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Deep Learning in Skin Disease Image Recognition: A Review

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ABSTRACT The application of deep learning methods to diagnose diseases has become a new research topic in the medical field. In the field of medicine, skin disease is one of the most common diseases, and its visual representation is more prominent compared with the other types of diseases. Accordingly, the use of deep learning methods for skin disease image recognition is of great significance and has attracted the attention of researchers. In this study, we review 45 research efforts on the identification of skin disease by using deep learning technology since 2016. We analyze these studies from the aspects of disease type, data set, data processing technology, data augmentation technology, model for skin disease image recognition, deep learning framework, evaluation indicators, and model performance. Moreover, we summarize the traditional and machine learning-based skin disease diagnosis and treatment methods. We also analyze the current progress in this field and predict four directions that may become the research topic in the future. Our results show that the skin disease image recognition method based on deep learning is better than those of dermatologists and other computer-aided treatment methods in skin disease diagnosis, especially the multi deep learning model fusion method has the best recognition effect.

INDEX TERMS Deep learning; Image recognition; Review; Skin disease;

I. INTRODUCTION

The skin, the largest organ of the human body, is an important barrier. The main function of the skin is to protect the human body from harmful substances from the outside world and prevent the outflow of various nutrients in the human body [1]. In human productive life, the skin health status is affected by many factors, such as solar radiation, smoking, drinking, sports activities, viruses, and working environment [2]. These factors not only affect the integrity of skin function but also cause certain damage to the skin, have an adverse effect on human health, and can even threaten human life in severe cases. Therefore, skin disease has become one of the common diseases of human beings. Skin disease covers all cultural regions and occurs in all ages. Approximately 30% to 70% of people are in

high-risk groups [3]. According to the British Skin Foundation Report in 2018, approximately 60% of the British people suffer from skin disease [4]; 5.4 million new cases of skin cancer are recorded in the United States every year; one in five Americans will be diagnosed with a cutaneous malignancy in their lifetime [43]. Skin disease brings not only a significant impact to human beings, such as daily activities damage, loss of the interpersonal relationship, and internal organ damage, but also death. This condition can also constitute mental illness, leading to isolation, depression, and even suicide [5]. Accordingly, skin disease has become one of the major topics in the field of medicine. Figure 1 shows the global hot situation of skin disease.

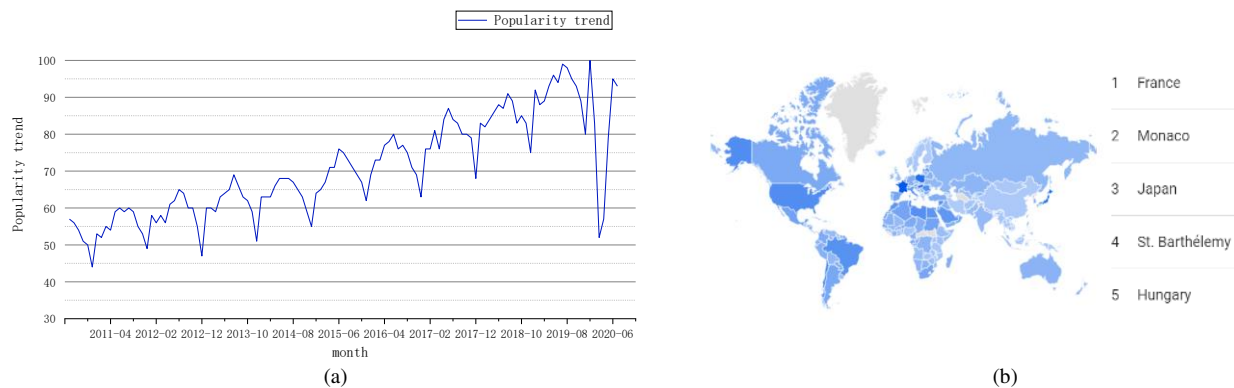


FIGURE 1. Global Skin Disease Thermal Map. Panel (a) is a global trend chart of skin disease search fever, and the horizontal axis represents the year's longitudinal axis. The chart shows that the global search fever for skin disease has started to increase since 2011, and the search fever has been increasing year by year. By 2020, the search fever for skin disease has reached its peak. Panel b is a global heat map of skin disease regions, starting from 2011. The map demonstrates skin disease problems in all parts of the world, except Greenland in the Arctic Circle, and the search for skin disease is hot near the equator. The highest search for skin disease in the world is France, followed by Monaco and Japan.

In the treatment of skin disease, early detection is the critical condition to cure the disease, effectively reduce its impact, and improve the survival rate. Take melanoma in skin disease as an example. In recent years, malignant neoplasms in skin diseases have increased significantly. Malignant melanoma (the deadliest type) is responsible for 10,000 deaths annually just in the United States [6]. Melanoma is a highly lethal but not incurable disease. If abnormal proliferation of skin melanocytes is detected in the early stage, then the survival rate is 96%; if it is detected in the late stage, then the survival rate is only reduced to 5% [7]. Therefore, early diagnosis and treatment of skin disease can minimize the damage caused by skin disease. However, the skin disease recognition accuracy is unideal due to the similarity between different skin diseases and the limited number of dermatologists with professional knowledge. The identification of skin disease has become a serious scientific challenge.

To address the issue of skin disease diagnosis and treatment, people used computer-aided diagnosis for automatic skin disease recognition based on the skin disease images earlier[26]. With the rapid development of the artificial intelligence technology, deep learning has quickly developed a computer vision. The medical image processing of skin disease has become an essential component and received great attention in the cross-field of image processing, machine science, and intelligent medicine. Many experts and scholars have been engaged in the image recognition of skin disease. The recent article published by Dick et al. is a good starting point. This article lists in detail the relevant articles on the diagnosis of melanoma in deep learning [8].

This study investigates the research status of skin disease recognition in recent years, summarizes the datasets used by researchers, and analyses from the aspects of image preprocessing, data augmentation, deep learning model, and framework performance indicators. On the one hand, this study provides a reference for deep learning methods for

dermatologists. On the other hand, this study facilitates researchers to quickly and accurately retrieve the literature related to dermatological image recognition. This survey's foundation is the rapidly developing artificial intelligence-based diagnosis technology in the medical field, which has become increasing popular among researchers. The application of artificial intelligence in other fields has shown its great potential. The fact that at least 45 studies have used deep learning to address skin disease identification issues and have achieved promising results encourages authors to prepare the survey.

This study mainly summarizes the research and application progress of skin disease image recognition based on deep learning. Section II briefly introduces the methods and status of skin disease recognition. Section III introduces the development history of skin disease detection. Section IV focuses on the research progress of skin disease image recognition based on deep learning. Section V summarizes the full text and discusses skin disease image recognition's future research trends based on deep learning.

II. METHOD

In recent years, deep learning has been given great attention to skin disease recognition, and research achievements increased. This study summarizes the relevant literature in the field of skin disease identification from 2016 to 2020. The distribution of the selected papers is shown in Figure 3. The three main steps in analyzing the literature in this field are as follows: (a) Use hierarchical search strategies to retrieve and collect relevant literature on each database, (b) conduct detailed review and analysis of collected literature, and (c) statistical analysis of relevant data.

The first step is to search for conference papers or journal articles from *CNKI* and the scientific databases *IEEE Xplore*, *ScienceDirect*, *Web of Science*, *Google Scholar*, *PubMed*, *arXiv*, and *Medline* by using hierarchical search strategies. The strategy is: First, the year limit is set, and the search year

is from 2016 to the present; then, the first-level keywords are used for the subject search for each database. The first-level keywords are as follows:

["Artificial intelligence" and "skin disease recognition"] or ["deep learning" and "skin disease recognition"] or ["deep learning" and "skin lesion"].

Approximately 2431 documents were retrieved through the first-level keyword search; then, the second-level keywords were used to retrieve the documents obtained at the upper level. The second-level search function was to obtain additional detailed granularity (such as melanoma, acne, and pigment lesions) to search for achieving the first level of missing and filling. In the field of deep learning image recognition, the mainstream method is to use convolutional neural networks (CNNs). Accordingly, the second level uses CNNs as keywords to narrow the search range. The second level keywords are as follows:

["Convolutional Neural Network" and "Melanoma Recognition"] or ["Convolutional Neural Network" and "Acne Classification"] or ["Convolutional Neural Network" and "Pigmented Skin Disease Classification"].

In this way, 312 papers related to deep learning are screened out and are suitable for the field of skin disease recognition. These papers are then sorted by "relevance" and "time". We quickly browse the downloaded and cited documents. The "Abstract, Introduction and Conclusions" "Exclude non-deep learning and non-dermatological identification documents". We also check the relevant literature cited in the paper and whether any literature meets skin disease recognition; and, if studies exist, we search them. During the browsing, if an expert or scholar is found to have unique insights in the field, then we will directly search the author's name for a literature search. Finally, 45 papers were retrieved through a hierarchical search strategy. The search strategy process is shown in Figure 2.

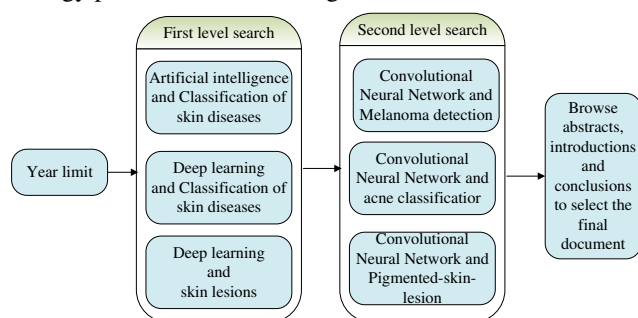


FIGURE 2. Search strategy flowchart. The search is mainly divided into two levels. First, the year is restricted. Second, the first-level search and second-level search are conducted.

In the second step, the 45 papers selected in the previous step are analyzed one by one. The following research questions are considered:

Which categories of skin disease have been identified in this study? What are the data sources and types used? Are the datasets public, and is it a dermoscopic image? Is the data preprocessed or enhanced? (4) What are the models used in deep learning, and what frameworks are chosen to build them?

(5) What are the innovations of this document and the main improvements of the model? Is model fusion used? (6) What are the performance indicators used, and how is the overall performance? Did the researchers test the performance of the models on different datasets?

The third step is to sort out the analyzed information, summarize the collected datasets, and classify the standard preprocessing techniques and deep learning neural network models in the literature. The network models are divided into single and multiple models, and the performance indicators used in the article are summarized. The main findings are discussed in detail in Section 4.

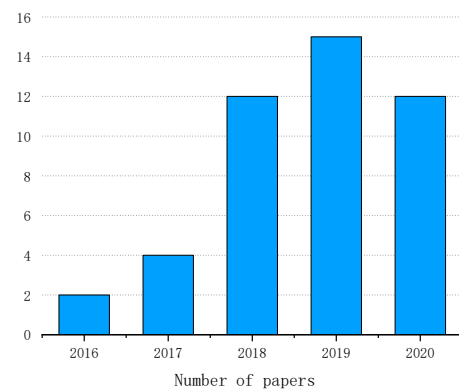


FIGURE 3. Distribution of selected papers. The horizontal axis represents the number of papers in the year along the vertical axis. The number of deep studies on skin disease recognition has rapidly increased since 2016. Twenty-one studies were published in 2018 and 2020, thereby accounting for more than 86.6%. From the content of the study, 91% (41 articles), 6% (3 articles), and less than 2% (1 article) were involved in skin disease identification, data generation, and interpretable studies of skin disease identification, respectively.

III. DEVELOPMENT OF SKIN DISEASE DIAGNOSIS TECHNOLOGY

A. TRADITIONAL MEDICAL DIAGNOSTIC PROCESS OF SKIN DISEASE

The traditional medical diagnosis of skin disease comes from the doctor according to his knowledge and experience or the characteristics and rules presented by the dermoscopic images to distinguish the status of the patient's skin lesions. The diagnostic process can be summarized as follows: first, through the doctor's visual observation, namely, visual diagnosis, to locate the necessary information of the patient, then dermoscopy and histopathological examination. Dermoscopy is a noninvasive skin imaging technology that can observe the skin structure at the junction of the lower epidermis and the superficial dermis, and it is a high-definition imaging technology [23]. Doctors analyzed the nature, distribution, arrangement, color, edge and boundary, shape and appearance of pigmented skin lesions according to dermoscopy detection methods such as seven-point checklist [12], ABCD rule [13], chaos and Clues [16], three-point checklists [14], and cash (color, architecture, symmetry, and homogeneity) [15]. However, only experienced dermatologists can accurately identify pigmented skin

disease' pathological features due to the similarity of skin lesions in color, texture, edge contour, and other features and the difference of pathological tissues between different patients. However, this method of relying on experience for diagnosis is far from meeting the patients' needs for medical resources. The process from a sampling test to a doctor's diagnosis, the histopathological examination, and then to the patient's report generally takes 4 to 5 days. This process requires a large amount of time and affects the patient's cure.

In summary, the traditional dermatological diagnosis has the following shortcomings: First, the lack of medical resources. Dermatologists with professional skin knowledge are limited. The mismatch between the dermatologists' growth rate and the incidence rate of skin disease has resulted in many patients with few professional dermatologists. Second, the accuracy of diagnosis is low. Dermatologists with professional knowledge have different work experiences and may have varying diagnoses for the same patient under subjective thoughts. Even the same doctor, affected by light, fatigue, and other factors, has different diagnostic results for the same skin disease picture. Third, the skin disease images

are complex due to the skin disease characteristics, and small gaps exist between categories and large gaps within categories. Accordingly, the diagnosis is prone to misdiagnosis or misses a diagnosis, leading reduction in the diagnostic accuracy.

B. SKIN DISEASE IMAGE RECOGNITION BASED ON MACHINE LEARNING

With the development of machine learning, the solving of the shortcomings of the traditional skin disease diagnostic process and image recognition technology of skin disease based on machine learning was established. Image recognition based on machine learning is an interdisciplinary field integrating medical skin disease imaging, mathematical modeling, and computer technology through feature engineering and machine learning classification algorithms to complete the recognition and diagnosis of skin disease. The flow chart is shown in Figure 4.

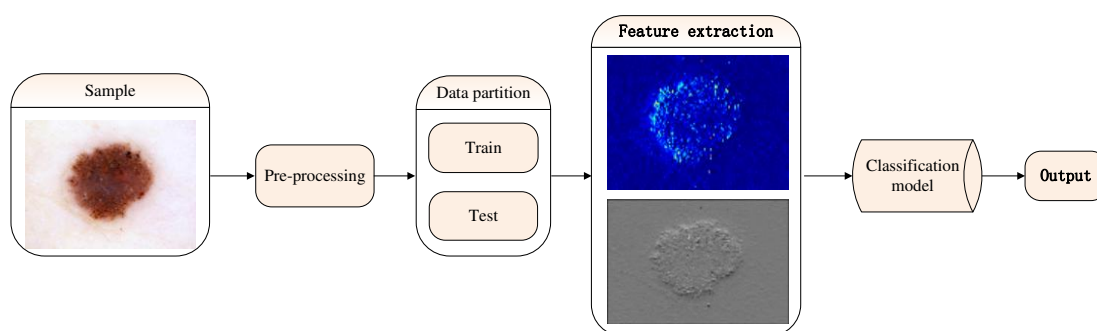


FIGURE 4. Flow chart of skin disease image recognition based on machine learning. Image processing is divided into image acquisition, image preprocessing, and dataset division. Image preprocessing includes image size adjustment, normalization, and noise removal. Image recognition mainly includes image feature extraction and classification models to classify the extracted features and then output the results.

The earliest study on the automatic classification of skin disease dates back to 1987 [9]. In 2007, Stanley et al. extracted melanomas' color characteristics, established color histograms, and classified them [10], [11]. In 2012, Rahil et al. used wavelet decomposition to derive texture features, modeled and analyzed lesion boundary sequences to derive boundary features, and based on shape indicators to derive geometric features. Finally, four classifiers, namely, Support Vector Machine, Random Forest, Logical Model Tree, and Hidden Naive Bayesian, are used for classification [17]. In 2013, Ballerini et al. proposed a hierarchical K-NN classifier algorithm for melanoma skin disease based on color and texture, which uses three classifiers for hierarchical combination and feature selection to adjust each classifier's feature set to suit its task. The recognition accuracy is more than 70% [18]. In the same year, Ning et al. used machine learning ID3 [19], classification and regression tree [20], and AdaBoost three different algorithms for feature extraction of their performance [21]. The AdaBoost performed well in

these algorithms [22]. In the 400 skin images collected with laser confocal scanning microscopy, the recognition accuracy was 94.75%, the specificity was 93%, and the sensitivity was 96.5%.

The image recognition method for skin disease based on machine learning is to extract the features of skin disease by manually setting extractors and classifying them by traditional machine learning methods. This method requires great professional medical knowledge to conduct deep exploratory data analysis and reduce its dimension. Finally, after a complex parameter adjustment, the results can be outputted, which requires a large amount of time and energy. This method has low portability, so the feature engineering is effective only in the same field. These shortcomings limit the development of machine learning in skin disease recognition. Deep learning has achieved good results in image recognition with its advancement. The deep learning method can automatically mine the deep-seated nonlinear relationship in medical images and do not need to establish feature

engineering compared with the traditional image recognition methods. The extraction efficiency is also efficient. Deep learning is adaptable and easy to transform, and the technology can be more easily adapted to different fields and applications.

IV. SKIN DISEASE IMAGE RECOGNITION BASED ON DEEP LEARNING

A. APPLIED SKIN DISEASE FIELD

The main application of deep learning in skin disease recognition is skin disease classification, that is, the quantitative feature extraction of lesion tissues through skin disease images. The classification is analyzed and judged. This direction is the mainstream application direction of skin disease recognition, and the main types of skin disease are benign neoplasms and malignant neoplasms. Benign neoplasms are a type of skin disease with a gradually increasing incidence, and the gap between the lesions is small,

and the recognition is low. Benign neoplasms commonly used for research includes nevus (23 articles) and seborrheic keratosis (17 articles).

Malignant neoplasms are another type of skin disease widely used in research. Malignant neoplasms are cellular dysplasia disease that occur in the skin, which are life-threatening through constant proliferation and metastasis. Malignant tumor identification in skin disease identification is exceptionally significant due to the high mortality rate of malignant neoplasms. The malignant neoplasms commonly used in research are basal cell carcinoma (13 articles), squamous cell carcinoma (seven articles), and malignant melanoma. Among the retrieved literature, the largest number of studies on melanoma recognition was 34. However, Non-neoplastic skin diseases are scarce, and only three articles on deep learning of eczema and psoriasis identification [46], [51], and [74] are collected in this study.

TABLE I
APPLICATION CATEGORIES OF SKIN DISEASE IDENTIFICATION

Type	Types of skin disease	Reference
Benign neoplasms	Nevus	[41],[44],[45],[47],[48],[50],[52],[53],[54],[55],[56],[57],[60],[61],[64],[65],[76],[77],[78],[79],[80],[82], and [84]
	Seborrheic keratosis	[44],[45],[47],[48],[50],[52],[53],[55],[60],[61],[62],[71],[75],[79],[80],[83], and [84]
Malignant neoplasms	Basal cell carcinoma	[38],[39],[41],[42],[54],[56],[57],[58],[62],[79],[82],[83], and [84]
	Squamous cell carcinoma	[38],[39],[41],[58],[79],[82], and [84]
	Melanoma	[38],[58],[59],[60],[61],[64],[65],[71],[72],[73],[76],[77],[78],[79],[80],[81],[82],[83],[84], and [85]
Non-neoplastic skin diseases	Eczema/Psoriasis	[46],[51], and [74]

B. DATA SOURCES

Deep learning requires a large amount of data to extract features during training. However, large-scale image data of skin disease are difficult to obtain due to certain aspects, such as the image of skin disease involves patients' privacy, variety of skin diseases, and the presence of some rare diseases. Skin disease images need to be labeled by experts with appropriate medical knowledge due to the similarity of lesion manifestations among various skin diseases, which limits the size of the skin disease dataset that is publicly available in academia. Currently, the acquisition of skin disease datasets is mainly divided into self-collected and public datasets. Self-collected datasets are currently less publicly available. Most published dermatological datasets are image data obtained by using dermoscopic imaging and collected from dermatological image databases [30], [32], [33], [34], [40], [97]. Some datasets are also collected by universities in collaboration with renowned hospitals [25], [26], [29], [37], [62]. Pathological sections of basal cell carcinoma and seborrheic keratosis studied by Meifeng et al. are obtained from the Second Affiliated Hospital of Xi'an Jiaotong University [62]. The HAM10000 dataset is a dermoscopic image collected from the Dermatology Department of Vienna Medical University in Austria and the dermatology practice of Cliff Rosendahl in Queensland,

Australia [29].

In this study, 18 datasets were collected, including 14 public datasets and 4 self-collected datasets [38], [49], [72], [75]. See Table II for details. The 18 datasets covered 17 types of skin diseases, among which melanoma has the most frequent occurrence, and 14 datasets contained relevant data. Eczema and psoriasis only appeared in DermIS, DemQuest, DermNZ, and Dermnet datasets, and all of them were nondermoscopic images. The ISIC competition dataset has become the widely used skin disease dataset due to its wide range of influence, massive data, rich types, and other characteristics. ISIC 2018 is the most widely used skin disease dataset, and 12 papers used it [42], [49], [53], [54], [56], [57], [66], [71], [76], [79], [84], [85]. Seven papers used the ISIC 2016 dataset [47], [52], [53], [55], [64], [65], [85], and 11 papers utilized the ISIC 2017 dataset [44], [45], [47], [48], [50], [52], [53], [55], [61], [68]. Researchers will use multiple datasets to conduct experiments due to the limited number of images in a single dataset. For example, researchers at Stanford University organized four skin disease databases into one database. One-third of the collected literature use the mixture of multiple datasets [38], [42], [43], [46], [47], [49], [51], [52], [53], [58], [73], [76], [80], [85].

TABLE II
SKIN DISEASE DATASET

Name	Number of images	Disease classification	Dermoscopic Image (\sqrt{x})	Reference
Atlas [24]	—	—	$\sqrt{}$	[73]
PH2 [25]	200	2	$\sqrt{}$	[46], [58], and [66]
BCN2000 [27]	19424	8	$\sqrt{}$	[42]
MSK [28]	—	8	$\sqrt{}$	[42] and [71]
HAM10000 [29]	10015	7	$\sqrt{}$	[42], [49], [53], [54], [56], [57], [66], [71], [76], [79], [84], and [85]
ISIC Archive [30]	23906	7	$\sqrt{}$	[46], [58], [59], [70], [77], [79], [80], and [81]
DermIS [32]	—	—	\times	[46], [51], [58], and [80]
DermQuest [33]	—	—	\times	[46], [51], and [80]
DermNZ [34]	—	—	\times	[46] and [51]
ISIC 2016 [35]	1279	2	$\sqrt{}$	[47], [52], [53], [55], [64], [65], [73], [78], and [85]
ISIC 2017 [36]	2000	3	$\sqrt{}$	[44], [45], [47], [48], [50], [52], [53], [55], [61], [68], and [85]
MED-NODE [37]	170	2	$\sqrt{}$	[38], [58], [76], [78], and [80]
*ASan [39]	17125	12	\times	[38]
Dermofit [40]	1300	10	\times	[38], [41], [66], and [71]
*MIH SBIR [49]	1636	5	\times	[49]
*Acral melanoma [72]	724	2	$\sqrt{}$	[72]
*Pigmentary dermatosis [75]	12816	6	\times	[75]
Dermnet [97]	—	—	\times	[74]

*Self-collected and nonpublic.

C. IMAGE PREPROCESSING

In deep learning image recognition, a deep learning model has high requirements for image quality because a good image quality can improve the model's generalization ability [108]. Image preprocessing is carried out before model training. The primary purpose is to eliminate the irrelevant information in the image, enhance the detectability of the useful and related information in the image, and simplify the data to the great extent to improve the model's feature extraction ability and recognition reliability. This work has 28 studies on image preprocessing, which are divided into data cleaning and data conversion. Table 3 shows the details.

1) DATA CLEANING

Data cleaning is mainly to ensure the integrity of data features of the skin disease images. In skin disease image recognition, the commonly used data cleaning approach is to remove the noise to reduce hair and shadow influence on skin disease recognition. The image's quality is affected by the skin's nature, the environment, the equipment, and the

lighting conditions. The image with a low quality will affect the recognition effect, resulting in the loss of accuracy and calculation cost. The commonly used denoising algorithms include spatial domain filtering, transform domain filtering, and partial differential equation. In the selected literature, four papers carried out noise removal on the selected datasets. Hameed et al. and Hagerty et al. removed hair from the skin disease images [46],[49]. Singhal et al. used filters to reduce the influence of noise [57]. Rahul et al. used nonlocal means deoiling method to remove noise [58]. Xiaoyu added noise to the skin disease image to study image noise's influence on skin disease recognition [38].

2) DATA CONVERSION

The purpose of data transformation is to transform data from one format or structure to another according to the requirements of a deep learning model. In the collected literature, the commonly used data conversion technologies are size adjustment (18 articles), normalization (10 articles), and graying (three articles). Table III shows the details.

TABLE III
DATA PREPROCESSING

Type	Objective	Method	Reference
Data cleaning	Noise removal	Hair removal	[46] and [49]
		Wave filter	[57]
		Nonlocal means denoising	[58]
Data conversion	Size adjustment	Zoom	[47], [50], [51], [52], [53], [58], and [65]
		Clipping	[38], [44], [49], [53], [55], [56], [57], and [75]
		—	[41], [42], [59], and [70]
	Normalization	Data normalization	[41], [47], [48], [52], [53], [54], [56], [58], [65], [70], and [75]

Grayscale conversion	Converts the original color image to a grayscale image	[47], [57], and [70]
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Size adjustment: its input size limits the network model of deep learning, and the size of its input image is mostly fixed. If the input size in the network model is large, then the abstract level of information cannot meet the needs of the network, and the amount of calculation will increase; if the input size is small, then the picture information will be lost; hence, the network model in deep learning mostly uses 224×224 or 227×227 fixed-size input. Seventeen papers carried out size adjustment, among which four papers did not introduce the size adjustment method used, while the other literature mainly utilized scaling (seven articles) and clipping (eight articles). Mahbod used zooming and clipping [53], while Mahbod and Zhang used bilinear interpolation method for scaling [47], [50].

Normalization: in the selected literature, 11 papers normalized the images. Normalization is to convert the samples' eigenvalues to the same dimension: to map the data to the interval of $[0,1]$ or $[-1,1]$, and the extreme value of variables only determines the data. This method aims to limit the preprocessing data to a specific range to eliminate the particular sample data's adverse effects.

Grayscale conversion: gray conversion operates on a single pixel of an image; its primary purpose is to improve the contrast of the image and threshold processing. In the collected literature, Mahbod et al. [47], Singhal et al. [57], and Hasan et al. [70] used grayscale transformation to process skin disease pictures.

D. IMAGE DATA AUGMENTATION

Skin disease data are difficult to collect due to the problems of personal privacy and professional equipment involved in the collection process of the medical skin disease dataset. Accordingly, less skin disease data has been collected. Some diseases' rarity makes the data collection of this category less, resulting in the uneven distribution of the collected datasets. In deep learning, small-scale datasets can easily lead to insufficient model learning and overfitting. To solve the problem of small skin disease dataset and improve the network model's generalization ability, researchers use data augmentation technology to expand the amount of training data. Data augmentation uses existing data to create new data under the guidance of task objectives. The traditional image data augmentation expands the dataset by introducing geometric transformation and image operation to the original data without changing the data label. The leading technologies are rotation, mirror image, adding noise, and dimension reduction. The new data amplification technology produces simulation data on the basis of the original data and by generating Gans' model [63]. The internal distribution law of pictures indicates that the generated confrontation network is not only limited to within-class information but also uses the information between categories to synthesize pictures.

Among the 17 papers discussed in this study, 17 papers used data augmentation, 14 of which used the traditional data

augmentation technology to expand the dataset. Eleven papers used the rotation method, and the commonly used rotation angles were 90° , 180° , and 270° . Given that the generation of confrontation network is a new technology, only three papers (Bissoto et al. [66], Albarqouni et al. [68], and Xianzhen et al. [69]) were collected for the generation of simulation data of skin disease image. Among these papers, Bissoto et al. and Albarqouni et al. used the generated countermeasure network to generate data from the ISIC 2017 dataset. See Table IV for details.

TABLE IV
DATA AUGMENTATION

Type	Method	Reference
Traditional data augmentation	Rotate	[41], [44], [45], [48], [50], [52], [56], [60], [64], [65], [72], and [81]
	Image	[64]
	Clipping	[60] and [81]
Generation of simulation data	Generate an antagonistic network	[66], [68], and [69]

Although the traditional data amplification technology will increase the number of images, the diversity of data has not been improved due to the lack of learning the image's characteristics, which is just a simple copy transformation. The method of model generation is to generate images by learning the features of images. The generated model has strong feature learning and expression abilities compared with the traditional image amplification technology. The generated image is different from the original image, and the image quality is also higher. This method can effectively solve the unbalanced data amount of various skin disease categories, and the diversity of generated images has been dramatically improved.

E. MODEL AND FRAMEWORK

The advantage of deep learning over traditional machine learning is that significant feature representations are automatically learned from raw data, such as image pixels, to no longer rely on traditional feature engineering. Currently, the primary method of skin disease image recognition is to use a convolution neural network in deep learning, convolution, and pooling operation of convolutional network in image recognition, which has translation, rotation, and scale invariances. The CNN has excellent superiority in feature representation. All the literature collected in this study is based on the CNN model.

From the type of CNN used, the research work collected in this study adopts popular CNN architecture, such as AlexNet, VGG, Inception, ResNet, and DensenNet [86]-[90]. Eight articles are about AlexNet ([41], [42], [47], [52], [59], [60], [74], and [80]), seven articles used VGG ([47], [52], [54], [55], [60], [61], [72], and [73]), and seven articles utilized the Inception ([38], [43], [54], [57], [60],

[78], and [85]). Among the collected literature, at most 18 papers focused on skin disease identification by ResNet ([38], [42], [47], [49], [50], [52], [53], [57], [60], [64], [65], [75], [76], [77], [79], [82], [84], and [85]). At least four papers used DenseNet ([56], [57], [75], and [85]). Certain papers are related to some relatively new networks, such as Zhuang et al. using SENet ([42] and [91]) and Hameed et al. utilizing SqueezeNet ([51] and [92]).

From the number of network models used by researchers, the collected model methods can be classified into two types: the first one is a single model method based on deep learning; the other type is a multimodel method based on deep learning. Twenty-seven papers focused on a single model method and 10 papers on the multimodel method.

1) SKIN DISEASE IMAGE RECOGNITION BASED ON A SINGLE MODEL

Twenty-seven pieces of literature used a single model in the literature, among which 10 papers ([38], [38], [43], [57], [59], [64], [73], [75], [76], and [77]) has no model improvement, and the other 15 articles have optimized the

model. The optimization methods mainly include: modifying the fully connected layer (five articles: [41], [44], [61], [74], and [80]), using a machine learning classifier (two articles: [65] and [81]), applying (1 article: [51]) the model to the mobile application, introducing attention mechanism (two articles: [50] and [55]), self-building network model (two articles: [58] and [70]), and multilevel classification (three articles: [46], [56], and [78]). Kawahara et al. [41] and Yuexiang et al. [44] formed a fully CNN based on CNNs and removed the fully connected layers. Xueying et al. set the weights on the Softmax loss function to improve recognition accuracy [61]. Yan et al. [55] and Zhang et al. [50] introduced the attention mechanism in the CNNs. Zhang et al. proposed a CNN (ARL-CNN) on the basis of attention residual learning. The network embeds the residual and attention learning mechanisms into each ARL module, builds an ARL-CNN model with an arbitrary depth by stacking multiple ARL blocks, and trains it in an end-to-end manner. See Table V for details.

TABLE V
A. SINGLE MODEL

Model	Network features	Reference
AlexNet	The full junction layer is changed into a convolution layer, and logistic regression classifier is trained to classify skin lesions.	[41]
	According to the thought of divide and rule, the classification of multiple categories is multilevel.	[46]
	The AlexNet model is built, and different parameters are set up for the two classifications.	[59]
	The model of AlexNet is established, and the Softmax classifier is used to detect acne, keratosis, eczema, and urticarial	[74]
	Transfer learning is used to pretrain the model, and the new Softmax function is utilized for classification	[80]
VGGNet	The attention mechanism is introduced to regularize the attention graph.	[55]
	The transfer learning pretraining model is used to set the Softmax loss function's weight to alleviate data imbalance.	[61]
	The VGG model is fine-tuned by using 3×3 convolution kernel and transfer learning	[72]
	The VGG model is pre-trained by transfer learning.	[73]
Inception	A classification algorithm based on ImageNet is designed and trained in detail.	[43]
	The effect of noise on the skin disease image recognition is studied by using the Inception V3 model.	[38]
	Google's Inception v4 CNN architecture was trained and validated using dermoscopic images and corresponding diagnoses. In a comparative cross-sectional reader study, a 100-image test-set was used (level I: dermoscopy only; level II: dermoscopy plus clinical information and images). The results are compared with 25 dermatologists.	[78]
	The ImageNet dataset is used for pretraining and fine-tuning the model.	[38]
ResNet	ARL block is proposed, which uses residual and attention learning mechanisms to stack model.	[50]
	The deep residual network is constructed to extract the high-dimensional features of the dermoscopic image. Residual learning is used to prevent the gradient degradation of the network and reduce network training difficulty.	[64]
	The ImageNet pretraining model is used to extract the features of skin lesions images, and SVM is utilized to classify the extracted features.	[65]
	Transfer learning is used to pre-train the model. The tasks are compared with the 145 dermatologists in clinical melanoma image classification.	[76]
	Pretrained ResNet50 is used to compare with dermatologists.	[77]
SqueezeNet	The model is pretrained in ImageNet and fine-tuned. The trained model is placed on the server, and the remote connection between the mobile application and the server is established. The model is applied.	[51]
DenseNet	In the ImageNet pretraining model, the classification is divided into two steps: first, three categories are identified; then, the other four categories are determined.	[56]

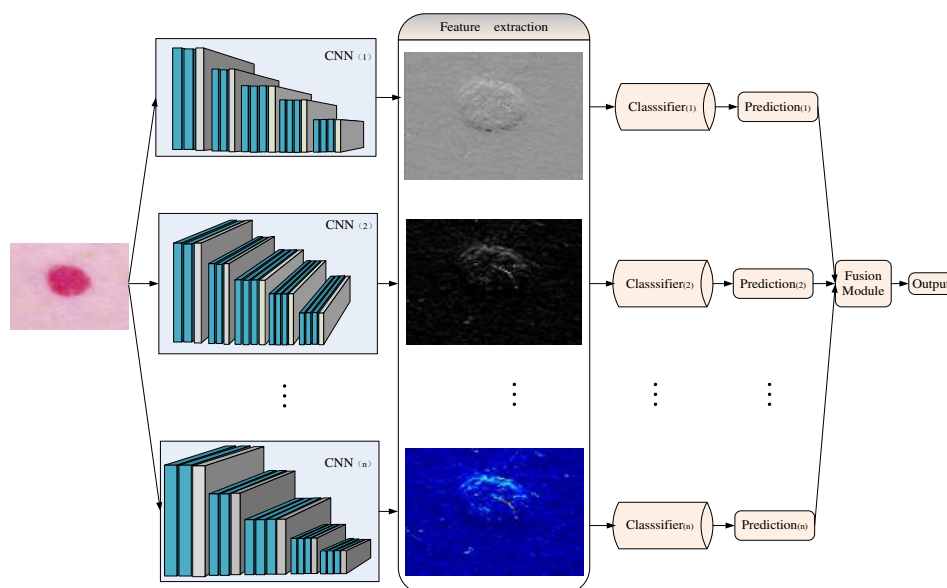
Others	The complete the convolutional residual network (FCRN) model can generate segmentation and rough classification results. A lesion index calculation unit is developed to refine the rough classification and calculate the distance thermal map to classify the results.	[44]
	Residual learning and separable convolution are used to identify the skin disease.	[58]
	Feature extraction is based on a CNN.	[70]
	The performance of these transfer learning models is compared by using four different latest architectures (Inception V3, ResNet 50, Densenet201, and Inception-ResNet V2)	[57]
	The objective is to compare ResNet-152 and DenseNet-96 in the pigmented skin disease	[75]
	The kernel of DePicT Melanoma Deep-CLASS is built upon a CNN composed of sixteen layers (excluding input and output layers), which can be recursively trained and learned. This method is a skin damage classification method that uses deep learning to detect melanoma early. A reasoning (CBR) system is proposed.	[81]
	Deep learning full resolution convolutional network is used. Then, a CNN classifier (i.e., Inception-v3, ResNet-50, Inception-ResNet-v2, and DenseNet-201) is applied on the segmented skin lesions for classification.	[85]

2) SKIN DISEASE IMAGE RECOGNITION BASED ON MULTIMODEL FUSION

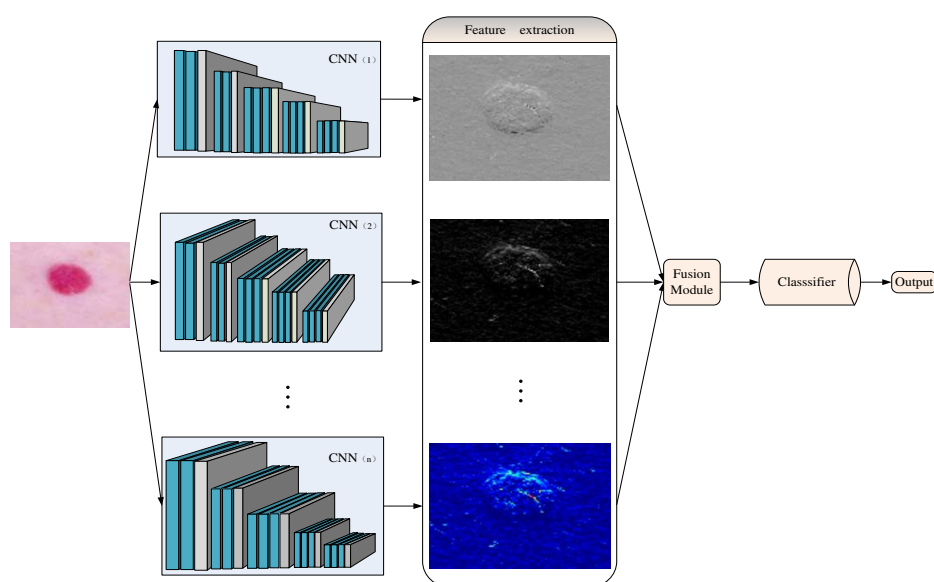
Multimodel fusion is mainly to learn through multiple learners and integrate multiple learners' results by using some rules. In the conventional multimodel fusion, the average method, weighting method, linear regression method, simple majority voting (SMV) rule, and maximum probability rule are used to fuse and recalculate the probability results of a certain type of output of each model. The final output of the fusion results is obtained. The other way is to fuse the features learned from multiple models to fuse the features and then output the final results. The flow chart of the above-mentioned two fusion methods is showed in Figure 5. Fourteen pieces of literature are about multimodel fusion collected in this study, and the details are shown in Table VI. The main methods are multi-input

model fusion (two articles), multimodel extraction feature SVM classification (two articles), fusion strategy combined with clinical information (four articles), the combination of human and artificial intelligence (two articles), and conventional multiple model fusion (four articles). AlexNet, VGG, Inception, and ResNet are often used for fusion.

A large number of experiments have proved that the multimodel fusion method is better than the single model method [60], [62] in the field of skin disease recognition. Given that the objective factors limit a single model, a model generalization bottleneck is commonly encountered when dealing with problems. The multimodel fusion can combine the excellent models and integrate each model's advantages to break through the bottleneck of the generalization ability of a single model



(a) Conventional multimodel fusion



(b) Multimodel fusion of feature fusion

FIGURE 5. Two fusion methods of multimodel fusion. The figure is the conventional multimodel fusion method. After the image is inputted, the feature is extracted by multiple CNNs and then categorized by the classifier to obtain the probability value of each sample image category. Finally, the probability value of the classification is fused by the fusion module to output the category of the maximum probability value. Panel B is the multimodel fusion of feature fusion, which mainly includes the following steps: After feature extraction, the CNN fuses the extracted features, and then the probability value of each category is outputted by the classifier. The output with the maximum probability value is selected.

TABLE VI
MULTIMODEL FUSION

Integration strategy	Model features	Reference
Multi-input model fusion	Different Gabor wavelets are used to represent the input, and some rules are used for decision fusion.	[45]
	Six different sizes of images are inputted by using a three-layer fusion scheme. The averaging strategy is used for each layer.	[53]
Multimodel feature extraction SVM classification	The deep features of multiple fine-tuned DNNs are extracted from the lesion image. Then, the features are integrated for classification by the SVM classifier.	[47]
	Three CNNs are utilized to extract features, and SVM is used to classify features.	[52]
Fusion strategy combined with clinical information	Multimodel fusion based on age and gender.	[48]
	Six graphic processing modules and one clinical information module were combined.	[49]
	A method that combines multiple imaging methods with patient metadata is proposed to improve the performance of the automatic skin lesion diagnosis.	[82]
	The output of two CNNs are merged, one of which has been trained on dermoscopy images, and the other is clinical close-ups and enhanced by extreme the gradient combination probability of XGBoost.	[83]
Conventional multimodel fusion	The cost-sensitive matrix is using to set different weights for model fusion.	[42]
	The averaging strategy is used to fuse multiple models.	[54]
	The sum of probability rule (SP), probability product rule (PP), SMV rule, and maximum probability rule (SMP) are used to fuse the model. Weights are calculating by using a simulated degeneration algorithm and a genetic algorithm [67].	[60]
	The maximum correlation entropy criterion is proposed as the objective function on the basis of the traditional fusion strategy.	[62]
Combination of human and artificial intelligence	Dermatologists and CNNs are fused on the basis of the ResNet50 architecture, and the fusion algorithm is XGBoost.	[79]
	This study presents a method of human-computer cooperation in the treatment of skin cancer	[84]

3) DEEP LEARNING FRAMEWORK

Among the collected literature, only 16 papers published the deep learning framework, including Keras [93], TensorFlow [94], MATLAB [95], and PyTorch [96]. Keras has become a popular framework owing to its consistent and concise API, which can significantly reduce the workload of users. Six articles have used the framework. Tensorflow is often used with Keras. The framework can deploy training models on various servers and mobile devices without executing a separate model decoder or

loading a Python interpreter. Pytorch is a deep learning framework released by Facebook AI research in 2017. This framework has the advantages of flexibility, ease of use, and fast speed. PyTorch is a rookie in the deep learning framework. Five articles collected use the framework. MATLAB has a long history of development and has advantages in the image, video, and audio data for accurate value annotation. Five papers use MATLAB. The details are shown in Table VII.

TABLE VII
DEEP LEARNING FRAMEWORK

Frame	Major function	Reference
PyTorch	GPU acceleration and automatic derivation function of the robust deep neural network	[42], [55], [56], [61], and [71]
MATLAB	Convenient results are viewed, statistics, and visualization	[47], [51], [60], [64], and [65]
Keras	It is applied to the programming implementation of quickly converting ideas into results.	[38], [44], [48], [53], [54], and [65]
TensorFlow	It is applied to the programming of various machine learning algorithms.	[53] and [54]

F. EVALUATING INDICATOR

To evaluate the performance of deep learning in skin disease image recognition, several performance indicators are used: accuracy (ACC) represents the percentage of correct prediction results in the total sample [100]; mean average precision (mAP) represents the average accuracy of all categories [101]; true positive rate (TPR), also known as sensitivity and Recall (R) [102], represents the probability of being predicted to be positive in the actual positive samples [103]; false positive rate (FPR) refers to the percentage of actual disease-free but judged to be disease-free; true

negative rate (TNR) [104], also known as specificity, indicates that the actual disease-free is correctly judged to be disease-free; area under the receiver operating characteristic (ROC) curve (AUC) refers to the probability that the classifier outputs positive and negative samples, and the likelihood that the classifier outputs a positive sample is greater than that of the negative sample; ROC is the working characteristic curve of subjects, which shows the performance of classification model under all classification thresholds [105]. The specific performance indicators are shown in Table VIII.

TABLE VIII
PERFORMANCE INDEX

Index	Formula	Explain	Reference
ACC	$Acc = \frac{TP + TN}{TP + TN + FN + FP}$	Accuracy of identification, classification, or prediction	[38], [41], [45], [50], [53], [56], [57], [59], [64], [65], [42], [44], [51], [58], [62], [71], [72], [74], [79], [80], [82], [84], and [85]
mAP	$mAP = 1 / C(\sum_{i=1}^N Acc(k) \Delta R(k))$	Accuracy of prediction target and category	—
TPR	$TPR = TP / (TP + FN)$	Among the positive samples, the correct proportion of samples was predicted.	[38], [44], [50], [51], [54], [61], [62], [64], [65], [71], [75], [76], [77], [78], [79], [80], [81], [83], [84], and [85]
FPR	$FPR = FP / (FP + FN)$	Among the negative samples, the proportion of samples with false predictions	—
TNR	$TNR = TN / (FP + TN)$	Among the negative samples, the correct proportion of samples was predicted	[44], [45], [50], [51], [64], [65], [71], [75], [76], [77], [78], [79], [80], [81], [83], and [85]

False negative rate (FNR)	$FNR = FN / (TP + FN)$	Among the positive samples, the proportion of false prediction samples	–
AUC	ROC AUC	ROC is the working characteristic of the curve of subjects, which shows a classification model's performance under all classification thresholds.	[47], [48], [49], [50], [52], [54], [55], [60], [64], [65], [72], [73], [75], [76], [78], [81], [82], and [83]

G. MODEL PERFORMANCE AND ANALYSIS

Two papers are difficult, or even impossible, to compare because the datasets used by researchers in various tasks, the types of skin disease identified, models, parameters, and performance indicators are different. Therefore, readers should carefully consider the comments in this section.

Among the 29 papers with accuracy indicators, four papers have an accuracy rate of less than 70%, 15 papers have an accuracy rate of 80% to 89%, and the other nine papers have an accuracy rate of more than 90%. This finding shows the good generalization ability of deep learning, and the recognition accuracy of the model built by Sarkar et al. is as high as 99.5% [58]. Among the 24 papers that use the AUC

as a measure of performance, nine papers have more than 90% of the AUC, and the highest area under the ROC curve reaches 97.5% [52]. Eleven articles have a range of 80% to 89%, and only four articles are below 80%. In terms of sensitivity, 28 papers are used as evaluation indicators, 10 papers have a sensitivity of more than 90%, six papers have a sensitivity of 80% to 89%, and 12 papers have a sensitivity lower than 80%. Twenty-four papers have specificity as an indicator, 11 papers have specificity higher than 90%, five papers are between 80% and 89%, and the other eight papers are less than 80%. See Table IX for details.

TABLE IX
PERFORMANCE IN THE LITERATURE

Source	ACC	Sensitivity	Specificity	AUC
Han et al. [39]	–	86.4 ± 3.5	85.5 ± 3.2	91% ± 0.01
Kawahara et al. [41]	85.8%	–	–	–
Zhuang et al. [42]	90.47%	–	–	–
Esteva et al. [43]	72.1 ± 0.9%	–	–	91%
Li et al. [44]	90.2%	69.3%	90.2%	84.8%
Serte et al. [45]	83%	62%	88%	96%
Hameed et al. [46]	96.47%	99.5%	98.17%	–
Mahbod et al. [47]	87.7%	95%	44.72%	89.85%
Matsunaga et al. [48]	–	–	–	92.4%
Hagerty et al. [49]	–	–	–	94%
Zhang et al. [50]	85%	65.8%	89.6%	87.5%
Hameed et al. [51]	97.21%	94.42%	98.14%	–
Mahbod et al. [52]	–	–	–	90.69%
Mahbod et al. [53]	86.2%	–	–	98.1%
Aldwgeri et al. [54]	80.1%	80%	98%	89%
Yan et al. [55]	–	–	–	85.2%
Moldovan et al. [56]	85%	–	–	–
Singhal et al. [57]	81.62%	–	–	97%
Sarkar et al. [58]	99.5%	99.31%	100%	99.49%
Bagadi et al. [59]	76.66%	87.9%	67%	–
Harangi et al. [60]	86.6%	55.6%	78.5%	89.1%
Xueying et al. [61]	71.34%	70.01%	–	–
Xu et al. [62]	97.07%	96.55%	97.33%	–
Qiu et al. [64]	87%	58.6%	96.7%	79%
Hang et al. [65]	86.28%	44%	96.71%	84.18%
Xianzher et al. [69]	69.25%	72.67%	–	–
Hasan et al. [70]	93.7%	84%	–	–
Haiting et al. [71]	93.4%	91.26%	100%	–
Yu et al. [72]	83.51%	92.57%	75.39	82%
Menegola et al. [73]	79.5%	47.6%	88.1%	80.7%
Shanthi et al. [74]	98.6%-99.04%	–	–	–
Yang et al. [75]	–	93.2%	98.9%	98%
Brinker et al. [76]	–	92.8%	61.1%	76.9%
Brinker et al. [77]	–	82.3%	77.9%	–
Haenssle et al. [78]	–	86.6% ± 9.3%	71.3 ± 11.2%	88.9%
Hekler et al. [79]	82.95%	84%	81.5%	–
Hosny et al. [80]	97.7%	97.34%	97.34%	–
Nasiri et al. [81]	–	73%	78%	75%
Yap et al. [82]	72.1%	–	–	86.6%

Tschandl et al. [83]	–	80.5%	37.6%	74.2%
Tschandl et al. [84]	81%	77.7%	–	–
Al-masni et al. [85]	89.28%	81%	87.16%	–

This paper analyses the collected literature and concludes that deep learning provides excellent performance in most related work. The deep learning-based methods used in 45 papers are effectively and correctly compared. The papers are difficult to summarize because the datasets, preprocessing techniques, indicators, models, and parameters involved in each article are different. Accordingly, the comparison of this study is strictly limited to the techniques used in each paper. Based on these constraints, we observed that deep learning performance is better than that of traditional methods, and the automatic feature extraction of deep learning models is more efficient than that of traditional methods (such as color [106], histogram [107], statistics, and texture [108]). Hang et al. used deep learning to extract the feature vectors of skin disease and categorized them by using SVM as a classifier [65]. The AUC of the model performance of the manual feature extraction method is 6.17% higher than those of traditional manual feature extraction methods. The multimodel fusion performance of a single convolution network is better than that of single-model skin disease image recognition. Harangi et al. and Xu Meifeng et al. compared the performance of multimodel fusion with that of a single model in their respective experiments [60], [62]. The results show that the performance of the multimodel fusion is better than that of a single network model. The multimodel approach combines the advantages of every single model to obtain the optimal solution. Another way of combining multiple models is to fuse the patient's clinical information (e.g., gender, age, and location of lesions.). Hagerty et al. used this method to resolve the patient's information [49], excluding the image recognition of skin disease. The ROC area increased from 83% to 94% for deep learning only.

V. SUMMARY AND PROSPECT

A. SUMMARY

In this article, we outline the development of skin disease diagnostic technology and the process of traditional medical diagnosis and machine learning-based image recognition of skin disease and investigate the results of deep learning-based research in the field of skin disease recognition. Forty-five relevant papers have been identified to obtain their concerns about the areas and skin disease types. These papers are utilized as basis to study the used data source, the preprocessing and data expansion techniques, the technical details of the models, and the performance indicators' overall performance. Deep learning models AlexNet, VGG, GoogleNet, and ResNet are widely used in skin disease recognition. Researchers often use the multimodel fusion technology to improve the performance of models. In future work, we plan to apply the concepts and best practices of

deep learning described in this survey to other medical fields that have not fully utilized this technology. This survey aims to encourage many researchers to conduct deep learning experiments and apply the model of deep learning in the field of computer vision involving medicine, thereby achieving smart and convenient development for the medical industry.

B. PROSPECT

Deep learning, a new field of machine learning, has good application prospects in computer vision and provides a new direction for medical image recognition. Currently, deep learning has achieved empirical progress in the field of medical image recognition [98]. However, deep learning is still in its infancy in medical image recognition. Additional developments and comprehensive attempts should be carried out in the field of skin disease recognition in the future. The following summarizes four possible directions for research development:

(1) Simulation data generation of skin disease image

Deep learning has good generalization ability in skin disease recognition; however, it requires many data in the learning process. Few data may cause insufficient feature extraction, which will affect the diagnosis and recognition of lesions. One of the significant problems in skin disease image recognition is the difficulty in obtaining a large number of data sets due to the complications in medical image collection and the rare occurrence of individual diseases. Generating simulated pictures of skin disease by using antinetwork is an excellent method to solve this problem. Such an approach also improves the performance of the network. This technology will have good development prospects in the field of skin disease recognition in the future.

(2) Clinical identification of skin disease

At present, the types of pictures recognized in the field of skin disease recognition are relatively single. Most graphics are based on the pictures generated by the dermoscopic imaging technology. Few studies have been conducted on the pictures taken in clinical or real life. The location of the lesion is difficult to identify due to several problems, such as low pixels and presence of shadows, and the lesion area only occupies a small part or incomplete in clinical or real scenes pictures. This aspect needs further study. In clinical recognition models, attention mechanisms or target detection methods can be introduced to focus on the lesion sites of skin disease for facilitating the feature extraction of lesion sites to improve the accuracy of clinical recognition models.

(3) Interpretability study on skin disease identification

Now, researchers focus on the performance indicators of models. However, various factors in the environment result in the transformation of model concepts and change of performance. The factors that drive the model should be

understood to make certain decisions. Currently, the interpretability research on skin disease identification models is a big gap in this field and must be explored. In the skin disease recognition field, deep learning models can be applied to identify skin disease to guide clinical automated medical diagnosis. However, these algorithms are still a "black box" to generate predictions on the basis of input data. There is no specific interpretation on what skin disease features does the deep learning model as a basis for judgment. The development of deep learning should be trustworthy and explainable. Making the algorithm public and transparent in decision-making will give users reliability and security. Interpretability research on skin disease identification can resolve prejudices and auditing brought about by artificial intelligence [109]. Interpretability makes artificial intelligence open and transparent in legal, moral, and philosophical aspects.

(4) AI diagnosis and treatment of skin disease

Mobile devices, such as smartphones, PDAs, and tablets, are becoming an essential part of human life [109]. Embedding AI diagnosis and treatment of skin disease on smart devices will be a significant trend in the future. However, most skin diseases are diagnosed on the basis of high-performance graphics processors. The computational complexity of the algorithm should be minimized while improving the algorithm recognition capability to ensure that it can be easily used on mobile phones and wearable intelligent devices [110]. This study is of great significance for AI diagnosis and treatment of skin diseases. In intelligent questions and answers on skin disease, AI assistants can replace dermatologists to inquire about patients and communicate with them about some repetitive contents, such as diagnosis, prescription, and health promotion. At present, some issues related to philosophy, law, and ethics still persist in addition to technical bottlenecks in AI diagnosis and treatment. Examples of the questions are as follows: Is the subject of AI diagnosis and treatment a human or a medical device on the legal level?; what is the legal standard in the clinical application?; what are the criteria for judging a medical defect or medical negligence?; and who is legally responsible for medical accidents in AI diagnosis and treatment?.

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REFERENCES

- [1] N. HameedK. Abuhassan and M. Hossain, "A comprehensive survey on image-based computer aided diagnosis systems for skin cancer," 2016.
- [2] H. N.S. A. and A.H. M., "A Computer-aided diagnosis system for classifying prominent skin lesions using machine learning, Proc. 2018 10th Computer Science and Electronic Engineering (CEECE), 2018, pp. 186-191.
- [3] M. Johnson and J. Roberts, "Skin conditions and related need for medical care among persons 1-74 years. United States, 1971-1974," Vital and health statistics. Series 11, Data from the national health survey, vol. 212, pp. 1, 1978.
- [4] "British skin foundation." [Online]. Available:http://www.britishtskinfoundation.org.uk.[Accessed: 20-May-2018].
- [5] A. PicardiI. Lega and E. Tarolla, "Suicide risk in skin disorders," Clin. Dermatol., vol. 31, pp. 47-56, 2013, doi: 10.1016/j.clindermatol.2011.11.006.
- [6] H.W. Rogers, M.A. Weinstock, S.R. Feldman and B.M. Coldiron, "Incidence Estimate of Nonmelanoma Skin Cancer (Keratinocyte Carcinomas) in the US Population, 2012," JAMA Dermatol., 2015.
- [7] K. Freedberg, A. Geller, D. Miller, R. Lew and H. Koh, "Screening for malignant melanoma: A cost-effectiveness analysis," J. Am. Acad. Dermatol., vol. 41, pp. 738-745, 1999, doi: 10.1016/S0190-9622(99)70010-1.
- [8] V. Dick, C. Sinz, M. Mittlböck, H. Kittler and P. Tschandl, "Accuracy of Computer-Aided Diagnosis of Melanoma: A Meta-analysis," JAMA Dermatol., vol. 155, 2019, doi: 10.1001/jamadermatol.2019.1375.
- [9] N. Cascinelli, M. Ferrario, T. Tonelli and E. Leo, "A possible new tool for clinical diagnosis of melanoma: The computer," J. Am. Acad. Dermatol., vol. 16, no. 2, Part 1, pp. 361-367, 1987, doi: https://doi.org/10.1016/S0190-9622(87)70050-4.
- [10] R. StanleyW. Stoecker and R. Moss, "A relative color approach to color discrimination for malignant melanoma detection in dermoscopy images," Skin research and technology : official journal of International Society for Bioengineering and the Skin (ISBS) [and] International Society for Digital Imaging of Skin (ISDIS) [and] International Society for Skin Imaging (ISSI), vol. 13, pp. 62-72, 2007, doi: 10.1111/j.1600-0846.2007.00192.x.
- [11] N. Xiong, Y. Shen, K. Yang, C. Lee and W. Chunxue, "Color sensors and their applications based on real-time color image segmentation for cyber physical systems," EURASIP J. Image Vide., vol. 2018, pp. 23, 2018, doi: 10.1186/s13640-018-0258-x.
- [12] W. StolzA. Reimann and A. Cognetta, "ABCD rule of dermoscopy: A new practical method for early recognition of malignant melanoma," vol. 4, 1993.
- [13] G. Argenziano, G. Fabbrocini, P. Carli, V. De Giorgi, E. Sammarco and M. Delfino, "Epiluminescence Microscopy for the Diagnosis of Doubtful Melanocytic Skin Lesions," Archives of Dermatology, vol. 134, 1998, doi: 10.1001/archderm.134.12.1563.
- [14] P. Soyer, et al., "Three-Point Checklist of Dermoscopy," Dermatology (Basel, Switzerland), vol. 208, pp. 27-31, 2004, doi: 10.1159/000075042.
- [15] J. Henning, et al., "The CASH (color, architecture, symmetry, and homogeneity) algorithm for dermoscopy," J. Am. Acad. Dermatol., vol. 56, pp. 45-52, 2007, doi: 10.1016/j.jaad.2006.09.003.
- [16] C. Rosendahl, A. Cameron, I. Mccoll and D. Wilkinson, "Dermoscopy in routine practice: Chaos and Clues," Aust. Fam. Physician, vol. 41, pp. 482-487, 2012.
- [17] R. GarnaviM. Aldeen and J. Bailey, "Computer-Aided Diagnosis of Melanoma Using Border- and Wavelet-Based Texture Analysis," IEEE transactions on information technology in biomedicine : a publication of the IEEE Engineering in Medicine and Biology Society, vol. 16, 2012, doi:

- 10.1109/TITB.2012.2212282.
- [18] L. Ballerini, R.B. Fisher, B. Aldridge and J. Rees, A Color and Texture Based Hierarchical K-NN Approach to the Classification of Non-melanoma Skin Lesions, *Color Medical Image Analysis, Lecture Notes in Computational Vision and Biomechanics*, M. E. Celebi and G. Schaefer, eds., Springer Netherlands, 2013, pp. 63-86.
 - [19] J. Vasquez and B. Comendador, "Competency Discovery System: Integrating the Enhanced ID3 Decision Tree Algorithm to Predict the Assessment Competency of Senior High School Students," *International Journal on Advanced Science, Engineering and Information Technology*, vol. 9, pp. 60, 2019, doi: 10.18517/ijaseit.9.1.7763.
 - [20] P.L. Gutierrez and S. Siva, "Classification and Regression Tree (CART)," Springer Netherlands, 2008.
 - [21] B. He, et al., "Fast automatic 3D liver segmentation based on a three-level AdaBoost-guided active shape model," *Med. Phys.*, 2016.
 - [22] Wang Ting, Zhang Ning, Hou Guirong. and Yu Xuefei, "Performance comparison of several machine learning methods for computer-aided diagnosis of melanoma," *Application Research of Computers*, vol. 30, no. 06, pp. 1731-1733, 2013.
 - [23] M.A. Marchetti, et al., "Results of the 2016 International Skin Imaging Collaboration International Symposium on Biomedical Imaging challenge: Comparison of the accuracy of computer algorithms to dermatologists for the diagnosis of melanoma from dermoscopic images," *J. Am. Acad. Dermatol.*, vol. 78, no. 2, pp. 270-277, 2018, doi: 10.1016/j.jaad.2017.08.016.
 - [24] H.S.V.G. G. Argenziano, "Dermoscopy: a tutorial," *Medical Publishing & New Media*, 2002.
 - [25] M. T., M.F. P., S.M. J., R.S.M. A. and R. J., PH2 - A dermoscopic image database for research and benchmarking, *Proc. 2013 35th Annual International Conference of the IEEE Engineering in Medicine and Biology Society (EMBC)*, 2013, pp. 5437-5440.
 - [26] Z.F. Mohammed and A.A. Abdulla, "An efficient CAD system for ALL cell identification from microscopic blood images," *Multimed. Tools Appl.*, 2020, doi: 10.1007/s11042-020-10066-6.
 - [27] M. Combalia, et al., BCN20000: Dermoscopic Lesions in the Wild, *Book BCN20000: Dermoscopic Lesions in the Wild*, ed., Editor ed., 2019, pp. 1908-2288.
 - [28] C.F.C. N., et al., Skin lesion analysis toward melanoma detection: A challenge at the 2017 International symposium on biomedical imaging (ISBI), hosted by the international skin imaging collaboration (ISIC), *Proc. 2018 IEEE 15th International Symposium on Biomedical Imaging (ISBI 2018)*, 2018, pp. 168-172.
 - [29] P. Tschandl, C. Rosendahl and H. Kittler, "The HAM10000 dataset, a large collection of multi-source dermoscopic images of common pigmented skin lesions," *Scientific Data*, vol. 5, no. 1, pp. 180161, 2018, doi: 10.1038/sdata.2018.161
 - [30] "ISIC project-ISIC Archive" [Online]. Available: <https://www.isic-archive.com>. [Accessed: 23-May-2019].
 - [31] R.H. J., et al., "Deep Learning and Handcrafted Method Fusion: Higher Diagnostic Accuracy for Melanoma Dermoscopy Images," *IEEE J. Biomed. Health*, vol. 23, no. 4, pp. 1385-1391, 2019, doi: 10.1109/JBHI.2019.2891049.
 - [32] "DermIS," [Online]. Available: <http://www.dermis.net/dermisrootle/home/index.htm>. [Accessed: 29-Jun-2017].
 - [33] "Derm101 Image Library", [Online]. Available: <https://www.derm101.com/image-library/>. [Accessed: 12-Jan-2018].
 - [34] "DermNZ-ImageLibrary", [Online]. Available: <https://www.dermnetnz.org/image-library/>. [Accessed: 13-Jan-2018].
 - [35] D. Gutman, et al., "Skin Lesion Analysis toward Melanoma Detection: A Challenge at the International Symposium on Biomedical Imaging (ISBI) 2016, hosted by the International Skin Imaging Collaboration (ISIC)," *arXiv:1605.01397 [cs.CV]*, 2016.
 - [36] N.C.F. Codella, et al., Skin Lesion Analysis Toward Melanoma Detection: A Challenge at the 2017 International Symposium on Biomedical Imaging (ISBI), Hosted by the International Skin Imaging Collaboration (ISIC), *Book Skin Lesion Analysis Toward Melanoma Detection: A Challenge at the 2017 International Symposium on Biomedical Imaging (ISBI)*, Hosted by the International Skin Imaging Collaboration (ISIC), ed., Editor ed., 2017, pp. 1710-5006.
 - [37] "Med-node dataset" [Online]. Available : http://www.cs.rug.nl/~imaging/databases/melanoma_naevi/. [Accessed: 13-Jan-2018].
 - [38] F. X., et al., "Effect of image noise on the classification of skin lesions using deep convolutional neural networks," *Tsinghua Science and Technology*, vol. 25, no. 3, pp. 425-434, 2020, doi: 10.26599/TST.2019.9010029.
 - [39] S.S. Han, M.S. Kim, W. Lim, G.H. Park, I. Park and S.E. Chang, "Classification of the Clinical Images for Benign and Malignant Cutaneous Tumors Using a Deep Learning Algorithm," *J. Invest. Dermatol.*, vol. 138, no. 7, pp. 1529-1538, 2018, doi: <https://doi.org/10.1016/j.jid.2018.01.028>.
 - [40] "American Cancer Society-Melanoma skin cancer" [Online]. Available : <http://www.cancer.org/acs/groups/cid/documents/webcontent/003120.pdf>. [Accessed: 23-Oct-2015].
 - [41] K. J.B. A. and H. G., Deep features to classify skin lesions, *Proc. 2016 IEEE 13th International Symposium on Biomedical Imaging (ISBI)*, 2016, pp. 1397-1400.
 - [42] D. Zhuang, K. Chen and J. Chang, "CS-AF: A Cost-sensitive Multi-classifier Active Fusion Framework for Skin Lesion Classification," 2020.
 - [43] A. Esteva, et al., "Dermatologist-level classification of skin cancer with deep neural networks," *Nature*, vol. 542, no. 7639, pp. 115-118, 2017, doi: 10.1038/nature21056.
 - [44] Y. Li and L. Shen, "Skin Lesion Analysis towards Melanoma Detection Using Deep Learning Network," *Sensors (Basel, Switzerland)*, vol. 18, no. 2, pp. 556, 2018, doi: 10.3390/s18020556.
 - [45] S. Serte and H. Demirel, "Gabor wavelet-based deep learning for skin lesion classification," *Comput. Biol. Med.*, vol. 113, pp. 103423, 2019, doi: <https://doi.org/10.1016/j.compbiomed.2019.103423>.
 - [46] N. Hameed, A. Shabut, M. Ghosh and A. Hossain, "Multi-Class Multi-Level Classification Algorithm for Skin Lesions Classification using Machine Learning Techniques," *Expert Syst. Appl.*, vol. 141, pp. 112961, 2019, doi: 10.1016/j.eswa.2019.112961.
 - [47] A. Mahbod, G. Schaefer, I. Ellinger, R. Ecker, A. Pitiot and C. Wang, "Fusing fine-tuned deep features for skin lesion classification," *Comput. Med. Imag. Grap.*, vol. 71, pp. 19-29, 2019, doi: <https://doi.org/10.1016/j.compmedimag.2018.10.007>.
 - [48] K. Matsunaga, A. Hamada, A. Minagawa and H. Koga, "Image Classification of Melanoma, Nevus and Seborrheic Keratosis by Deep Neural Network Ensemble," 2017.
 - [49] R.H. J., et al., "Deep Learning and Handcrafted Method Fusion: Higher Diagnostic Accuracy for Melanoma Dermoscopy Images," *IEEE J. Biomed. Health*, vol. 23, no. 4, pp. 1385-1391, 2019, doi: 10.1109/JBHI.2019.2891049.
 - [50] Z. J., X. Y., X. Y. and S. C., "Attention Residual Learning for Skin Lesion Classification," *IEEE T. Med. Imaging*, vol. 38, no. 9, pp. 2092-2103, 2019, doi: 10.1109/TMI.2019.2893944.
 - [51] N. Hameed, A. Shabut, F. Hameed, S. Cirstea, S. Harriet and A. Hossain, "Mobile-based Skin Lesions Classification Using Convolution Neural Network," *Annals of Emerging Technologies in Computing*, vol. 4, pp. 26-37, 2020, doi: 10.33166/AETIC.2020.02.003.
 - [52] A. Mahbod, R. Ecker and I. Ellinger, "Skin Lesion Classification Using Hybrid Deep Neural Networks," 2017.
 - [53] A. Mahbod, G. Schaefer, C. Wang, G. Dorfner, R. Ecker and I. Ellinger, "Transfer learning using a multi-scale and multi-network ensemble for skin lesion classification," *Comput. Meth. Prog. Bio.*, vol. 193, pp. 105475, 2020, doi:

- https://doi.org/10.1016/j.cmpb.2020.105475.
- [54] A. Aldwgeri and N. Abubacker, Ensemble of Deep Convolutional Neural Network for Skin Lesion Classification in Dermoscopy Images, 2019, pp. 214-226.
- [55] Y. YanJ. Kawahara and G. Hamarneh, Melanoma Recognition via Visual Attention, 2019, pp. 793-804.
- [56] M. D., Transfer Learning Based Method for Two-Step Skin Cancer Images Classification, *Proc. 2019 E-Health and Bioengineering Conference (EHB)*, 2019, pp. 1-4.
- [57] A. Singhal, R. Shukla, P. Kankar, S. Dubey, S. Singh and R. Pachori, "Comparing the capabilities of transfer learning models to detect skin lesion in humans," *Proceedings of the Institution of Mechanical Engineers Part H Journal of Engineering in Medicine*, 2020, doi: 10.1177/0954411920939829.
- [58] R. SarkarC. Chatterjee and A. Hazra, "Diagnosis of melanoma from dermoscopic images using a deep depthwise separable residual convolutional network," *IET Image Process.*, vol. 13, 2019, doi: 10.1049/iet-ipr.2018.6669.
- [59] A. U.J.D. I. and R. R., Convolutional Neural Network based Skin Lesion Classification and Identification, *Proc. 2020 International Conference on Inventive Computation Technologies (ICICT)*, 2020, pp. 264-270.
- [60] Balazs and Harangi, "Skin lesion classification with ensembles of deep convolutional neural networks.," *J. Biomed. Inform.*, 2018.
- [61] H.E. XueyingH. Zhongyi and W. Benzhen, "Pigmented skin lesion recognition and classification based on deep convolutional neural network," *Journal of Computer Applications*, 2018.
- [62] X.U. Meifeng, et al., "Skin Disease Recognition Method Based on Multi-Model Fusion of Convolutional Neural Network," *Journal of Xi'an Jiaotong University*, 2019.
- [63] K. Wang, C. Gou, Y.J. Duan, L. Yilun and X.H. Zheng, "Generative Adversarial Networks: The State of the Art and Beyond," *Zidonghua Xuebao/Acta Automatica Sinica*, vol. 43, pp. 321-332, 2017, doi: 10.16383/j.aas.2017.y000003.
- [64] GUAN Qiu,LI Jiang,HU Haigen, et al. "Melanoma skin lesion recognition based on deep residual network," *Journal of Zhejiang University of Technology*, no. 4, 2019.
- [65] L. Hang, Y. Zhen, N. Dong, B. Lei and W. Tianfu, "Melanoma recognition in dermoscopy images via deep residual network," *Chinese Journal of Biomedical Engineering*, vol. 37, pp. 274-282, 2018, doi: 10.3969/j.issn.0258-8021.2018.03.003.
- [66] A. Bissoto, F. Perez, E. Valle and S. Avila, "Skin Lesion Synthesis with Generative Adversarial Networks," 2019.
- [67] F. Long, N. Xiong, A.V. Vasilakos, L.T. Yang and F. Sun, "A sustainable heuristic QoS routing algorithm for pervasive multi-layered satellite wireless networks," *Wirel. Netw.*, vol. 16, no. 6, pp. 1657-1673, 2010, doi: 10.1007/s11276-009-0220-z.
- [68] C. BaurS. Albarqouni and N. Navab, "MelanoGANs: High Resolution Skin Lesion Synthesis with GANs," 2018.
- [69] H.M.S.Y. SHANG Xianzhen, "Skin Diseases Diagnosis Method Based on Generative Adversarial Networks and Naive Bayes," *Journal of Frontiers of Computer Science and Technology*, vol. 13, no. 6, pp. 1005-1015, 2019, doi: 10.3778/j.issn.1673-9418.1806019.
- [70] M. Hasan, S. Barman, S. Islam and A. Reza, "Skin Cancer Detection Using Convolutional Neural Network," 2019, pp. 254-258.
- [71] Z.Q.C.M. JING Haiting, "Few-Shot Domain Adaptation for the Identification of Clinical Imaging for Dermatology," *Journal of Xi'an Jiaotong University*, pp. 1-8, 2020.
- [72] C. Yu, et al., "Acral melanoma detection using a convolutional neural network for dermoscopy images," *PLoS One*, vol. 13, pp. e193321, 2018, doi: 10.1371/journal.pone.0193321.
- [73] M. A., F. M., P. R., V.B. F., A. S. and V. E., Knowledge transfer for melanoma screening with deep learning, *Proc. 2017 IEEE 14th International Symposium on Biomedical Imaging (ISBI 2017)*, 2017, pp. 297-300.
- [74] T. ShanthiR.S. Sabeenian and R. Anand, "Automatic diagnosis of skin diseases using convolution neural network," *Microprocess. Microsy.*, vol. 76, pp. 103074, 2020, doi: https://doi.org/10.1016/j.micpro.2020.103074.
- [75] Y. Yang, et al., "Development and validation of two artificial intelligence models for diagnosing benign, pigmented facial skin lesions," *Skin Res. Technol.*, vol. n/a, no. n/a, 2020, doi: 10.1111/srt.12911.
- [76] T.J. Brinker, et al., "A convolutional neural network trained with dermoscopic images performed on par with 145 dermatologists in a clinical melanoma image classification task," *Eur. J. Cancer*, vol.111, pp. 148-154, 2019, doi: https://doi.org/10.1016/j.ejca.2019.02.005.
- [77] T. Brinker, et al., "Deep neural networks are superior to dermatologists in melanoma image classification," *Eur. J. Cancer*, vol. 119, pp. 11-17, 2019, doi: 10.1016/j.ejca.2019.05.023.
- [78] H.A. Haenssle, et al., "Man against machine: diagnostic performance of a deep learning convolutional neural network for dermoscopic melanoma recognition in comparison to 58 dermatologists," *Ann. Oncol.*, vol. 29, no. 8, pp. 1836-1842, 2018, doi: https://doi.org/10.1093/annonc/mdy166.
- [79] A. Hekler, et al., "Superior skin cancer classification by the combination of human and artificial intelligence," *Eur. J. Cancer*, vol. 120, pp. 114-121, 2019, doi: 10.1016/j.ejca.2019.07.019.
- [80] K. HosnyM. Kassem and M. Fouad, "Classification of skin lesions using transfer learning and augmentation with Alex-net," *PLoS One*, vol. 14, pp. e217293, 2019, doi: 10.1371/journal.pone.0217293.
- [81] S. Nasiri, J. Helsper, M. Jung and M. Fathi, "DePicT Melanoma Deep-CLASS: a deep convolutional neural networks approach to classify skin lesion images," *BMC Bioinformatics*, vol. 21, 2020, doi: 10.1186/s12859-020-3351-y.
- [82] Jordan, Yap, William, Yolland, Philipp and Tschandl, "Multimodal skin lesion classification using deep learning.," *Exp. Dermatol.*, 2018.
- [83] P. Tschandl, et al., "Expert-Level Diagnosis of Nonpigmented Skin Cancer by Combined Convolutional Neural Networks," *JAMA Dermatol.*, vol. 155, no. 1, pp. 58-65, 2019, doi: 10.1001/jamadermatol.2018.4378.
- [84] P. Tschandl, et al., "Human-computer collaboration for skin cancer recognition," *Nat. Med.*, vol. 26, no. 8, pp. 1229-1234, 2020, doi: 10.1038/s41591-020-0942-0.
- [85] M. Al-masniD. Kim and T. Kim, "Multiple Skin Lesions Diagnostics via Integrated Deep Convolutional Networks for Segmentation and Classification," *Comput. Meth. Prog. Bio.*, vol. 190, pp. 105351, 2020, doi: 10.1016/j.cmpb.2020.105351.
- [86] A. KrizhevskyI. Sutskever and G. Hinton, "ImageNet Classification with Deep Convolutional Neural Networks," *Neural Information Processing Systems*, vol. 25, 2012, doi: 10.1145/3065386.
- [87] K. Simonyan and A. Zisserman, "Very Deep Convolutional Networks for Large-Scale Image Recognition," *arXiv 1409.1556*, 2014.
- [88] C. Szegedy, et al., "Going Deeper with Convolutions.," 2014.
- [89] K. He, X. Zhang, S. Ren and J. Sun, "Deep Residual Learning for Image Recognition.," 2016, pp. 770-778.
- [90] G. HuangZ. Liu and K. Weinberger, "Densely Connected Convolutional Networks.," pp. 12, 2016.
- [91] H. J., S. L., A. S., S. G. and W. E., "Squeeze-and-Excitation Networks," *IEEE T. Pattern Anal.*, vol. 42, no. 8, pp. 2011-2023, 2020, doi: 10.1109/TPAMI.2019.2913372.
- [92] F.N. Iandola, S. Han, M.W. Moskewicz, K. Ashraf, W.J. Dally and K. Keutzer, SqueezeNet: AlexNet-level accuracy with 50x fewer parameters and <0.5MB model size,ed., Editor ed., 2016, pp. 1602-7360.
- [93] "Keras Documentation." [Online]. Available: https://keras.io/applications/. [Accessed: 12-Feb-2020].
- [94] M. Abadi, et al., "TensorFlow: Large-Scale Machine Learning on Heterogeneous Distributed Systems.," 2016.
- [95] A. Vedaldi and K. Lenc, "MatConvNet - Convolutional Neural Networks for MATLAB.," 2014.
- [96] Paszke, A., et al, "Automatic differentiation in pytorch," In: NIPS Workshop Autodiff, MA , USA ,2017.
- [97] "Dermnet-Skin Disease Atlas," [Online]. Available: http://www.dermnet.com/. [Accessed: 13-Dec-2016].
- [98] S.S. S. and P.N. S., Blood cell segmentation from microscopic blood images, *Proc. 2015 International Conference on*

Information Processing (ICIP), 2015, pp. 502-505.

- [99] Y. Y., X. N., Y.C. N. and D. X., "A Decentralized and Adaptive Flocking Algorithm for Autonomous Mobile Robots, Proc. 2008 The 3rd International Conference on Grid and Pervasive Computing - Workshops, 2008, pp. 262-268.
- [100] Y.S. VangZ. Chen and X. Xie, Deep Learning Framework for Multi-class Breast Cancer Histology Image Classification, Springer International Publishing, 2018, pp. 914-922.
- [101] T. Su, S. Mu, A. Shi, Z. Cao and M. Dong, "A CNN-LSVM MODEL FOR IMBALANCED IMAGES IDENTIFICATION OF WHEAT LEAF," Neural Netw. World, vol. 29, pp. 345-361, 2019, doi: 10.14311/NNW.2019.29.021.
- [102] D. BardouK. Zhang and S. Ahmad, "Classification of Breast Cancer Based on Histology Images Using Convolutional Neural Networks," IEEE Access, vol. PP, pp. 1, 2018, doi: 10.1109/ACCESS.2018.2831280.
- [103] Z. Hameed, S. Zahia, B. Zapirain, J.J. Aguirre Anda and A. Vanegas, "Breast Cancer Histopathology Image Classification Using an Ensemble of Deep Learning Models," 2020.
- [104] S. AsareF. You and O. Tetey, "Efficient, Ultra-facile Breast Cancer Histopathological Images Classification Approach Utilizing Deep Learning Optimizers," International Journal of Computer Applications, vol. 177, pp. 1-9, 2020, doi: 10.5120/ijca2020919875.
- [105] X. Wang, K. Mao, L. Wang, P. Yang, D. Lu and P. He, "An Appraisal of Lung Nodules Automatic Classification Algorithms for CT Images," Sensors-Basel, vol. 19, pp. 194, 2019, doi: 10.3390/s19010194.
- [106] H. R., L. K., X. N. and Z. Y., "Garment Image Retrieval on the Web with Ubiquitous Camera-Phone, Proc. 2008 IEEE Asia-Pacific Services Computing Conference, 2008, pp. 1584-1589.
- [107] Z. Xia, X. Wang, X. Sun, Q. Liu and N. Xiong, "Steganalysis of LSB matching using differences between nonadjacent pixels," Multimed. Tools Appl., vol. 75, no. 4, pp. 1947-1962, 2016, doi: 10.1007/s11042-014-2381-8.
- [108] J. Yang, Y. Jiao, N. Xiong and D.S. Park, "Fast Face Gender Recognition by Using Local Ternary Pattern and Extreme Learning Machine," KSII Transactions on Internet and Information Systems(TIIS), 2013.
- [109] W. Fang, Y. Li, H. Zhang, N. Xiong, J. Lai and A.V. Vasilakos, "On the throughput-energy tradeoff for data transmission between cloud and mobile devices," Inform. Sciences, 2014.
- [110] Y. ZhouDi Zhang and N. Xiong, "Post-Cloud Computing Paradigms: A Survey and Comparison," Tsinghua Science and Technology, vol. 22, no. 06, pp. 714-732, 2017.



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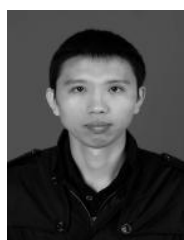
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