



合肥

VALSE 2019

April 11 – 14, 2019

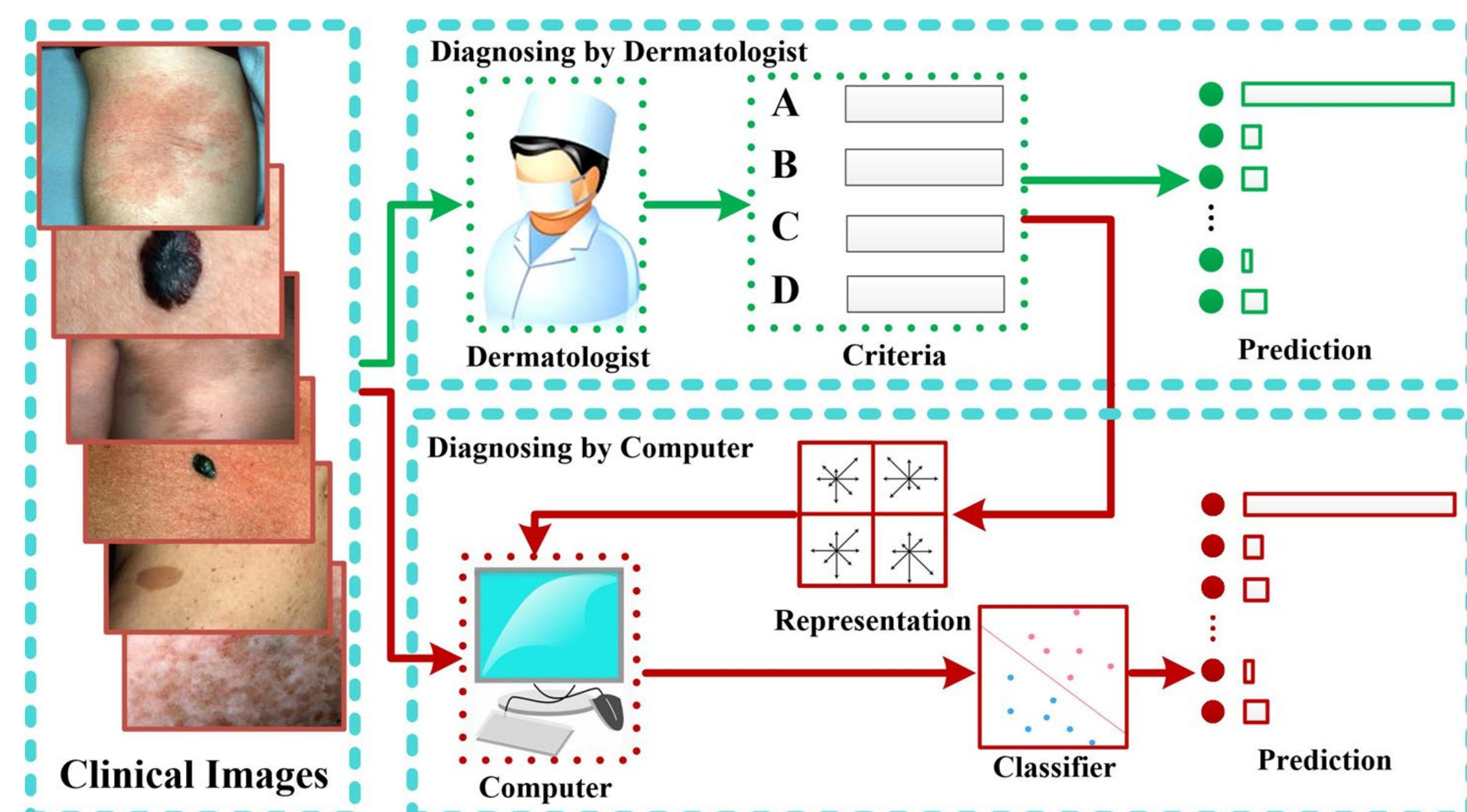
Abstract

We proposed medical representations inspired by the dermatological criteria for diagnosing clinical skin lesions. The dermatological criteria are highly correlated with measurable visual components, which cover three aspects: structure, color and shape of each skin disease. Major improvements includes:

- the interpretable representation for effectively capturing the manifestations of skin lesions;
- the integrated diagnostics of the diagnosis system for accurate recognizing.

The final performance on clinical images with 198 categories of skin disease is comparable with dermatologists

Motivation



For common skin disease (the **left** box):

- (1) dermatologists make a diagnosis by observing the appearance of the lesions (the top right box).
- (2) The designed skin disease recognition system (the bottom right box) based on clinical images and dermatological criteria has two major steps: firstly, the medical information observed by the doctors during diagnosis is exploited. Then, measurable medical representations for skin lesions are designed for diagnosis.

Accordingly, the designed representations for skin lesions relate to three aspects, i.e., **structure**, **color** and **shape** of the lesion.

Skin Lesion Diagnosis using Representations

Inspired by Dermatologist

Email: xpwu95@163.com Website: cv.nankai.edu.cn

Jufeng Yang¹, Xiaoxiao Sun¹, Jie Liang¹, Paul L. Rosin²

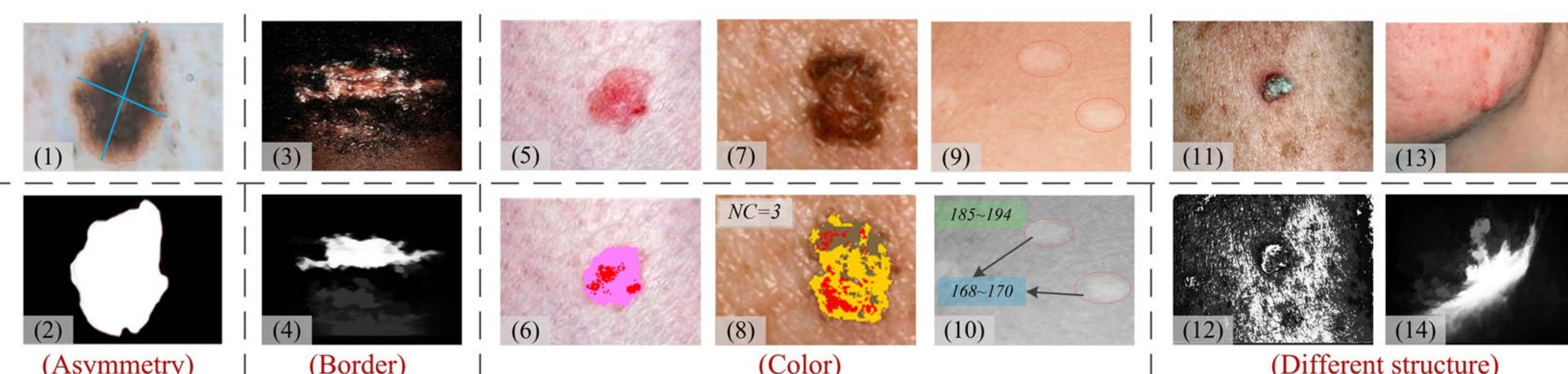
¹Nankai University, ²Cardiff University

This paper has been accepted by CVPR 2018

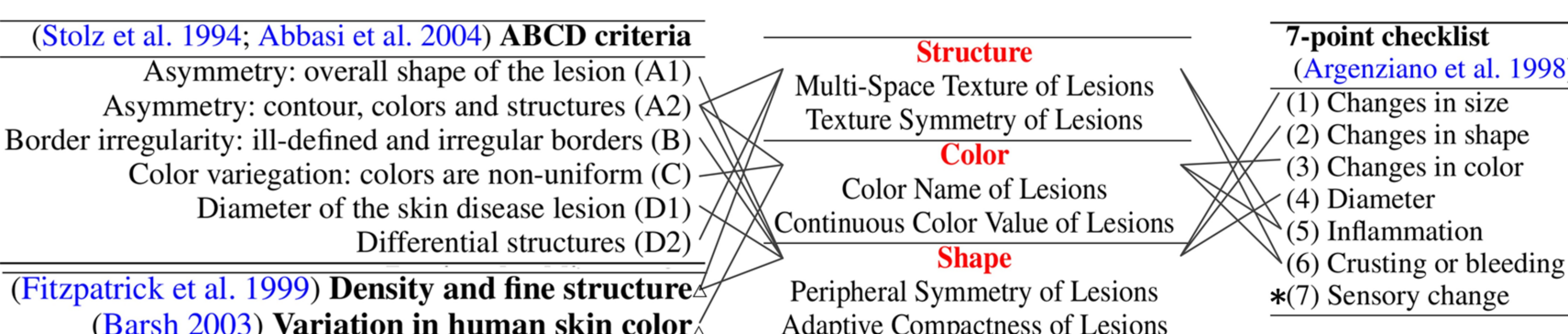


Criteria of Skin Disease

ABCD[1] criteria: (A)symmetry: the property on shape, color, contour and structures; (B)order; (C)olor; (D)ifferent structure (Diameter): pustule, wrinkle, inflammation and so on:



The mapping to visual components from dermatological criteria: **ABCD**[1-2], **7-point**[3], and the **two specialized criteria**[4-5]:



Medical Representations

Structure Representation:

(1) Multi-Space Texture of Lesion (MST-L): $MST(x) = [G_i(x)]_{i=1}^K$
($G_i(x)$ is the set of the texture features extracted from the i -th color channels and K denotes the number of spaces.)

(2) Texture Symmetry of Lesion (TS-L):

$$TS_i(x) = [G_i(L(x)_1), G_i(L(x)_2), S_i(x)]_{i=1}^K$$

($S_i(x) = \{g_{ij}^1 - g_{ij}^2\}_{j=1}^d$, d is the dimension of the extracted features, g_{ij}^1 and g_{ij}^2 are the j -th entry of $G_i(L(x)_1)$ and $G_i(L(x)_2)$).

The lesion region detected by MBD+ is divided into two parts $L(x)_1$ and $L(x)_2$ along principal axes.)

Color Representation:

(1) Color Name of Lesion (CN-L): $CN(x) = \arg\max_{c_l} [p(c_l|c)]_{l=1}^M$

($[p(c_l|c)]_{l=1}^M \propto \sum_{i=1}^N p(c_l|c_i) g^\sigma(|c_i - c|_{Lab})$ where c denotes the original value of the color bin, c_l is $L * a * b$ -value for c , $N = 389$ is the total number of the color bins and C is the set of basic colors used.

(2) Continuous Color Values of Lesion (CCV-L): $CCV(c) \propto p(C, c) \times \theta(c)$

where $p(C, c)$ indicates the probability of mapping the color bin c into its nearest color name C . The weighting value of the pixel $\theta(c) = \sum_{i=1}^n n(c) u(c)$. $n(c)$ is the frequency of the corresponding color in the image, $u(c)$ is the color value of c in the RGB space.

Shape Representation:

(1) Peripheral Symmetry of Lesion (PS-L): $PS(x) = F(A(L(x)^1), A(L(x)^2))$

(2) Adaptive Compactness of Lesion (AC-L):

$$AL = \sum_{z \in L(x)} p(C|c, z), Com = \frac{4\pi AL}{p^2},$$

where z denotes the pixel in the lesion $L(x)$. $p(C|c, z)$ is the probability of mapping color to a specific color category in the Color Name feature. $A(\cdot)$ denotes the extracted feature of lesions and $F(\cdot, \cdot)$ denotes the concatenation function applied to the two features.

Experiment Results

	Components	#	Features	Dimension	KNN		SVM		RF	
					ACC	SE	ACC	SE	ACC	SE
Baseline	Texture	1	SIFT	21000	20.35	19.17	25.55	24.75	21.42	21.25
		2	HOG	12400	19.14	17.85	17.62	14.45	10.54	10.66
		3	LBP	23200	15.13	14.80	18.89	14.69	14.61	13.24
		4	BRIEF	19200	16.74	15.62	12.21	8.39	15.67	15.03
		5	SURF	38400	17.47	16.50	31.17	25.35	27.34	26.52
		6	Wavelet	256	15.94	15.52	14.82	12.73	13.37	14.02
	Color	7	ORB	19200	20.53	21.44	23.21	22.94	18.86	17.46
		8	CH	256	12.33	12.58	4.19	4.41	18.77	16.81
		9	CN	21000	20.02	20.10	20.23	21.62	27.64	28.73
	Border	10	ColorSIFT	21000	21.29	19.62	22.51	21.43	28.49	27.24
		11	GIST	512	21.93	21.52	16.49	17.19	15.01	12.33
		12	Gabor	4000	13.67	13.00	10.15	8.62	13.73	12.43
		13	Prewitt	900	12.55	13.14	11.91	10.76	11.27	10.87
		14	Sobel	10000	12.27	12.03	10.42	10.18	13.46	12.46
Ours	Integration	15	Canny	10000	15.22	17.16	13.91	14.51	16.46	15.20
		16	1&10&11	2500	47.36	47.23	46.84	47.24	48.06	46.73
	Structure	17	MST-L	21000	44.99	45.62	48.06	46.38	43.23	42.73
		18	TS-L	21000	47.30	47.80	48.94	47.21	43.92	43.07
	Color	19	CN-L	21000	42.50	43.24	38.91	39.78	44.59	46.21
		20	CCV-L	21000	42.80	43.97	40.13	39.22	45.32	45.70
	Shape	21	PS-L	10000	30.04	30.47	38.58	38.29	38.94	36.87
		22	AC-L	10000	31.50	29.75	39.73	38.92	37.61	35.42
Ours	Integration	23	18&20&22	3000	57.62	56.41	56.47	53.15	57.81	56.65

GT:	5	GT:	46	GT:	31	GT:	1	GT:	15	GT:	15	GT:	92	GT:	56
General D:	5	General D:	67	General D:	68	General D:	155	General D:	35	General D:	44	General D:	81	General D:	167
Expert:	5	Expert:	46	Expert:	55	Expert:	1	Expert:	35	Expert:	15	Expert:	92	Expert:	56
Rank1:	5	Rank1:	46	Rank1:	68	Rank1:	155	Rank1:	35	Rank1:	44	Rank1:	92	Rank1:	56
Rank2:	119	Rank2:	39	Rank2:	178	Rank2:	87	Rank2:	115	Rank2:	124	Rank2:	27	Rank2:	176
Rank3:	49	Rank3:	67	Rank3:	31	Rank3:	1	Rank3:	15	Rank3:	26	Rank3:	115	Rank3:	67

	Method	ACC		SE
		ACC	SE	
Deep features [33]	CaffeNet	42.31	41.57	
	CaffeNet + ft	46.69	45.18	
	VGGNet	37.91	37.25	
	VGGNet + ft	50.27	48.25	
	GoogleNet	35.33	35.21	
	GoogleNet + ft	46.48	45.86	
	ResNet	48.78	47.62	
Doctors	ResNet + ft	53.35	51.24	
	General D	49.00	47.50	
	Junior D	52.00	53.40	
	Expert	83.29	85.00	
Ours		56.47	53.15	

'General D' is general doctor who does not focus on one specific kind of disease.

'Junior D' is junior dermatologist.

'Expert' is an expert for diagnosing skin lesions.

Dataset & Demo



- [1] N. R. Abbasi, H. M. Shaw, D. S. Rigel, R. J. Friedman, W. H. Mc Carthy, I. Osman, A. W. Kopf, and D. Polsky. Early diagnosis of cutaneous melanoma: revisiting the ABCD criteria. The Journal of the American Medical Association, 292(22), 2004.
- [2] W. Stolz, A. Riemann, A. Cognetta, L. Pillet, W. Abmayr, D. Holzel, P. Bilek, F. Nachbar, and M. Landthaler. ABCD rule of dermatoscopy- a new practical method for early recognition of malignant-melanoma. European Journal of Dermatology, 4(7), 1994.
- [3] G. Argenziano, G. Fabbrocini, P. Carli, V. De Giorgi, E. Sammarco, and M. Delfino. Epiluminescence microscopy for the diagnosis of doubtful melanocytic skin lesions: comparison of the ABCD rule of dermatoscopy and a new 7-point checklist based on pattern analysis. Archives of Dermatology, 134(12), 1998.
- [4] T. Fitzpatrick, J. Bernhard, T. Cropley, et al. The structure of skin lesions and fundamentals of diagnosis. Dermatology in General Medicine, 5, 1999
- [5] G. S. Barsh. What controls variation in human skin color? PLoS Biology, 1(1), 2003.