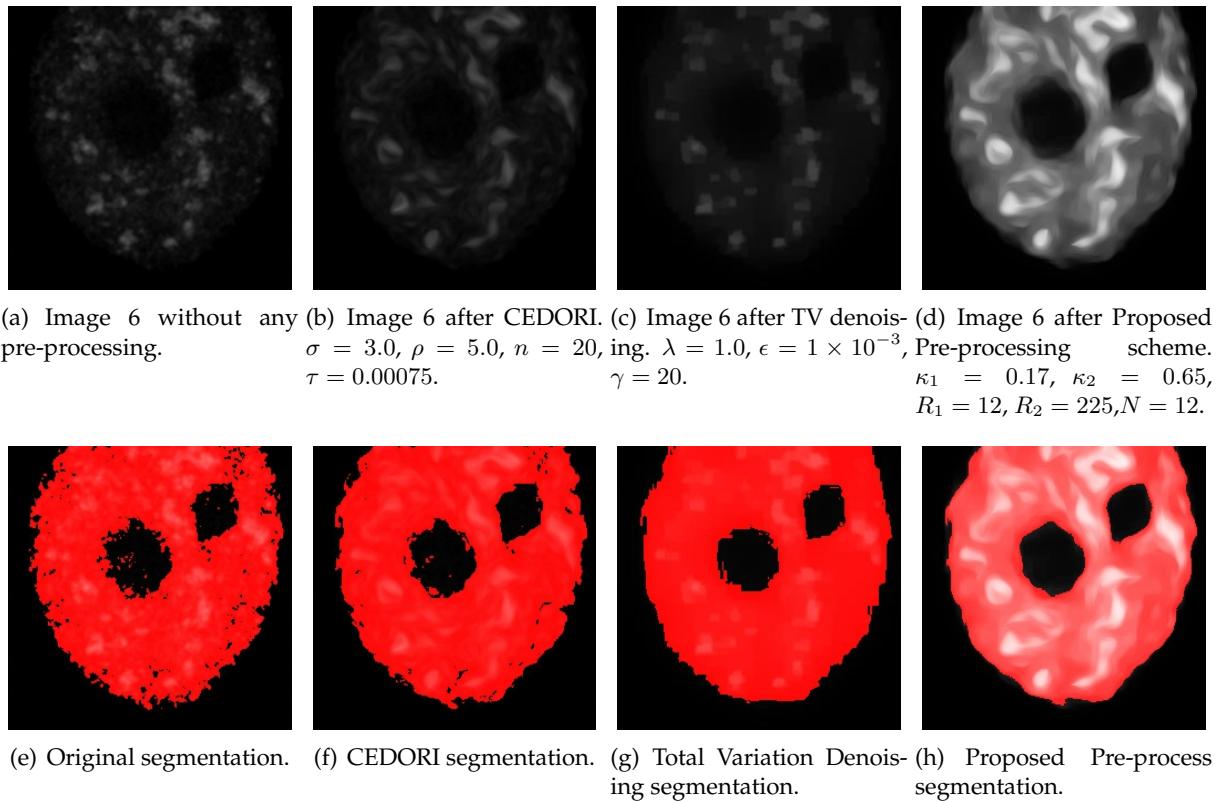
c - Image after Coherence Enhancing Diffusion with Optimised Rotataional Invariance (CEDORI).

t - Image after Total Variation denoising.

p - Image after Proposed pre-processing scheme.

FIGURE 5.8: Image 1 from test set Appendix B pre-processing segmentation results. $\mu = 1, \lambda_0 = 240, \lambda_1 = 8$.FIGURE 5.9: Image 2 from test set Appendix B pre-processing segmentation results. $\mu = 1, \lambda_0 = 5, \lambda_1 = 1$.

FIGURE 5.10: Image 3 from test set Appendix B pre-processing segmentation results. $\mu = 1, \lambda_0 = 240, \lambda_1 = 8$.FIGURE 5.11: Image 4 from test set Appendix B pre-processing segmentation results. $\mu = 1, \lambda_0 = 300, \lambda_1 = 5$.

FIGURE 5.12: Image 5 from test set Appendix B pre-processing segmentation results. $\mu = 1, \lambda_0 = 220, \lambda_1 = 11$.FIGURE 5.13: Image 6 from test set Appendix B pre-processing segmentation results. $\mu = 1, \lambda_0 = 1600, \lambda_1 = 80$.

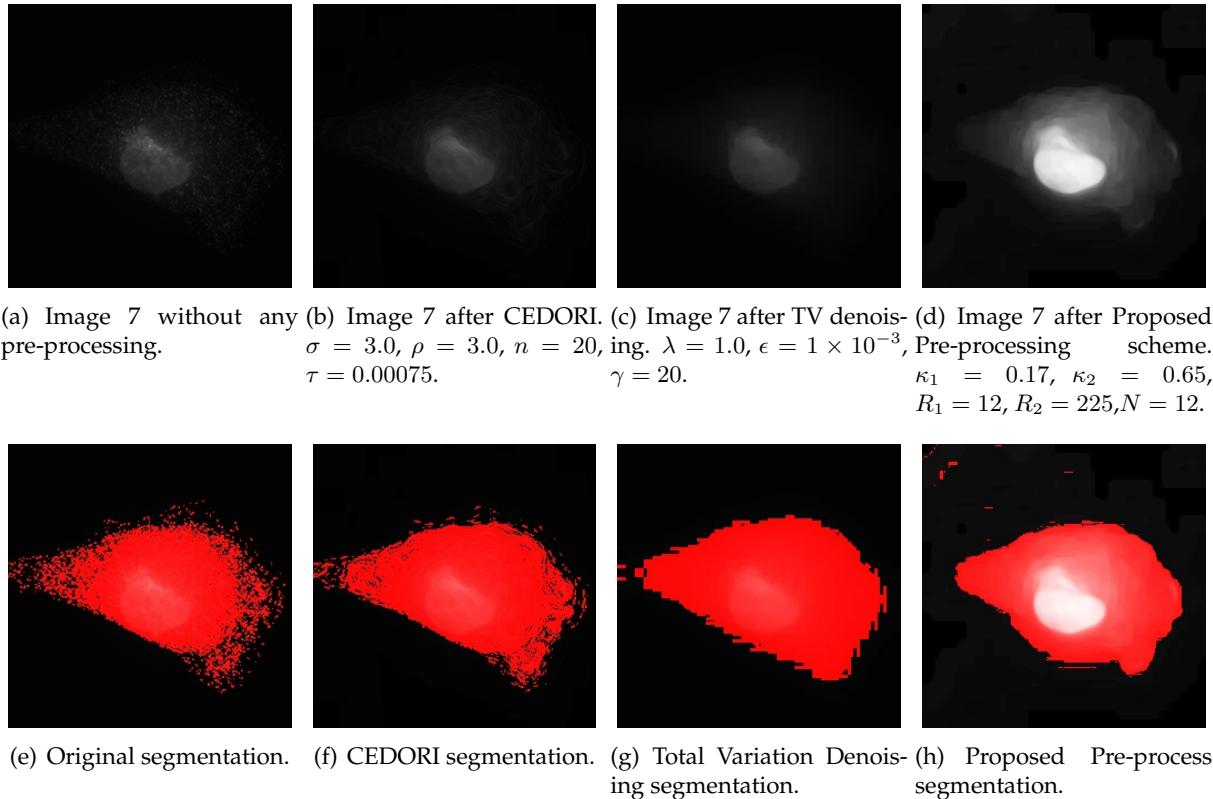


FIGURE 5.14: Image 7 from test set Appendix B pre-processing segmentation results. $\mu = 1, \lambda_0 = 10, \lambda_1 = 1$.

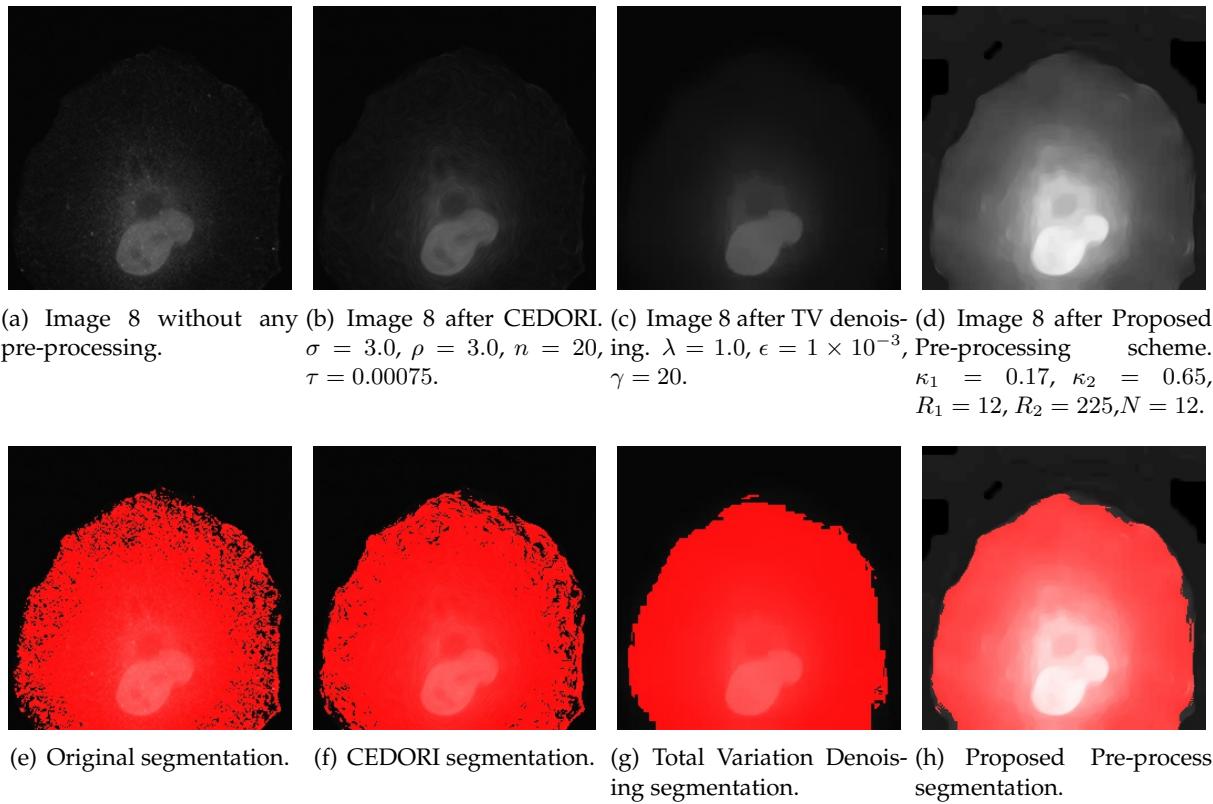


FIGURE 5.15: Image 8 from test set Appendix B pre-processing segmentation results. $\mu = 1, \lambda_0 = 10, \lambda_1 = 1$.

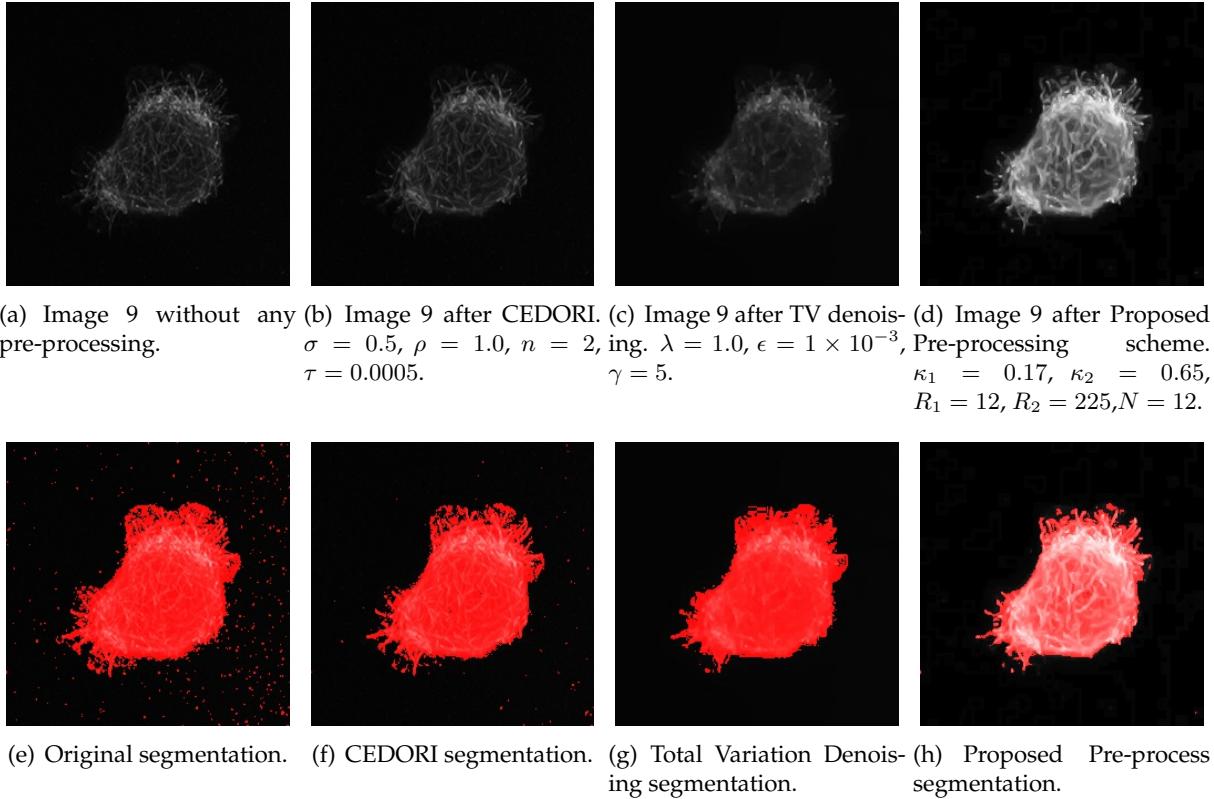


FIGURE 5.16: Image 9 from test set Appendix B pre-processing segmentation results. $\mu = 1, \lambda_0 = 2700, \lambda_1 = 90$.

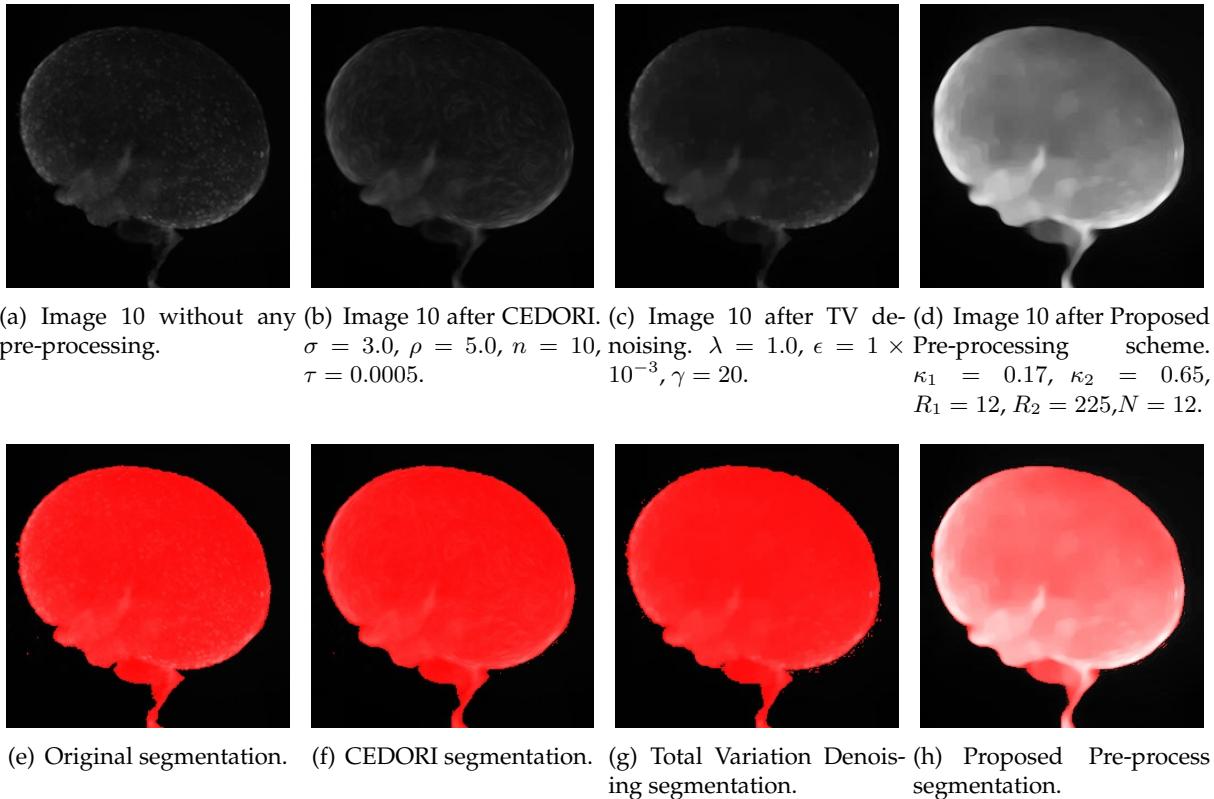
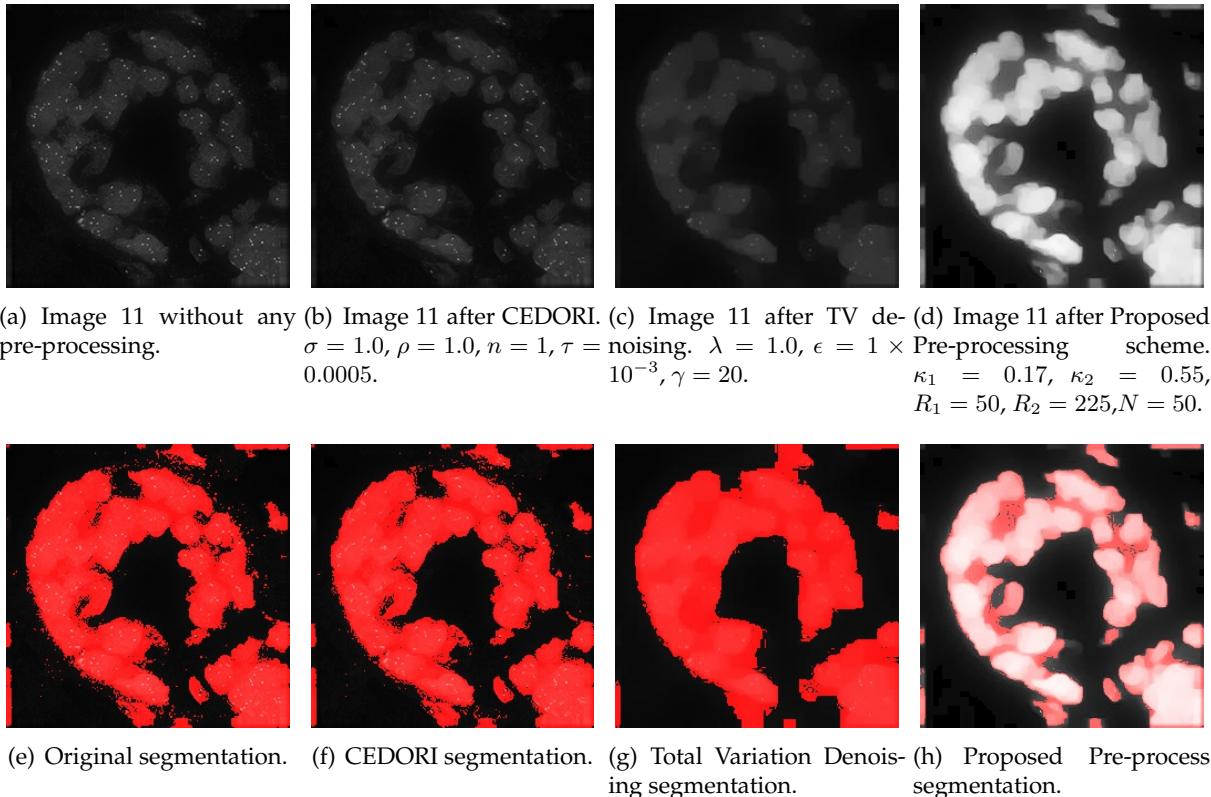
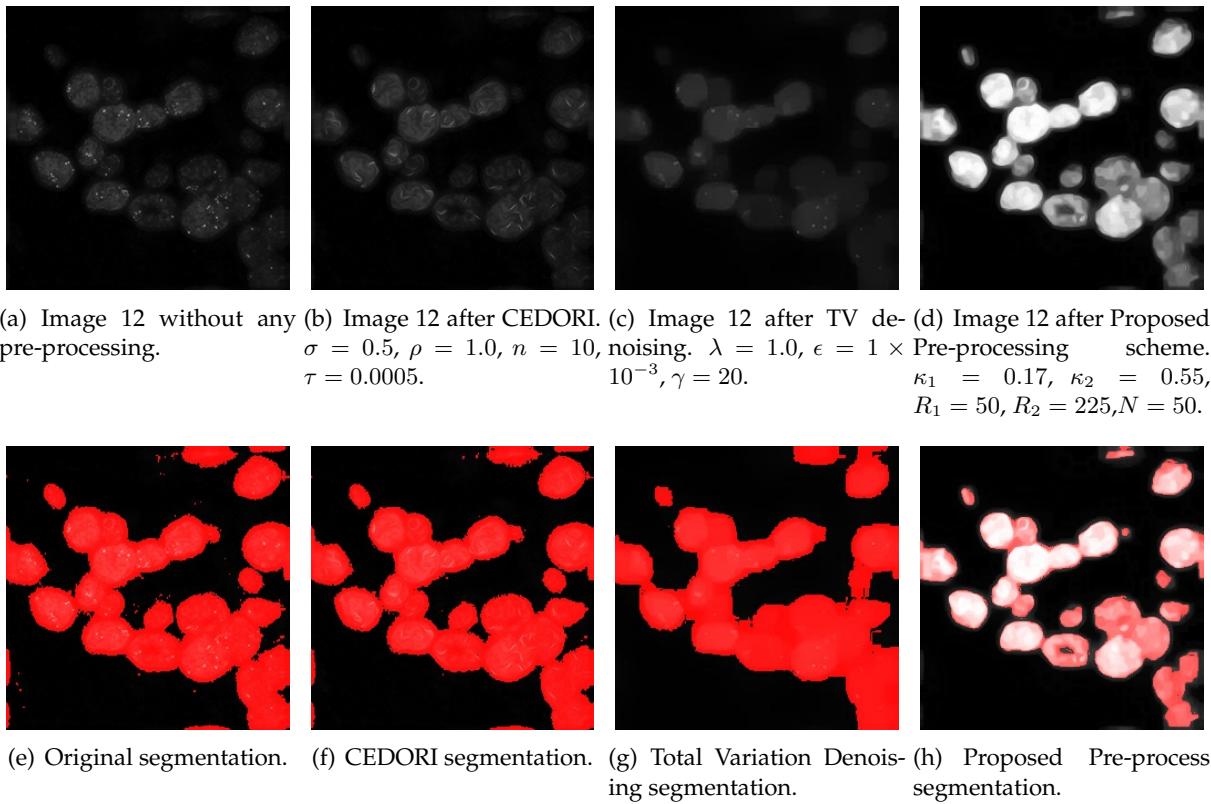
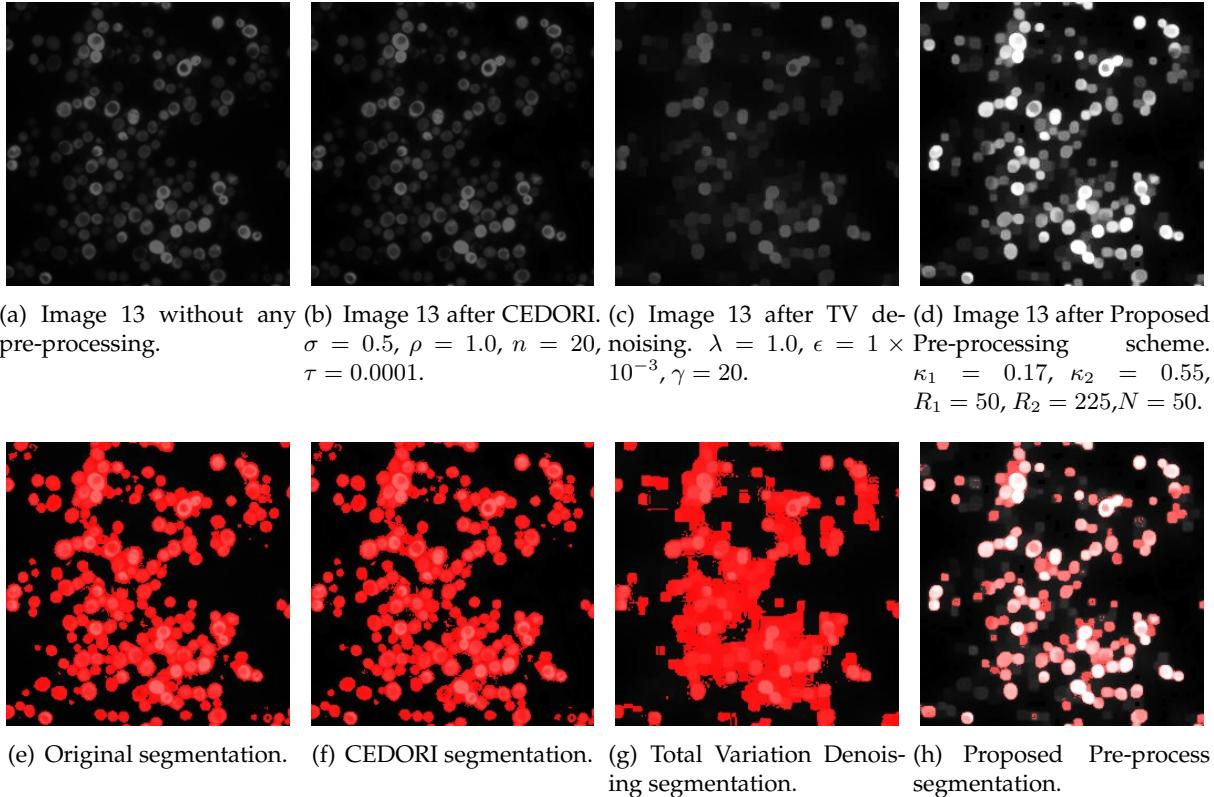
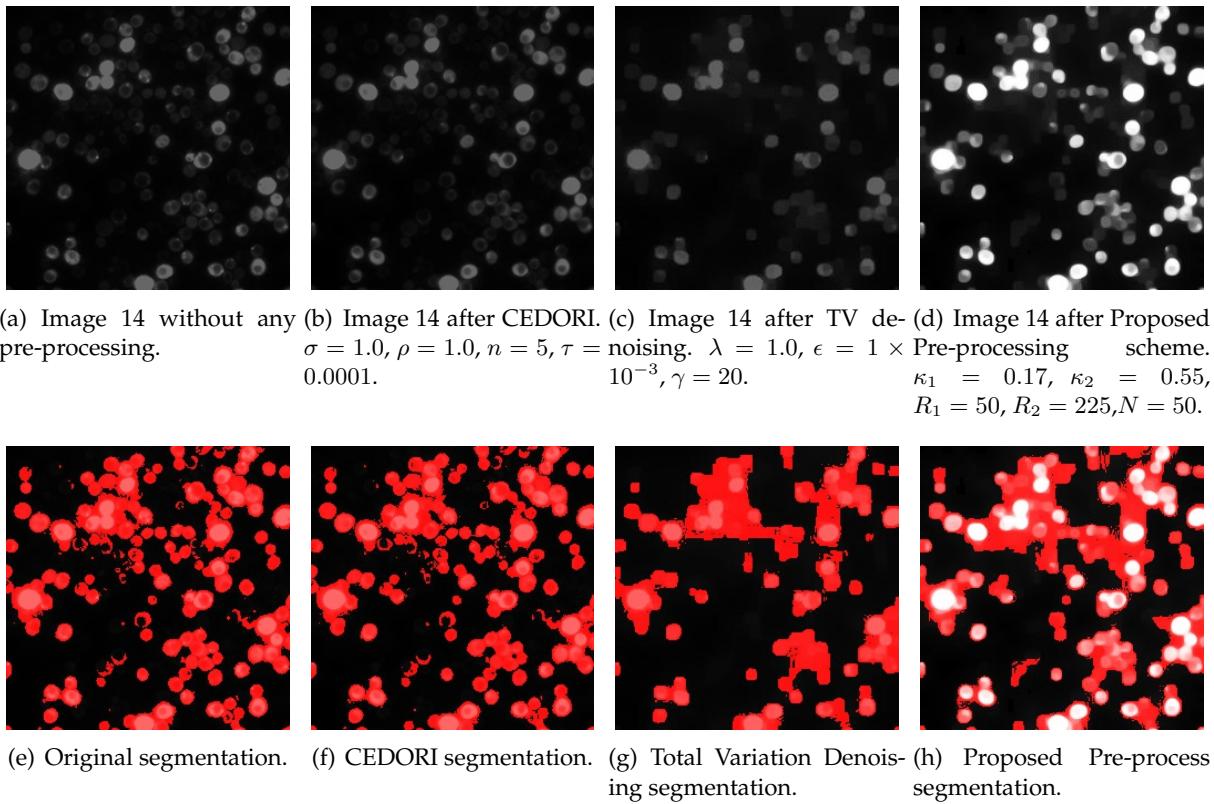
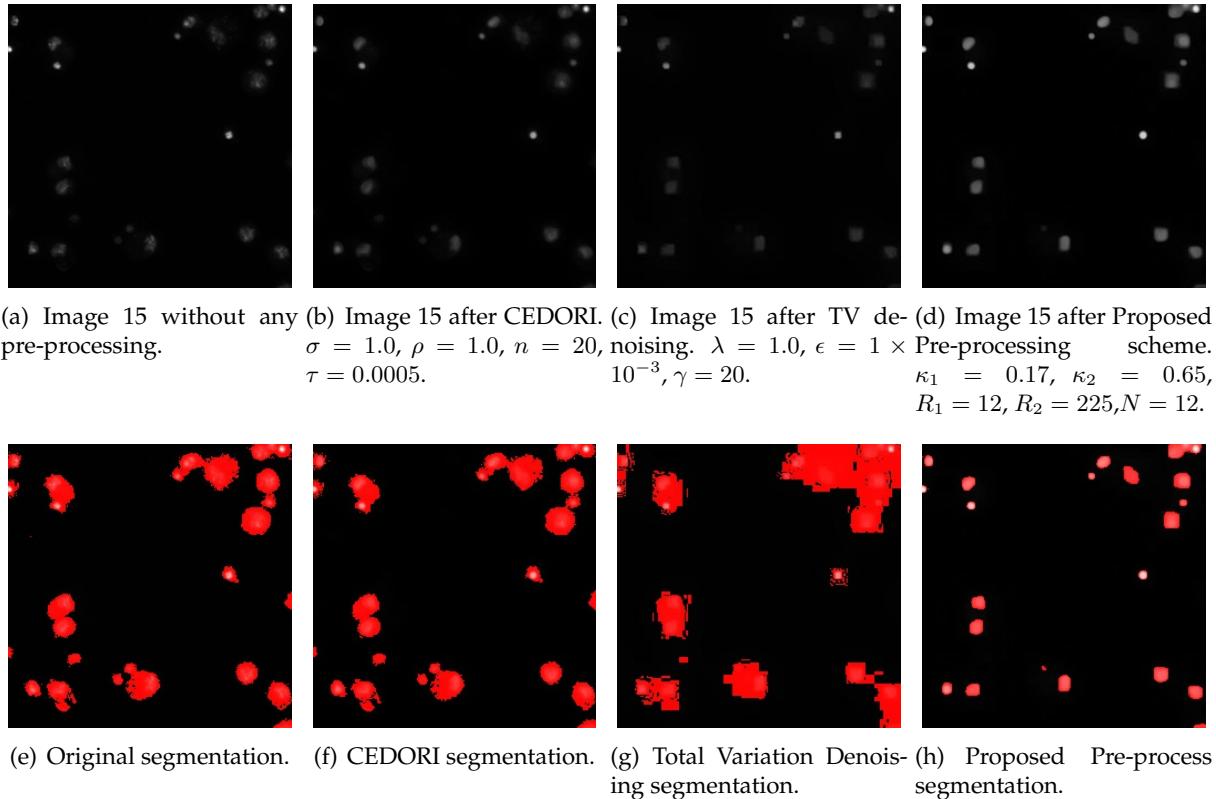
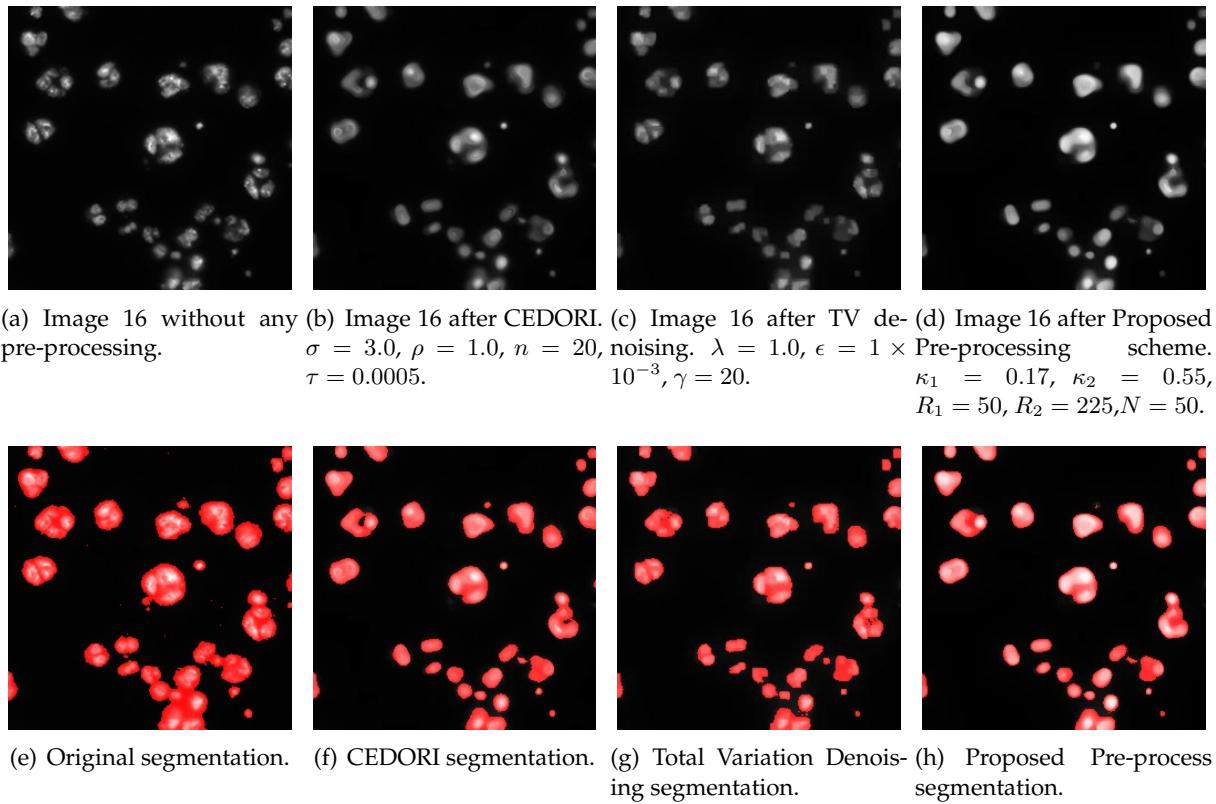


FIGURE 5.17: Image 10 from test set Appendix B pre-processing segmentation results. $\mu = 1, \lambda_0 = 4050, \lambda_1 = 135$.

FIGURE 5.18: Image 11 from test set Appendix B pre-processing segmentation results. $\mu = 1, \lambda_0 = 10, \lambda_1 = 1$.FIGURE 5.19: Image 12 from test set Appendix B pre-processing segmentation results. $\mu = 1, \lambda_0 = 10, \lambda_1 = 1$.

FIGURE 5.20: Image 13 from test set Appendix B pre-processing segmentation results. $\mu = 1, \lambda_0 = 10, \lambda_1 = 1$.FIGURE 5.21: Image 14 from test set Appendix B pre-processing segmentation results. $\mu = 1, \lambda_0 = 3, \lambda_1 = 1$.

FIGURE 5.22: Image 15 from test set Appendix B pre-processing segmentation results. $\mu = 1, \lambda_0 = 10, \lambda_1 = 1$.FIGURE 5.23: Image 16 from test set Appendix B pre-processing segmentation results. $\mu = 1, \lambda_0 = 10, \lambda_1 = 1$.

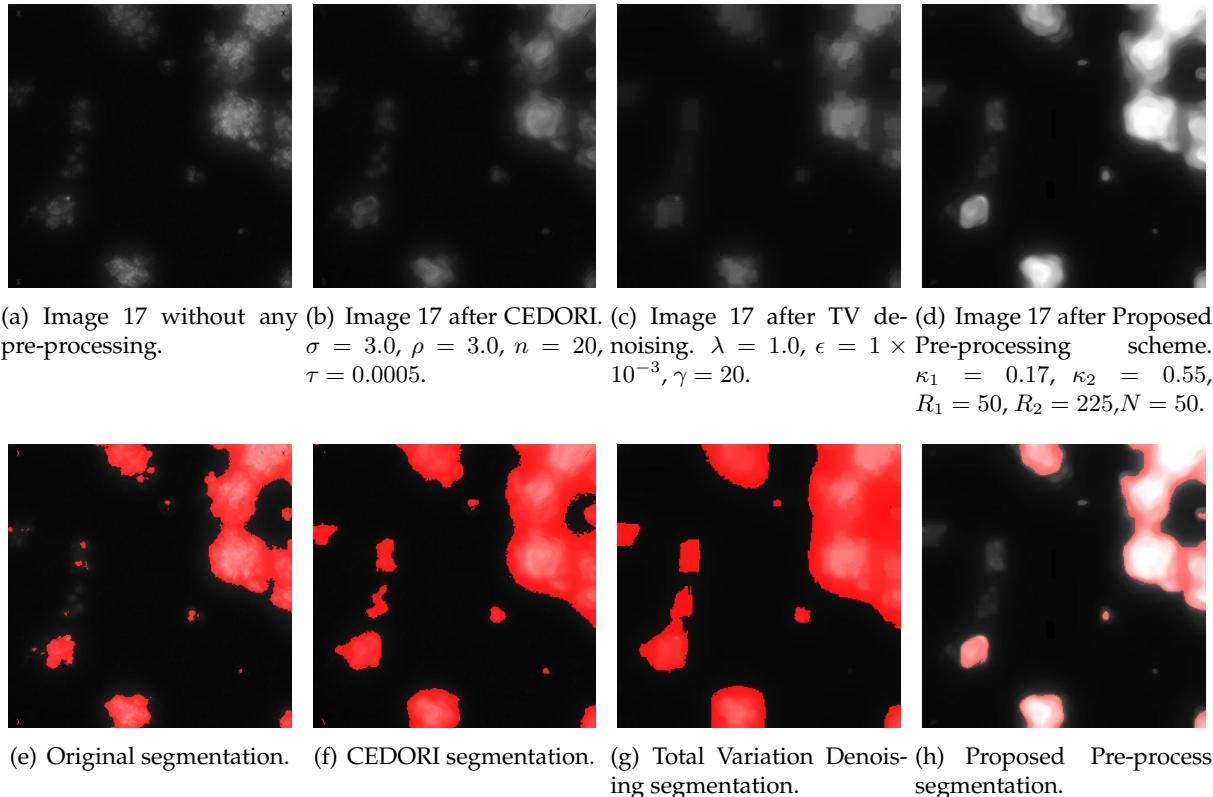


FIGURE 5.24: Image 17 from test set Appendix B pre-processing segmentation results. $\mu = 1, \lambda_0 = 5, \lambda_1 = 1$.

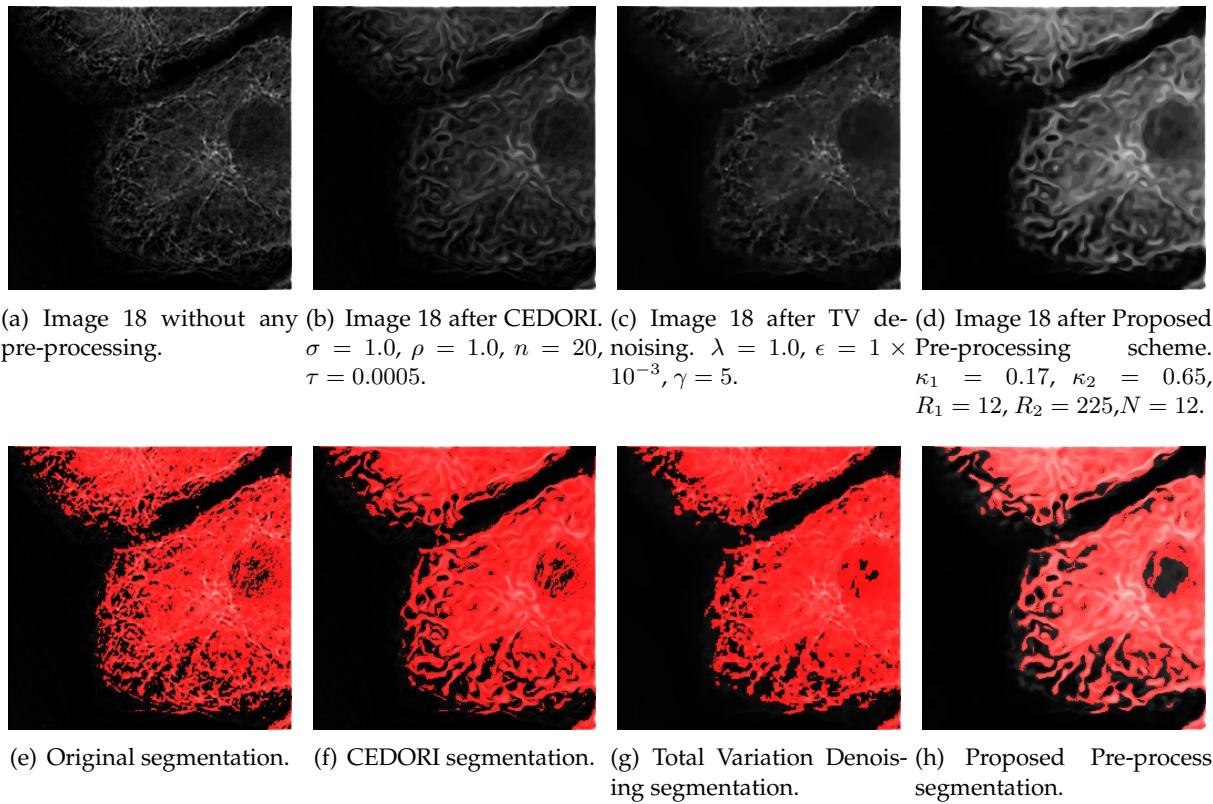


FIGURE 5.25: Image 18 from test set Appendix B pre-processing segmentation results. $\mu = 1, \lambda_0 = 5, \lambda_1 = 1$.

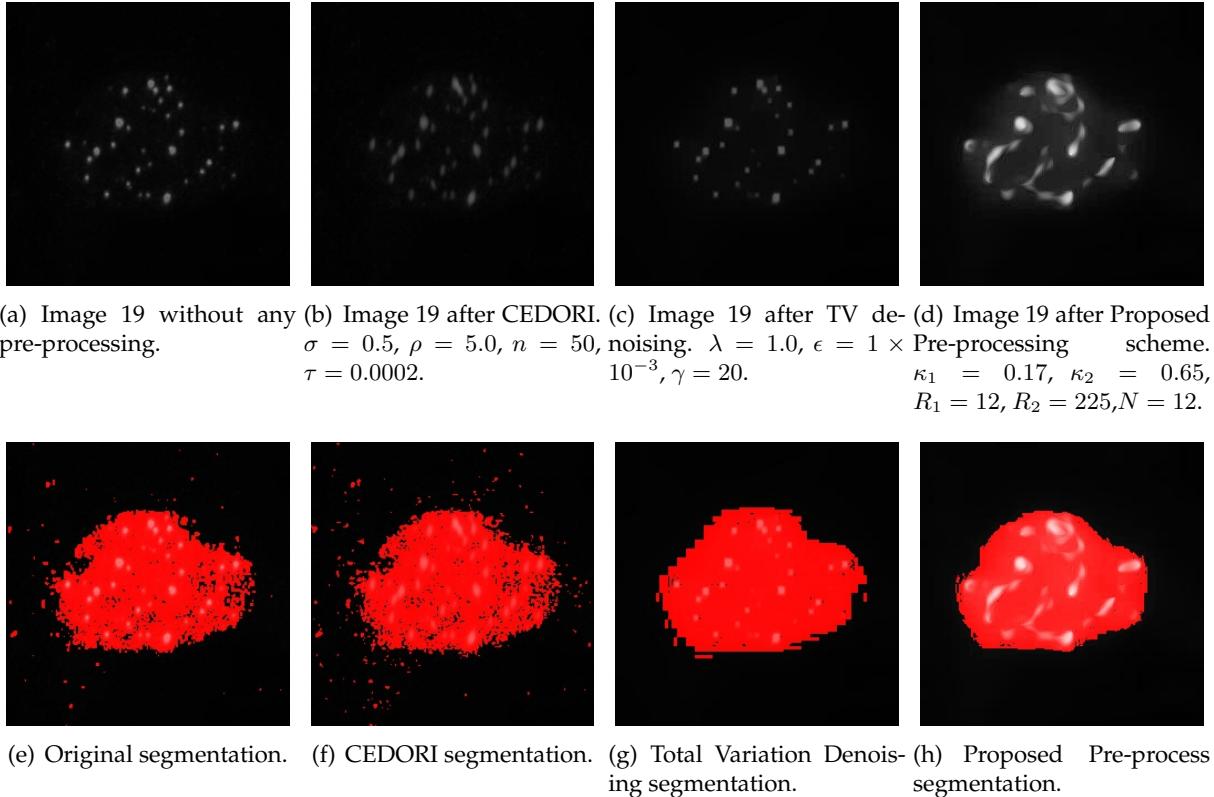


FIGURE 5.26: Image 19 from test set Appendix B pre-processing segmentation results. $\mu = 1, \lambda_0 = 6, \lambda_1 = 2$.

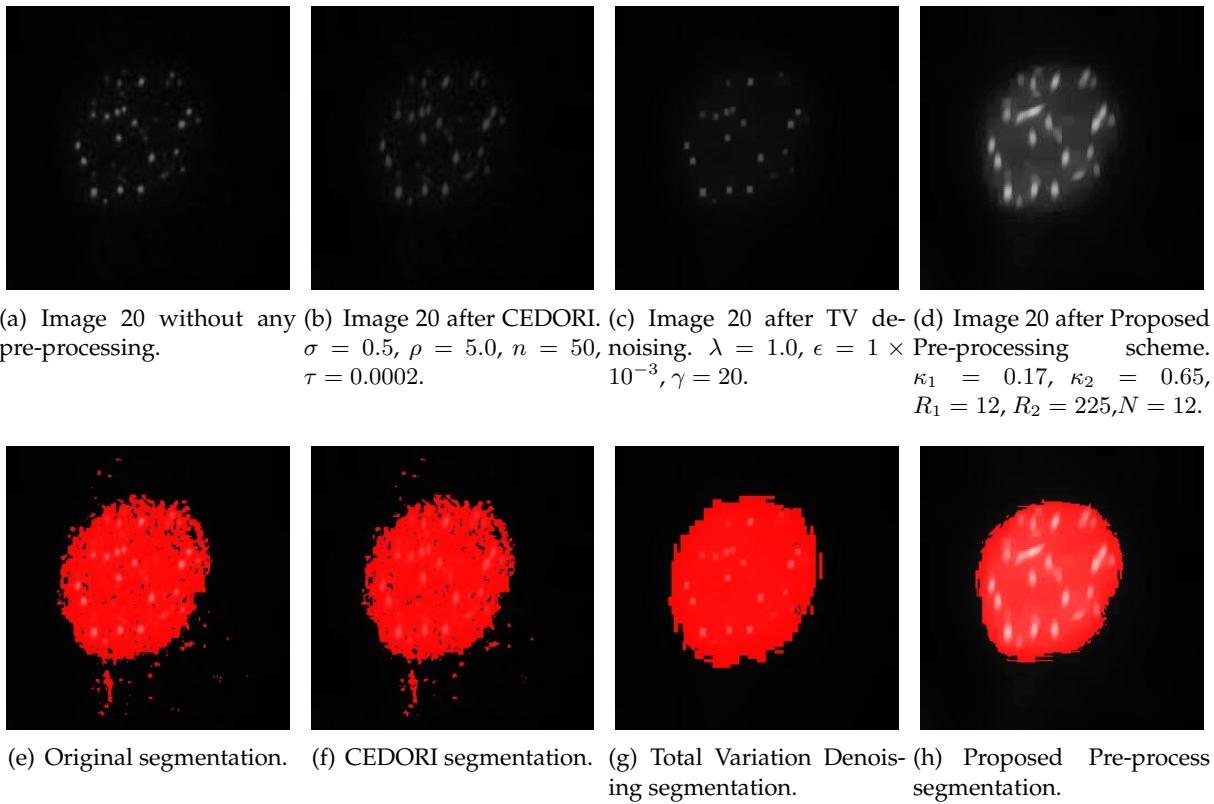


FIGURE 5.27: Image 20 from test set Appendix B pre-processing segmentation results. $\mu = 1, \lambda_0 = 6, \lambda_1 = 2$.

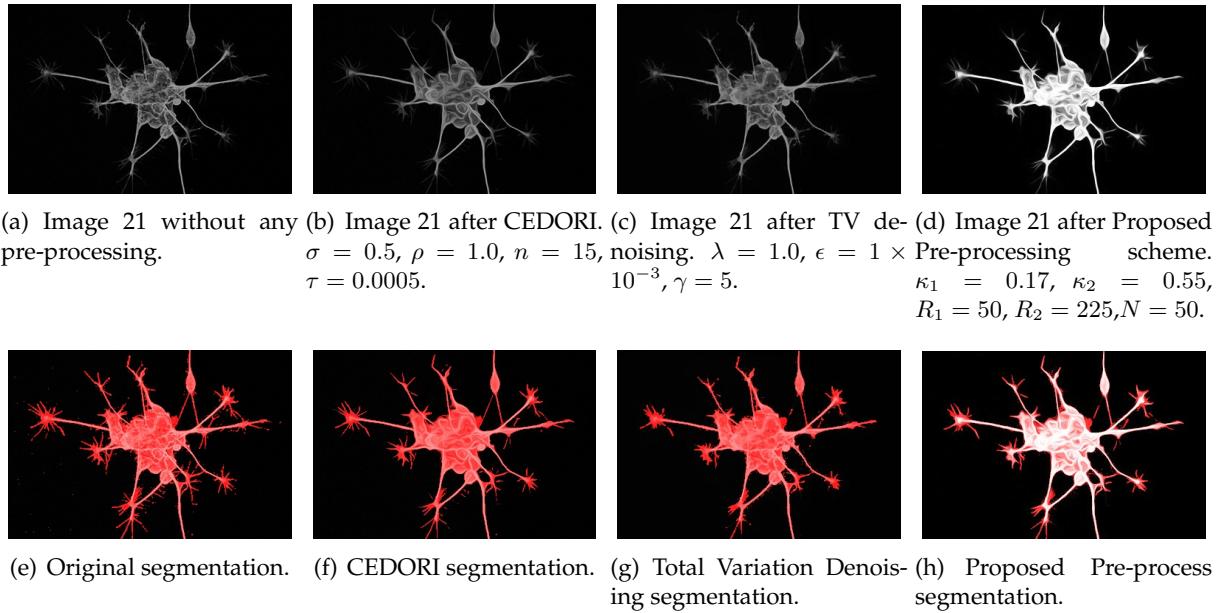


FIGURE 5.28: Image 21 from test set Appendix B pre-processing segmentation results. $\mu = 1, \lambda_0 = 300, \lambda_1 = 30$.

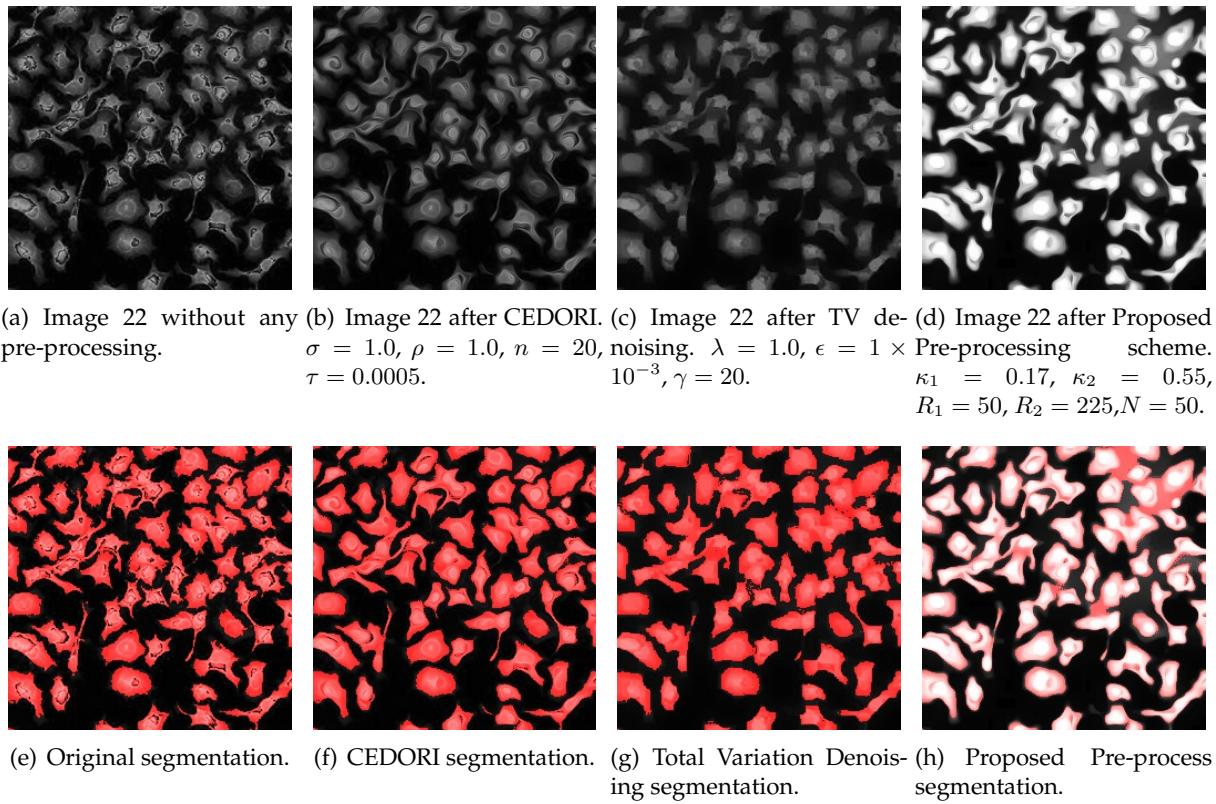


FIGURE 5.29: Image 22 from test set Appendix B pre-processing segmentation results. $\mu = 1, \lambda_0 = 5, \lambda_1 = 1$.

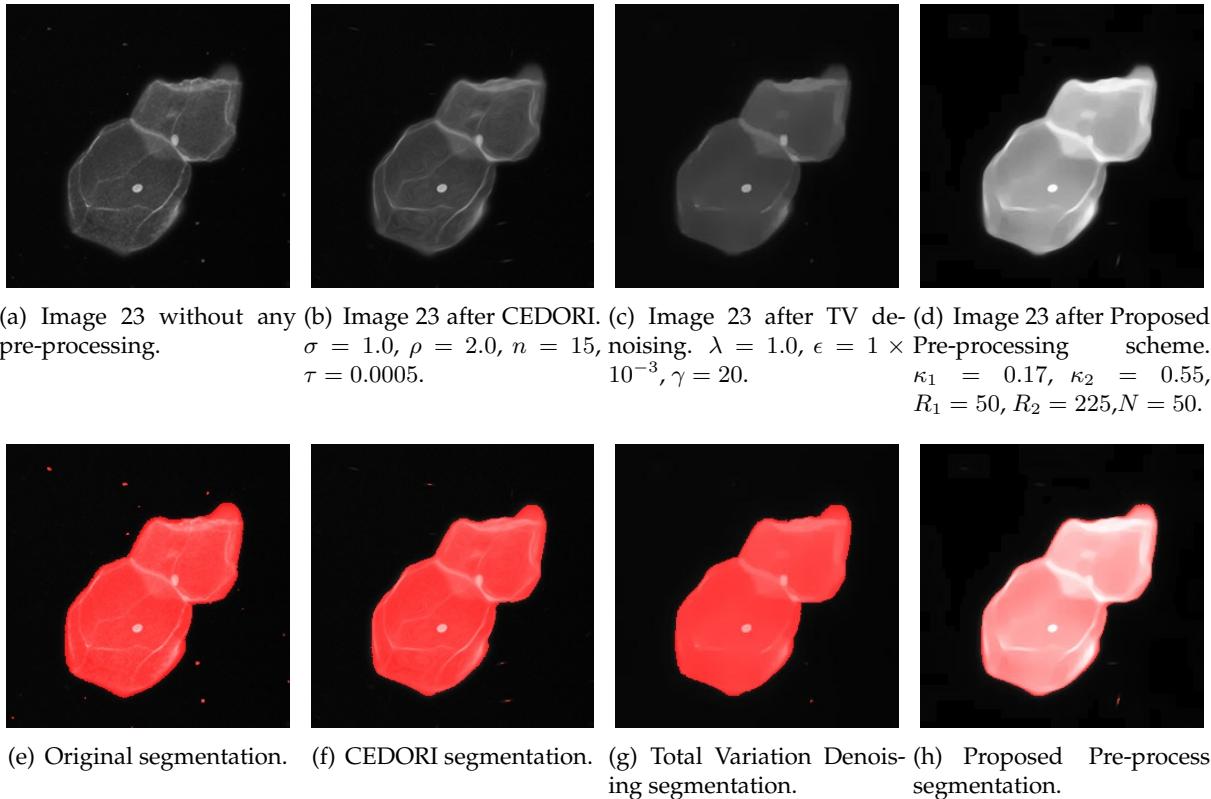


FIGURE 5.30: Image 23 from test set Appendix B pre-processing segmentation results. $\mu = 1, \lambda_0 = 5, \lambda_1 = 1$.

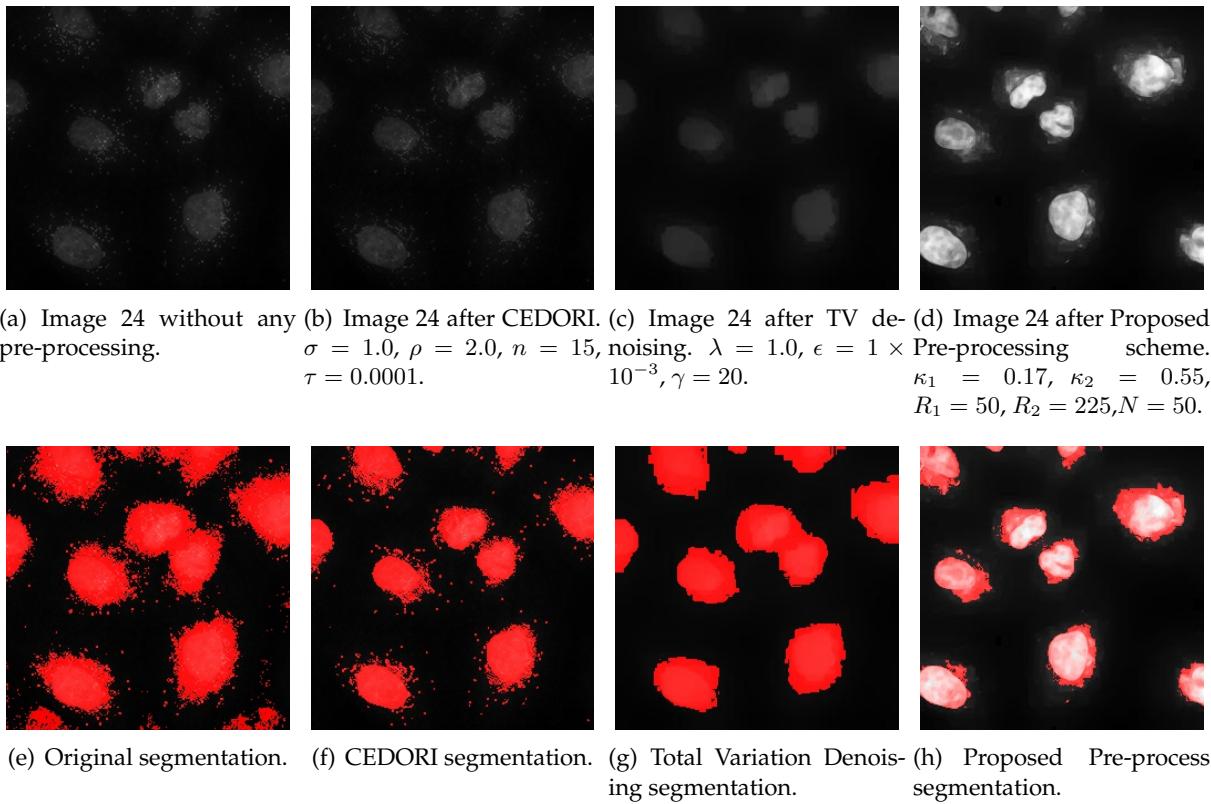


FIGURE 5.31: Image 24 from test set Appendix B pre-processing segmentation results. $\mu = 1, \lambda_0 = 5, \lambda_1 = 1$.

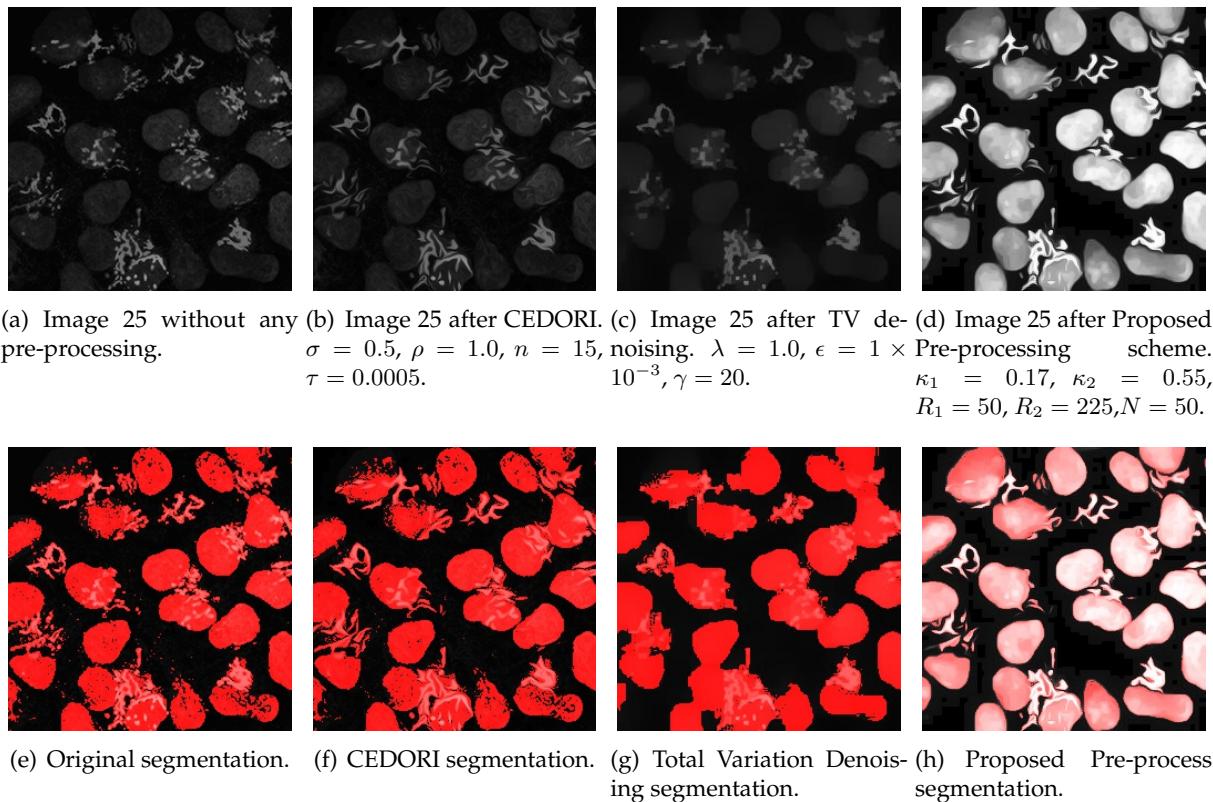


FIGURE 5.32: Image 25 from test set Appendix B pre-processing segmentation results. $\mu = 1, \lambda_0 = 5, \lambda_1 = 1$.

TABLE 5.1: Pre-processing Scheme Segmentation Results.

Image	TP	TN	FP	FN	Precision	Recall	Accuracy	MCC
1-o	38030	25503	13	1990	0.999658	0.950275	0.969437	0.938453
1-c	37673	25510	6	2347	0.999841	0.941354	0.964096	0.928266
1-t	37520	25511	5	2500	0.999867	0.937531	0.961777	0.923879
1-p	39215	25225	291	805	0.992634	0.979885	0.983276	0.965088
2-o	21312	39624	3966	634	0.843105	0.971111	0.929810	0.853334
2-c	21275	39136	4454	671	0.826888	0.969425	0.921799	0.838151
2-t	21250	36946	6644	696	0.761813	0.968286	0.888000	0.778750
2-p	20292	40378	3212	1654	0.863342	0.924633	0.925751	0.837361
3-o	11674	49067	4795	0	0.708847	1.000000	0.926834	0.803581
3-c	10105	53253	2178	0	0.822682	1.000000	0.966766	0.889020
3-t	9491	55234	197	614	0.979666	0.939238	0.987625	0.952035
3-p	11674	53280	582	0	0.952513	1.000000	0.991119	0.970681
4-o	15593	45595	4348	0	0.781957	1.000000	0.933655	0.844914
4-c	15593	46990	2953	0	0.840774	1.000000	0.954941	0.889416
4-t	15588	47537	2406	5	0.866289	0.999679	0.963211	0.907842
4-p	15593	47397	2546	0	0.859639	1.000000	0.961151	0.903226
5-o	24669	33229	7638	0	0.763581	1.000000	0.883453	0.787952
5-c	24669	35234	5633	0	0.814105	1.000000	0.914047	0.837789
5-t	24669	35134	5733	0	0.811427	1.000000	0.912521	0.835222
5-p	24669	35801	5066	0	0.829628	1.000000	0.922699	0.852517
6-o	37231	25337	418	2550	0.988897	0.935899	0.954712	0.908495
6-c	37685	25459	296	2096	0.992207	0.947312	0.963501	0.925927
6-t	38189	25218	537	1592	0.986133	0.959981	0.967514	0.932921
6-p	38072	25560	195	1709	0.994904	0.957040	0.970947	0.940811
7-o	14955	46580	771	3230	0.950973	0.822381	0.938950	0.845166
7-c	16706	46374	977	1479	0.944749	0.918669	0.962524	0.905905
7-t	17704	44904	2447	481	0.878567	0.973550	0.955322	0.894514
7-p	16833	45609	1742	1352	0.906218	0.925653	0.952789	0.883128
8-o	38556	22106	52	4822	0.998653	0.888838	0.925629	0.852381
8-c	39639	22114	44	3739	0.998891	0.913804	0.942276	0.882573
8-t	38925	22153	5	4453	0.999872	0.897344	0.931976	0.864207
8-p	40658	22123	35	2720	0.999140	0.937295	0.957962	0.912393

9-o	12324	50443	2767	2	0.816646	0.999838	0.957748	0.879778
9-c	12312	51825	1385	14	0.898883	0.998864	0.978653	0.934988
9-t	12301	51571	1639	25	0.882425	0.997972	0.974609	0.923572
9-p	12258	52314	896	68	0.931884	0.994483	0.985291	0.953825
10-o	34387	31026	122	1	0.996465	0.999971	0.998123	0.996243
10-c	34376	30955	193	12	0.994417	0.999651	0.996872	0.993742
10-t	34372	30809	339	16	0.990234	0.999535	0.994583	0.989183
10-p	34374	30829	319	14	0.990805	0.999593	0.994919	0.989851
11-o	24010	34854	5998	674	0.800120	0.972695	0.898193	0.803199
11-c	24060	34824	6028	624	0.799654	0.974720	0.898499	0.804291
11-t	24173	33511	7341	511	0.767056	0.979298	0.880188	0.775454
11-p	22670	37071	3781	2014	0.857056	0.918409	0.911575	0.815628
12-o	19444	39384	6577	131	0.747243	0.993308	0.897644	0.795294
12-c	19382	40090	5871	193	0.767513	0.990140	0.907471	0.811032
12-t	19171	39115	6846	404	0.736864	0.979361	0.889374	0.776794
12-p	15920	44976	985	3655	0.941733	0.813282	0.929199	0.828371
13-o	10441	42330	12619	146	0.452775	0.986210	0.805222	0.583053
13-c	10442	42277	12672	145	0.451761	0.986304	0.804428	0.582072
13-t	9583	39968	14981	1004	0.390124	0.905167	0.756088	0.480901
13-p	3818	46078	8871	6769	0.300891	0.360631	0.761353	0.185529
14-o	12749	44255	7640	892	0.625288	0.934609	0.869812	0.690498
14-c	12758	44119	7776	883	0.621311	0.935269	0.867874	0.687447
14-t	10713	43713	8182	2928	0.566975	0.785353	0.830475	0.562565
14-p	13338	39865	12030	303	0.525781	0.977788	0.811813	0.621765
15-o	3370	59038	3128	0	0.518621	1.000000	0.952271	0.701802
15-c	3370	58988	3178	0	0.514661	1.000000	0.951508	0.698821
15-t	3289	55293	6873	81	0.323657	0.975964	0.893890	0.528042
15-p	2316	62153	13	1054	0.994418	0.687240	0.983719	0.819597
16-o	7443	53278	4815	0	0.607195	1.000000	0.926529	0.746236
16-c	7388	57196	897	55	0.891732	0.992611	0.985474	0.932971
16-t	7365	57307	786	78	0.903570	0.989520	0.986816	0.938376
16-p	7228	57294	799	215	0.900461	0.971114	0.984528	0.926545
17-o	9824	54780	358	574	0.964840	0.944797	0.985779	0.946353
17-c	10398	49941	5197	0	0.666752	1.000000	0.920700	0.777115

17-t	10398	47423	7715	0	0.574063	1.000000	0.882278	0.702666
17-p	10064	55134	4	334	0.999603	0.967878	0.994843	0.980609
18-o	31266	28291	84	5895	0.997321	0.841366	0.908768	0.831616
18-c	31543	28021	354	5618	0.988902	0.848820	0.908875	0.829087
18-t	32157	28158	217	5004	0.993297	0.865343	0.920334	0.850014
18-p	28470	35580	377	1109	0.986931	0.962507	0.977325	0.954360
19-o	15285	46767	2533	951	0.857840	0.941426	0.946838	0.863586
19-c	14435	47043	2257	1801	0.864786	0.889074	0.938080	0.835567
19-t	15359	47526	2459	192	0.861993	0.987654	0.959549	0.897312
19-p	16195	49250	50	41	0.996922	0.997475	0.998611	0.996275
20-o	12628	50195	2224	489	0.850256	0.962720	0.958603	0.879548
20-c	12624	50209	2210	493	0.851018	0.962415	0.958755	0.879889
20-t	13015	49425	2994	102	0.812980	0.992224	0.952759	0.870803
20-p	12889	50599	1820	228	0.876266	0.982618	0.968750	0.909043
21-o	11423	53657	372	84	0.968461	0.992700	0.993042	0.976311
21-c	10725	53839	190	782	0.982593	0.932041	0.985168	0.948192
21-t	10113	53340	689	1394	0.936216	0.878856	0.968216	0.888145
21-p	9883	52360	1669	1624	0.855523	0.858869	0.949753	0.826708
22-o	26975	36034	56	2471	0.997928	0.916084	0.961441	0.924093
22-c	26883	35756	334	2563	0.987728	0.912959	0.955795	0.912233
22-t	26684	35248	842	2762	0.969411	0.906201	0.945007	0.889781
22-p	27007	34403	1687	2439	0.941207	0.917170	0.937042	0.872713
23-o	15978	48669	870	19	0.948362	0.998812	0.986435	0.964467
23-c	15848	49274	265	149	0.983554	0.990686	0.993683	0.982935
23-t	15904	49184	355	93	0.978166	0.994186	0.993164	0.981635
23-p	15852	49003	536	145	0.967293	0.990936	0.989609	0.972197
24-o	7859	43669	14006	2	0.359433	0.999746	0.786255	0.521556
24-c	7808	52834	4841	53	0.617282	0.993258	0.925323	0.748597
24-t	7850	48994	8681	11	0.474865	0.998601	0.867371	0.634457
24-p	502	46486	11189	7359	0.042939	0.063860	0.716980	-0.110446
25-o	28607	31722	307	4900	0.989382	0.853762	0.920547	0.849861
25-c	29110	31437	592	4397	0.980069	0.868774	0.923874	0.853820
25-t	29615	30347	1682	3892	0.946257	0.883845	0.914948	0.831958
25-p	29825	31755	274	274	0.990897	0.890113	0.939636	0.884272

TABLE 5.2: Overall Pre-processing Segmentation Efficiency.

Scheme	Precision		Recall		Accuracy	
	Mean	Std. Dev.	Mean	Std. Dev.	Mean	Std. Dev.
None	0.821354	0.186952	0.956262	0.054109	0.928629	0.052098
CED	0.844110	0.159666	0.958646	0.045520	0.939639	0.042925
TV	0.817916	0.203910	0.951100	0.057106	0.927104	0.056975
Proposed	0.859945	0.232488	0.883139	0.218280	0.940026	0.072479

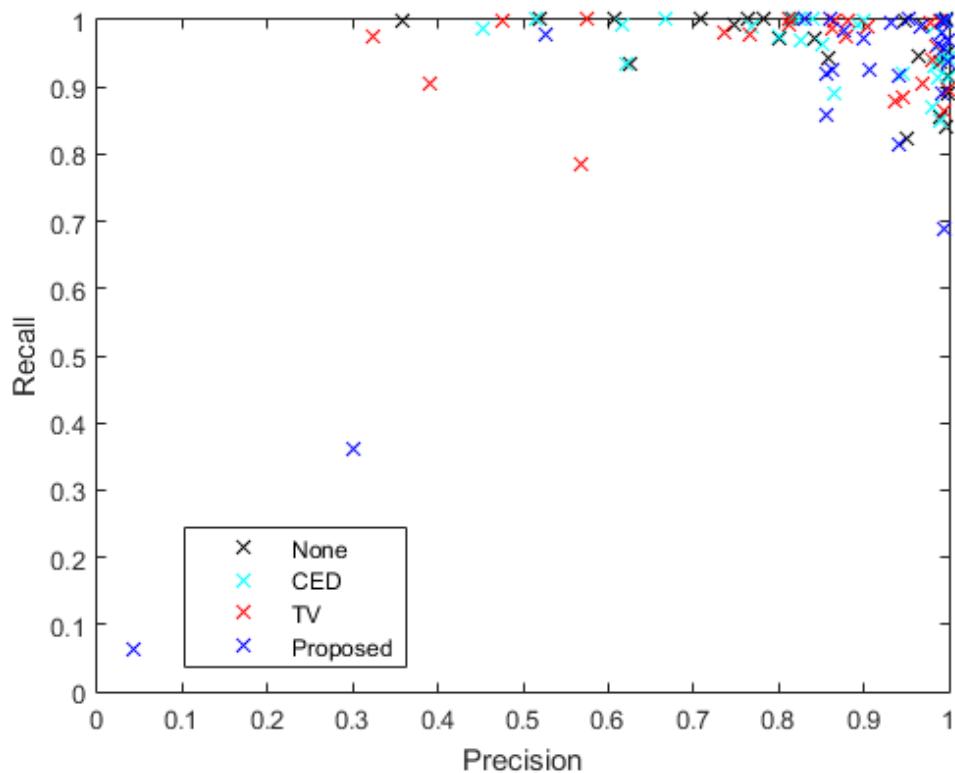


FIGURE 5.33: Pre-processing scheme and methods segmentation precision against recall over test set.

5.3 Discussion

We have presented a scheme that adjusts the properties of an image such that more accurate segmentation can be obtained. Experiments performed on 2D fluorescence microscopy image data show that the proposed scheme outperforms the existing techniques. One limitation of the scheme is the tuning of the various parameters. The running time can also be reduced by using parallel computing or by optimizing algorithms used. From the precision versus recall plot in Figure 5.33, we can see that all methods, even the original image after segmentation, produce very high precision and recall. This is due to the robustness of the Chan-Vese segmentation algorithm. However, over a large set we can see that the proposed method does allow for more accurate segmentation as seen in Table 5.2. From the

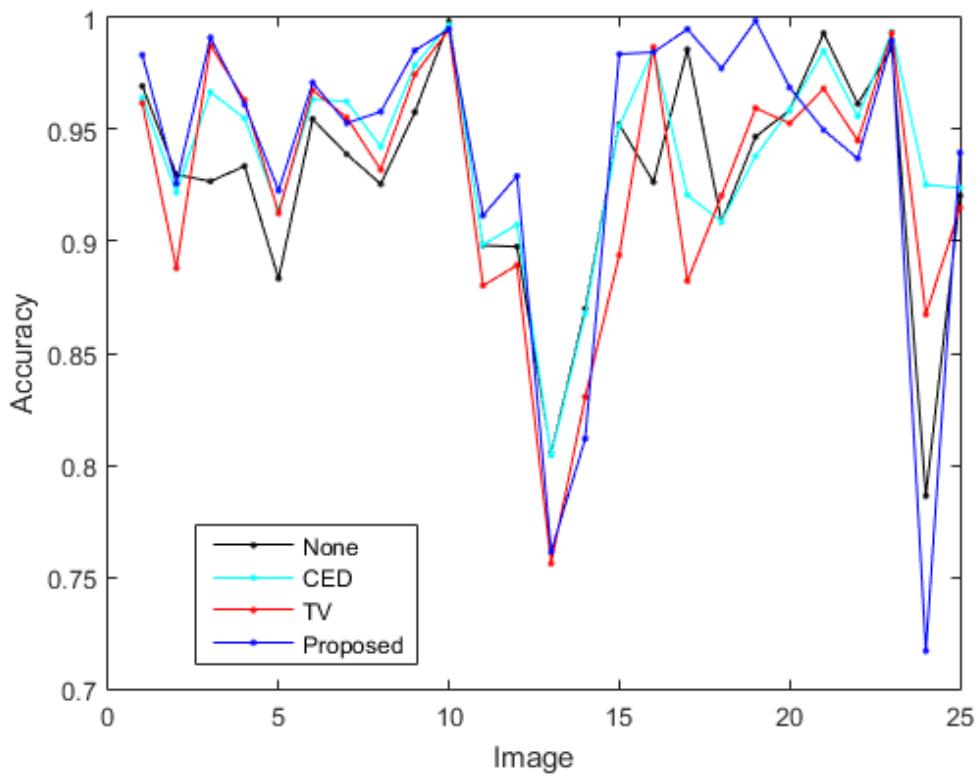


FIGURE 5.34: Pre-processing scheme and methods segmentation accuracy over test set.

accuracy plot in Figure 5.34, we can see that all methods suffer from producing inconsistent results; however, due to the fact that they're so close to the ground truth, this is negligible, for the most part.

5.4 Conclusion

Fluorescence imaging is a low light and low contrast imaging technique. Therefore, segmentation algorithms often have to work on images that have low SNR. To reduce the inaccuracies in the segmentation step, the image is normally pre-processed to allow for enhanced visualisation and segmentation accuracy.

For the segmentation technique used, which is already very robust against low contrast, the pre-processing scheme is there to improve the quality of the boundary and remove segmentation artefacts. We have presented a novel pre-processing scheme, which is a hybrid of algorithms that are designed to combat the problems typically found in fluorescence images. We also design a novel contrast enhancing function based on the non-linear remapping from a quadratic Bezier curve with intuitive parameters. The proposed scheme boosted the average segmentation accuracy to 94.0026%, which is a 1.1397% increase compared to an image with no pre-processing. The ground truth was developed on the original images. However, after the pre-processing, much of the hidden data surfaced which was captured by the segmentation algorithm. If the ground truth was developed on the enhanced image, then much better performance from the proposed method would have been seen.

Chapter 6

Parameter Estimation for ACWE Chan-Vese Segmentation

The Mumford-Shah functional [148] is used to find optimal sub-regions of an image. The image is modelled to as a piece-wise smooth function. This functional is difficult to optimise and is usually done by optimising an approximation. Chan and Vese [171] optimised an approximation of the Mumford-Shah functional by using variational level sets. Hence, the curves are not explicitly manipulated. Instead, they are implicitly manipulated by evolving the level set function. This was found to be highly robust and accurate in practice, even when algorithms terminated at near optimums. In the following section, we cover how this functional can be adapted for graph cut optimisation, for global optimisation. We then tweak the energy function for easier mathematical analysis and tune the parameters specifically for fluorescence image segmentation in Section 6.2. We present the results of our proposed energy function and parameter estimation method against other published parameter settings in Section 6.3. A discussion is presented in Section 6.4 which is followed by a conclusion in Section 6.5.

6.1 Graph Cut Model for Chan-Vese Segmentation

In this section, we briefly introduce the graph cut formulation for the Chan-Vese formulation of the Mumford-Shah evolution energy function for image segmentation. The Mumford-Shah model uses gradient descent techniques to obtain a minimum; but as previously discussed in Section 3.2.2, they usually terminate at local minima. By reformulating the energy function in a discrete form that allows for appropriate graph representability, we can use graph cuts which are able to terminate at a global minimum, to iteratively converge to the optimal solution. For an in-depth exposition into this technique, see [148, 171, 239].

The level set representation of the Mumford-Shah energy function is

$$\begin{aligned}
 F(c_1, c_2, \phi) = & \mu \int_{\Omega} \delta(\phi(x, y)) |\nabla \phi(x, y)| dx dy \\
 & + \nu \int_{\Omega} H(\phi(x, y)) dx dy \\
 & + \lambda_1 \int_{\Omega} |u(x, y) - c_1|^2 H(\phi(x, y)) dx dy \\
 & + \lambda_2 \int_{\Omega} |u(x, y) - c_2|^2 (1 - H(\phi(x, y))) dx dy,
 \end{aligned} \tag{6.1}$$

where $u(x, y)$ is the image, $H(\cdot)$ is the Heaviside step function, $\delta(\cdot)$ is the Dirac delta function and $\phi : \Omega \rightarrow \mathbb{R}$ is the level set function, such that:

$$\begin{aligned}\omega &= \{(x, y) \in \Omega | \Phi(x_p) > 0\} && \text{Inside the boundary} \\ \bar{\omega} &= \{(x, y) \in \Omega | \Phi(x_p) < 0\} && \text{Outside the boundary} \\ C &= \partial\omega = \{(x, y) \in \Omega | \Phi(x_p) = 0\} && \text{Along the boundary,}\end{aligned}\tag{6.2}$$

c_1 and c_2 are the arithmetic means given by:

$$c_1(\phi) = \frac{\int_{\Omega} u(x, y) H(\phi(x, y)) dx dy}{\int_{\Omega} H(\phi(x, y)) dx dy},\tag{6.3}$$

$$c_2(\phi) = \frac{\int_{\Omega} u(x, y) (1 - H(\phi(x, y))) dx dy}{\int_{\Omega} (1 - H(\phi(x, y))) dx dy}.\tag{6.4}$$

The piece-wise smooth approximation of the image is then

$$u(x, y) = c_1 H(\phi(x, y)) + c_2 (1 - H(\phi(x, y))).\tag{6.5}$$

Discrete Approximation of Contour Length For the energy function to be represented as a graph, one of the requirements is that it must be in a discrete representation. This means that the length of the contour, the first term in Equation (6.1), must be approximated discretely and be graph representable. This work has already been done by Kolmogorov and Boykov in [240, 241] where they used the Cauchy-Crofton theorem. The theorem states that the length of a curve can be approximated by drawing a large number of straight lines from 0 to 2π and counting the number of intersections between the lines and the contour. The mathematical representation is

$$\int_L n_L dL = \int_0^\pi \int_{-\infty}^\infty n_L d\rho d\theta = 2\|C\|_E,\tag{6.6}$$

where n_L is the number of intersections between the contour C and the line L , $\|C\|_E$ is the Euclidean length of the contour, $0 < \rho < \infty$ and $0 < \theta < 2\pi$. From this, the discrete approximation used by Boykov and Zabih is

$$\|C\|_E = \frac{1}{2} \sum_k n_k \frac{\delta^2 \Delta \theta_k}{|e_k|} = \frac{1}{2} \sum_k n_k w_k\tag{6.7}$$

An example of approximating the contour by two grids is illustrated in Figure 6.1(a) using four families of parallel lines which are 45° apart.

Discrete Representation of Mumford-Shah Function With the exception of the second term in Equation (6.1), the remaining terms are represented easily discretely. For each pixel $p \in \Omega$, let x_p be a binary variable such that

$$x_p = \begin{cases} 0 & \phi(p) \leq 0 \\ 1 & \phi(p) > 0 \end{cases}\tag{6.8}$$

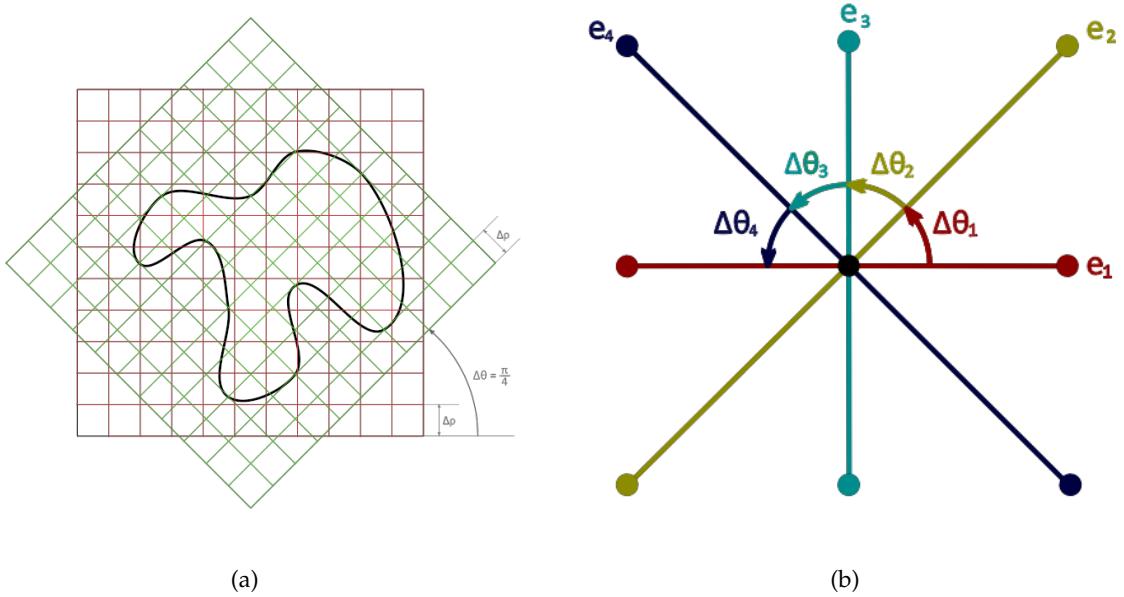


FIGURE 6.1: (a) Cauchy-Crofton length approximation. (b) 8-connected neighbourhood system.

The means can now be calculated using

$$c_1 = \frac{\sum_p u(x, y)x_p}{\sum_p x_p}, \quad (6.9)$$

$$c_2 = \frac{\sum_p u(x, y)(1 - x_p)}{\sum_p (1 - x_p)}. \quad (6.10)$$

For simplification, $\nu = 0$. To determine contour length using an 8-neighbourhood system, as illustrated in Figure 6.1(b), we set $\Delta\rho = 1$. The weight w_k is assigned to its corresponding edge e_k . The Euclidean length of the edges is $|e_1| = |e_3| = 1$ and $|e_2| = |e_4| = \sqrt{2}$. Therefore the corresponding weights, which are determined using Equation (6.7), is $w_1 = w_3 = \frac{\pi}{8}$ and $w_2 = w_4 = \frac{\pi}{8\sqrt{2}}$. To calculate n_k we need to count the intersections between the lines and the contour. An intersection between two pixels p and q exists if and only if x_p and x_q have different labels.

$$n_k = x_p(1 - x_q) + x_q(1 - x_p); k = (pq) \in \mathcal{N}_p. \quad (6.11)$$

The contour length can now fully be expressed discretely as

$$\|C\|_E = \sum_{p,q \in e_k} w_k(x_p(1 - x_q) + x_q(1 - x_p)). \quad (6.12)$$

The discrete representation of Equation (6.1) is

$$\begin{aligned} F(x_1, \dots, x_n) = & \mu \sum_{p,q \in e_k} w_k(x_p(1-x_q) + x_q(1-x_p)) \\ & + \lambda_1 \sum_p |u(x, y) - c_1|^2 x_p \\ & + \lambda_2 \sum_p |u(x, y) - c_2|^2 (1 - x_p) \end{aligned} \quad (6.13)$$

Graph Representation The discrete energy function Equation (6.13) has been shown that it obeys the submodularity constraint for graph representability. Therefore, the data energy and regularisation energy is

$$E^p(x_p) = \lambda_1 |u(x, y) - c_1|^2 x_p + \lambda_2 |u(x, y) - c_2|^2 (1 - x_p) \quad (6.14)$$

$$E^{pq}(x_p, x_q) = (x_p + x_q - 2x_p x_q) w_{pq} \quad (6.15)$$

The graph for the energy function is constructed as in [175].

6.2 Modified Weighting and Parameter Estimation

In this section, we introduce a novel idea to weighting a graph for image segmentation based on the Chan-Vese formulation of the Mumford-Shah evolution function. Previous parameter estimation schemes focused very specifically on certain images or image characteristics and this resulted in hard-coded parameters settings. In [239], where this method was first devised, they used the parameter settings $\mu = 0.1 \times 255^2$, $\lambda_1 = \lambda_2 = 1$. This worked very well on their synthetic images proving a strong resilience to noise and initial conditions. However, for fluorescence microscopy images, the results tend to be a bit too over-segmented for practical use. These parameters were used to segment the images in the sample set, Figure B.1, and the segmentation results are shown in Figure 6.2.

Masaka *et al.* [237] proposed a segmentation and tracking scheme for whole fluorescent cells in a time-lapse series. The result was another set of hard-coded parameters which were based on maximising the Jaccard coefficient of the automatically segmented cells in each frame of the time-lapse series which was also manually segmented. The optimal parameter settings were $\mu = 0.01$, $\lambda_1 = 1$ and $\lambda_2 = 85$. These parameters were used on the sample set in Figure B.1 and the results proved to be less useful as shown in Figure 6.3. It can be deduced that the parameters were specifically tuned to the image type that was studied in the time-lapse series.

It can be seen that hard-coded parameter values do not produce very consistent results over a large range of image types as that found in fluorescence microscopy imaging. In Sections 6.2.1 and 6.2.2, we devise a novel weighting scheme, and parameter estimation; as well as a new way to think of the parameters involved, in terms of *proxy relational parameters*. We then specifically focus on tuning these parameters for fluorescence images in Section 6.2.3.

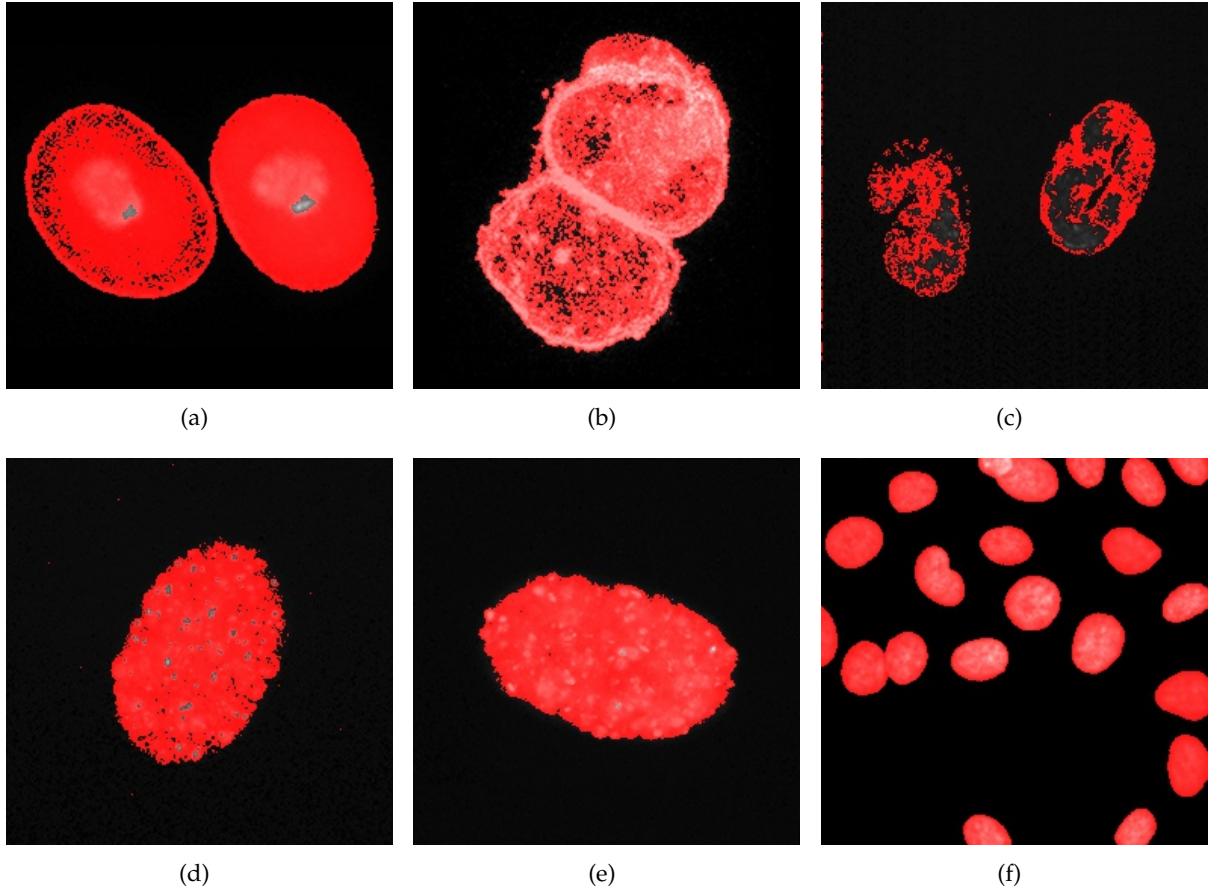


FIGURE 6.2: Segmentation results on fluorescence images using the default parameters settings provides by El-Zehiry *et al.* [239].

6.2.1 Graph Weighting

The first thing we do is normalise the weighting for both the data and smoothing connections. For the weighting of the neighbourhood connections we use the Euclidean distance between adjacent nodes. This results in neighbourhood connections as illustrated in Figure 6.4. The range of pixel intensities is also normalised i.e. $p \in [0, 1]$. The weight of the connection from the source to the node p is given by $E^i(0)|_{i=p} = \lambda_0|p - c_0|^2$. This is seen as how far away the pixel is from c_0 . Similarly, the weight of the connection from the node to the sink is given by $E^i(1)|_{i=p} = \lambda_1|p - c_1|^2$, i.e. how far away the pixel is from c_1 . The fully connected graph for a single node in the 8-connected neighbourhood system is illustrated in Figure 6.4.

6.2.2 Analysis of Weighting System and Parameter Relationships

To better understand the relationship between λ_0 and λ_1 and its impact on the final solution we explicitly formalise the dependency and set

$$\lambda_0 = \alpha\lambda_1. \quad (6.16)$$

Forcing this relation between λ_0 and λ_1 makes further analysis simpler and more intuitive. We can immediately see a constraint on α . Since, we require data connections to be positive, i.e. $E^i(0), E^i(1) \geq$

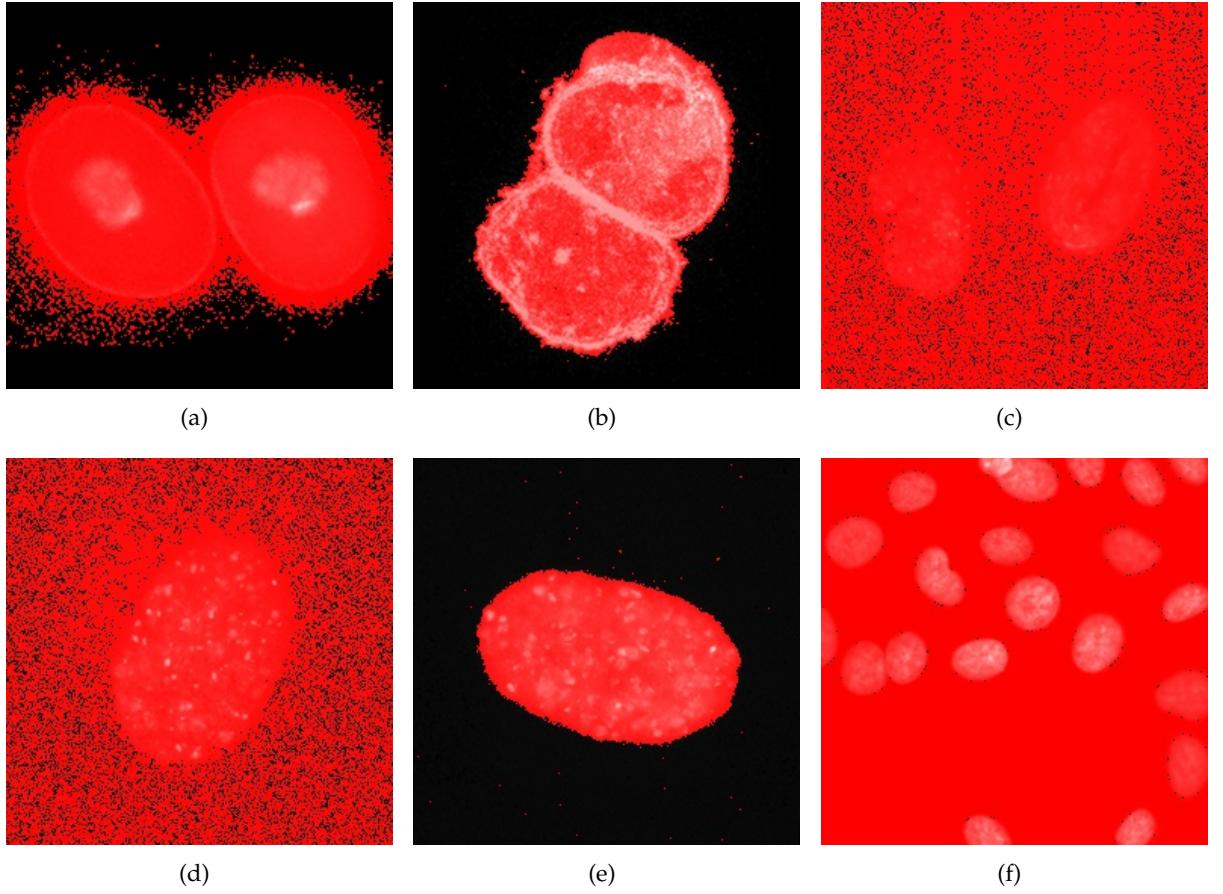


FIGURE 6.3: Segmentation results on fluorescence images using the parameters settings provides by Maska *et al.* [237].

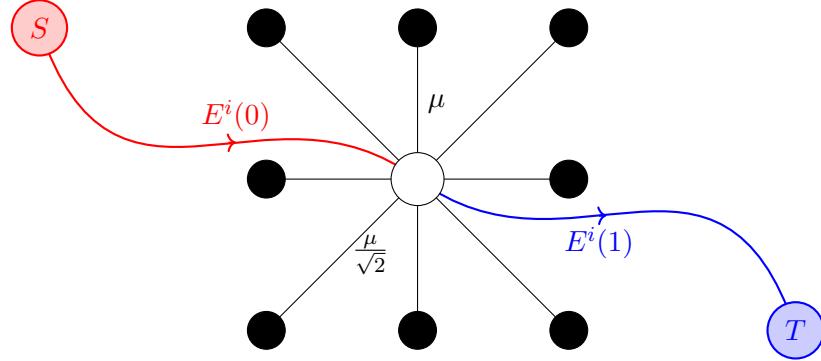


FIGURE 6.4: Fully connected single node.

, see Positivity constraint in Equation (3.3), this gives us a lower-bound on α for positive concavity of the energy functions

$$\alpha > 0 \text{ lower-bound on } \alpha \quad (6.17)$$

We will now analyse the flow through a single node. We use Figure 6.4 to facilitate our explanation. From the neighbourhood connections, in an 8-connected neighbourhood construction, the

maximum flow into or out of a node to its neighbours is

$$f_{max} = 4\mu + 4\frac{\mu}{\sqrt{2}} = \mu(2\sqrt{2} + 4). \quad (6.18)$$

To guarantee that a node p will be placed in the source set, $p \in S$, we know that the incoming flow from the source must completely saturate all flow outlets, this can be expressed as

$$E^i(0) > E^i(1) + \mu(2\sqrt{2} + 4). \quad (6.19)$$

This can be read as "*The source saturates the sink and all neighbourhood connections*". Similarly, to guarantee the node will be in the sink set, $p \in T$

$$E^i(1) > E^i(0) + \mu(2\sqrt{2} + 4). \quad (6.20)$$

This can be read as "*The sink is larger than the source and all neighbourhood connections*". To aid in understanding the energies, we use Figure 6.5.

For quadratic energies with $0 < c_0 < c_1 < 1$, there is a point, between c_0 and c_1 , where the incoming flow from the source completely saturates the sink with no excess remaining. This point, where the energies are equal, we call p_e , i.e. $E_0(p_e) = E_1(p_e)$. This point of zero net flow can be found as follows

$$\begin{aligned} E^{i=p_e}(1) &= E^{i=p_e}(0) \\ \lambda_1(p_e - c_1)^2 &= \lambda_0(p_e - c_0)^2 \\ \frac{(p_e - c_0)^2}{(p_e - c_1)^2} &= \frac{\lambda_1}{\lambda_0} \\ \frac{p_e - c_0}{p_e - c_1} &= \sqrt{\frac{\lambda_1}{\lambda_0}} \quad \text{or} \quad \frac{p_e - c_0}{p_e - c_1} = -\sqrt{\frac{\lambda_1}{\lambda_0}} \end{aligned}$$

We know that

$$\begin{aligned} c_0 &< p_e < c_1 \\ \therefore p_e - c_0 > 0 &\quad \text{and} \quad p_e - c_1 < 0 \end{aligned}$$

It follows directly that

$$\begin{aligned} \frac{p_e - c_1}{p_e - c_0} &= -\sqrt{\frac{\lambda_1}{\lambda_0}} \\ \frac{(p_e - c_0) + (c_0 - c_1)}{p_e - c_0} &= -\sqrt{\frac{\lambda_1}{\lambda_0}} \\ \frac{c_0 - c_1}{p_e - c_0} &= -\left(\sqrt{\frac{\lambda_1}{\lambda_0}} + 1\right) \\ p_e = c_0 + \frac{c_1 - c_0}{\sqrt{\frac{\lambda_1}{\lambda_0}} + 1} & \end{aligned}$$

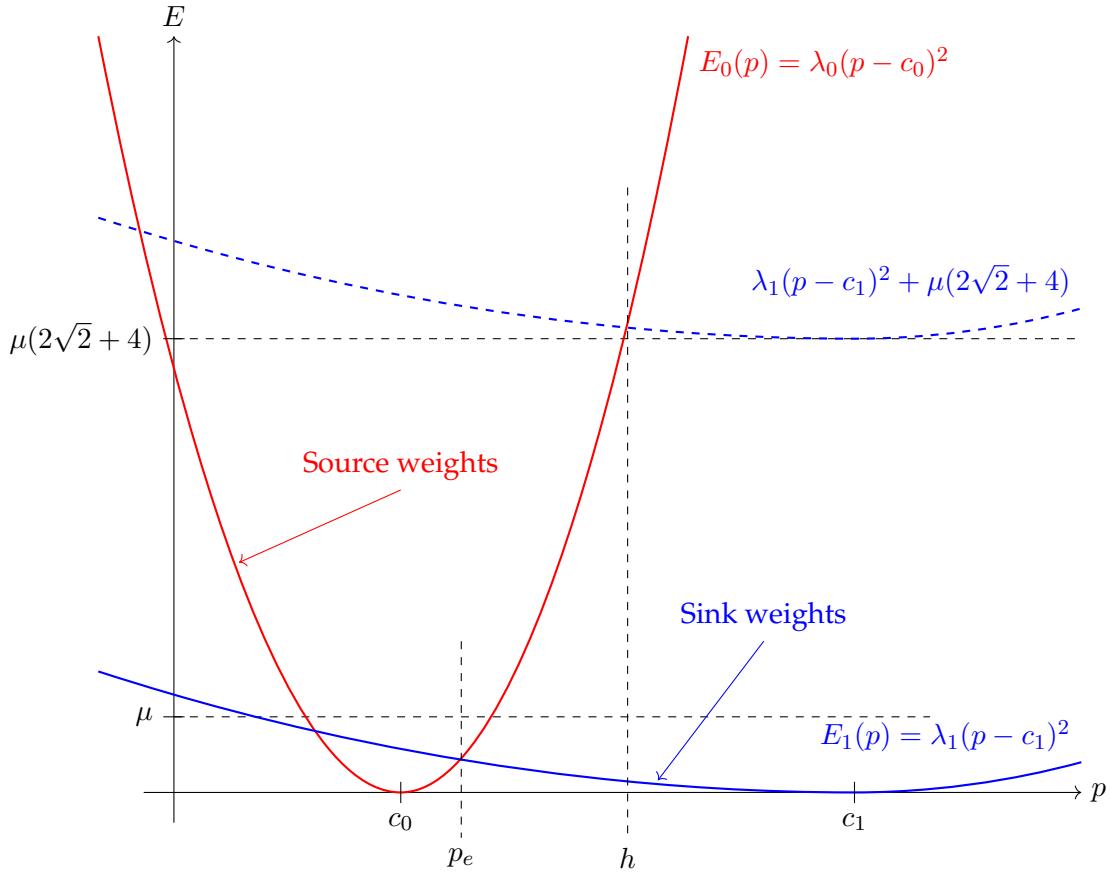


FIGURE 6.5: Data energy functions plot.

After substituting the relation in Equation (6.16) we get

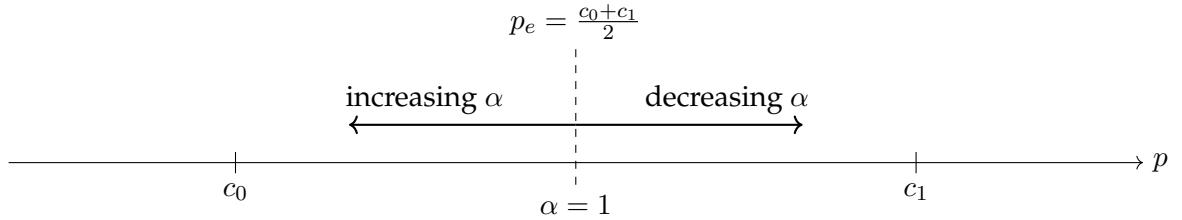
$$p_e = c_0 + \frac{c_1 - c_0}{\sqrt{\alpha} + 1} \quad (6.21)$$

The point where the energies are equal, p_e , is shown in Figure 6.5.

Analysis of the relationship between p_e and α From Equation (6.21) we note that there is one tuneable parameter, i.e. α . We can see that p_e and α are inversely related. This is expressed mathematically as

$$\begin{aligned} \text{if } \alpha = 1, p_e &= c_0 + \frac{c_1 - c_0}{1 + \sqrt{1}} = \frac{c_0 + c_1}{2} && \text{(midpoint between } c_0 \text{ and } c_1\text{)} \\ \lim_{\alpha \rightarrow \infty} p_e &= c_0 && \text{(maximum } \alpha \text{ yields lower-bound on } p_e\text{)} \\ \lim_{\alpha \rightarrow 0} p_e &= c_1 && \text{(minimum } \alpha \text{ yields upper-bound on } p_e\text{)} \end{aligned}$$

The relationship between p_e and α is illustrated in Figure 6.6.

FIGURE 6.6: Relationship between α and p_e .

Figuring α If we are able to make good estimates on p_e , c_0 and c_1 for the final segmented image, then it is possible to calculate α as follows:

$$\begin{aligned} p_e &= c_0 + \frac{c_1 - c_0}{\sqrt{\alpha} + 1} \\ 1 + \sqrt{\alpha} &= \frac{c_1 - c_0}{p_e - c_0} \\ \alpha &= \left(\frac{c_1 - c_0}{p_e - c_0} - 1 \right)^2 \end{aligned} \tag{6.22}$$

Lower-bound on μ When we found the point, p_e , where the energies are equal in Equation (6.21), we ignored the other solution as it was not within the range from c_0 to c_1 . Let this point be p_{e^*} . If this point is positive and $0 < p_{e^*} < c_0$ then we must ensure that at no point within this range must the source flow saturate all outgoing edges. This forces a limit on how low μ can be. This is only of significant concern when $\alpha > 1$. We only need to concern ourselves with the point $p = 0$ as this is the point where the difference $E^i(0) - E^i(1)$ is the largest. The lower-bound on μ can be obtained as follows

$$\begin{aligned} E^i(0)|_{p_i=0} &< E^i(1)|_{p_i=0} + \mu (2\sqrt{2} + 4) \\ \lambda_0 c_0^2 &< \lambda_1 c_1^2 + \mu (2\sqrt{2} + 4) \\ \therefore \mu (2\sqrt{2} + 4) &> \lambda_0 c_0^2 - \lambda_1 c_1^2 \\ \mu &> \frac{\lambda_0 c_0^2 - \lambda_1 c_1^2}{(2\sqrt{2} + 4)} \end{aligned}$$

Taking into account the relation in Equation (6.16) this becomes

$$\mu > \frac{\lambda_1 (\alpha c_0^2 - c_1^2)}{(2\sqrt{2} + 4)} \tag{6.23}$$

Absolutely in the source set From Equation (6.19) we can see that there is a point beyond which all nodes which correspond to pixel value higher than that point will be saturated and have excess flow which means that they will be in the source set. We will call this point the *saturation point* and denote

it by h . This is shown in Figure 6.5. This point can be determined as follows:

$$\begin{aligned}\lambda_0(h - c_0)^2 &> \lambda_1(h - c_1)^2 + f_{max} \\ \lambda_0(h - c_0)^2 - \lambda_1(h - c_1)^2 &> f_{max} \\ (\lambda_0 - \lambda_1)h^2 + (-2\lambda_0c_0 + 2\lambda_1c_1)h + (\lambda_0c_0^2 - \lambda_1c_1^2 - f_{max}) &> 0\end{aligned}$$

The solutions to h are

$$h = \frac{(2\lambda_0c_0 - 2\lambda_1c_1) \pm \sqrt{(-2\lambda_0c_0 + 2\lambda_1c_1)^2 - 4(\lambda_0 - \lambda_1)(\lambda_0c_0^2 - \lambda_1c_1^2 - f_{max})}}{2(\lambda_0 - \lambda_1)}$$

Substituting the relation in Equation (6.16)

$$\begin{aligned}h &= \frac{(\alpha c_0 - c_1) \pm \sqrt{(c_1 - \alpha c_0)^2 - (\alpha - 1)(\alpha c_0^2 - c_1^2 - \frac{f_{max}}{\lambda_1})}}{\alpha - 1} \\ &= \frac{(\alpha c_0 - c_1) \pm \sqrt{\alpha(c_0 - c_1)^2 + \frac{f_{max}}{\lambda_1}(\alpha - 1)}}{\alpha - 1}\end{aligned}$$

If the μ is greater than the lower-bound in Equation (6.23) then there is only one solution to h which is of importance. This is the positive solution for h which is

$$h = \frac{(\alpha c_0 - c_1) + \sqrt{\alpha(c_0 - c_1)^2 + \frac{\mu(2\sqrt{2}+4)}{\lambda_1}(\alpha - 1)}}{\alpha - 1} \quad (6.24)$$

This point is marked off in Figure 6.5.

Determining λ_1 Given good approximations for c_0 , c_1 , α , h and μ , we can calculate the appropriate value for λ_1 . We proceed from Equation (6.19) as follows

$$\begin{aligned}\lambda_0(h - c_0)^2 &= \lambda_1(h - c_1)^2 + \mu(2\sqrt{2} + 4) \\ \lambda_1(\alpha(h - c_0)^2 - (h - c_1)^2) &= \mu(2\sqrt{2} + 4) \\ \lambda_1 &= \frac{\mu(2\sqrt{2} + 4)}{\alpha(h - c_0)^2 - (h - c_1)^2} \quad (6.25)\end{aligned}$$

Parameter estimation process The parameter estimation is based on the assumption that sufficiently good approximations for c_0 , c_1 , p_e and h can be obtained. By sufficiently good we are referring to the closeness to the values these parameters would be for an ideal segmentation. From these approximations, we calculate the approximation for α using Equation (6.22). The parameters μ and λ_1 are related and are not separable, therefore we choose to set μ . We can then calculate λ_1 using Equation (6.25). For the chosen μ we can calculate the upper-bound on λ_1 to ensure that the constraint Equation (6.23) is met. The constraint on λ_1 is calculated as follows

$$\mu(2\sqrt{2} + 4) > \lambda_1(\alpha c_0^2 - c_1^2)$$

$$\lambda_1 < \frac{\mu(2\sqrt{2} + 4)}{\alpha c_0^2 - c_1^2} \quad (6.26)$$

Finally λ_0 can be calculated using Equation (6.16).

6.2.3 Tuning Parameters for Fluorescence Microscopy

The properties of the images obtained in fluorescence microscopy imaging can be used to guide the parameter estimation process. We focus specifically on black background images. Due to the fact that the predominant form of noise in the imaging system is Poisson distributed, we can further assume that the darker the background, the less noise that is present therein. The Poisson process also tells us that brighter regions exhibit a greater intensity variation due to the sampling process. Therefore, the curve for $E^i(1)$ is less convex than $E^i(0)$ as in Figure 6.5 and, resultantly, the value for p_e , in Figure 6.6, is shifted to the left. This places a new lower-bound on α for fluorescence images

$$\alpha \geq 1. \quad (6.27)$$

Manual Tuning Before moving further into analysis of the relationships between the parameters, we first perform a manual tuning of parameters and observing the effect on the segmented results. This allows us to understand how strongly correlated the parameter is to the final result. We note that the curves, $E^i(0)$ and $E^i(1)$, can be tuned relative to a fixed value for μ and this would not impact significantly on the range of possible solution sets. Therefore, we set $\mu = 1$ in all our manual parameter tuning. We use a stopping criterion of $\epsilon = 1 \times 10^{-3}$.

We cover a relatively wide range of parameter settings and note the output, i.e. over-segmented, under-segmented, almost ideal, etc. At this point, categorising the segmented output is largely subjective. We also use various initial conditions. Specifically, we use the segmented output from Otsu binarization, K-means ($k = 2$) and Expectation-Maximisation for Gaussian Mixture Modelling (EMGMM) with ($k = 2$). The initial and final means are of significant importance in this study.

For Figure B.1(a), we show the output for the following combinations of α and λ_1 . $(\alpha, \lambda_1) = \{(30, 150), (40, 50), (40, 100), (40, 150), (40, 200), (45, 150), (50, 150)\}$. The initial segmentation masks are shown in Figure 6.7. The segmented output for the combinations are shown in Figure 6.8.

For Figure B.1(b), we show the output for the following combinations of α and λ_1 . $(\alpha, \lambda_1) = \{(30, 150), (40, 50), (40, 100), (40, 150), (40, 200), (45, 150), (50, 150)\}$. The initial segmentation masks are shown in Figure 6.9. The segmented output for the combinations are shown in Figure 6.10.

For Figure B.1(c), we show the output for the following combinations of α and λ_1 . $(\alpha, \lambda_1) = \{(1, 150), (10, 150), (10, 200), (10, 400), (10, 800), (20, 150), (30, 150), (40, 150), (50, 150)\}$. The initial segmentation masks are shown in Figure 6.11. The segmented output for the combinations are shown in Figure 6.12.

For Figure B.1(d), we show the output for the following combinations of α and λ_1 . $(\alpha, \lambda_1) = \{(30, 50), (30, 100), (30, 150), (40, 100), (40, 150), (50, 150)\}$. The initial segmentation masks are shown in Figure 6.13. The segmented output for the combinations are shown in Figure 6.14.

For Figure B.1(e), we show the output for the following combinations of α and λ_1 . $(\alpha, \lambda_1) = \{(30, 50), (30, 100), (30, 150), (40, 100), (40, 150), (50, 150)\}$. The initial segmentation masks are shown in Figure 6.15. The segmented output for the combinations are shown in Figure 6.16.

For Figure B.1(f), we show the output for the following combinations of α and λ_1 . $(\alpha, \lambda_1) = \{(20, 20), (20, 150), (30, 150), (40, 50), (40, 150), (50, 150)\}$. The initial segmentation masks are shown in Figure 6.17. The segmented output for the combinations are shown in Figure 6.18.

The means and standard deviations for the initial masks obtained are shown in Table 6.1.

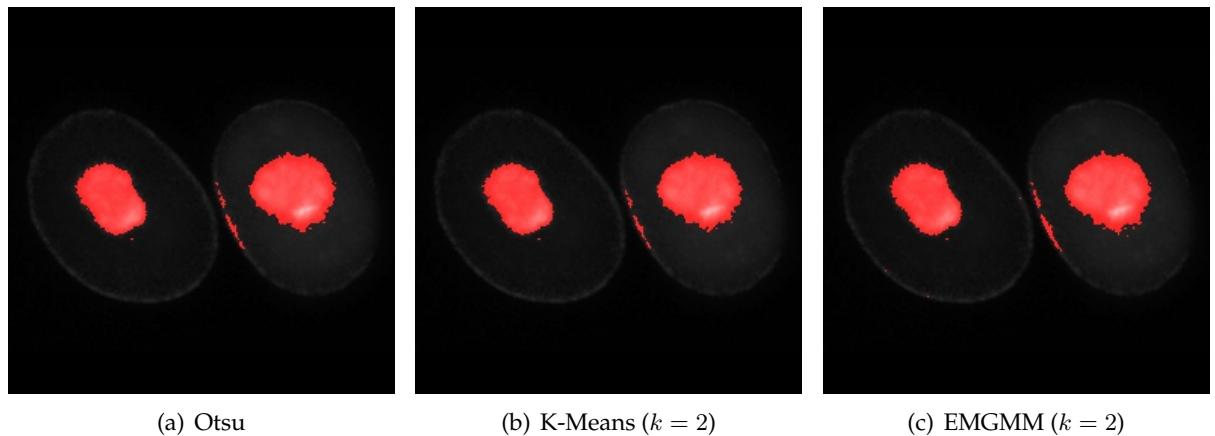
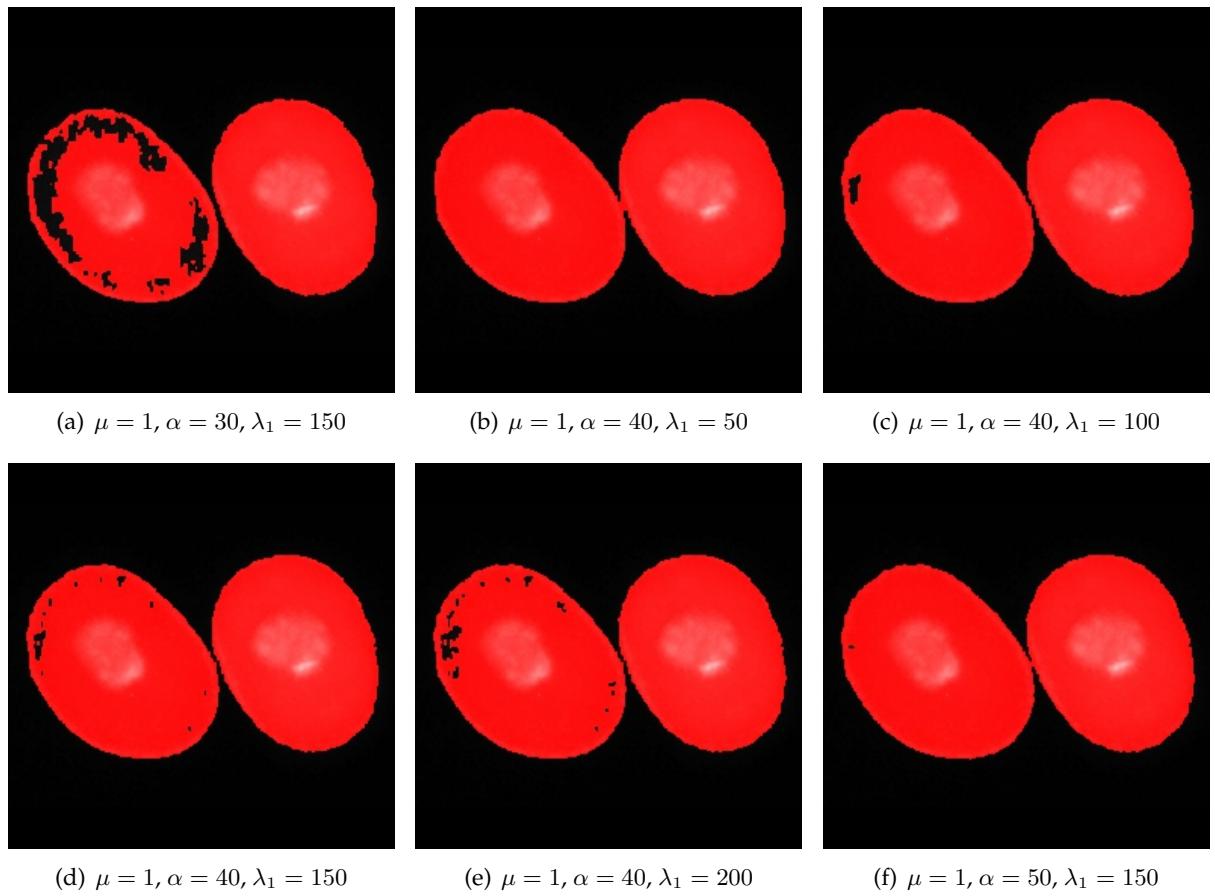


FIGURE 6.7: Image 1 from sample set Figure B.1 initial masks.

FIGURE 6.8: Segmented output for various combination of α and λ_1 for Image 1 in the sample set.

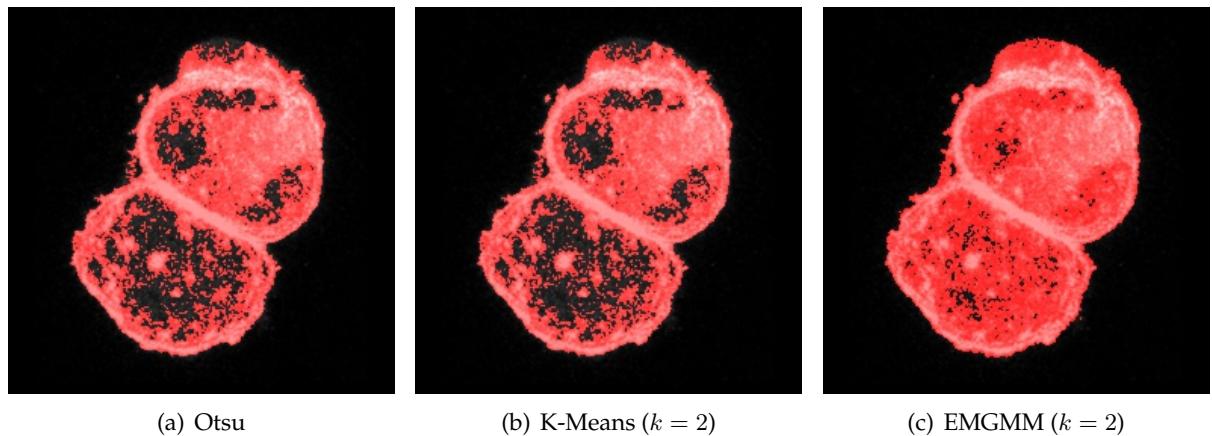
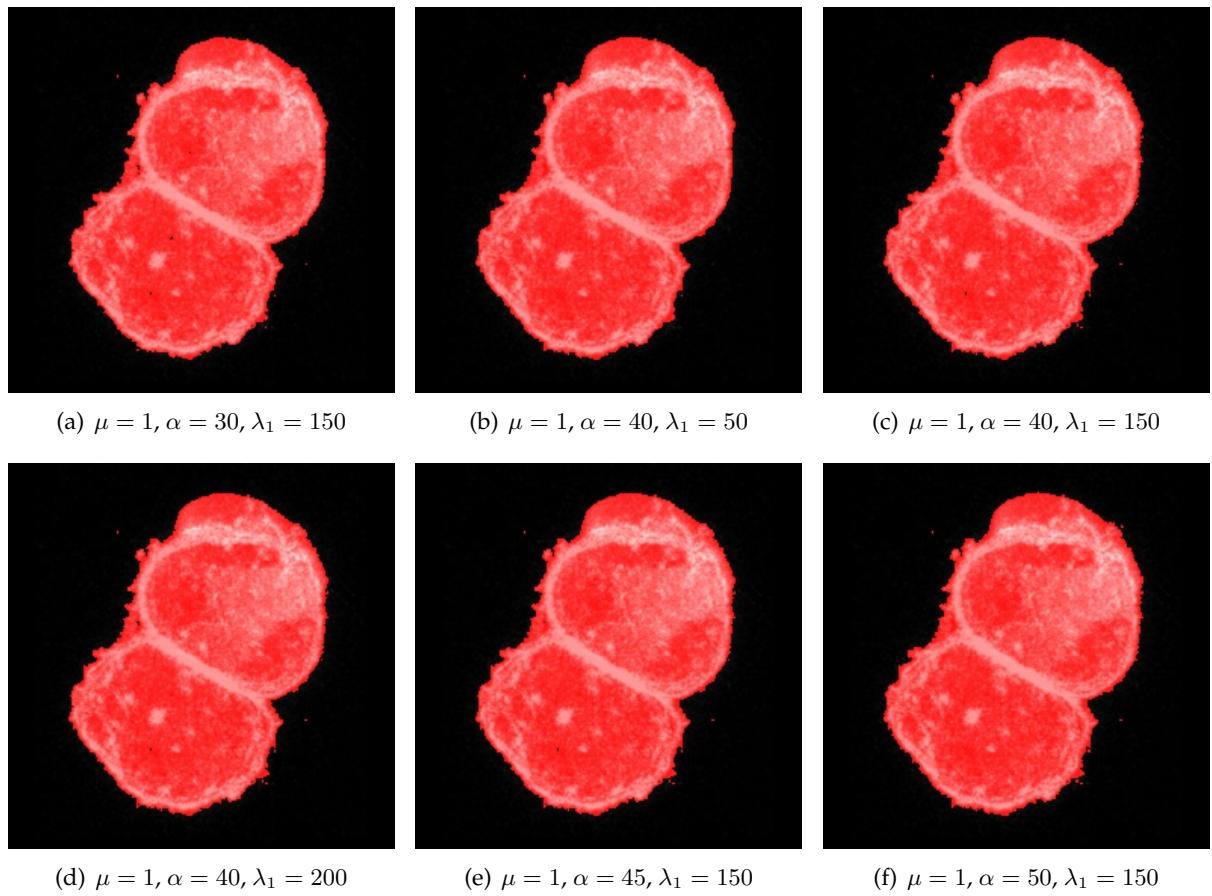


FIGURE 6.9: Image 2 from sample set Figure B.1 initial masks.

FIGURE 6.10: Segmented output for various combination of α and λ_1 for Image 2 in the sample set.

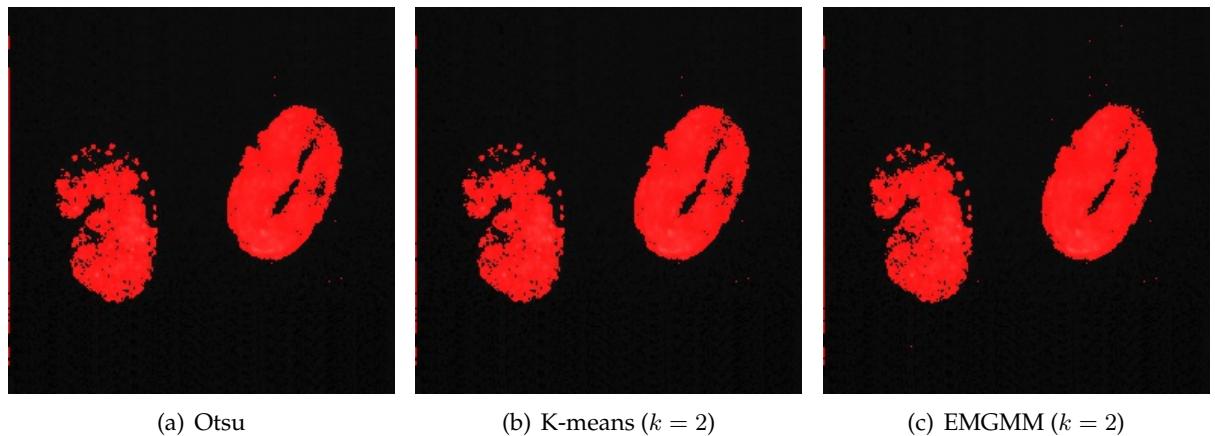
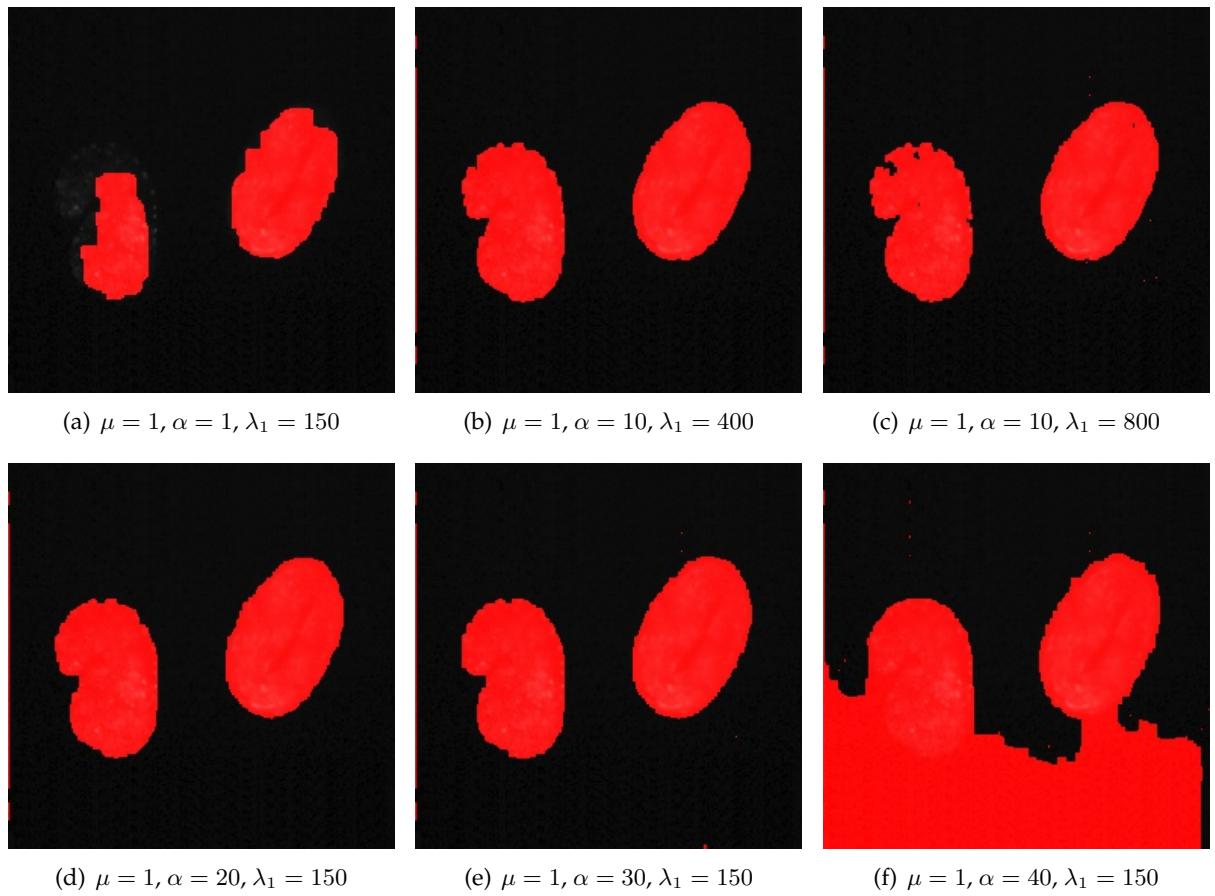


FIGURE 6.11: Image 3 from sample set Figure B.1 initial masks.

FIGURE 6.12: Segmented output for various combination of α and λ_1 for Image 3 in the sample set.

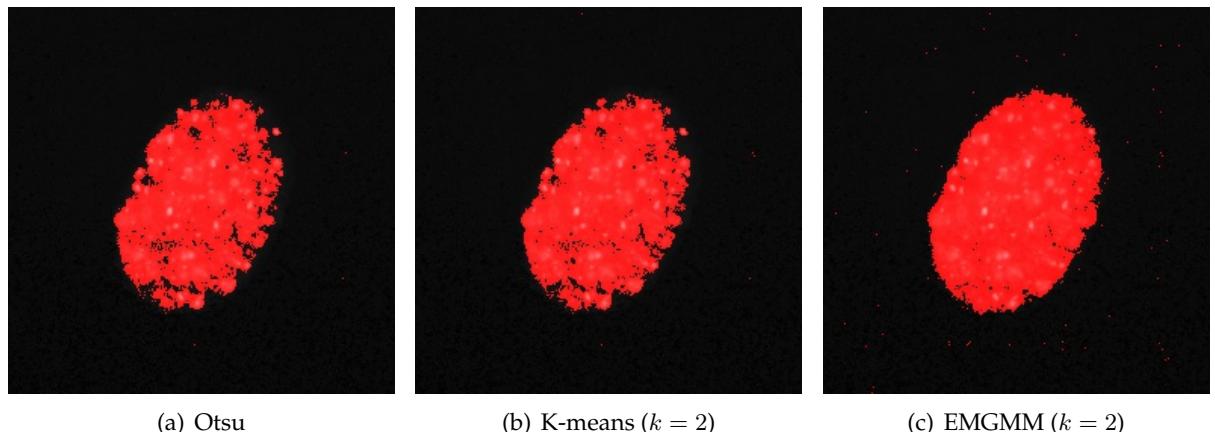
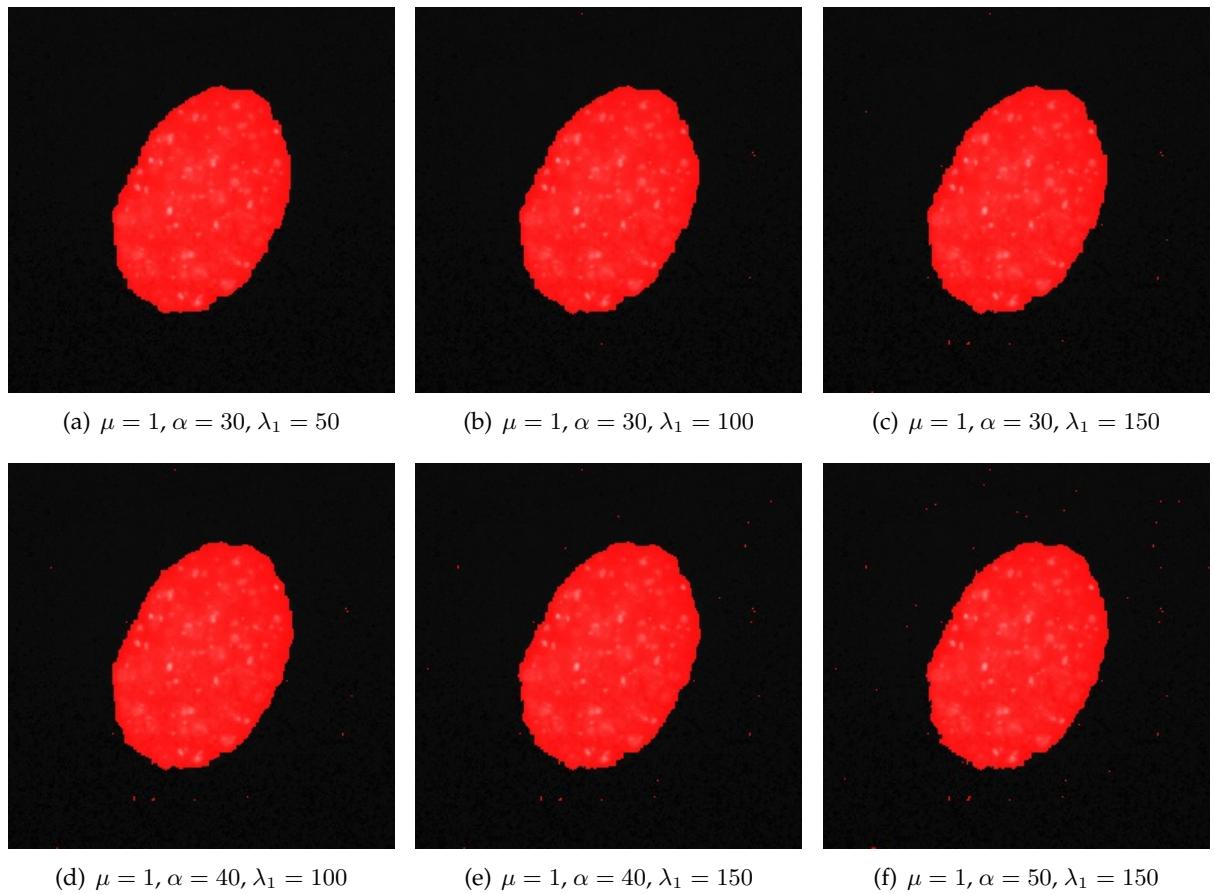


FIGURE 6.13: Image 4 from sample set Figure B.1 initial masks.

FIGURE 6.14: Segmented output for various combination of α and λ_1 for Image 4 in the sample set.

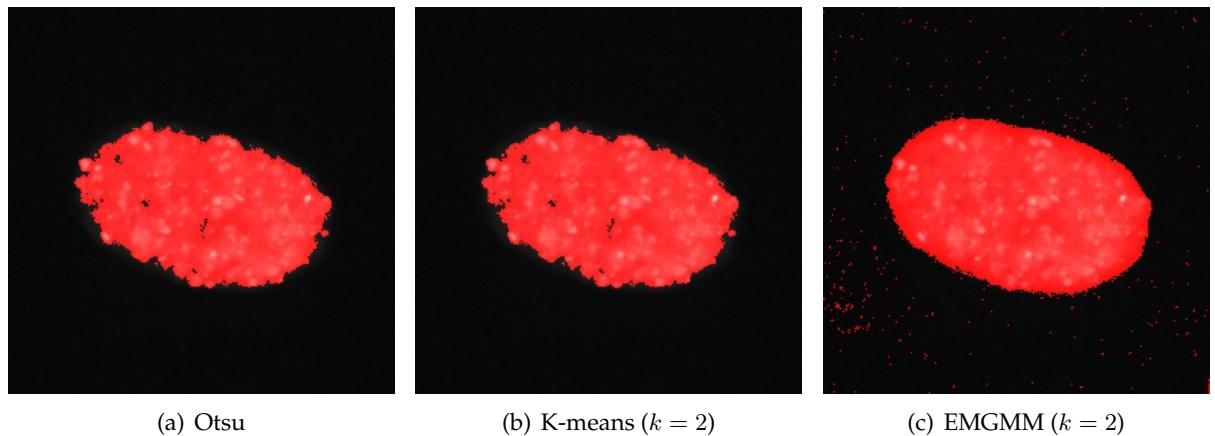
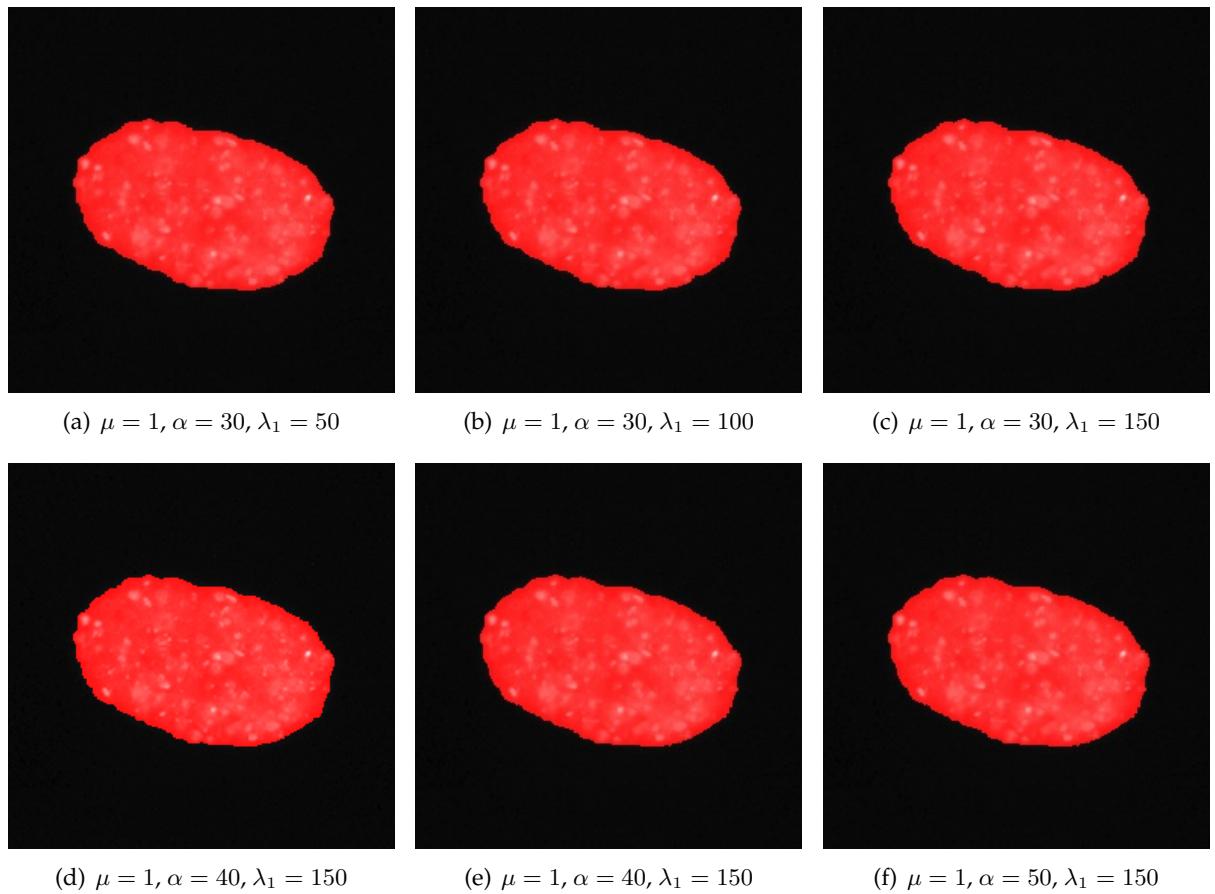


FIGURE 6.15: Image 5 from sample set Figure B.1 initial masks.

FIGURE 6.16: Segmented output for various combination of α and λ_1 for Image 5 in the sample set.

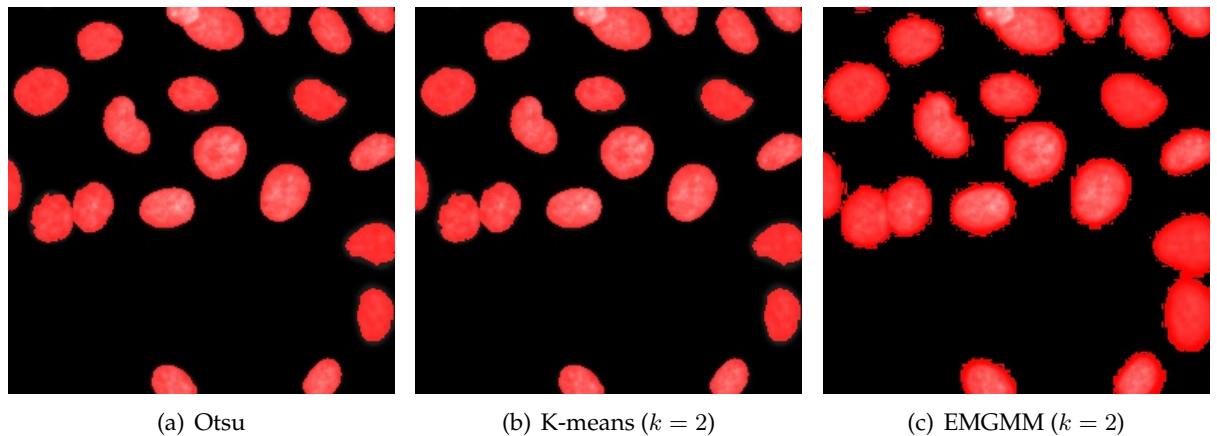


FIGURE 6.17: Image 6 from sample set Figure B.1 initial masks.

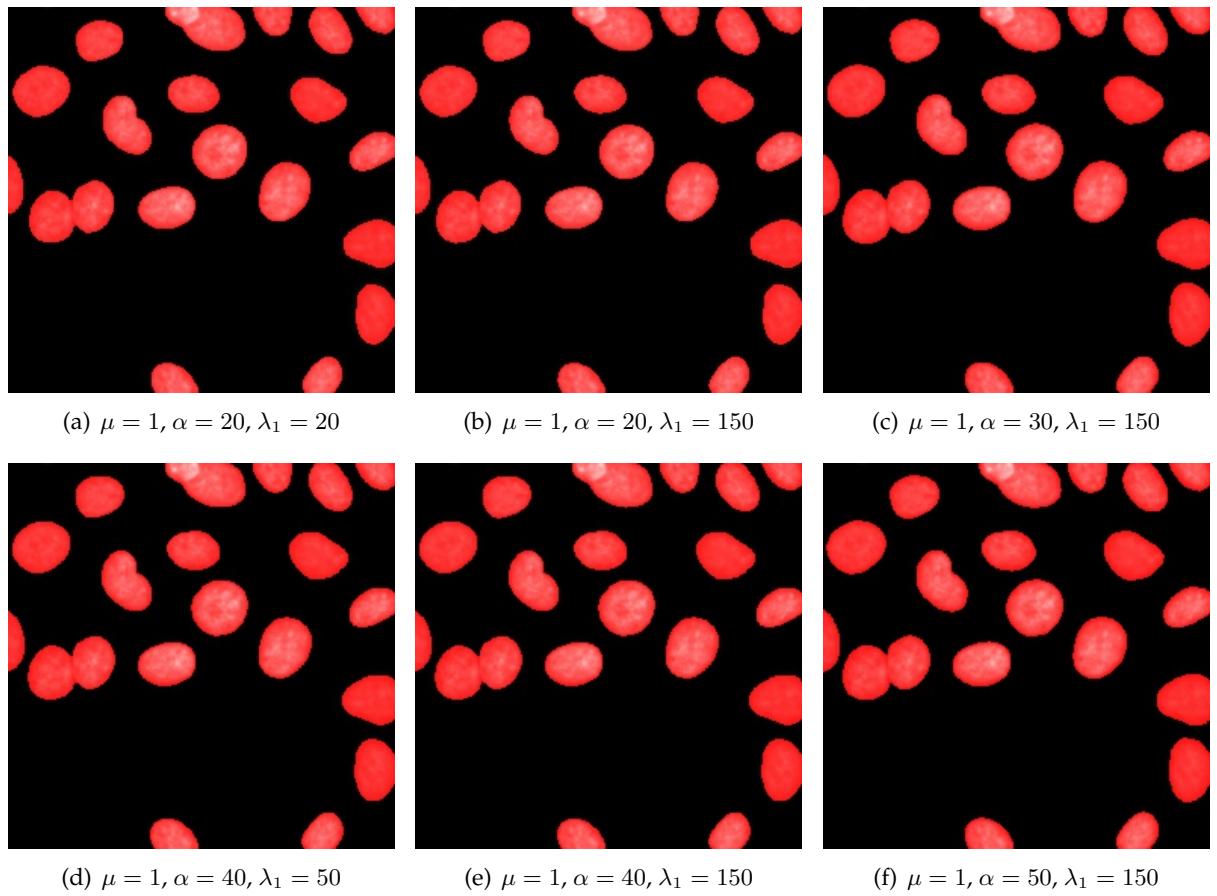
FIGURE 6.18: Segmented output for various combination of α and λ_1 for Image 6 in the sample set.

TABLE 6.1: Initial means and standard deviations for all images in the sample set for Otsu, K-means and EMGMM clustering.

	c_0	c_1	s_0	s_1
Image 1				
Otsu	0.0307515	0.26466	0.0385001	0, 0803506
K-means($k = 2$)	0.0308984	0.267192	0.0387009	0.0793621
EMGMM ($k = 2$)	0.0305648	0.261392	0.0382529	0.081608
Image 2				
Otsu	0.0293218	0.373293	0.054652	0.124376
K-means($k = 2$)	0.0293218	0.373293	0.054652	0.124376
EMGMM ($k = 2$)	0.010789	0.321058	0.0211362	0.140498
Image 3				
Otsu	0.0422978	0.105043	0.0104066	0.0269333
K-means($k = 2$)	0.0422978	0.105043	0.0104066	0.0269333
EMGMM ($k = 2$)	0.042026	0.103124	0.0100784	0.0273395
Image 4				
Otsu	0.0424602	0.147303	0.0133024	0.0513289
K-means($k = 2$)	0.0420671	0.145145	0.0125633	0.0513602
EMGMM ($k = 2$)	0.0407566	0.136167	0.0102742	0.052472
Image 5				
Otsu	0.037086	0.216513	0.0138978	0.0626164
K-means($k = 2$)	—	—	—	—
EMGMM ($k = 2$)	0.0343141	0.190501	0.00456055	0.0782783
Image 6				
Otsu	0.270483	0.00657006	0.0228327	0.0916426
K-means($k = 2$)	0.270483	0.00657006	0.0228327	0.0916426
EMGMM ($k = 2$)	0	0.187883	0	0.133022

The final values for the segmentation results are shown in Table 6.2. The table highlights extremely poor results in red, between 70%-90% accuracy in orange and remainder is left as is. We take the final results are the means of object and the background. From the final results, we calculate the values for p_e , Equation (6.21), and h , Equation (6.24). The means for each image can vary greatly. To put the values of p_e and h into a relative perspective, they are also shown as a fraction of the distance between c_0 and c_1 . Let $k_p \in (0, 1)$ be the fraction of the distance $p_e - c_0$ and $c_1 - c_0$ as illustrated in Figure 6.19. Let $k_h \in (k_p, 1)$ be the fraction of the distance $h - c_0$ and $c_1 - c_0$ as illustrated in Figure 6.20, we have $0 < k_p < k_h$.

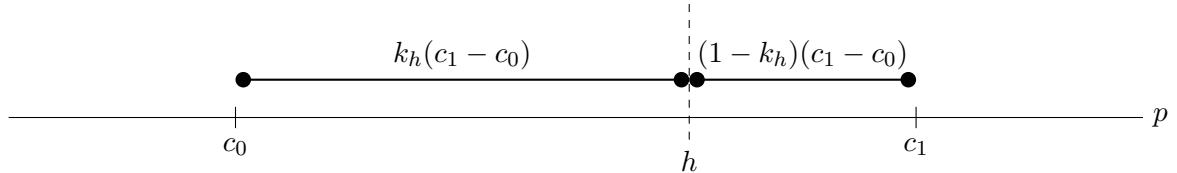
FIGURE 6.19: p_e as a fraction of the distance between c_0 and c_1 .FIGURE 6.20: h as a fraction of the distance between c_0 and c_1 .

TABLE 6.2: Results from manual tuning.

Image	α	λ_1	c_0	c_1	p_e	h	k_p	k_h
1-O	30	100	0.009171	0.117209	0.025851	0.058086	0.154387	0.452755
1-O	40	100	0.006909	0.110953	0.021114	0.049359	0.136527	0.407996
1-O	40	50	0.006803	0.110618	0.020976	0.065665	0.136527	0.566987
1-O	40	150	0.006953	0.111119	0.021175	0.042396	0.136527	0.340248
1-O	40	200	0.007093	0.111541	0.021353	0.038508	0.136527	0.300772
1-O	45	150	0.006832	0.110758	0.020314	0.040326	0.129732	0.322287
1-O	50	150	0.006681	0.110301	0.019519	0.038516	0.123899	0.307235
1-K	30	100	0.009557	0.118306	0.026346	0.058499	0.154387	0.450049
1-K	40	100	0.006928	0.111013	0.021138	0.049379	0.136527	0.407847
1-K	40	50	0.006818	0.110658	0.020995	0.065680	0.136527	0.566856
1-K	40	150	0.007063	0.111430	0.021312	0.042514	0.136527	0.339679
1-K	40	200	0.007176	0.111789	0.021459	0.038600	0.136527	0.300384
1-K	45	150	0.006852	0.110824	0.020340	0.040348	0.129732	0.322165
1-K	50	150	0.006685	0.110311	0.019524	0.038521	0.123899	0.307219
1-E	30	150	0.008876	0.116478	0.025489	0.049694	0.154387	0.379343
1-E	40	100	0.006687	0.110890	0.020913	0.049142	0.136527	0.407426
1-E	40	50	0.006788	0.110583	0.020959	0.065649	0.136527	0.567093
1-E	40	150	0.006944	0.111087	0.021163	0.042385	0.136527	0.340313
1-E	40	200	0.007058	0.111434	0.021308	0.038469	0.136527	0.300940
1-E	45	150	0.006800	0.110654	0.020273	0.040291	0.129732	0.322481
1-E	50	150	0.006676	0.110283	0.019512	0.038510	0.123899	0.307268
2-O	30	150	0.007640	0.309246	0.054204	0.066628	0.154387	0.195579

2-O	40	50	0.007449	0.308057	0.048489	0.076410	0.136527	0.229407
2-O	40	100	0.007428	0.307890	0.048449	0.063950	0.136527	0.188118
2-O	40	150	0.007424	0.307975	0.048457	0.059239	0.136527	0.172400
2-O	40	200	0.007441	0.308094	0.048488	0.056764	0.136527	0.164052
2-O	45	150	0.007423	0.307976	0.046415	0.056577	0.129732	0.163545
2-O	50	150	0.007297	0.307166	0.044450	0.054107	0.123899	0.156103
2-K	30	150	0.007640	0.309246	0.054204	0.066628	0.154387	0.195579
2-K	40	50	0.007449	0.308057	0.048489	0.076410	0.136527	0.229407
2-K	40	100	0.007428	0.307890	0.048449	0.063950	0.136527	0.188118
2-K	40	150	0.007424	0.307975	0.048457	0.059239	0.136527	0.172400
2-K	40	200	0.007441	0.308094	0.048488	0.056764	0.136527	0.164052
2-K	45	150	0.007423	0.307976	0.046415	0.056577	0.129732	0.163545
2-K	50	150	0.007297	0.307166	0.044450	0.054107	0.123899	0.156103
2-E	30	150	0.007556	0.308779	0.054061	0.066498	0.154387	0.195674
2-E	40	50	0.007568	0.308796	0.048694	0.076578	0.136527	0.229095
2-E	40	100	0.007564	0.308815	0.048693	0.064163	0.136527	0.187881
2-E	40	150	0.007566	0.308838	0.048697	0.059458	0.136527	0.172244
2-E	40	200	0.007558	0.308800	0.048686	0.056948	0.136527	0.163952
2-E	45	150	0.007467	0.308260	0.046489	0.056646	0.129732	0.163496
2-E	50	150	0.007343	0.307470	0.044528	0.054178	0.123899	0.156053
3-O	10	150	0.041366	0.096327	0.054571	0.108955	0.240253	1.229755
3-O	10	200	0.041364	0.096323	0.054568	0.099806	0.240253	1.063370
3-O	10	400	0.041359	0.096339	0.054568	0.082894	0.240253	0.755457
3-O	10	800	0.041371	0.096708	0.054666	0.071643	0.240253	0.547049
3-O	20	150	0.041323	0.095848	0.051287	0.089056	0.182743	0.875426
3-O	30	150	0.041267	0.094999	0.049563	0.080313	0.154387	0.726674
3-O	40	150	0.044802	0.057616	0.046551	0.078701	0.136527	2.645434
3-K	10	150	0.041366	0.096327	0.054571	0.108955	0.240253	1.229755
3-K	10	200	0.041364	0.096323	0.054568	0.099806	0.240253	1.063370
3-K	10	400	0.041359	0.096339	0.054568	0.082894	0.240253	0.755457
3-K	10	800	0.041371	0.096708	0.054666	0.071643	0.240253	0.547049
3-K	20	150	0.041323	0.095848	0.051287	0.089056	0.182744	0.875426
3-K	30	150	0.041267	0.094999	0.049563	0.080313	0.154387	0.726674
3-K	40	150	0.044802	0.057616	0.046551	0.078701	0.136527	2.645434

3-E	10	150	0.041362	0.096283	0.054557	0.108952	0.240253	1.230667
3-E	10	200	0.041357	0.096260	0.054548	0.099799	0.240253	1.064463
3-E	10	400	0.041356	0.096309	0.054559	0.082890	0.240253	0.755812
3-E	10	800	0.041367	0.096932	0.054717	0.071656	0.240253	0.545117
3-E	20	150	0.041321	0.095836	0.051284	0.089054	0.182744	0.875599
3-E	30	150	0.041262	0.094923	0.049545	0.080307	0.154387	0.727635
3-E	40	150	0.044849	0.057396	0.046562	0.078753	0.136527	2.702195
4-O	30	50	0.040339	0.131121	0.054355	0.107943	0.154387	0.744683
4-O	30	100	0.040328	0.131038	0.054332	0.088659	0.154387	0.532819
4-O	30	150	0.040319	0.130976	0.054315	0.080354	0.154387	0.441615
4-O	40	100	0.040240	0.129694	0.052453	0.082233	0.136527	0.469438
4-O	40	150	0.040264	0.130244	0.052549	0.075108	0.136527	0.387237
4-O	50	150	0.040232	0.129761	0.051324	0.071509	0.123899	0.349362
4-K	30	50	0.040356	0.131362	0.054406	0.107962	0.154387	0.742875
4-K	30	100	0.040343	0.131304	0.054386	0.088682	0.154387	0.531430
4-K	30	150	0.040339	0.131307	0.054383	0.080387	0.154387	0.440244
4-K	40	100	0.040233	0.129556	0.052428	0.082222	0.136527	0.470084
4-K	40	150	0.040260	0.130177	0.052536	0.075101	0.136527	0.387482
4-K	50	150	0.040223	0.129553	0.051291	0.071494	0.123899	0.350061
4-E	30	50	0.040262	0.129873	0.054097	0.107852	0.154387	0.754261
4-E	30	100	0.040254	0.129884	0.054091	0.088555	0.154387	0.538901
4-E	30	150	0.040237	0.129708	0.054049	0.080225	0.154387	0.446937
4-E	40	100	0.040236	0.129624	0.052439	0.082227	0.136527	0.469763
4-E	40	150	0.040225	0.129532	0.052418	0.075043	0.136527	0.389868
4-E	50	150	0.040201	0.129085	0.051213	0.071456	0.123899	0.351638
5-O	30	50	0.034756	0.201631	0.060519	0.104517	0.154387	0.418045
5-O	30	150	0.034757	0.201659	0.060525	0.079633	0.154387	0.268872
5-O	40	150	0.034675	0.200630	0.057333	0.073912	0.136527	0.236429
5-O	50	150	0.034487	0.197806	0.054722	0.069683	0.123899	0.215506
5-K	30	50	0.034756	0.201631	0.060519	0.104517	0.154387	0.418045
5-K	30	150	0.034757	0.201659	0.060525	0.079633	0.154387	0.268872
5-K	40	150	0.034675	0.200630	0.057333	0.073912	0.136527	0.236429
5-K	50	150	0.034487	0.197806	0.054722	0.069683	0.123899	0.215506
5-E	30	50	0.034683	0.200718	0.060316	0.104407	0.154387	0.419937

5-E	30	150	0.034681	0.200701	0.060313	0.079483	0.154387	0.269859
5-E	40	150	0.034582	0.199297	0.057069	0.073726	0.136527	0.237648
5-E	50	150	0.034580	0.199280	0.054986	0.069870	0.123899	0.214269
6-O	20	20	0.002964	0.251959	0.048466	0.136161	0.182744	0.534939
6-O	20	150	0.002932	0.251781	0.048407	0.066168	0.182744	0.254113
6-O	30	50	0.001533	0.228802	0.036620	0.074639	0.154387	0.321672
6-O	30	150	0.001521	0.238690	0.038137	0.053144	0.154387	0.217666
6-O	40	50	0.001817	0.242106	0.034623	0.066509	0.136527	0.269226
6-O	40	150	0.001815	0.242097	0.034619	0.047477	0.136527	0.190034
6-O	50	50	0.001259	0.235130	0.030236	0.059146	0.123899	0.247513
6-O	50	150	0.001239	0.234857	0.030184	0.041921	0.123899	0.174135
6-K	20	20	0.002964	0.251959	0.048466	0.136161	0.182744	0.534939
6-K	20	150	0.002932	0.251781	0.048407	0.066168	0.182744	0.254113
6-K	30	50	0.001533	0.228802	0.036620	0.074639	0.154387	0.321672
6-K	30	150	0.001521	0.238690	0.038137	0.053144	0.154387	0.217666
6-K	40	50	0.001817	0.242106	0.034623	0.066509	0.136527	0.269226
6-K	40	150	0.001814	0.242097	0.034619	0.047476	0.136527	0.190034
6-K	50	50	0.001259	0.235130	0.030236	0.059146	0.123899	0.247513
6-K	50	150	0.001239	0.234857	0.030185	0.041921	0.123899	0.174135
6-E	20	20	0.002643	0.249529	0.047759	0.135753	0.182744	0.539155
6-E	20	150	0.002584	0.249164	0.047645	0.065530	0.182744	0.255277
6-E	30	50	0.002252	0.246370	0.039940	0.076508	0.154387	0.304183
6-E	30	150	0.002246	0.246324	0.039928	0.054615	0.154387	0.214558
6-E	40	50	0.001527	0.238746	0.033914	0.066025	0.136527	0.271893
6-E	40	150	0.001519	0.238667	0.033896	0.046879	0.136527	0.191277
6-E	50	50	0.001655	0.240298	0.031222	0.059817	0.123899	0.243720
6-E	50	150	0.001645	0.240193	0.031201	0.042755	0.123899	0.172337

Upon comparing the initial means in Table 6.1 and final means for the acceptable and good segmentation results in Table 6.2, the values of the initial means are larger. This is due to over-segmentation produced by Otsu, K-means and EMGMM clustering. A naïve approach to shifting the initial means closer to the final means is to dilate the initial mask. This pushes the boundaries of the contour for the object to accept the lower intensity neighbouring pixels, as well as remove these relatively higher values from the background mask.

To determine the optimal dilation size, we compare the difference of the mean values for each image in Table 6.2, for the good segmentation results only. These values are shown in Table 6.3. We use an elliptical element for dilation. The results of the dilation and the difference from the respective final means is shown in Table 6.4, Table 6.5 and Table 6.6 for the initial masks obtained from Otsu binarization, K-means and EMGMM clustering respectively. The closest values to the average final mean are highlighted in blue. From the tables, it can be seen that a dilation size of 3 for a elliptical dilation element results in mean values that are closest to the average final means.

TABLE 6.3: Average of final means for good segmentations.

Image	\bar{c}_0	\bar{c}_1
1	0,006824	0,110693
2	0,007468	0,308217
3	0,041334	0,095952
4	0,040282	0,130360
5	0,034656	0,200287
6	0,001926	0,241672

TABLE 6.4: Dilation after Otsu binarization.

Image	Size	c_0	c_1	$ c_0 - \bar{c}_0 $	$ c_1 - \bar{c}_1 $	$\sum_i c_i - \bar{c}_i $
1	3	0.027716	0.205161	0.020892	0.094468	0.115360
1	5	0.026188	0.173038	0.019364	0.062345	0.081709
1	7	0.024682	0.151369	0.017858	0.040676	0.058534
1	9	0.023158	0.136273	0.016334	0.025580	0.041914
1	11	0.021564	0.124931	0.014739	0.014238	0.028978
1	13	0.019981	0.116536	0.013157	0.005843	0.018999
1	15	0.018341	0.109705	0.011517	0.000988	0.012505
2	3	0.007563	0.301761	9.47e-05	0.006456	0.006551
2	5	0.006025	0.286271	0.001443	0.021946	0.023389
2	7	0.005386	0.273472	0.002082	0.034745	0.036827
2	9	0.004936	0.262053	0.002532	0.046164	0.048696
2	11	0.004587	0.250695	0.002881	0.057522	0.060403
2	13	0.004277	0.240651	0.003191	0.067566	0.070757
2	15	0.004041	0.231146	0.003428	0.077071	0.080498
3	3	0.041796	0.090729	0.000462	0.005223	0.005685
3	5	0.041641	0.084544	0.000307	0.011408	0.011715
3	7	0.041535	0.080118	0.000201	0.015834	0.016034
3	9	0.042983	0.076541	0.001649	0.019411	0.021059

3	11	0.041439	0.073293	0.000105	0.022659	0.022764
3	13	0.041397	0.070676	6.27e-05	0.025276	0.025339
3	15	0.041369	0.068354	3.48e-05	0.027599	0.027633
4	3	0.041060	0.133918	0.000778	0.003558	0.004336
4	5	0.040715	0.128101	0.000433	0.002259	0.002692
4	7	0.040534	0.122898	0.000252	0.007462	0.007714
4	9	0.040429	0.118341	0.000147	0.012019	0.012166
4	11	0.040354	0.113853	7.17e-05	0.016507	0.016579
4	13	0.040297	0.109960	1.53e-05	0.020400	0.020415
4	15	0.040235	0.106414	4.70e-05	0.023946	0.023993
5	3	0.035322	0.203137	0.000666	0.002850	0.003516
5	5	0.034726	0.193711	7.02e-05	0.006576	0.006646
5	7	0.034434	0.184644	0.000222	0.015643	0.015865
5	9	0.034281	0.176477	0.000375	0.023810	0.024185
5	11	0.034188	0.168599	0.000468	0.031688	0.032156
5	13	0.034120	0.161732	0.000536	0.038555	0.039091
5	15	0.034056	0.155296	0.000600	0.044991	0.045591
6	3	0.000457	0.209125	0.001469	0.032547	0.034016
6	5	7.29e-05	0.171123	0.001853	0.070549	0.072402
6	7	1.84e-05	0.144280	0.001908	0.097392	0.099299
6	9	1.08e-05	0.125501	0.001915	0.116171	0.118086
6	11	1.15e-05	0.110706	0.001914	0.130966	0.132880
6	13	1.27e-05	0.100486	0.001913	0.141186	0.143099
6	15	1.44e-05	0.092725	0.001912	0.148947	0.150858

TABLE 6.5: Dilation after K-means clustering.

Image	Size	c_0	c_1	$ c_0 - \bar{c}_0 $	$ c_1 - \bar{c}_1 $	$\sum_i c_i - \bar{c}_i $
1	3	0.027716	0.205161	0.020892	0.094468	0.115360
1	5	0.026188	0.173038	0.019364	0.062345	0.081709
1	7	0.024682	0.151369	0.017858	0.040676	0.058534
1	9	0.023158	0.136273	0.016334	0.025580	0.041914
1	11	0.021564	0.124931	0.014739	0.014238	0.028978
1	13	0.019981	0.116536	0.013157	0.005843	0.018999
1	15	0.018341	0.109705	0.011517	0.000988	0.012505

2	3	0.007563	0.301761	9.47e-05	0.006456	0.006551
2	5	0.006025	0.286271	0.001443	0.021946	0.023389
2	7	0.005386	0.273472	0.002082	0.034745	0.036827
2	9	0.004936	0.262053	0.002532	0.046164	0.048696
2	11	0.004587	0.250695	0.002881	0.057522	0.060403
2	13	0.004277	0.240651	0.003191	0.067566	0.070757
2	15	0.004041	0.231146	0.003427	0.077071	0.080499
3	3	0.041796	0.090729	0.000462	0.005223	0.005685
3	5	0.041641	0.084544	0.000307	0.011408	0.011715
3	7	0.041535	0.080118	0.000201	0.015834	0.016034
3	9	0.041480	0.076541	0.000146	0.019411	0.019557
3	11	0.041439	0.073293	0.000105	0.022659	0.022764
3	13	0.041397	0.070676	6.27e-05	0.025276	0.025339
3	15	0.041369	0.068354	3.48e-05	0.027599	0.027633
4	3	0.041060	0.133918	0.000778	0.003558	0.004336
4	5	0.040715	0.128101	0.000433	0.002259	0.002692
4	7	0.040534	0.122898	0.000252	0.007462	0.007714
4	9	0.040429	0.118341	0.000147	0.012019	0.012166
4	11	0.040354	0.113853	7.17e-05	0.016507	0.016579
4	13	0.040297	0.109960	1.53e-05	0.020400	0.020415
4	15	0.040235	0.106414	4.70e-05	0.023946	0.023993
5	3	0.035322	0.203137	0.000666	0.002850	0.003516
5	5	0.034726	0.193711	7.02e-05	0.006576	0.006646
5	7	0.034434	0.184644	0.000222	0.015643	0.015865
5	9	0.034281	0.176477	0.000375	0.023810	0.024185
5	11	0.034188	0.168599	0.000468	0.031688	0.032156
5	13	0.034120	0.161732	0.000536	0.038555	0.039091
5	15	0.034056	0.155296	0.000600	0.044991	0.045591
6	3	0.000457	0.209125	0.001469	0.032547	0.034016
6	5	7.29e-05	0.171123	0.001853	0.070549	0.072402
6	7	1.84e-05	0.144280	0.001908	0.097392	0.099299
6	9	1.08e-05	0.125501	0.001915	0.116171	0.118086
6	11	1.15e-05	0.110706	0.001914	0.130966	0.132880
6	13	1.27e-05	0.100486	0.001913	0.141186	0.143099

6	15	1.44e-05	0.092725	0.001912	0.148947	0.150858
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TABLE 6.6: Dilation after EM-GMM clustering.

Image	Size	c_0	c_1	$ c_0 - \bar{c}_0 $	$ c_1 - \bar{c}_1 $	$\sum_i c_i - \bar{c}_i $
1	3	0.027251	0.198511	0.020427	0.087818	0.108245
1	5	0.025597	0.166707	0.018773	0.056014	0.074787
1	7	0.023968	0.146071	0.017144	0.035378	0.052522
1	9	0.022313	0.131983	0.015489	0.021290	0.036779
1	11	0.020595	0.121338	0.013771	0.010645	0.024416
1	13	0.018866	0.113441	0.012042	0.002748	0.014789
1	15	0.017084	0.107025	0.010260	0.003668	0.013928
2	3	0.006516	0.294763	0.000952	0.013454	0.014406
2	5	0.005719	0.280590	0.001748	0.027627	0.029375
2	7	0.005165	0.268079	0.002303	0.040138	0.042441
2	9	0.004774	0.256867	0.002694	0.051350	0.054044
2	11	0.004438	0.245779	0.003030	0.062438	0.065468
2	13	0.004165	0.235995	0.003303	0.072222	0.075525
2	15	0.003947	0.226736	0.003521	0.081481	0.085002
3	3	0.041729	0.089955	0.000395	0.005997	0.006392
3	5	0.041603	0.083968	0.000269	0.011984	0.012252
3	7	0.041502	0.079529	0.000168	0.016423	0.016590
3	9	0.041435	0.075914	0.000101	0.020038	0.020138
3	11	0.041384	0.072717	5.00e-05	0.023236	0.023285
3	13	0.041335	0.070134	1.30e-06	0.025819	0.025819
3	15	0.041299	0.067855	3.52e-05	0.028097	0.028132
4	3	0.040788	0.114645	0.000506	0.015715	0.016221
4	5	0.040886	0.099155	0.000604	0.031205	0.031809
4	7	0.041119	0.087943	0.000838	0.042417	0.043255
4	9	0.041369	0.080176	0.001087	0.050184	0.051271
4	11	0.041665	0.074350	0.001383	0.056010	0.057393
4	13	0.042029	0.070365	0.001747	0.059995	0.061742
4	15	0.042324	0.067586	0.002042	0.062774	0.064816
5	3	0.034368	0.147707	0.000288	0.052580	0.052868
5	5	0.034298	0.119630	0.000358	0.080657	0.081015

5	7	0.034222	0.102563	0.000434	0.097724	0.098158
5	9	0.034181	0.092259	0.000475	0.108028	0.108504
5	11	0.034143	0.085251	0.000513	0.115036	0.115549
5	13	0.034105	0.080911	0.000551	0.119377	0.119928
5	15	0.034075	0.077899	0.000581	0.122388	0.122969
6	3	0.000000	0.136655	0.001926	0.105017	0.106943
6	5	0.000000	0.116893	0.001926	0.124779	0.126705
6	7	0.000000	0.103734	0.001926	0.137938	0.139864
6	9	0.000000	0.094463	0.001926	0.147209	0.149135
6	11	0.000000	0.087325	0.001926	0.154347	0.156273
6	13	0.000000	0.082486	0.001926	0.159186	0.161112
6	15	0.000000	0.079177	0.001926	0.162495	0.164421

When defining the values for p_e and h implicitly and k_p and k_h respectively, we can find the updated equation for determining α following from Equation (6.22)

$$\begin{aligned}\alpha &= \left(\frac{c_1 - c_0}{p_e - c_0} - 1 \right)^2 \\ &= \left(\frac{c_1 - c_0 - p_e + c_0}{p_e - c_0} \right)^2 \\ &= \left(\frac{c_1 - c_0 - c_0 - k_p(c_1 - c_0) + c_0}{c_0 + k_p(c_1 - c_0) - c_0} \right)^2 \\ &= \left(\frac{c_1 - c_0 - k_p(c_1 - c_0)}{k_p(c_1 - c_0)} \right)^2 \\ &= \left(\frac{(1 - k_p)(c_1 - c_0)}{k_p(c_1 - c_0)} \right)^2\end{aligned}$$

Therefore, given k_p , the equation to calculate α is

$$\alpha = \left(\frac{1 - k_p}{k_p} \right)^2 \quad (6.28)$$

$$\alpha - 1 = \frac{1 - 2k_p + k_p^2 - k_p^2}{k_p^2} = \frac{1 - 2k_p}{k_p^2} \quad (6.29)$$

Following from Equation (6.25)

$$h = c_0 + k_h(c_1 - c_0) = c_1 - (1 - k_h)(c_1 - c_0)$$

$$\begin{aligned}
& \alpha(c_0 + k_h(c_1 - c_0) - c_0)^2 - (c_1 - (1 - k_h)(c_1 - c_0) - c_1)^2 \\
&= \alpha k_h^2(c_1 - c_0)^2 + (k_h - 1)^2(c_1 - c_0)^2 \\
&= \alpha k_h^2(c_1 - c_0)^2 - (k_h - 1)^2(c_1 - c_0)^2 \\
&= (c_1 - c_0)^2(\alpha k_h^2 - (k_h - 1)^2) \\
&= (c_1 - c_0)^2(\alpha k_h^2 - k_h^2 + 2k_h - 1) \\
&= (c_1 - c_0)^2(k_h^2(\alpha - 1) + 2k_h - 1) \\
&= (c_1 - c_0)^2\left(\left(\frac{1-2k_p}{k_p^2}\right)k_h^2 + 2k_h - 1\right) \\
\lambda_1 &= \frac{\mu(2\sqrt{2} + 4)}{(c_1 - c_0)^2\left(\left(\frac{1-2k_p}{k_p^2}\right)k_h^2 + 2k_h - 1\right)}
\end{aligned} \tag{6.30}$$

6.3 Experimental Results

The parameters α and λ_1 , in the proposed parameter estimation method, depend greatly on predicting good final means, c_0 and c_1 , as well as p_e and h for the ideal segmentation. For determining α we find the average of all k_p in Table 6.2 for all good segmentations. This is calculated to be

$$k_p = 0.154494.$$

From this we can calculate the value for α immediately using Equation (6.28). This turns out to be

$$\alpha = 29.9509. \tag{6.31}$$

Similarly, to determine h we find the average of all k_h in Table 6.2 for all good segmentations. This is calculated to be

$$k_h = 0.412737.$$

It is rarely the case where the final means are known or can be pre-calculated, if ever at all. Instead, we use the results from three popular unsupervised clustering algorithms and compare the final segmented result against a ground truth. The algorithms chosen to generate an initial mask, from which we calculate c_0 and c_1 are: *Otsu* [206], *k*-means [242] with $k = 2$ and Expectation Maximisation Gaussian Mixture Modelling (EMGMM) [243] with $k = 2$. From observations we see that these algorithms tend to generate over-segmented results. In light of this, we generate a fourth mask which is an EMGMM output with dilation. For dilation, we used an elliptical shape with a radius of $3px$. Once we have the initial mask and have calculated the initial c_0 and c_1 from this initial mask, we calculate λ_1 using Equation (6.30). Now that we have α and λ_1 we calculate λ_0 using Equation (6.16). We set the termination criterion at $\epsilon = 1 \times 10^{-3}$.

We also compare our results to two previously published parameter settings. The first is the parameter settings used El Zehiry *et al.* [239], which is where this technique was first published. Their results showed excellent segmentation output on synthetic images and mammography images with very high robustness against noise. They do not specify the noise type. The second parameter setting which we test against is presented by Masaka *et al.* [237]. Their parameter setting was based

on a time-lapse series of fluorescence images. Their scheme is a hybrid of algorithms designed to segment whole fluorescent cells; however, we use the parameter setting they have presented for segmentation only. Their parameter setting was obtained by minimising the Jaccard coefficient over the time-lapse series. Their results show a greater area of cell detection and smoother boundaries; although, the smoother contours are a result of CED-ORI [235] which is part of the scheme before segmentation.

The initial masks used in subsequent parameter estimation is shown in Figures 6.21, 6.23, 6.25, 6.27, 6.29, 6.31, 6.33, 6.35, 6.37, 6.39, 6.41, 6.43, 6.45, 6.47, 6.49, 6.51, 6.53, 6.55, 6.57, 6.59, 6.61, 6.63, 6.65, 6.67 and 6.69.

The segmented outputs are shown in Figures 6.22, 6.24, 6.26, 6.28, 6.30, 6.32, 6.34, 6.36, 6.38, 6.40, 6.42, 6.44, 6.46, 6.48, 6.50, 6.52, 6.54, 6.56, 6.58, 6.60, 6.62, 6.64, 6.66, 6.68 and 6.70.

The parameter settings and corresponding results are summarised in Table 6.7. The last column, labelled “*Diff*”, is a measure of how far off the final means are from the ideal. It is calculated as

$$Diff = |c_0^{Final} - c_0^{Ideal}| + |c_1^{Final} - c_1^{Ideal}|.$$

In Table 6.7 differentiate between methods on the same image as follows:

[imageno]-[method],

where *imageno* goes from 1 to 25 and *method* is defined as follows:

- n** - using parameter setting presented in [239].
- m** - using parameter setting presented in [237].
- o** - Proposed method with parameters estimated from an initial Otsu segmentation.
- k** - Proposed method with parameters estimated from an initial K-means segmentation with $k = 2$.
- e** - Proposed method with parameters estimated from an initial EMGMM segmentation with $k = 2$.
- d** - Proposed method with parameters estimated from an initial EMGMM segmentation with $k = 2$ and an elliptical dilation of $3px$.

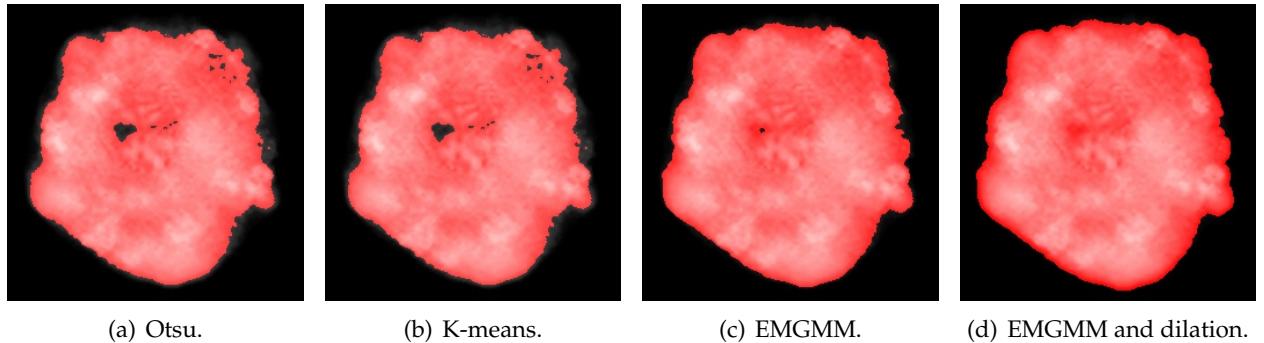


FIGURE 6.21: Image 1 from test set Appendix B initial masks.

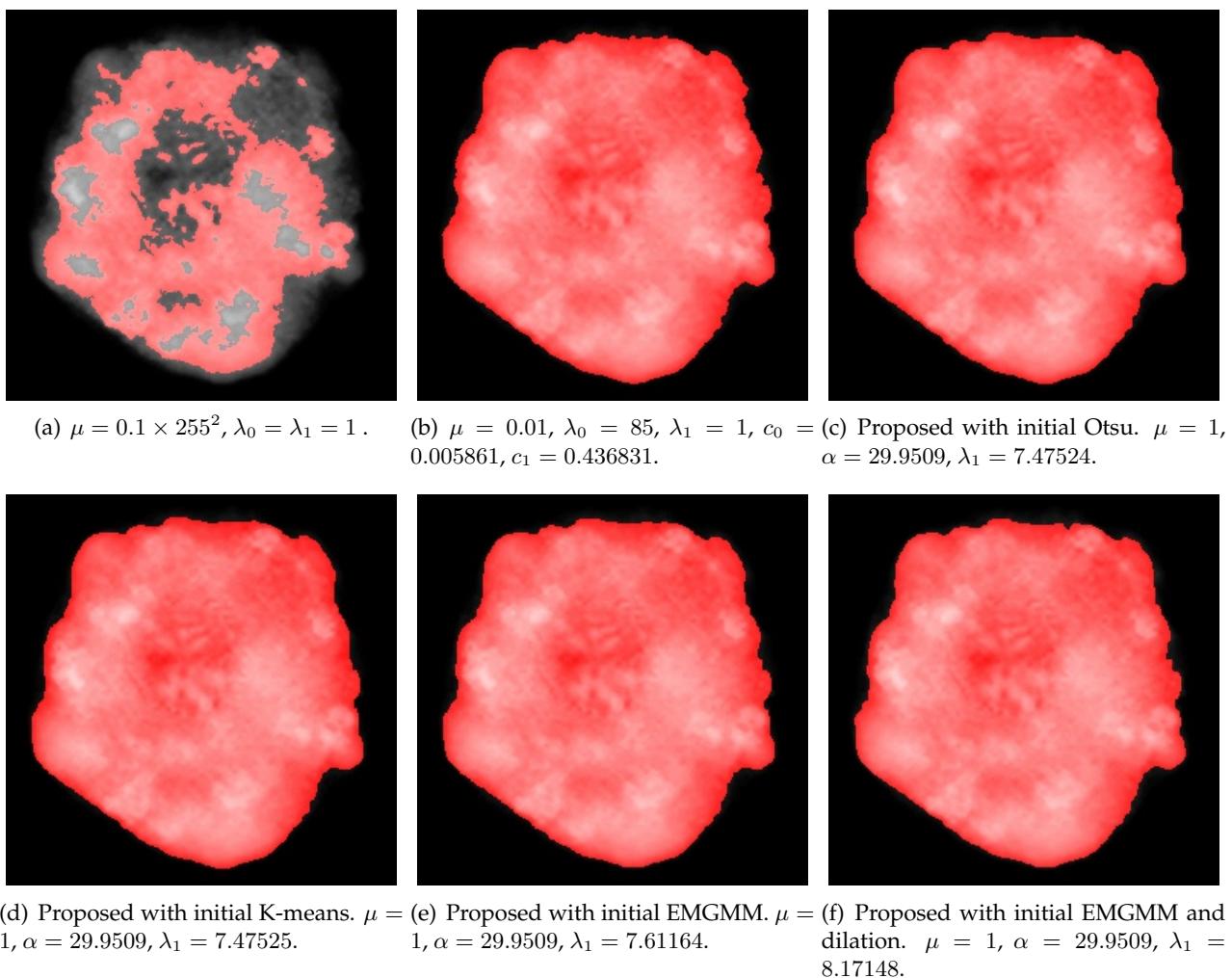


FIGURE 6.22: Image 1 from test set Appendix B segmentation results.

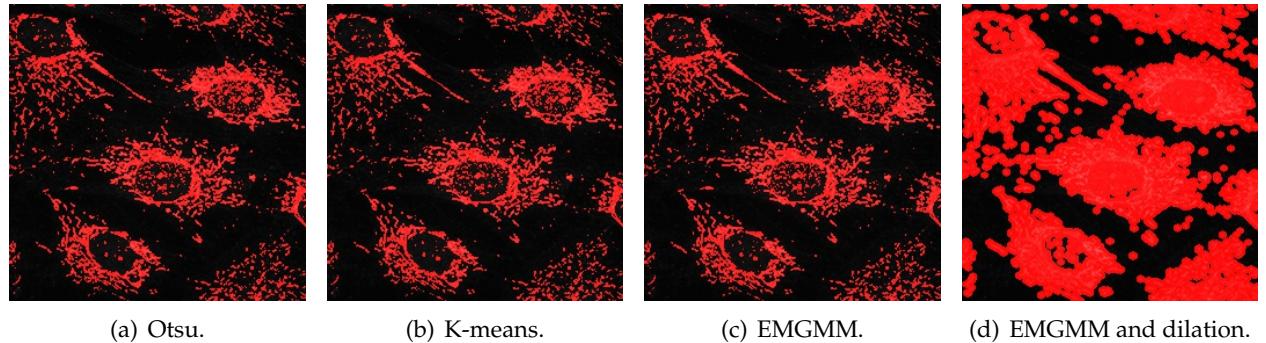


FIGURE 6.23: Image 2 from test set Appendix B initial masks.

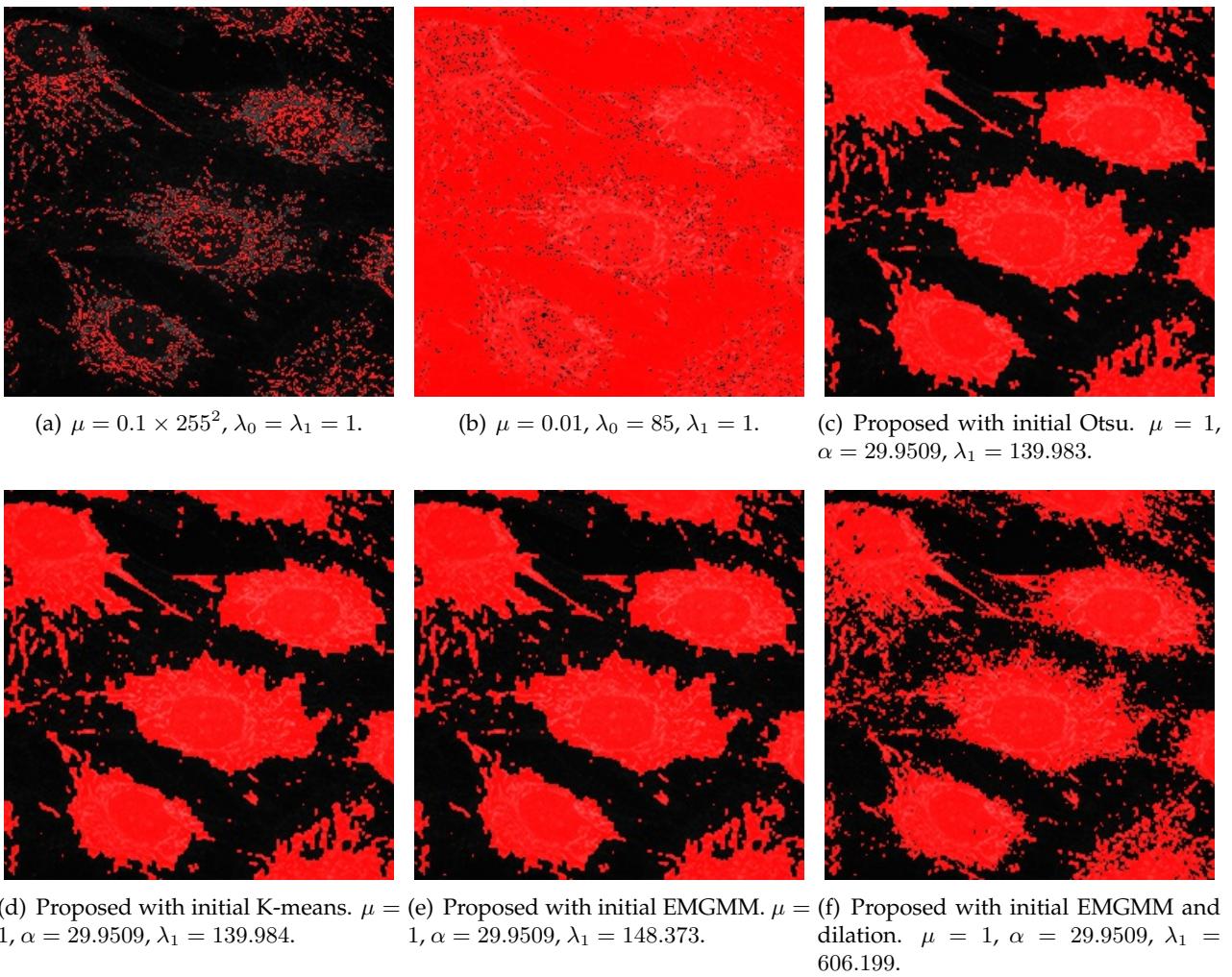


FIGURE 6.24: Image 2 from test set Appendix B segmentation results.

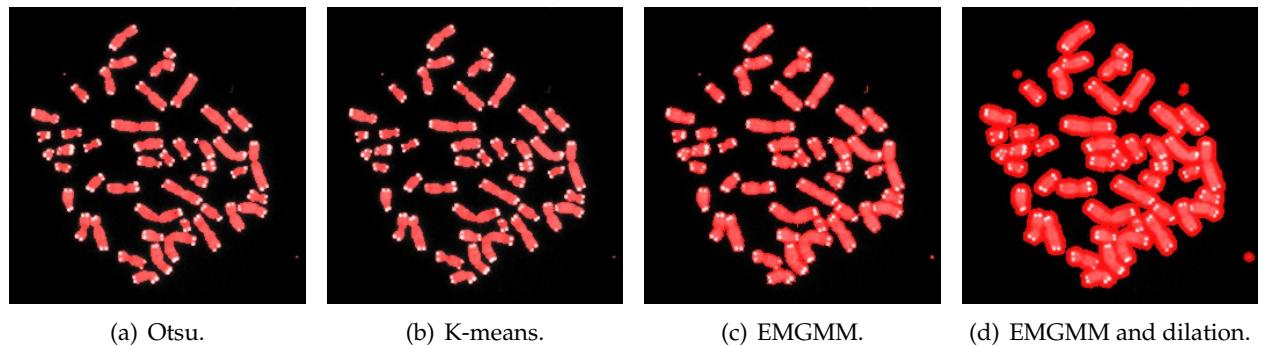


FIGURE 6.25: Image 3 from test set Appendix B initial masks.

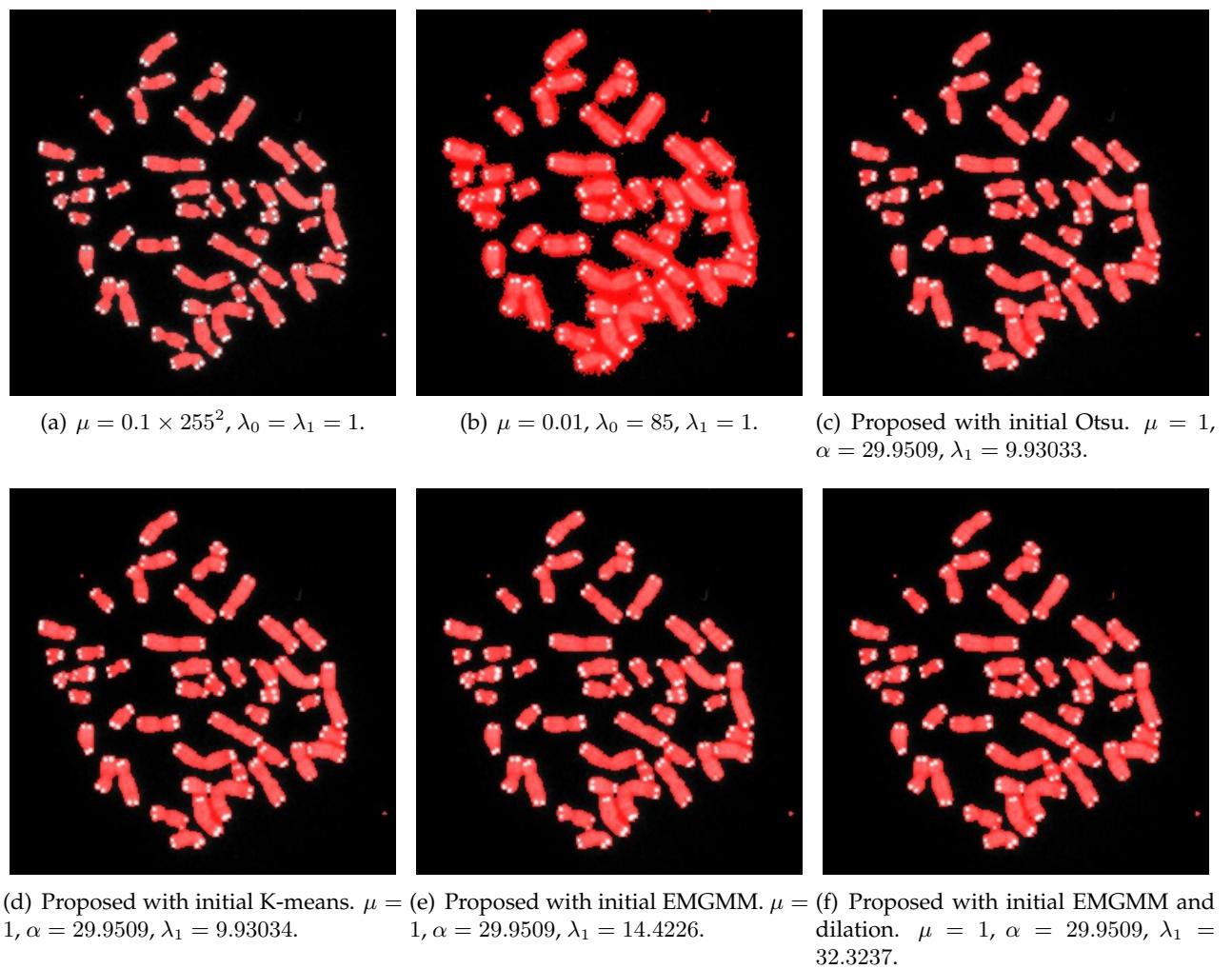


FIGURE 6.26: Image 3 from test set Appendix B segmentation results.

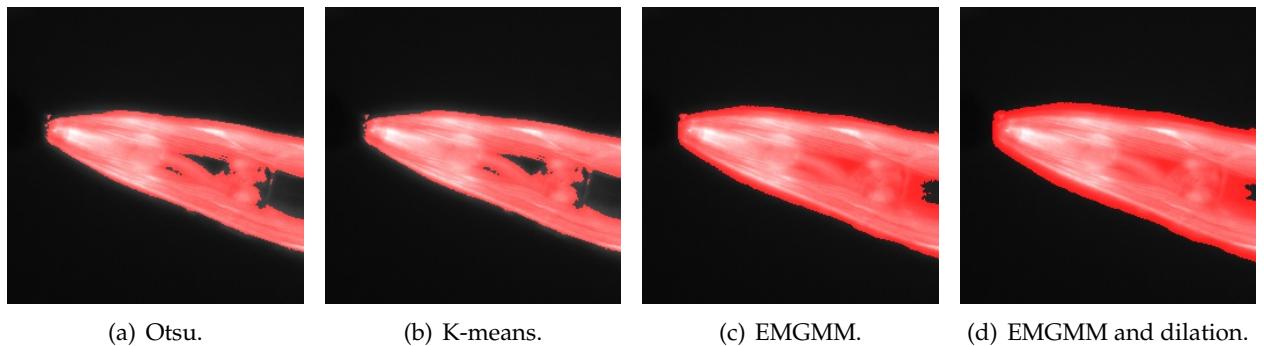


FIGURE 6.27: Image 4 from test set Appendix B initial masks.

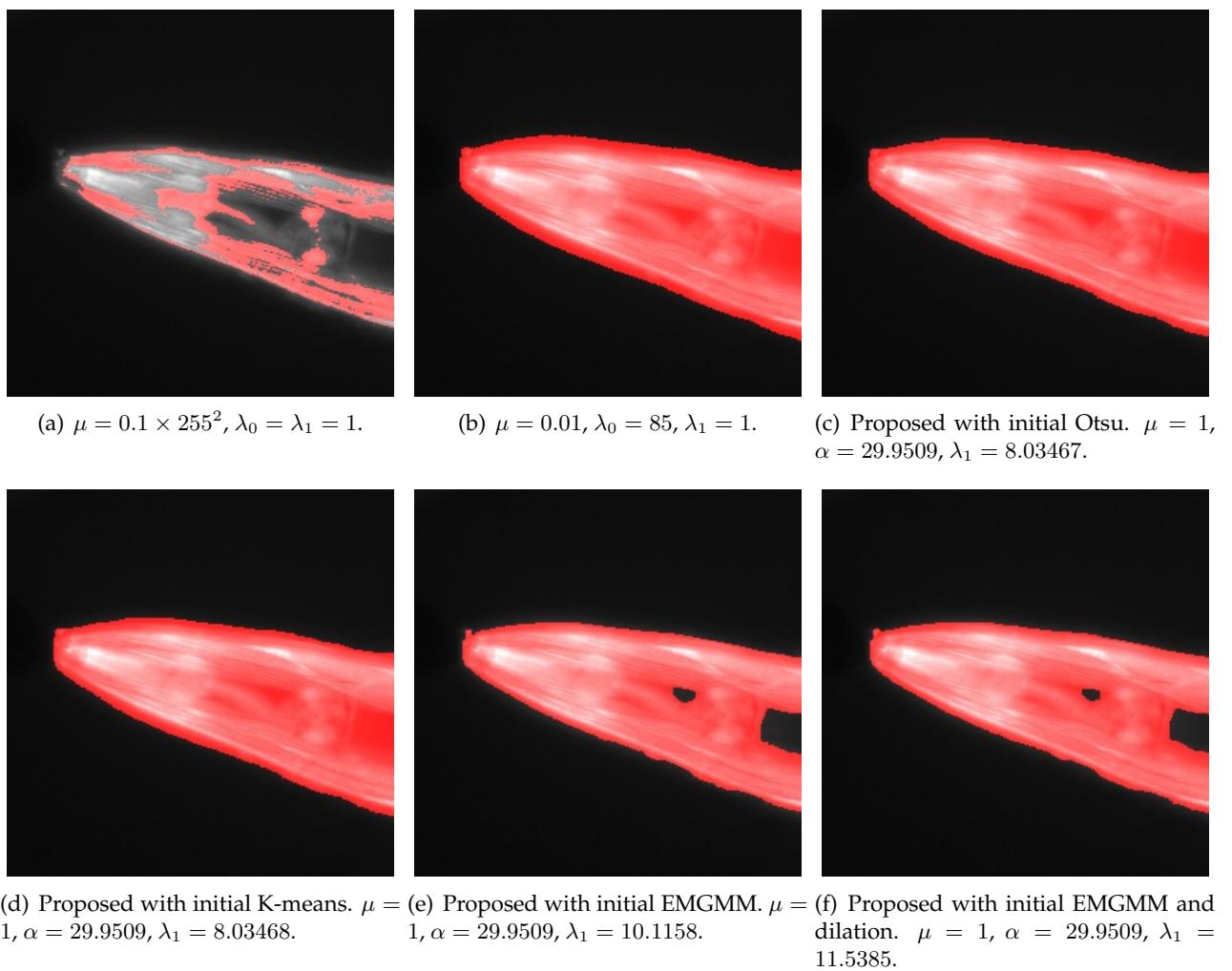


FIGURE 6.28: Image 4 from test set Appendix B segmentation results.

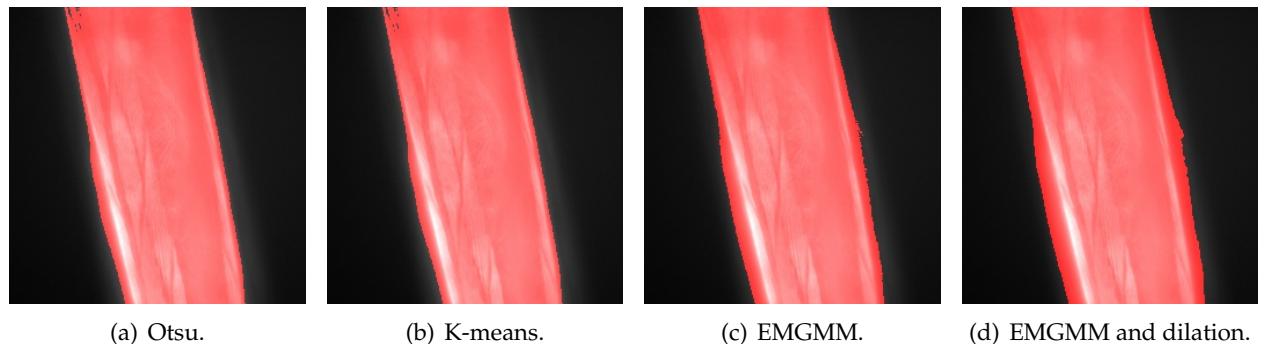


FIGURE 6.29: Image 5 from test set Appendix B initial masks.

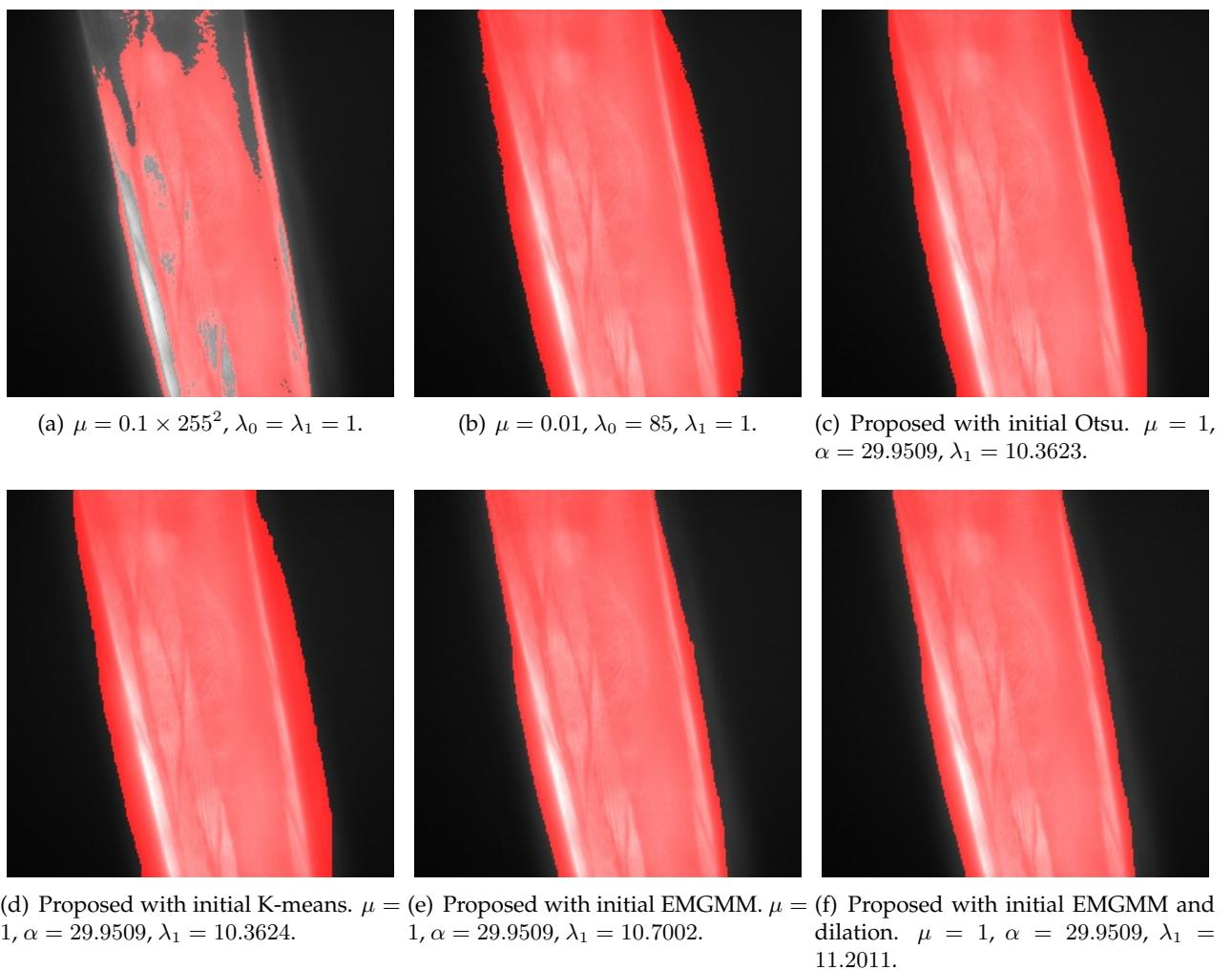


FIGURE 6.30: Image 5 from test set Appendix B segmentation results.

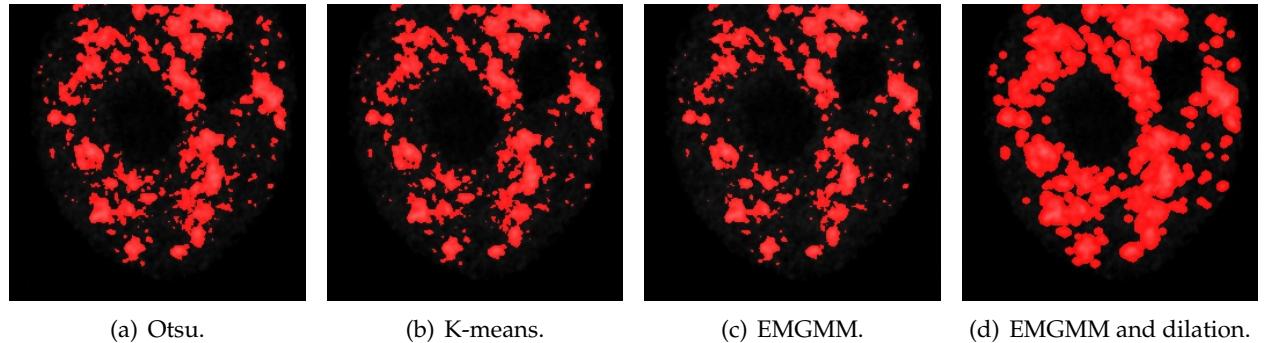


FIGURE 6.31: Image 6 from test set Appendix B initial masks.

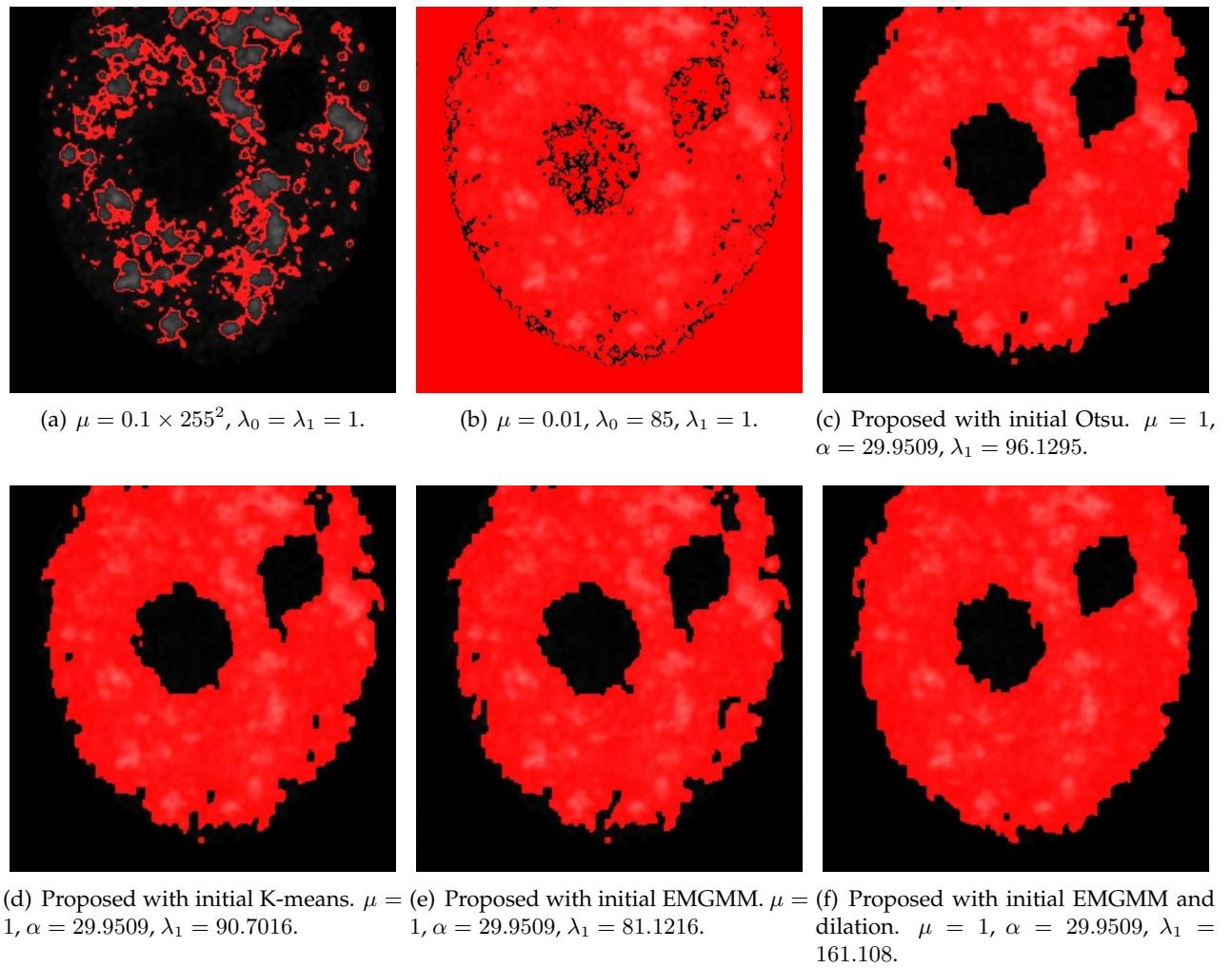


FIGURE 6.32: Image 6 from test set Appendix B segmentation results.

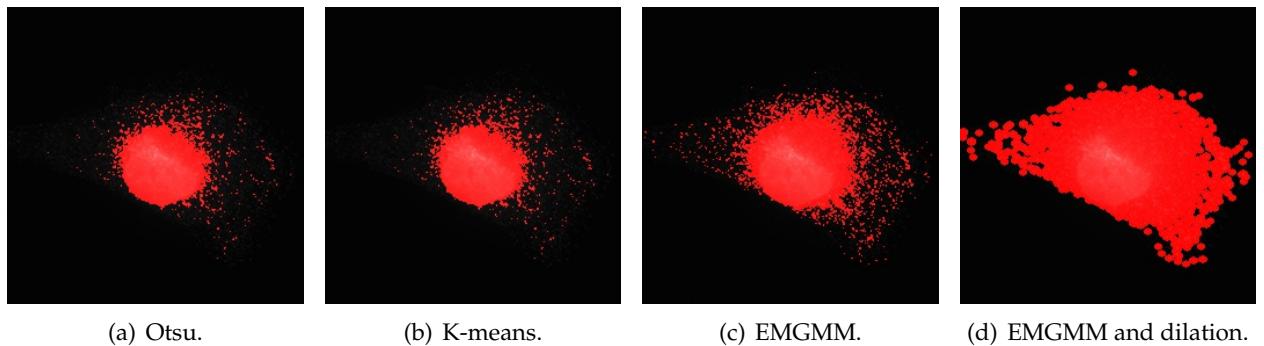


FIGURE 6.33: Image 7 from test set Appendix B initial masks.

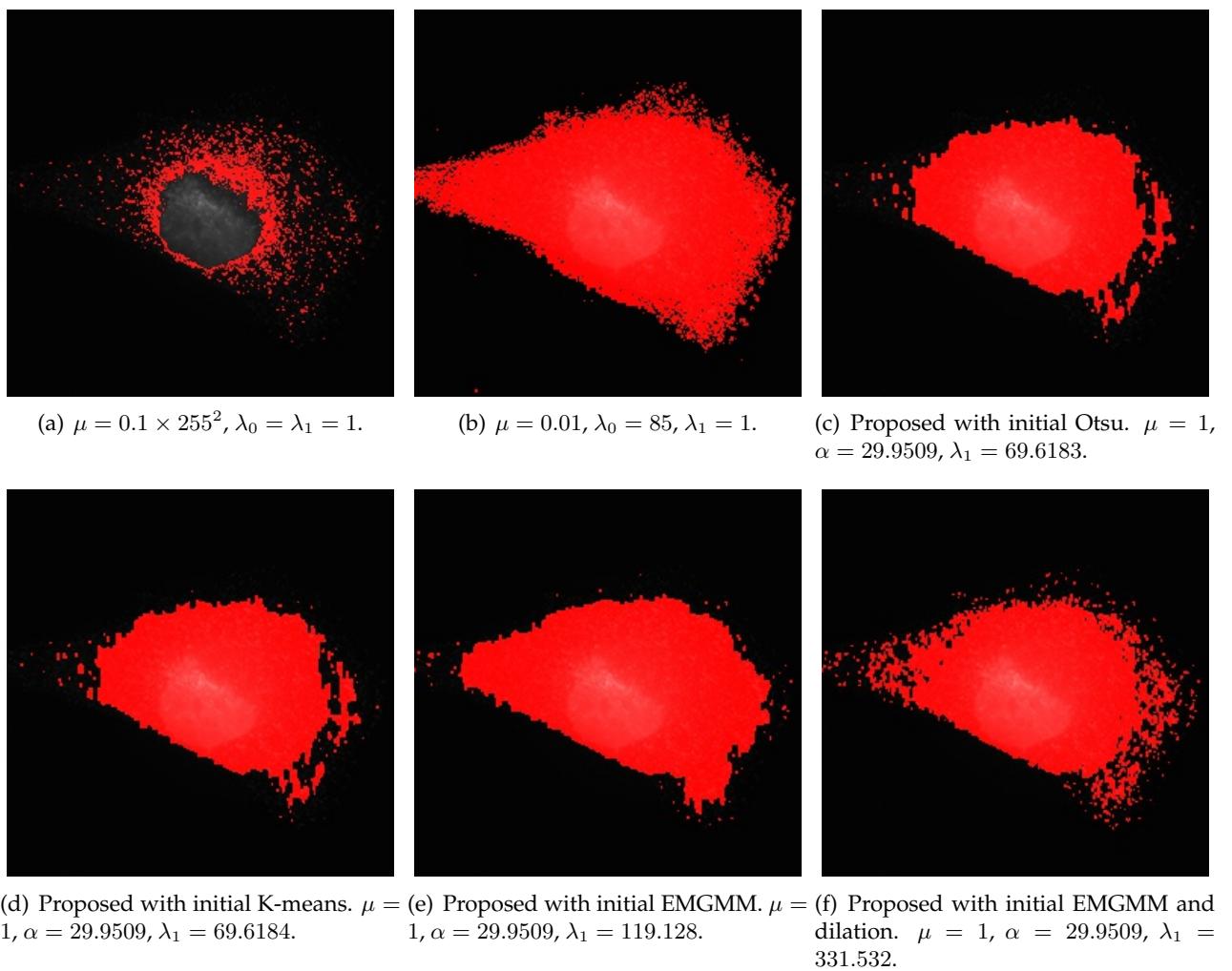


FIGURE 6.34: Image 7 from test set Appendix B segmentation results.

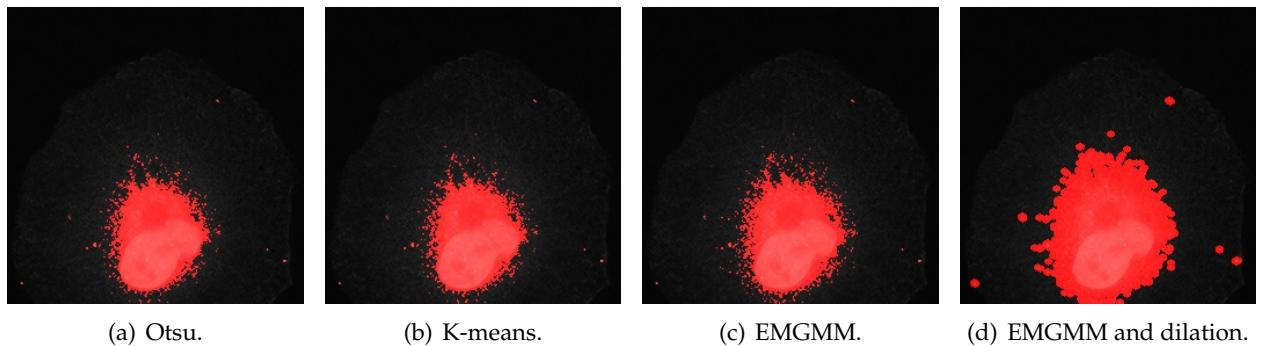


FIGURE 6.35: Image 8 from test set Appendix B initial masks.

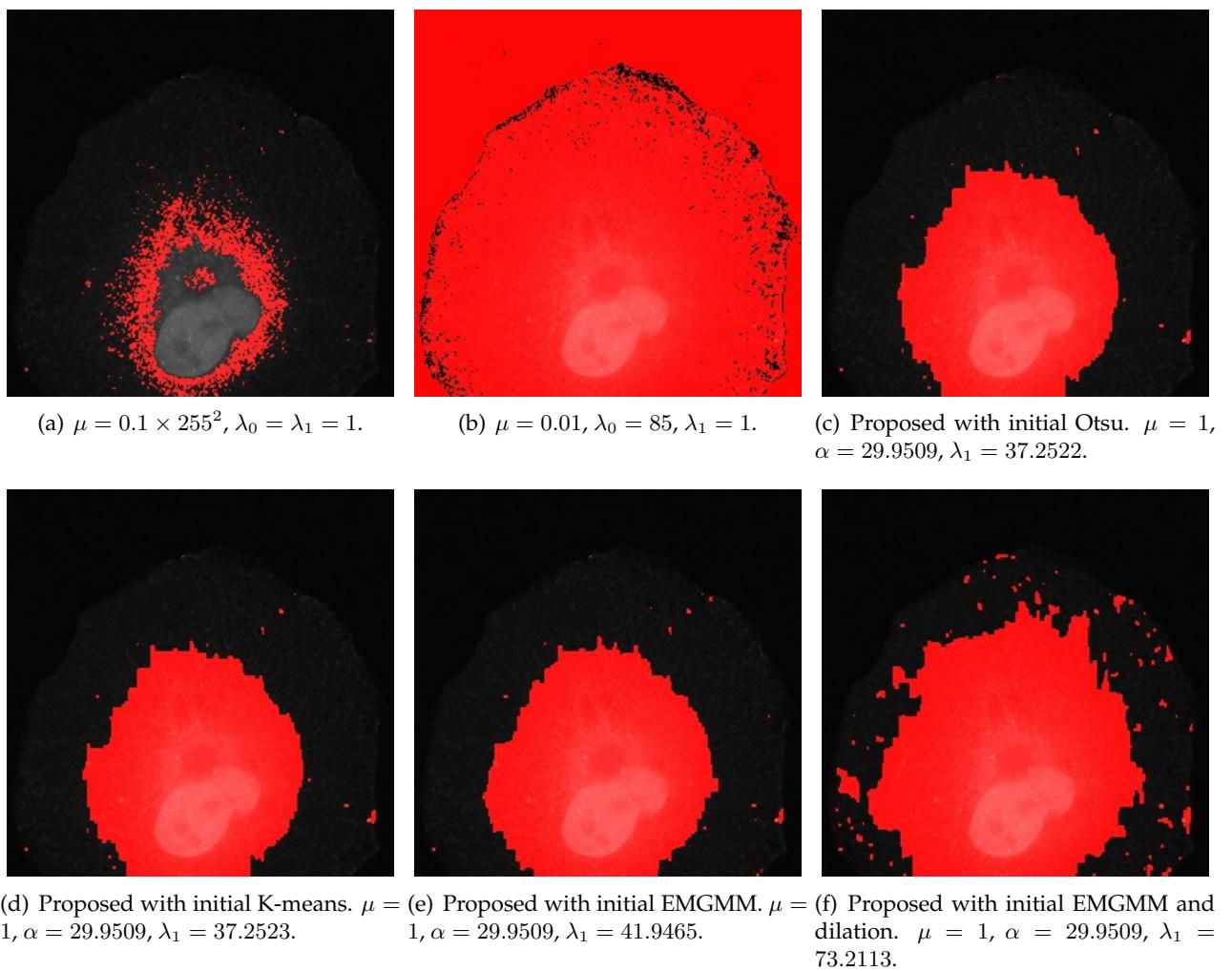


FIGURE 6.36: Image 8 from test set Appendix B segmentation results.

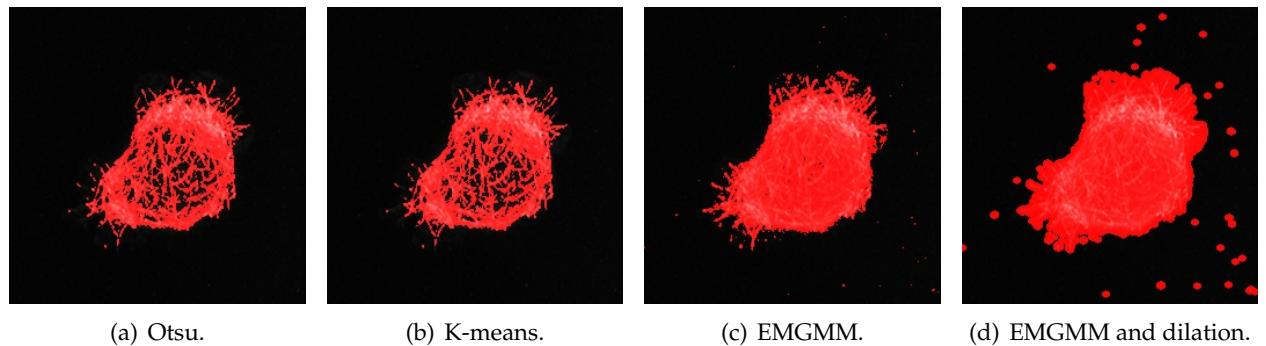


FIGURE 6.37: Image 9 from test set Appendix B initial masks.

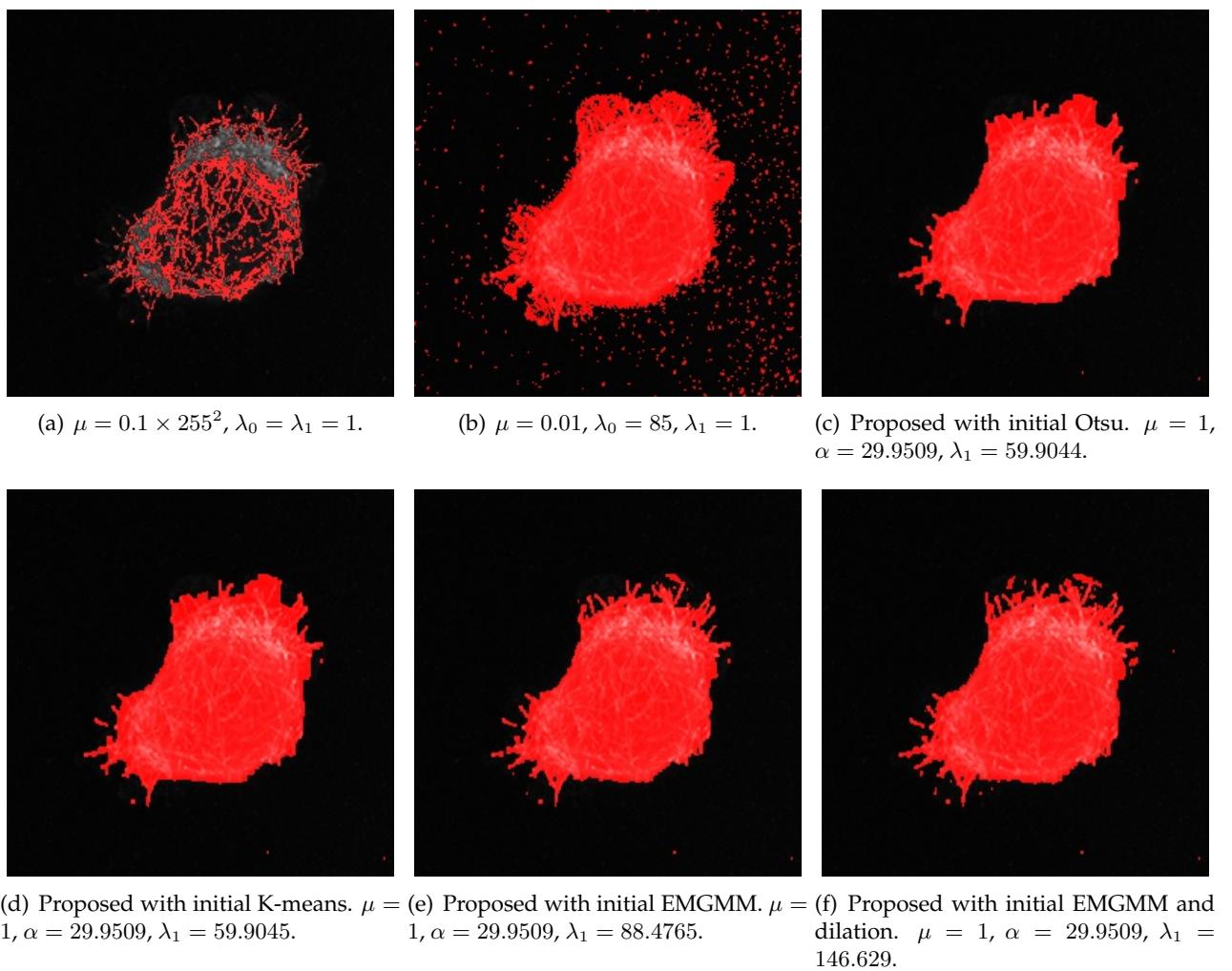


FIGURE 6.38: Image 9 from test set Appendix B segmentation results.

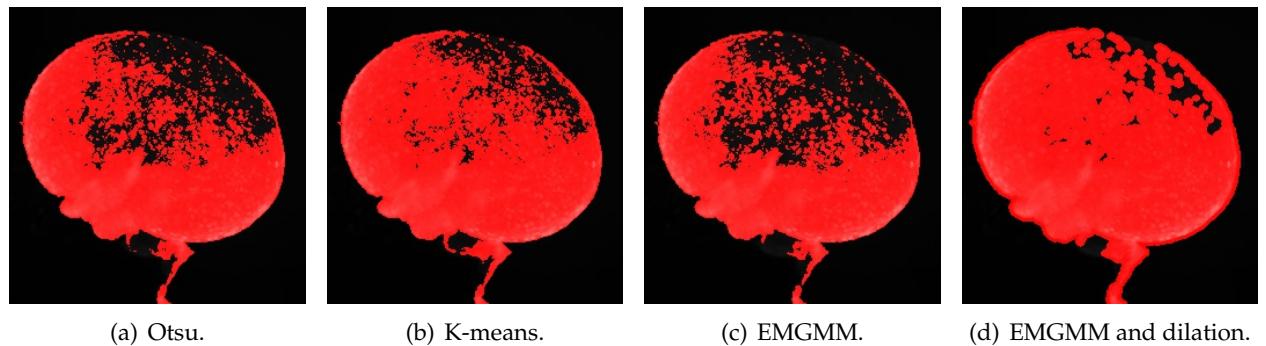


FIGURE 6.39: Image 10 from test set Appendix B initial masks.

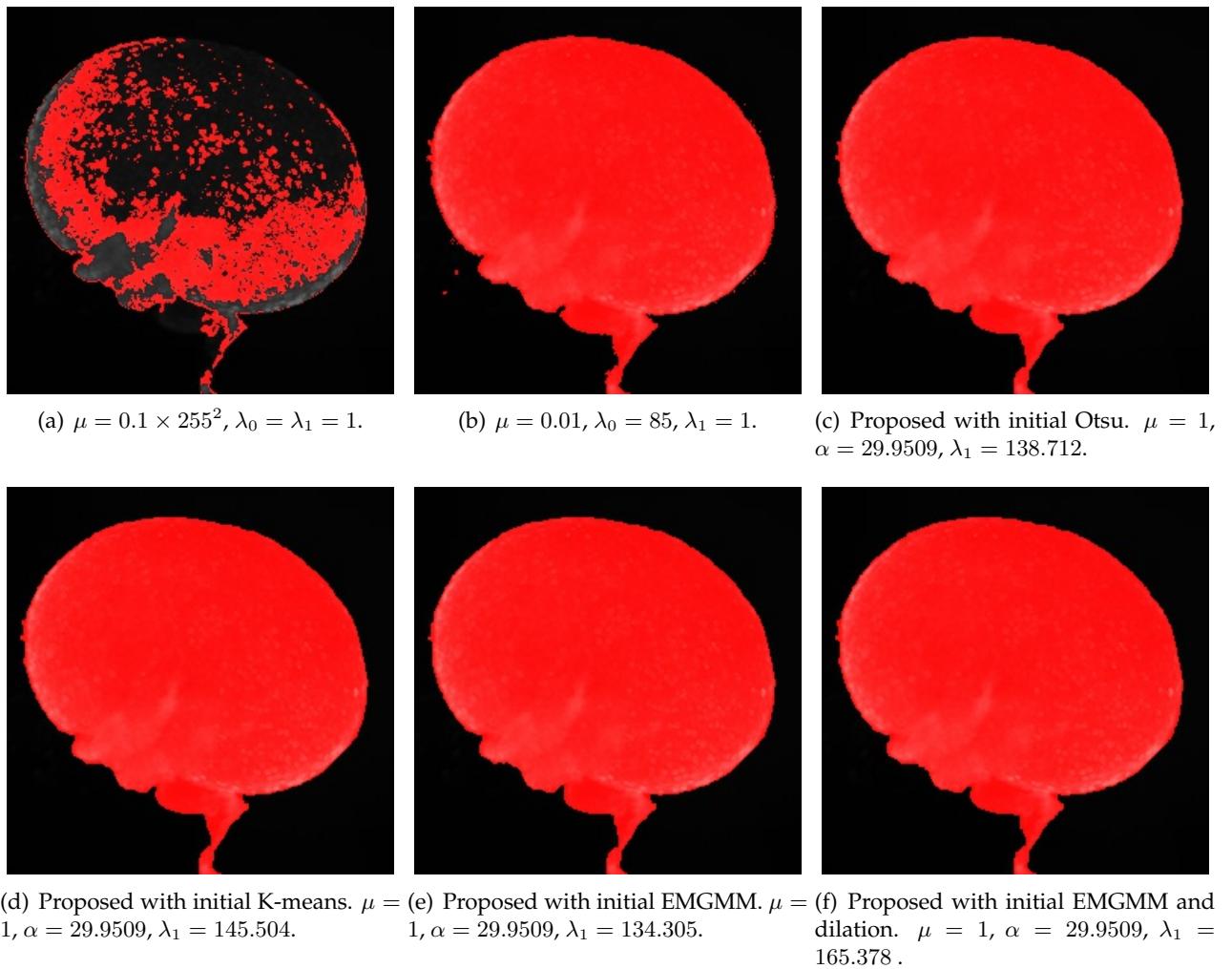


FIGURE 6.40: Image 10 from test set Appendix B segmentation results.

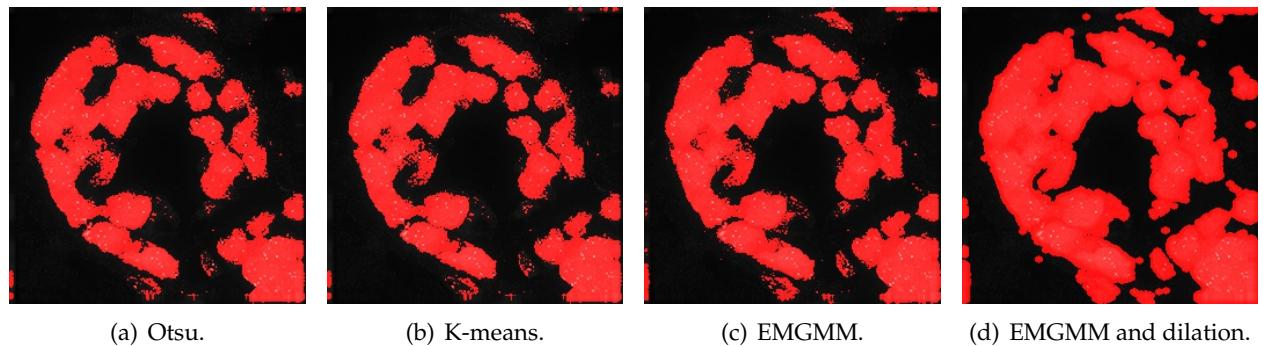


FIGURE 6.41: Image 11 from test set Appendix B initial masks.

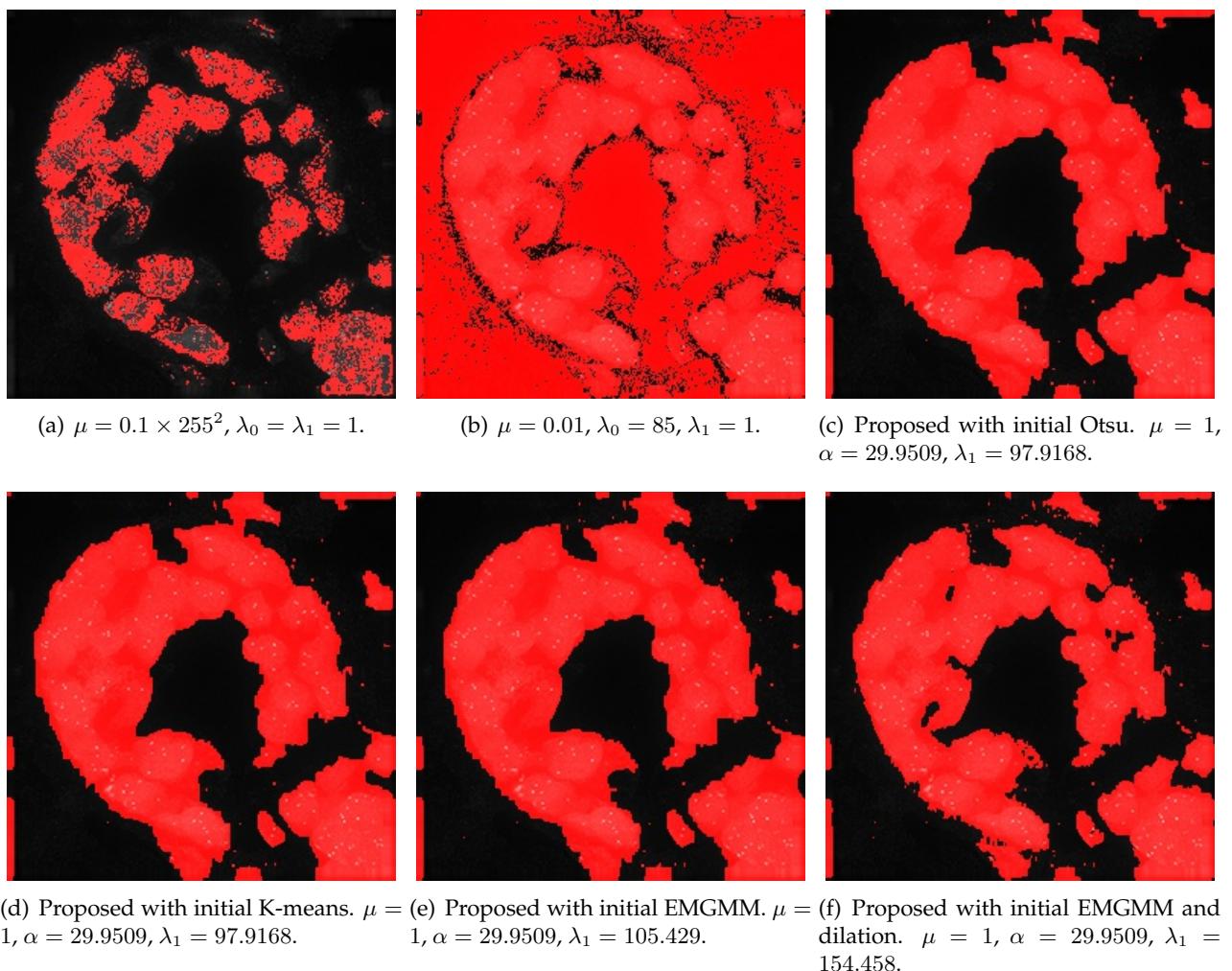


FIGURE 6.42: Image 11 from test set Appendix B segmentation results.

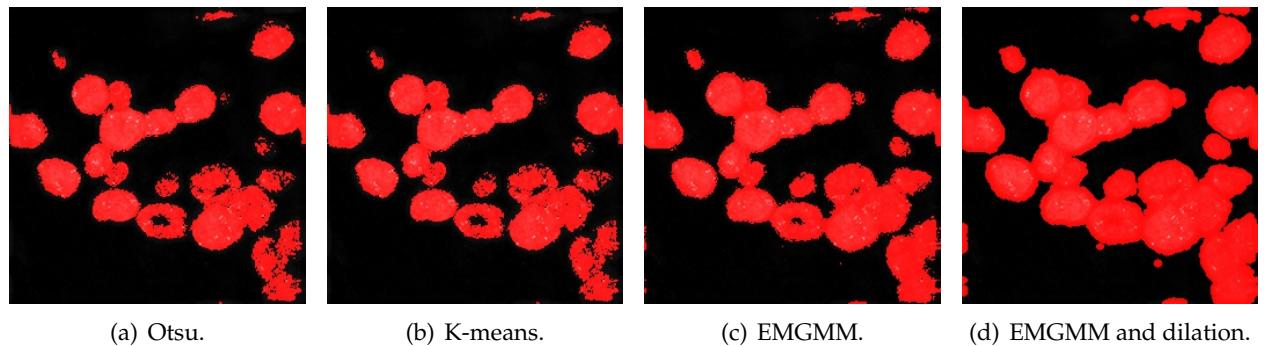


FIGURE 6.43: Image 12 from test set Appendix B initial masks.

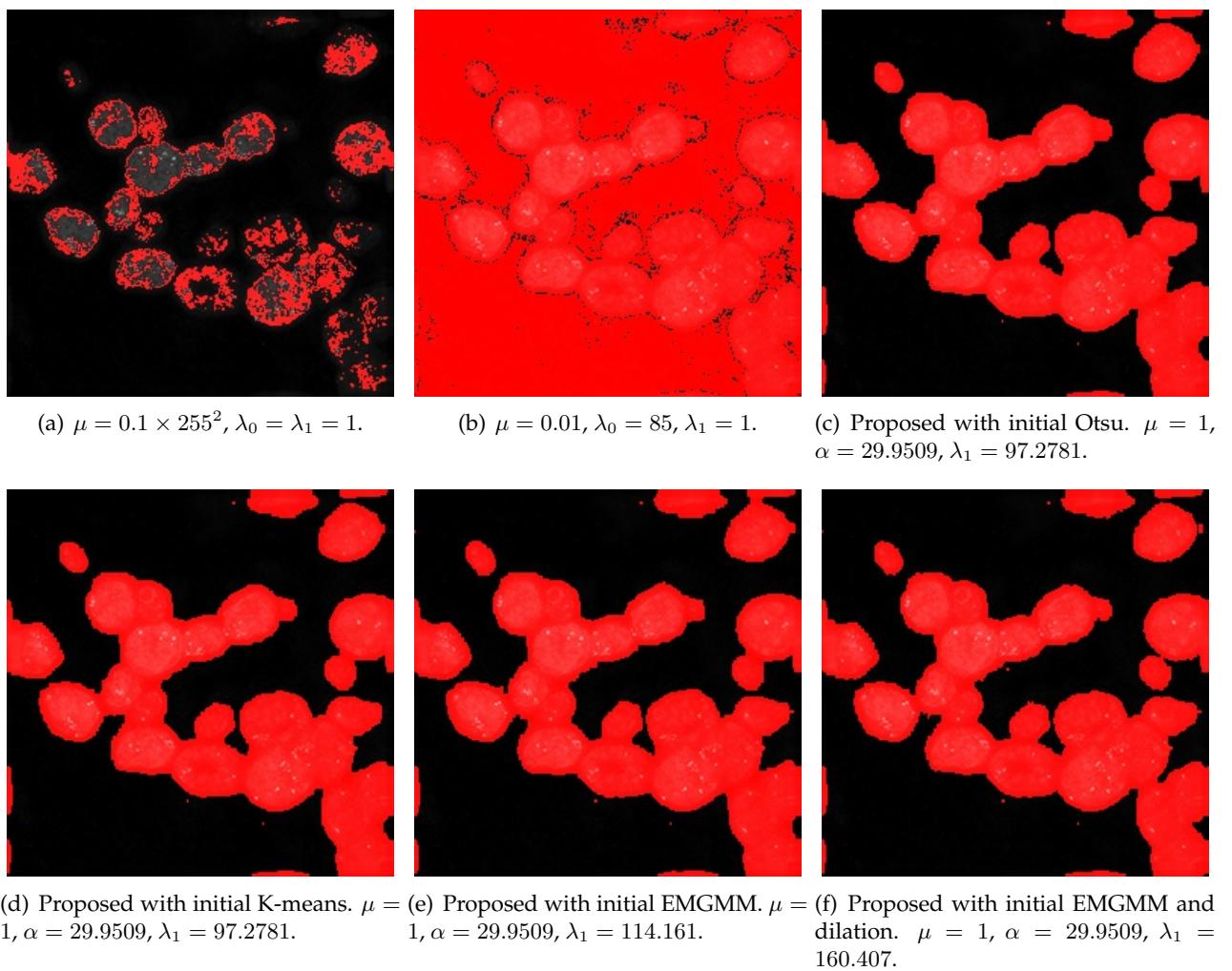


FIGURE 6.44: Image 12 from test set Appendix B segmentation results.

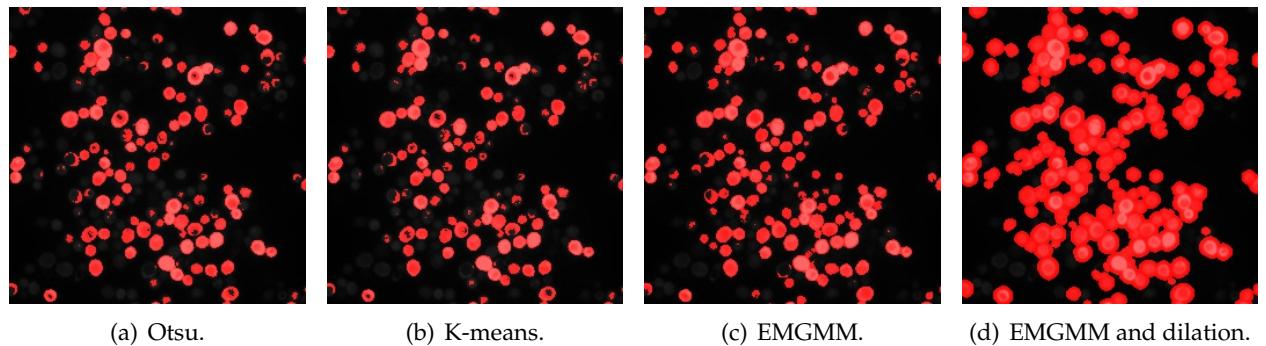


FIGURE 6.45: Image 13 from test set Appendix B initial masks.

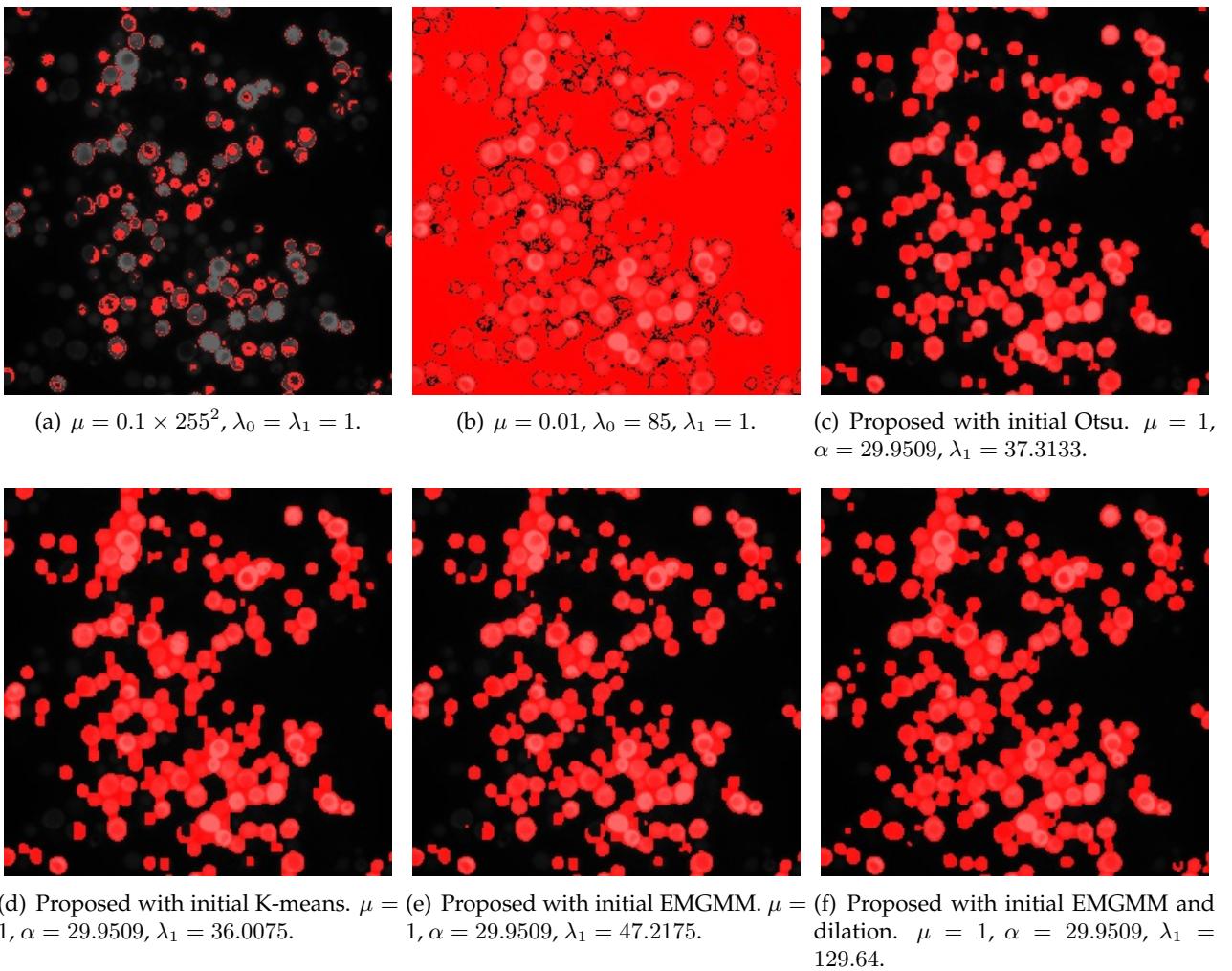


FIGURE 6.46: Image 13 from test set Appendix B segmentation results.

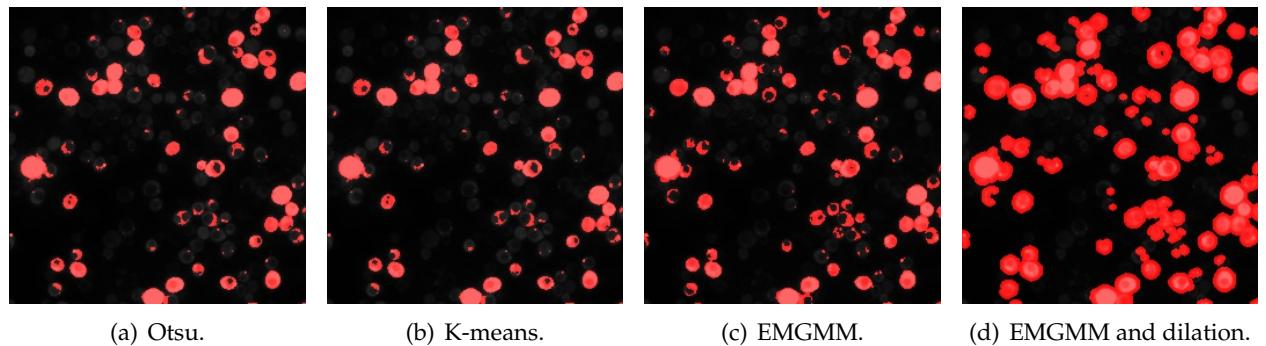


FIGURE 6.47: Image 14 from test set Appendix B initial masks.

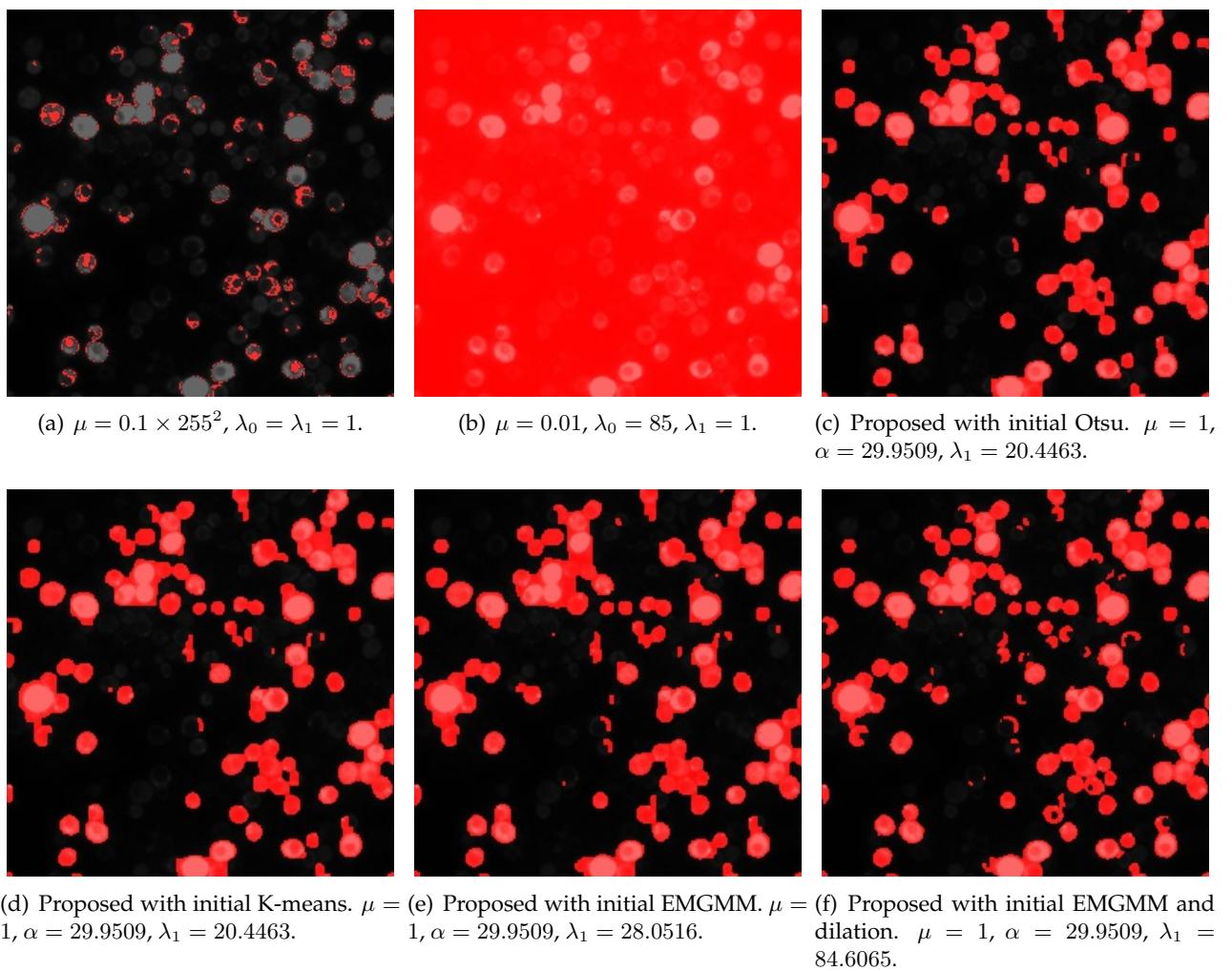


FIGURE 6.48: Image 14 from test set Appendix B segmentation results.

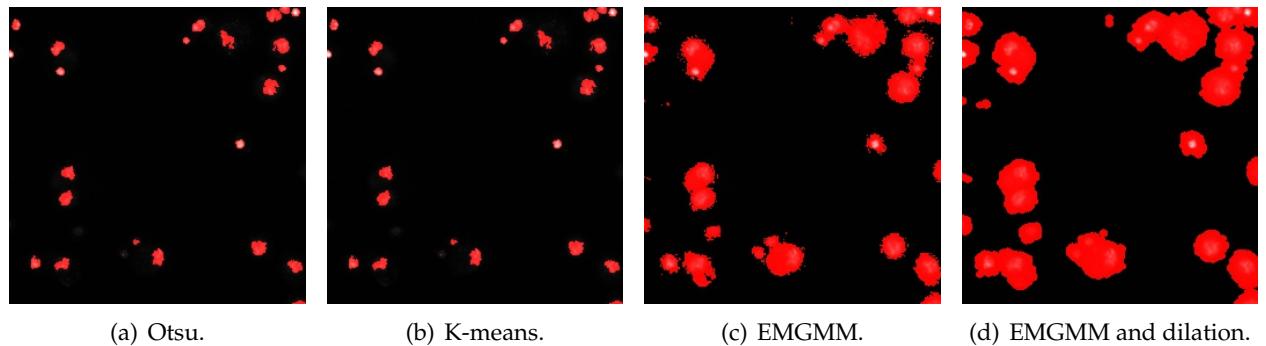


FIGURE 6.49: Image 15 from test set Appendix B initial masks.

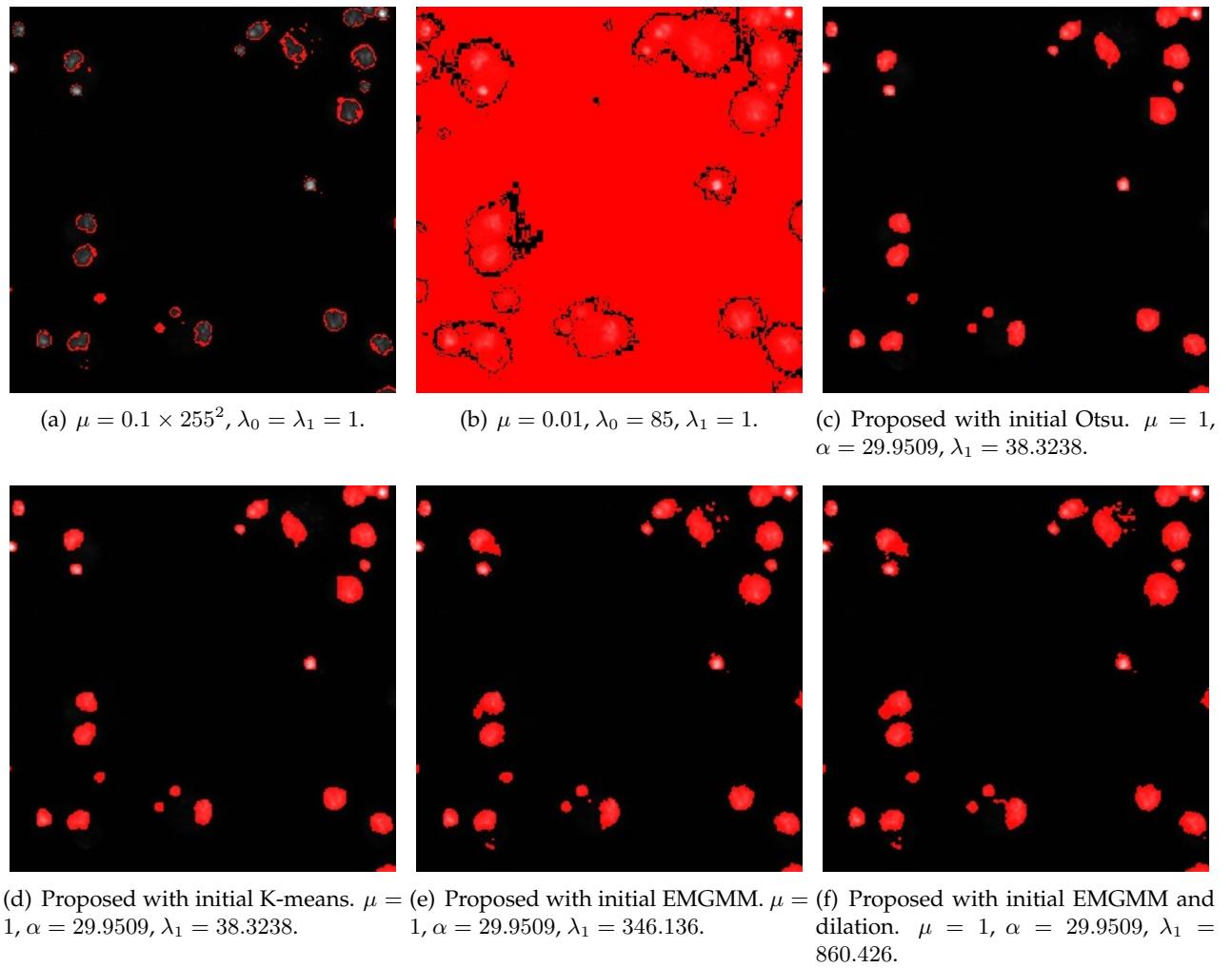


FIGURE 6.50: Image 15 from test set Appendix B segmentation results.

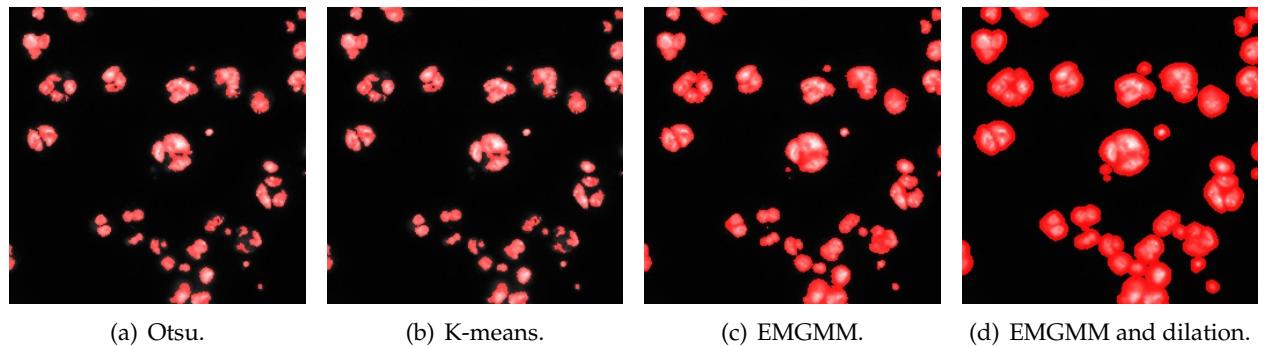


FIGURE 6.51: Image 16 from test set Appendix B initial masks.

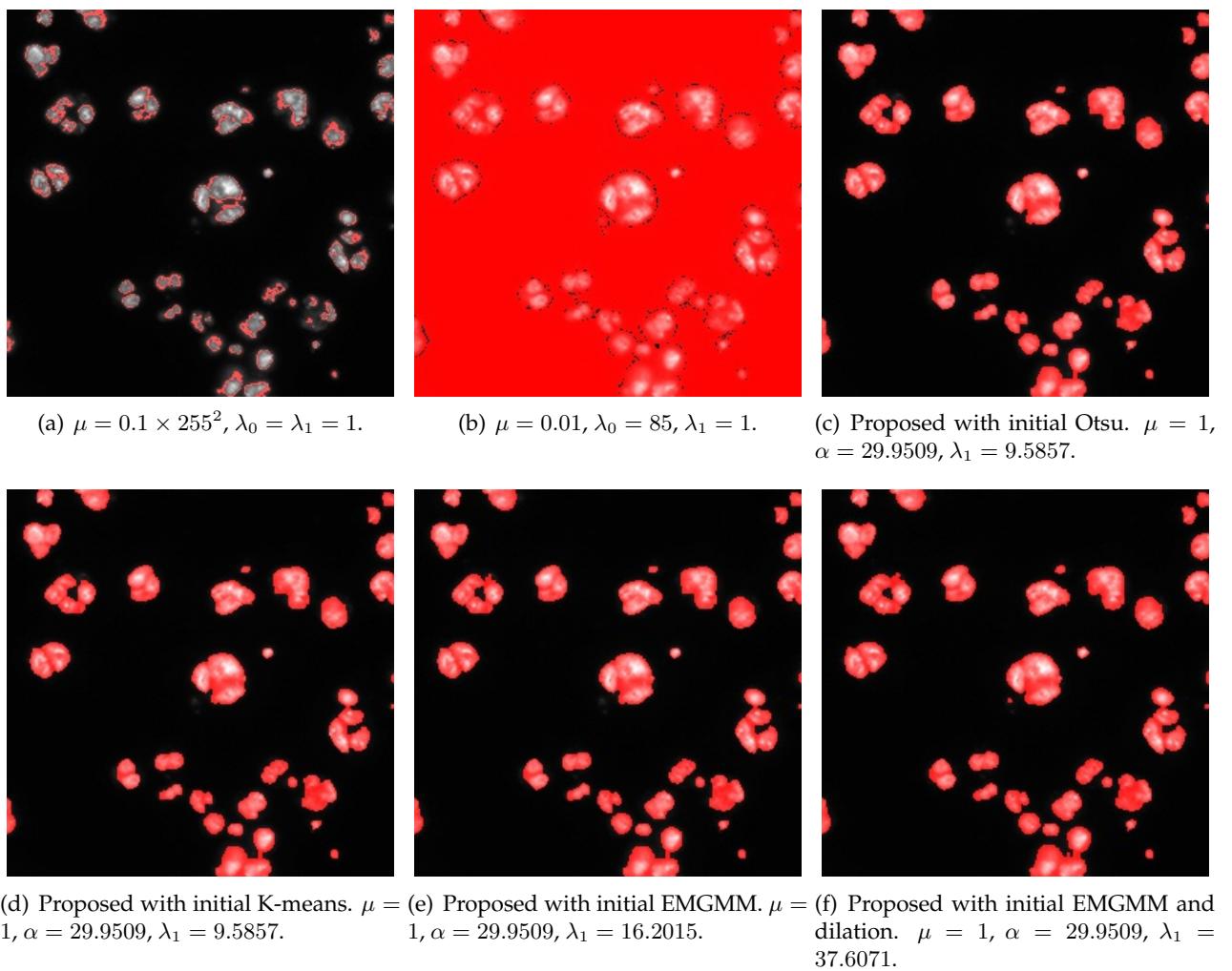


FIGURE 6.52: Image 16 from test set Appendix B segmentation results.

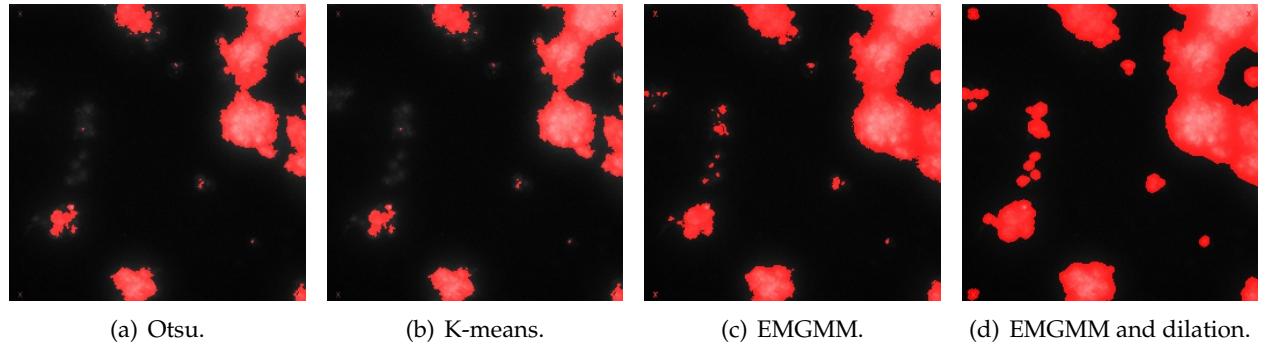


FIGURE 6.53: Image 17 from test set Appendix B initial masks.

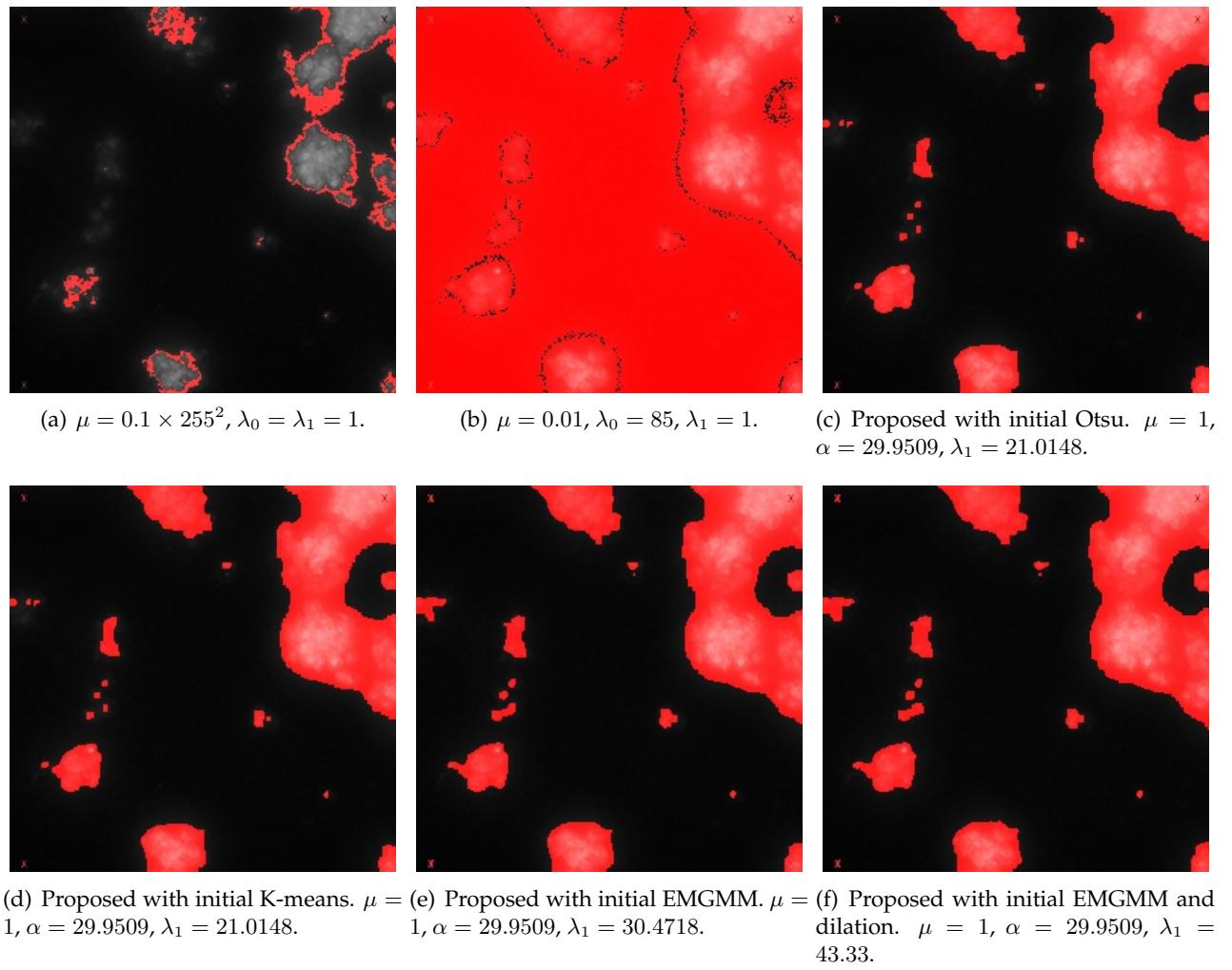


FIGURE 6.54: Image 17 from test set Appendix B segmentation results.

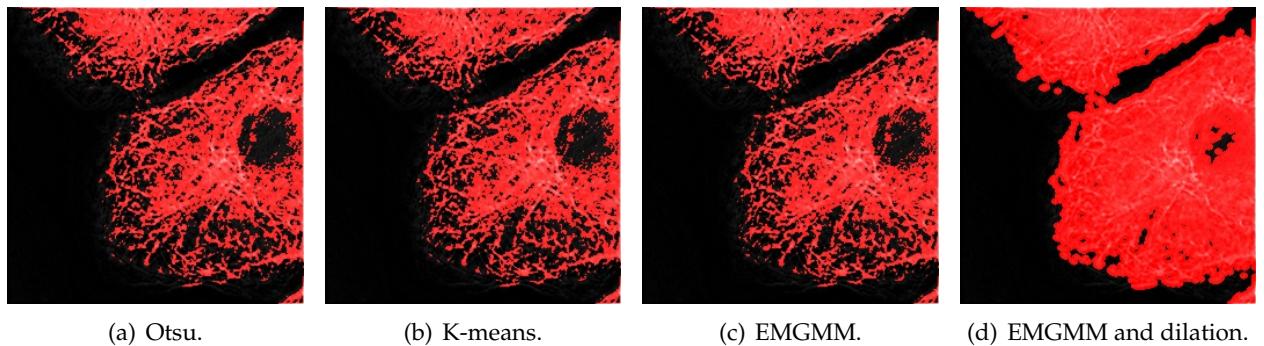


FIGURE 6.55: Image 18 from test set Appendix B initial masks.

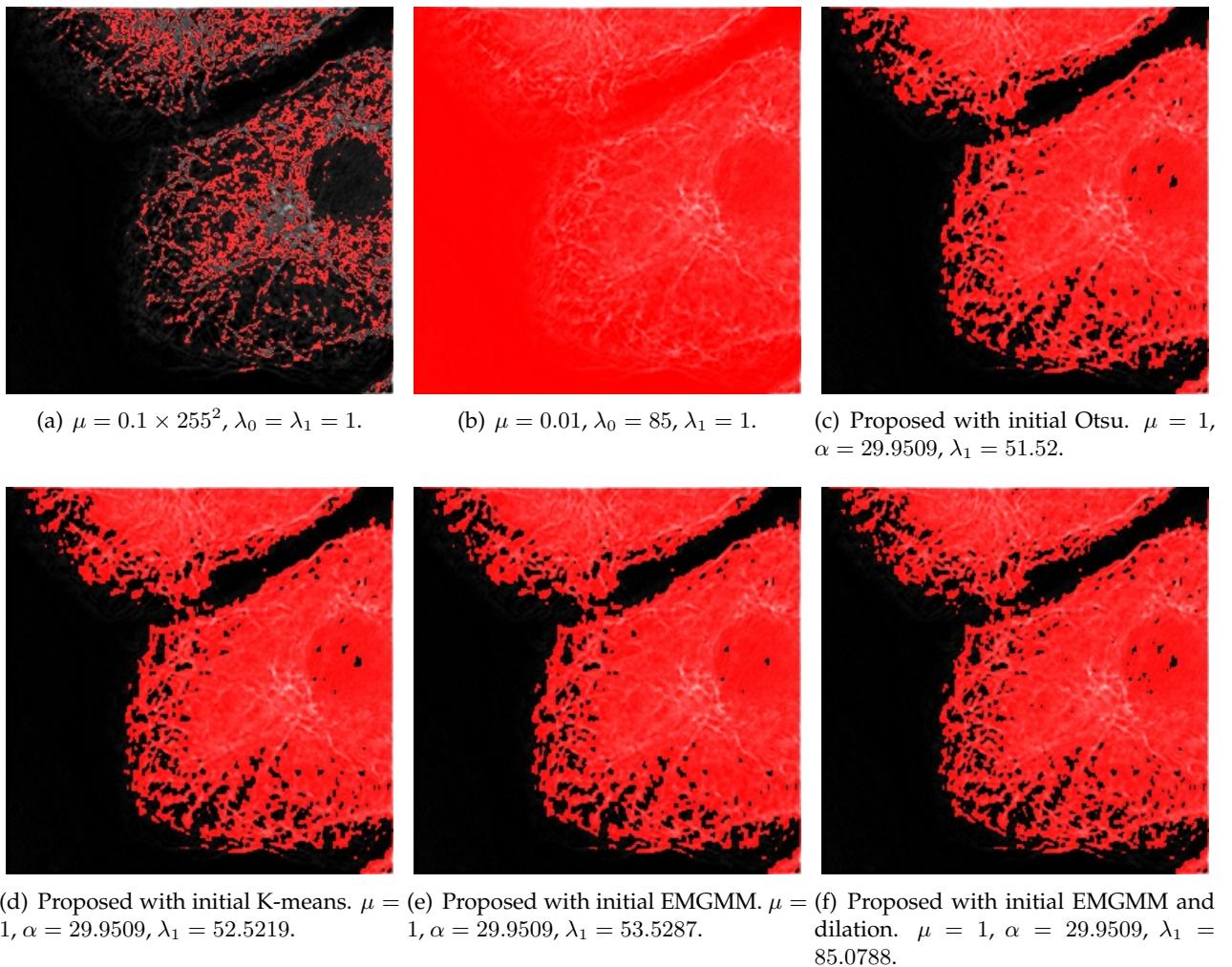


FIGURE 6.56: Image 18 from test set Appendix B segmentation results.

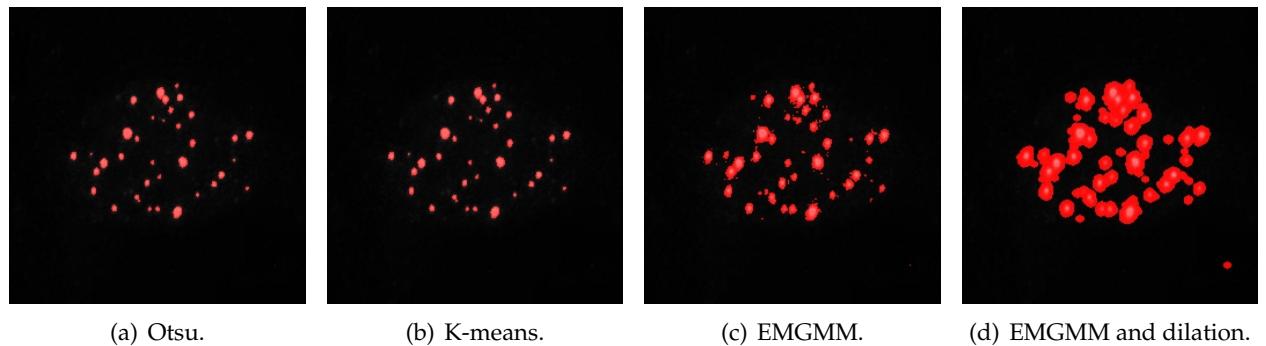


FIGURE 6.57: Image 19 from test set Appendix B initial masks.

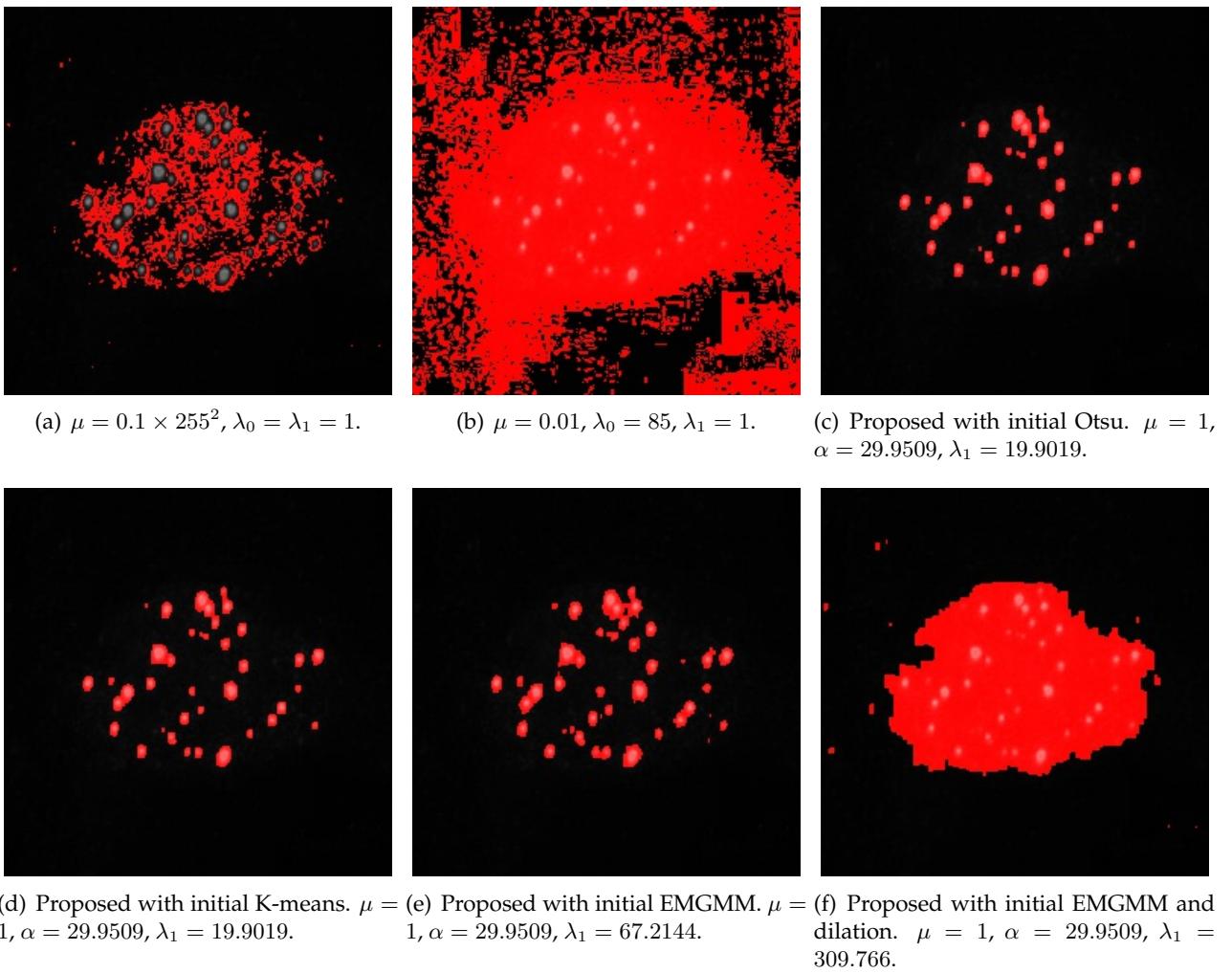


FIGURE 6.58: Image 19 from test set Appendix B segmentation results.

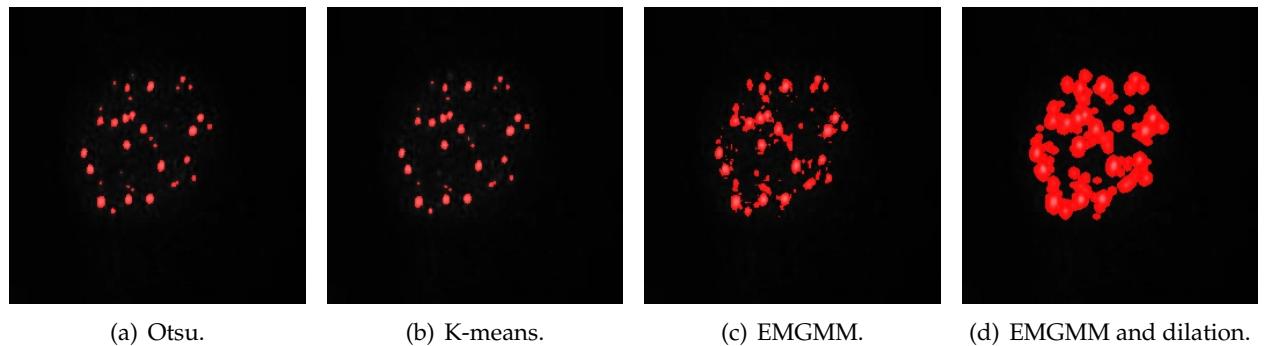


FIGURE 6.59: Image 20 from test set Appendix B initial masks.

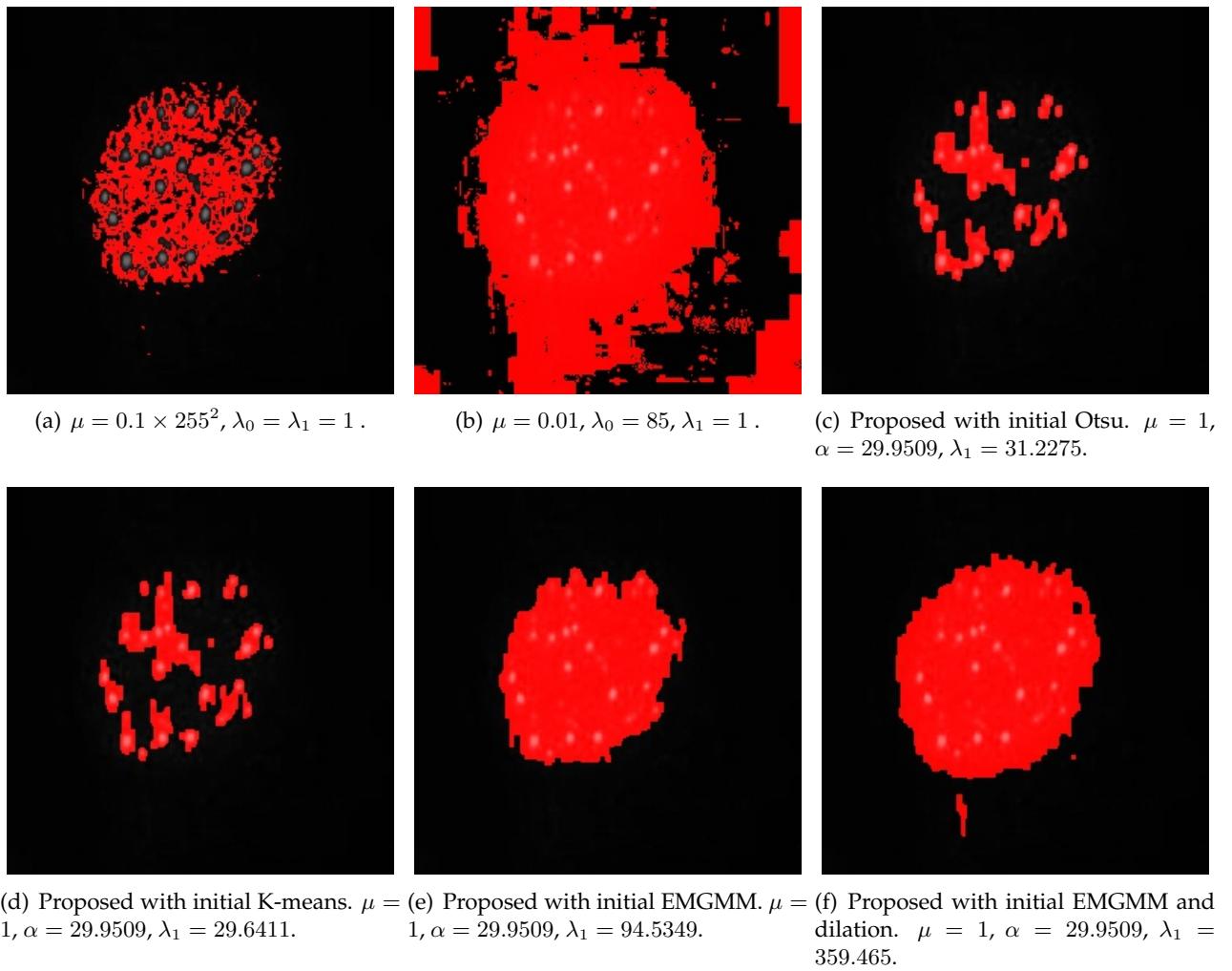


FIGURE 6.60: Image 20 from test set Appendix B segmentation results.

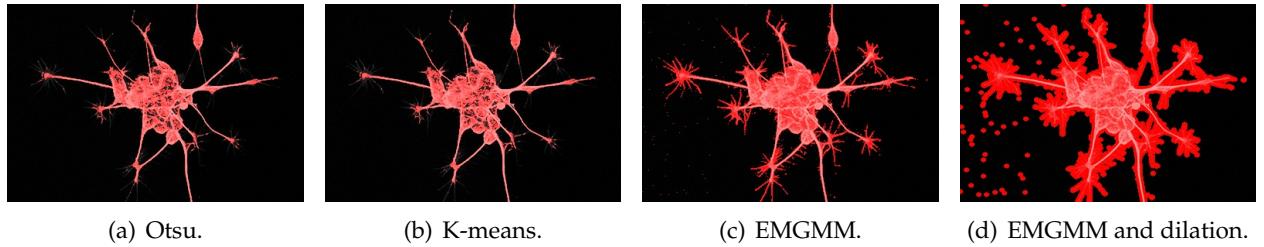


FIGURE 6.61: Image 21 from test set Appendix B initial masks.

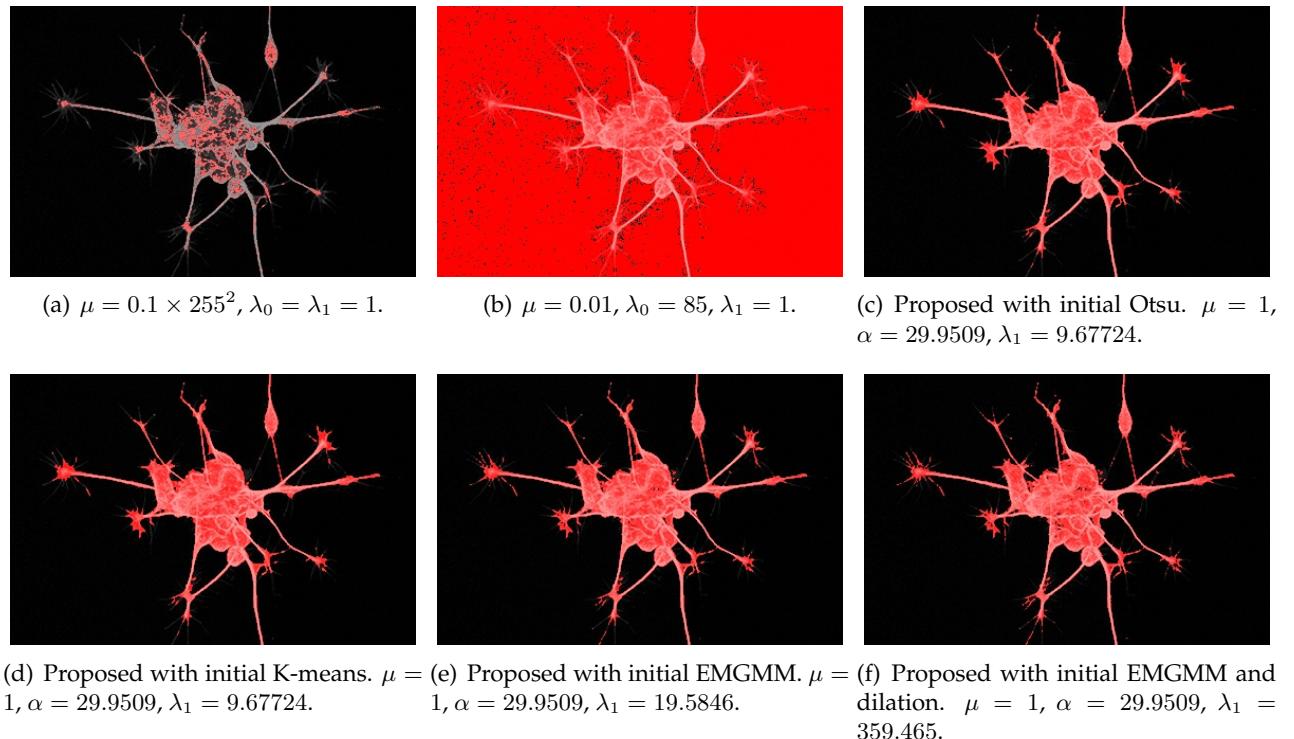


FIGURE 6.62: Image 21 from test set Appendix B segmentation results.

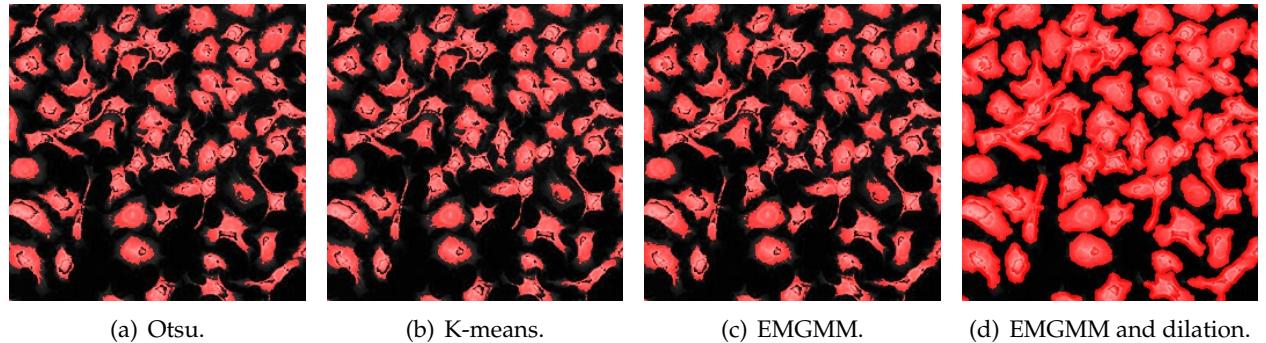


FIGURE 6.63: Image 22 from test set Appendix B initial masks.

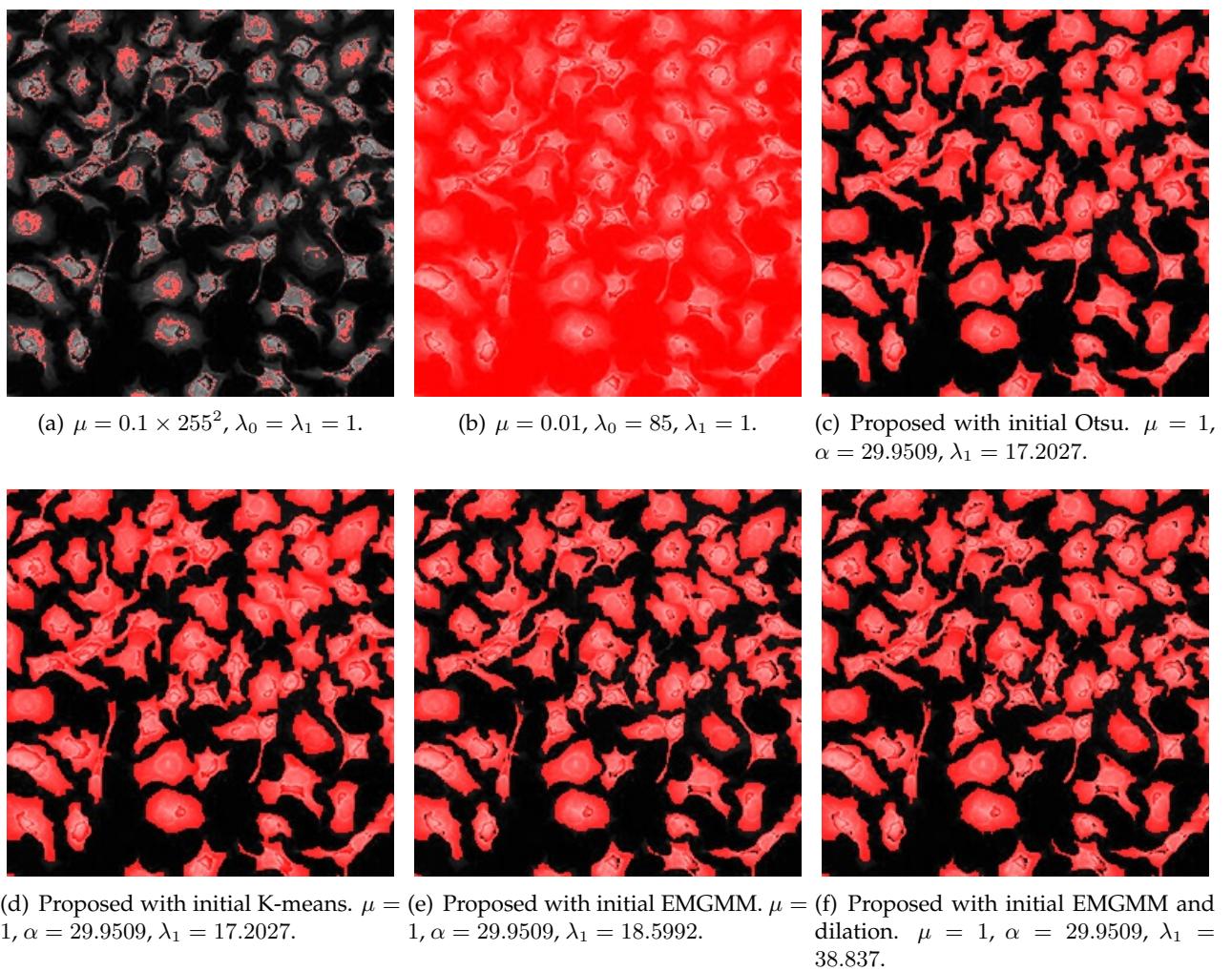


FIGURE 6.64: Image 22 from test set Appendix B segmentation results.

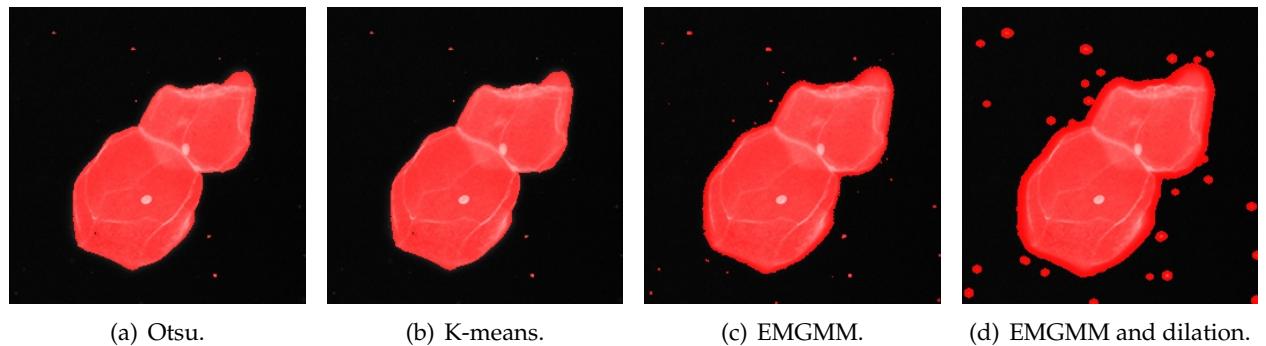


FIGURE 6.65: Image 23 from test set Appendix B initial masks.

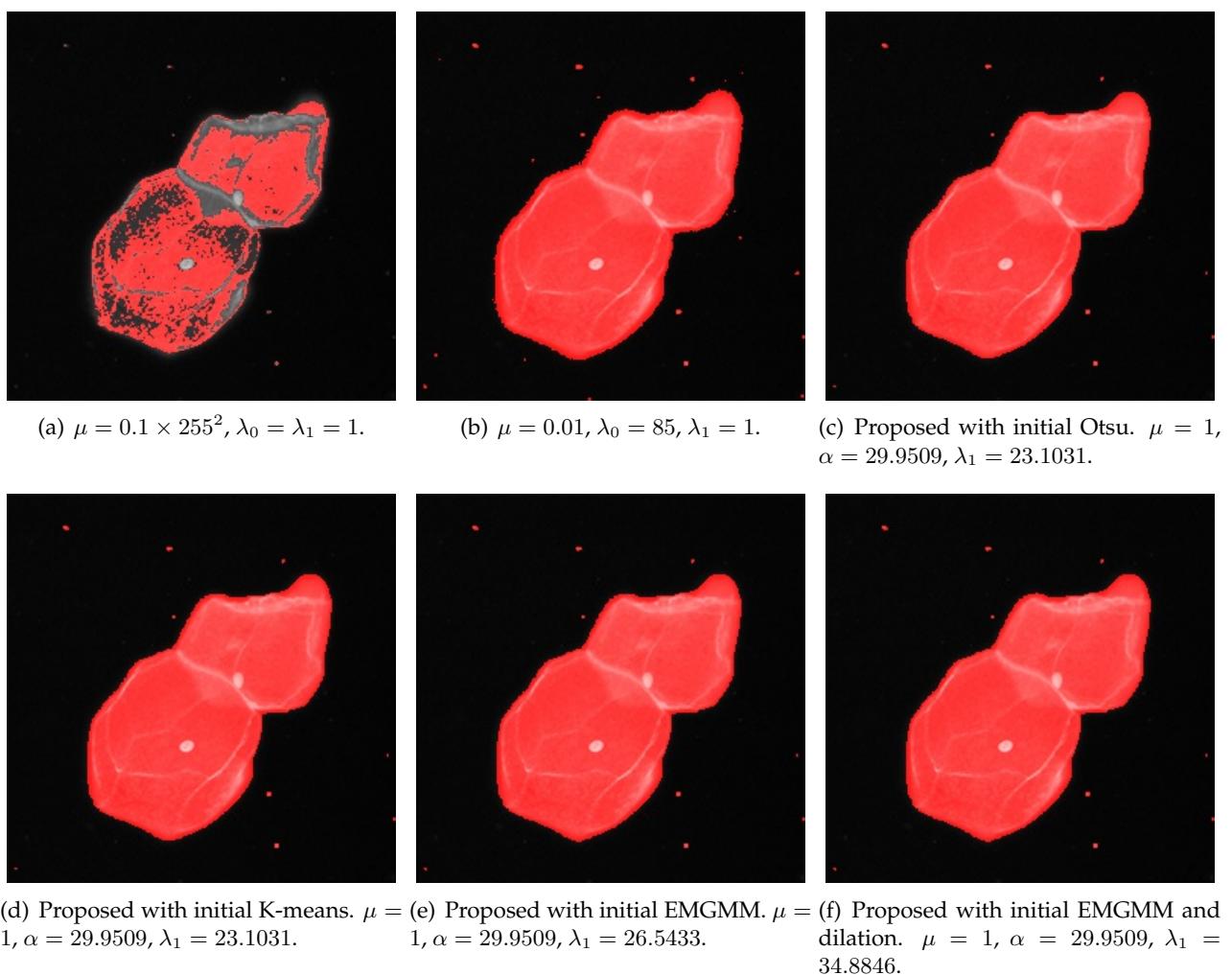


FIGURE 6.66: Image 23 from test set Appendix B segmentation results.

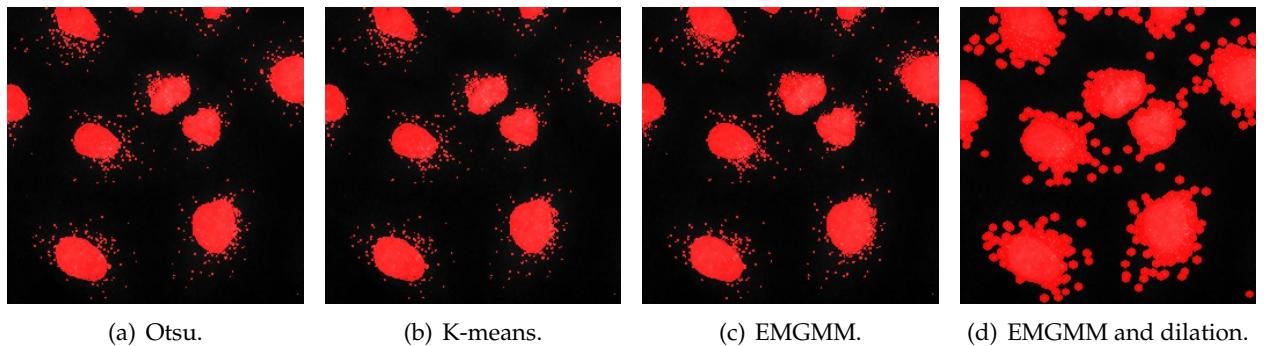


FIGURE 6.67: Image 24 from test set Appendix B initial masks.

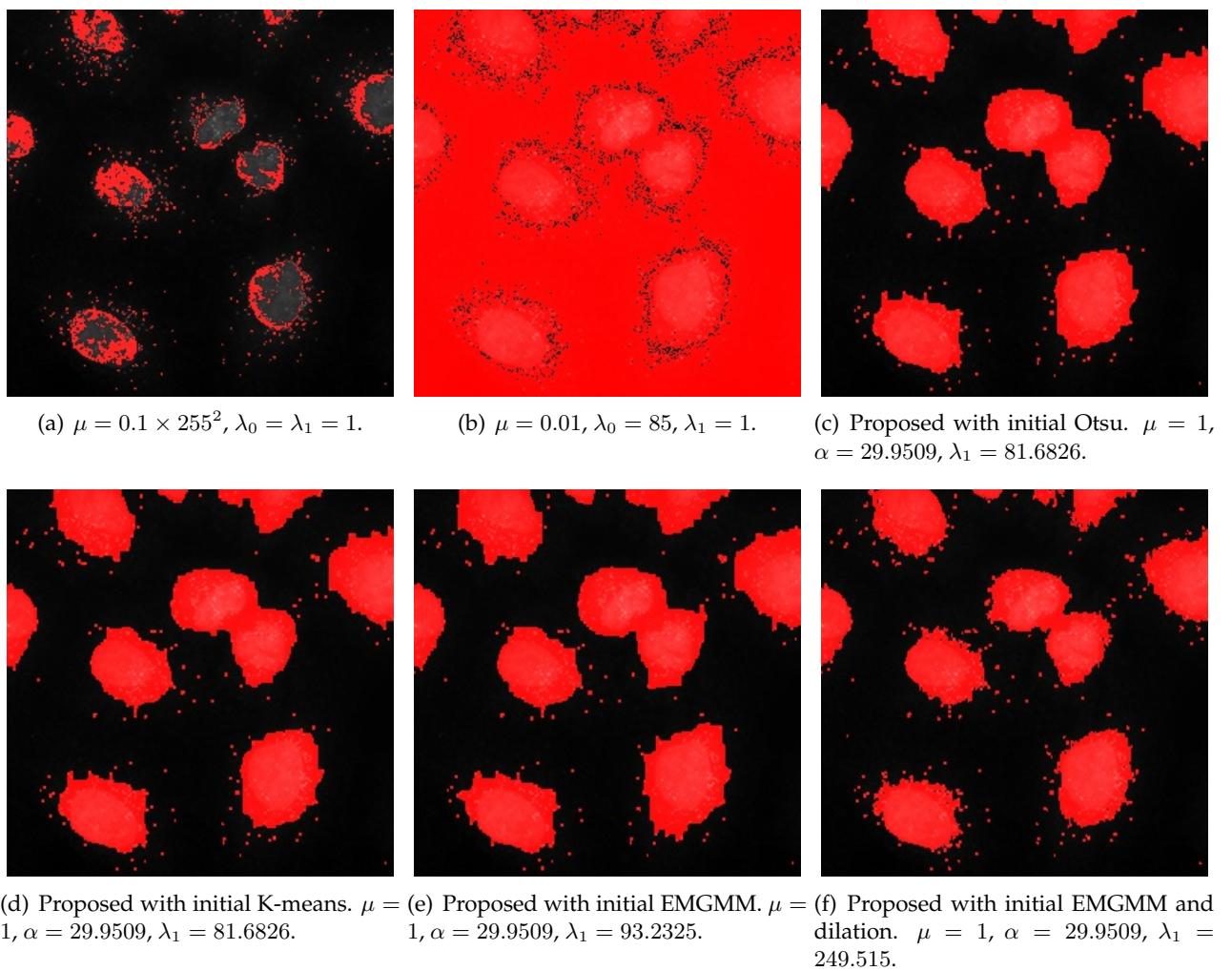


FIGURE 6.68: Image 24 from test set Appendix B segmentation results.

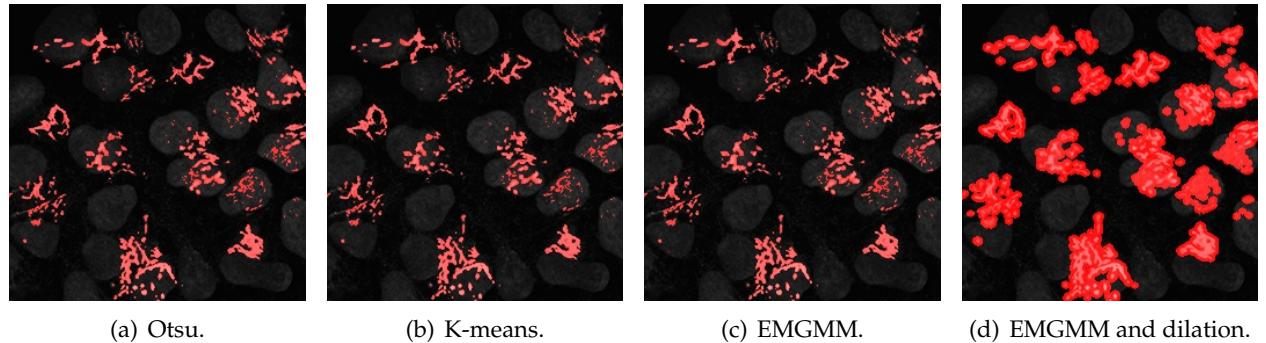


FIGURE 6.69: Image 25 from test set Appendix B initial masks.

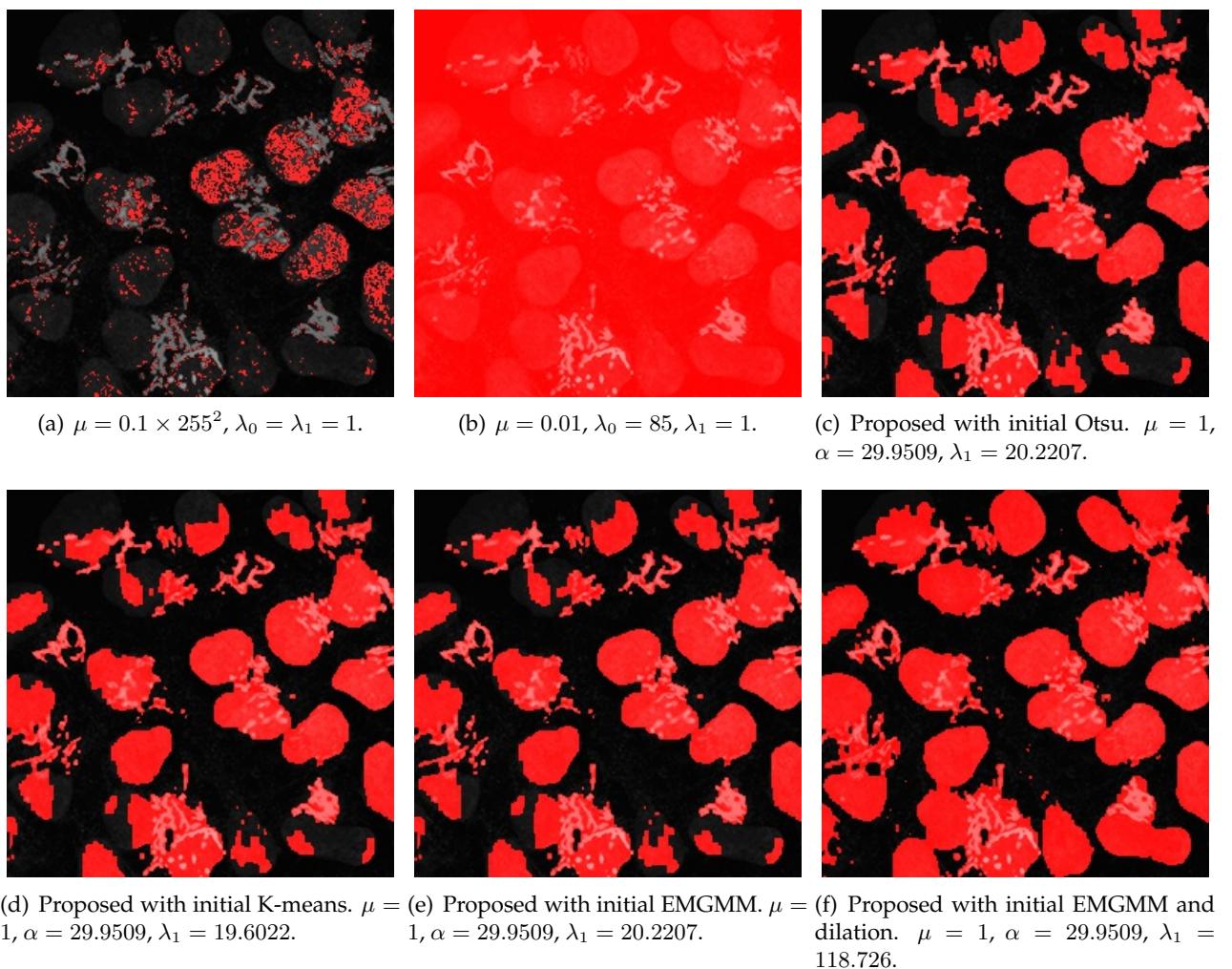


FIGURE 6.70: Image 25 from test set Appendix B segmentation results.

TABLE 6.7: Parameter Settings and Segmentation Results.

Image	Initial		Calculated				Final		Ideal		$ \Delta c_0 + \Delta c_1 $
	c_0	c_1	p_e	h	α	λ_1	c_0	c_1	c_0	c_1	
1-n	0.00895	0.40423	-	-	-	-	0.13967	0.50221	0.00165	0.41653	0.22371
1-m	0.00895	0.40423	-	-	-	-	0.00586	0.43683			0.02452
1-o	0.02625	0.46444	0.09395	0.20711	29.9509	7.47524	0.00556	0.43593			0.02332
1-k	0.02625	0.46444	0.09395	0.20711	29.9509	7.47525	0.00556	0.43593			0.02333
1-e	0.00873	0.44298	0.07582	0.18796	29.9509	7.61164	0.00638	0.43808			0.02629
1-d	0.00256	0.42168	0.06732	0.17555	29.9509	8.17148	0.00785	0.44130			0.03098
2-n	0.05448	0.06033	-	-	-	-	0.05264	0.11120	0.03371	0.10663	0.02352
2-m	0.05448	0.06033	-	-	-	-	0.05490	0.05822			0.06960
2-o	0.04080	0.14206	0.05645	0.08259	29.9509	139.983	0.03037	0.09425			0.00905
2-k	0.04080	0.14206	0.05645	0.08259	29.9509	139.984	0.03037	0.09425			0.00905
2-e	0.04001	0.13836	0.05521	0.08061	29.9509	148.373	0.03014	0.09362			0.00945
2-d	0.02952	0.07818	0.03704	0.04961	29.9509	606.199	0.02838	0.09184			0.00945
3-n	0.42108	0.11303	-	-	-	-	0.03498	0.33562	0.01567	0.37442	0.05812
3-m	0.42108	0.11303	-	-	-	-	0.00591	0.24458			0.12009
3-o	0.01897	0.39916	0.07772	0.17589	29.9509	9.93033	0.01364	0.35543			0.01697
3-k	0.01897	0.39916	0.07772	0.17589	29.9509	9.93034	0.01364	0.35543			0.01697
3-e	0.01118	0.32665	0.05992	0.14139	29.9509	14.4226	0.01312	0.34998			0.02190
3-d	0.00485	0.21558	0.03741	0.09183	29.9509	32.3237	0.01235	0.34116			0.02995
4-n	0.07545	0.20713	-	-	-	-	0.12290	0.47460	0.06335	0.45869	0.07546
4-m	0.07545	0.20713	-	-	-	-	0.05485	0.41407			0.03612
4-o	0.06630	0.48896	0.13160	0.24075	29.9509	8.03467	0.05593	0.42575			0.02552
4-k	0.06630	0.48896	0.13170	0.24076	29.9509	8.03468	0.05593	0.42575			0.02552
4-e	0.05670	0.43339	0.11490	0.21218	29.9509	10.1158	0.05996	0.45888			0.00319
4-d	0.05442	0.40712	0.10892	0.19999	29.9509	11.5385	0.05930	0.45449			0.00015
5-n	0.13533	0.32821	-	-	-	-	0.15583	0.47288	0.12897	0.46482	0.03493
5-m	0.13533	0.32821	-	-	-	-	0.07446	0.41944			0.00914
5-o	0.09483	0.46700	0.15233	0.24844	29.9509	10.3623	0.07522	0.42195			0.01088
5-k	0.09483	0.46701	0.15234	0.24844	29.9509	10.3624	0.07522	0.42196			0.01088
5-e	0.08698	0.45324	0.14357	0.23815	29.9509	10.7002	0.08706	0.45341			0.03049
5-d	0.08161	0.43958	0.13692	0.22936	29.9509	11.2011	0.08709	0.45348			0.03054
6-n	0.02387	0.06961	-	-	-	-	0.04522	0.10423	0.00365	0.08386	0.06194
6-m	0.02387	0.06961	-	-	-	-	0.02346	0.05387			0.04979
6-o	0.03077	0.15296	0.04965	0.08121	29.9509	96.1295	0.00650	0.09013			0.00913
6-k	0.03173	0.15753	0.05117	0.08366	29.9509	90.7016	0.00733	0.09131			0.01113
6-e	0.03348	0.16650	0.05404	0.08839	29.9509	81.1216	0.00798	0.09229			0.01276
6-d	0.02350	0.11789	0.03809	0.06246	29.9509	161.108	0.00446	0.08688			0.00384
7-n	0.01226	0.04400	-	-	-	-	0.02896	0.09064	0.01346	0.08035	0.02579
7-m	0.01226	0.04400	-	-	-	-	0.01214	0.07147			0.00755

7-o	0.02098	0.16457	0.04317	0.08025	29.9509	69.6183	0.01476	0.09537				0.01632
7-k	0.02098	0.16457	0.04317	0.08025	29.9509	69.6184	0.01477	0.09538				0.01634
7-e	0.01766	0.12742	0.03462	0.06297	29.9509	119.128	0.01353	0.08515				0.00488
7-d	0.02163	0.16513	0.02354	0.04053	29.9509	331.532	0.01393	0.09152				0.01165
8-n	0.04166	0.09509	-	-	-	-	0.07004	0.14628	0.02345	0.10122		0.09166
8-m	0.04166	0.09509	-	-	-	-	0.04011	0.07601				0.04187
8-o	0.05632	0.25262	0.08666	0.13735	29.9509	37.2522	0.04505	0.16167				0.08207
8-k	0.05632	0.25262	0.08666	0.13735	29.9509	37.2523	0.04505	0.16167				0.08207
8-e	0.05497	0.23995	0.08355	0.13133	29.9509	41.9465	0.04440	0.15773				0.07747
8-d	0.04971	0.18973	0.07134	0.10750	29.9509	73.2113	0.03964	0.13450				0.04949
9-n	0.01870	0.06039	-	-	-	-	0.03729	0.14899	0.02003	0.15095		0.01921
9-m	0.01870	0.06039	-	-	-	-	0.01830	0.12179				0.02743
9-o	0.02349	0.17829	0.04741	0.08739	29.9509	59.9044	0.01959	0.14422				0.00629
9-k	0.02349	0.17829	0.04741	0.08739	29.9509	59.9045	0.01959	0.14422				0.00629
9-e	0.01962	0.14699	0.03929	0.07219	29.9509	88.4765	0.01988	0.14957				0.00124
9-d	0.01864	0.11758	0.03393	0.05948	29.9509	146.629	0.01965	0.14696				0.00361
10-n	0.01651	0.08788	-	-	-	-	0.04651	0.11558	0.01009	0.10699		0.04501
10-m	0.01651	0.08788	-	-	-	-	0.00991	0.10639				0.00042
10-o	0.02193	0.12366	0.03765	0.06392	29.9509	138.712	0.01007	0.10694				0.00003
10-k	0.01674	0.11606	0.03209	0.05774	29.9509	145.504	0.01005	0.10689				0.00006
10-e	0.02429	0.12767	0.04027	0.06696	29.9509	134.305	0.01008	0.10697				0.00001
10-d	0.01316	0.10633	0.02756	0.05162	29.9509	165.378	0.01000	0.10671				0.00019
11-n	0.06341	0.10813	-	-	-	-	0.07140	0.17037	0.04957	0.16022		0.03199
11-m	0.06341	0.10813	-	-	-	-	0.06248	0.09365				0.07948
11-o	0.05184	0.17292	0.07055	0.10182	29.9509	97.9168	0.04129	0.14344				0.00849
11-k	0.05184	0.17292	0.07055	0.10182	29.9509	97.9168	0.04129	0.14344				0.00849
11-e	0.04848	0.16516	0.06651	0.09664	29.9509	105.429	0.04061	0.14129				0.00997
11-d	0.06373	0.18362	0.05598	0.08088	29.9509	154.458	0.04301	0.14940				0.00426
12-n	0.02937	0.05704	-	-	-	-	0.03715	0.12645	0.01438	0.12225		0.02699
12-m	0.02937	0.05704	-	-	-	-	0.02924	0.04705				0.09006
12-o	0.01644	0.13791	0.03521	0.06658	29.9509	97.2781	0.00783	0.10746				0.00824
12-k	0.01644	0.13791	0.03521	0.06658	29.9509	97.2781	0.00783	0.10746				0.00824
12-e	0.01226	0.12439	0.02958	0.05854	29.9509	114.161	0.00746	0.10557				0.00975
12-d	0.02173	0.14265	0.02199	0.04642	29.9509	160.407	0.00784	0.10775				0.00796
13-n	0.03887	0.07238	-	-	-	-	0.05347	0.16217	0.03707	0.17735		0.03158
13-m	0.03887	0.07238	-	-	-	-	0.03714	0.06115				0.11628
13-o	0.03368	0.22981	0.06398	0.11463	29.9509	37.3133	0.02422	0.16750				0.00299
13-k	0.03430	0.23396	0.06515	0.11671	29.9509	36.0075	0.02267	0.15618				0.00677
13-e	0.02970	0.20405	0.05664	0.10167	29.9509	47.2175	0.02342	0.16248				0.00122
13-d	0.02115	0.12638	0.03742	0.06459	29.9509	129.640	0.02045	0.14440				0.01633
14-n	0.06029	0.06139	-	-	-	-	0.05777	0.16960	0.03323	0.16654		0.02761
14-m	0.06029	0.06139	-	-	-	-	0.00000	0.06097				0.07232
14-o	0.03772	0.30268	0.07866	0.14709	29.9509	20.4463	0.02669	0.19334				0.02027

14-k	0.03772	0.30268	0.07866	0.14709	29.9509	20.4463	0.02669	0.19334			0.02027
14-e	0.03359	0.25979	0.06854	0.12696	29.9509	28.0516	0.02523	0.18031			0.00578
14-d	0.02592	0.15617	0.04605	0.07969	29.9509	84.6065	0.02585	0.18815			0.01425
15-n	0.01840	0.00893	-	-	-	-	0.01161	0.07348			0.06484
15-m	0.01840	0.00893	-	-	-	-	0.00784	0.01275			0.12179
15-o	0.00746	0.20099	0.03736	0.08734	29.9509	38.3238	0.00627	0.14850	0.00599	0.13270	0.01609
15-k	0.00746	0.20099	0.03736	0.08734	29.9509	38.3238	0.00627	0.14850			0.01609
15-e	0.00454	0.06893	0.01449	0.03112	29.9509	346.136	0.00584	0.12732			0.00523
15-d	0.00407	0.04491	0.01038	0.02093	29.9509	860.426	0.00564	0.11797			0.01438
16-n	0.04415	0.06767	-	-	-	-	0.05110	0.31654			0.08116
16-m	0.04415	0.06767	-	-	-	-	0.04313	0.05892	0.01943	0.36603	0.33081
16-o	0.02332	0.41028	0.08311	0.18304	29.9509	9.58570	0.01792	0.34656			0.01796
16-k	0.02332	0.41028	0.08311	0.18304	29.9509	9.58570	0.01792	0.34656			0.01796
16-e	0.01595	0.31360	0.06194	0.13881	29.9509	16.2015	0.01751	0.34066			0.02346
16-d	0.01240	0.20776	0.04258	0.09304	29.9509	37.6071	0.01674	0.32819			0.03515
17-n	0.07135	0.09023	-	-	-	-	0.07254	0.22327			0.06720
17-m	0.07135	0.09023	-	-	-	-	0.07058	0.07857	0.04673	0.26467	0.20994
17-o	0.04768	0.30903	0.08806	0.15555	29.9509	21.0148	0.03922	0.24421			0.01294
17-k	0.04768	0.30903	0.08806	0.15555	29.9509	21.0148	0.03922	0.24421			0.01294
17-e	0.04090	0.25794	0.07444	0.13049	29.9509	30.4718	0.03779	0.23286			0.02287
17-d	0.03681	0.21881	0.06493	0.11193	29.9509	43.3300	0.03736	0.22938			0.02592
18-n	0.08499	0.09648	-	-	-	-	0.07755	0.18418			0.09629
18-m	0.08499	0.09648	-	-	-	-	0.00000	0.09214	0.01243	0.15301	0.04843
18-o	0.03657	0.20349	0.06237	0.10547	29.9509	51.5200	0.01491	0.16035			0.00982
18-k	0.03498	0.20029	0.06053	0.10322	29.9509	52.5219	0.01479	0.16009			0.00944
18-e	0.03336	0.19711	0.05866	0.10095	29.9509	53.5287	0.01364	0.15719			0.00539
18-d	0.01379	0.14367	0.03386	0.06740	29.9509	85.0788	0.01438	0.16005			0.00900
19-n	0.00892	0.02787	-	-	-	-	0.01849	0.04199			0.01771
19-m	0.00892	0.02787	-	-	-	-	0.00784	0.02717	0.01094	0.05214	0.02187
19-o	0.01731	0.28587	0.05881	0.12816	29.9509	19.9019	0.01611	0.18929			0.14232
19-k	0.01731	0.28587	0.05881	0.12816	29.9509	19.9019	0.01611	0.18929			0.14232
19-e	0.01568	0.16181	0.03826	0.07599	29.9509	67.2144	0.01563	0.15783			0.11039
19-d	0.01388	0.08195	0.02439	0.04198	29.9509	309.766	0.01059	0.05115			0.00064
20-n	0.00778	0.02631	-	-	-	-	0.01592	0.04527			0.01634
20-m	0.00778	0.02631	-	-	-	-	0.00784	0.02987	0.01017	0.05587	0.02367
20-o	0.01658	0.23098	0.04971	0.10508	29.9509	31.2275	0.01410	0.10095			0.04902
20-k	0.01666	0.23671	0.05066	0.10749	29.9509	29.6411	0.01420	0.10368			0.05185
20-e	0.01507	0.13829	0.03411	0.06594	29.9509	94.5349	0.01065	0.06009			0.00471
20-d	0.01287	0.07606	0.02264	0.03896	29.9509	359.465	0.00954	0.05207			0.00317
21-n	0.03017	0.07699	-	-	-	-	0.03501	0.36663			0.08947
21-m	0.03017	0.07699	-	-	-	-	0.02604	0.04685	0.00965	0.30254	0.27208
21-o	0.01359	0.39872	0.07309	0.17256	29.9509	9.67724	0.01025	0.35268			0.05076

21-k	0.01359	0.39872	0.07309	0.17256	29.9509	9.67724	0.01025	0.35268				0.05076
21-e	0.00772	0.27844	0.04955	0.11946	29.9509	19.5846	0.01018	0.35304				0.05104
21-d	0.00717	0.13728	0.02264	0.03896	29.9509	359.465	0.00954	0.05207				0.25036
22-n	0.14636	0.15438	-	-	-	-	0.13589	0.34221				0.15033
22-m	0.14636	0.15438	-	-	-	-	0.00000	0.15135				0.09591
22-o	0.06587	0.35472	0.11050	0.18509	29.9509	17.2027	0.03676	0.27528	0.04026	0.28753		0.00874
22-k	0.06587	0.35472	0.11050	0.18509	29.9509	17.2027	0.03676	0.27528				0.00874
22-e	0.05808	0.33588	0.10100	0.17274	29.9509	18.5992	0.04178	0.29272				0.00673
22-d	0.03248	0.22473	0.06219	0.11184	29.9509	38.8370	0.03865	0.28447				0.00145
23-n	0.02449	0.12763	-	-	-	-	0.05654	0.26228				0.04183
23-m	0.02449	0.12763	-	-	-	-	0.02594	0.26215				0.01136
23-o	0.02805	0.27730	0.06656	0.13093	29.9509	23.1031	0.02650	0.26851	0.02827	0.27584		0.00556
23-k	0.02805	0.27730	0.06656	0.13093	29.9509	23.1031	0.02650	0.26851				0.00556
23-e	0.02569	0.25823	0.06162	0.12167	29.9509	26.5433	0.02649	0.26838				0.00568
23-d	0.02501	0.22786	0.05635	0.10874	29.9509	34.8846	0.02649	0.26839				0.00567
24-n	0.04744	0.05351	-	-	-	-	0.04570	0.14237				0.03186
24-m	0.04744	0.05351	-	-	-	-	0.04705	0.05137				0.12421
24-o	0.03258	0.16514	0.05307	0.08729	29.9509	81.6826	0.02706	0.11987	0.03581	0.16433		0.03571
24-k	0.03258	0.16514	0.05307	0.08729	29.9509	81.6826	0.02706	0.11987				0.03571
24-e	0.03115	0.15523	0.05033	0.08237	29.9509	93.2325	0.02639	0.11461				0.04031
24-d	0.02616	0.10201	0.03788	0.05747	29.9509	249.515	0.02781	0.12694				0.02940
25-n	0.07693	0.09011	-	-	-	-	0.07952	0.15395				0.06571
25-m	0.07693	0.09011	-	-	-	-	0.00000	0.08513				0.02585
25-o	0.06532	0.33175	0.10648	0.17529	29.9509	20.2207	0.03901	0.16680	0.02839	0.13938		0.03806
25-k	0.06567	0.33628	0.10749	0.17737	29.9509	19.6022	0.03990	0.16887				0.04101
25-e	0.06532	0.33175	0.10640	0.17529	29.9509	20.2207	0.03901	0.16680				0.03806
25-d	0.08493	0.40334	0.07377	0.10217	29.9509	118.726	0.02844	0.14221				0.00289

As can be seen, the parameter settings presented previously in literature are not robust over a large range of image types that commonly occur in fluorescence microscopy whereas the proposed method produces more objectively consistent and visually accurate segmentations. From the table we can also see that the final means from the proposed method are closer to the ideal means obtained from the ground truth.

Measuring Segmentation Efficiency We now objectively compare the methods under critique. We use a confusion matrix to quantify the number of correct and incorrect classifications. We relate the classification of each pixel as:

TP (True Positives) An object pixel that is correctly classified as an object.

TN (True Negatives) A background pixel that is correctly classified as a background.

FP (False Positives) A background pixel that is incorrectly classified as an object.

FN (False Negatives) An object pixel that is incorrectly classified as a background.

From these counts, we calculate the following binary classification measures:

$$precision = \frac{TP}{TP + FP}$$

$$recall = \frac{TP}{TP + FN}$$

$$accuracy = \frac{TP + TN}{TP + TN + FP + FN}$$

$$MCC = \frac{TP \times TN - FP \times FN}{\sqrt{(TP + FP)(TP + FN)(TN + FP)(TN + FN)}}$$

Accuracy is the fraction of the pixels that are correctly classified from among all pixels. MCC is the *Matthews Correlation Coefficient* and is more accurate measure of accuracy when comparing classes whose sizes differ greatly. The result is a value between -1 and $+1$, where -1 refers to a complete opposite classification, i.e. all background pixels are labelled as "object" and all object pixels are labelled as "background", and $+1$ means a perfect classification, 0 is equivalent to a random classification. The results for each image in the test set are tabulated in Table 6.8. The overall efficiency for each method is tabulated in Table 6.9. For each table, rows that are highlighted in blue show the method that performed the best for each image in Table 6.8 and over the test set in Table 6.9.

TABLE 6.8: Segmentation Efficiency.

Image	TP	TN	FP	FN	Precision	Recall	Accuracy	MCC
1-n	24954	25516	0	15066	1.000000	0.623538	0.770111	0.626140
1-m	37875	25506	10	2145	0.999736	0.946402	0.967117	0.934010
1-o	37973	25506	10	2047	0.999737	0.948851	0.968613	0.936880
1-k	37973	25506	10	2047	0.999737	0.948851	0.968613	0.936880
1-e	37736	25510	6	2284	0.999841	0.942929	0.965057	0.930095
1-d	37364	25513	3	2656	0.999920	0.933633	0.959427	0.919468
2-n	5561	43411	179	16385	0.968815	0.253395	0.747253	0.416180
2-m	21864	120	43470	82	0.334650	0.996264	0.335449	-0.008374
2-o	21856	37015	6575	90	0.768738	0.995899	0.898300	0.804725
2-k	21856	37015	6575	90	0.768738	0.995899	0.898300	0.804725
2-e	21878	36617	6973	68	0.758310	0.996901	0.892563	0.795678
2-d	21867	34669	8921	79	0.710244	0.996400	0.862671	0.748686
3-n	9113	55407	24	992	0.997373	0.901831	0.984497	0.939774
3-m	10105	47657	7774	0	0.565188	1.000000	0.881378	0.697081
3-o	10104	54541	890	1	0.919047	0.999901	0.986404	0.950885
3-k	10104	54541	890	1	0.919047	0.999901	0.986404	0.950885

3-e	10105	54277	1154	0	0.897504	1.000000	0.982391	0.937454
3-d	10105	53847	1584	0	0.864488	1.000000	0.975830	0.916397
4-n	10654	49844	99	4939	0.990793	0.683255	0.923126	0.783314
4-m	15593	46825	3118	0	0.833360	1.000000	0.952423	0.883930
4-o	15593	47552	2391	0	0.867048	1.000000	0.963516	0.90859
4-k	15593	47552	2391	0	0.867048	1.000000	0.963516	0.90859
4-e	14839	48771	1172	754	0.926800	0.951645	0.970612	0.919839
4-d	14920	48593	1350	673	0.917025	0.956840	0.969131	0.916491
5-n	21195	38591	2276	3474	0.903029	0.859175	0.912262	0.811918
5-m	24669	31160	9707	0	0.717623	1.000000	0.851883	0.739708
5-o	24669	31482	9385	0	0.724408	1.000000	0.856796	0.747027
5-k	24669	31482	9385	0	0.724408	1.000000	0.856796	0.747027
5-e	24669	35418	5449	0	0.819078	1.000000	0.916855	0.842536
5-d	24669	35427	5440	0	0.819323	1.000000	0.916992	0.842769
6-n	8041	25755	0	31740	1.000000	0.202132	0.515686	0.300907
6-m	38173	1358	24397	1608	0.610085	0.959579	0.603195	0.028915
6-o	35876	25713	42	3905	0.998831	0.901838	0.939774	0.883440
6-k	35096	25729	26	4685	0.999260	0.882230	0.928116	0.863032
6-e	34459	25734	21	5322	0.999391	0.866218	0.918472	0.846506
6-d	37815	25503	252	1966	0.993380	0.950579	0.966156	0.931253
7-n	3213	47332	19	14972	0.994121	0.176684	0.771255	0.364534
7-m	17876	43120	4231	309	0.808613	0.983008	0.930725	0.846322
7-o	13903	47235	116	4282	0.991726	0.764531	0.932892	0.832125
7-k	13903	47235	116	4282	0.991726	0.764531	0.932892	0.832125
7-e	16478	46937	414	1707	0.975491	0.906131	0.967636	0.918642
7-d	14740	46846	505	3445	0.966874	0.810558	0.939728	0.847704
8-n	4494	22158	0	38884	1.000000	0.103601	0.406677	0.193924
8-m	42118	133	22025	1260	0.656627	0.970953	0.644699	-0.075582
8-o	16780	22158	0	26598	1.000000	0.386832	0.594147	0.419288
8-k	16780	22158	0	26598	1.000000	0.386832	0.594147	0.419288
8-e	17641	22158	0	25737	1.000000	0.406681	0.607285	0.433758
8-d	24348	22141	17	19030	0.999302	0.561298	0.709366	0.548682
9-n	4890	53207	3	7436	0.999387	0.396722	0.886490	0.589732
9-m	12325	49186	4024	1	0.753869	0.999919	0.938583	0.834732

9-o	12301	52336	874	25	0.933662	0.997972	0.986282	0.957060
9-k	12301	52336	874	25	0.933662	0.997972	0.986282	0.957060
9-e	12258	52948	262	68	0.979073	0.994483	0.994965	0.983657
9-d	12298	52633	577	28	0.955184	0.997728	0.990768	0.970635
10-n	14187	31146	2	20201	0.999859	0.412557	0.691727	0.500151
10-m	34387	30887	261	1	0.992467	0.999971	0.996002	0.992013
10-o	34373	31109	39	15	0.998867	0.999564	0.999176	0.998348
10-k	34381	31095	53	7	0.998461	0.999796	0.999084	0.998165
10-e	34363	31115	33	25	0.999041	0.999273	0.999115	0.998226
10-d	34384	31013	135	4	0.996089	0.999884	0.997879	0.995755
11-n	14671	40720	132	10013	0.991083	0.594353	0.845200	0.684969
11-m	24500	3572	37280	184	0.396568	0.992546	0.428345	0.166735
11-o	24284	33091	7761	400	0.757809	0.983795	0.875473	0.769468
11-k	24284	33091	7761	400	0.757809	0.983795	0.875473	0.769468
11-e	24347	32248	8604	337	0.738885	0.986347	0.863571	0.751768
11-d	24030	35179	5673	654	0.809009	0.973505	0.903458	0.812402
12-n	7204	45666	295	12371	0.960661	0.368020	0.806732	0.519903
12-m	19530	776	45185	45	0.301785	0.997701	0.309845	0.060018
12-o	19415	39881	6080	160	0.761522	0.991826	0.904785	0.806923
12-k	19415	39881	6080	160	0.761522	0.991826	0.904785	0.806923
12-e	19465	39285	6676	110	0.744616	0.994381	0.896454	0.793664
12-d	19408	39947	6014	167	0.763433	0.991469	0.905685	0.808358
13-n	2608	53651	1298	7979	0.667691	0.246340	0.858444	0.346226
13-m	10541	2533	52416	46	0.167432	0.995655	0.199493	0.079031
13-o	9284	47991	6958	1303	0.571604	0.876925	0.873947	0.639563
13-k	9606	46365	8584	981	0.528092	0.907339	0.854050	0.617332
13-e	9525	47363	7586	1062	0.556659	0.899688	0.868042	0.638175
13-d	10320	44499	10450	267	0.496870	0.974780	0.836472	0.620618
14-n	1461	51387	508	12180	0.742001	0.107104	0.806396	0.231433
14-m	13641	0	51895	0	0.208145	1.000000	0.208145	NaN
14-o	9703	48117	3778	3938	0.719754	0.711311	0.882263	0.641301
14-k	9703	48117	3778	3938	0.719754	0.711311	0.882263	0.641301
14-e	10329	47119	4776	3312	0.683813	0.757203	0.876587	0.641224
14-d	10154	47865	4030	3487	0.715877	0.744374	0.885300	0.657278

15-n	936	62133	33	2434	0.965944	0.277745	0.962357	0.507270
15-m	3370	3061	59105	0	0.053942	1.000000	0.098129	0.051537
15-o	2847	62143	23	523	0.991986	0.844807	0.991669	0.911385
15-k	2847	62143	23	523	0.991986	0.844807	0.991669	0.911385
15-e	3345	61918	248	25	0.930977	0.992582	0.995834	0.959144
15-d	3363	61528	638	7	0.840540	0.997923	0.990158	0.911074
16-n	4543	58089	4	2900	0.999120	0.610372	0.955688	0.762067
16-m	7443	99	57994	0	0.113743	1.000000	0.115082	0.013923
16-o	7434	57377	716	9	0.912147	0.998791	0.988937	0.948497
16-k	7434	57377	716	9	0.912147	0.998791	0.988937	0.948497
16-e	7434	57155	938	9	0.887960	0.998791	0.985550	0.934020
16-d	7437	56680	1413	6	0.840339	0.999194	0.978348	0.905052
17-n	2858	55353	634	6691	0.818442	0.299298	0.888229	0.452365
17-m	9536	298	55689	13	0.146202	0.998639	0.150055	0.020336
17-o	8509	51944	4043	1040	0.677900	0.891088	0.922440	0.734195
17-k	8509	51944	4043	1040	0.677900	0.891088	0.922440	0.734195
17-e	8915	51233	4754	634	0.652206	0.933606	0.917786	0.736986
17-d	9006	50960	5027	543	0.641773	0.943135	0.915009	0.733933
18-n	10662	28372	3	26499	0.999719	0.286914	0.595612	0.384991
18-m	37161	0	28375	0	0.567032	1.000000	0.567032	NaN
18-o	34582	28099	276	2579	0.992082	0.930599	0.956436	0.914421
18-k	34653	28086	289	2508	0.991729	0.932510	0.957321	0.916017
18-e	35373	27881	494	1788	0.986227	0.951885	0.965179	0.930209
18-d	34845	28118	257	2316	0.992678	0.937677	0.960739	0.922580
19-n	6080	49795	190	9471	0.969697	0.390972	0.852585	0.559970
19-m	15551	21811	28174	0	0.355655	1.000000	0.570099	0.393942
19-o	1743	49985	0	13808	1.000000	0.112083	0.789307	0.296349
19-k	1743	49985	0	13808	1.000000	0.112083	0.789307	0.296349
19-e	2340	49985	0	13211	1.000000	0.150473	0.798416	0.344988
19-d	15144	48771	1214	407	0.925786	0.973828	0.975266	0.933388
20-n	7631	52325	94	5486	0.987832	0.581764	0.914856	0.719637
20-m	13112	31433	20986	5	0.384539	0.999619	0.679703	0.479944
20-o	3934	52419	0	9183	1.000000	0.299916	0.859879	0.505181
20-k	3742	52419	0	9375	1.000000	0.285279	0.856949	0.491933

20-e	11467	52403	16	1650	0.998607	0.874209	0.974579	0.919788
20-d	12993	50360	2059	124	0.863208	0.990547	0.966690	0.904878
21-n	5859	54029	0	5648	1.000000	0.509168	0.913818	0.678954
21-m	11504	410	53619	3	0.176650	0.999739	0.181793	0.035231
21-o	9354	54019	10	2153	0.998932	0.812896	0.966995	0.883569
21-k	9354	54019	10	2153	0.998932	0.812896	0.966995	0.883569
21-e	9367	54019	10	2140	0.998934	0.814026	0.967194	0.884286
21-d	9386	54016	13	2121	0.998617	0.815677	0.967438	0.885155
22-n	9992	36090	0	19454	1.000000	0.339333	0.703156	0.469557
22-m	29446	0	36090	0	0.449310	1.000000	0.449310	NaN
22-o	29301	33903	2187	145	0.930545	0.995076	0.964417	0.930373
22-k	29301	33903	2187	145	0.930545	0.995076	0.964417	0.930373
22-e	28465	35927	163	981	0.994306	0.966685	0.982544	0.964943
22-d	29163	35176	914	283	0.969611	0.990389	0.981735	0.963345
23-n	11850	49497	42	4147	0.996468	0.740764	0.936081	0.824684
23-m	15994	48127	1412	3	0.918879	0.999812	0.978409	0.944698
23-o	15978	48676	863	19	0.948756	0.998812	0.986542	0.964737
23-k	15978	48676	863	19	0.948756	0.998812	0.986542	0.964737
23-e	15978	48664	875	19	0.948080	0.998812	0.986359	0.964275
23-d	15978	48665	874	19	0.948137	0.998812	0.986374	0.964313
24-n	3284	57098	577	4577	0.850557	0.417759	0.921356	0.562635
24-m	7861	1055	56620	0	0.121912	1.000000	0.136047	0.047223
24-o	7853	48469	9206	8	0.460344	0.998982	0.859406	0.621496
24-k	7853	48469	9206	8	0.460344	0.998982	0.859406	0.621496
24-e	7858	47094	10581	3	0.426162	0.999618	0.838501	0.589715
24-d	7844	50004	7671	17	0.505575	0.997837	0.882690	0.661018
25-n	5597	32002	27	27910	0.995199	0.167040	0.573715	0.296608
25-m	33507	0	32029	0	0.511276	1.000000	0.511276	NaN
25-o	23435	31813	216	10072	0.990867	0.699406	0.843018	0.720953
25-k	22771	31815	214	10736	0.990690	0.679589	0.832916	0.704884
25-e	23435	31813	216	10072	0.990867	0.699406	0.843018	0.720953
25-d	31767	31142	887	1740	0.972836	0.948071	0.959915	0.920148

From the marked-up rows, we can clearly see that the proposed method supersedes the competing methods. This is better shown in Table 6.9. From the data in Table 6.8 we plot Precision vs

Recall in Figure 6.71. From Figure 6.71 we can see that the parameter settings proposed by Masaka *et al.* tend to have low precision but a very high recall. We can also see that the parameter settings proposed by El Zehiry *et al.* tend to have very low recall but a high precision. The proposed method, for all initialisation methods, have a relatively high precision and recall. We plot the accuracy over the test set which is shown in Figure 6.72. We can clearly see that the performance of the competitor parameter settings are very erratic.

TABLE 6.9: Overall Segmentation Efficiency.

Method	Precision		Recall		Accuracy	
	Mean	Std. Dev.	Mean	Std. Dev.	Mean	Std. Dev.
El-Zehiry <i>et al.</i> [239]	0.951912	0.088710	0.421993	0.226299	0.805732	0.151199
Maska <i>et al.</i> [237]	0.485812	0.292765	0.993592	0.014007	0.547369	0.321099
Prop. - Otsu	0.876652	0.153530	0.845668	0.240808	0.911657	0.087720
Prop. - K-means	0.874892	0.157316	0.844808	0.242856	0.909905	0.088468
Prop. - EMGMM	0.875713	0.161034	0.883279	0.202419	0.918983	0.087190
Prop. - EMGMM+Dilate	0.860245	0.149355	0.939366	0.103397	0.935329	0.064493

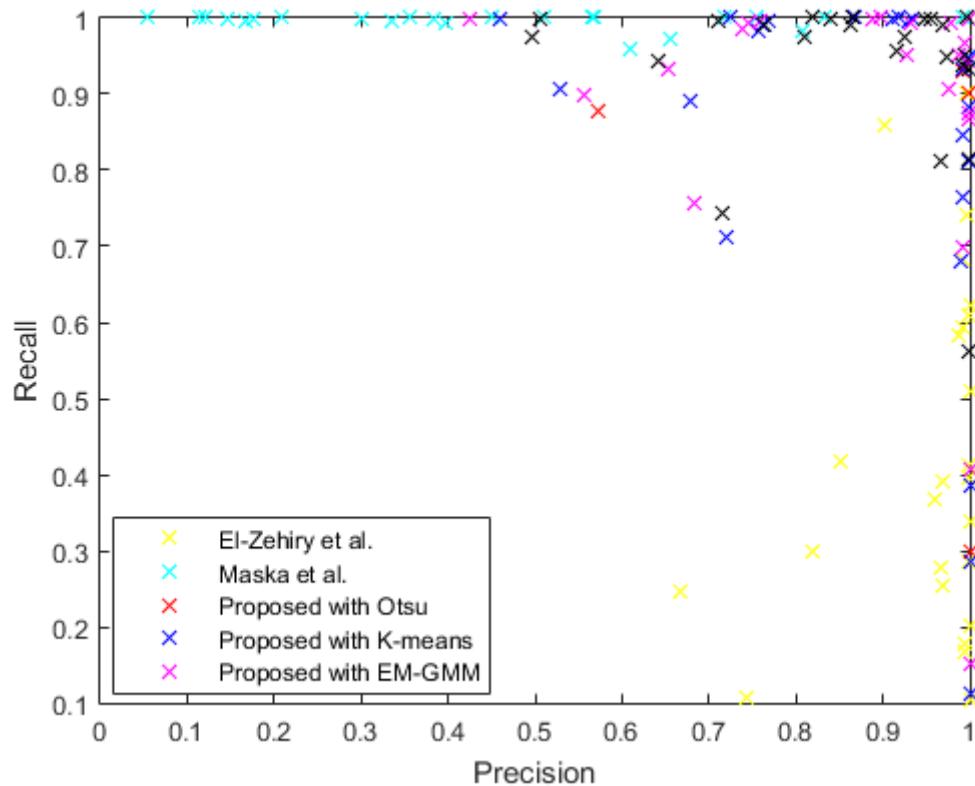


FIGURE 6.71: Precision vs Recall over test set.

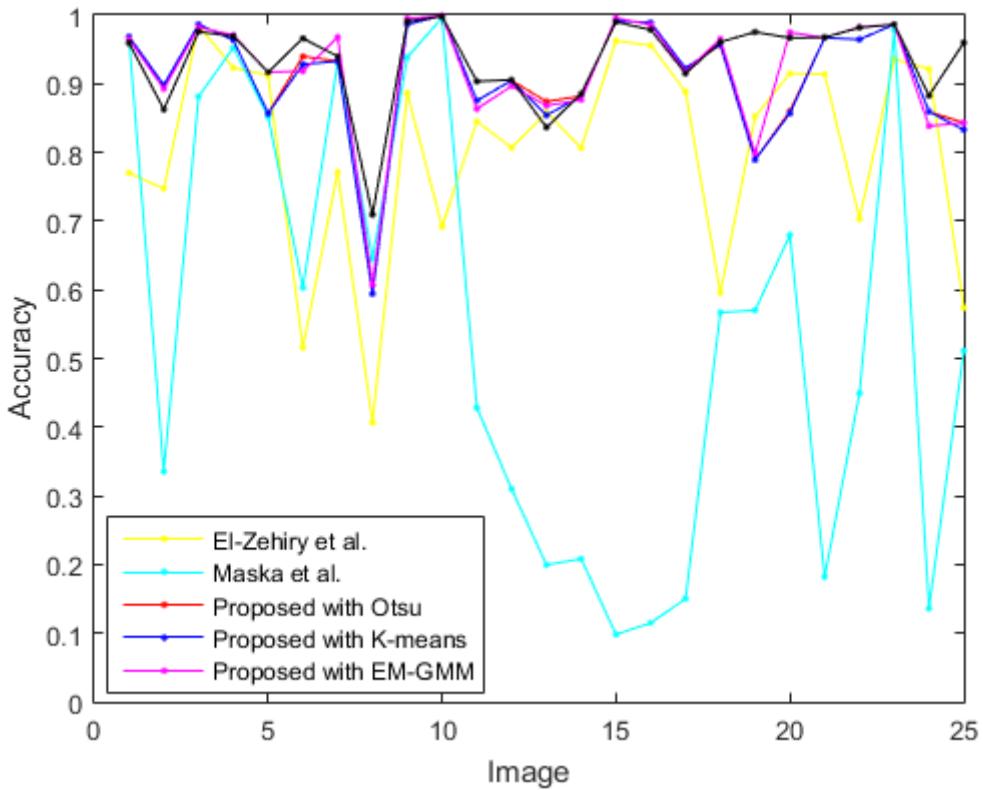


FIGURE 6.72: Accuracy over test set.

6.4 Discussion

The general parameter settings by El-Zehiry *et al.* [239] clearly outperform the fluorescence imaging specific parameter settings presented by Masaka *et al.* [237] by 25.8363%. However, even the least accurate proposed parameter estimation technique outperforms El-Zehiry's parameter settings by 10.5925% while the most accurate proposed method boasts an increase of 12.9597%. The proposed method is also much more stable over a greater variety of fluorescence images, as can be seen by the standard deviation of accuracy shown in Table 6.9.

6.5 Conclusion

Fluorescence images are hugely diverse; even so of the images within the fields of cellular biology and medicine. The design of a general and automatic segmentation method is not a trivial exercise; however, it is much needed.

We have presented a novel parameter estimation technique which is able to tune the parameters of the graph-cut application of the Chan-Vese segmentation method, to the specific image. This is a huge contrast in comparison to the fixed parameter settings as proposed by El-Zehiry *et al.* [239] and Masaka *et al.* [237]. Our method, does however, rely on the strong assumption that the initial means obtained from the initial unsupervised segmentation scheme is relatively close to the final means. Fortunately though, in the domain of biological and medical fluorescence images, the unsupervised segmentation results obtained from an Otsu, K-means or EMGMM clustering is close enough to

outperform the competitor parameter settings. We have shown that an EMGMM clustering and a dilation of 3 pixels, with an elliptical dilation element, provides the best approximation to the final means which results in an average of 93.5329% classification accuracy. Given the diversity of fluorescence images in the test set, this is a very pleasing result, especially when compared to the competitor parameter settings by El-Zehiry *et al.* [239] which showed an average of 80.5732% and Masaka *et al.* [237] which showed an average of 54.7369%. Furthermore, considering the fact that it is not possible, for even the same professional, to consistently manually segment an image the same everytime, and that greater variation in groundtruths will result from many professionals, we can see that even a 10% variation in groundtruths will result in the proposed scheme superseding the other schemes. This further shows the robustness of the algorithm.

Once the initial means have been obtained, the parameters are adapted to the image by direct calculation by means of the *relationship variables and formulae*. These relationship variables and formulae encode the important relation of the parameters with each other. This allows the ability to adapt the parameter settings according to the properties of the image. These resulting parameters only allow high segmentation accuracy if the image has the properties that are encoded in the relationship variables and formulae.

The relationship formulae only take into account the information from the intensity of the image. Other information, such as texture, can be incorporated into the formulae as well. This will include adapting the graph to hold this information so that it can be accounted for in the energy function.

Chapter 7

Fluorescence Imaging Specific Energy Function for Interactive Segmentation

In the previous section, we worked with an automatic segmentation technique. As motivated in the introduction, it is ideal to remove as much manual input as possible, however, automatic segmentation isn't a silver bullet solution and at times it is necessary for the user to get involved. The goal of reducing user interaction still remains. In this section, we design a novel energy function specifically for the images acquired in fluorescence microscopy for interactive segmentation.

In Section 7.1 we cover two very popular and widely used graph weighting schemes for general graph cut purposes. We then focus on the proposed weighting system in Section 7.2. We determine the optimal parameters for all weighting schemes in Section 7.3 and present the results in Section 7.4. We present a discussion in Section 7.5 and close the chapter with a conclusion in Section 7.6.

7.1 Popular Weighting Systems

In this section, we briefly cover the two weighting systems that are very popular for interactive graph cut segmentation.

Seeding only Eriksson *et al.* [26] presented a weighting system that incorporates prior knowledge into the image partitioning process. The prior knowledge is given by the user by marking image regions that are either "object" (\mathcal{O}) or "background" (\mathcal{B}). From the histogram of the labelled regions, the intensity distribution is modelled as a Gaussian Mixture Model (GMM) which is fitted through Expectation Maximisation (EM). The terminal links are weighted as

$$D_p("obj") = \frac{\Pr(I_p|\mathcal{O})}{\Pr(I_p|\mathcal{O}) + \Pr(I_p|\mathcal{B})}, \quad (7.1)$$

$$D_p("bkg") = \frac{\Pr(I_p|\mathcal{B})}{\Pr(I_p|\mathcal{O}) + \Pr(I_p|\mathcal{B})}. \quad (7.2)$$

The neighbourhood interaction function they used was

$$V_{\{p,q\}} = \exp\left(\frac{r(p,q)}{\sigma_R}\right) \exp\left(-\frac{(I_p - I_q)^2}{\sigma_W}\right), \quad (7.3)$$

where $r(p, q)$ is the Euclidean distance between pixels p and q . Eriksson *et al.* did not explicitly state the number of Gaussian functions they've used in their intensity distribution model, therefore we've defaulted to using two curves.

Seeding with hard constraints Boykov and Jolly [25] proposed a general purpose interactive segmentation scheme for multi-dimensional images. As in the previous scheme, the user marks image regions that are either "object" (\mathcal{O}) or "background" (\mathcal{B}). An intensity distributions based on the histogram from the seeded regions are calculated for each labelled region. Data weights for unseeded nodes are weighted as the negative log-likelihood from the intensity distribution. For the pixel p , the terminal weighting is

$$D_p(\text{"obj"}) = -\ln \Pr(I_p | \mathcal{O}), \quad (7.4)$$

$$D_p(\text{"bkg"}) = -\ln \Pr(I_p | \mathcal{B}). \quad (7.5)$$

Boykov and Jolly did not explicitly state how the intensity distribution is to be derived from the histogram. Therefore, we've used a GMM of two Gaussian curves which approximates the histogram. The neighbourhood interaction function they've proposed is

$$V_{\{p,q\}} = \exp\left(-\frac{(I_p - I_q)^2}{2\sigma^2}\right) \cdot \frac{1}{r(p, q)}, \quad (7.6)$$

where $r(p, q)$ is the distance between pixels p and q . The distance function they used wasn't explicitly stated. Therefore, we've used the Euclidean norm.

For the marked pixels, in addition to serving as seeds, they're also hard constraints. To ensure that they retain the label that they're marked with, they're given a terminal link weight that cannot be saturated by the neighbourhood nodes. This unsaturable weight is calculated as

$$K = 1 + \max_{p \in \mathcal{P}} \sum_{q \in N_p} V_{\{p,q\}}. \quad (7.7)$$

Then the object-marked node p is data weighted as $c(S, p) = K$ and $c(p, T) = 0$, and background-marked node p is data weighted as $c(S, p) = 0$ and $c(p, T) = K$, where $c(u, v)$ is the capacity of the edge uv .

7.2 Proposed Weighting

In this section, we introduce a novel weighting system that is based on the general properties of black background fluorescence images.

Object weighting function We first normalise the intensity distribution. Let the maximum value of the intensity distribution be $k_{max} = \max(P_{FG})$. Now $P_{FG}^1 = \frac{P_{FG}}{k_{max}}$.

Let the maximum intensity value in the image occur at I_{max} . Let the weight for this intensity value be F . The value of F in the final function is the same as that of the intensity distribution from the Gaussian Mixture Model. Therefore $F = P_{FG}^1(I_{max})$.

Let the intensity value at which P_{FG}^1 is a maximum be denoted by ι_{FG} . The intensity distribution after the point where the maximum value occurs is replaced by a parabola where the turning point maximum is $(\iota_{FG}, P_{FG}^1(\iota_{FG}))$ and the end point is (I_{max}, F) . The parabola defined by these points is

$$P_{FG}^2(x) = \frac{F - 1}{(I_{max} - \iota_{FG})^2}(x - \iota_{FG})^2 + 1 \quad (7.8)$$

where $x \in [\iota_{FG}, I_{max}]$. The final object weighting function is

$$D_p("obj") = \begin{cases} P_{FG}^1(x) & x \in [0, \iota_{FG}] \\ P_{FG}^2(x) & x \in [\iota_{FG}, I_{max}] \end{cases} \quad (7.9)$$

This function is plotted in Figure 7.1.

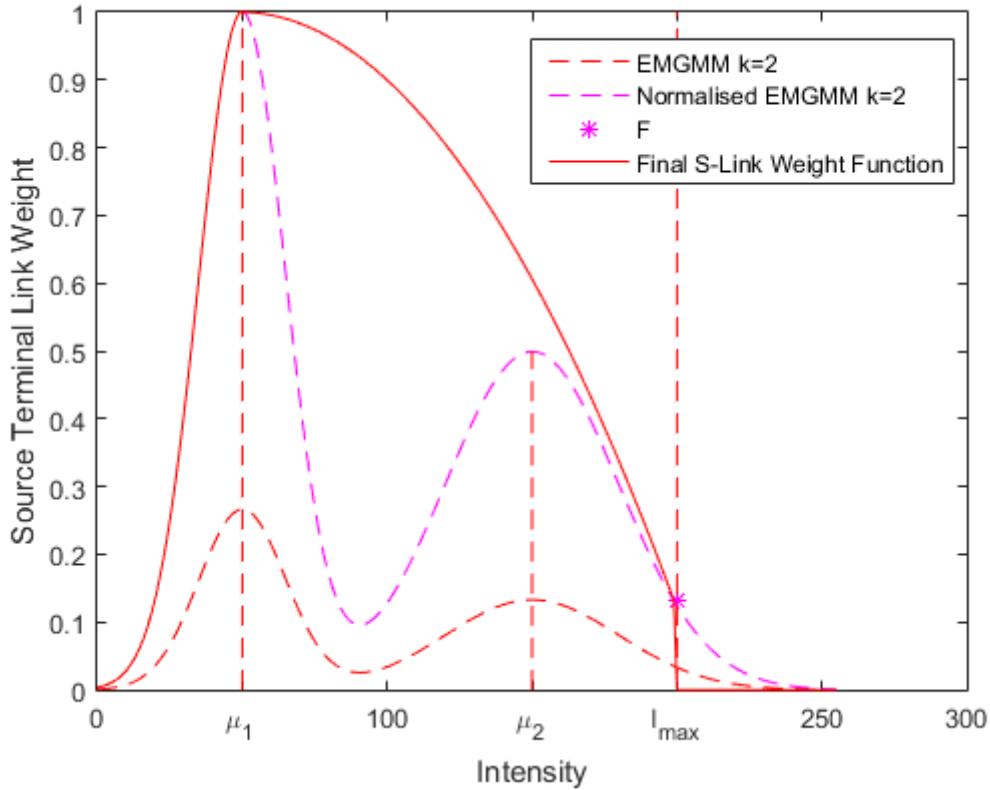


FIGURE 7.1: Object weighting function.

Background weighting function Similarly, the background weighting function is derived in the same way as the object weighting function. Let the maximum value of the intensity distribution be $k_{max} = P_{BG}$. Now $P_{BG}^1 = \frac{P_{BG}}{k_{max}}$.

Let the intensity value at which P_{BG}^1 is a maximum be denoted by ι_{BG} . The intensity distribution before the point where the maximum value occurs is replaced by a parabola where the turning point maximum is $(\iota_{BG}, P_{BG}^1(\iota_{BG}))$ and the end point is $(0, B)$. The parabola defined by these points is

$$P_{BG}^2(x) = \frac{B - 1}{(\iota_{BG})^2}(x - \iota_{BG})^2 + 1 \quad (7.10)$$

where $x \in [0, \iota_{BG}]$. The final background weighting function is

$$D_p("bkg") = \begin{cases} P_{BG}^2(x) & x \in [0, \iota_{BG}) \\ P_{BG}^1(x) & x \in [\iota_{BG}, I_{max}] \end{cases} \quad (7.11)$$

This function is plotted in Figure 7.2.

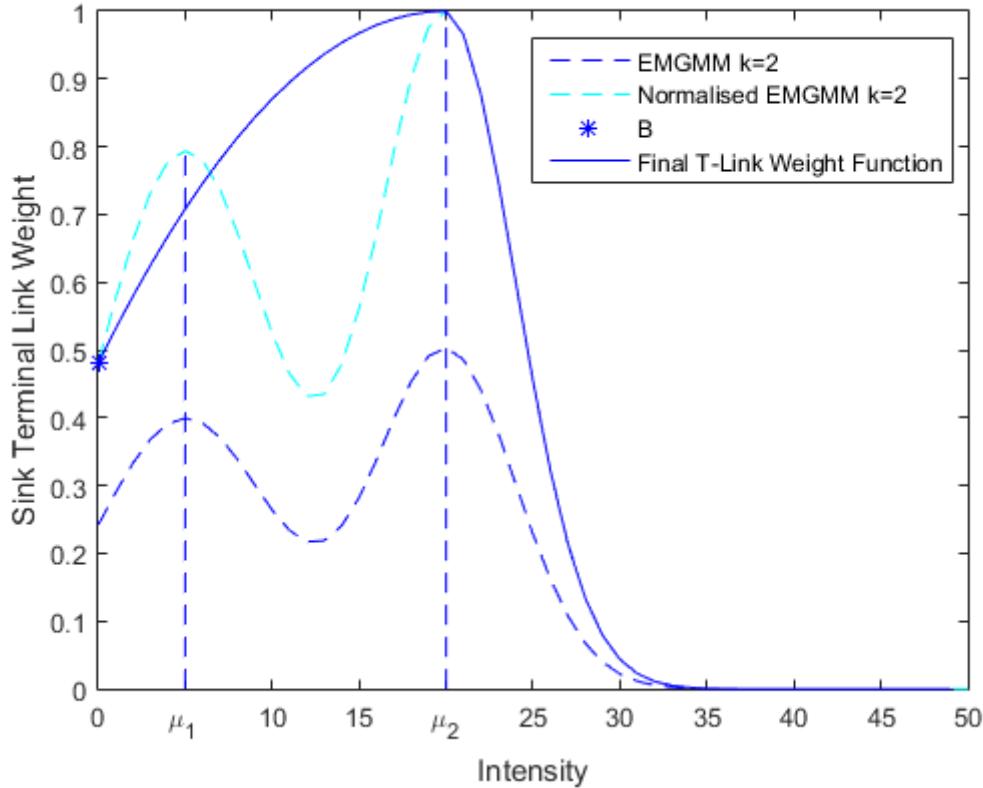


FIGURE 7.2: Background weighting function.

Hard Constraints To enable the seeds marked by the user as hard constraints, we use the same technique in [25]. Background and object seeds are weighted using Equation (7.7).

Neighbourhood Interaction Function The neighbourhood interaction weights from the energy functions proposed by Boykov and Jolly [25] and Eriksson *et al.* [26] both encode the dissimilarity between adjacent pixels as an exponential squared difference fall-off. This is compounded even further by reducing it by a function that is the difference of the distance between pixels. This is a very steep fall-off for fluorescence images as pixels over a longer range still have a relatively high correlation. An appropriate neighbourhood weighting function is possible but requires very specific setting of the parameters as a slight change in the parameters results in a steep reaction of the neighbourhood energy function.

To diminish the sharp reaction of the neighbourhood energy function of the previous functions, we use the following energy function which encourages greater correlation over longer distances and

milder reaction to parameter changes

$$V_{\{p,q\}} = \exp\left(-\frac{(I_p - I_q)}{\sigma}\right) \cdot \frac{1}{r(p,q)}, \quad (7.12)$$

where $r(p, q)$ is the distance between pixels p and q . We use the Euclidean distance.

The motivation for replacing part of the function with a parabolic function is to reduce the exponential decay of probability. The probability of pixels beyond the highest point, for object data, do not decay so quickly. Similarly, for the decay rate of pixels below the highest point for background data. It is also counter-intuitive to have a wavering probability distribution at higher or lower pixel intensity values. The modified energy function is still submodular and is derived off the energy model in Equation (3.13), therefore its global optimum can be found using graph cuts.

7.3 Determining Optimal Parameter Settings

Although the weighting schemes presented by Eriksson [26] and Boykov [25] are widely and probably one of the most common weighting schemes in use for interactive segmentation, there aren't any published parameters settings that work well specifically for fluorescent images. Therefore, we varied the tuning parameters for each weighting system and compared the segmentation results against the ground truth for the sample set in Figure B.1. We ran the same test for the proposed scheme with and without hard constraints. We used the following parameter settings

$$\lambda = \{0.5, 1.25, 2.5, 5.0, 7.5, 10.0\}$$

$$\sigma = \sigma_W = \{0.5, 1.25, 2.5, 5, 7.5\}$$

$$\sigma_R = 1.$$

To maintain as much commonality between the different weighting techniques we've used a common seed as illustrated in Figure 7.3. An extra seed is needed for the weighting systems by Eriksson *et al.* and Boykov *et al.* since, for all parameter settings, they were not able to register the cell nuclei as an object label. The results of the mean accuracy over the set for all parameter settings is plotted in Figure 7.4. The parameter setting that performed the best is marked by a red asterisk. The optimal parameters over the range were found to be

$(\lambda, \sigma_W, \sigma_R) = (10.0, 7.5, 1.0)$	Eriksson <i>et al.</i>
$(\lambda, \sigma) = (7.5, 5.0)$	Boykov <i>et al.</i>
$(\lambda, \sigma) = (0.5, 7.5)$	Proposed (No hard constraints)
$(\lambda, \sigma) = (0.5, 7.5)$	Proposed (Hard constraints)

The segmentation results with the optimal parameters for each weighting system is shown in Figures 7.5 to 7.10.

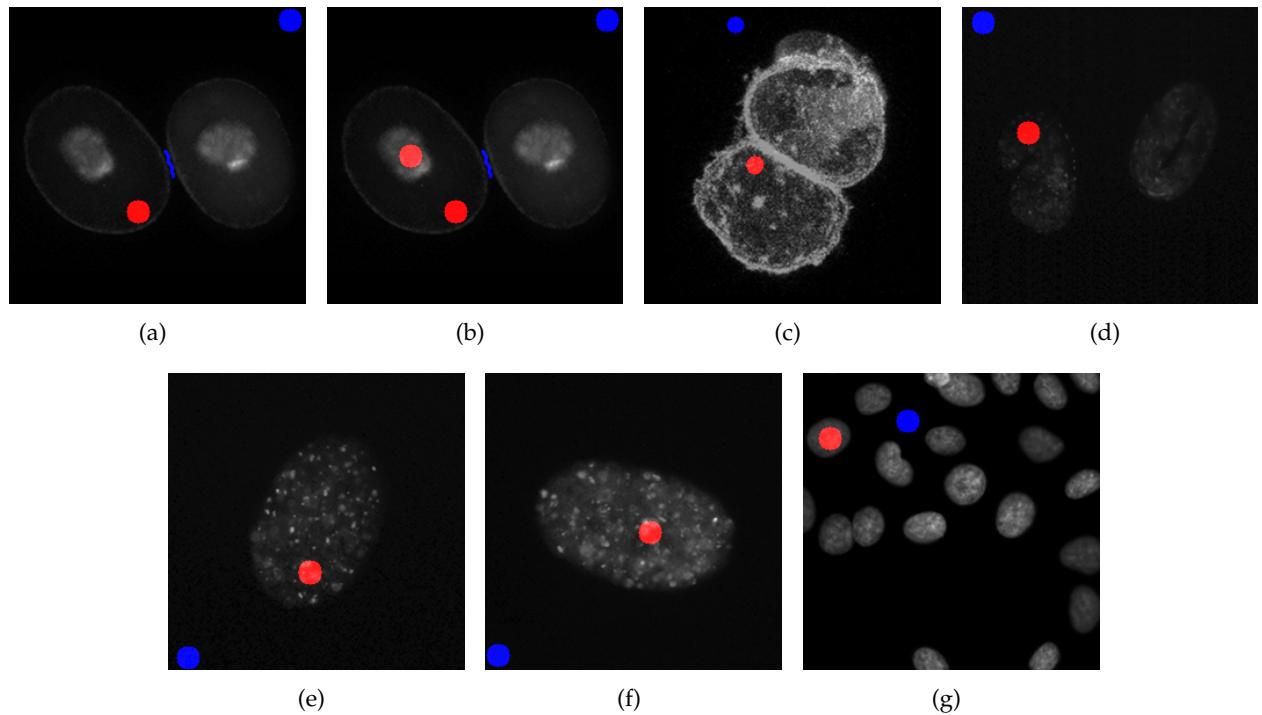


FIGURE 7.3: Seeds used over the sample set. The seeds for the first image differ in **(a)**, used for the proposed scheme, and **(b)** used for the weighting systems by Boykov and Eriksson as these were not able to grab the cell nuclei as part of the object.

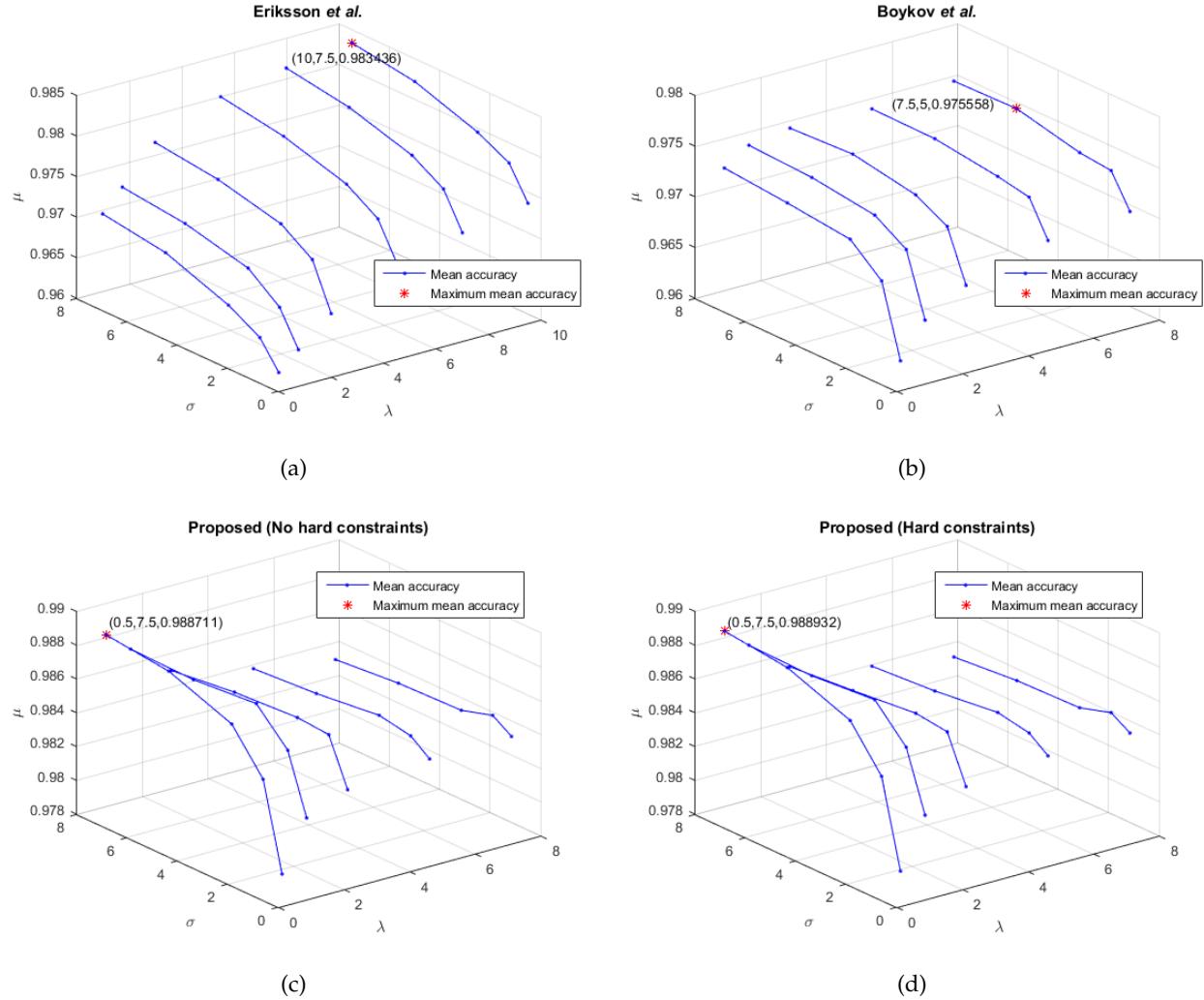


FIGURE 7.4: Plots of mean segmentation accuracy against the corresponding method tuning parameters. **(a)** Weighting system proposed by Eriksson *et al.* [26]. The remaining parameter is set to $\sigma_R = 1$. **(b)** Weighting system with hard constraints proposed by Boykov *et al.* [25]. **(c)** Proposed weighting system without hard constraints. **(d)** Proposed weighting system with hard constraints.

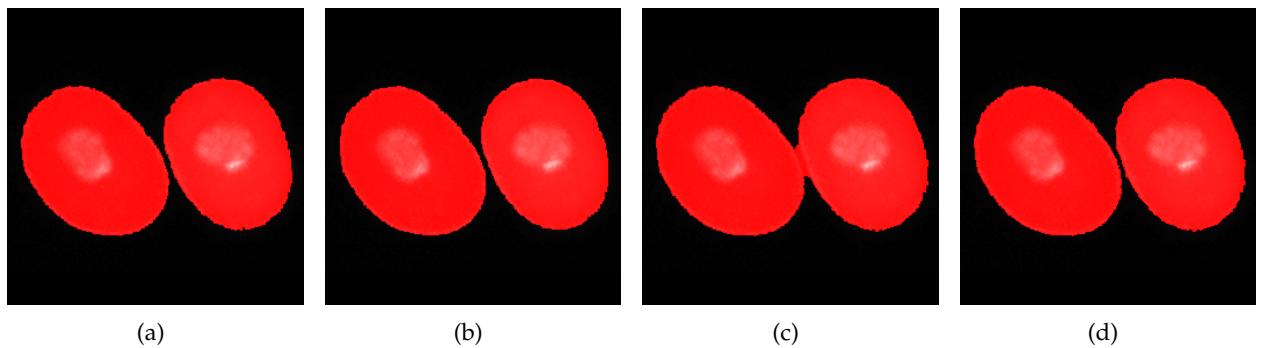


FIGURE 7.5: Interactive segmentation of Image 1 in sample set (Figure B.1). **(a)** Eriksson *et al.* [26]. **(b)** Boykov *et al.* [25]. **(c)** Proposed (No hard constraints) **(d)** Proposed with hard constraints.

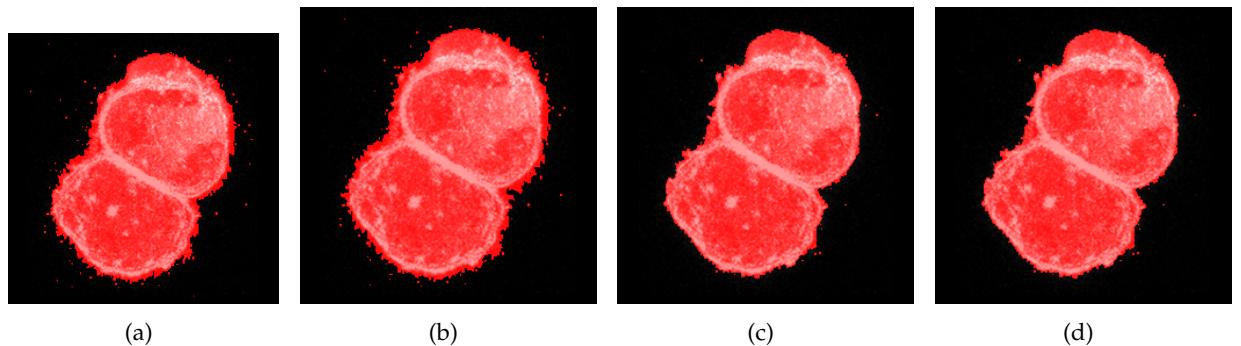


FIGURE 7.6: Interactive segmentation of Image 2 in sample set (Figure B.1). **(a)** Eriksson *et al.* [26]. **(b)** Boykov *et al.* [25]. **(c)** Proposed (No hard constraints) **(d)** Proposed with hard constraints.

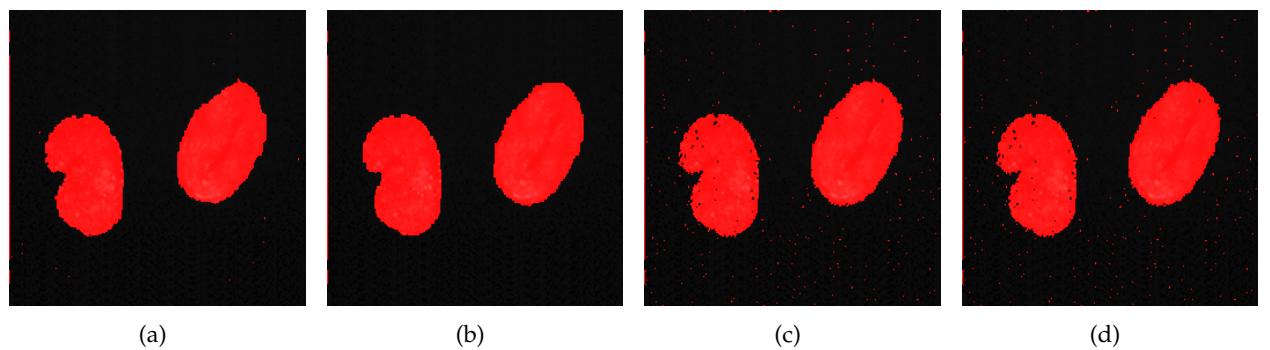


FIGURE 7.7: Interactive segmentation of Image 3 in sample set (Figure B.1). **(a)** Eriksson *et al.* [26]. **(b)** Boykov *et al.* [25]. **(c)** Proposed (No hard constraints) **(d)** Proposed with hard constraints.

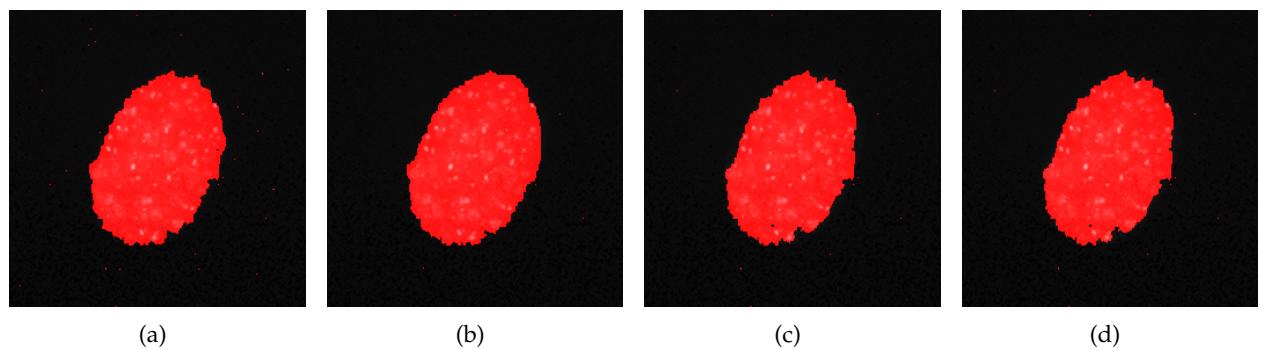


FIGURE 7.8: Interactive segmentation of Image 4 in sample set (Figure B.1). **(a)** Eriksson *et al.* [26]. **(b)** Boykov *et al.* [25]. **(c)** Proposed (No hard constraints) **(d)** Proposed with hard constraints.

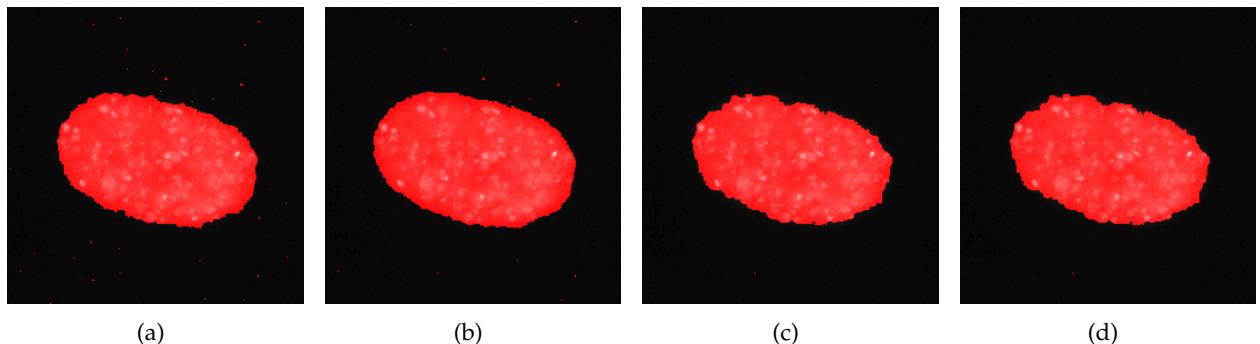


FIGURE 7.9: Interactive segmentation of Image 5 in sample set (Figure B.1). **(a)** Eriksson *et al.* [26]. **(b)** Boykov *et al.* [25]. **(c)** Proposed (No hard constraints) **(d)** Proposed with hard constraints.

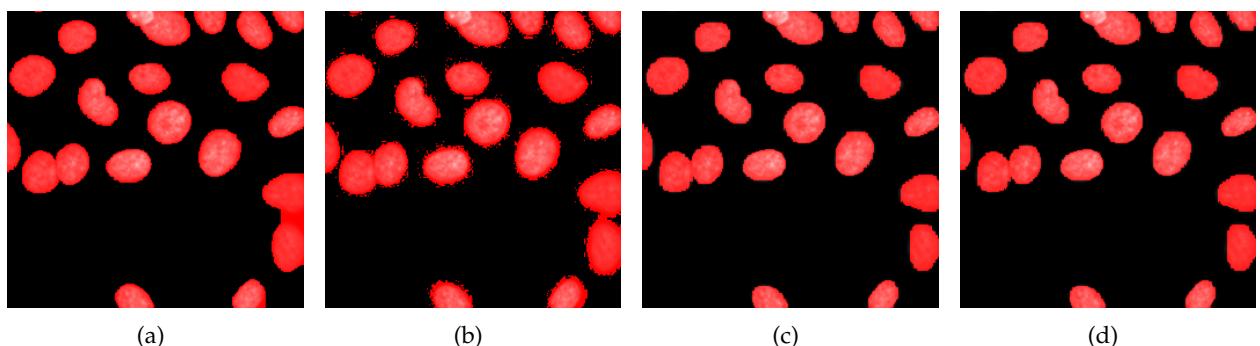


FIGURE 7.10: Interactive segmentation of Image 6 in sample set (Figure B.1). **(a)** Eriksson *et al.* [26]. **(b)** Boykov *et al.* [25]. **(c)** Proposed (No hard constraints) **(d)** Proposed with hard constraints.

7.4 Experimental Results

We compared the segmentation results for the optimal parameters found in Section 7.3. We used a common seed for all images which is shown in Figures 7.11(a), 7.12(a), 7.13(a), 7.14(a), 7.15(a), 7.16(a), 7.17(a), 7.18(a), 7.19(a), 7.20(a), 7.21(a), 7.22(a), 7.23(a), 7.24(a), 7.25(a), 7.26(a), 7.27(a), 7.28(a), 7.29(a), 7.30(a), 7.31(a), 7.32(a), 7.33(a), 7.34(a) and 7.35(a). From the seeds, we calculated a probability distribution based on EMGMM for $k = 2$, each for the background and the object seeds.

The segmentation results were compared using the same method as in Section 6.3. That is, we have compiled a label for label comparison on the segmentation mask and the ground truth. The parameter settings with the corresponding segmentation results are shown in Table 7.1 including the efficiency measures *precision*, *recall*, *accuracy* and *Matthews Correlation Coefficient (MCC)*. The overview results are shown in Table 7.2. For each image, we have highlighted the method which performs the best in blue and the worst in red.

We differentiate between methods on the same image as follows:

[imageno]-[method],

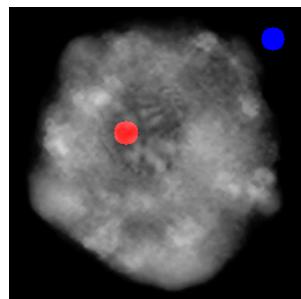
where *imageno* goes from 1 to 25 and *method* is defined as follows:

e - using the optimal parameter setting for the weighting described by Eriksson *et al.* [26].

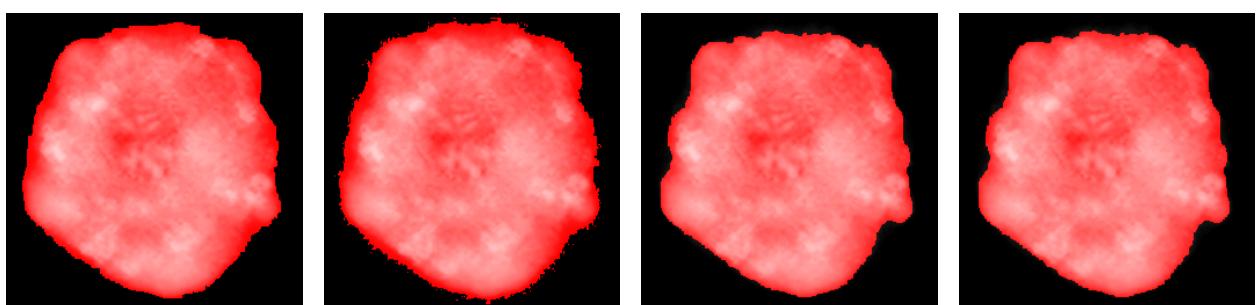
b - using the optimal parameter setting for the weighting described by Boykov *et al.* [25].

r - Proposed method with optimal parameters found on the sample set.

rh - Proposed method with hard constraints for optimal parameters found on the sample set.

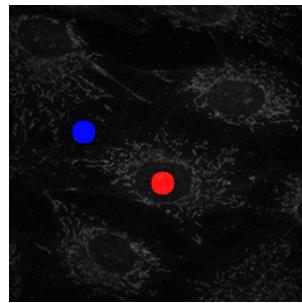


(a) Image 1 with seeds.



(b) Eriksson *et al.* [26] $\lambda = 10$, (c) Boykov *et al.* [25] $\lambda = 7.5$, (d) Proposed $\lambda = 0.5$, $\sigma = 7.5$, $\sigma_W = 7.5$, $\sigma_R = 1$. (e) Proposed with hard constraints $\lambda = 0.5$, $\sigma = 7.5$.

FIGURE 7.11: Image 1 from test set Appendix B interactive segmentation results.



(a) Image 2 with seeds.

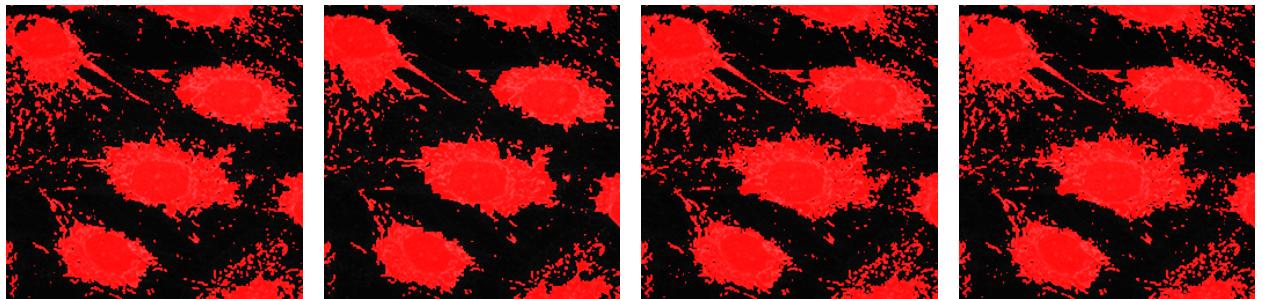
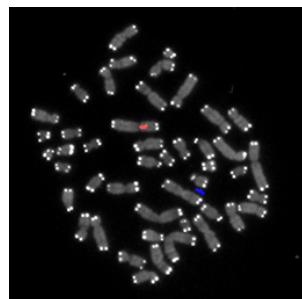
(b) Eriksson *et al.* [26] $\lambda = 10$, (c) Boykov *et al.* [25] $\lambda = 7.5$, (d) Proposed $\lambda = 0.5$, $\sigma = 7.5$.
 $\sigma_W = 7.5$, $\sigma_R = 1$. (e) Proposed with hard constraints $\lambda = 0.5$, $\sigma = 7.5$.

FIGURE 7.12: Image 2 from test set Appendix B interactive segmentation results.



(a) Image 3 with seeds.

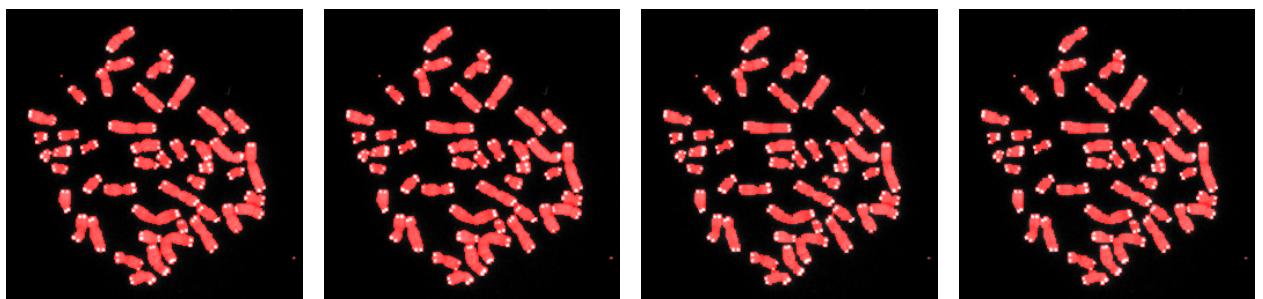
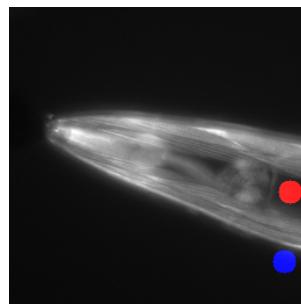
(b) Eriksson *et al.* [26] $\lambda = 10$, (c) Boykov *et al.* [25] $\lambda = 7.5$, (d) Proposed $\lambda = 0.5$, $\sigma = 7.5$.
 $\sigma_W = 7.5$, $\sigma_R = 1$. (e) Proposed with hard constraints $\lambda = 0.5$, $\sigma = 7.5$.

FIGURE 7.13: Image 3 from test set Appendix B interactive segmentation results.



(a) Image 4 with seeds.

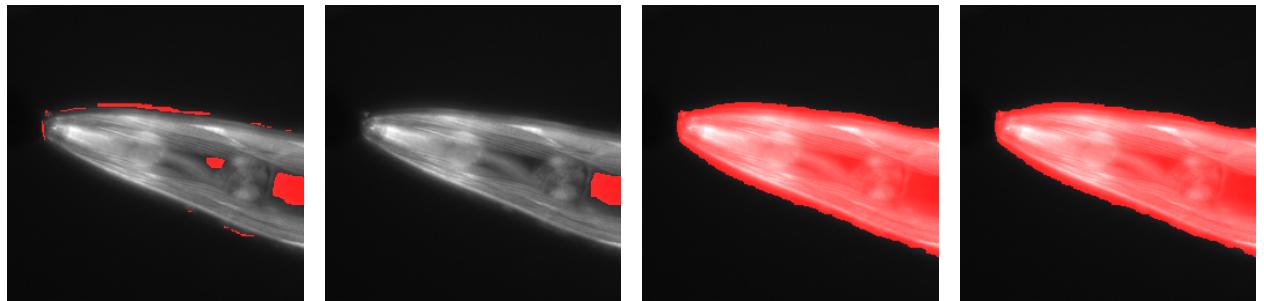
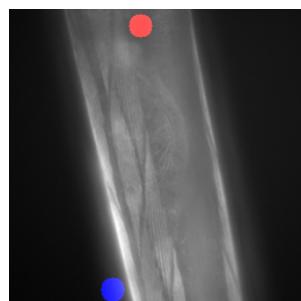
(b) Eriksson *et al.* [26] $\lambda = 10$, (c) Boykov *et al.* [25] $\lambda = 7.5$, (d) Proposed $\lambda = 0.5$, $\sigma = 7.5$.
 $\sigma_W = 7.5$, $\sigma_R = 1$. (e) Proposed with hard constraints $\lambda = 0.5$, $\sigma = 7.5$.

FIGURE 7.14: Image 4 from test set Appendix B interactive segmentation results.



(a) Image 5 with seeds.

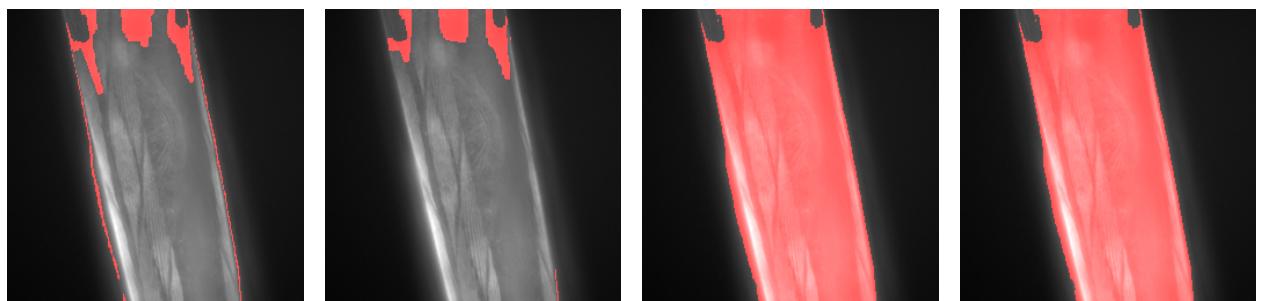
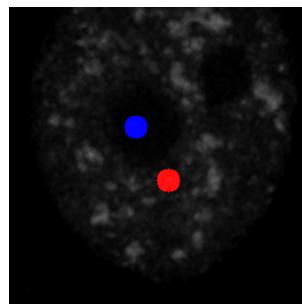
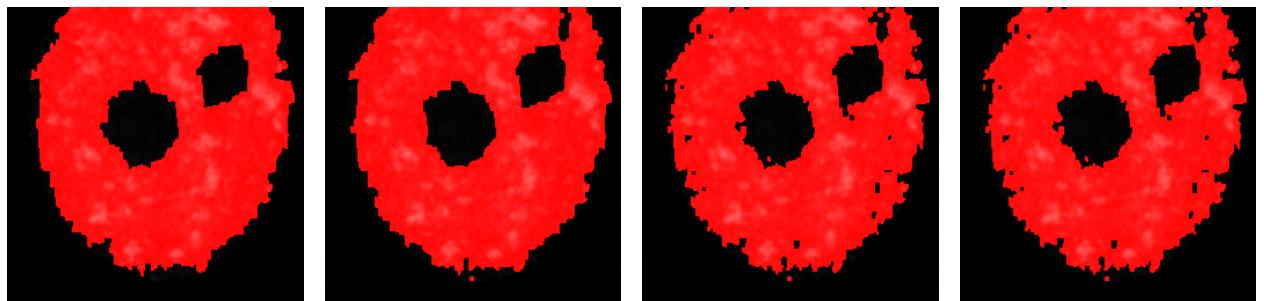
(b) Eriksson *et al.* [26] $\lambda = 10$, (c) Boykov *et al.* [25] $\lambda = 7.5$, (d) Proposed $\lambda = 0.5$, $\sigma = 7.5$.
 $\sigma_W = 7.5$, $\sigma_R = 1$. (e) Proposed with hard constraints $\lambda = 0.5$, $\sigma = 7.5$.

FIGURE 7.15: Image 5 from test set Appendix B interactive segmentation results.

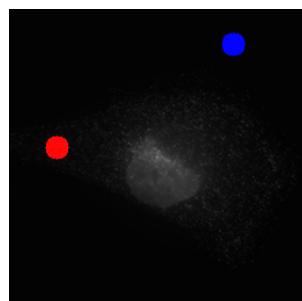


(a) Image 6 with seeds.

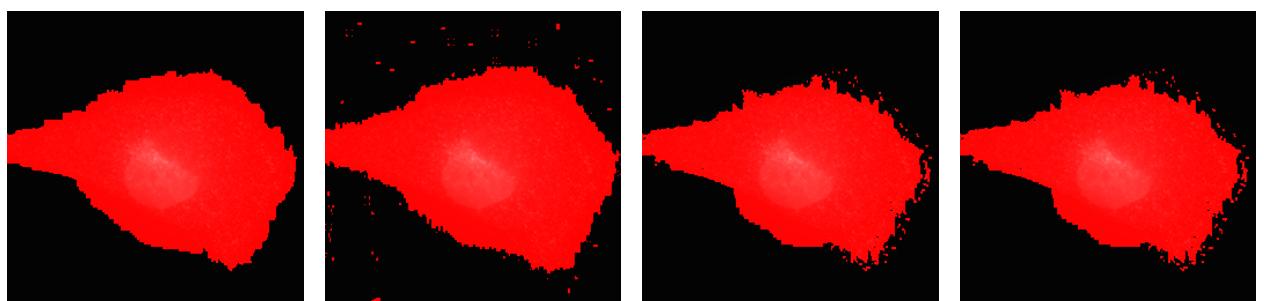


(b) Eriksson *et al.* [26] $\lambda = 10$, (c) Boykov *et al.* [25] $\lambda = 7.5$, (d) Proposed $\lambda = 0.5$, $\sigma = 7.5$.
 $\sigma_W = 7.5$, $\sigma_R = 1$. (e) Proposed with hard constraints $\lambda = 0.5$, $\sigma = 7.5$.

FIGURE 7.16: Image 6 from test set Appendix B interactive segmentation results.

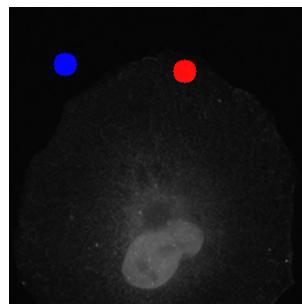


(a) Image 7 with seeds.

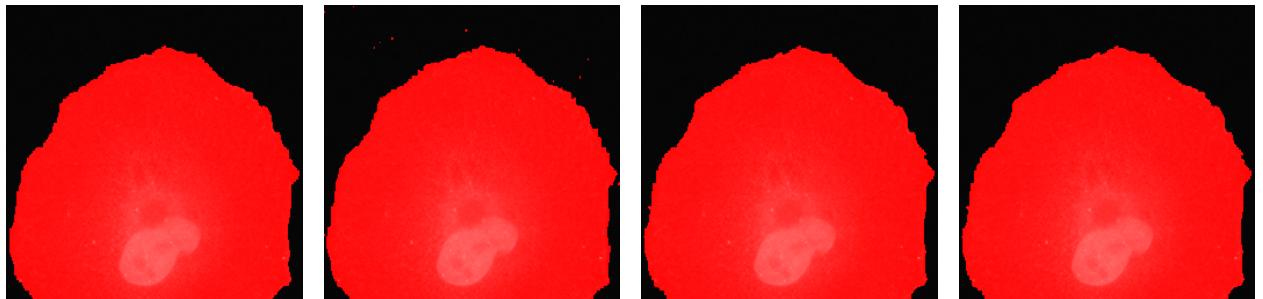


(b) Eriksson *et al.* [26] $\lambda = 10$, (c) Boykov *et al.* [25] $\lambda = 7.5$, (d) Proposed $\lambda = 0.5$, $\sigma = 7.5$.
 $\sigma_W = 7.5$, $\sigma_R = 1$. (e) Proposed with hard constraints $\lambda = 0.5$, $\sigma = 7.5$.

FIGURE 7.17: Image 7 from test set Appendix B interactive segmentation results.

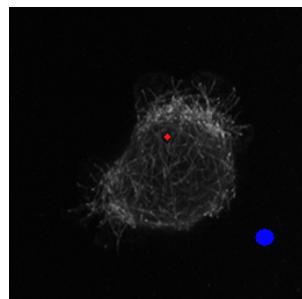


(a) Image 8 with seeds.

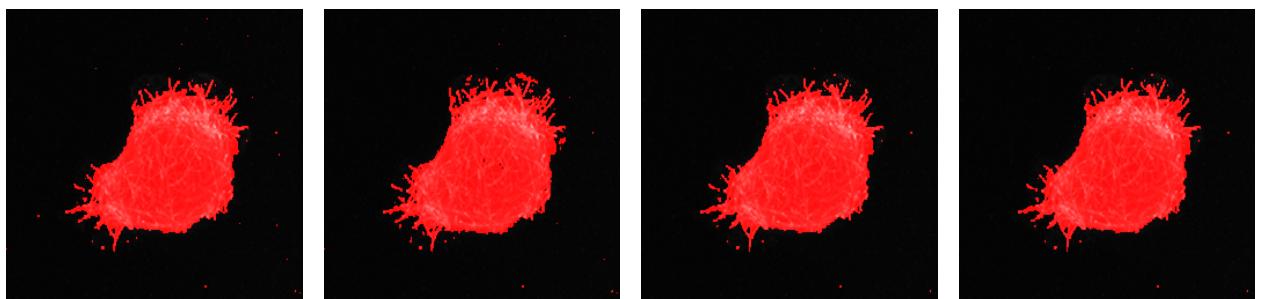


(b) Eriksson *et al.* [26] $\lambda = 10$, (c) Boykov *et al.* [25] $\lambda = 7.5$, (d) Proposed $\lambda = 0.5$, $\sigma = 7.5$.
 $\sigma_W = 7.5$, $\sigma_R = 1$. (e) Proposed with hard constraints $\lambda = 0.5$, $\sigma = 7.5$.

FIGURE 7.18: Image 8 from test set Appendix B interactive segmentation results.

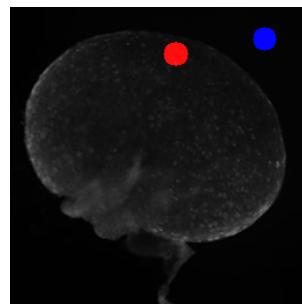


(a) Image 9 with seeds.



(b) Eriksson *et al.* [26] $\lambda = 10$, (c) Boykov *et al.* [25] $\lambda = 7.5$, (d) Proposed $\lambda = 0.5$, $\sigma = 7.5$.
 $\sigma_W = 7.5$, $\sigma_R = 1$. (e) Proposed with hard constraints $\lambda = 0.5$, $\sigma = 7.5$.

FIGURE 7.19: Image 9 from test set Appendix B interactive segmentation results.



(a) Image 10 with seeds.

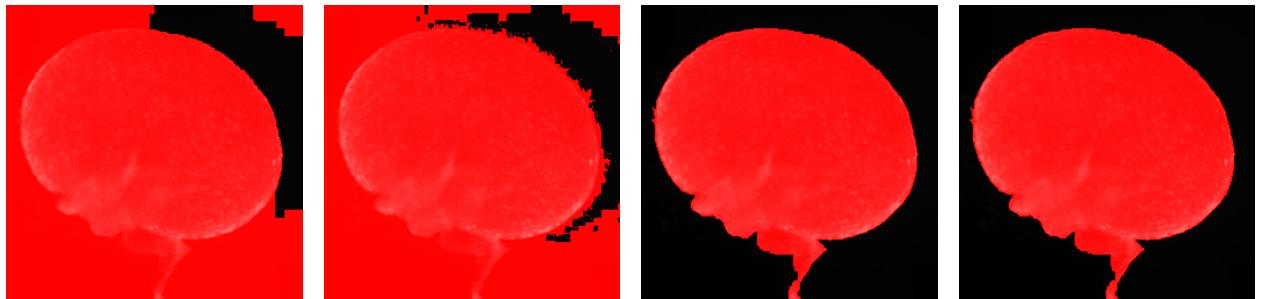
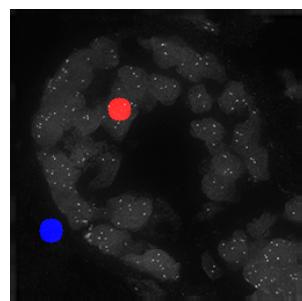
(b) Eriksson *et al.* [26] $\lambda = 10$, (c) Boykov *et al.* [25] $\lambda = 7.5$, (d) Proposed $\lambda = 0.5$, $\sigma = 5$.
 $\sigma_W = 7.5$, $\sigma_R = 1$. (e) Proposed with hard constraints $\lambda = 0.5$, $\sigma = 7.5$.

FIGURE 7.20: Image 10 from test set Appendix B interactive segmentation results.



(a) Image 11 with seeds.

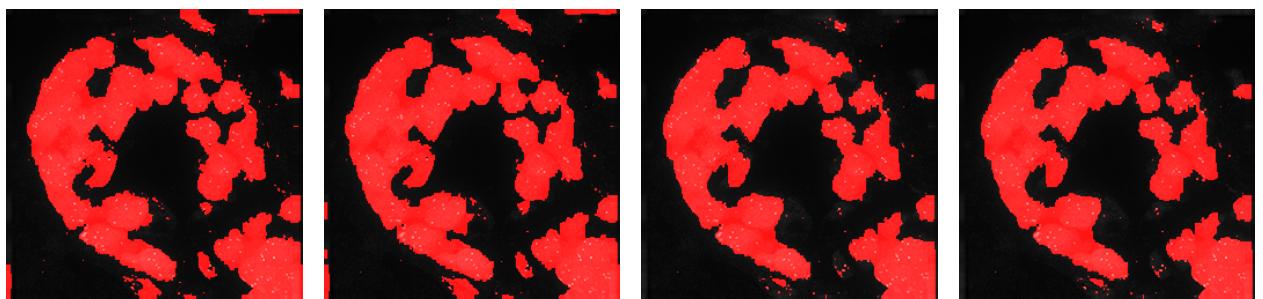
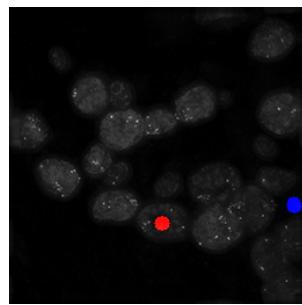
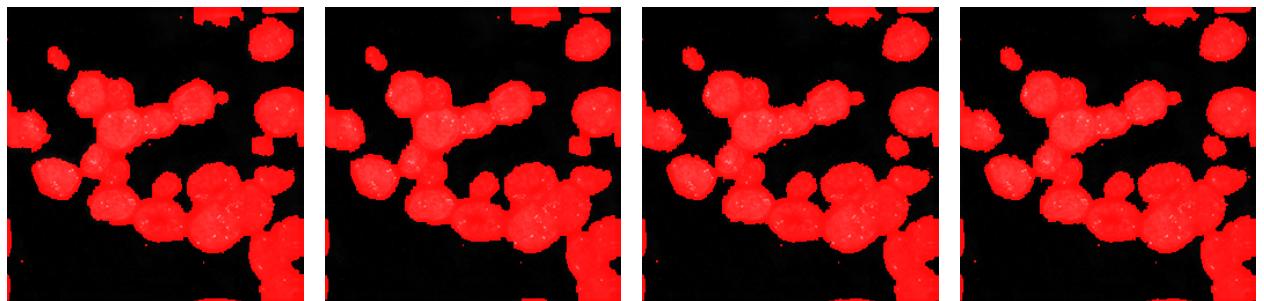
(b) Eriksson *et al.* [26] $\lambda = 10$, (c) Boykov *et al.* [25] $\lambda = 7.5$, (d) Proposed $\lambda = 0.5$, $\sigma = 5$.
 $\sigma_W = 7.5$, $\sigma_R = 1$. (e) Proposed with hard constraints $\lambda = 0.5$, $\sigma = 7.5$.

FIGURE 7.21: Image 11 from test set Appendix B interactive segmentation results.

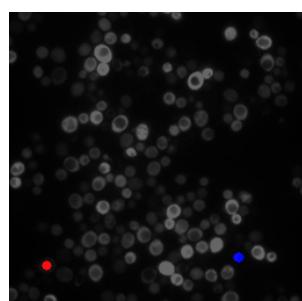


(a) Image 12 with seeds.

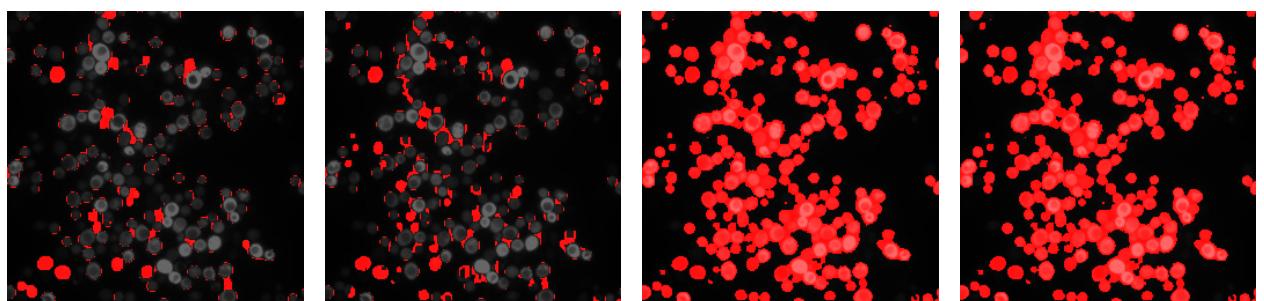


(b) Eriksson *et al.* [26] $\lambda = 10$, (c) Boykov *et al.* [25] $\lambda = 7.5$, (d) Proposed $\lambda = 0.5$, $\sigma = 7.5$.
 $\sigma_W = 7.5$, $\sigma_R = 1$. (e) Proposed with hard constraints $\lambda = 0.5$, $\sigma = 7.5$.

FIGURE 7.22: Image 12 from test set Appendix B interactive segmentation results.

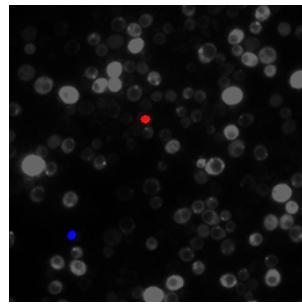


(a) Image 13 with seeds.



(b) Eriksson *et al.* [26] $\lambda = 10$, (c) Boykov *et al.* [25] $\lambda = 7.5$, (d) Proposed $\lambda = 0.5$, $\sigma = 7.5$.
 $\sigma_W = 7.5$, $\sigma_R = 1$. (e) Proposed with hard constraints $\lambda = 0.5$, $\sigma = 7.5$.

FIGURE 7.23: Image 13 from test set Appendix B interactive segmentation results.



(a) Image 14 with seeds.

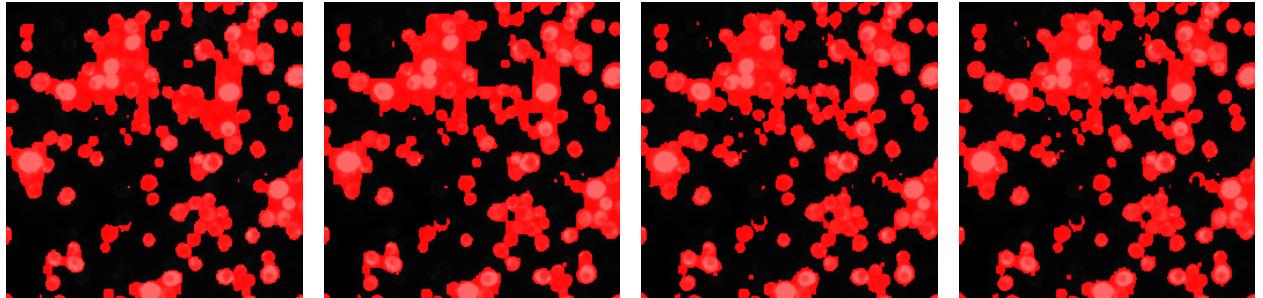
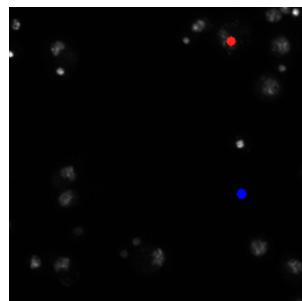
(b) Eriksson *et al.* [26] $\lambda = 10$, (c) Boykov *et al.* [25] $\lambda = 7.5$, (d) Proposed $\lambda = 0.5$, $\sigma = 7.5$.
 $\sigma_W = 7.5$, $\sigma_R = 1$. (e) Proposed with hard constraints $\lambda = 0.5$, $\sigma = 7.5$.

FIGURE 7.24: Image 14 from test set Appendix B interactive segmentation results.



(a) Image 15 with seeds.

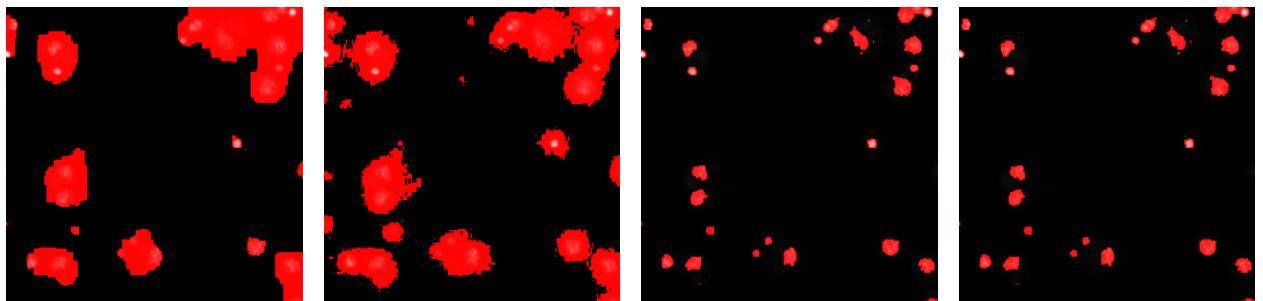
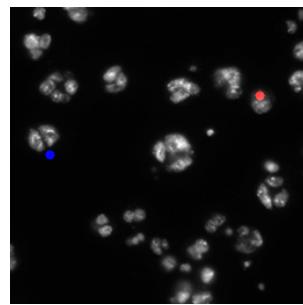
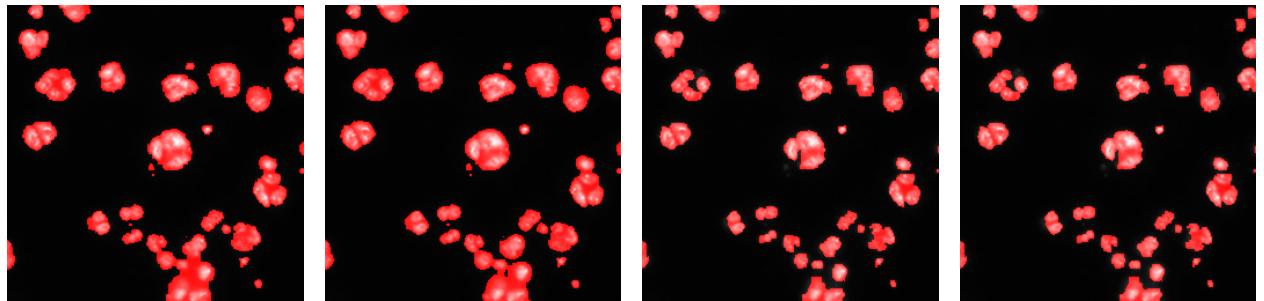
(b) Eriksson *et al.* [26] $\lambda = 10$, (c) Boykov *et al.* [25] $\lambda = 7.5$, (d) Proposed $\lambda = 0.5$, $\sigma = 7.5$.
 $\sigma_W = 7.5$, $\sigma_R = 1$. (e) Proposed with hard constraints $\lambda = 0.5$, $\sigma = 7.5$.

FIGURE 7.25: Image 15 from test set Appendix B interactive segmentation results.

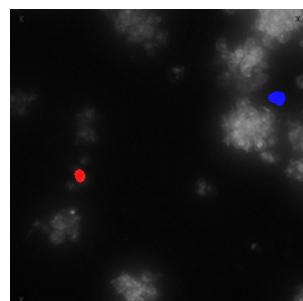


(a) Image 16 with seeds.

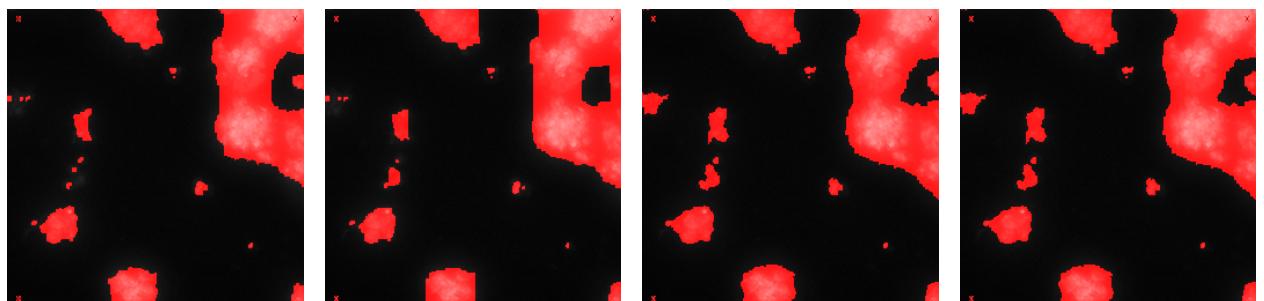


(b) Eriksson *et al.* [26] $\lambda = 10$, (c) Boykov *et al.* [25] $\lambda = 7.5$, (d) Proposed $\lambda = 0.5$, $\sigma = 7.5$.
 $\sigma_W = 7.5$, $\sigma_R = 1$. (e) Proposed with hard constraints $\lambda = 0.5$, $\sigma = 7.5$.

FIGURE 7.26: Image 16 from test set Appendix B interactive segmentation results.

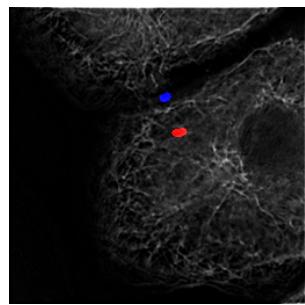


(a) Image 17 with seeds.



(b) Eriksson *et al.* [26] $\lambda = 10$, (c) Boykov *et al.* [25] $\lambda = 7.5$, (d) Proposed $\lambda = 0.5$, $\sigma = 7.5$.
 $\sigma_W = 7.5$, $\sigma_R = 1$. (e) Proposed with hard constraints $\lambda = 0.5$, $\sigma = 7.5$.

FIGURE 7.27: Image 17 from test set Appendix B interactive segmentation results.



(a) Image 18 with seeds.

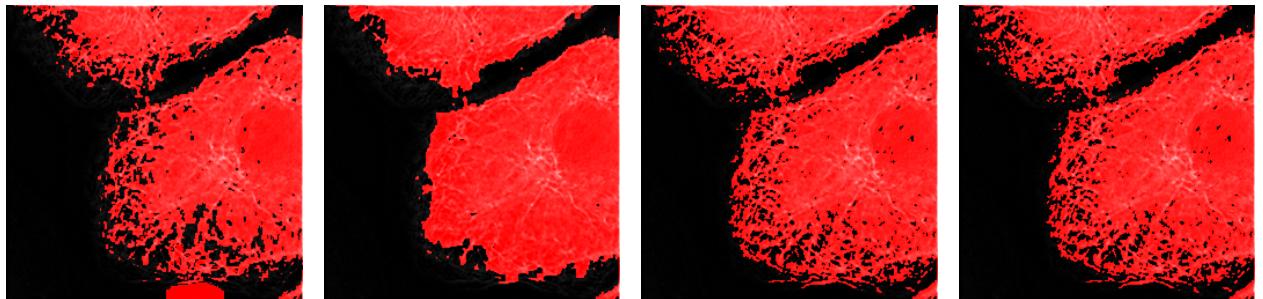
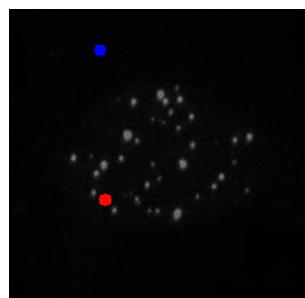
(b) Eriksson *et al.* [26] $\lambda = 10$, (c) Boykov *et al.* [25] $\lambda = 7.5$, (d) Proposed $\lambda = 0.5$, $\sigma = 7.5$.
 $\sigma_W = 7.5$, $\sigma_R = 1$. (e) Proposed with hard constraints $\lambda = 0.5$, $\sigma = 7.5$.

FIGURE 7.28: Image 18 from test set Appendix B interactive segmentation results.



(a) Image 19 with seeds.

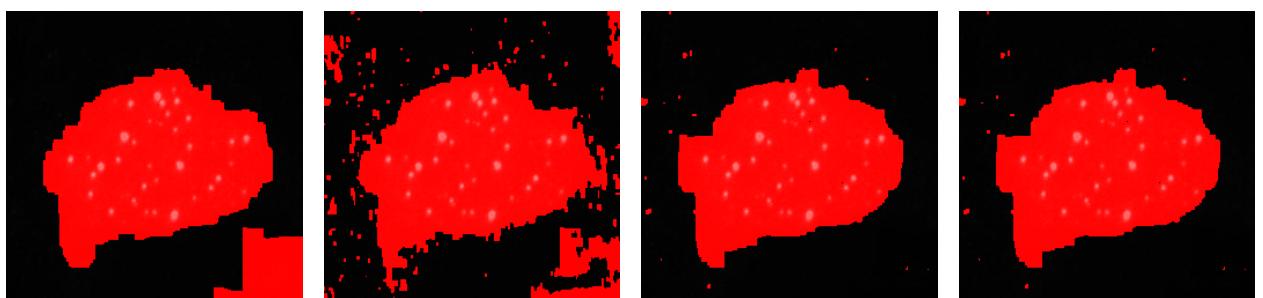
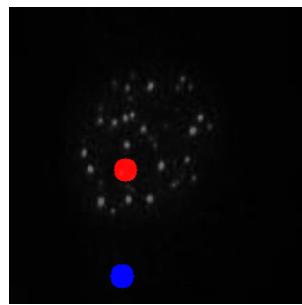
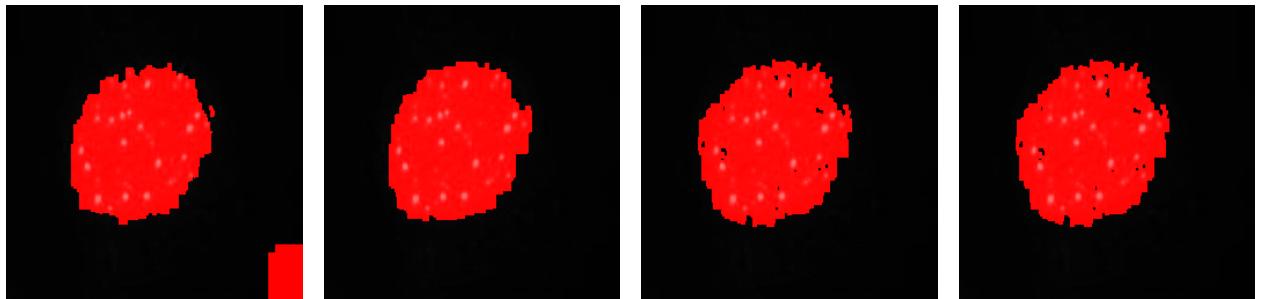
(b) Eriksson *et al.* [26] $\lambda = 10$, (c) Boykov *et al.* [25] $\lambda = 7.5$, (d) Proposed $\lambda = 0.5$, $\sigma = 7.5$.
 $\sigma_W = 7.5$, $\sigma_R = 1$. (e) Proposed with hard constraints $\lambda = 0.5$, $\sigma = 7.5$.

FIGURE 7.29: Image 19 from test set Appendix B interactive segmentation results.

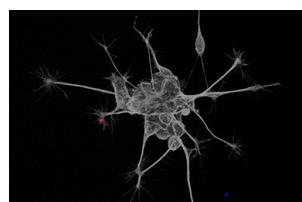


(a) Image 20 with seeds.

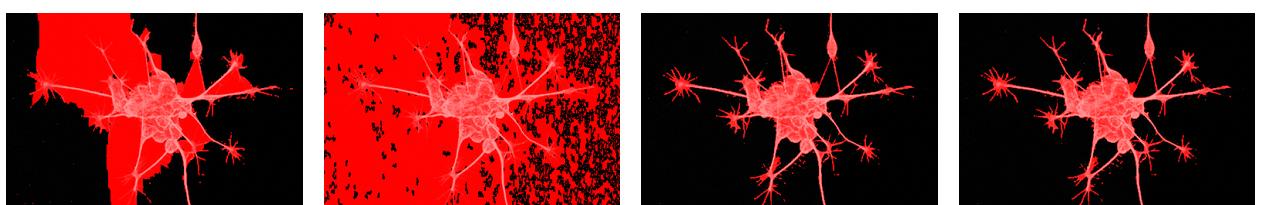


(b) Eriksson *et al.* [26] $\lambda = 10$, (c) Boykov *et al.* [25] $\lambda = 7.5$, (d) Proposed $\lambda = 0.5$, $\sigma = 7.5$.
 $\sigma_W = 7.5$, $\sigma_R = 1$. (e) Proposed with hard constraints $\lambda = 0.5$, $\sigma = 7.5$.

FIGURE 7.30: Image 20 from test set Appendix B interactive segmentation results.

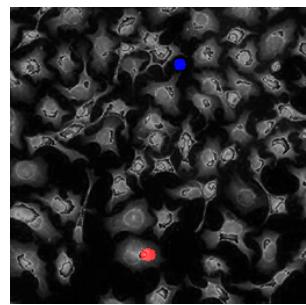


(a) Image 21 with seeds.



(b) Eriksson *et al.* [26] $\lambda = 10$, (c) Boykov *et al.* [25] $\lambda = 7.5$, (d) Proposed $\lambda = 0.5$, $\sigma = 7.5$.
 $\sigma_W = 7.5$, $\sigma_R = 1$. (e) Proposed with hard constraints $\lambda = 0.5$, $\sigma = 7.5$.

FIGURE 7.31: Image 21 from test set Appendix B interactive segmentation results.



(a) Image 22 with seeds.

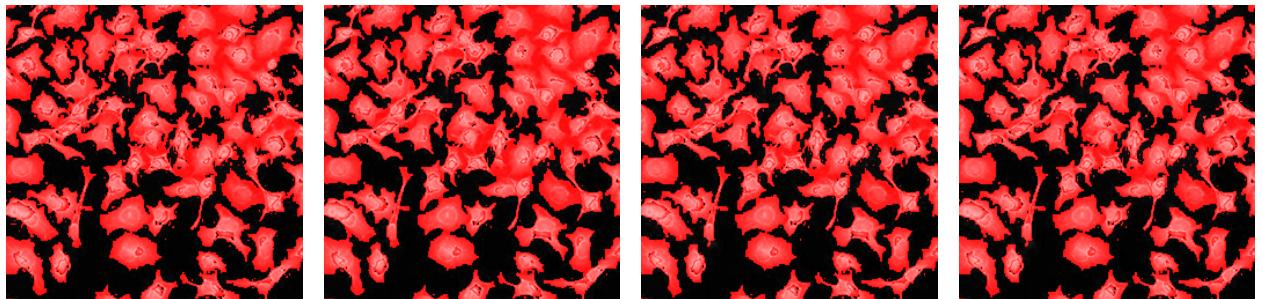
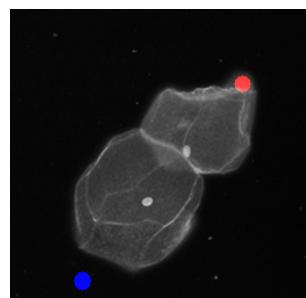
(b) Eriksson *et al.* [26] $\lambda = 10$, (c) Boykov *et al.* [25] $\lambda = 7.5$, (d) Proposed $\lambda = 0.5$, $\sigma = 7.5$.
(e) Proposed with hard constraints $\lambda = 0.5$, $\sigma = 7.5$.

FIGURE 7.32: Image 22 from test set Appendix B interactive segmentation results.



(a) Image 23 with seeds.

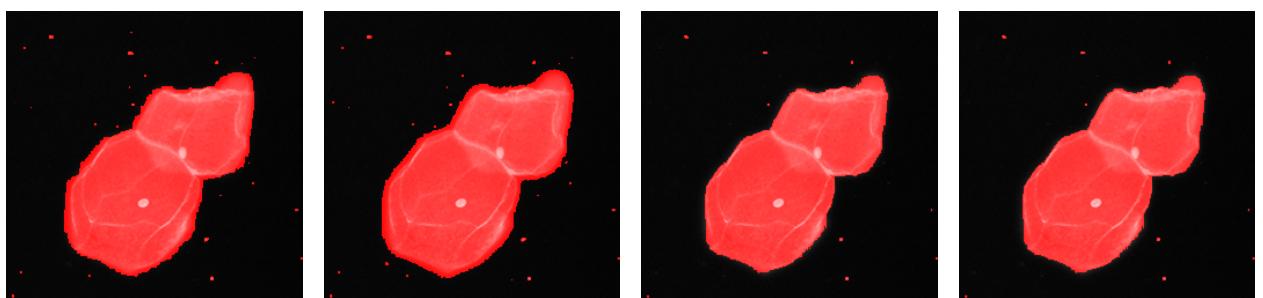
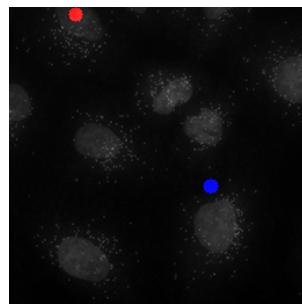
(b) Eriksson *et al.* [26] $\lambda = 10$, (c) Boykov *et al.* [25] $\lambda = 7.5$, (d) Proposed $\lambda = 0.5$, $\sigma = 7.5$.
(e) Proposed with hard constraints $\lambda = 0.5$, $\sigma = 7.5$.

FIGURE 7.33: Image 23 from test set Appendix B interactive segmentation results.



(a) Image 24 with seeds.

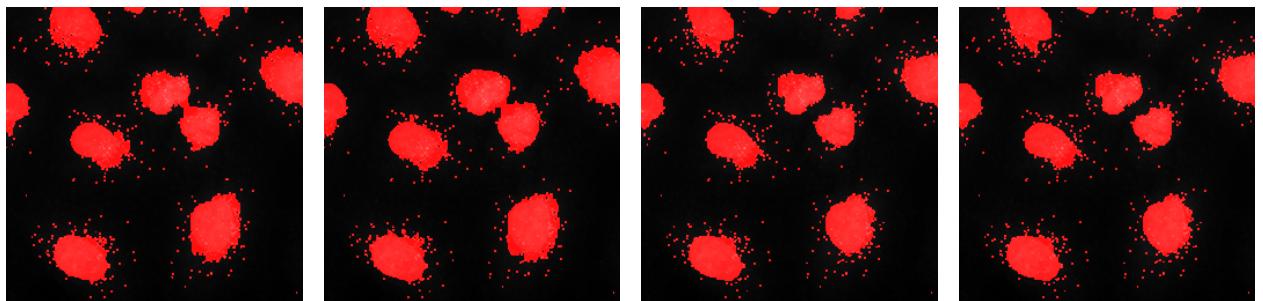
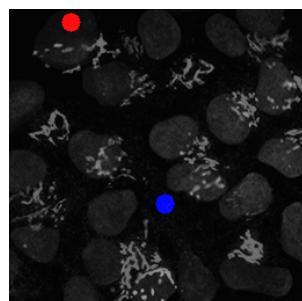
(b) Eriksson *et al.* [26] $\lambda = 10$, (c) Boykov *et al.* [25] $\lambda = 7.5$, (d) Proposed $\lambda = 0.5$, $\sigma = 7.5$.
 $\sigma_W = 7.5$, $\sigma_R = 1$. (e) Proposed with hard constraints $\lambda = 0.5$, $\sigma = 7.5$.

FIGURE 7.34: Image 24 from test set Appendix B interactive segmentation results.



(a) Image 25 with seeds.

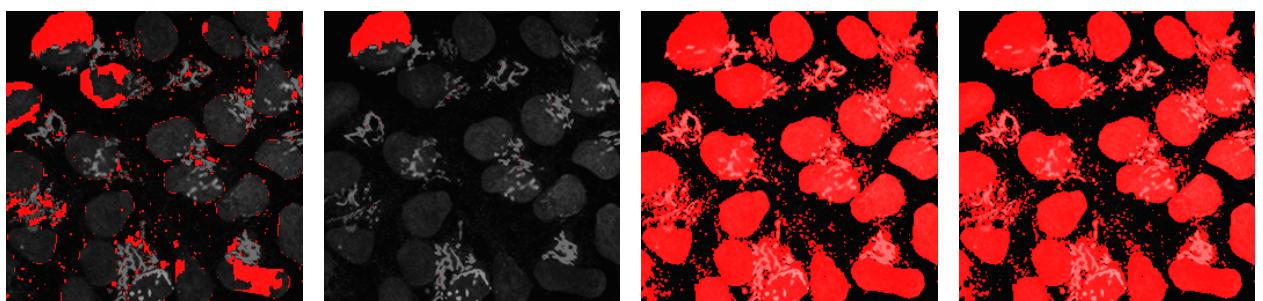
(b) Eriksson *et al.* [26] $\lambda = 10$, (c) Boykov *et al.* [25] $\lambda = 7.5$, (d) Proposed $\lambda = 0.5$, $\sigma = 7.5$.
 $\sigma_W = 7.5$, $\sigma_R = 1$. (e) Proposed with hard constraints $\lambda = 0.5$, $\sigma = 7.5$.

FIGURE 7.35: Image 25 from test set Appendix B interactive segmentation results.

TABLE 7.1: Interactive Segmentation Results.

Image	TP	TN	FP	FN	Precision	Recall	Accuracy	MCC
1-e	39901	24637	879	119	0.978445	0.997026	0.984772	0.968092
1-b	40020	23280	2236	0	0.947084	1.000000	0.965881	0.929565
1-r	38125	25489	27	1895	0.999292	0.952649	0.970673	0.940780
1-rh	38125	25489	27	1895	0.999292	0.952649	0.970673	0.940780
2-e	20880	40848	2742	1066	0.883922	0.951426	0.941895	0.873376
2-b	20417	40801	2789	1529	0.879816	0.930329	0.934113	0.854946
2-r	21798	38965	4625	148	0.824963	0.993256	0.927170	0.853529
2-rh	21798	38965	4625	148	0.824963	0.993256	0.927170	0.853529
3-e	10043	55027	404	62	0.961329	0.993864	0.992889	0.973300
3-b	10105	55056	375	0	0.964218	1.000000	0.994278	0.978619
3-r	9387	55215	216	718	0.977507	0.928946	0.985748	0.944652
3-rh	9387	55215	216	718	0.977507	0.928946	0.985748	0.944652
4-e	726	49547	396	14867	0.647059	0.046559	0.767105	0.126807
4-b	606	49943	0	14987	1.000000	0.038864	0.771317	0.172896
4-r	15593	47243	2700	0	0.852403	1.000000	0.958801	0.897953
4-rh	15593	47246	2697	0	0.852542	1.000000	0.958847	0.898056
5-e	2001	40188	679	22668	0.746642	0.081114	0.643753	0.157788
5-b	1709	40781	86	22960	0.952089	0.069277	0.648346	0.199395
5-r	24112	37964	2903	557	0.892541	0.977421	0.947205	0.892121
5-rh	24112	37968	2899	557	0.892673	0.977421	0.947266	0.892237
6-e	36339	25698	57	3442	0.998434	0.913476	0.946609	0.895655
6-b	36441	25708	47	3340	0.998712	0.916040	0.948318	0.898843
6-r	35795	25670	85	3986	0.997631	0.899801	0.937881	0.879705
6-rh	35795	25670	85	3986	0.997631	0.899801	0.937881	0.879705
7-e	18125	39494	7857	60	0.697598	0.996701	0.879196	0.760449
7-b	18184	37646	9705	1	0.652013	0.999945	0.851898	0.719945
7-r	17865	42666	4685	320	0.792239	0.982403	0.923630	0.832668
7-rh	17865	42666	4685	320	0.792239	0.982403	0.923630	0.832668
8-e	43065	21953	205	313	0.995262	0.992784	0.992096	0.982368
8-b	43262	21549	609	116	0.986118	0.997326	0.988937	0.975287
8-r	43148	21880	278	230	0.993598	0.994698	0.992249	0.982674
8-rh	43148	21880	278	230	0.993598	0.994698	0.992249	0.982674

9-e	12118	52960	250	208	0.979787	0.983125	0.993011	0.977150
9-b	12263	52689	521	63	0.959246	0.994889	0.991089	0.971480
9-r	12085	53035	175	241	0.985726	0.980448	0.993652	0.979179
9-rh	12085	53035	175	241	0.985726	0.980448	0.993652	0.979179
10-e	34360	7816	23332	28	0.595577	0.999186	0.643555	0.384800
10-b	34388	6963	24185	0	0.587096	1.000000	0.630966	0.362275
10-r	34297	31105	43	91	0.998748	0.997354	0.997955	0.995902
10-rh	34297	31105	43	91	0.998748	0.997354	0.997955	0.995902
11-e	22982	38696	2156	1702	0.914233	0.931048	0.941132	0.875182
11-b	23205	38353	2499	1479	0.902778	0.940083	0.939301	0.872254
11-r	21035	39776	1076	3649	0.951336	0.852171	0.927902	0.846315
11-rh	21035	39776	1076	3649	0.951336	0.852171	0.927902	0.846315
12-e	19092	40749	5212	483	0.785550	0.975326	0.913101	0.816694
12-b	19172	40683	5278	403	0.784131	0.979413	0.913315	0.818206
12-r	19243	40656	5305	332	0.783893	0.983040	0.913986	0.820421
12-rh	19243	40656	5305	332	0.783893	0.983040	0.913986	0.820421
13-e	877	53053	1896	9710	0.316264	0.082837	0.822906	0.088365
13-b	1249	52310	2639	9338	0.321245	0.117975	0.817245	0.108974
13-r	10343	44375	10574	244	0.494478	0.976953	0.834930	0.619385
13-rh	10343	44375	10574	244	0.494478	0.976953	0.834930	0.619385
14-e	13304	42783	9112	337	0.593505	0.975295	0.855820	0.684384
14-b	13517	41200	10695	124	0.558277	0.990910	0.834915	0.660146
14-r	13518	41344	10551	123	0.561635	0.990983	0.837128	0.663360
14-rh	13518	41344	10551	123	0.561635	0.990983	0.837128	0.663360
15-e	3322	53662	8504	48	0.280906	0.985757	0.869507	0.487566
15-b	3370	50812	11354	0	0.228878	1.000000	0.826752	0.432523
15-r	2439	62155	11	931	0.995510	0.723739	0.985626	0.842398
15-rh	2439	62155	11	931	0.995510	0.723739	0.985626	0.842398
16-e	7439	55023	3070	4	0.707869	0.999463	0.953094	0.818543
16-b	7443	54644	3449	0	0.683346	1.000000	0.947372	0.801733
16-r	7320	57448	645	123	0.919021	0.983474	0.988281	0.944220
16-rh	7320	57448	645	123	0.919021	0.983474	0.988281	0.944220
17-e	8541	51662	4325	1008	0.663843	0.894439	0.918625	0.725841
17-b	8611	50915	5072	938	0.629321	0.901770	0.908295	0.704143

17-r	9180	50237	5750	369	0.614869	0.961357	0.906631	0.722288
17-rh	9180	50237	5750	369	0.614869	0.961357	0.906631	0.722288
18-e	31177	27384	991	5984	0.969193	0.838971	0.893570	0.796921
18-b	34038	26721	1654	3123	0.953659	0.915960	0.927109	0.853331
18-r	35776	27921	454	1385	0.987469	0.962730	0.971939	0.943464
18-rh	35776	27921	454	1385	0.987469	0.962730	0.971939	0.943464
19-e	15551	39888	10097	0	0.606324	1.000000	0.845932	0.695591
19-b	15551	38006	11979	0	0.564875	1.000000	0.817215	0.655364
19-r	15545	43925	6060	6	0.719509	0.999614	0.907440	0.794909
19-rh	15545	43925	6060	6	0.719509	0.999614	0.907440	0.794909
20-e	12134	50512	1907	983	0.864183	0.925059	0.955902	0.866613
20-b	12496	51789	630	621	0.952004	0.952657	0.980911	0.940397
20-r	12751	51106	1313	366	0.906641	0.972097	0.974380	0.922984
20-rh	12751	51106	1313	366	0.906641	0.972097	0.974380	0.922984
21-e	11464	32586	21443	43	0.348376	0.996263	0.672150	0.456093
21-b	11498	5333	48696	9	0.191016	0.999218	0.256821	0.136162
21-r	11199	53750	279	308	0.975693	0.973234	0.991043	0.969032
21-rh	11199	53750	279	308	0.975693	0.973234	0.991043	0.969032
22-e	29332	27618	8472	114	0.775897	0.996129	0.868988	0.766566
22-b	29377	27242	8848	69	0.768528	0.997657	0.863937	0.759217
22-r	29359	28654	7436	87	0.797907	0.997045	0.885208	0.792940
22-rh	29359	28654	7436	87	0.797907	0.997045	0.885208	0.792940
23-e	15993	47936	1603	4	0.908900	0.999750	0.975479	0.937647
23-b	15997	47310	2229	0	0.877702	1.000000	0.965988	0.915538
23-r	15754	49292	247	243	0.984563	0.984810	0.992523	0.979741
23-rh	15754	49292	247	243	0.984563	0.984810	0.992523	0.979741
24-e	7791	53310	4365	70	0.640918	0.991095	0.932327	0.765182
24-b	7823	52446	5229	38	0.599372	0.995166	0.919632	0.735852
24-r	7705	54773	2902	156	0.726407	0.980155	0.953339	0.820244
24-rh	7705	54773	2902	156	0.726407	0.980155	0.953339	0.820244
25-e	4850	30953	1076	28657	0.818427	0.144746	0.546310	0.193738
25-b	1301	32010	19	32206	0.985606	0.038828	0.508286	0.136046
25-r	33203	28333	3696	304	0.899835	0.990927	0.938965	0.882349
25-rh	33203	28337	3692	304	0.899932	0.990927	0.939026	0.882461

TABLE 7.2: Overall Interactive Segmentation Efficiency.

Method	Precision		Recall		Accuracy	
	Mean	Std. Dev.	Mean	Std. Dev.	Mean	Std. Dev.
Eriksson <i>et al.</i> [26]	0.747138	0.212337	0.827658	0.331833	0.869989	0.123919
Boykov <i>et al.</i> [25]	0.757085	0.246706	0.831052	0.342374	0.846089	0.173522
Proposed	0.865337	0.147243	0.961572	0.059908	0.945771	0.046453
Proposed with Hard Constraints	0.865351	0.147245	0.961572	0.059908	0.945778	0.046454

From the accuracy and Matthews Correlation Coefficient measures shown in Table 7.1, it seems as if there is a close four-way "tug-o' war" between the methods. All methods seem to produce very good results over the test set. However, closer inspection reveals that the proposed method with hard constraints outperforms the least efficient weighting method by Boykov *et al.* [25] by 9.9689% and the second least efficient weighting method by Eriksson *et al.* [26] by 7.5789%. This is shown in Table 7.2. The graph of precision versus recall for all weighting methods is shown in Figure 7.36. As can be seen, the proposed method is much more stable, producing high precision and recall consistently. The accuracy variation over the test set is shown in Figure 7.37. This further shows the consistency and efficiency of the proposed weighting method with hard constraints over more general fluorescent images.

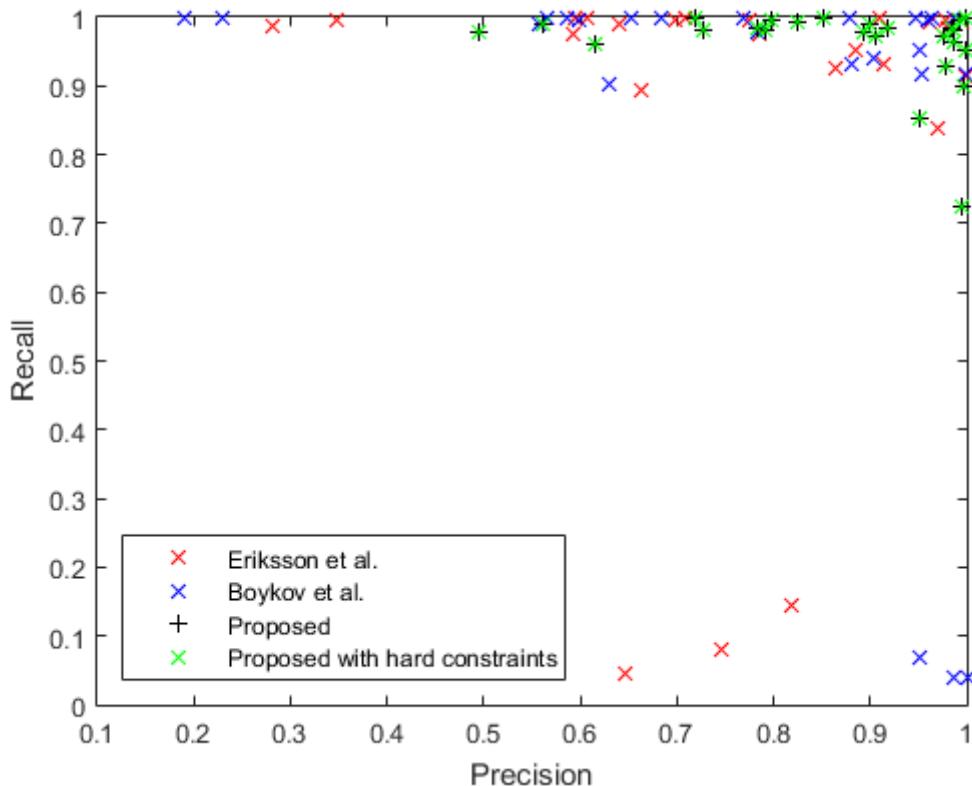


FIGURE 7.36: Interactive segmentation precision against recall over test set.

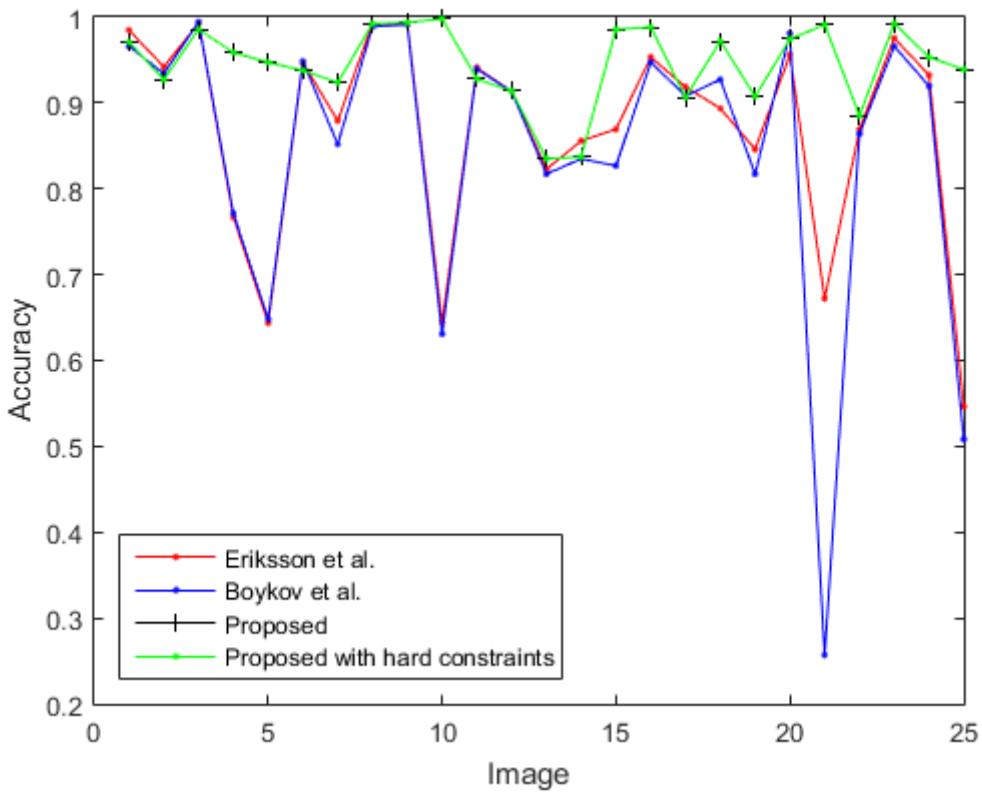


FIGURE 7.37: Interactive segmentation accuracy over test set.

7.5 Discussion

We have experimented, very thoroughly, with the two of the most common graph weighting systems and have derived optimal parameters settings for each system which work very well on general fluorescence images. The worst performing of these was the weighting system proposed by Boykov and Jolly [25] which still showed a very high average of 84.6089% classification accuracy being outperformed by the weighting system proposed by Eriksson *et al.* [26] by 2.39%. However, these two systems fall staggeringly short of the proposed scheme with hard constraints which showed an average classification accuracy at 94.5778%; a 7.5789% increase in comparison to the best competitor weighting system, which was proposed by Eriksson *et al.* [26].

7.6 Conclusion

It is not an ideal case to allow human interaction in segmentation; as such, it is a strongly sort after ideal to make segmentation entirely automatic. Consequently, automatic segmentation has come a long way; however, there still are cases where human involvement is necessary. The goal of increasing segmentation accuracy and diminishing human interaction is still prime.

We have presented a novel energy function for interactive segmentation of fluorescence images, specifically black background fluorescent images. The function accounts for the fact that the probability distribution intensity values do not exponentially decay as the intensity values move away

from the mean of the corresponding region i.e. background or object. We have replaced the exponential decay with a quadratic decay and rigorously compared the new energy function to two widely used energy functions which were proposed by Eriksson *et al.* [26] and Boykov and Jolly [25]. The proposed energy function can be implemented with and without hard constraints, and both showed very high performance in accuracy and consistency. The competitor energy functions exhibited a fairly high average segmentation accuracy of 84.6089% for Boykov and Jolly [25] and 86.9989% for Eriksson *et al.* [26]. However, they were trumped by the proposed energy function which boasted an average accuracy of 94.5778% with hard constraints and 94.5771% without hard constraints.

These results are subject to the seeds chosen. Given additional, the competitor algorithms could have exhibited much better performance. This comes at the cost of increased user interaction. From the graphs of the proposed energy functions, illustrated in Figure 7.1 and Figure 7.2, we can see that all that is required is for the foreground/object seed to mark the lower range of object and the background seed to mark the higher range of the background.

Chapter 8

Conclusion

This dissertation explores and investigates fluorescence image segmentation focussing specifically on the instances of the most common and widely used energy minimisation models that can be optimised via the graph cut framework. Fluorescence imaging covers a wide range of disciplines; we restrict our concern to biomedical and biological images.

A rudimentary understanding of fluorescence imaging concepts are covered in Chapter 2, along with factors that affect image quality, common segmentation techniques and what these segmentations are used for in the higher stages of analysis. In Chapter 3, we present the mathematical foundation of the energy models used in graph cut segmentation. Our own contributions and efforts are documented in the next three chapters.

Image Preparation - Chapter 5 Due to our current understanding of the physical process involved in capturing fluorescent images, we typically cannot produce high quality images. This is a problem for segmentation algorithms which will readily produce inaccuracies and segmentation artefacts. To abate the problems, fluorescence images are usually pre-process in some way to allow for more accurate segmentation results.

We propose a pre-processing segmentation scheme that pushes a great deal of the object data to the foreground which allows for enhanced visualisation and increased accuracy in segmentation. This is due to our novel data amplification function. Segmentation contours, after pre-processing with the proposed scheme, are smoother and follow the contour of the cell more tightly. We achieved an average accuracy of 94.0026%, which is a 1.1397% increase compared to an image with no pre-processing.

Parameter Estimation for Discrete Chan-Vese Segmentation - Chapter 6 Fluorescence images in cellular biology alone are greatly diverse and it seems as if there are very few common characteristics traits that are common to all fluorescence images. Previous researchers have tried to find parameter settings that work well, however; these are very limited in the general application of fluorescence image segmentation.

We propose a parameter estimation method that tunes the parameters to the corresponding image. The derived method focuses not on the parameters directly but on the relationship between parameters. When suitable approximations of the final means, for the background and the object, are obtained, the parameters are calculated based on the relationship between parameters which, in this case, were tuned for cellular fluorescence images. The proposed parameter estimation scheme was tested against two published parameter settings by El-Zehiry *et al.* [239] and Masaka *et al.* [237]. The proposed scheme exhibited high and consistent accuracy over a large range of images which average

at 93.5329%. This is a significant boost in comparison to the parameter setting by El-Zehiry *et al.* [239] which averaged at 80.5732% and Masaka *et al.* [237] which averaged at 54.7369%.

Interactive Segmentation - Chapter 7 User interaction allows for specific marking of the objects of interest which sometimes presents too difficult a problem for automatic recognition or if the automatic seeding is not reliable. While fully automatic segmentation is the ultimate goal of image segmentation it is not always possible in industry especially when the properties of the image is not well known. While interactive segmentation is still widely used, the goal of reduced user interaction and increased segmentation accuracy has not changed.

We propose an energy weighting system that is tailored to fluorescence image segmentation. There are two variants i.e. with and without hard constraints. The energy function is based on the general properties of intensity variation and pixel probability. The energy functions are based on the intensity variation of fluorescence images. The proposed energy functions were tested against two common energy systems for interactive graph cut segmentation by Boykov and Jolly [25] and Eriksson *et al.* [26]. The proposed scheme shows an increase in accuracy and greater consistency over a large range of images which averaged at 94.5778% with hard constraints and 94.5771% without hard constraints. This is a significant boost in comparison to the general energy functions proposed by Boykov and Jolly [25] which averaged at 84.6089% and Eriksson *et al.* [26] at 86.9989%.

The results from the research undertaken has confirmed that the aim of this dissertation has been achieved. We have studied the fluorescence imaging process and fluorescence images, and leveraged the understanding thereof to design fluorescence-image-specific segmentation schemes and functions that allow for more accurate segmentation and reduced user interaction.

8.1 Limitations

1. The remapping function presented in Section 5.1.2 has 5 tuning parameters. Although there is greater freedom in tuning the curve, the added complexity in parameter finding is not welcomed.
2. The pre-processing scheme presented in Chapter 5 has a total of 14 tuning parameters. This makes it difficult to devise parameter estimation methods.
3. In the proposed parameter settings for the proposed parameter estimation method for ACWE graph cut segmentation in Section 6.2, we maintained a proportional relation between λ_0 and λ_1 . Similarly, the α parameter remains constant and is not subject to appropriate variation with the image.

8.2 Future Work and Extensions

1. The parameters for the remapping function can be estimated from the image. This will allow the pre-processing scheme to be more adaptable over a greater variety of images and reduce manual parameter tuning.

2. For the proposed parameter estimation scheme, we used simple mathematical relations. More sophisticated, comprehensive and accurate mathematical relations might allow for better segmentation results and applicability to a large class of images.
3. Devise a method for automatic seeding and use those seeds as input for the proposed method in Section 7.2. This will allow an interactive segmentation technique to become automatic.
4. The parameter estimation technique in Section 6.2 can be extended to colour images.
5. The energy function proposed Section 7.2 can be extended to colour images.

Appendix A

Introduction to Graph Theory

Graph A graph G is a pair (V, E) , where V is the set of nodes/vertices and E is the set of edges consisting of pairs (u, v) where $u, v \in V$. The graph is assumed to be finite i.e. $|V| = n$ and $|E| = m$.

In an **undirected graph**, the edge (u, v) and (v, u) are not distinct. That is, they refer to the same edge. However, in a **directed graph**, the two edges are now distinct. In a directed graph with edge (u, v) , u is known as the **tail** and v is known as the **head**. In directed graphs, edges, also known as arcs, are depicted by placing arrowheads at the head of the edge. Given an edge $e = (u, v)$, u and v are said to be **incident** on e . A graph is said to be **simple** if it does not contain any self-loops. A **self-loop** is an edge with each of its end points being the same vertex.

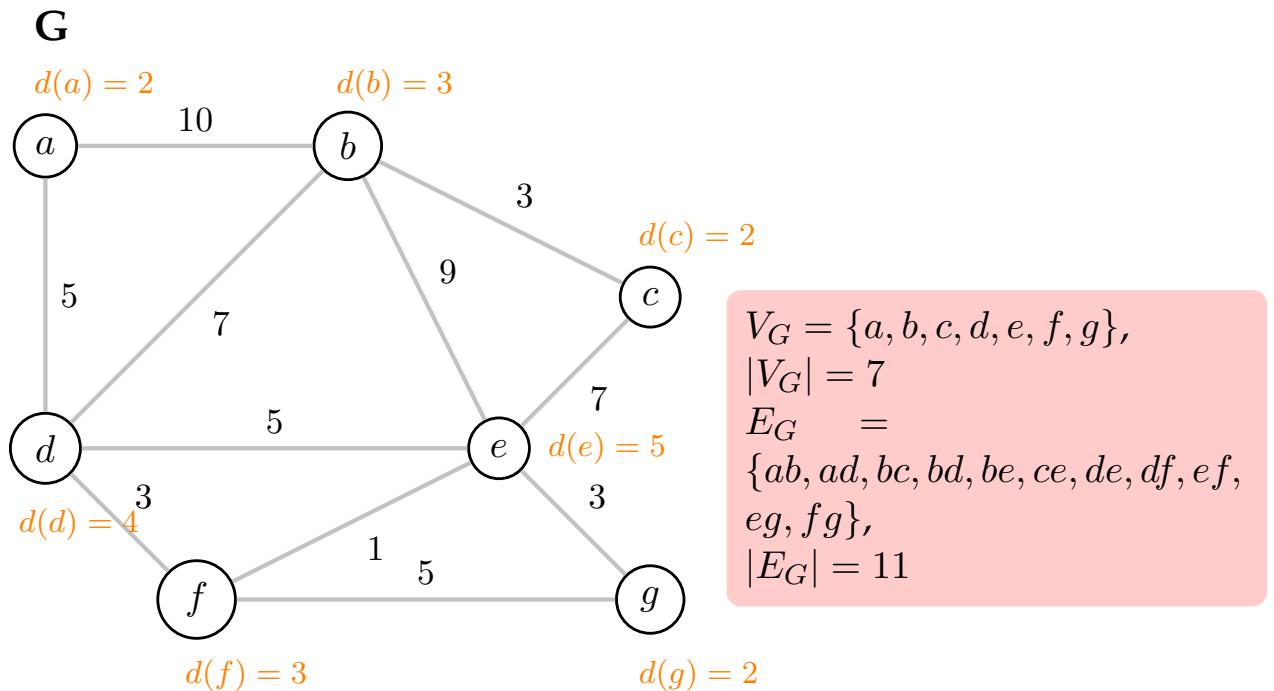


FIGURE A.1: Undirected weighted graph G . The degree of each node is shown next to the corresponding node. The graph is simple. The red box shows the vertex set, V_G , and edge set, E_G , and their corresponding norm.

Degree The degree of a vertex v is the number of edges incident on it. $\deg(v) = |\{(u, v), (v, u) \in E\}|$. A self-loop counts for 2.

If a graph is directed, it is also known as a **digraph**, then a node v has an **in-degree** $d_{in}(v)$ and an **out-degree** $d_{out}(v)$. A digraph is said to be **balanced** if $d_{in}(v) = d_{out}(v), \forall v \in V$.

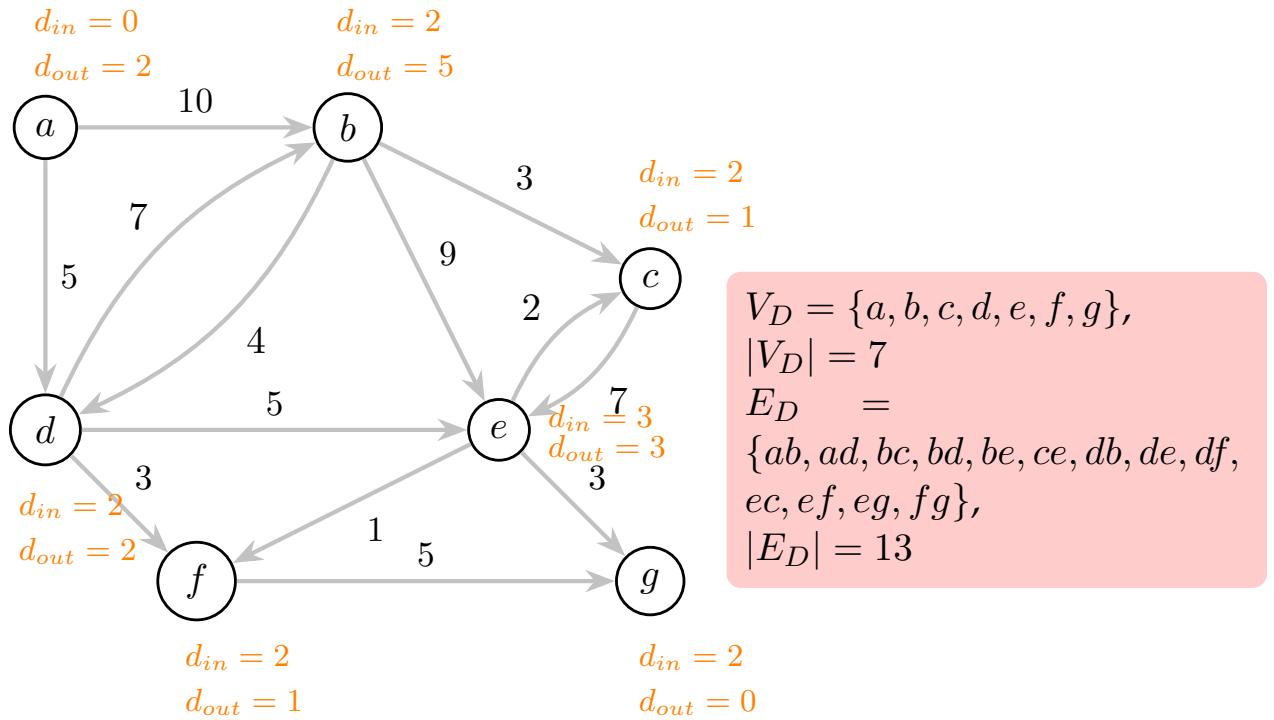
D

FIGURE A.2: Directed weighted graph (Digraph) **D**. The in-degree and out-degree is shown next to each node. The graph is simple and not balanced. The red box shows the vertex set, V_D , and edge set, E_D , and their corresponding norm.

Subgraph A graph $G' = (V', E')$ is said to be a sub-graph of $G = (V, E)$, denoted as $G' \subseteq G$, if $V' \subseteq V$ and $E' \subseteq E$.

Clique A clique is a maximal subgraph.

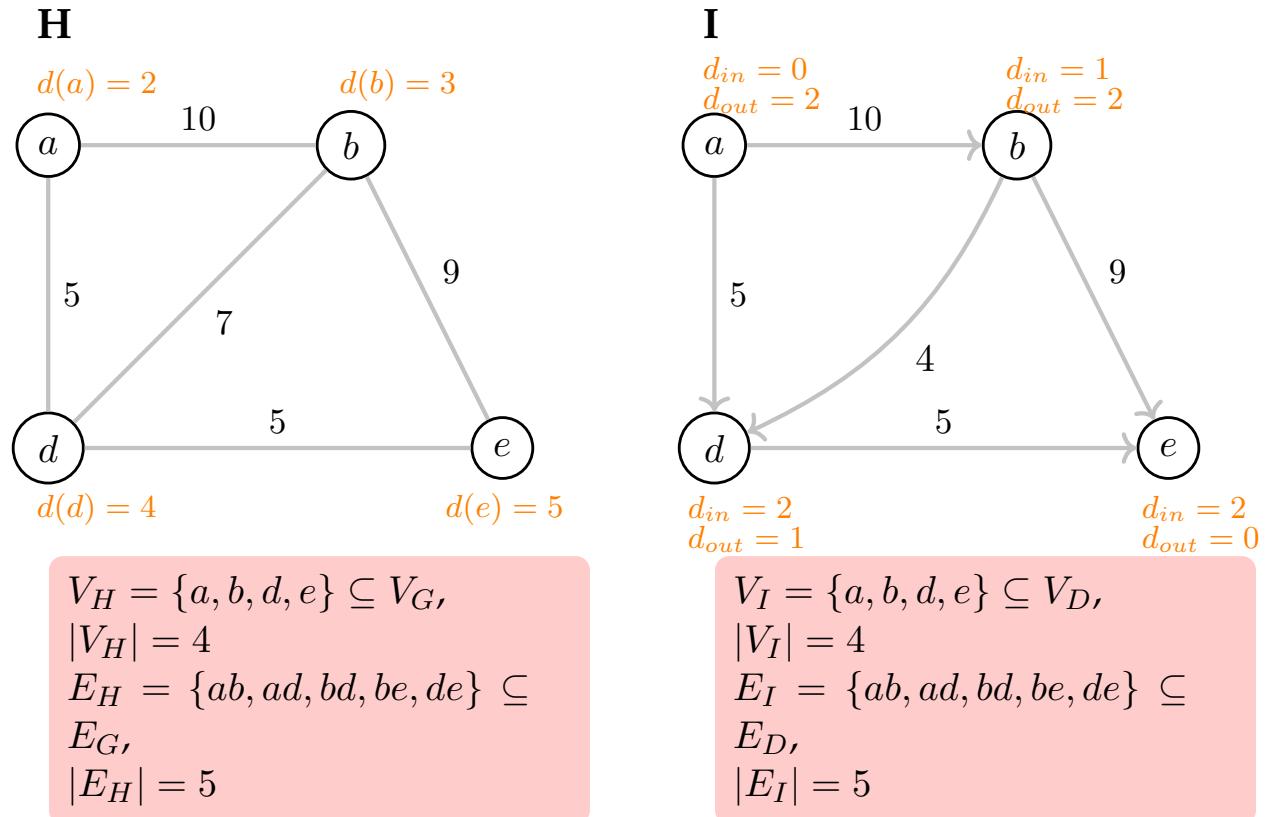


FIGURE A.3: Undirected weighted graph \mathbf{H} is a subgraph of \mathbf{G} in Figure A.1, $\mathbf{H} \subseteq \mathbf{G}$. Directed weighted graph \mathbf{I} is a subgraph of \mathbf{D} in Figure A.2, $\mathbf{I} \subseteq \mathbf{D}$. The degree of each node is shown next to the corresponding node. The red box shows the vertex set, the edge set and their corresponding norms.

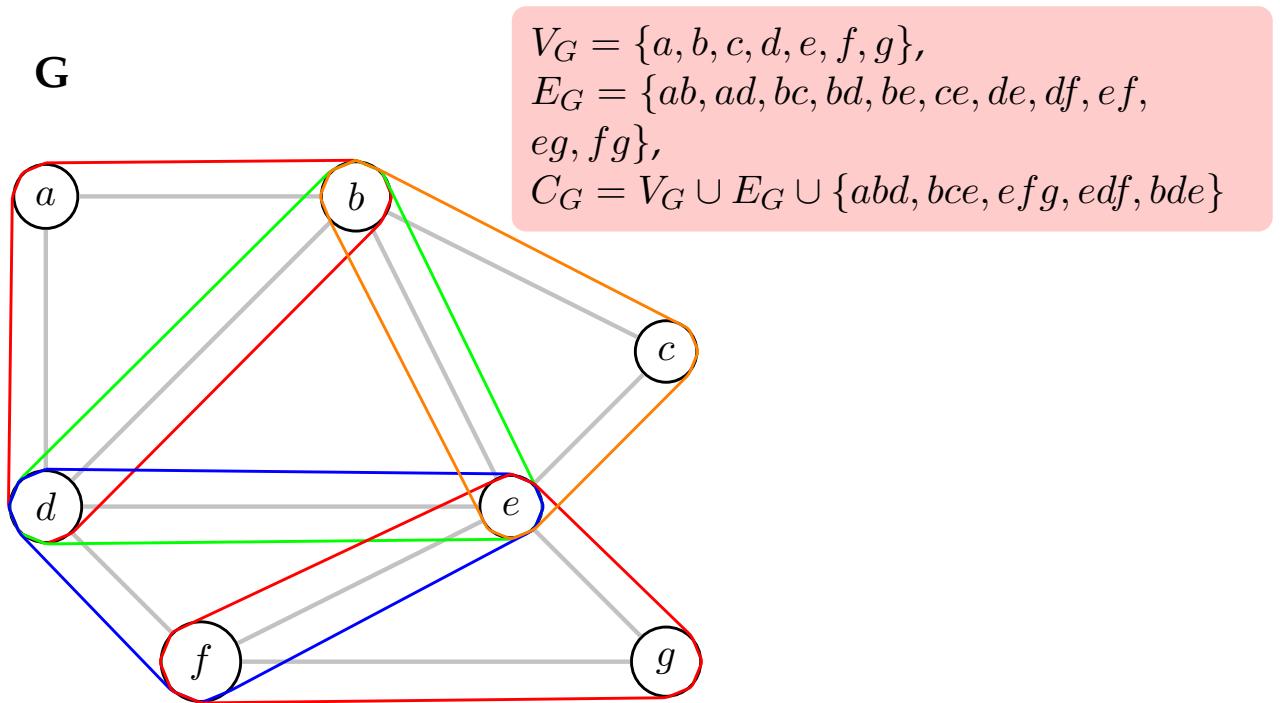


FIGURE A.4: Cliques of the undirected weighted graph \mathbf{G} . The maximal cliques are shown by the hyper-edges that encompass the nodes of that clique.

Appendix B

Cell Images Dataset

The dataset is composed of two subsets. One as the sample set and one as the test set. The sample set is used for tuning parameters and testing theories or predictions. This dataset is composed of images that are relatively simple but still try to maintain some of the variation of images obtained in fluorescence microscopy. The other is the test set. The test set contains more complex images and is used to test the robustness of the segmentation schemes or techniques. We aim for a larger coverage of the types of images that are frequently obtained in fluorescence microscopy.

B.1 Sample Set

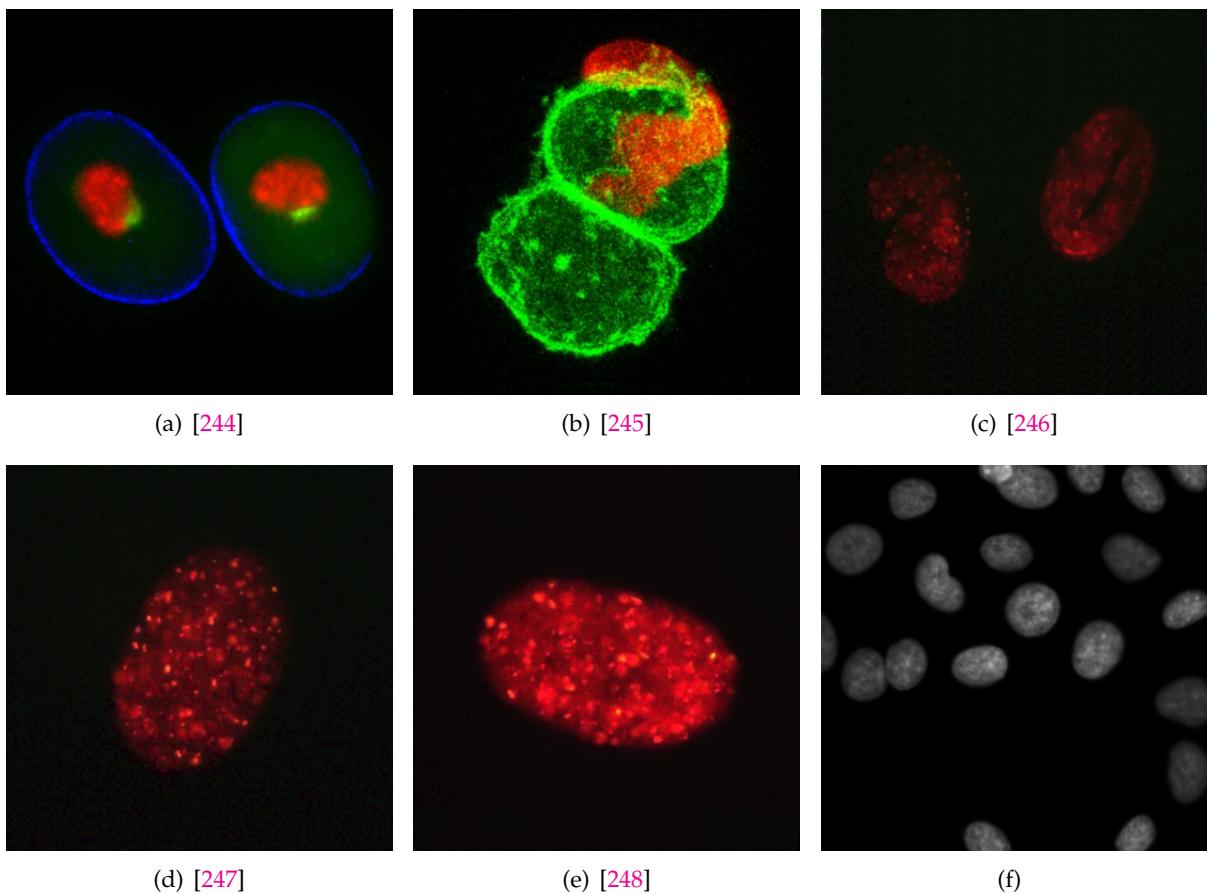
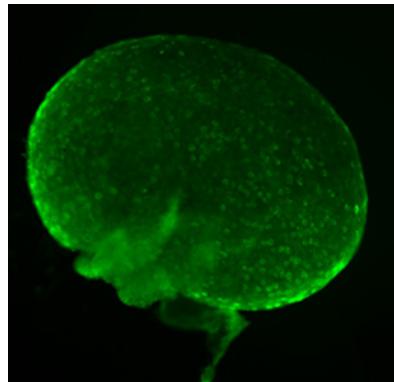


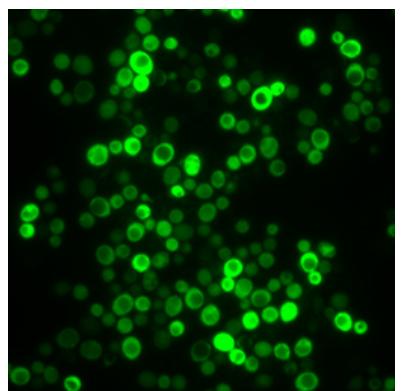
FIGURE B.1: Sample set.

B.2 Test Set

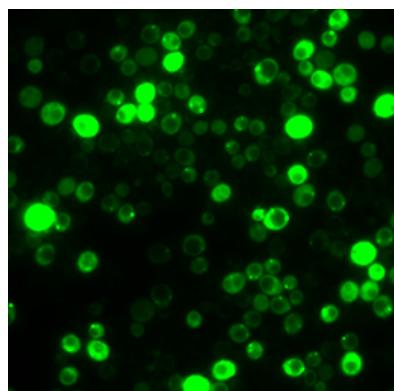


(a) [249]

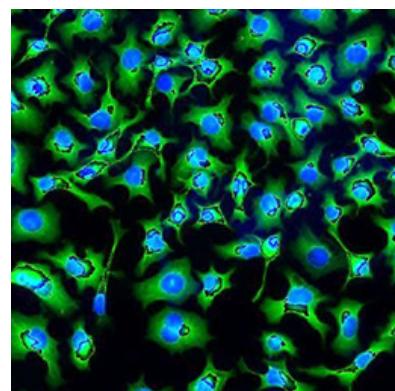
FIGURE B.2: Uneven Illumination



(a) [250]



(b) [251]



(c) [252]

FIGURE B.3: High cell density

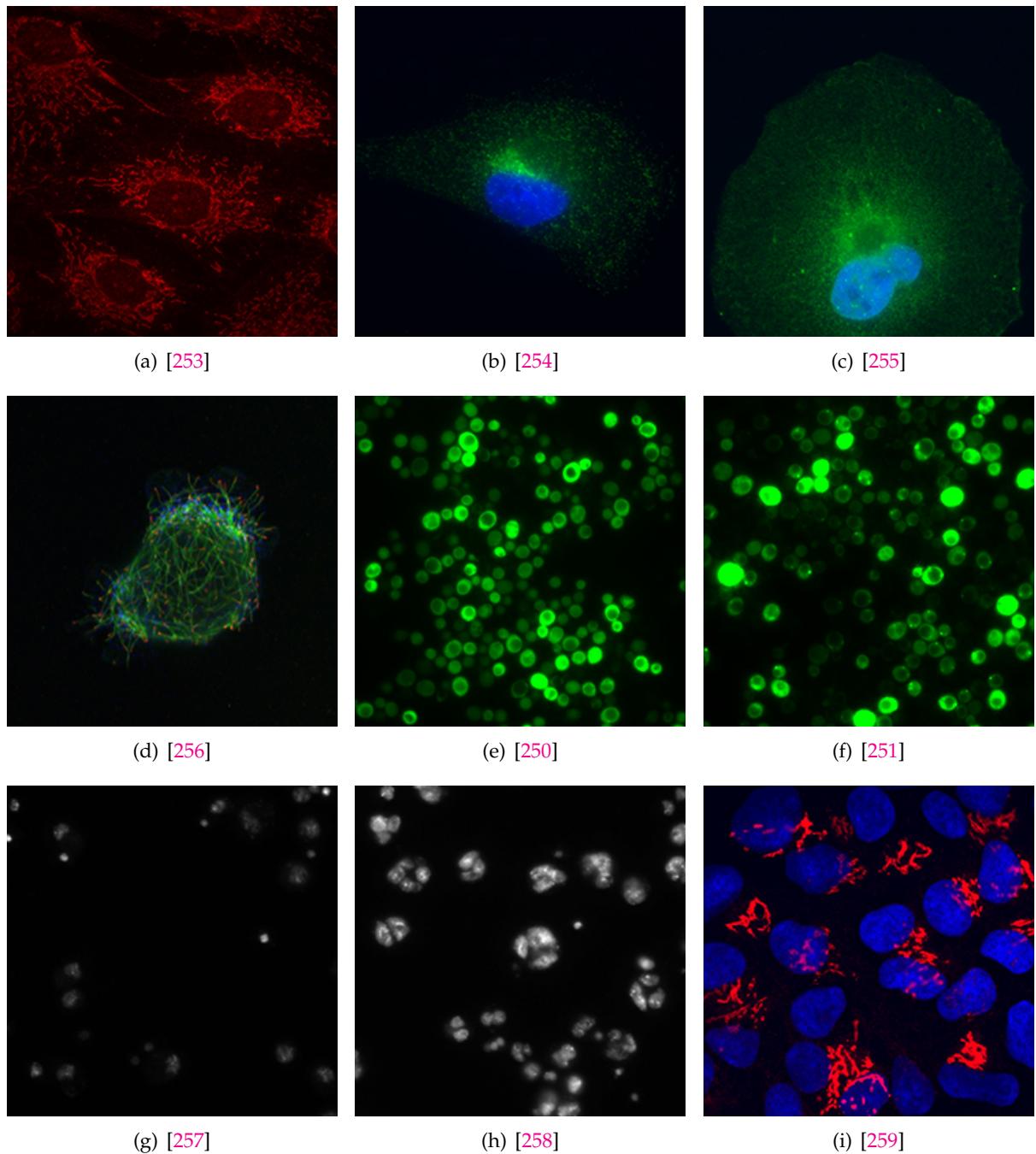


FIGURE B.4: Multi-modal (non-bi-modal)

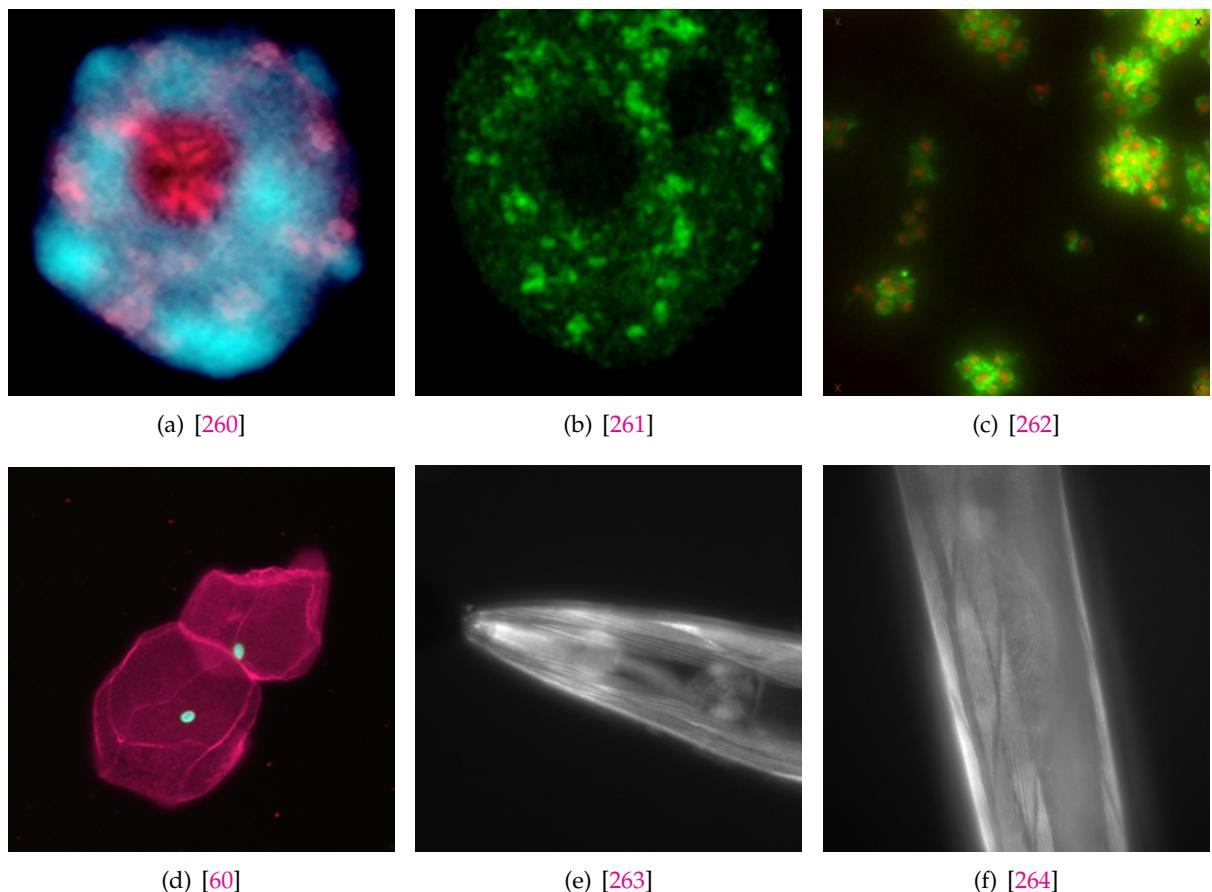


FIGURE B.5: Hazy/Glowing Edges

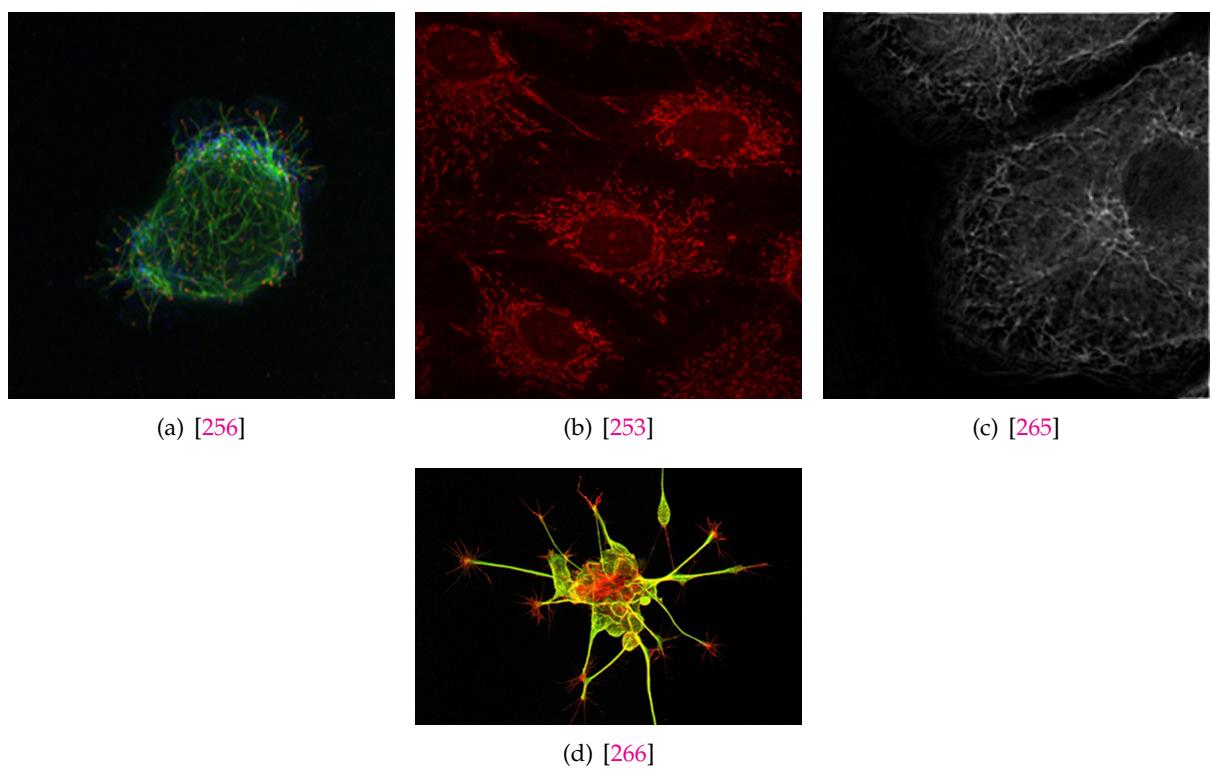


FIGURE B.6: Thin Tentacles

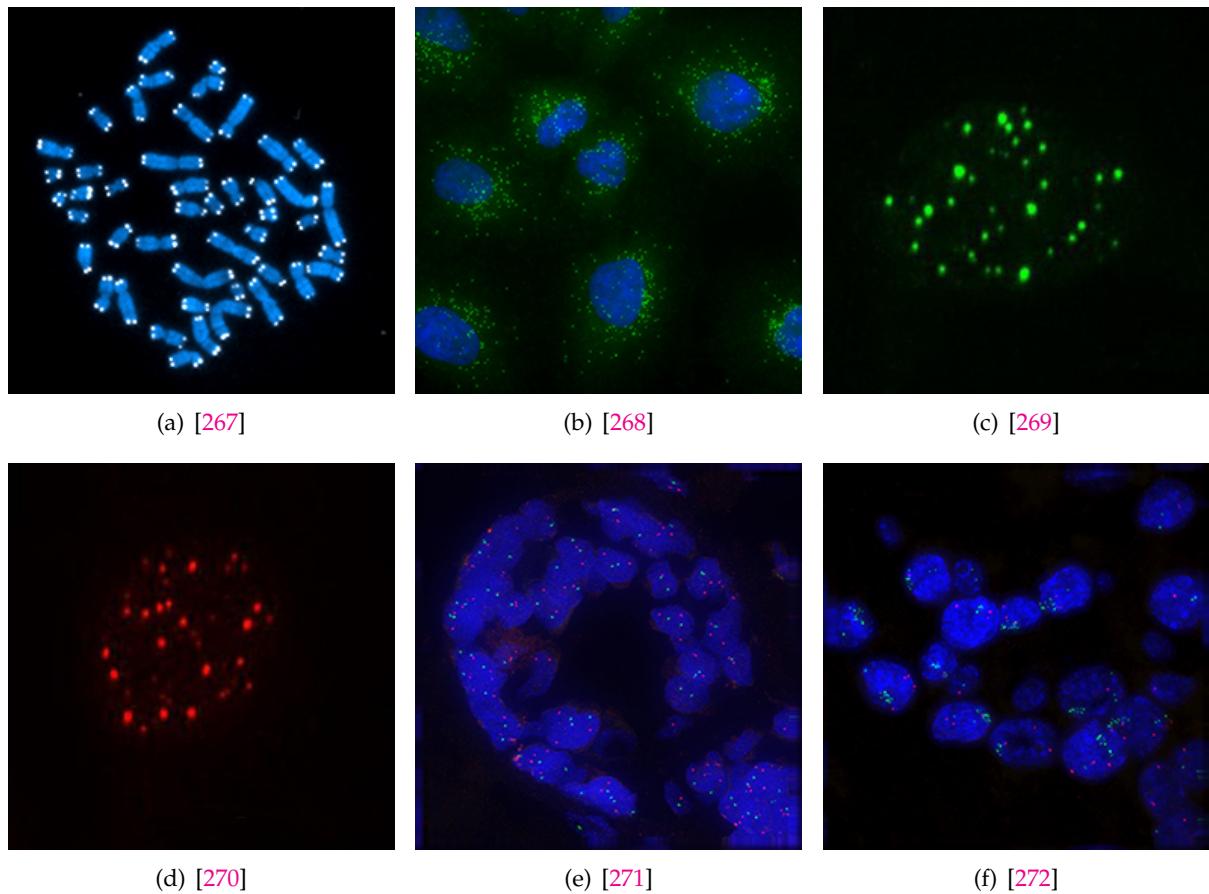


FIGURE B.7: Bright Spots and Speckles

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