ORIGINAL RESEARCH

Skin Cancer Segmentation Based on Triangular Intuitionistic Fuzzy Sets

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Abstract

Malignant Melanoma is a dangerous form of skin cancer, and its detection is a challenging task as it appears in numerous ranges of size, shape, and shading with various skin tones. Also, artefacts like hairs, outlines, blood vessels, and boils add further complexity. A simplifed clustering method is proposed in this paper to improve melanoma detection while reducing time complexity.The triangular membership function (TMF) is used to detect the initial regions for obtaining initial centroids. These initial centroids are used to apply intuitionistic fuzzy c-means clustering. The TMF helps in identifying the initial clusters and regions and reduces the number of iterations needed for segmentation. The proposed method efectively detects skin cancer regions with an average accuracy of 90% and performs well.

Keywords Fuzzy sets · Intuitionistic fuzzy sets · Medical image segmentation · Image processing · Skin cancer

Introduction

According to the World Health Organization, skin cancer is one of the most common types of cancer worldwide. It ranks 19th among all diseases in men and women, with 324,635 cases reported in the year 2020. With the increased cases, early detection and timely treatment for melanoma are critical for patient survival.

Dermoscopy is a well-known in vivo unobtrusive imaging tool that uses captivated light to help dermatologists inspect pigmented skin sores based on a variety of morphological highlights. In spite of the fact that dermoscopy has been shown to prompt expanded demonstrative exactness, appropriate understanding of dermoscopic images is typically tedious, complex, and costly and depends on the observer's perception. The traditional approaches are carried out with

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low-quality image processing methods that fall under image analysis in the medical feld. The initial methods carried out in cleaning the raw data are done by pre-processing methods that adjust the intensity, contract, redundant data, scaling, and binarisation $[1, 2]$ $[1, 2]$ $[1, 2]$ $[1, 2]$ in the morphological structure of the image.

Recent advances in information technology have resulted in high-performance results in detecting and diagnosing various types of cancer, particularly skin cancer. The unwanted noises are removed with pre-processing methods proposed by $[3]$ $[3]$. The quality of the image is enhanced by pre-processing methods, which reduce the processing time and complexity.

Image resizing plays an important role while processing the data. As diferent datasets have varied sizes of images, it is necessary to resize them to the same size to apply the algorithms. The pre-processing is followed by image segmentation to detect the region of interest (ROI). The ROI may be anything that we need to analyze. In the medical image, ROI could be the diseased region that is separated by the unafected region [\[4\]](#page-12-3). This segments the unafected tissue, and then the required features are considered from the afected region so that the result will be accurate and clear $[5]$ $[5]$.

The classical way of segmenting is performed by thresholding, clustering, and other edge detection methods, and the ABCDE method is used to diagnose melanoma in the

skin during the analysis [[6\]](#page-13-1). Even though many methods are incorporated in detecting and diagnosing cancer, there are still many challenges in dealing with real-time image data. It contained many unexpectedly complex data points, and the image's ocular emergence in the afected skin area made separating the desired ROI of melanoma with an accurate contour measure from the image difficult. [\[7](#page-13-2)].

Automated state-of-the-art techniques have been developed to help dermatologists improve their efectiveness and objectivity in the visual understanding of dermoscopic images[[8](#page-13-3)]. Automatic image segmentation is performed mainly using classifcation and clustering. Classifcation is supervised segmentation that requires prior information about the classes to be classifed. Clustering is preferred because it is unsupervised and does not require prior information. However, clustering algorithms require initial centroids to obtain clusters. Wrongly chosen clusters can result in local minima producing invalid segmentation regions, deeming the effort unfruitful.

Many researchers proposed automated methods to overcome these challenges. The broad category of these methods includes thresholding, clustering, classifcation, and contours and snakes, each having its merits and demerits [\[9](#page-13-4)].

Skin lesion segmentation based on thresholding namely, Otsu's[[10](#page-13-5)], fuzzy logic [\[11\]](#page-13-6), adaptive thresholding [[12](#page-13-7)] are available in literature. The threshold methods lead to irregular edges with varied thresholds. The lesions of a melanoma tend to have fuzzy borders and irregular shapes, leading to uncertainty in identifying the exact lesion from skin images. Researchers have used fuzzy sets and rough sets to address the uncertainty and vagueness in the image. Skin cancer with fuzzy logic approaches are proposed in [\[13](#page-13-8)[–16\]](#page-13-9). Fuzzybased methods have the advantage of representing the intermediate data in intervals, allowing qualitative analysis of the data [[3\]](#page-12-2). A fuzzy intensity threshold with type2 fuzzy logic proposed in [[11\]](#page-13-6) was able to detect the pixel intensity either belongs to ROI or the background. These fuzzy-based methods failed to group lesion pixels with low contrast as lesions.

Garcia et al. used fuzzy classifcation of pixels and histogram threshold with innovative methods applied on skin images for segmentation in [[17](#page-13-10)]. The method used Otsu's threshold with fuzzifed regions determination and applied morphological operators to extract the tissues in case of artefacts. Castillejos et al. [\[14](#page-13-11)] presented predefned cluster selection using fuzzy c-means. This work specifes automatically, the number of clusters needed by an image for segmentation. The clustering algorithms are highly infuenced by the initial centroids and the number of clusters needed for clustering.

Many algorithms based on active contours and snakes are proposed for skin lesion segmentation [[18](#page-13-12)–[23\]](#page-13-13). These algorithms usually depend on the initial curves and positioning of curves that undergo deformation to detect the boundaries of lesions. An automated method for adapting the contour to skin lesions is proposed in [[23\]](#page-13-13), but still, the contour selection is needed.

In recent days, many deep learning models for segmenting skin lesions are proposed in the literature based on convolutional neural networks, generative adversarial models [[24,](#page-13-14) [25\]](#page-13-15), deep autoencoders, stacked autoencoders, convolutional autoencoders, restricted Boltzmann's machine, and recurrent neural nets are reviewed and analyzed in [[26](#page-13-16)[–31\]](#page-13-17). Deep learning methods have the ability to obtain the features from images that are useful in segmentation. The performance of the deep learning models has improved when compared to existing methods. However, Graphics Processing Units (GPU) are used in parallel to analyze the features and to detect the skin lesions[\[8,](#page-13-3) [32,](#page-13-18) [33](#page-13-19)].

Adrian et al. use the trapezoidal intuitionistic fuzzy method for transferring the data into interval-valued trapezoidal multi-criteria for decision-making $[34]$ $[34]$. To efficiently handle the data, intuitionistic fuzzy sets(IFS) provide a triple vector to make appropriate decision making. Aribarg has introduced a modifed fuzzy ant-miner (MFAM) that employs attributes and case weighting for training [\[35](#page-13-21)]. IFS was successfully applied for brain image segmentation for classifying tissues and tumors in presence of noise and artefacts [[36,](#page-13-22) [37](#page-13-23)]. This motivated to use IFS for segmenting skin lesions in presence of artefacts.

Intuitionistic fuzzy set (IFS) theory for medical images has been proposed to handle images with a lot of uncertainties, as they are badly illumined with fuzzy. Direct segmentation of results often creates unusable results. In [\[38](#page-13-24)] Charia et al. used IFS theory to produce accurate diagnosing of diseases using cellular images. FCM clustering method has been proposed by Huang et al. [[39\]](#page-13-25) for image segmentation with rough set theory. The segmentation results of various clustering numbers using FCM where the image is distributed into various small regions based on the indistinguishable attributes and their relationship, have reduced error rates and improved segmentation of fuzzy boundary regions.

Advanced fuzzy set-theoretic techniques are very important and play a major role in the area of medical images. Image segmentation is performed on digital pathology images using the IFS method. This method results in strong results compared to other segmentation methods [\[40,](#page-13-26) [41](#page-13-27)]. Shaw et al. show that triangular intuitionistic fuzzy numbers with arithmetic procedures can be used to check the reliability of the fuzzy systems. To calculate imprecise failure, each component failure is taken into considerations by the triangular intuitionistic fuzzy numbers [\[42](#page-13-28), [43](#page-13-29)].

Tilson et al. use FCM clustering programs for data analysis problems, and this method generate fuzzy classifcation for any set of given numerical data. The clustering method used to group the subsets and generalized objective with least square method [[44\]](#page-13-30) is proposed.

Intuitionistic fuzzy c-means(IFCM) algorithm has been proposed for image segmentation in [[45\]](#page-14-0) for Brian images. Verma et al. use IFCM to handle the uncertainty. However, the results show that the method is very sensitive to noise and does not incorporate any local spatial information. From this, IIFCM (improved intuitionistic fuzzy c-means) has been proposed to handle the insensitive to noise and freedom of choosing the parameter when tuning is performed. IIFCM method results in signifcant performance increase compared to other existing methods [[37](#page-13-23), [46](#page-14-1)].

The clustering algorithms are based on IFS also suffer from initial centroid selection. In this paper, a novel algorithm is presented to obtain the melanoma regions from skin images. The proposed method identifes the initial regions using triangular fuzzy sets. The obtained regions are considered for diferent membership functions for IFS and the centroids are also initialized with mean values of the regions. The updating of the centroids and the membership functions of IFS are iterated until clusters are stable and exactly identify the melanoma regions.

The major contribution of the paper are:

- The paper has studied and analyzed triangular Intuitionistic Fuzzy Sets and its application to Skin cancer segmentation.
- Every fuzzy based clustering algorithm depends on initial centroid. In this paper, using the peaks and valleys of the histogram of the input image, the centroids are computed. The successive distance between valleys and peaks are considered for the parameters to defne TMF.
- Applying TMF, the initial regions are computed that in turn are used to compute the initial centroids.
- TMF helps in identifying the initial clusters and regions and reduce the number of iterations needed for segmentation.
- TMF converges faster due to the initial centroids selection and the accuracy of the segmentation is increased.

The paper has diferent sections that explains the background in "[Background"](#page-2-0), Triangular Intuitionistic Fuzzy C-Means (TIFCM) in ["Proposed Methodology"](#page-4-0), implementation and experimental results analysis in ["Implementation](#page-8-0) [and Experimental Results"](#page-8-0) and fnally conclusions and the future scope in "[Conclusion"](#page-12-4).

Background

Challenges of Skin Cancer Detection

Skin cancer detection is a challenging task due to numerous artefacts associated with the skin medical images.

Irregular Shape, Size and Shade

Large changes in skin lesions have made accurate detection and segmentation of skin lesions difficult. Numerous factors, like lesion localization, its size, shape, skin hair, bubbles, blood vessels, uneven boundaries with fuzziness, and tone of the skin, need to be handled with preprocessing for accurate segmentation.

Image Acquisition Artefacts

The artefacts related to image acquisition include rule markers, imaging frames, low contrast, and illumination. Figure [1](#page-3-0) depicts the artefacts that make skin cancer detection more complex.

Pre‑processing and Post‑processing

Preprocessing is performed on the data sets in diferent ways. The images are frst converted to grayscale, and then resized because diferent data sets contain images of varying sizes.There is no additional pre-processing in this case because intuitionistic sets are used to obtain lesion belonging and non-belonging as well as fuzzy boundaries based on the membership function. Binarization is performed on the lesion-belonging region as part of the post-processing binarization to obtain the lesion region.

Fuzzy Sets and Triangular Fuzzy Membership Function

Definition 1 The definition of fuzzy set is notated by a membership function $\mu_{\bar{a}}(e)$ where *E* is the universe of discourse. This function establishes a mapping between each element $e \in E$ to a real valued number in the range [0, 1]. This membership function reveals the grade of membership of the element *e* in $\mu_{\bar{a}}(e)$. As the value of the function gets closer to unity, the higher will be the membership grade.

Definition 2 The function $\mu_{\overline{\partial}}(e)$ for a triangular fuzzy number $\tilde{a} = (a1, a2, a3)$ is established as:

Fig. 1 Images from HAM10000 data set **a** skin tone, **b** irregular shape, **c** hair, **d** air bubble images, **e** fuzzy boundaries **f** cracked skin tones, **g** rule markers, **h** imaging frames

$$
\mu_{\bar{a}}(e) = \begin{cases}\n0 & \text{if } e < a1 \\
\frac{e-a1}{a^2 - a1} & \text{if } a1 < e \leq a2 \\
\frac{a3 - e}{a^3 - a2} & \text{if } a2 < e < a3 \\
0 & \text{Otherwise}\n\end{cases} \tag{1}
$$

The values of the triplet (*a*1, *a*2, *a*3) are real numbers that confines to $a1 \le a2 \le a3$. The value of *e* at the triplet element *a*2 is the most likely value of the evaluation data and it is also observed as the supreme grade of $\mu_{\bar{a}}(e)$. On the other hand, the minimum grade of *e* at $\mu_{\bar{a}}(e)$ is the least likely value to be evaluated from the membership function i.e., $\mu_{\bar{a}}(e) = 0$. The range of evaluation data is bounded between the constants [*a*1, *a*3], which is an implication of degree of fuzziness of the evaluation data.

Intuitionistic Fuzzy Sets

Generalization of fuzzy sets is characterised by membership, non-membership and hesitancy values [\[47](#page-14-2)]. The degree of belongingness to a cluster determines the membership and non-membership property of the elements. The dilemma of a particular pixel belonging to a specifc cluster is determined by the hesitancy. Hesitancy adds precision to the imperfect knowledge obtained from the fuzzy sets.

$$
\begin{array}{c} \text{SN Computer Science} \\ \text{A SPRINGER NATURE\,Journal} \end{array}
$$

$$
A = \{ (e, \mu_A(e), \pi_A(e)) \mid e \in E \}
$$
 (2)

with $\gamma_A(e) = 1 - (\mu_A(e) + \pi_A(e))$ where the functions $\mu_A(e), \pi_A(e)$ indicate the degree to which an element belongs or does not belong to the fnite set *E*. The intuitionistic fuzzy index is determined by $\gamma_A(e)$ signifies the degree of hesitation of an element. The necessary conditions to be satisfied by an element $e \in E$ that is defined as IFS is given by $0 \leq \mu_A(e), \gamma_A(e), \pi_A(e) \leq 1$. The bias, noise and assignment of an element to a particular cluster is precisely represented by the three functions.

Data Sets

Diferent data sets available to work with skin cancer algorithms are explained below and are presented in Table [1](#page-3-1).

PH2 [[48\]](#page-14-3) consists of epiluminoscopy images, also known as dermoscopic images having the size of 765X574 pixel values. There are 200 images in this dataset of melanocytic

Table 1 Data sets used for simulation

Dimension	Dataset	Number of images	Image size
2D	PH ₂	200	765×574
2D	ISIC 2019	25331	varying \times varying
2D	HAM10000	10015	600×450

lesions, including nevi, sensitive nevi, and melanomas.This dataset also contains numerous image descriptors.

ISIC 2019 dataset is categorized into 8 diferent classes of dermoscopic images for training and testing. The datasets are available with and without metadata and contain a total of 25,331 images. Groundtruths are also available for testing on standard images of lesions.This dataset is derived from the BCN20000, HAM10000, and MSK datasets [[46,](#page-14-1) [49\]](#page-14-4).

HAM10000 [[50](#page-14-5)] as the name indicates, it contains 10,015 images for training that were classifed in to diferent dermatoscopic images in the realm of pigmented lesions collected from varied populations. They were collected, confrmed and stored through diferent modalities rate [[50\]](#page-14-5). They are available for the research and public use in the ISIC collection.

Proposed Methodology

This section mainly concentrates on the flow of segmentation performed on skin images using triangular intuitionistic fuzzy sets.

Triangular Membership Based Fuzzy Region Determination

The fuzzy functions highly impact the segmentation performance as these are necessary for producing clusters. The proposed method presents a novel approach by identifying the initial regions with the triangular membership functions, which act as the input to the intuitionistic fuzzy

 (a)

Fig. 2 a, **c** ISIC sample images, **b**, **d** valleys and peaks of the images

C-means. The observed image having the size $m \times n$ is denoted as $E = \{e_i/e_i\}$ represents *i*th pixel value in the image }. To apply triangular membership function *a*, *b* and *c* values are needed. These values are calculated from the gray values associated with the peaks and valleys of the histogram of the image [[51](#page-14-6)]. Figure [2](#page-4-1) show the ISIC images and their corresponding peaks and valleys. Every sequence of valley, peak and valley will be treated as *a*1, *a*2 and *a*3 that will extract the membership $\mu_{\bar{a}}(e)$ for the regions using Eq. [1.](#page-3-2)

The regions obtained from the triangular membership function are given as input to the intuitionistic fuzzy c-means. Figure [3](#page-5-0) shows the regions extracted using triangular membership function.

Triangular Intuitionistic Fuzzy c‑Means (TIFCM)

The regions obtained from the triangular membership function are used to fnd the centroids needed to perform intuitionistic fuzzy c-means. Many image segmentation works based on IFS are proposed in [[37,](#page-13-23) [52](#page-14-7), [53\]](#page-14-8). The image *E* is correlated to intuitionistic fuzzy with the below equation

$$
A = \{ (e_i, \mu(e_i), \pi(e_i)) \mid e_i \in E \}
$$
\n(3)

To find $\mu(e_i)$, $\pi(e_i)$ and $\gamma(e_i)$, membership functions, the three regions obtained with *a*, *b*, *c* of triangular membership function is considered as deterministic region $D(e_i)$, $H(e_i)$ and other region as indeterminacy region $I(e_i)$. Compute the mean value of every region to obtain the initial centroids *Cj* where $1 < j < n$ (*n* is the 3 for three regions).

Fig. 3 a, e, i Images from ISIC data set, b, f, j regions extracted for values $a1 < e_i \le a2$, c, g, k regions extracted for values $a2 < e_i \le a3$, d, h, **l** other region for values $e_i \leq a \cdot 1$ and $e_i \geq a \cdot 3$

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The computation of membership value $\mu(e_i)$ is performed with the deterministic region $D(e_i)$ and the centroids with the below Eq. [4.](#page-6-0)

$$
\mu(e_i) = \frac{1}{\sum_{j=1}^n \left(\frac{||D(e_i) - C_j||}{\sum_{c=1}^n ||D(e_i) - C_c||} \right)^{\frac{2}{m-1}}}
$$
(4)

where C_j is the centroid of *j*th cluster (here, mean value of $(D(e_i))$ and C_c is the centroid values of regions other than *j*th cluster (here, $H(e_i)$, $I(e_i)$) are obtained from triangular membership function). Compute $\pi(e_i)$, and $\gamma(e_i)$ with the help of $I(e_i)$ and $H(e_i)$ by making use of Eqs.[\(5](#page-6-1))-([6\)](#page-6-2).

$$
\pi(e_i) = \frac{1}{\sum_{j=1}^n \left(\frac{||I(e_i) - C_j||}{\sum_{c=1}^n ||I(e_i) - C_c||} \right)^{\frac{2}{m-1}}}
$$
(5)

$$
\gamma(e_i) = \frac{1}{\sum_{j=1}^n \left(\frac{||H(e_i) - C_j||}{\sum_{c=1}^{n_c} ||H(e_i) - C_c||} \right)^{\frac{2}{m-1}}}
$$
(6)

The IFS partition matrix $\mu_{IFS}(e_i)$ and cluster centroid $C(\mu(e_i))$ are updated using the below equations.

$$
\mu_{IFS}(e_i) = \mu(e_i) + \pi(e_i) + \gamma(e_i)
$$
\n⁽⁷⁾

$$
C(\mu(e_i)) = \frac{\sum_{e_i \in E} (\mu(e_i))^m(e_i)}{\sum_{e_i \in E} (\mu(e_i))^m}
$$
(8)

Find the other centroids using the Eq. [8](#page-6-3) for $\pi(e_i)$ and $\gamma(e_i)$. Here, *m* indicates the fuzzification constant given by user. The membership function converts to be crisp and binary if the values *m* nears to 1 otherwise fuzzy and blurred with increased value [\[54\]](#page-14-9). Good segmentation results are produced for vast data with $1.5 < m < 3$ and generally 2 is considered[[55,](#page-14-10) [56\]](#page-14-11).

Triangular intuitionistic fuzzy c-means is also iterative similar to other clustering algorithms. It fnds the three regions using Eqs. [4,](#page-6-0) [5](#page-6-1), [6](#page-6-2) and obtains the new centroids using Eq. [8](#page-6-3) repeatedly and stops when the clusters are stable. The stability of cluster is computed with the similarity coefficient based on the Hamming distance between the previous clusters $\mu(e_i)$, $\pi(e_i)^l$ and $\gamma(e_i)^l$ and present cluster $\mu(e_i)^{l+1}, \pi(e_i)^{l+1}$ and $\gamma(e_i)^{l+1}$. The hamming distance is given with the equation

$$
sc = \sum_{i=1}^{M*N} | \mu_i e_i |^l - \mu_i e_i |^{l+1} |
$$

+
$$
| \pi(e_i)^l - \pi(e_i)^{l+1} |
$$

+
$$
| \gamma(e_i)^l - \gamma(e_i)^{l+1} |
$$
 (9)

Triangular Intuitionistic Fuzzy c‑Means Clustering Algorithm (TIFCM)

The step-wise algorithm of Triangular intuitionistic fuzzy is listed below.

Algorithm 1 Segmentation algorithm for images with generalized triangular intuitionistic fuzzy sets

INPUT: TIFCM image E and ϵ exit criteria with 0.01.

OUTPUT: *n* number of clusters to be generated for representing the regions of skin image contain a, b, c values for each regions using method defined by [51]

- 1 Obtain a,b,c values for each regions using method defined by [51].
- 2 Obtain the three regions $D(e_i)$, $H(e_i)$ and $I(e_i)$ as per Eq. 1.
- 3 Find the cluster centroids C_j from the regions by finding the mean value of the region.
- 4 while true
- 4.1 for every *j* in *n* clusters and every *eⁱ* in *E* obtain three intuitionistic membership function values using Eq. 4-6
- 4.2 Update the partition matrix $\mu_{IFS}(e_i)$ and the cluster centroid C_j using Eq. 7 and Eq. 8.

4.3 Estimate the similarity coefficient(*sc*) using Eq.9

4.4 If $sc \leq \epsilon$ then break else compute $\mu_{IFS}(e_i)^l = \mu_{IFS}(e_i)^{l+1}$ and repeat 4.

5 with stable $\mu(e_i)$, $\gamma(e_i)$ and $\pi(e_i)$ obtain *n* clusters.

Fig. 4 Segmentation results of proposed algorithm TIFCM for diferent images of ISIC data set. (Column 1) Original images of ISIC data sets, (Column 2) Deterministic region depict ing cancer region, (Column 3) Indeterminacy regions (Column 4) Hesitance regions

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Implementation and Experimental Results

Experimental Setup

To find the efficacy of the proposed algorithm TIFCMM, The ISIC 2016, HAM10000 2019 data set [\[49\]](#page-14-4) and PH2 [[48\]](#page-14-3) data set is used. The data set contains the skin images with diagnostic, clinical, technical and database attributes based images. The proposed TIFCM algorithm is worked out with MATLAB software on i5 processor. The total number of clusters *n* is assigned as 3 in the proposed algorithm to determine the three regions. The centroid values are initialised through the triangular membership functions. The diference between the subsequent clusters is fxed as the exit criteria, which is 0.01.

As part of pre-processing the images are resized to 512×640 as the image sizes vary for ISIC, PH2, HAM10000 data sets. In post processing of the proposed algorithm, the deterministic fnal region is converted to black and white to compare with the ground truths.

The segmentation results of the TIFCM algorithm on various skin images is shown in Fig. [4.](#page-7-0) Column 1shows the Original images of ISIC data sets. The images (a) ISIC_0000000,(e) ISIC_0000008 (i)ISIC_0000049 (m) ISIC_0000008 (q) ISIC_0000002 (u) ISIC_0000043 are shown in Column 1 and are used to extract malignant region using proposed TIFCM. Column 2 shows the Deterministic region depicting cancer region, Column 3 depicting Indeterminacy regions and Column 4 showing the Hesitance regions extracted from the stable clusters of TIFCM. Further, in Fig. [5](#page-9-0) the segmentation results of images ISIC0034329, ISIC0034332, ISCI0034337 and ISIC0034343 of ISIC 2019 data set are presented. The subjective analysis show that the lesion was extracted efficiently using the proposed method.

Figure [6](#page-10-0) exhibits the results of TIFCM algorithm for different images like malignant melanoma, melanoma, benign melanoma, common nevus, a typical nevus image that are commonly identifed skin diseases of PH2 data set. The lesion detection of these images are compared with their ground truths [\[7\]](#page-13-2). The results showed that the proposed algorithm performs very well when compared to other algorithms. The comparative analysis of the proposed algorithms with other state of art techniques namely KM [[57\]](#page-14-12), FCM [\[55\]](#page-14-10), RFCM [\[58\]](#page-14-13), and GRIFCM [\[37\]](#page-13-23) is given in Fig [7.](#page-11-0) The detailed analysis with other segmentation algorithms reveals that the proposed algorithm gives improved cancer region segmentation.

Quantitative Analysis

The segmentation results are analysed quantitatively using the performance measures namely, segmentation accuracy (SA), Jaccard's coefficient (JC) and Dice coefficient (DC).

Jaccard coefficient =
$$
\frac{E1 \bigcap E2}{E1 \bigcup E2}
$$
 (10)

$$
\text{Dice coefficient} = \frac{2 \mid E1 \bigcap E2 \mid}{\mid E1 \mid + \mid E2 \mid}
$$
\n⁽¹¹⁾

where $E1$ is the ground truth image. $E2$ is the resultant segmented image,

Segmentation Accuracy =
$$
\frac{TP + FN}{TP + FN + TN + FN}
$$
 (12)

where,

- 1. True positive (TP): Ratio of sum of true pixels in the E2 detected in E1.
- 2. True negative (TN): Ratio of sum of true pixels in the E2 falsely segmented as negative in E1.
- 3. False positive (FP): Ratio of sum of false pixels in E2 falsely segmented as positive in E1.
- 4. False negative (FN): Ratio of sum of false pixels in E2 detected as negative in E1.

The SA, JC, DC, precision, specificity and sensitivity measures are given in Table [2](#page-12-5). These performance measures are calculated for diferent skin disease, namely, malignant melanoma, melanoma, benign melanoma, common nevus, typical nevus images of ISIC data set images. The results proved that the proposed algorithms have efficiently extracted the lesion of an cancerous tissues. The average JC value of 0.85, SA of 0.90 is achieved for with the proposed TIFCM. The quality of segmentation can be assessed by observing the segmentation JC value with 0.8 or above indicating the visual "correctness". It is observed that for 100 images from ISIC 2019 the observant agreement for JC value is considered as 0.786[[49\]](#page-14-4). It is observed that, the fall in JC value below 0.7, the "correctness" of the segmentation can be debated. With the proposed method,78 out of 100 images fell above JC of 0.7, 10 images fell at or below 0.7, rest images fell below 0.6.

Fig. 5 Segmentation results of proposed algorithm TIFCM for ISIC_0034329, ISIC_0034332, ISCI_0034337 and ISIC_0034343 of ISIC 2019 data set. (Column 1) Original images of ISIC data sets,

(Column 2) Deterministic region depicting cancer region, (Column 3) Indeterminacy regions (Column 4) Hesitance regions

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Fig. 6 Lesion segmentation of diferent PH2 data set images. (Row 1) **a** Malignant melano mas image IMD063, **b** Ground Truth(GT), **c** extracted using TIFCM. (Row 2) **d** Mela noma image IMD058, **e** GT of IMD058, **f** extracted using TIFCM. (Row 3) **g** A typical Nevus image IMD048, **h** GT of IMD048, (i)extracted using TIFCM. (Row 4) **j** Common Nevus image ISIC_0012253, **k** GT of (**j**), **l** extracted using TIFCM.(row 5) **m** Benign mela noma image IMD075, **n** GT of IMD075, **o** extracted using TIFCM. (row 6) **p** Benign mela noma image ISIC_0000125, **q** GT of **p**, **r** extracted using TIFCM

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Fig. 7 Comparison of Algorithms applied to ISIC_0000006 for Benign nevus detection (Row 1) Segmentation with KM, (Row 2) Segmentation with FCM, (Row 3) Segmentation with RFCM, (Row 4) Segmentation with GRIFCM, (Row 5) Segmentation with TIFCM

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Table 2 Comparisons of performance measures for the images in Fig. [3](#page-5-0)

Images	SA	JC	DC	Precision	Sensitivity	Specificity
Malignant melanoma	0.9059	0.8799	0.8438	0.8797	0.9206	0.8935
Melanoma	0.9308	0.9341	0.7939	0.9396	0.9530	0.8915
A typical nevus	0.9185	0.7424	0.8071	0.7468	0.7868	0.9585
Common nevus	0.8719	0.8323	0.8123	0.8623	0.9880	0.8639
Benign melanoma	0.8689	0.7921	0.8528	0.8121	0.9853	0.8520

Table 3 Comparisons of performance measures for diferent segmentation algorithms

Algorithm	SА	JC	Sensitivity	Specificity	DC
Yolo et al. $[30]$	0.929	0.795	0.836	0.940	0.881
Yuan et al. $[8]$	0.930	0.765	0.825	0.975	0.849
Li et al. $[32]$	0.932	0.7620	0.820	0.978	0.847
Lin et al. $[33]$		0.620			0.770
FCM [59]	0.884	0.665	0.869	0.923	0.884
TIFCM	0.905	0.856	0.952	0.862	0.830

Table 4 Execution time comparison for diferent algorithms

To depict the efficiency and robustness of the proposed TIFCM method, a comparison with the latest methods in the literature, are shown in Table [3](#page-12-6). The methods used in the comparison are executed on ISIC 2017 challenge and are ranked based on the Jaccard's coefficient. It is clear from the table that the JC values produced with the proposed method has an marginal increment over other lesion segmentation algorithms.Also, the improved sensitivity values are observed with the proposed method. In contrast, the deep learning methods have exhibited high segmentation accuracy and specifcity. The proposed method dice coefficients are inline with the results of the deep learning methods (Table [3](#page-12-6)). In Table [4](#page-12-7) the execution times of proposed method in comparision to the existing algorithms are presented. The proposed TIFCM method shows improved performace as the initial centroids are chosen with the histrogram peaks and valleys.

The efficacy of the proposed method is also compared with the method FCM classification of pixels with histogram thresholds [[59\]](#page-14-14). The SA, JC and sensitivity values are signifcantly improved with the proposed methods while the specificity and DC values are less with the proposed method.

Conclusion

Computer-aided detection of malignant melanoma skin cancer with the use of skin images has tremendously assisted clinicians. Skin cancer detection has become more difficult due to a variety of factors such as artefacts in skin images, non-homogeneous intensities, and low contrast images.In this paper, we present a novel TIFCM algorithm for skin image segmentation. The triangular membership function is used to extract the initial regions and centroids. This avoids the initial selection of centroids, which is the drawback of every clustering algorithm. The triangular membership based centroids and regions assist in the faster stability of clusters, reducing the execution time. The regions extracted are used as initial inputs to calculate the membership functions of the intuitionistic fuzzy c-means that efectively handle the uncertainty in the skin images.

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Data availability The data will be made available on request from the author.

Declarations

Conflict of interest The authors declare that there is no confict of interest.

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