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# Recent advances in deep learning applied for skin cancer detection

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## Abstract

1 Skin cancer is a major public health problem around the world. Its early detection  
2 is very important to increase patient prognostics. However, the lack of qualified  
3 professionals and medical instruments is a significant issue in this field. For this  
4 reason, over the past few years, deep learning models applied to automated skin  
5 cancer detection has become a trend. In this paper, we present an overview of the  
6 recent advances reported in this field as well as a discussion about the challenges  
7 and opportunities for improvement in the current models. In addition, we also  
8 present some important aspects regarding the use of these models in smartphones  
9 and indicate future directions we believe the field will take.

## 10 1 Introduction

11 Skin cancer is the most common cancer worldwide. The World Health Organization (WHO) estimates  
12 that one in every three cancers diagnosed is a skin cancer [1]. In countries such as USA, Canada, and  
13 Australia, the number of people diagnosed with skin cancer has been increasing at a fairly constant  
14 rate over the past decades [2, 3, 4]. The deadliest type of skin cancer is the melanoma and its early  
15 detection greatly improves the prognosis of patients [5]. Nonetheless, there is a lack of medical  
16 instruments and qualified professionals to assist the population, especially in rural areas [6] and in  
17 economically emerging countries [7]. In this sense, over the past decades, different computer-aided  
18 diagnosis (CAD) systems have been proposed to tackle skin cancer detection. These systems are  
19 mostly based on traditional computer vision algorithms to extract various features, such as shape,  
20 color, and texture in order to feed a classifier [8, 9, 10, 11, 12]. Recently, machine learning techniques  
21 became a trend to handle this task. Deep learning models, in particular, Convolutional Neural  
22 Networks (CNN), have been achieving remarkable results in this field. Yu et al. [13] presented a  
23 very deep CNN and a set of schemes to learn under limited training data. Esteva et al. [14] used a  
24 pre-trained CNN model to train more than 120 thousand images and achieve a dermatologist-level  
25 diagnostic. Haenssle et al. [15] and Brinker et al. [16] presented CNN models that have shown  
26 competitive or outperformed the dermatologists. Other efforts have been made using deep learning to  
27 detect skin cancer, such as ensemble of models [17, 18], feature aggregation of different models [19],  
28 among others [20, 21, 22].

29 The recent progress achieved by the machine learning methodologies has been leading to the acces-  
30 sion of smartphone-based applications as a tool to handle the lack of dermatoscopes<sup>1</sup> available to  
31 dermatologists and general practitioners. According to the Ericsson mobile report [23], there are  
32 around 7.9 billion smartphones around the world. Thereby, a CAD system embedded in smartphones  
33 seems to be a low-cost approach to tackle this problem. However, even though this technology has  
34 the potential to be widely used in dermatology, there are important aspects that must be addressed

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<sup>1</sup>a medical instrument that allows the visualization of the subsurface structures of the skin revealing lesion details in colors and textures

35 such as target users and how to present the system predictions. In addition, there are important ethical  
36 concerns regarding patient confidentiality, informed consent, transparency of data ownership, and  
37 data privacy protection [24].

38 Since the impact of machine learning in dermatology will increase in the next few years, the goal  
39 of this paper is to critically review the latest advances in this field as well as to reflect about the  
40 challenges and aspects that need to improve. To this end, first, we present the main methodologies  
41 and results reported for the task. Then, we provide a discussion about general limitations regarding  
42 the machine learning methods and about the smartphone application issues. Lastly, we conclude this  
43 paper with our perspectives about this field for the future.

## 44 **2 Automated skin cancer detection**

### 45 **2.1 Recent advances**

46 The automated skin cancer detection is a challenging task due to the variability of skin lesions in the  
47 dermatology field. The recent advances reported for this task have been showing that deep learning  
48 is the most successful machine learning technique addressed to the problem. In this sense, the  
49 International Skin Imaging Collaboration (ISIC) has been playing an important role by maintaining  
50 the ISIC Archive, an international repository of dermoscopic skin images, which includes, skin  
51 diseases and skin cancer [25]. This archive have been proving data for different deep learning  
52 methodologies such as the ones proposed by Yu et al. [13], Codella et. al. [17], Haenssle et al. [15],  
53 and Briker et al. [16]. Currently, the ISIC archive contains 25,331 images for training and 8,238 test  
54 available for research purposes.

55 While developing approaches using the ISIC archive is important, it constrains its use for dermoscopic  
56 images. It means this system cannot be used, for example, in smartphone apps, except if the device  
57 has a special dermoscope attached to it. In this context, it is necessary to expand the models to also  
58 handle clinical images. However, for this case, there is no large public archive available like ISIC.  
59 Thereby, Han et al. [20] combined clinical images from 5 repositories, public and private, in order  
60 to differentiate benign and malignant cutaneous tumors. Nonetheless, a breakthrough work was  
61 presented by Esteva et al. [14] in which the authors collected 129,450 clinical images and trained  
62 a convolutional neural network (CNN) that achieved a dermatologist level in the benign/malignant  
63 identification. Unfortunately, this dataset is private and it is not available for the research community.

64 Another trend in this field is to adopt an ensemble of deep models instead of a single method. The  
65 main goal of this method is to make the predictions more effective and reliable. Codella et al. [17]  
66 employed an ensemble of different deep models, including deep residual networks and convolutional  
67 neural networks (CNNs), in order to detect malignant melanomas, the deadliest type of skin cancer.  
68 Similarly, Gessert et al. [26] adopted several types of CNN architectures in order to classify 7  
69 different types of skin diseases. In general, the ensemble of models have been achieving landmark  
70 results, in particular for ISIC archive [27].

71 In Table 1, we summarize all previously mentioned methods and their main contributions. It  
72 is important to note that all those models use only images to output their diagnostics. In fact,  
73 dermatologists do not trust only on the image screening, they also use the patient clinical information  
74 in order to provide a more reliable diagnostic. Pieces of information such as the patient's age,  
75 sex, ethnicity, if the lesion hurts or itches, among many others, are relevant clues towards a better  
76 prediction [28]. Thence, another breakthrough work has been recently proposed by Google Health  
77 researches in which they developed a deep learning system that is able to combine one or more images  
78 with the patient metadata in order to classify 26 skin conditions [29]. The addition of metadata  
79 provided a 4-5% consistent improvement in their model. They also report a result that is on par with  
80 U.S. board-certified dermatologists. Nonetheless, the authors indicate that is necessary to investigate  
81 prospectively the clinical impact of using this tool in actual clinical workflows.

82 To conclude this section, it is worth noting the recent work developed by Faes et al. [31]. In this  
83 work, the authors, who do not have any experience with algorithm development, used the Google  
84 Cloud AutoML to design several deep learning models for medical images, including skin cancer.  
85 They use a partition of the ISIC archive and reported a result that is comparable to other elementary  
86 classification tasks in this section. For one hand, it is a democratization of deep learning techniques.  
87 However, it also raise some questions about ethical principles when using these automated models.

Table 1: A summary of the recent deep learning models proposed to skin cancer detection

Ref.	Objective	Model	Main findings
[13]	Diagnose melanoma and non-melanoma using dermoscopic image	A two-stage framework composed of a fully convolutional residual network (FCRN) and a Deep Residual Network (DRN)	It was one of the first deep learning models applied to skin cancer detection and experimental results demonstrate the significant performance gains of the proposed framework compared to handcrafted feature models
[15]	Diagnose melanomas and nevus using dermoscopic images	Inception v4 CNN model	The authors compared the model performance to a group of 58 dermatologists using 100 images in the test set. The model AUC was greater than the average AUC of the dermatologists
[30]	Diagnose melanomas and nevus using dermoscopic images	ResNet50 CNN model	The authors compared the model to a group of 157 dermatologists using 100 images. The model outperformed 136 of them in terms of average specificity and sensitivity
[20]	Diagnose benign and malignant cutaneous tumors among 12 types of skin diseases using clinical images	ResNet-152 CNN model	The results achieved by the model was comparable to the performance of 16 dermatologists. The authors also affirm it is necessary to collect images with a broader range of ages and ethnicities in order to improve the model
[14]	Diagnose 757 types of skin diseases using clinical images	GoogleNet Inception v3 CNN model	The model achieved performance on par with 21 dermatologists considering the binary classification of the most common and the deadliest cases of skin cancer
[17]	Diagnose melanoma and non-melanoma using dermoscopic images	An ensemble composed of DRNs, CNNs and Fully CNNs	The ensemble was compared to the average of 8 dermatologists on a subset of 100 test images, and provided a higher accuracy and specificity, and an equivalent sensitivity
[26]	Diagnose 7 different types of skin diseases using dermoscopic images	An ensemble composed of ResNets, Densenets and Senets	The authors presented a new strategy based on a vast amount of unscaled image crops to generate final predictions. This approach outperforms most of the current models proposed for the ISIC archive

88 **2.2 Challenges and opportunities**

89 The models and results summarized in the previous section indicate the potential of CAD systems  
90 based on deep learning models applied for skin cancer detection. Nonetheless, there are several  
91 concerns that must be addressed in order to improve those systems. In this context, the goal of this  
92 section is to present a discussion about these concerns as well as indicate challenges and opportunities  
93 in this field.

94 **2.2.1 Dataset, bias and uncertainty**

95 It is known that to apply deep learning approaches it is necessary a large amount of data. However,  
96 collecting medical data, in particular from skin cancer, is a challenge task. Therefore, one of the main  
97 concerns of applying deep learning for this task is the lack of training data [20, 13]. As stated before,  
98 the ISIC archive is very important to tackle this issue. However, the number of samples available  
99 is still insufficient and very imbalanced among the classes. In order to tackle these issues, several  
100 approaches have been proposing such as transfer learning, data augmentation, up/downsampling and

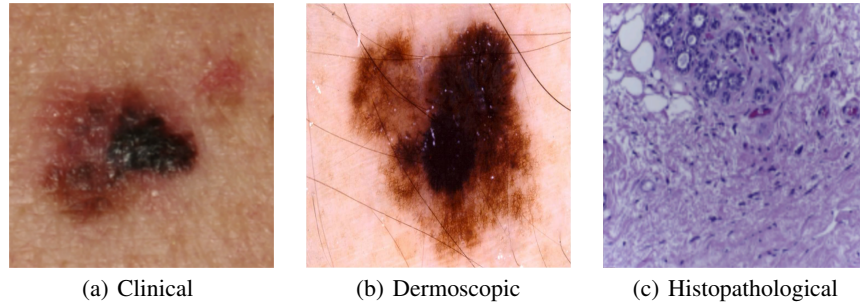


Figure 1: The difference between the clinical [20], dermoscopic [25] and histopathological [34] images of a skin cancer

101 weighted loss [32, 33]. Nevertheless, there is still room for improvement and approaches to learn  
 102 with limited data and based on weak supervision seem to be good choices to deal with it.

103 It is also important to note that the lack of open clinical data is a limiting factor for this task. As shown  
 104 in Figure 1, dermoscopic and clinical images present significant differences related to the level of  
 105 details available in each image. For this reason, reuse a model trained using only dermoscopic images  
 106 to predict clinical images is not possible. The previously described works that deal with clinical data  
 107 either combined some small datasets [20] or have access a private ones [14, 29]. In this sense, a  
 108 concerted effort is needed in order to build a clinical image archive such as ISIC. Furthermore, it is  
 109 important to include, along with the images, the patient information (metadata). As Liu et al. [29] has  
 110 shown, the use of metadata may help the deep learning systems deal with the lack of a large number  
 111 of images.

112 Another challenge regarding the skin cancer detection is to understand the current bias that distort the  
 113 performance of the models. Bissoto et al. [35] carried out a study that suggests spurious correlations  
 114 guiding the models. Moreover, some datasets, such as the used by Liu et al. [29], contain just a few  
 115 samples of skin types IV and V [28], which contribute to the bias. All these points must be considered  
 116 in order to deploy a model that is able to detect skin cancer for a more diverse group of people.

117 Beyond the bias, the patient metadata may contain uncertain information. Pieces of information  
 118 such as family cancer history, if the lesion is painful or itching, among many others, are surrounded  
 119 by uncertainty. Currently, the models do not take it into account, but it is an issue that should be  
 120 addressed in the future.

## 121 2.2.2 Presenting the predicted diagnosis

122 Currently, the most common way the models provide the diagnostic is choosing the label that produces  
 123 the highest probability. Some models also provide a ranking or a threshold for suspicious lesions  
 124 [20, 29]. However, how can a clinician interpret a low probability assigned to a melanoma? In fact,  
 125 they require more explanations than only the model’s predictions [36]. Instead of focus only on the  
 126 final accuracy, we need to improve how we present the results to the users. In this context, it is very  
 127 important to determine the target user. Dermatologists, general practitioners, medical students, or  
 128 even patients, have different levels of knowledge, hence, different needs.

129 In general, a clinician is interested in CAD systems that support their diagnostic by presenting insights  
 130 and visual explanations of the features used by a model in classification process [36]. They want  
 131 to know why the model is selecting such label. In this sense, we need to also focus on models that  
 132 are able to output not only the labels probabilities but the pattern analysis as well. Kawahara and  
 133 Hamarneh [37] proposed a model to detect dermoscopic feature classification, but it needs to be  
 134 improved and extended to clinical data. In Figure 2 is depicted an example of the 7-point checklist,  
 135 an algorithm based on pattern analysis commonly used to dermatologists to detect skin cancer [38].  
 136 As we can note, the expert is able to identify known patterns in the image in order to determine the  
 137 final diagnosis. While it is a very challenging task, it should be the ultimate goal of a CAD system  
 138 employed to skin cancer detection.

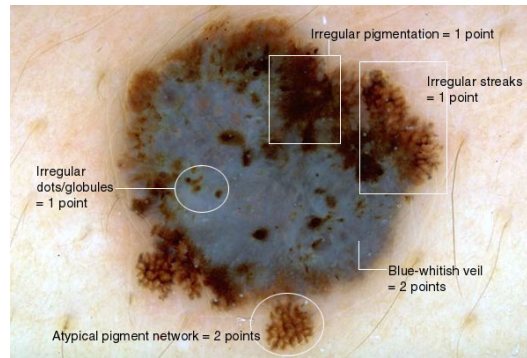


Figure 2: An example of the pattern analysis according to the 7-point checklist [39]

### 139 3 Skin cancer detection using smartphones

140 Due to the recent progress achieved by the CAD systems for skin cancer detection, currently, there  
 141 are several smartphone-based applications that aims to deal with this task. As stated before, embed  
 142 a skin cancer detection in a smartphone is a low-cost approach to tackle the lack of dermatoscopes  
 143 in remote places. It is clear that this technology has the potential to impact positively on people's  
 144 life. It may accelerate and help clinicians to provide a reliable diagnosis. However, developing such  
 145 technology is not only deploy the model in a smartphone, there are important ethical aspects that  
 146 must be addressed. The amount of those apps available for general users has drawn the attention of  
 147 different researchers that claim several issues regarding its use. Kassianos et al. [40] carried out a  
 148 study that identified 40 smartphone apps available to detect or prevent melanoma by nonspecialist  
 149 users. Half of them enabled patients to capture and store images of their skin lesions either for review  
 150 by a dermatologist or for self-monitoring. Chao et al. [24] conducted a similar study and concluded  
 151 that only a few apps have involved the input of dermatologists. In addition, most of them do not  
 152 provide a disclosure of authorship and credentials. As such, the application should make it clear  
 153 how it handles user data. It must ensure patient confidentiality as well as let them know what the  
 154 application does with their data after the processing. It may sound obvious, but as Chaos et al. [24]  
 155 have shown, researchers/developers are not respecting that.

156 Beyond the problems regarding to patient confidentiality and privacy, the lack of regulation for  
 157 those apps may result in harm to the patient or mislead them with an incorrect diagnostic. Let us  
 158 consider a hypothetical situation of a false negative for melanoma to a given user. It may delay  
 159 their treatment and, in the worst scenario, it may lead them to death. This is a serious problem that  
 160 we, machine learning researchers, need to confront. First of all, it is quite important the opinion of  
 161 dermatologists to improve the effectiveness of this technology. Then, those applications must be  
 162 exhaustively tested before deployed. Lastly, in our opinion, they should not be allowed to general  
 163 users before certification of a board of experts. To this end, it is necessary regulation and we need to  
 164 advocate for this.

165 To conclude, in addition to the challenges described in the previous section, in particular, the target  
 166 users and the way to present the results, there is an important technological issue about deploying deep  
 167 learning models in smartphones that should be discussed. The main use of this kind of applications  
 168 will be in remote places such as rural areas. In this sense, it is expected no internet access in those  
 169 places. However, the current apps do not process the data inside the smartphone, but in a server,  
 170 which demands internet. There are some fair reasons for this characteristic: the classification is  
 171 based on more than one model, i.e., an ensemble; the models are computationally expensive, which  
 172 demands better hardware than the ones usually found in smartphones; and the model's weights are  
 173 large files, which may not fit in the smartphone memory. In summary, this is an important aspect that  
 174 we could not find any discussion about it. In our opinion, this may lead to the development of lighter  
 175 models in order to deal with it.



Figure 3: An example of the VQA problem applied to skin cancer detection

#### 176 4 Final considerations and future directions

177 The recent advances in deep learning models for skin cancer detection have been showing the  
 178 potential of this technique to deal with this task. Nonetheless, there are some limitations and  
 179 important aspects that need to be addressed. In this paper, we presented a discussion about the  
 180 state-of-the-art approaches as well as the main challenges and opportunities related to the problem.  
 181 Despite the remarkable results reported, we indicated that there are rooms for improvement, especially  
 182 for the way the results should be presented. In this context, we believe that in the future this task  
 183 needs to be addressed as a variant of the visual and question answering (VQA) problem [41]. In  
 184 Figure 3 is illustrated an example of the VQA problem applied to skin cancer detection. The main  
 185 goal is to allow clinicians to make questions about the lesion in order to understand the predicted  
 186 diagnosis outputted by the model. This approach is in accordance with the interest of the clinicians,  
 187 which we described in Section 2.2.2. It is clear that addressing the skin cancer detection as a VQA  
 188 problem increases the difficulty of the problem. However, it is an efficient way toward the goal of  
 189 delivering a more useful tool for doctors.

190 Another aspect we believe will become a trend in the near future is the use of three types of skin cancer  
 191 images: clinical, dermoscopic and histopathological. As we can see in Figure 1, each image presents  
 192 different characteristics, which may help to correlate features in order to improve the predicted  
 193 diagnostic. In addition, the CAD systems will be able to act from the clinical diagnostic to the biopsy,  
 194 which makes it more desirable and useful. To conclude, regarding the deployment of deep models  
 195 in smartphones, as noticed earlier, the use of lighter models is necessary in order to make the apps  
 196 available in remote places. In this context, investigating better ways to improve the transfer learning  
 197 and considering not only the image but also the patient metadata are important aspect to be explored  
 198 in the future.

#### 199 References

- 200 [1] World Health Organization (WHO). How common is the skin cancer?, 2019. <https://www.who.int/uv/faq/skincancer/en/index1.html> [Last accessed: 15 September 2019].  
 201
- 202 [2] Cancer Council Australia. Understanding skin cancer - a guide for people with cancer, their families and friends, 2018. <https://www.cancer.org.au/about-cancer/types-of-cancer/skin-cancer.html> [Last accessed: 15 September 2019].  
 203  
 204
- 205 [3] Canadian Cancer Society’s Advisory Committee on Cancer Statistics. Canadian cancer statistics  
 206 2014 - special topic: Skin cancers, 2014. <https://www.cancer.ca/statistics>. [Last  
 207 accessed: 15 September 2019].
- 208 [4] Rebecca L Siegel, Kimberly D Miller, and Ahmedin Jemal. Cancer statistics, 2019. *CA: a  
 209 cancer journal for clinicians*, 69(1):7–34, 2019.
- 210 [5] Anthony F Jerant, Jennifer T Johnson, Catherine Demastes Sheridan, and Timothy J Caffrey.  
 211 Early detection and treatment of skin cancer. *American family physician*, 62(2), 2000.

- 212 [6] Hao Feng, Juliana Berk-Krauss, Paula W Feng, and Jennifer A Stein. Comparison of derma-  
213 tologist density between urban and rural counties in the united states. *JAMA dermatology*,  
214 154(11):1265–1271, 2018.
- 215 [7] Richard M Scheffler, Jenny X Liu, Yohannes Kinfu, and Mario R Dal Poz. Forecasting the  
216 global shortage of physicians: an economic-and needs-based approach. *Bulletin of the World*  
217 *Health Organization*, 86:516–523B, 2008.
- 218 [8] Fikret Ercal, Anurag Chawla, William V Stoecker, Hsi-Chieh Lee, and Randy H Moss. Neural  
219 network diagnosis of malignant melanoma from color images. *IEEE Transactions on Biomedical*  
220 *Engineering*, 41(9):837–845, 1994.
- 221 [9] M Emre Celebi, Hassan A Kingravi, Bakhtiyar Uddin, Hitoshi Iyatomi, Y Alp Aslandogan,  
222 William V Stoecker, and Randy H Moss. A methodological approach to the classification of  
223 dermoscopy images. *Computerized Medical Imaging and Graphics*, 31(6):362–373, 2007.
- 224 [10] Paul Wighton, Tim K Lee, Harvey Lui, David I McLean, and M Stella Atkins. Generalizing  
225 common tasks in automated skin lesion diagnosis. *IEEE Transactions on Information Technology*  
226 *in Biomedicine*, 15(4):622–629, 2011.
- 227 [11] Ilias Maglogiannis and Konstantinos K Delibasis. Enhancing classification accuracy utiliz-  
228 ing globules and dots features in digital dermoscopy. *Computer Methods and Programs in*  
229 *Biomedicine*, 118(2):124–133, 2015.
- 230 [12] Catarina Barata, M Emre Celebi, and Jorge S Marques. Improving dermoscopy image classifi-  
231 cation using color constancy. *IEEE Journal of Biomedical and Health Informatics*, 19(3):1146–  
232 1152, 2014.
- 233 [13] Lequan Yu, Hao Chen, Qi Dou, Jing Qin, and Pheng-Ann Heng. Automated melanoma  
234 recognition in dermoscopy images via very deep residual networks. *IEEE Transactions on*  
235 *Medical Imaging*, 36(4):994–1004, 2017.
- 236 [14] Andre Esteva, Brett Kuprel, Roberto A Novoa, Justin Ko, Susan M Swetter, Helen M Blau, and  
237 Sebastian Thrun. Dermatologist-level classification of skin cancer with deep neural networks.  
238 *Nature*, 542(7639):115, 2017.
- 239 [15] Holger A Haenssle, Christine Fink, R Schneiderbauer, Ferdinand Toberer, Timo Buhl, A Blum,  
240 A Kalloo, A Ben Hadj Hassen, L Thomas, A Enk, et al. Man against machine: diagnostic  
241 performance of a deep learning convolutional neural network for dermoscopic melanoma  
242 recognition in comparison to 58 dermatologists. *Annals of Oncology*, 29(8):1836–1842, 2018.
- 243 [16] Titus J Brinker, Achim Hekler, Alexander H Enk, Joachim Klode, Axel Hauschild, Carola Berk-  
244 ing, Bastian Schilling, Sebastian Haferkamp, Dirk Schadendorf, Tim Holland-Letz, et al. Deep  
245 learning outperformed 136 of 157 dermatologists in a head-to-head dermoscopic melanoma  
246 image classification task. *European Journal of Cancer*, 113:47–54, 2019.
- 247 [17] Noel CF Codella, Q-B Nguyen, Sharath Pankanti, DA Gutman, Brian Helba, AC Halpern, and  
248 John R Smith. Deep learning ensembles for melanoma recognition in dermoscopy images. *IBM*  
249 *Journal of Research and Development*, 61(4):5–1, 2017.
- 250 [18] Balazs Harangi. Skin lesion classification with ensembles of deep convolutional neural networks.  
251 *Journal of Biomedical Informatics*, 86:25–32, 2018.
- 252 [19] Zhen Yu, Xudong Jiang, Feng Zhou, Jing Qin, Dong Ni, Siping Chen, Baiying Lei, and Tianfu  
253 Wang. Melanoma recognition in dermoscopy images via aggregated deep convolutional features.  
254 *IEEE Transactions on Biomedical Engineering*, 66(4):1006–1016, 2019.
- 255 [20] Seung Seog Han, Myoung Shin Kim, Woohyung Lim, Gyeong Hun Park, Ilwoo Park, and  
256 Sung Eun Chang. Classification of the clinical images for benign and malignant cutaneous  
257 tumors using a deep learning algorithm. *Journal of Investigative Dermatology*, 138(7):1529–  
258 1538, 2018.

- 259 [21] Mohamed Attia, Mohamed Hossny, Saeid Nahavandi, and Anousha Yazdabadi. Skin melanoma  
260 segmentation using recurrent and convolutional neural networks. *IEEE 14th International*  
261 *Symposium on Biomedical Imaging*, pages 292–296, 2017.
- 262 [22] Nudrat Nida, Aun Irtaza, Ali Javed, Muhammad Haroon Yousaf, and Muhammad Tariq Mah-  
263 mood. Melanoma lesion detection and segmentation using deep region based convolutional  
264 neural network and fuzzy c-means clustering. *International journal of medical informatics*,  
265 124:37–48, 2019.
- 266 [23] Ericsson Inc. Ericsson mobility report, 2019. [https://www.ericsson.com/en/  
267 mobility-report/reports/june-2019](https://www.ericsson.com/en/mobility-report/reports/june-2019). [Last accessed: 15 September 2019].
- 268 [24] Elizabeth Chao, Chelsea K Meenan, and Laura K Ferris. Smartphone-based applications for  
269 skin monitoring and melanoma detection. *Dermatologic clinics*, 35(4):551–557, 2017.
- 270 [25] International Skin Imaging Collaboration (ISIC). Isic archive, 2019. [https://www.  
271 isic-archive.com](https://www.isic-archive.com). [Last accessed: 15 September 2019].
- 272 [26] Nils Gessert, Thilo Sentker, Frederic Madesta, Rüdiger Schmitz, Helge Kniep, Ivo Baltruschat,  
273 René Werner, and Alexander Schlaefer. Skin lesion diagnosis using ensembles, unscaled  
274 multi-crop evaluation and loss weighting. *arXiv preprint arXiv:1808.01694*, 2018.
- 275 [27] Fábio Perez, Sandra Avila, and Eduardo Valle. Solo or ensemble? choosing a cnn architecture  
276 for melanoma classification. In *Proceedings of the IEEE Conference on Computer Vision and*  
277 *Pattern Recognition Workshops*, pages 0–0, 2019.
- 278 [28] Klaus Wolff, Richard Allen Johnson, Arturo P. Saavedra, and Ellen K. Roh. *Fitzpatrick’s color*  
279 *atlas and synopsis of clinical dermatology*. McGraw-Hill Education, New York, USA, 8 edition,  
280 2017.
- 281 [29] Yuan Liu, Ayush Jain, Clara Eng, David H Way, Kang Lee, Peggy Bui, Kimberly Kanada,  
282 Guilherme de Oliveira Marinho, Jessica Gallegos, Sara Gabriele, et al. A deep learning system  
283 for differential diagnosis of skin diseases. *arXiv preprint arXiv:1909.05382*, 2019.
- 284 [30] Titus Josef Brinker, Achim Hekler, Jochen Sven Utikal, Niels Grabe, Dirk Schadendorf, Joachim  
285 Klode, Carola Berking, Theresa Steeb, Alexander H Enk, and Christof von Kalle. Skin cancer  
286 classification using convolutional neural networks: systematic review. *Journal of Medical*  
287 *Internet Research*, 20(10):e11936, 2018.
- 288 [31] Livia Faes, Siegfried K Wagner, Dun Jack Fu, Xiaoxuan Liu, Edward Korot, Joseph R Ledsam,  
289 Trevor Back, Reena Chopra, Nikolas Pontikos, Christoph Kern, et al. Automated deep learning  
290 design for medical image classification by health-care professionals with no coding experience:  
291 a feasibility study. *The Lancet Digital Health*, 1(5):e232–e242, 2019.
- 292 [32] Cristina Nader Vasconcelos and Bárbara Nader Vasconcelos. Experiments using deep learning  
293 for dermoscopy image analysis. *Pattern Recognition Letters*, 2017. . In press.
- 294 [33] Fábio Perez, Cristina Vasconcelos, Sandra Avila, and Eduardo Valle. Data augmentation  
295 for skin lesion analysis. In *OR 2.0 Context-Aware Operating Theaters, Computer Assisted*  
296 *Robotic Endoscopy, Clinical Image-Based Procedures, and Skin Image Analysis*, pages 303–311.  
297 Springer, 2018.
- 298 [34] Irene Fondón, Auxiliadora Sarmiento, Ana Isabel García, María Silvestre, Catarina Eloy,  
299 António Polónia, and Paulo Aguiar. Automatic classification of tissue malignancy for breast  
300 carcinoma diagnosis. *Computers in biology and medicine*, 96:41–51, 2018.
- 301 [35] Alceu Bissoto, Michel Fornaciali, Eduardo Valle, and Sandra Avila. (de) constructing bias on  
302 skin lesion datasets. In *Proceedings of the IEEE Conference on Computer Vision and Pattern*  
303 *Recognition Workshops*, pages 0–0, 2019.
- 304 [36] George A Zakhem, Catherine C Motosko, and Roger S Ho. How should artificial intelligence  
305 screen for skin cancer and deliver diagnostic predictions to patients? *JAMA dermatology*,  
306 154(12):1383–1384, 2018.



- 307 [37] Jeremy Kawahara and Ghassan Hamarneh. Fully convolutional neural networks to detect clinical  
308 dermoscopic features. *IEEE journal of biomedical and health informatics*, 23(2):578–585,  
309 2018.
- 310 [38] Giuseppe Argenziano, Gabriella Fabbrocini, Paolo Carli, Vincenzo De Giorgi, Elena Sammarco,  
311 and Mario Delfino. Epiluminescence microscopy for the diagnosis of doubtful melanocytic skin  
312 lesions: comparison of the ABCD rule of dermoscopy and a new 7-point checklist based on  
313 pattern analysis. *Archives of Dermatology*, 134(12):1563–1570, 1998.
- 314 [39] Dermoscopy. 7-point checklist, 2019. <http://www.dermoscopy.org/consensus/2d.asp>  
315 [Last accessed: 17 September 2019].
- 316 [40] AP Kassianos, JD Emery, P Murchie, and Fiona Mary Walter. Smartphone applications for  
317 melanoma detection by community, patient and generalist clinician users: a review. *British*  
318 *Journal of Dermatology*, 172(6):1507–1518, 2015.
- 319 [41] Stanislaw Antol, Aishwarya Agrawal, Jiasen Lu, Margaret Mitchell, Dhruv Batra,  
320 C Lawrence Zitnick, and Devi Parikh. Vqa: Visual question answering. In *Proceedings*  
321 *of the IEEE international conference on computer vision*, pages 2425–2433, 2015.