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## AN AUTOMATED DENTAL CARIES DETECTION AND SCORING SYSTEM FOR OPTIC IMAGES OF TOOTH OCCLUSAL SURFACE

A dissertation submitted in partial fulfillment of the requirements for the degree of Doctor of Philosophy at Virginia Commonwealth University.

by

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Virginia Commonwealth University Richmond, VA June, 2014



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#### ABSTRACT

### AN AUTOMATED DENTAL CARIES DETECTION AND SCORING SYSTEM FOR OPTIC IMAGES OF TOOTH OCCLUSAL SURFACE

by Leila Ghaedi, Ph.D.

# A dissertation submitted in partial fulfillment of the requirements for the degree of Doctor of Philosophy at Virginia Commonwealth University

Virginia Commonwealth University, 2014

Advisor: Rosalyn Hobson Hargraves, Associate Professor, Department of Electrical and Computer Engineering

Dental caries are one of the most prevalent chronic diseases. Worldwide 60 to 90 percent of school children and nearly 100 percent of adults experienced dental caries. The management of dental caries demands detection of carious lesions at early stages. The research of designing diagnostic tools in caries has been at peak for the last decade. This research aims to design an automated system to detect and score dental caries according to the International Caries Detection and Assessment System (ICDAS) guidelines using the optical images of the occlusal tooth surface. There have been numerous works that address the problem of caries detection by using new imaging technologies or advanced measurements. However, no such study has been done to detect and score caries with the use of optical images of the tooth surface. The aim of this dissertation is to develop image processing and machine learning algorithms to address the problem of detection and scoring the caries by the use of optical image of the tooth surface.



#### **Novelty and Contribution**

Dental caries are one of the most prevalent chronic diseases in the world. According to World Health Organization report on oral health nearly 100 percent of adults experienced dental caries. Scientific evidences show that the early stages of caries can be arrested and possibly reversed by noninvasive intervention such as reduction of cariogenic diet, oral hygiene improvement and fluoride therapy in various delivery modalities. The opportunity of reversing the caries development noninvasively introduced an everyday challenge to the dentists to determine whether noninvasive intervention or restorative intervention is required based on severity and activity of carious lesion; the diagnostic tools can help with decision making in this stage.

A number of existing technologies for caries diagnosis include devices based on laser fluorescence or infrared, electrical conductance measurements, direct digital radiography, etc. are available. These technologies have relatively high prices and also are user sensitive and require several steps in order to perform a clinical reading correctly. The value given by these caries detection devices is subjectively interpreted by the clinician and thus requires a trained dental professional to make a diagnostic or treatment decision. This study uses the optical images of the tooth surface taken by intraoral cameras, which are relatively easy to use, widely available and inexpensive hardware imaging technique, to give a quantitative score of caries severity as well as visual feedback. This may easily augment the decision making process of treatment provided to patients.

The design of this diagnostic tool is very challenging due variation in image quality, presence of natural pits and fissure areas in tooth surface and presence of several other organs and textures in the images. Described below, the methods presented in this dissertation have several novel components that address the above-mentioned challenges.



1. A novel multi stage image segmentation algorithm is created, which incorporates shape, color and gradient specifications of the tooth image.

The presence of normal or carious pits and fissure areas in the tooth surface especially when the change of color occurs near the tooth boundary fails any conventional image segmentation method to segment the tooth surface properly. The proposed segmentation method uses the particular shape of the tooth to find a unique seed point; then uses a top down approach to find out the tooth boundary based on color information and finally refines the output of previous step to the actual tooth surface using a bottom up approach which is applied to the gradient of gray scale image.

2. A modified version of Circular Hough Transform (CHT) is created which uses the tooth shape to find the initial seed point.

Original CHT finds too many false circles in the images. The proposed method applies a series of modification to CHT to address the false circles detection. The calculation of accumulation array is limited to a set of minimum and maximum possible radius. The limits defined based on the application to reduce the computational cost and avoid finding false circles. A level of thresholding applied to the gradient values and another threshold applied to the accumulation array values to reduce the false detected circles.

3. A novel algorithm is proposed to define presence of irregularities in the tooth occlusal surface; irregular regions are the region of interest for dentists while they examine the tooth.

While the irregular regions are defined qualitatively by the dentists based on different color, translucency and porosity; the proposed method utilizes texture analysis and morphological operators to map the irregular regions.



4. The novel feature extraction algorithm was developed to incorporate the texture and morphological information as well as color and intensity levels in the feature space.

The proposed method utilizes a novel approach by calculation of the color and intensity based features on two different masks (irregular regions and tooth surface except irregular regions). Since the irregular region mask encompasses the texture and morphological information, by separate calculation of color and intensity levels for these regions; the final feature space not only has the color and intensity level information but also the texture information.



### Chapter 1

### Introduction

#### 1.1 Aim

The aim of this dissertation is to design an automated system to detect and score dental caries according to the International Caries Detection and Assessment System (ICDAS) guidelines using the optical images of the occlusal tooth surface. The imaging technologies and advanced measurements for caries detection have been an active area of research for the last decade. The final goal of early caries detection tools is to provide an adjacent to clinical decision making and support preventive treatment planning in conjunction with caries risk assessment which finally reduce the risk of premature restoration intervention. However, no such study has been done to detect and score caries with the use of optical images. All the available imaging technologies for caries detection have relatively high prices. Any of the current technologies do not consider information present in the optical images. This study has been designed to incorporate digital images acquired from off-the-shelf commercially available intraoral cameras which are inexpensive in comparison to other dental imaging modalities. By applying image processing techniques, several features extracted from the image of the tooth surface and those features will provide the measures for scoring the tooth according to ICDAS guidelines. These features reveal the spatial information along with texture parameters of the whole tooth area as well as the detected irregular regions.



#### **1.2 Motivation**

#### **1.2.1** Dental Caries Detection Impact

Dental caries are one of the most prevalent chronic diseases in the world. According to World Health Organization report on oral health at April 2012, worldwide 60 to 90 percent of school children and nearly 100 percent of adults experienced dental caries [1]. A significant general reduction in caries lesions has been noted in the United States in the last several decades with the increased availability of fluoride in public water supply, tooth paste and mouth rinse [2]. The widespread use and availability of fluoride has changed the behavior of carious lesions dramatically. The resulted slower progression of carious lesions has afforded the dental profession the opportunity to diagnosis and manage caries at an early stage [3]. Scientific evidences show that the early stages of caries can be arrested and possibly reversed by noninvasive intervention such as reduction of cariogenic diet, oral hygiene improvement and fluoride therapy in various delivery modalities. The opportunity of reversing the caries development noninvasively introduced an everyday challenge to the dentists to determine whether noninvasive intervention or restorative intervention is required based on severity and activity of carious lesion; the diagnostic tools can help with decision making in this stage [3]. Clinical standards for diagnosing caries include visual examination, tactile sensation, aided by radiography combined with patient's individual caries risk levels. Visual examination assesses color, translucency and porosity while tactile examination evaluates hardness and porosity using explorers. When using traditional examination for caries detection; the end result is low sensitivity and high specificity, meaning a large number of lesions may be missed. In addition, using traditional diagnostic methods for diagnosing pit and fissure caries on occlusal surfaces

have a high false positive and false negative rate [4-5]. On the other hand, the greatest reduction



in caries has been noted in smooth tooth surfaces and this type of interproximal lesion can be more easily identified by radiographic techniques [6]. Occlusal lesions have become the largest proportion of the total caries burden [4]. In addition, the current diagnostic methods have a high false positive and false negative rate when diagnosing pit and fissure caries on occlusal surfaces [4-5]. Existing technologies for caries diagnosis include devices based on laser fluorescence (LF) such as LF device, LF pen, LF camera, or infrared (IR) laser fluorescence, referred to as quantitative laser or light fluorescence (QLF). Electrical conductance measurements (ECM), direct digital radiography, Digital Imaging Fiber-Optic Trans-Illumination (DIFOTI) and simple Fiber Optic Trans-Illumination (FOTI), LED-based caries detector and less popular fluorescence spectrophotometer, MicroCT and heat induced detection technique [3, 7-12]. Data shows varying degrees of sensitivity and specificity for In-Vitro and In-Vivo studies [3, 8-9, 13-15]. These technologies have relatively high prices and also are user sensitive and require several steps in order to perform a clinical reading correctly. The value given by these caries detection devices is subjectively interpreted by the clinician and thus requires a trained dental professional to make a diagnostic or treatment decision.

Accurate detection of dental caries, in particular at the early stages, can greatly improve the quality of dental care. The method uses the optical images of the tooth surface taken by intraoral cameras, which are relatively easy to use, widely available and inexpensive hardware imaging technique. This may easily augment the decision making process of treatment provided to patients and their overall impression of the quality of dental care they are receiving.

#### 1.2.2 Objectives

The objective of this thesis dissertation is to design an automated system to detect and score dental caries. The input of the system is the optical image of occlusal tooth surface which has



been taken with the intraoral camera and the output of the system is an ICDAS score which quantify the presence and severity of caries on that tooth surface. Solving this particular problem needs several stages. The first stage of this work is to design an image segmentation method to segment the image into background, regular tooth surface and region of interest for the dentists, which are called; irregular regions. The second stage is extracting features from the segmented areas. The last stage is a classification problem which assigns a score to each image with regard to the severity of the caries on the tooth surface.

The objectives of this dissertation are summarized as follows:

- Create a novel segmentation method, to effectively segment the tooth surface images (both in-vitro and in-vitro images) into background, regular tooth surface and irregular regions according to the guidelines for clinical caries detection.
- Design a feature extraction algorithm that allows for the accurate classification of the dental carries. The extraction of features in medical image applications is a very crucial task. The method utilizes the extracted features and selects only the predominant features through a multi-stage feature selection process in order to automatically score the caries.
- Create a novel classification technique classifies the features extracted from tooth images into the clinical scores. For computing the classification model, an ensemble classifier has been developed which essentially encompasses four different classification methods.

#### **1.3 Overview of Dissertation**

This thesis dissertation is organized as follows.

Chapter 2 provides an overview for the background of the problem. First, International Caries Detection and Assessment System (ICDAS) guidelines are introduced. Then, image



segmentation methods are presented followed by an overview of feature selection and classification methods.

Chapter 3 describes the multi staged segmentation method which results the segmentation of tooth boundary from the complicated background.

Chapter 4 presents the application of texture analysis to segment the irregular regions inside the tooth.

Chapter 5 presents feature extraction, feature selection and feature classification methods which used to classify each tooth to different caries severity classes.

Chapter 6 describes specification of the data sets.

Chapter 7 provides the results and the discussion to assess the performance of the methods. Chapter 8 describes a summary of the work and the future work for this study.



### Chapter 2

### Background

#### **2.1 Introduction**

While carries detection is paramount in the field of cardiology, most in practice methods utilize traditional visual inspection. No similar study has been done to provide a decision support system in the field of cariology with the use of optical images. Moreover none of the current caries detection technologies provide a quantitative feedback for caries management along with visual feedback. Some existing technologies for caries diagnosis are using other types of images such as radiographic images, laser fluorescence images, Fiber-Optic Trans-Illumination images and simple Fiber Optic Trans-Illumination images. Due to different nature of these types of images and optical images of the tooth surface which is the subject of this study, and also different appearance of caries lesions, the image processing methods used in current technologies are not applicable to our problem. However none of the current caries detection devices provided a dental decision support system with the application of machine learning tools. These devices are subjectively interpreted by the clinician and thus require a trained dental professional. The other types of caries diagnosis technologies, such as Electrical conductance measurements (ECM) do not provide any visual feedback [3, 7-12].



With the lack of background in the field of study in dental applications, this study rely on the other areas of medical decision support systems' background, especially those with medical image processing components.

This chapter provides an overview of the ICDAS standard as well as image segmentation, classification and feature extraction algorithms which have been used in the medical decision support systems.

#### **2.2 Caries Detection and ICDAS Guideline**

Clinical standards for diagnosing carious lesions of teeth include visual inspection of tooth surface for color and translucency evaluation, analysis of radiographic images, evaluation of dental surface porosity or hardness; visually or using tactile sense combined with patient's individual caries risk levels [7,16]. As the understanding of dental caries progressed, the clinical criteria systems remained focused on assessment the disease process at only one stage, the so called 'decayed' status. In April and August 2002, a group of caries researchers, epidemiologists, and restorative dentists, met to integrate the different definitions. The group selected a foundation for a new system and proposed a new system which was named the International Caries Detection and Assessment System (ICDAS). A study in 2004, reviewed 29 caries detection criteria systems concluded that the majority of the caries detection systems were ambiguous and did not measure the disease process at its different stages [5]. In 2005, the Rationale and Evidence for the International Caries Detection and Assessment System was presented, followed by the publication of the modified International Caries Detection and Assessment System Criteria Manual [17]. The ICDAS integrated several criteria systems into one standard system for caries detection and assessment [17]. The ICDAS measures the surface changes and potential histological depth of carious lesions by relying on surface characteristics.



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The ICDAS evaluation of pit and fissure caries is based on biological processes of demineralization followed by re-mineralization manifested clinically as changes in color or cavitation. According to ICDAS the dental examiners evaluate the tooth surface and classify the carious status of each tooth surface using a seven-point ordinal scale ranging from sound to extensive cavitation. The classification of the carious status based upon ICDAS is as follows. Sound Tooth Surface (Score 0): There should be no evidence of caries, either no or questionable change in enamel translucency after prolonged air drying for 5 seconds. First visual change in enamel (Score 1): When seen wet there is no evidence of any change in color attributable to carious activity, but after prolonged air drying for 5 seconds carious opacity or discoloration (white or brown lesion) is visible that is not consistent with the clinical appearance of sound enamel. Distinct visual change in enamel (Score 2): The tooth must be viewed wet. There is a carious opacity, white spot lesion and/or brown carious discoloration which are wider than the natural fissure that is not consistent with the clinical appearance of sound enamel; the lesion must still be visible when dry. Localized enamel breakdown because of caries with no visible dentin or underlying shadow (Score 3): The tooth viewed wet may have a clear carious opacity, white spot lesion and/or brown carious discoloration which is wider than the natural fissure that is not consistent with the clinical appearance of sound enamel. After drying for approximately 5 seconds there is carious loss of tooth structure at the entrance to, or within, the pit or fissure area. Underlying dark shadow from dentin with or without localized enamel **breakdown** (Score 4): This lesion appears as a shadow of discolored dentin visible through an apparently intact enamel surface which may or may not show signs of localized breakdown, loss of continuity of the surface that is not showing the dentin. The shadow appearance is often seen more easily when the tooth is wet. The darkened area is an intrinsic shadow which may appear as



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grey, blue or brown in color. The shadow must clearly represent caries that started on the tooth surface being evaluated. Distinct cavity with visible dentin (Score 5): Cavitation in opaque or discolored enamel exposing the dentin beneath. The tooth viewed wet may have darkening of the dentin visible through the enamel. Once dried for 5 seconds there is visual evidence of loss of tooth structure at the entrance to or within the pit or fissure frank cavitation. There is visual evidence of demineralization such as opaque (white), brown or dark brown walls at the entrance to or within the pit or fissure and in the examiner judgment dentin is exposed. The WHO/CPI/PSR probe can be used to confirm the presence of a cavity apparently in dentin. This is achieved by sliding the ball end along the suspect pit or fissure and a dentin cavity is detected if the ball enters the opening of the cavity and in the opinion of the examiner the base is in dentin. Extensive distinct cavity with visible dentin (Score 6): Obvious loss of tooth structure, the cavity is both deep and wide and dentin is clearly visible on the walls and at the base. An extensive cavity involves at least half of a tooth surface or possibly reaching the pulp [17]. The description of the scoring system has been provided in table 2.1. Figure 2.1 provides a schematic section of the tooth structure, which illustrates dentin and enamel locations.

Often this process is not detectable using the current technology available for *In-Vivo* use [17]. *In-Vitro* studies of ICDAS validated the practicality of the system and its correlation with histological examination of pits and fissures in occlusal surfaces of extracted teeth [18]. Studies assessed inter- and intra-examiner reproducibility and accuracy in the detection and assessment of occlusal caries in extracted teeth using ICDAS, using histology as 'gold standard'. ICDAS-II presented good reproducibility and accuracy in detecting occlusal caries [19] and was able to especially identify caries lesions in the outer half of the enamel [8]. More recently, the accuracy of ICDAS was measured *In-Vivo* to compare performance of automated caries detection devices.



The teeth were then extracted and ICDAS was validated based on histological findings [8-20]. ICDAS demonstrated good performance in helping detect occlusal caries *In-Vivo* and moreover, better accuracy was achieved in detecting early lesions [20]. A recent study assessed the agreement among four techniques used as gold standard for the validation of methods for occlusal caries detection and concluded that the outcome of caries diagnostic tests may be influenced by the validation method applied [23], hence the difference in ICDAS accuracy between studies. Based on this evidence of the validity of the ICDAS in caries diagnosis, ICDAS scores were used in this study as the gold standard.

ICDAS Score	Description
0	Sound Tooth
1	First visual Change in Enamel
2	Distinct Visual Change in Enamel
3	Localized Enamel Breakdown
4	Underlying Dentin Shadow
5	Distinct Cavity with Visible Dentin
6	Extensive Cavity with Visible Dentin

Table 2.1: ICDAS scores' description

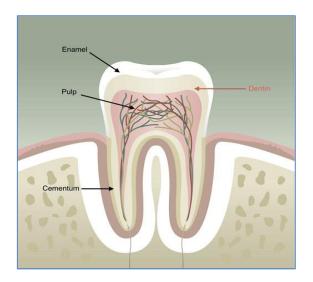


Figure 2.1: Schematic section tooth [22]



#### 2.3 Image Segmentation Methods

Image segmentation plays a crucial role in many medical imaging applications by automating or facilitating the delineation of anatomical structure or other regions of interest [23]. In medical imaging, typically the task of segmentation corresponds to different organs, biological structures or pathologies. Segmentation methods use either discontinuity or homogeneity of gray level values in a region to define the segments. Partitioning based approaches form the segments by detecting isolated points, lines and edges according to abrupt changes in gray levels. Homogeneity based algorithms include thresholding, clustering, region growing, and region splitting and merging.

#### 2.3.1 Threshold-Based Methods

These methods are among the simplest methods used for segmentation. Threshold based image segmentation techniques discriminate regions on the basis of intensity value difference between pixels. The pixels in the image are classified into two classes based on some predefined threshold value [23-30]. Threshold for image segmentation has been calculated based on maximum entropy, interclass variation or histogram. Threshold based segmentation does not account for spatial characteristics of an image, making it sensitive to noise and intensity inhomogenities. The threshold based segmentation techniques perform well for images which have only two components; for complex images, these methods are often used as an initial step in a sequence of image processing operations [31].

#### 2.3.2 Region Growing Methods

The idea of region based algorithms comes from the observation that pixels inside a structure tend to have similar intensities. Region growing techniques are used to segment regions based on



some similarity criteria. Each region of interest (ROI) requires its own seed initialization, after selecting the initial seeds, algorithm searches for the neighborhood pixels which have intensities within a predefined interval [23-24]. To eliminate the need for manual seed initialization, some algorithms used the statistical information and a prior knowledge of the ROIs to select the seeds semi automatically or fully automatically. The drawbacks of these methods are that they are sensitive to the seed selection and also sensitive to the noise, sometimes the similarity criterion is not exactly defined, also the algorithm mainly relies on the image intensity information. In addition, these techniques are dominated by the growth of the current region. Region growing methods are simple techniques that provide good results especially with smaller region segmentation once all mentioned challenges are properly addressed.

#### 2.3.3 Active Contour Models (Snakes)

Active contour models (ACMs) or snakes employ model-based methods that use a prior model to try to find the best match for the model within the image. Active contour models are often called snakes because they appear to slither across image edges. ACMs are one example of the general technique of matching a deformable model to an image using energy minimization. From any starting point, subject to certain constraints, ACM will deform into alignment with the nearest salient feature in the image; such features correspond to local minima in the energy generated by processing the image. ACMs provide a low-level mechanism that seeks appropriate local minima rather than searching for a global solution. In comparison to bottom-up image processing techniques, this technique uses a top-down approach. The ACM algorithm makes use of the identification of local structures such as edges, points and other low-level structures in the image that are assembled into groups to find the objects. The ACM algorithm creates a model of the shape that uses two opposing energy terms, an internal term which works towards smoothing the



curve, and an external term which moves the curves towards image features, to locate the outline of an object. ACMs are good for amorphous objects like cells, but they tend not to perform well with objects that have a known shape. The ACM algorithm does not try to solve the entire problem of finding salient image features; they rely on high-level mechanisms to place them somewhere near a desired solution (a prior knowledge). For example, automatic initialization procedures can use standard image processing techniques to locate features of interest that are then refined using snakes [32, 33].

#### 2.3.4 Color Image Segmentation

Color segmentation presents its own unique challenges. Color segmentation approaches are based on monochrome segmentation approaches operating in different color spaces. There is no uniquely superior technique, as each application presents its own specific challenges and all of the existing color image segmentation approaches are strongly application dependent. An image segmentation problem is basically one of psychophysical perception, and it is essential to supplement any mathematical solutions by a priori knowledge about the image in specific application. Most gray level image segmentation techniques could be extended to color image, such as histogram thresholding, clustering, region growing, edge detection and fuzzy based approaches. They can be directly applied to each component of a color space, and then the results can be combined in some way to obtain the final segmentation result. However, one of the problems is how to employ the color information as a whole for each pixel. When color is projected onto three components, the color information is so scattered that the color image becomes simply a multispectral image and the color information that human can perceive is lost. Another problem is how to choose the color representation for segmentation, since each color representation has its advantages and disadvantages [34]. In most of the existing color image



segmentation approaches, the definition of a region is based on similar color. This assumption often makes it difficult for many algorithms to separate the objects with highlights, shadows, shading or texture which cause inhomogeneous colors of the objects' surface.

#### 2.3.4.1 Color Space Presentation

Color is perceived by humans as a combination of tristimuli R (red), G (green), and B (blue) which usually called the three primary colors. Several color representations are defined by linear or nonlinear transformations of RGB space. Several color spaces, such as RGB, HSV and CIE are utilized in color image segmentation, but none of them outperforms the others for all kinds of color images [34]. The Hue Saturation Value (HSV) is more intuitive to human vision [34]. The HSV is one of the most common cylindrical coordinate representations of color. HSV system separates color information of an image from its intensity information. Color information is represented by Hue and Saturation, while Value describes the brightness of an image, is determined by the amount of the light. Hue represents basic colors, and is determined by the dominant wavelength in the spectral distribution of light wavelengths. It is the location of the peak in the spectral distribution. The saturation is a measure of the purity of the color, and signifies the amount of white light mixed with the hue. It is the height of the peak relative to the entire spectral distribution. The HSV color space can be described geometrically as in Figure 2.2.

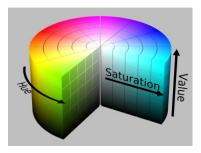


Figure 2.2: HSV color space [35]



#### **2.4 Classification Methods**

The task of assigning an input vector, to several classes is called a classification problem. The input vector of N components is called a *pattern* and each component of the input vector is called a *feature*. The task of classifying data is to decide class membership y' of an unknown data item x based on a data set  $D = (x_1, y_1), \dots, (x_n, y_n)$  data items  $x_i$  with known class memberships y<sub>i</sub>. For ease of discussion, only dichotomous classification problems are considered, where the class labels y are either 0 or 1. The  $x_i$  are usually N-dimensional vectors, the components of which are called covariates and independent variables in statistics parlance or *features* by the machine learning community. In most problem domains, there is no functional relationship y=f(x) between y and x. In this case, the relationship between x and y has to be described more generally by a probability distribution P(x,y); one then assumes that the data set D contains independent samples from P. From statistical decision theory, it is well known that the optimal class membership decision is to choose the class labely that maximizes the posterior distribution P(y|x). In this dissertation the features are statistical measures of the tooth image and the classes are caries scores according to ICDAS. The design of this study is based on supervised learning paradigm. There are several machine learning algorithms to choose from, where the choice simply depends on the type of dataset and its complexity. Four popular classification methods which has been used in medical decision support applications introduces in this session [36].

#### 2.4.1 Support Vector Machine (SVM)

Support vector machines are algorithmic implementations of ideas from statistical learning theory. Statistical learning theory solves the problem of building consistent estimators from data, meaning by having only characteristics of the model, and performance on a training set, how the



performance of a model on an unknown data set can be estimated. SVMs build optimal separating boundaries between data sets by solving a constrained quadratic optimization problem. By using different kernel functions, varying degrees of nonlinearity and flexibility can be included in the model. Because they can be derived from advanced statistical ideas, and bounds on the generalization error can be calculated for them, support vector machines have received considerable research interest over the past years. The disadvantage of support vector machines is that the classification result is purely dichotomous, and no probability of class membership is given.

#### 2.4.2 C4.5 Decision Tree

The C4.5 algorithm builds decision trees from a set of training data in the, using the concept of information entropy. The input vector of N components is called a *pattern* and each component of the input vector is called a *feature*. The task of classifying data is to decide class membership y' of an unknown data item x' based on a training data set  $D=(x_1,y_1),...,(x_n,y_n)$  of data items xi with known class memberships yi. At each node of the tree, C4.5 chooses the feature of the data that most effectively splits its set of samples into subsets enriched in one class or the other. The splitting criterion is the normalized information gain (difference in entropy). The feature with the highest normalized information gain is chosen to make the decision.

#### 2.4.3 Random Forest Tree

Random forests are a combination of tree classifiers such that each tree depends on the values of a random vector sampled independently and with the same distribution for all trees in the forest. Random forest uses multiple trees or a forest to develop decisions and classifications. Random forest can be used for both supervised and unsupervised data learning problems. In this method many classification trees are grown to develop the rules for decisions and classifications. The



generalization error for forests converges to a limit as the number of trees in the forest becomes larger. The generalization error of a forest of tree classifiers depends on the strength of the individual trees in the forest and the correlation between them. A random forest is a classifier consisting of a collection of tree-structured classifiers { $h(\mathbf{x}, k)$ , k = 1, ...} where the { $_k$ } are independent identically distributed random vectors and each tree casts a unit vote for the most popular class at input x [37].

To classify a new object from an input vector, the input vector is applied to each of the trees in the forest. Each tree gives a classification, and the tree votes for that class. Over all the grown tress, the forest chooses the classification having the most votes. When the training set for the current tree is drawn by sampling with replacement, about one-third of the cases are left out of the sample. This left out data is used to get a running unbiased estimate of the classification error as trees are added to the forest. It is also used to get estimates of variable importance. After each tree is built, all of the data are run through the tree, and proximities are computed for each pair of cases.

#### 2.4.4 Neural Network Classifier

Artificial Neural Networks (ANN) represents a paradigm for machine learning. The most widely applied use of ANNs in medical imaging is as a classifier [23-24]. ANNs are parallel networks of processing elements that simulate biological learning. Each node in an ANN is capable of performing elementary computations. Learning is achieved through the adaptation of weights assigned to connections between nodes. These networks have high parallel ability and high interaction among the processing units enabling it to model any kind of process. Because of many interconnections used in a neural network, spatial information can be easily incorporated to its classification procedure.



#### **2.5 Feature Extraction**

To be able to apply the machine learning algorithms to an image, feature extraction is needed to aggregate the image specification. Exploration of spatial information and textural information of the images are crucial in this study. Both global and regional features should be extracted. Although a clear definition of texture does not exist, it can be understood as a group of image properties that relate to our intuitive notions of coarseness, rugosity, smoothness etc. [34]. Texture features can be grouped into transform-based and statistical techniques. Transform approaches comprise all methods based on frequency or scale transforms such as Fourier Wavelet; they attempt to describe the image regions using their frequency content or their frequency and scale content. The statistical approaches use the pixel gray level distribution to extract texture information from the image and are the most used for medical images analysis which seems reasonable given the irregularity of shapes and variety of texture types found in medical images [34].

#### 2.6 Feature Selection Methods

In many machine learning applications, it is not only important to be able to classify the data sets, but also to determine which features are the most relevant for achieving this separation. A large number of algorithms have been proposed for feature subset selection and many methods have been introduced to measure feature strength. Such methods can be divided into two broad categories: heuristic-based methods and wrapper-based methods. Heuristic methods utilize a predefined measure of feature strength with respect to the class variable. An example is *information gain ratio*, defined as follows.

Information Gain (*Class*, *Feature*) = H(*Class*) - H(*Class* | *Feature*)



Wrapper based methods utilize an induction algorithm to create a model. Then, according to the performance of the model, the features are either ranked through some measure of contribution to the model or best subsets are found. The task of feature selection can be categorized under the task of parameter optimization for a Maximum Likelihood algorithm. For most induction algorithms certain parameters are not tuned /optimized automatically. While the weights assigned to each feature are necessarily optimized when building, for instance, a logistic regression model or a neural network model, other constant parameters such as the number of hidden neurons, learning rate, misclassifications allowed, etc. remain constant. During additive logistic regression, as the weights assigned to some features may approach *zero*, it would result in automatic feature selection. As such, feature selection can be seen as the task of optimizing a utility vector *U* that selects/discards each of *m* features.

$$U = \{u_{f1}, ..., u_{fm}\}, \text{ where } u_{fi} \subset \{0, 1\}$$

Since the wrapper approach involves building numerous models/mappings, only the fastest algorithms can be used in wrappers. Simple decision trees, logistic regression, naive bayes are a few examples; the implementations of SVMs are too slow for use in wrappers for feature selection. However, the combination of linear SVMs and feature ranking has been used successfully for this purpose [38].

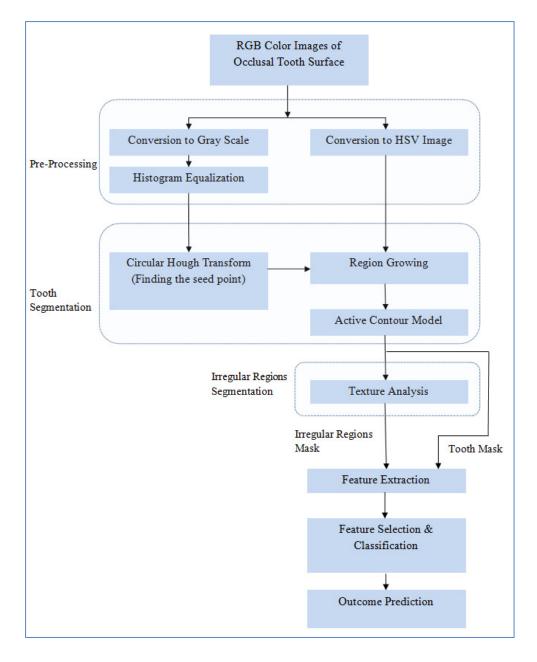
#### 2.7 Overview of the Method

The methodology is a multi-stage hierarchical technique that applies some of the methods discussed above. In particular, the algorithm provides a novel approach for texture feature extraction based on both color and gray level image information. The technique also introduces a



multi-stage tooth segmentation technique that deals with the variations typically observed in biomedical images.

The ultimate goal of the project is to segment the tooth and irregular regions and extract features from these images and provide recommendations that can be directly used for diagnosis and treatment planning. The block diagram of the system components is presented in figure 2.3.







### Chapter 3

### Tooth Surface Segmentation

#### **3.1 Introduction**

The designed computational method herein analyzes photographs captured by digital cameras and produces predictions as to the existence and the severity of caries. The method segments the tooth image into background, healthy enamel surface, and any irregular regions. Irregular regions, in this study, are the regions of interest for the dentists, which show different color, translucency and porosity. Segmentation is performed in two stages; at the first stage the tooth surface is segmented from the background and the second step is to determine the irregular regions within the tooth boundaries. The first step of segmentation process is described in this chapter and the irregular region segmentation methodology is described in the next chapter.

Segmentation of tooth from the complex background is the first step in order to design a practical dental decision support system. Given the complex backgrounds (gum, tongue, adjacent teeth, etc.) as well as variety of tooth shapes, the tooth boundaries detection cannot be achieved by applying either a top-down or a bottom-up approach alone. By combining a top-down and a bottom-up approach, this method is capable of accurate detection of the tooth boundary. The



methodology is a novel multi-stage technique that applies a single seeded color image region growing method and an active contour model to find the tooth boundary.

#### **3.2 Pre-Processing**

The initial color image is in RGB format. Both grayscale and HSV information will be used in the multi stage segmentation technique; so a color space transformation is needed. In order to convert RGB to grayscale the standard NTSC (National Television System Committee) formula is used. The intensity is calculated directly from gamma-compressed primary intensities as a weighted sum as described in equation 3.1.

$$I = 0.2990R + 0.5870G + 0.1140B \tag{3.1}$$

The Hue Saturation Value (HSV) system separates color information of an image from its intensity information. Color information is represented by Hue and Saturation, while Value describes the brightness of an image. Hue represents basic colors, and is determined by the dominant wavelength in the spectral distribution of light wavelengths. It is the location of the peak in the spectral distribution. The saturation is a measure of the purity of the color, and signifies the amount of white light mixed with the hue. It is the height of the peak relative to the entire spectral distribution. RGB to HSV transformation is as described in equations 3.2 to 3.4.

$$H = \arctan\left(\frac{\sqrt{3} (G-B)}{(R-G)+(R-B)}\right)$$
(3.2)

$$V = \frac{(R+G+B)}{3} \tag{3.3}$$

$$S = 1 - \frac{\min(R, G, B)}{V} \tag{3.4}$$



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Histogram equalization is applied to the gray level image to improve performance in the subsequent image processing steps. Histogram equalization reduces the effect of under and over exposure. Histogram equalization accomplishes this by effectively spreading out the most frequent intensity values. The gray level transformation function, T(x) is given by equations 3.5 and 3.6.

$$y = T(x) \tag{3.5}$$

$$y_{k} = T(x_{k}) = \sum_{j=0}^{k} P_{x}(x_{j}) = \sum_{j=0}^{k} \frac{n_{j}}{n}$$
(3.6)

Where, x is the input image and y is the output image and k=0,1,...,L-1; L is the total number of gray levels in the image (in this case 256);  $n_j$  is the number of occurrence of a pixel with gray level j and n is the total number of pixels in the image, so  $P_x(j)=n_j/n$  is the probability of occurrence of a pixel with gray level j [39].

### 3.3 Initial Single Seed Selection

Conventional image segmentation techniques using region growing require initial seed selection and recursive partitioning/ merging which has high computational cost and execution time. Also with the pits and fissure areas and possible existence of caries inside the tooth, the conventional image segmentation will partition the tooth area to more than one region. The tooth background also consists of gums, tongue and adjacent teeth with different color, intensity and texture, so with conventional region growing methods (seed selection, partitioning and merging) the image will be partitioned into more than two regions (one for tooth area and one for background). By selecting a single seed inside the tooth, the desired segmentation and reduction of computational cost is possible.



#### 3.3.1 Modified Circular Hough Transform

Circular Hough Transform used to find a single seed inside the tooth boundary. Circular Hough Transform (CHT) [40-41] detects presence of circular shapes inside an image based on gradient field of the image. The semicircular shape of tooth occlusal surface makes it possible to use CHT to find circles which nearly contain the tooth boundary. During the process of finding the centers and radii some inaccuracies can happen. In this application; finding the accurate circle center is important not the radius of the circle. With the specific use of CHT some modification has been done to adapt the original CHT to suit this research problem.

The original CHT is used to transform a set of feature points in the image space into a set of accumulated votes in a parameter space. Then, for each feature point, votes are accumulated into an accumulator array for all parameter combinations; the accumulation array has the same dimension as the input image. The local maxima of accumulation array that contain the highest number of votes indicate the presence of the circular shape.

A circle pattern is described by equation 3.6. Where  $(x_c, y_c)$  are the coordinates of the center and  $(x_p, y_p)$  are the coordinates of any point on the circle and r is the radius of the circle.

$$(x_p - x_c)^2 + (y_p - y_c)^2 = r^2$$
(3.6)

The CHT utilizes the drawing of perpendicular lines to the edge of a curve or circle, these lines will cross at the center of the circle. Therefore a "hot spot" is achieved at the center of that circle; the accumulation array is calculated to identify that hot spot. In order to get the edges the gradient of the gray scale image as described in equation 3.7 is used.

The gradient of a two-variable function (in this case intensity function f(x,y)) at each image point is a 2D vector, with the components given by the derivatives in the horizontal (x) and vertical (y)



directions. At each image point f(x,y), the gradient vector points in the direction of largest possible intensity increase, and the length of the gradient vector corresponds to the rate of change in that direction. These gradients are less susceptible to lighting and camera changes, so matching errors are reduced.

Gradient = 
$$\Delta f = \frac{\Delta f}{\Delta x} + \frac{\Delta f}{\Delta y}$$
 (3.7)

Gradient Magnitude = 
$$\sqrt{\left(\frac{\Delta f}{\Delta x}\right)^2 + \left(\frac{\Delta f}{\Delta y}\right)^2}$$
 (3.8)

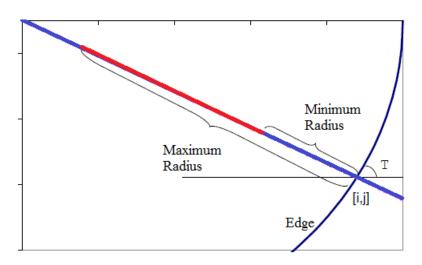


Figure 3.1: Calculating the line segment perpendicular to the edge- limited by minimum and maximum possible radius- for any detected edge, any pixel with the coordinates of red line in accumulation array will get a value

A gradient threshold has been applied to the remove the uniform intensities. The threshold is defined as 6% of maximum intensity of the image which is applicable to all three sets of images. Figure 3.1 describes how the accumulation array is determined. One of the most common uses of gradient is in edge detection. After gradient images have been computed, pixels with large gradient values become possible edge pixels. The pixels with the largest gradient values in the direction of the gradient become edge pixels. The pixels which are perpendicular to any detected



edge and are within (Minimum Radius, Maximum Radius) range will get a value in accumulation array. Another level of thresholding applied to accumulation array value, where any value which is less than mean of accumulation array values is removed. With the application of this threshold local minima with small weights which cause false hot spots and thus false detected circles are avoided. The output of CHT will be a set of circle centers and circle radii. Basically the detected circle's radius can be anything, even larger than the image size. A smaller range of radii would save computational time and memory. In this application the minimum and maximum radii of circles are defined as follows:

Minimum Radius = 
$$\frac{1}{10}$$
 smallest dimension of the image (3.9)

$$Maximum Radius = largest dimension of the image$$
(3.10)

The definition of minimum radius is heuristic and based on this idea that the tooth of interest should be "conceptually obvious" in the image. The output of CHT is a set of N different parameter triplets ( $x_c$ ,  $y_c$ , r), where N is the number of detected circles and ( $x_c$ ,  $y_c$ , r) are circle center dimensions and radius.

To set the seed point, first any circle center which is outside the borders of the image has been removed; the rationale behind this is that the tooth of interest with semi-circular shape should be inside the tooth and any circular shape with a center outside the borders of the image is either a part of an adjacent tooth which should not be used. Then the vector sum of all remained circle centers fall inside the tooth and will be used as the initial seed for region growing step. The CHT applied to the grayscale conversion of the tooth image.



#### Modification to original CHT for the application:

- 1- Minimum and maximum radius definition with regards to the application to reduce the computational cost.
- 2- The method of selection and calculation of final seed point with regards to the definition of tooth of interest.
- 3- Apply two levels of thresholding; first to the gradient values and second to the accumulation array values to reduce the false hot spots with regards to the application.

# 3.4 Color Image Seeded Region Growing

Seeded region growing (SRG) is a hybrid method. It starts with an assigned seed, and the region is grown by merging a pixel into its nearest neighboring seeded region. Considering local information such as regions similarity, boundaries and smoothness makes SRG robust to a large variety of images. Each ROI requires its own seed initialization; in this application there is one ROI, the whole tooth surface. Thus, one initial seed is requires, which is the output of CHT. Once the seed is determined, then the region is grown in the neighborhood of the pixels from the seed. HSV color model which is corresponding to human color perception has been used for region growing. For any pixel at (x,y) a 4- pixel neighbourhood N(x, y)={(x-1,y), (x, y+1), (x+1, y), (x, y-1)} is defined and used for region growing.

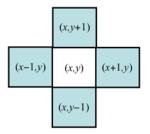


Figure 3.2: 4-Neighbourhood



The region is iteratively grown by comparing all unallocated neighboring pixels to the region. A measure of similarity explained later for HSV space. The pixel with the smallest difference measured is allocated to the respective region. This process stops when the difference measure between region mean and new pixel become larger than a certain threshold (MaxDistance). The obtained image is the initial segmented tooth image [31].

MaxDistance = mean (hsv image) + standard deviation (hsv image) (3.12)

#### 3.4.1 Measure of Similarity for HSV Space

For a pixel at (x,y) the color information is (h(x,y), s(x,y), v(x,y)). The average value of color over the neighborhood N(x,y) is  $(\overline{h(x,y)}, \overline{s(x,y)}, \overline{v(x,y)})$ .

Equation 3.12 computes the distance between (h(x, y), s(x, y), v(x, y))and  $(\overline{h(x, y)}, \overline{s(x, y)}, \overline{v(x, y)})$ .

$$d(x,y) = \sqrt{(v-\bar{v})^2 + (s\cos(h) - \bar{s}\cos(\bar{h}))^2 + (s\sin(h) - \bar{s}\sin(\bar{h}))^2}$$
(3.12)

The value of d(x,y) over N(x,y) defined as a measure of smoothness. The output of region growing is almost near the tooth boundary but it needs another refinement to exactly locate the true boundary of the tooth.

#### **3.5 Active Contour Model**

For almost all the in-vitro images, the two last steps of segmentation were able to segment the tooth but for in-vivo images, yet another step is needed to segment the tooth. The output of region growing has been used as the initial active contour and the gray level image has been used as the input image. An active contour model is a parametric contour that deforms over a series of iterations. A parameter x(s,t) along with the contour therefore depends on two parameters s (contour space parameter) and t (time parameter). The contour is influenced by internal and external constraints, and by image forces. Internal forces constraints give the model tension and



stiffness. External constraints come from high-level sources such as human (in this case region growing algorithm). Image energy is used to drive the model towards salient features such as light and dark regions, edges, and terminations.

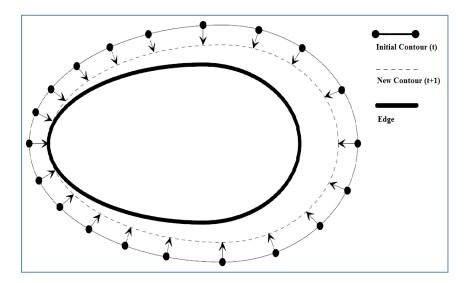


Figure 3.3: An Active Contour Model, over a series of iterations, the active contour moves into alignment with the nearest salient feature, in this case an edge

A final solution is given by the minimum total energy of the snake, which is the result of equation 3.13. Where  $E_{int}$  and  $E_{ext}$  are the internal and external energy of ACM, respectively. The internal energy is given by the membrane energy sum. u(s) is the curve of ACM which has been created by sampling 50 points over the edge of region growing algorithm output.

$$E_{ACM} = \int_0^1 [E_{int}(u(s)) + E_{ext}(u(s))] ds \qquad (3.13)$$

$$u(s) = (x(s), y(s))$$
 (3.14)



# Chapter 4

# Irregular Region Identification

### 4.1 Introduction

The irregular regions are defined by spatial statistics as well as texture analysis, adding texture information empowers the system to not only focus on visible changes in the enamel, which is the region of interest for the dentists but also focus on the textural changes which are not visible and usually are only detectable with tactile examination. These image processing features are designed to best represent visual irregularities examined by the dentists during visual/visuo-tactile examination. These features are then used to detect the existence and severity of caries in the identified irregular regions.

#### **4.2 Texture Analysis**

The irregular regions within the tooth boundaries were segmented by the application of texture assessment through the use of morphology operators. After finding the tooth boundaries, the irregular regions are identified. Haar Discrete Wavelet Transform (DWT) is used to do the texture analysis [42]. The Haar wavelet's mother wavelet function  $\psi$  (t) described in equation 4.1 and its scaling function  $\Phi$ (t) described in equation 4.2.

$$\Psi(t) = \begin{cases} 1 & 0 \le t \le \frac{1}{2} \\ -1 & \frac{1}{2} \le t \le 1 \\ 0 & otherwise \end{cases}$$
(4.1)



$$\psi_{n,k}(t) \!\!=\! 2^{\frac{n}{2}} \psi(2^n t - k), t \in \pmb{R} \tag{4.2}$$

To start irregular region detection, a background mask applied to the color image, meaning all the operations just applied to the tooth boundary. The mean value of the first component of color space (Mean of Hue) is calculated then a black and white mask generated by applying the Mean of RED threshold value. This black and white mask is convolved by a low pass filter of size  $9 \times 9$ to smooth it and reduce the number of connected components. Morphological operations are applied to remove the spurious edges and objects with area smaller than  $3\times3$  squares. The resulted black and white mask then applied to the gray scale image. Morphological operations are a collection of non-linear operations related to the shape or morphology of features in an image. Morphological operations rely only on the relative ordering of pixel values, not on their numerical values, and therefore are especially suited to the processing of binary images. Morphological operations probe an image with a small shape or template called a structuring element. The structuring element is positioned at all possible locations in the image and it is compared with the corresponding neighborhood of pixels then test whether the element fits within the neighborhood or not. In this application the structuring element is a  $3\times3$  square. The  $3 \times 3$  square probes the whole binary image; each of its pixels is associated with the corresponding pixel of the neighborhood under the structuring element. The structuring elements will set to ones if the majority of corresponding pixel of the neighborhood are ones (5 or more ones) otherwise they will set to zeros.

Wavelet transform applied to the output of the previous step in order to reconstruct the image using only the approximation matrix. Haar wavelet is selected as the mother wavelet because of its discontinuity and intrinsic ability to accentuate transitions between gray levels. Then the mean



value ( $\alpha$ ) and standard deviation ( $\beta$ ) of this image is calculated to be used for automatic thresholding.

$$Threshold = \alpha + \beta \tag{4.3}$$

The binary mask is created by selecting the pixels that have gray levels greater than the threshold. Another step of morphological operations again applied to the binary mask to remove the objects smaller than  $3\times3$  squares. The resulting black and white mask is used for identifying the irregular regions.

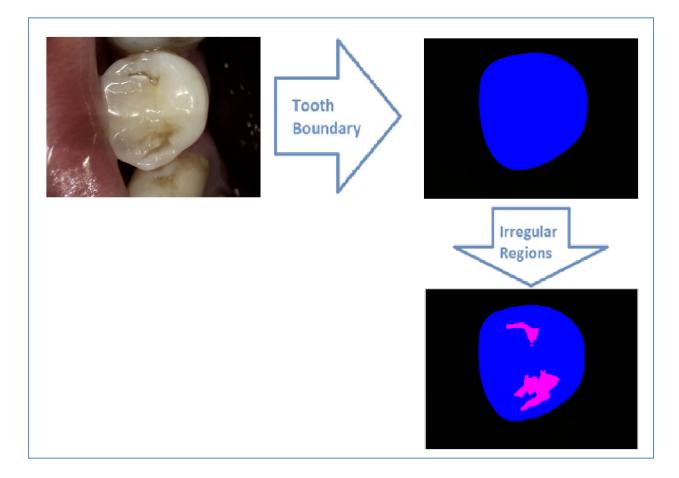


Figure 4.1: Segmentation workflow



# Chapter 5

# Feature Selection and Classification

### **5.1Feature Extraction**

The features are measures calculated from  $10 \times 10$  windows scrolled over the entire enamel surface and the detected irregular region, separately. Feature extraction over windows presents local information in feature space. Experimental testing revealed that  $10 \times 10$  window size performed best for this application. Window sizes ranging from  $7 \times 7$  to  $12 \times 12$  were tested and  $10 \times 10$  had the best performance by visual evaluation and also final accuracy. The designed system extracted 87 region-based and pixel-based features from both enamel (as control) and the irregular regions separately based on color space and Fourier transforms. Each feature is described below.

Mean of matrix elements in a  $10 \times 10$  window calculated as described in equation 5.1.

Mean of matrix elements 
$$= \frac{1}{100} \sum_{i=1}^{10} \sum_{j=1}^{10} f(i,j)$$
 (5.1)

An image gradient, which described in equation 5.2 is a directional change in the intensity or color in the image.

Gradient = 
$$\Delta f = \frac{\Delta f}{\Delta x} + \frac{\Delta f}{\Delta y}$$
 (5.2)



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Table 5.1 describes how the features created in 10 by 10 windows level and how their statistical measures create the final feature pool for each image. Subscript 2 means the operation has been done in 10 by 10 windows level and subscript 1 means has been done in global level. The table shows 43 possible features for an image. These 43 features were calculated for the tooth surface except irregular regions mask; an example of such a mask is the blue regions in figure 7.15. The other 43 features were calculated for the irregular regions mask; an example of such a mask is the red regions in figure 7.15. The ratio of the total area of irregular regions to the total tooth area is the last feature .Finally 87 features have been calculated for each image.

	First Component	Second Component	First Component	First Component	Second Component	First Component	Fourier Transform
	of RGB	of RGB	of RGB	of HSV	of HSV	of HSV	Tansiorm
	Color Space	Color Space	Color Space	Color Space	Color Space	Color Space	
	(RED)	(GREEN)	(BLUE)	(Hue)	(Saturation)	(Value)	
Mean <sub>1</sub> (Mean <sub>2</sub> )	Х	Х	Х	Х	Х	Х	X
Std <sub>1</sub> (Mean <sub>2</sub> )	X	X	X	X	X	X	
Median <sub>1</sub> (Mean <sub>2</sub> )	X	X	X	X	X	X	
Maximum <sub>1</sub> (Mean <sub>2</sub> )	X	X	X	X	X	X	
Maximum <sub>1</sub> (Gradient <sub>2</sub> )	Х	X	X	X	X	X	
Mean <sub>1</sub> (Maximum <sub>2</sub> )	X	X	X	X	X	X	
Mean <sub>1</sub> (Minimum <sub>2</sub> )	Х	X	X	X	X	X	

#### **Table 5.1: Feature Extraction**

#### **5.2 Feature Selection and Classification**

While the dataset for this research has representation in each of the ICDAS categories, it does not have sufficient examples of some of ICDAS scores to warrant individual score classification. ICDAS Scoring System has seven scores, defining a state of caries development. 0: Sound Tooth – 1: First visual Change in Enamel – 2: Distinct Visual Change in Enamel- 3: Localized Enamel Breakdown- 4: Underlying Dentin Shadow- 5: Distinct Cavity with Visible Dentin- 6: Extensive Cavity with Visible Dentin. Thus the traditional ICDAS classification has been modified for this



research to specify: Sound occlusal (ICDAS Score 0), Initial caries (ICDAS Score 1 or 2), Moderate caries (ICDAS Score 3-5), severe caries (ICDAS Score 6) [43].

ICDAS scores are available for 94 images; including 88 images of two in-vitro data sets and 6 images of 6 unique teeth of in-vitro data set. Based on the criteria for caries treatment planning and a re-classification of ICDAS scores, the ICDAS scores in this study were grouped into three overview classes (Figure 5.1). Other studies also re-categorized ICDAS scores to fewer categories than the initial 7 Scores [8, 15, 20-21, 40-41, 43-44].

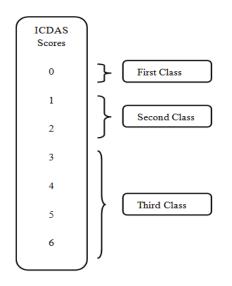


Figure 5.1: Re-categorization map of seven ICDAS scores into three classes

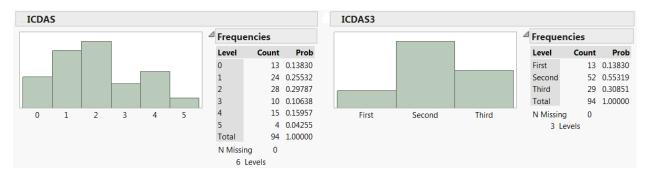


Figure 5.2: The histogram of ICDAS and reduced ICDAS3 for 94 images

Figure 5.2 shows the histogram of ICDAS scores and the reduced ICDAS classes; ICDAS3 for



the classification data set (94 images). It is clear that ICDAS3 is an imbalance data set. The classifier constructed to minimize the overall error rate; it will tend to focus more on the prediction accuracy of the majority class, which often results in poor accuracy for the minority classes. There are multiple approaches to cope with imbalance data sets, such as under sampling, over sampling and cost sensitive learning. For our size of data set, cost sensitive learning is the best approach. Since the classifier tends to be biased towards the majority class, a heavier penalty on misclassifying the minority class should be defined. A weight has been assigned to each class, with the minority class given larger weight (i.e., higher mis- classification cost). A weighted random forest classifier used to train the final classification model.

In order to compare performance of the ICDAS based caries detection system, another system has been trained which uses the histological scores instead of ICDAS. Histological assessment is known as the gold standard in the cariology field. The histological scores are as follows; 0: No enamel demineralization or a narrow surface zone of opacity, 1: Enamel demineralization (opacity) limited to the outer 50% of the enamel layer, 2: Demineralization (brown discoloration) involving between 50% of the enamel and 1/3 of the dentin, 3: Demineralization (brown discoloration) involving the middle third of the dentin, 4: Deminerlization (brown discoloration) involving the inner third of dentin. Due to small number of teeth in each class and based on the criteria for caries treatment planning and a re-classification of histological scores, the histological scores in this study were grouped into three overview classes. Histological scores 0 and 1 re-named to classes Histo1 and Histo2 respectively and histological scores is presented in Figure 5.3.

Basically feature classification should have been done separately for two systems. The first



system uses ICDAS scores and the second system uses histological scores.

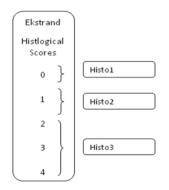


Figure 5.3: Re-categorization map for Ekstrand histological scores: five histological scores mapping into three classes

To reduce the feature space dimension, a filter based method and a super classifier applied to the extracted features respectively. First the Wilcoxon signed-rank test used to evaluate all the 87 features with respect to three ICDAS classes. Wilcoxon is a non-parametric statistical hypothesis test used to assess whether the population mean ranks differ within the classes or not [33]. It can be used as an alternative to the paired Student's t-test, t-test for matched pairs, or the t-test for dependent samples when the population cannot be assumed to be normally distributed [33]. Wilcoxon signed-rank test has been used instead of paired Student's t-test because some of the 87 features were not normally distributed.

With the application of Wilcoxon signed-ranked test 32 features out of 87 features showed statistical difference within the three classes. The rest of the features; meaning 55 other features filtered out at this point due to not showing statistical difference among three classes. The remaining 32 features called "reduced feature pool" [45]. The details of Wilcoxon signed-ranked test have been provided in the appendices.



A heuristic super classifier method used to select the high ranked features as well as a classification model [45]. Super Classifier encompasses four classification methods to perform the classification task including C4.5 decision tree [46], Support Vector Machine (SVM) [47], Random Forest classifier [48] and Artificial Neural Network classifier [30]. These four classification methods showed successful performance in medical decision support systems. The super classifier uses the ten-fold cross validation to avoid over-fitting. For each image a reduced feature pool of 32 are available. Feature ranking was done by using information gain ratio method; ranking has been assigned to the features according to relevance to the categories. An extensive search has been performed to find the best features and the best classifier. Through the extensive search the number of high ranked features varied from 5 to 32 for each as well as the classification methods and classification parameters. Five was chosen as the minimum number of selected features because any less than that would not provide sufficient information to classify items into three classes. Figure 5.4 describes the multi stage feature selection and classification algorithm. Ten-Fold cross validation was used to validate the results in all cases. Based on the best performance, Random Forest Classifier with the reported setting was selected for both systems.

Ten-fold cross validation, which is a standard process to assess the accuracy and reliability of the models developed from data, was used to validate the resulting computational model. In ten-fold cross-validation, the combined dataset was randomly partitioned into 10 subsets. Of the 10 subsets, a single subset was kept as the validation set for testing the model, and the remaining 9 subsets were used for training of the model. Cross-validation was then repeated 10 times (the folds) in a "round-robin" process, with each of the 10 subsets used exactly once as the validation sets.



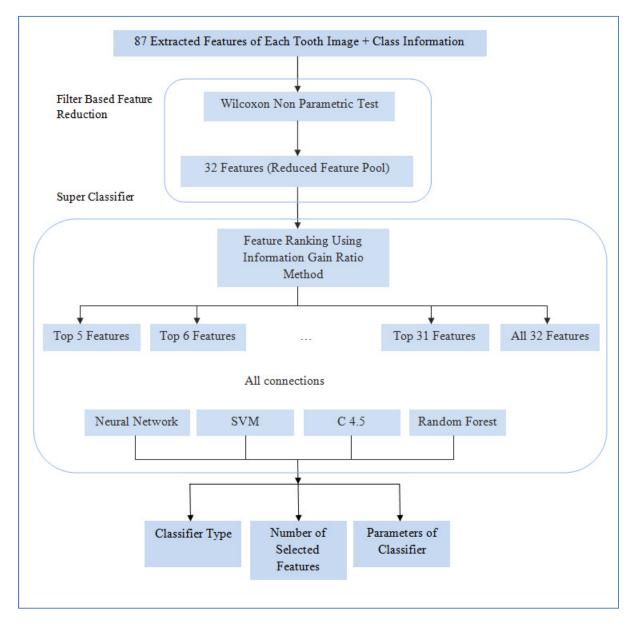


Figure 5.4: Filter based feature reduction and super classifier diagram



# Chapter 6

# Description of Data Set

### **6.1 Introduction**

The data used in this study consist of three sets of color images; while all sets of images contain occlusal surface of a molar or pre-molar teeth. All the teeth are free from restoration on occlusal surface. The first two data sets are in-vitro images taken from extracted teeth in different experiments. The third data set consists of in-vivo images of a molar or pre-molar tooth of interest and the adjacent teeth.

#### 6.2 In-Vitro Data Sets

The in-vitro data set consists of 88 images of the occlusal surface of an extracted tooth, one image per tooth surface. The data set consists of images of teeth with different stages of caries. In-vitro data set consists of 72 and 16 images taken in two different experiments.

#### 6.2.1 First In-Vitro Data Set

The first in-vitro data set includes 72 images of size 1440×2160×3 with a light blue background. Where the first two image size dimensions represent pixel size and the third dimension represents RGB (Red, Green, Blue). The images were taken with digital camera with 1440×2160 resolution. Each extracted tooth was examined and scored by ICDAS experts in a previous cariology study conducted in 2001 [19]. Each image was scored based on ICDAS [19] by



multiple experts after visual and tactile examination of the extracted tooth, where the scores were different, the final ICDAS score was determined by consensus decision making after a discussion session. The Ekstrand method histological assessment scores are available for 57 images of this data set, which has been used to develop a separate scoring system to verify the ability of the system to be trained by histological assessment instead of ICDAS assessment.



Figure 6.1: One example of First In-Vivo Data Set

#### 6.2.2 Second In-Vitro Data Set

This data set consists of 16 images of size 768×1024×3 with a dark background. This image set was taken by a KODAK a1500 intraoral camera which has a ½.5 inch CMOS with 1024×768 pixels and lighting is provided by super white LEDs / class 1 LED according to IEC 60825-1. Each extracted tooth of this data set was analyzed and scored at VCU School of Dentistry during the VCU-ICDAS Training Workshop in 2011 [43]. Each image was scored based on the International Caries Detection and Assessment System (ICDAS) [19] by multiple experts after visual and tactile examination, where the scores were different, the final ICDAS score was determined by consensus decision making after a discussion session.





Figure 6.2: One example of Second In-Vivo Data Set



Figure 6.3: KODAK A1500 Intraoral Camera [55]

### 6.3 In-Vivo Data Set

This data set consists of 50 images of size 768×1024×3 taken with the same KODAK a1500 intraoral camera. The images were taken from one person's molar and premolar teeth. There are several images of each tooth which the percentage of the tooth of interest's appearance and the adjacent teeth and also the tongue and lip are different from one image to another. This data set consists of six unique teeth. Each unique tooth is examined and scored by ICDAS expert in VCU School of Dentistry.

The whole in-vivo data set (50 in-vivo images) has been used in order to develop the segmentation method for in-vivo image; but for training and testing of the classification



algorithm only one image per tooth (6 images) was used to keep the consistency with in-vitro images. As it was discussed earlier in-vivo data set consists of 50 images of 6 unique teeth.

The tooth of interest should be "conceptually obvious" in the image, which means an image can include the tooth of interest's occlusal surface and adjacent teeth and tissues but the image should not include two whole occlusal surfaces of two teeth in a way that it's hard to tell which tooth is the subject of evaluation.





Figure 6.4: Example of three different views of one unique tooth in In-Vivo Data Set



# Chapter 7 Results and Discussion

Chapters 3 and 4 described the methodologies used for tooth segmentation with regards to cariology guidelines. Chapter 5 presented the methodology used to extract and select the feature in order to predict the caries severity class. The results obtained using all these methodologies are given in this chapter. These results include results obtained at several stages starting from preprocessing to outcome prediction. The sample results presented in this chapter are the results obtained from three different sets of images of tooth surface which were described in chapter 6. Three different data sets which encompass in-vitro and in-vivo images and represent different lighting and background texture allow for the better assessment and evaluation of the method's performance.

# 7.1 Pre-Processing Results

This section provides the example results for pre-processing steps; one example per data set. This includes the RGB image mapping to HSV space, RGB image mapping to gray scale and histogram equalization of gray scale image.



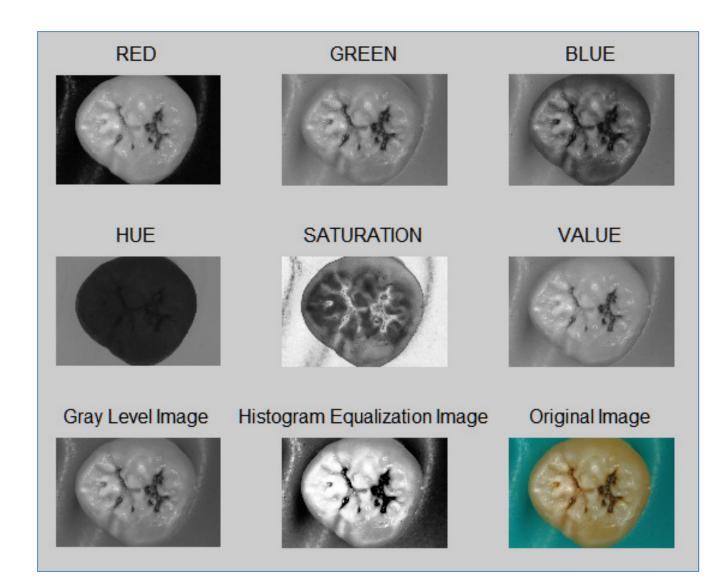


Figure 7.1: Pre-processing- Tooth #24 - First In-Vitro Data Set



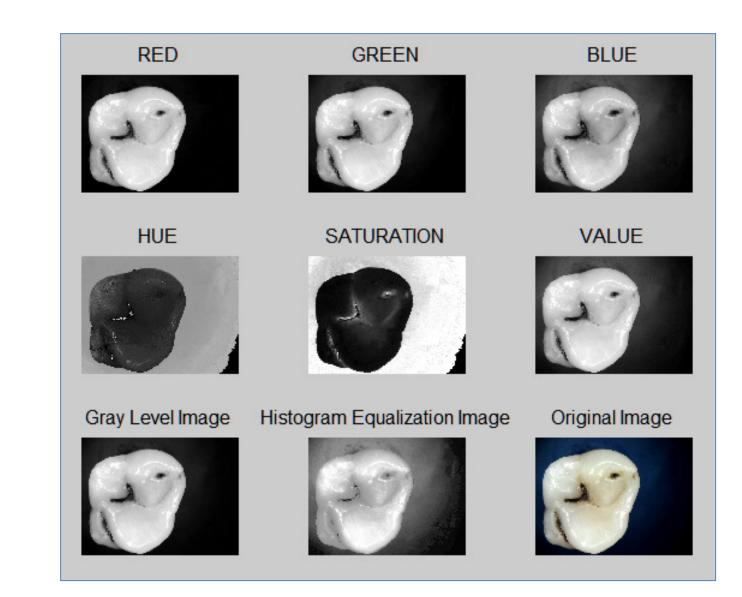


Figure 7.2: Pre-processing- Tooth #14- Second In-Vitro Data Set



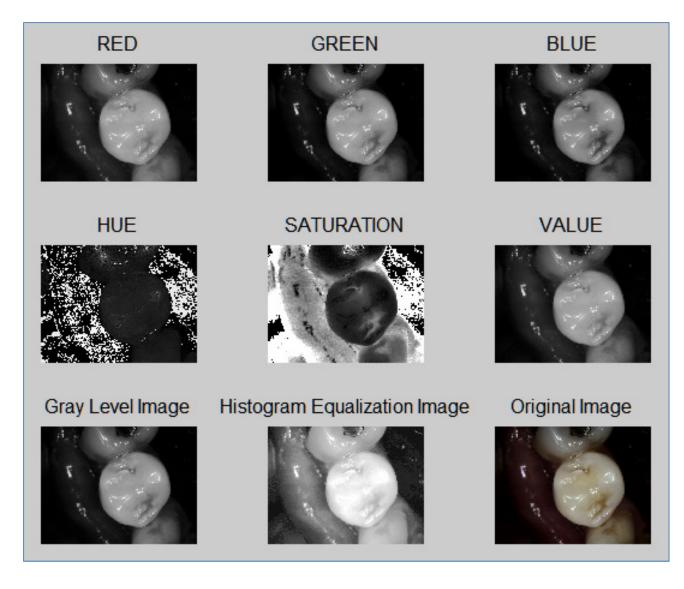


Figure 7.3: Pre-processing- Tooth #23 - In-Vivo Data Set

# 7.2 Initial Seed Selection Results

Figure 7.4 illustrates the accumulation array which is calculated for one example image of the second in-vitro data set. Figure 7.5 shows the graphical illustration of CHT output which is a set of circle centers and radii, finally the mean value of circle dimensions array selected as the seed point which is illustrated in figure 7.6. The high values in the accumulation array (hot spots) denote the possible centers of detected circle centers.



Figures 7.7 to 7.9 are showing the same information as figures 7.4 to 7.6 for an example of invivo data set.

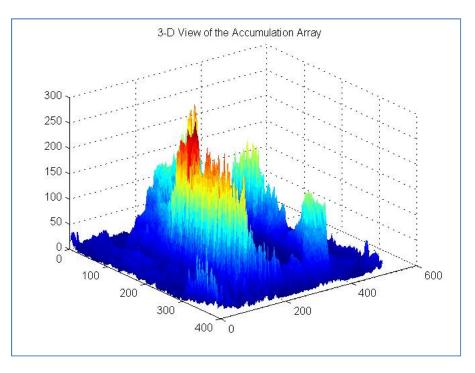


Figure 7.4: Accumulation array of tooth #4 of Second In-Vitro Data Set



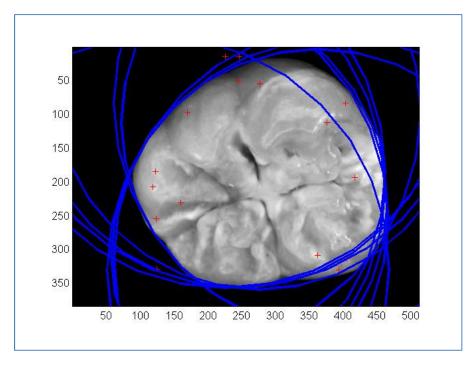


Figure 7.5: Output of CHT (multiple circle centers and radii) for the same image as figure 7.4

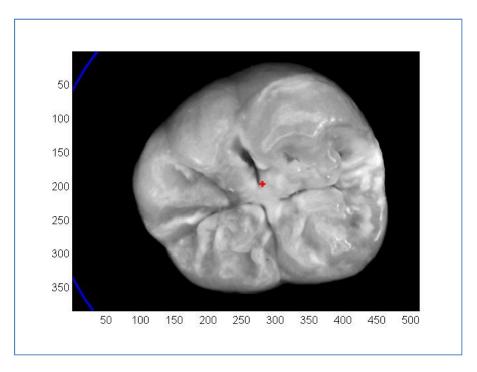


Figure 7.6: Final seed for the same image as figure 7.4



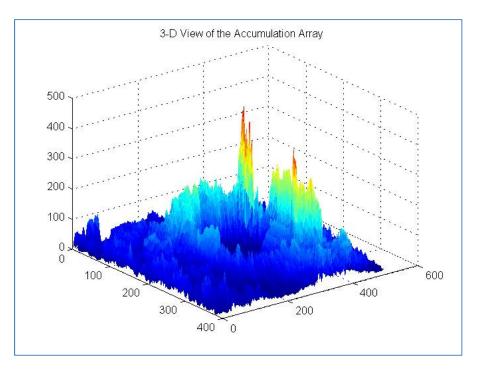


Figure 7.7: Accumulation array of tooth #12 of In-Vivo Data Set

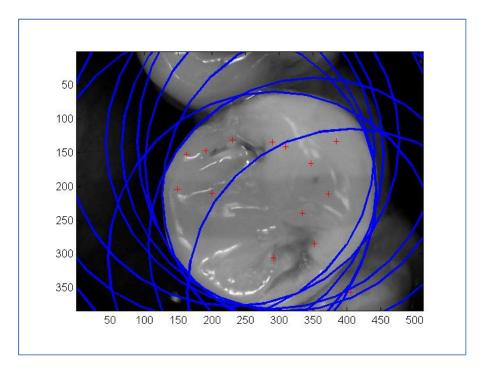


Figure 7.8: Output of CHT (multiple circle centers and radii) for the same image as figure 7.7



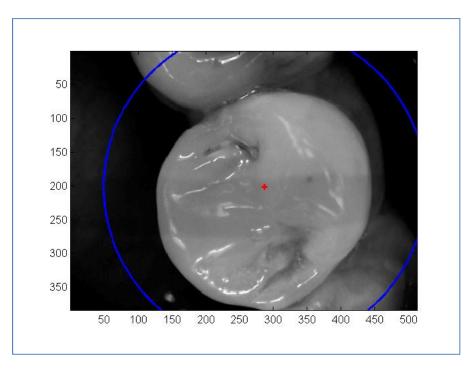


Figure 7.9: Final seed for the same image as figure 7.7

### 7.3 Region Growing and Active Contour Model Results

In this section multiple examples of region growing performance and active contour model performance are presented. The segmentation of in-vivo images is more challenging when compared to in-vitro images. There are multiple examples in which the region growing algorithm grows to a part of adjacent tooth or stops at a change of texture or color especially near the tooth borders. In addition then the active contour model tends to merge to the tooth edge. Figure 7.10 represents one of these examples. Two different images of one particular tooth are shown in figure 7.11 and figure 7.12 with the segmentation results. While strict guidelines for clinical image acquisition may be defined, the accurate performance of the method in presence of various adjacent organs and multiple views of the same surface is important.



Figure 7.13 and figure 7.14 are also illustrating two images of one tooth with their segmentation results. The image in figure 7.14 is not the ideal image to capture the occlusal surface but still the segmentation is successful.

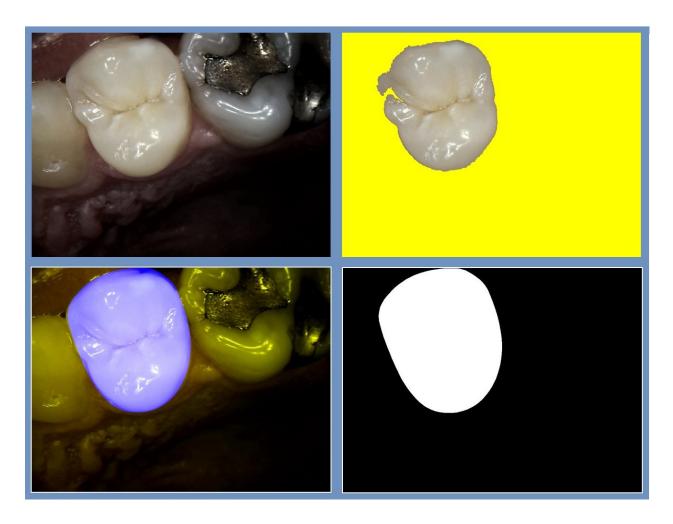


Figure 7.10: Image #2 -In-Vitro Data Set; From up left to down right

1: Original image, 2:Region growing output, 3: Active contour model Output, 4: Active contour model

output black and white mask



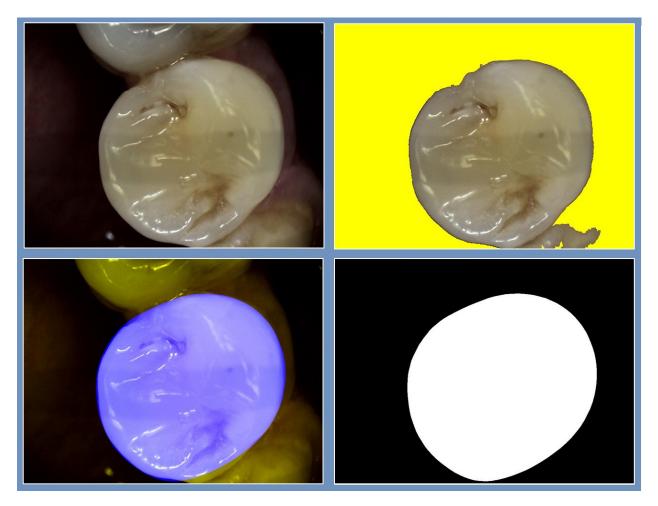


Figure 7.11: Image #12 -In-Vitro Data Set; From up left to down right

1: Original image, 2:Region growing output, 3: Active contour model Output, 4: Active contour model

output black and white mask



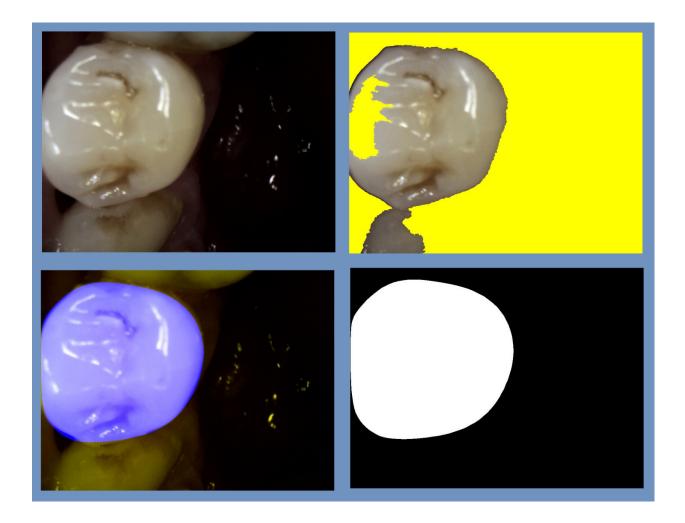


Figure 7.12: Image #6 -In-Vitro Data Set (The same tooth with image # 12 in Figure 7.11 ); From up left to down right; 1: Original image, 2:Region growing output, 3: Active contour model Output, 4: Active contour model output black and white mask



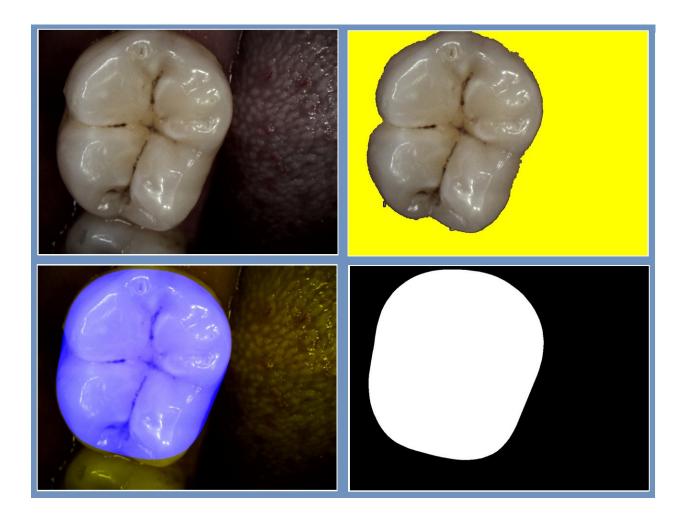


Figure 7.13: Image #39 -In-Vitro Data Set; From up left to down right

1: Original image, 2:Region growing output, 3: Active contour model Output, 4: Active contour model

output black and white mask



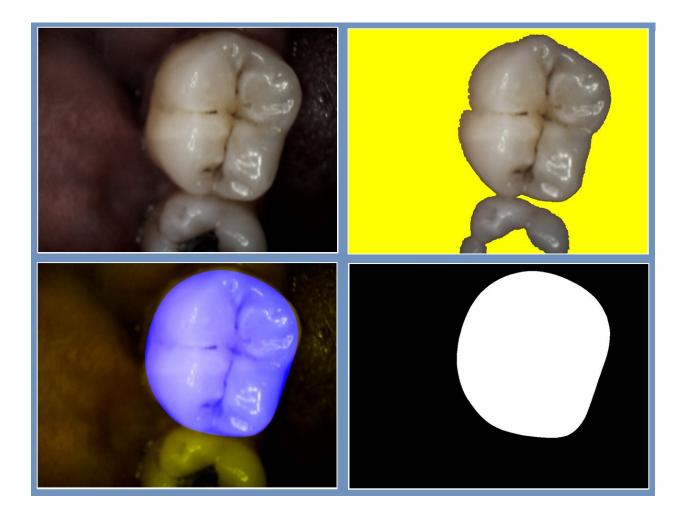


Figure 7.14: Image #42 -In-Vitro Data Set (The same tooth with image # 39 in Figure 7.13 ); From up left to down right; 1: Original image, 2:Region growing output, 3: Active contour model output, 4: Active contour model output black and white mask

Verma et al. reported 15 second segmentation time for image of size 481×321 on a system configured with Intel processor 2.63 GHz and 1 Gigabyte of RAM. The designed system segments the image of size 768×1024 in about 50 seconds with the same system configuration.

The following criteria were defined to compare segmentation results qualitatively : **Good** segmentation means mis-segmentation is less than or equal to 2% of tooth surface, **Acceptable** segmentation means mis-segmentation is greater than 2% and less than or equal to 5% of tooth surface and **Unacceptable** segmentation is defined by mis-segmentation greater than 5% of tooth



surface. The two following figures show the performance of tooth segmentation algorithm for 88 in-vitro images and 50 in-vivo images respectively. As hypothesized, segmentation algorithm performed much better for in-vitro images comparing to in-vivo images. The final step of tooth segmentation (S3), creates 98% Good segmentation results for in-vitro images where this is 82% for in-vivo image.

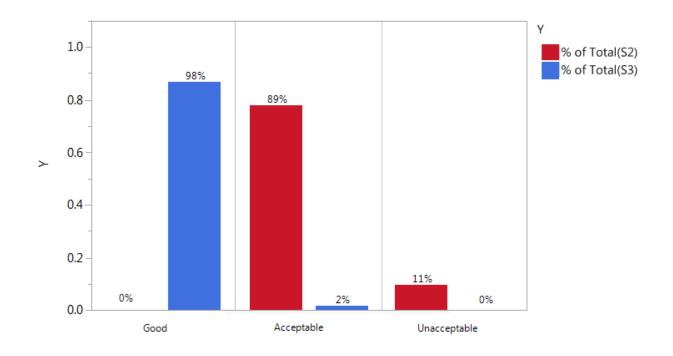
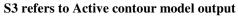


Figure 7.15: Tooth segmentation results for 88 in-vitro images- S2 refers to region growing step output and





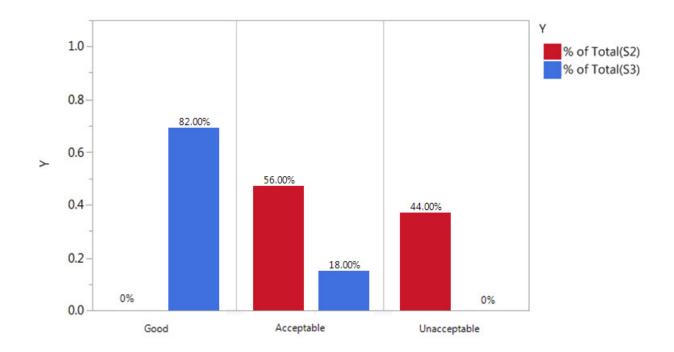


Figure 7.16: Tooth segmentation results for 50 in-vivo images- S2 refers to region growing step output and S3 refers to Active contour model output

## 7.4 Irregular Region Segmentation Results

The system segments the image into background, regular tooth surface and region of interest for the dentists. Once the segmentation of tooth and the background is performed, irregular region segmentation finds the irregularities inside the tooth. Figures 7.17 to 7.19 illustrate the background, regular tooth surface and irregular regions for one example image per dataset. In three color mask of the image, black defines background, blue is the regular tooth image and red maps the irregular regions. The features then will be calculated on the blue mask and red mask separately based on table 1.5 definition.



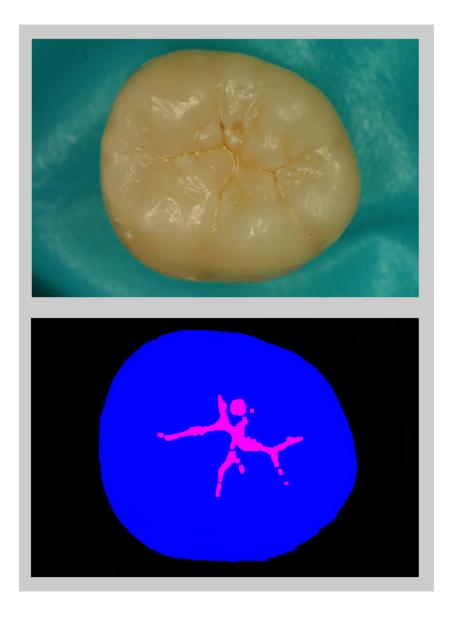


Figure 7.17: Image #23 - First In-Vitro Data Set – Irregular regions



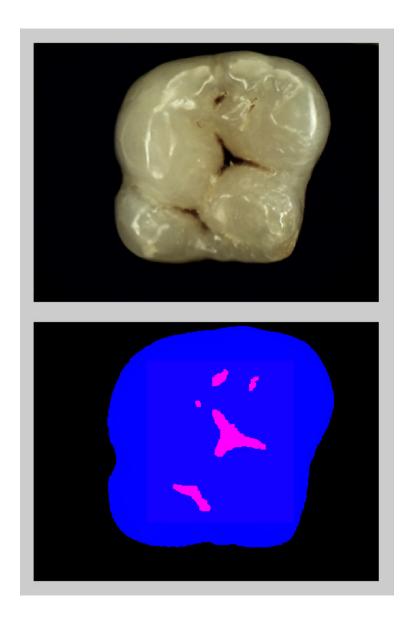


Figure 7.18: Image #2 - Second In-Vitro Data Set – Irregular regions



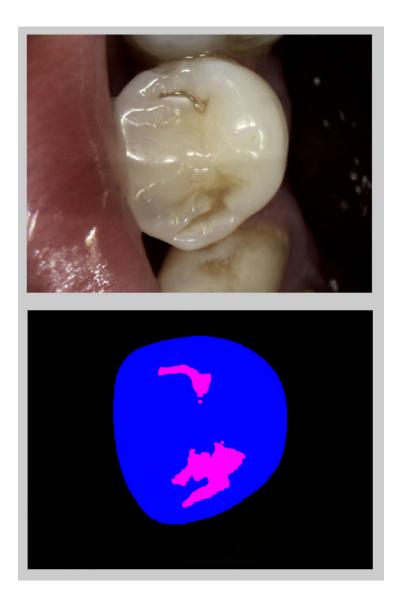
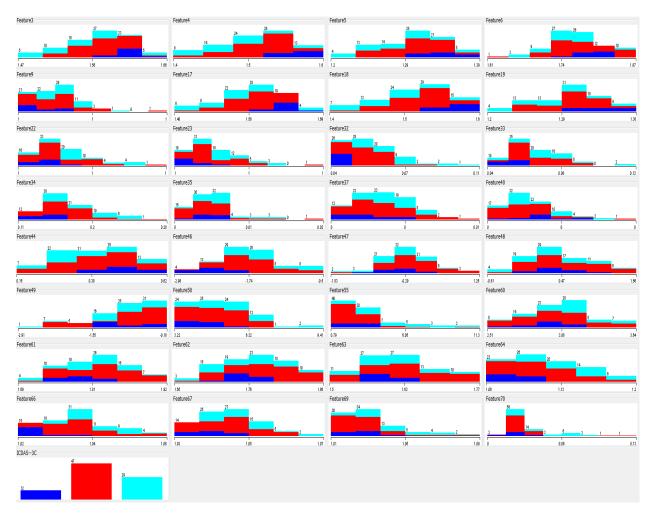


Figure 7.19: Image #24 - In-Vivo Data Set – Irregular regions

## 7.5 Feature Selection and Classification Results

The feature selection and classification performed on in-vitro data sets. 87 features extracted for each image. Wilcoxon signed-rank test used to evaluate all the 87 features with respect to three ICDAS classes and reduced the feature pool to 32 features. These 32 features applied to four different classifiers to select the classifier. Figure 7.20 illustrates the reduced feature pool for





each of the three ICDAS classes. These 32 features are showing statistically significant difference by three ICDAS classes.

Figure 7.20: Reduced feature pool (32 features) vs 3 ICDAS classes; dark blue class (0), red class (1,2) and

cyan class(3-6)

The final set of the features' box plot versus 3 ICDAS classes are shown in figure 7.21. Table 1 describes the final set of features.



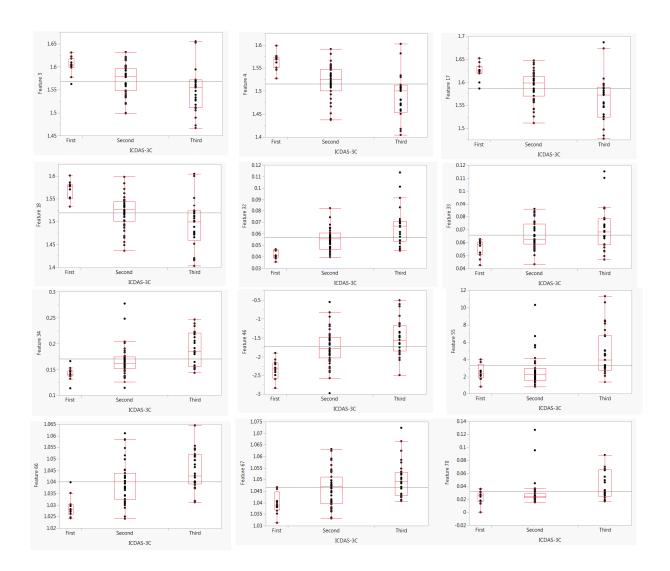


Figure 7.21: Final selected features (12 features) box plot grouped by 3 ICDAS classes



Row Number	Feature Number	Feature Description
1	3	Mean <sub>1</sub> (Mean <sub>2</sub> ) of Hue over tooth except irregular region
2	4	Mean <sub>1</sub> (Mean <sub>2</sub> ) of Value over tooth except irregular region
3	17	Median <sub>1</sub> (Mean <sub>2</sub> ) of Hue over tooth except irregular region
4	18	Median <sub>1</sub> (Mean <sub>2</sub> ) of Value over tooth except irregular region
5	32	Maximum <sub>1</sub> (Gradient <sub>2</sub> ) of Hue over tooth except irregular region
6	33	Mean1(Maximum2) of Hue over tooth except irregular region
7	34	Mean <sub>1</sub> (Maximum <sub>2</sub> ) of Value over tooth except irregular region
8	46	Mean <sub>1</sub> (Mean <sub>2</sub> ) of Hue over irregular region
9	55	Mean <sub>1</sub> (Mean <sub>2</sub> ) of Saturation over irregular region
10	66	Mean <sub>1</sub> (Maximum <sub>2</sub> ) of Hue over irregular region
11	67	Mean <sub>1</sub> (Maximum <sub>2</sub> ) of Value over irregular region
12	70	Irregular region pixel numbers to total tooth surface pixel numbers

 Table 7. 1: Final selected features description (12 features)

	0 (Calculated score)	1-2 (Calculated score)	3-6 (Calculated score)	Sum of Rows
0 (ICDAS experts' score)	11	2	0	13
1-2 (ICDAS experts' score)	2	47	3	52
3-6 (ICDAS experts' score)	0	4	25	29
Sum of Columns	13	53	28	

Table 7.2: Confusion Matrix; this table shows the confusion matrix for the designed system, comparing it to the ICDAS scores. The designed system's scores has been generated by the system and where ICDAS experts' scores are the scores that ICDAS experts assigned to each tooth after examining the extracted tooth as well as looking at the images of occlusal surface

The final performance of the random forest classifier for 94 of combined data sets described here. The extracted features of each segmentation method were used to train and test Random Forest Classifier, selecting the top 12 high-ranked features that gave us the best performance. The 12 selected features represented each image and the Random Forest Classifier categorized



each image to one of three classes using Ten Fold Cross Validation. For three-classes, classification resulted in 88.3% of accuracy, 92.0% of specificity, and 86.0% of sensitivity. Analysis of variance (ANOVA) on selected features regarding the output categories, showed that the selected features were statistically significant (p-value=0.05).

The classification of the subset of images which histology assessment is available for them (57 images) resulted in 76.4% of accuracy, 76.6% of specificity, and 76.4% of sensitivity.

The confusion table, Table 7.2, further exhibits the designed system's performance, by comparing it to ICDAS experts' score, which was treated as the gold standard in this study. Table 7.1 shows system performance based on three classes. In our study, ICDAS score "0" matched the system's score 11 out of 13 times. The additional two images were scored "1-2" by the system. ICDAS scores "1-2" matched the system's score 47 out of 52 times, with the rest of the images divided between scores "0" and "3-6". ICDAS scores "3-6" matched the system's score 25 out of 29 times, with additional two images scoring at "1-2" by the system. So the class accuracy for ICDAS score "0" is 84.6%, the class accuracy for ICDAS score "1-2" is 90.4% and the class accuracy for ICDAS score "3-6" is 86.2%. The class accuracies are still different and higher for the majority class but with the use of weighted random forest classifier, the class accuracy difference is acceptable.

### 7.6 Alternative System

In the designed system, grayscale image was used as the input of modified circular Hough transform algorithm to find the initial seed for region growing algorithm and the rest of image processing algorithms applied to HSV representation of the image (see figure 2.3). During the investigation, Value- which is the third component of HSV space- appeared to be very similar to the intensity of grayscale image. An effort made to investigate the effect of substitution of Value



in HSV space instead of intensity of the grayscale image. The alternative system would not significantly change the performance of the classifier nor the time complexity of the segmentation. The alternative system just reduces space complexity and can be considered as a solution in case space complexity is important for the application.

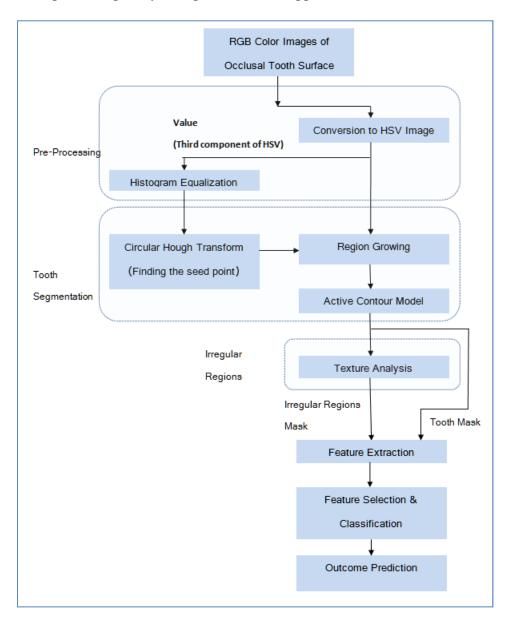


Figure 7.22: Diagram of the alternative system components



# Chapter 8

# Summary and Future Work

### 8.1 Summary

In this work, an automated system to detect and score caries lesions on a set of in-vitro and invivo images is developed and tested. An automated, user friendly system that eliminates the need for manual land-marking while using advanced feature extraction methods leading to a more reliable system for detection of early caries lesions has been developed. Two and three classes of caries conditions were observed and compared.

The designed system may provide an advantage over existing systems, which require expensive imaging sensors and external hardware. Another study which assessed performance of laser fluorescence device (LF) and LF pen and Fluorescence camera (FC) in detecting occlusal caries (Carious versus Non-Carious) by using histological gold standard described specificity and sensitivity levels of 100% and 85% for LF, 80% and 89% for LF pen, 80% and 74% for FC respectively [11], these are comparable to specificity and sensitivity of 88.3% and 86.0% in this study when classifying to three classes. The system's performance is better than LF pen and FC and almost similar to LF.

Other existing technologies for caries diagnosis include devices based on laser fluorescence (LF device, LF pen, F camera) or infrared (IR) laser fluorescence, referred to as quantitative laser or light fluorescence (QLF). Electrical conductance measurements (ECM), direct digital



radiography, Digital Imaging Fiber-Optic Trans-Illumination (DIFOTI) and simple Fiber Optic Trans-Illumination (FOTI), LED-based caries detector and less popular fluorescence spectrophotometer, MicroCT and heat induced detection technique [3, 7-12]. Data show varying degrees of sensitivity and specificity for *In-Vitro* and *In-Vivo* studies [3, 8-9, 13-15]. In addition to the unpredictable accuracy and the relatively high price, the existing technologies are user sensitive and require several steps in order to perform a clinical reading correctly. The value given by these caries detection devices is subjectively interpreted by the clinician and thus requires a trained dental professional to make a diagnostic or treatment decision.

Relatively easy-to-use, widely available and inexpensive hardware imaging techniques, such as an intraoral camera, may easily augment the decision making process of treatment provided to patients (surgical or preventive) and their overall impression of dental care they are receiving. The use of computer-aided systems to detect caries lesions based on different types of tooth images may be a way to increase the sensitivity of caries detection. There is promising evidence

that the automated caries detection system using intraoral photographs can provide high sensitivity and specificity rates when compared with ICDAS scores [39].

## 8.2 Future Work

The future work of this dissertation is as follows.

- Evaluation of the method on a larger in-vitro data set, following with the histological assessment of caries.
- Evaluation of the method for a larger in-vivo data set.



# **Bibliography**

- [1] http://www.who.int/mediacentre/factsheets/fs318/en/
- [2] Caries Annual Report NIDCR/CDC Dental, Oral and Craniofacial Data Resource Center (DRC), http://drc.hhs.gov/report/1\_3.htm accessed on 8/29/13.
- [3] A.F. Zandona, D.T. Zero, Diagnostic tools for early caries detection, The Journal of the American Dental Association 137 (12) (2006) 1675-1684.
- [4] M. Toraman Alkurt, I. Peker, H. Deniz Arishu, O. Bala, B. Altunkaynak, *In-Vivo* comparison of laser fluorescence measurements with conventional methods for occlusal caries detection, Lasers in medical science 23 (3) (2008) 307-312.
- [5] A.I. Ismail, Visual and Visuo-tactile Detection of Dental Caries, Journal of Dental Research83 (2004) C56-C66.
- [6] B. Senel, K. Kamburoglu, O. Ucok, S.P. Yuksel, T. Ozen , H. Avsever, Diagnostic accuracy of different imaging modalities in detection of proximal caries, Dentomaxillofacialradiology 39 (8) (2010) 501-511.
- [7] G.K. Rochlen , M.S. Wolff, Technological advances in caries diagnosis, Dental clinics of North America 55 (3) (2011) 441-452.
- [8] M.B. Diniz, T. Boldieri, J.A. Rodrigues, L. Santos-Pinto, A. Lussi, R.C. Cordeiro, The performance of conventional and fluorescence-based methods for occlusal caries detection: an *In-Vivo* study with histologic validation, Journal of the American Dental Association 143 (4) (2012) 339-350.



- [9] J.H. Berg, E.J Swift, Current caries detection devices, Journal of Esthetic and Restorative Dentistry 22 (5) (2010) 464-470.
- [10] L. Duruturk, A. Ciftci, S. Baharoglu, D. Oztuna, Clinical evaluation of DIAGNOdent in detection of occlusal caries in newly erupted noncavitated first permanent molars in cariesactive children, Operative Dentistry 36 (4) (2011) 348-355.
- [11] M. Graye, K. Markowitz, M. Strickland, G. Guzy, M. Burke, M. Houpt, *In-Vitro* evaluation of the Spectra early caries detection system, The Journal of clinical dentistry 23 (1) (2012) 1-6.
- [12] O. Samek, H.H. Telle, D.C.S. Beddows, Laser-induced breakdown spectroscopy: a tool for real-time, *In-Vitro* and *In-Vivo* identification of carious teeth, BMC Oral Health 1 (2001) 1-1.
- [13] A.M. Aktan, M.A. Cebe, M.E. Ciftci, E.S. Karaarslan, A novel LED-based device for occlusal caries detection, Lasers in Medical Science. 27 (2012) 1-7.
- [14] A. Jablonski-Momeni, D.N. Ricketts, S. Rolfsen, R. Stoll, M. Heinzel-Gutenbrunner, V. Stachniss, K. Pieper, Performance of laser fluorescence at tooth surface and histological section, Lasers in Medical Science 26 (2) (2011) 171-178.
- [15] K.W. Neuhaus, C. Longbottom, R. Ellwood, A. Lussi, Novel lesion detection aids, Monographs in oral science 21 (2009) 52-62.
- [16] V. Baelum, H. Hintze, A. Wenzel, B. Danielsen, B. Nyvad, Implications of caries diagnostic strategies for clinical management decisions, Community Dentistry and Oral Epidemiology 40 (3) (2012) 257-266.



- [17] International Caries Detection and Assessment System (ICDAS) Coordinating Committee, Criteria Manual International Caries Detection and Assessment System (ICDAS II), (2009).
- [18] A.I. Ismail, W. Sohn, M. Tellez, A. Amaya, A. Sen, H. Hasson, N.B. Pitts, The International Caries Detection and Assessment System (ICDAS): an integrated system for measuring dental caries, Community Dentistry and Oral Epidemiology 35 (3) (2007) 170-178.
- [19] A. Jablonski-Momeni, V. Stachniss, D.N. Ricketts, M. Heinzel-Gutenbrunner, K. Pieper, Reproducibility and accuracy of the ICDAS-II for detection of occlusal caries *In-Vitro*, Caries Research 42 (2) (2008) 79-87.
- [20] M.M. Braga, M.S. de Benedetto, J.C. Imparato, F.M. Mendes, New methodology to assess activity status of occlusal caries in primary teeth using laser fluorescence device, Journal of Biomedical Optics 15 (4) (2010) 047005.
- [21] 40J.A. Rodrigues, K.W. Neuhaus, M.B. Diniz, I. Hug, H. Stich, L. Karlsson, A. Lussi, Comparison among gold standard techniques used for the validation of methods for occlusal caries detection, Microscopyresearch and technique 75 (5) (2012) 605-608.
- [22] <u>http://faculty.mu.edu.sa/public/uploads/1380556025.9751The%20tooth%20supporting%</u> 20structures.pdf
- [23] D. L. Pham, C. Xu, and J. L. Prince. Current Methods in Medical Image Segmentation. *Annu. Rev. Biomed. Eng.*,315-337, 2000.
- [24] Z. Ma, J. M. Tavares, R. N. Jorge, and T. Mascarenhas. A review of algorithms for medical image segmentation and their applications to the female pelvic cavity. *Comput Methods Biomech Biomed Engin*.235-46, 2010.



- [25] N. Otsu. A Threshold selection method from gray-level histograms. *IEEE Trans. On Systems, Man, and Cybernetics*, 9(1): 62-66, 1979.
- [26] M. Sezgin. Survey over image thresholding techniques and quantitative performance evaluation. *Journal of Electronic Imaging*, 2004.
- [27] S. Wang, F. Chung, and F. Xiong. A novel image thresholding method based on parzen window estimate. *Pattern Recognition*, 41: 117-129, 2008.
- [28] M. Maitra and A. Chatterjee. A hybrid cooperative comprehensive learning based PSO algorithm for image segmentation using multilevel thresholding. *Expert Systems with Applications*, 34(2): 1341-1350, 2008.
- [29] K. Hammouche, M. Diaf, and P. Siarry. A multilevel automatic thresholding method based on genetic algorithm for fast image segmentation. *Computer Vision and Image Understanding*, 109: 163-175, 2008.
- [30] D. Y. Kim and J. W. Park. Connectivity-based local adaptive thresholding for carotid artery segmentation using MRA images. *Image and Vision Computing*, 23(14):1277-1287, 2005.
- [31] O. P. Verma, M. Hanmandlu, S. Susan, M. Kulkarni and P. Kumar. A Single Seeded Region Growing Algorithm for Color Image Segmentation Using Adaptive Thresholding. *IEEE International Conference on Communication Systems and Network Technologies*. 500-503, 2011.
- [32] M. Kass, A. Witkin and D. Terzopoulos. Snakes: Active contour models. *International Journal of Computer Vision*. (1) 321-331, 1988.



- [33] E. A. Gehan. A generalized Wilcoxon test for comparing arbitrarily singly-censored samples. *Biometrika* 52(1) 203-223, 1965.
- [34] H. D. Cheng, X.H. Jiang, Y. Sun and J. Wang. Color image Segmentation: advances and prospects. *Pattern Recognition* 34(12) 2259-2281, 2001.
- [35] <u>http://en.wikipedia.org/wiki/HSL\_and\_HSV</u>
- [36] S. Dreiseutl, L. Ohno-Machado, Logistic regression and artificial neural network classification models: a methodology review, Journal of Biomedical Informatics 35(5) (2002) 352-359.
- [37] L. Breiman, Random Forests, Machine Learning 45 (2001) 5-32.
- [38] I. Guyon, J. Weston, S. Barnhill, and V. Vapnik. Gene selection for cancer classification using support vector machines. *Machine Learning*, 46:389–422, 2002.
- [39] R. C. Gonzalez and R. E. Woods. Digital Image Processing, 2008.
- [40] R.O. Duda, P.E. Hart, Use of the Hough Transformation to Detect Lines and Curves in Pictures, Communication of the ACM 15 (1) (1972) 11-15.
- [41] M. Rizon, H. Yazid, P. Saad, A.Y.M. Shakaff, A.R. Saad, S. Masanori, S. Yaccob, M.R. Mamat, M. Karthigayan, Object detection Using Circular Hough Transform, American Journal of Applied Sciences 2 (12) (2005) 1606-1609.
- [42] I. Daubechies, Ten lectures on wavelets, CBMS-NSF conference series in applied mathematics, SIAM Ed (1992).
- [43] A.I. Ismail, M. Tellez, N.B. Pitts, K.R. Ekstrand, D. Ricketts, C. Longbottom, H. Eggertsson, C. Deery, J. Fisher, D.A. Young, J.D.B. Featherstone, R.W. Evans, GG. Zeller, D. Zero, S. Martignon, M. Fontana, A. Zandona, Caries management pathways preserve dental tissues and promote oral health, Community Dent Oral Epidemiol 41 (2013) e12–e40.



- [44] L. Ghaedi, R. Gottlieb, K. Najarian, Towards An Automated Caries Detection System Using Intra-Oral Photographs, AADR Annual Meeting Tampa, Florida, March 2012 Proceeding (2012).
- [45] A. Belle, M. Pfaffenberger, R. H. Hargraves and K. Najarian. An Automated Decision Making System for Detecting Loss of Attention in Individuals Using Real Time Processing of Electroencephalogram. *Biosignal Interpretetion-7<sup>th</sup> International Workshop*, 2012.
- [46] J.R. Quinlan, J. Ross, C4.5: programs for machine learning, Morgan Kaufmann Publishers (1943).
- [47] H.E. Strassler, L.G. Sensi, Least Square Support Vector Machine Classifiers, Neural Processing letters 9 (1999) 293-300.
- [48] M.S. de Oliveira, L. E. Betting, S. B. Mory, F. Cendes, G. Castellano. Texture analysis of magnetic resonance images of patients with juvenile myoclonic epilepsy. *Epilepsy & Behavior*, 27:22–28, 2013.
- [49] Sarkar, M.K. Biswas, and K. M. S. Sharma. A simple unsupervised MRF model based image segmentation approach. *IEEE Transactions on Image Processing*, 9, 2000.
- [50] R. Adams and L. Bischof, Seeded Region Growing, IEEE Transaction on Pattern Analysis and Machine Intelligence 16 (6) (1994)641-647.
- [51] Dean's blog, Virginia Commonwealth University, School of Dentistry, http://wp.vcu.edu/dentistrydean/2012/01/07/icdas-training-and-caries-managementstrategies/, accessed on 8/29/2013.
- [52] G.F. Olsen, S.S. Brilliant, D. Primeaux, K. Najarian, An Image-Processing Enabled Dental Caries Detection System, ICME International Conference on Complex Medical Engineering (2009).



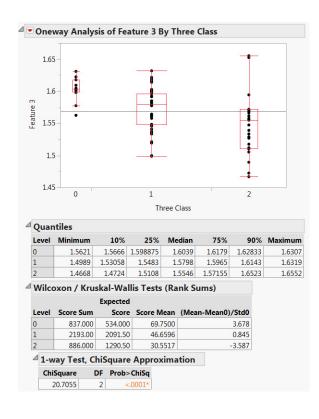
75

- [53] C. Huang, Q. Liu and X. Li. Color Image Segmentation by Seeded Region Growing and Region Merging. *IEEE Seventh International conference on Fuzzy Systems and Knowledge Discovery*. 533-536, 2010.
- [54] J. M. Pardo, D. Cabello and J. Heras. A snake for model-based segmentation of biomedical images. *Pattern Recognition Letters*. (18) 1529-1538, 1997.
- [55] <u>http://www.eurekafamilydental.com/services/oral-exams-x-rays/</u>

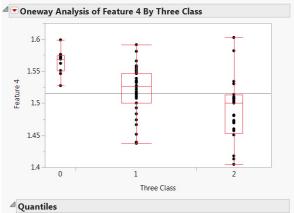


#### **APPENDICES**

The following are Wilcoxon signed-rank test analysis results for all extracted features which showed statistical significant difference among three ICDAS classes (reduced feature pool).





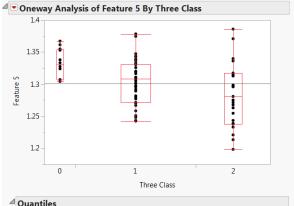


Level	Minimum	10%	25%	Median	75%	90%	Maximum
0	1.5272	1.53305	1.5508	1.569	1.574625	1.5919	1.5988
1	1.4372	1.46638	1.4999	1.5258	1.547	1.55656	1.5913
2	1.4047	1.4177	1.45365	1.5005	1.5137	1.5819	1.6027

<sup>⊿</sup> Wilcoxon / Kruskal-Wallis Tests (Rank Sums)

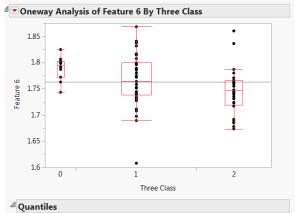
		Expected		
Level	Score Sum	Score	Score Mean	(Mean-Mean0)/Std0
0	889.000	534.000	74.0833	4.311
1	2164.00	2091.50	46.0426	0.602
2	863.000	1290.50	29.7586	-3.791



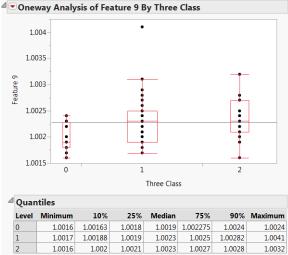


Quan	ittles						
Level	Minimum	10%	25%	Median	75%	90%	Maximun
0	1.3047	1.3047	1.30735	1.33065	1.3543	1.36551	1.367
1	1.2426	1.25048	1.2717	1.308	1.3305	1.34296	5 1.378
2	1.1987	1.2209	1.23805	1.2809	1.3178	1.371	1.385
Wilco	oxon / Kru	skal-Wall	is Tests	(Rank S	ums)		
		Expected					
Level	Score Sum	Score	Score M	ean (Mea	an-Mean0)	/Std0	
0	762.000	534.000	63.5	000		2.766	
1	2116.00	2091.50	45.0	213		0.201	
2	1038.00	1290.50	35.7	931		-2.237	
⊿ <b>1-</b> w	ay Test, C	hiSquare	Approx	imation			
Chi	Square I	DF Prob>	ChiSq				
1	10.0268	2 0.	0066*				





Level	Minimum		10%	25%	Me	dian	75%	9	0%	Maximum
0	1.7437	1.	74928	1.7718	1.7	9505	1.802275	1.81	899	1.8246
1	1.6078	1.	70582	1.7387	1	7643	1.7991	1.83	836	1.8686
2	1.6738	1	L.6775	1.71965	1	7478	1.7656	1.7	865	1.8599
Wilco	oxon / Kr	uska	al-Wal	lis Test	s (Ra	nk S	ums)			
		Ð	pected							
Level	Score Sun	1	Score	Score N	/lean	(Mea	n-Mean0)	)/Std0		
0	744.00	0	534.000	62.	0000			2.548		
1	2196.0	0	2091.50	46.	7234			0.870		
2	976.00	0	1290.50	33.	6552			-2.788		
⊿ <b>1-</b> w	ay Test,	ChiS	quare	Appro	xima	tion				
Chi	Square	DF	Prob>	ChiSq						
	11.2143	2	0	0037*						

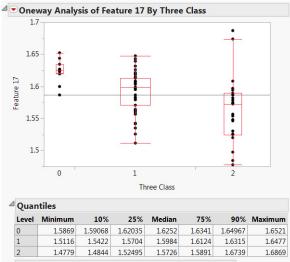


2	1.0017 1.0016	1.00188 1.002	1.0019 1.0021	1.0023 1.0023		1.00282 1.0028	1.0041 1.0032
⊿ Wilco	oxon / Kru	iskal-Wa	llis Tests	(Rank S	ums)		
		Expected	d				

Level	Score Sum	Score	Score Mean	(Mean-Mean0)/Std0
0	291.500	534.000	24.2917	-2.955
1	2114.00	2091.50	44.9787	0.185
2	1510.50	1290.50	52.0862	1.957
⊿ <b>1-</b> w	ay Test, Ch	iSquare	Approxima	ntion
Chi	Square D	F Prob>	ChiSa	

10.1715 2 0.0062\*

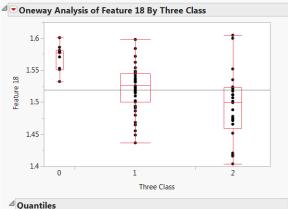




Wilcoxon / Kruskal-Wallis Tests (Rank Sums)

		Expected		
Level	Score Sum	Score	Score Mean	(Mean-Mean0)/Std0
0	849.500	534.000	70.7917	3.831
1	2184.50	2091.50	46.4787	0.774
2	882.000	1290.50	30.4138	-3.622

21 8144	2	< 0001



£							
Level	Minimum	10%	25%	Median	75%	90%	Maximum
0	1.5323	1.53779	1.55105	1.5774	1.580925	1.59647	1.601
1	1.4362	1.46742	1.5002	1.5256	1.5445	1.56388	1.5981
2	1.4033	1.418	1.4586	1.4996	1.52245	1.6001	1.6046
Wilco	oxon / Kru	skal-Wa	llis Tests	(Rank S	ums)		

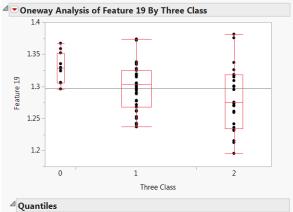
re Sum	Score	Score Mean	(Mean-Mean0)/Std0
		Score mean	(Weari-Weario)/ Stud
893.000	534.000	74.4167	4.359
2131.00	2091.50	45.3404	0.326
892.000	1290.50	30.7586	-3.533
	2131.00	2131.00 2091.50	2131.00 2091.50 45.3404

1-way Test, ChiSquare Approximation

 ChiSquare
 DF
 Prob>ChiSq

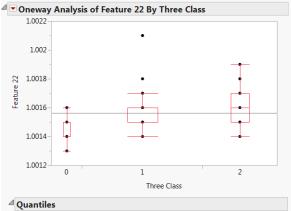
 24.9002
 2
 <.0001\*</td>





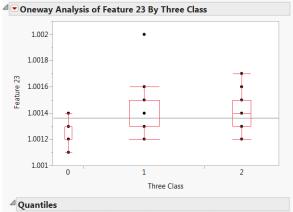
Level	Minimum	10%	25%	Median	75%	90%	Maximur
0	1.2959	1.2959	1.3056	1.3284	1.352	1.36449	1.367
1	1.2372	1.2426	1.2676	1.3036	1.3247	1.33784	1.373
2	1.1953	1.2154	1.2341	1.2753	1.3184	1.3757	1.38
Wilco	oxon / Kru	skal-Wall	lis Tests	(Rank S	ums)		
		Expected					
Level	Score Sum	•		ean (Mea	an-Mean0)	/Std0	
Level	Score Sum 772.000	•	Score M		an-Mean0)	<b>/Std0</b> 2.888	
		Score	Score M	333			

1	207	9.50	2091.50	44.2447		-0.096
2	106	54.50	1290.50	36.7069		-2.002
⊿	1-way Te	ation				
	ChiSquare	DF	Prob>	ChiSq		
	9,9374	2	0.0	0070*		



el Minim	num	10%	25	% Me	dian	75%	90	0%	Maximum
1.0	0013	1.0013	1.001	14	.0015	1.0015	1.001	.57	1.0016
1.0	0014	1.0014	1.001	15 1	.0015	1.0016	1.00	17	1.0021
1.0	0014	1.0014	1.001	15 1	.0016	1.0017	1.00	19	1.0019
lcoxon /	Krusk	al-Wa	llis Tes	sts (Ra	nk Su	ıms)			
	E	xpected	I I						
el Score	Sum	Score	Score	Mean	(Mea	n-Mean0)	/Std0		
31	2.500	534.000	) 2	26.0417			2.787		
19	82.50	2091.50	) 4	2.1809			0.941		
16	21.00	1290.50	) 5	5.8966			3.038		
-way Te	st, Chi	Square	e Appr	oxima	tion				
ChiSquare	DF	Prob>	ChiSq						
13.3584	2	C	.0013*						
	el Minin 1.0 1.0 1.0 1.0 1.0 1.0 1.0 1.0	el Minimum 1.0013 1.0014 1.	el Minimum 10% 1.0013 1.0013 1.0014 1.0014 1.0014 1.0014 1.0014 I.0014 Icoxon / Kruskal-Wal Expected Score Sum Score 312.500 534.000 1982.50 2091.50 1621.00 1290.50 -way Test, ChiSquare DF Prob	Iminimum         10%         25           1.0013         1.0013         1.001           1.0014         1.0014         1.001           1.0014         1.0014         1.001           1.0014         1.0014         1.001           1.0014         1.0014         1.001           Icoxon / Kruskal-Wallis Test         Expected           el         Score Sum         Score           312.500         534.000         2           1982.50         2091.50         4           1621.00         1290.50         5           -way Test, ChiSquare         DF         Prob>ChiSq	Iminimum         10%         25%         Me           1.0013         1.0014         1         1.0014         1           1.0014         1.0014         1.0015         1           1.0014         1.0014         1.0015         1           1.0014         1.0014         1.0015         1           Icoxon / Kruskal-Wallis Tests (Ra         Expected         E           el         Score Sum         Score Mean         312.500           312.500         534.000         26.0417           1982.50         2091.50         42.1809           1621.00         1290.50         55.8966           -way Test, ChiSquare Approximat         ChiSquare         DF	Minimum         10%         25%         Median           1.0013         1.0013         1.0014         1.0015           1.0014         1.0014         1.0015         1.0016           1.0014         1.0014         1.0015         1.0016           1.0014         1.0014         1.0015         1.0016           Icores results           Score Score Mean (Mean 312.500           312.500         534.000         26.0417           1982.50         2091.50         42.1809           1621.00         1290.50         55.8966          way Test, ChiSquare Approximation	Iminimum         10%         25%         Median         75%           1.0013         1.0014         1.0014         1.0015         1.0015           1.0014         1.0014         1.0015         1.0015         1.0016           1.0014         1.0014         1.0015         1.0016         1.0017           Icoxon / Kruskal-Wallis Tests         (Rank Sums)           Expected         Score         Score Mean         (Mean-Mean0)           312.500         534.000         26.0417         -           1982.50         2091.50         42.1809         -           -way Test, ChiSquare Approximation         -         -	Minimum         10%         25%         Median         75%         99           1.0013         1.0013         1.0014         1.0015         1.0015         1.0015         1.0016         1.0016           1.0014         1.0014         1.0015         1.0015         1.0016         1.0016         1.001           1.0014         1.0014         1.0015         1.0016         1.0017         1.001           Icoxon / Kruskal-Wallis Tests (Rank Sums)         Expected         Expected         Score Sum         Score Score Mean (Mean-Mean0)/Std0         312.500         534.000         26.0417         -2.787         1982.50         2091.50         42.1809         -0.941         1621.00         1290.50         55.8966         3.038         -way Test, ChiSquare Approximation         ChiSquare DF Prob>ChiSq	Iminimum         10%         25%         Median         75%         90%           1.0013         1.0013         1.0014         1.0015         1.0015         1.0015         1.0017           1.0014         1.0014         1.0015         1.0015         1.0016         1.0017           1.0014         1.0014         1.0015         1.0016         1.0017         1.0019           Icoxon / Kruskal-Wallis Tests (Rank Sums)         Expected         Expected         Score Sum         Score Score Mean (Mean-Mean0)/Std0           312.500         534.000         26.0417         -2.787         1982.50         2091.50         42.1809         -0.941           1621.00         1290.50         55.8966         3.038         -         -           -way Test, ChiSquare         DF         Prob>ChiSq         -         -

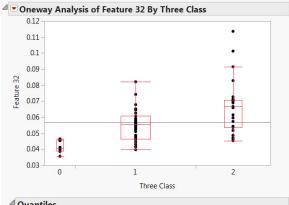
المنسارات



Level	Minimum	10%	25%	Median	75%	90%	Maximum
0	1.0011	1.0011	1.0012	1.0012	1.0013	1.00137	1.0014
1	1.0012	1.0012	1.0013	1.0013	1.0015	1.00152	1.002
2	1.0012	1.0013	1.0013	1.0014	1.0015	1.0017	1.0017

Wilcoxon / Kruskal-Wallis Tests (Rank Sums)

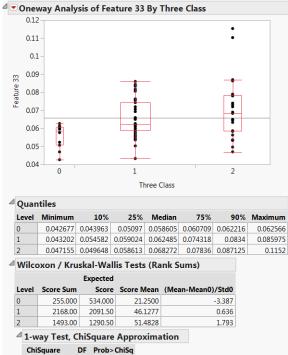
		E	xpected					
Level	Score S	Sum	Score	Score	Mean	(Mean-	Mean0)/Sto	dO
0	226	.500	534.000	1	8.8750		-3.8	52
1	204	6.50	2091.50	4	3.5426		-0.3	84
2	164	3.00	1290.50	5	6.6552		3.2	25
⊿ <b>1-</b> w	vay Tes	tion						
Chi	Square	DF	Prob>	ChiSq				
	19.9177	2	<.	0001*				



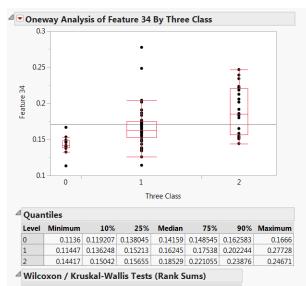
Level	Minimum	10%	25%	Median	75%	90%	Maximum
0	0.03558	0.036497	0.038985	0.041029	0.045662	0.046311	0.046412
1	0.03976	0.044351	0.046501	0.055522	0.060837	0.065762	0.0821
2	0.045302	0.047286	0.053796	0.066698	0.070724	0.09135	0.1136
		Expected					
Level	Score Sum	Score	Score N	lean (Mea	an-Mean0)	/StdO	
						Juli	
	116.000	534.000	) 9.0	5667		-5.077	
0	116.000 2030.00					-	
0 1 2		2091.50	) 43.	5667		-5.077	
0 1 2	2030.00	2091.50 1290.50	) 43. ) 61.0	5667 1915 0345		-5.077 -0.510	







12.2978 2 0.0021\*



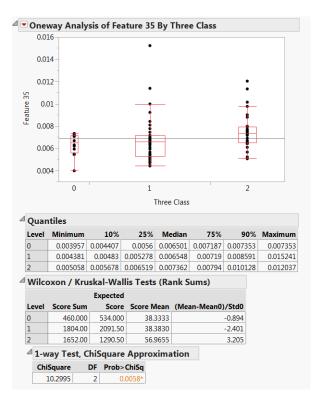
		Expected							
Level	Score Sum	Score	Score Mean	(Mean-Mean0)/Std0					
0	201.000	534.000	16.7500	-4.043					
1	2040.00	2091.50	43.4043	-0.427					
2	1675.00	1290.50	57.7586	3.409					
4 .									

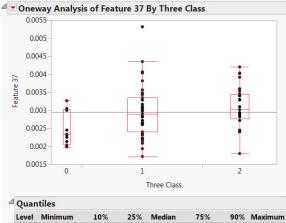
#### 1-way Test, ChiSquare Approximation

ChiSquare DF Prob>ChiSq

22.0588 2 <.0001



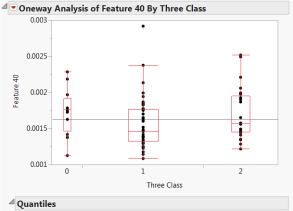




Level	Minimum	10%	25%	Median	75%	90%	Maximum
0	0.001976	0.00199	0.002053	0.002326	0.003032	0.003271	0.003271
1	0.001717	0.002144	0.0024	0.0029	0.003354	0.004045	0.005312
2	0.001792	0.002388	0.00277	0.003019	0.00345	0.004017	0.0042
· · · · · ·	oxon / Kru	isitar vva	mo resta	(Italia S	units)		
		Expected	1				
Level	Score Sum			lean (Mea	an-Mean0)	/Std0	
Level 0	Score Sum 314.000	Score	Score N	lean (Mea		/ <b>Std0</b> -2.669	

<sup>4</sup> 1-way Test, ChiSquare Approximation
ChiSquare DF Prob>ChiSq

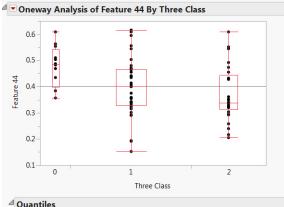




Level	Minimum	10%	25%	Median	75%	90%	Maximum
0	0.001127	0.001203	0.001469	0.001767	0.001921	0.002256	0.002287
1	0.001088	0.001239	0.001328	0.001456	0.001768	0.001952	0.002919
2	0.001213	0.001339	0.001449	0.001576	0.00195	0.002204	0.002521

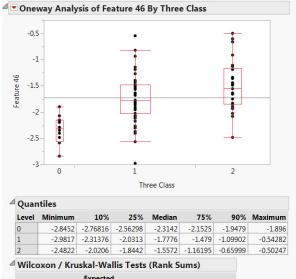
Wilcoxon / Kruskal-Wallis Tests (Rank Sums)

		E	xpected			
Level	Score	Sum	Score	Score Mean	(Mean-	Mean0)/Std0
0	659	.000	534.000	54.9167		1.514
1	180	3.00	2091.50	38.3617		-2.409
2	145	4.00	1290.50	50.1379		1.447
⊿ <b>1-</b> w	ation					
Chi	Square	DF	Prob>	ChiSq		
	6.1215	2	0.0	0469*		



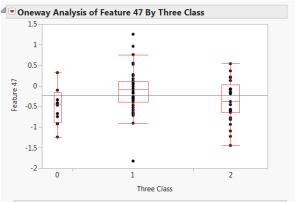
ittles						
Minimum	10%	25%	Median	75%	90%	Maximun
0.35653	0.35653	0.397623	0.48526	0.543363	0.59569	0.6095
0.15173	0.27089	0.33018	0.39926	0.46682	0.563718	0.6166
0.20527	0.23983	0.31247	0.33766	0.443765	0.55149	0.6086
oxon / Kru	skal-Wa	llis Tests	(Rank S	ums)		
	Expected	i				
Score Sum	Score	Score M	lean (Me	an-Mean0)	/Std0	
779.500	534.000	64.9	9583		2.979	
2070 50	2001 50	147	1447		1000000000	
2079.50	2091.50	44.2	2447		-0.096	
1057.00	1290.50		447		-0.096 -2.069	
1057.00	1290.50	36.4	1483			
1057.00 /ay Test, C	1290.50 hiSquare	36.4	1483			
	0.35653 0.15173 0.20527 ••••••••••••••••••••••••••••••••••••	Minimum         10%           0.35653         0.35653           0.15173         0.27089           0.20527         0.23983           Dxoor / Kruskal-War         Expected           Score Sum         Score           779.500         534.000	Ninimum         10%         25%           0.35653         0.35653         0.397623           0.15173         0.27089         0.33018           0.20527         0.23983         0.31247           DXDON / Krussel-Wallstein         Tests           Expected         Score M         Score M           779.500         534.000         64.9	Minimum         10%         25%         Median           0.35653         0.397623         0.48526           0.15173         0.27089         0.33018         0.39926           0.20527         0.23983         0.31247         0.33766           Dxoor / Kruskal-Wallis Tests (Rank S         Carrier (Rank S)         Carrier (Rank S)           Score Sum         Score Sum         Score (Marcine (Rank S))         Carrier (Rank S)           779.500         534.00         64.9583         Carrier (Rank S)	Minimum         10%         25%         Median         75%           0.35653         0.35653         0.397623         0.48526         0.543363           0.15173         0.27089         0.33018         0.39926         0.46682           0.20527         0.23983         0.31247         0.33766         0.443765           Expected           Score Sum         Kenes/Kenes	Minimum         10%         25%         Median         75%         90%           0.35653         0.35653         0.39762         0.48526         0.54336         0.59569           0.15173         0.27089         0.33018         0.39926         0.46682         0.56318           0.20527         0.23983         0.31247         0.33766         0.443765         0.5149           xxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxx





		Expected		
Level	Score Sum	Score	Score Mean	(Mean-Mean0)/Std0
0	155.000	534.000	12.9167	-4.602
1	2114.00	2091.50	44.9787	0.184
2	1647.00	1290.50	56.7931	3.160

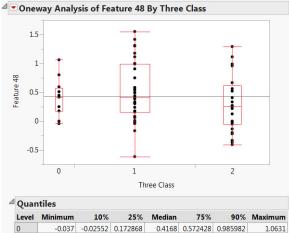
25.0748 2 <.0001\*



Quantiles									
Level	Minimum	10%	25%	Mediar	n 75%	90%	Maximum		
0	-1.2347	-1.14132	-0.87221	-0.4464	3 -0.16787	0.32295	0.32295		
1	-1.8311	-0.67041	-0.39956	-0.0957	2 0.12035	0.592576	1.2538		
2	-1.4487	-1.0893	-0.6447	-0.3763	9 0.030906	0.36168	0.54197		
Wilcoxon / Kruskal-Wallis Tests (Rank Sums)									
Expected									
Level	Score Sum	Score	Score N	lean (M	ean-Mean0	)/Std0			
0	378.000	534.000	31.	5000		-1.891			
1	2396.00	2091.50	50.9	9787		2.543			
2	1142.00	1290.50	39.3	3793		-1.314			
⊿ <b>1-</b> w	ay Test, C	hiSquare	Approx	cimatio	n				
Chi	Square	DF Prob>	ChiSq						

7.2959 2 0.0260\*

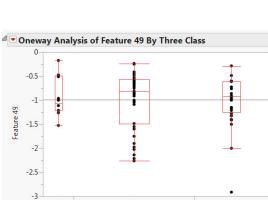


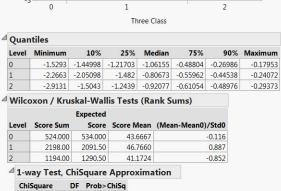


0	-0.037	-0.02552	0.172868	0.4168	0.572428	0.985982	1.0631
1	-0.61064	-0.02552	0.15844	0.40539	0.98758	1.3029	1.5568
2	-0.40862	-0.33789	-0.05639	0.25559	0.614505	0.98885	1.2965

Wilcoxon /	Kruskal-Wallis	Tests	(Rank Sums)
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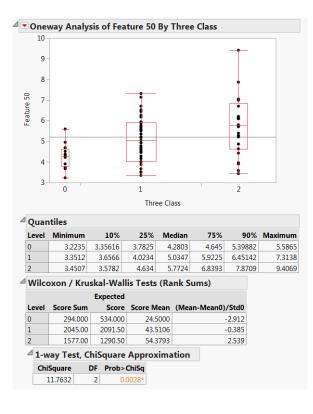
		E	pected			
Level	Score Su	m	Score	Score Mea	an (Mean-Mean0)/St	d0
0	532.0	00	534.000	44.333	-0.0	18
1	2307.	00	2091.50	49.085	51 1.7	99
2	1077.	00	1290.50	37.137	-1.8	91
⊿ <b>1</b> -w	/ay Test,	Chis	quare	Approxin	nation	
Chi	Square	DF	Prob>	ChiSq		
	3.9232	2	0.	1406		

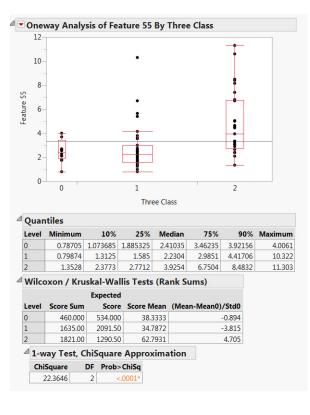




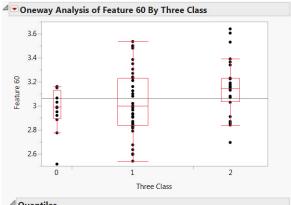
0.8746 2 0.6458











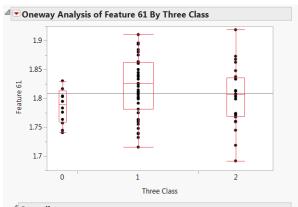
Quar	itiles						
Level	Minimum	10%	25%	Median	75%	90%	Maximum
0	2.5137	2.59275	2.895925	2.9883	3.1297	3.15838	3.1615
1	2.5395	2.6349	2.8415	2.9996	3.2311	3.4025	3.5347
2	2.6929	2.8536	3.0325	3.1427	3.2311	3.5315	3.6383

Wilcoxon / Kruskal-Wallis Tests (Rank Sums)

		Expected		
Level	Score Sum	Score	Score Mean	(Mean-Mean0)/Std0
0	415.000	534.000	34.5833	-1.441
1	1928.50	2091.50	41.0319	-1.360
2	1572.50	1290.50	54.2241	2.499

_	1-way Test,	Chis	quare Approximation
	ChiSquare	DE	Proh > ChiSa

6.8789 2 0.0321\*

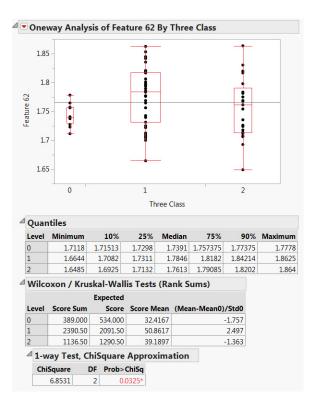


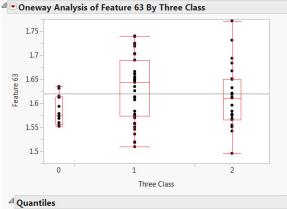
Level	Minimum	10%	25%	Median	75%	00%	Maximun
Level	winning	10 %	23/0	Weulan	13/0	50 %	Waximun
0	1.7404	1.74187	1.75915	1.78915	1.8133	1.82606	1.830
1	1.716	1.74646	1.781	1.8265	1.8625	1.88614	1.910
2	1.6918	1.7183	1.7691	1.806	1.83505	1.8679	1.918

#### Wilcoxon / Kruskal-Wallis Tests (Rank Sums)

		E	cpected				
Level	Score Su	ım	Score	Score	e Mean	(Mean-N	lean0)/Std0
0	376.5	00	534.000	3	31.3750		-1.909
1	2391	.00	2091.50		50.8723		2.501
2	1148	.50	1290.50	3	39.6034		-1.256
⊿ <b>1</b> -w	ay Test	, Chi	Square	Аррі	roxima	tion	
Chi	Square	DF	Prob>(	ChiSq			
	71580	2	0.0	1279*			







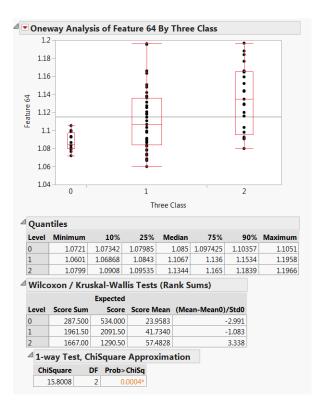
Level	Minimum	10%	25%	Median	75%	90%	Maximum
0	1.5523	1.55296	1.55585	1.57625	1.613975	1.63401	1.63
1	1.5098	1.544	1.5734	1.6442	1.6906	1.7246	1.739
2	1.4961	1.5425	1.5668	1.6109	1.6506	1.6942	1.771
whice	oxon / Kru	Expected	13 10303	(Rank 5	unisy		
Level	Score Sum	Score	Score M	ean (Mea	an-Mean0)	/Std0	
	374.000	534.000	31.1	667		-1.939	
0	374.000	554.000	51.1	007		-1.939	
0	2372.50		50.4			2.346	

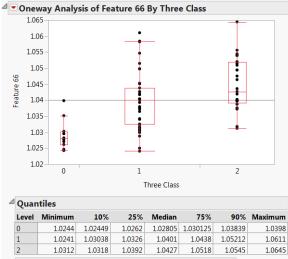
#### <sup>4</sup> 1-way Test, ChiSquare Approximation

ChiSquare DF Prob>ChiSq

6.6172 2 0.0366<sup>s</sup>







 Wilcoxon / Kruskal-Wallis Tests (Rank Sums)

 Expected

 Level
 Score Sum
 Core Gene Mean
 Mean-Mean0)/Std0

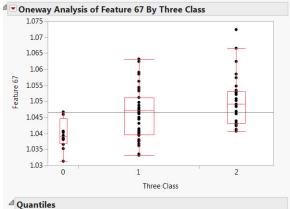
 0
 149.500
 534.000
 12.4583
 -4.670

1		2105.50	2091.50	44.7979	0.113			
2		1661.00	1290.50	57.2759	3.285			
⊿	<sup>4</sup> 1-way Test, ChiSquare Approximation							

ChiSquare DF Prob>ChiSq





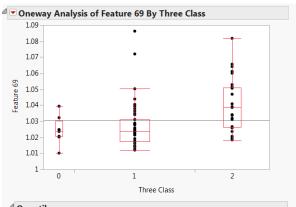


Level	Minimum	10%	25%	Median	75%	90%	Maximum
0	1.0314	1.03257	1.036975	1.03895	1.0446	1.0466	1.0466
1	1.0331	1.03666	1.0395	1.047	1.0512	1.05862	1.0631
2	1.0407	1.0412	1.0432	1.0491	1.0531	1.0625	1.0723

Wilcoxon / Kruskal-Wallis Tests (Rank Sums)

		Expected		
Level	Score Sum	Score	Score Mean	(Mean-Mean0)/Std0
0	250.500	534.000	20.8750	-3.441
1	2060.00	2091.50	43.8298	-0.259
2	1605.50	1290.50	55,3621	2.792

Chiloquare	UF	Flob>clibq
15,5401	2	0.0004*



Quan	ntiles										
Level	Minimum	10%	25%	Median	75%	90	%	Maximum			
0	1.0103	1.01327	1.020475	1.02345	1.030525	1.03	73	1.0394			
1	1.012	1.0141	1.0172	1.0236	1.0311	1.0412	28	1.0862			
2	1.0185	1.0205	1.0261	1.0388	1.051	1.064	43	1.0819			
<sup>⊿</sup> Wilco	<sup>4</sup> Wilcoxon / Kruskal-Wallis Tests (Rank Sums)										
	Expected										
Level	Score Sum	Score	Score M	lean (Me	an-Mean0)	/Std0					
0	418.500	534.000	34.8	3750		-1.398					
1	1751.00	2091.50	37.2	2553		-2.844					
2	1746.50	1290.50	60.2	241		4.044					

2	1746.50		1290.50		60.2241		4.	
<sup>4</sup> 1-way Test, ChiSquare Approximation								
Chi	Square	DF	Prob>	ChiSq				

 ChiSquare
 DF
 Prob>ChiSq

 16.4730
 2
 0.0003\*



