

Available Online at http://www.recentscientific.com

CODEN: IJRSFP (USA)

International Journal of Recent Scientific Research Vol. 10, Issue, 07(J), pp. 34000-34003, July, 2019 International Journal of Recent Scientific Re*r*earch

DOI: 10.24327/IJRSR

Research Article

A STUDY OF ROLE OF DERMOSCOPY IN DISEASE ACTIVITY OF VITILIGO PATIENTS

Ambresh S Badad., Anusha Chowdry* and Ashok Hogade

Department of DVL, MRMC Kalaburagi

DOI: http://dx.doi.org/10.24327/ijrsr.2019.1007.3797

ARTICLE INFO ABSTRACT

Article History: Received 13th April, 2019 Received in revised form 11th May, 2019 Accepted 8th June, 2019 Published online 28th July, 2019

Key Words:

Role of Dermoscopy In Disease Activity

Background: Vitiligo, an autoimmune disorder of pigmentation characterized by loss of functional melanocytes and melanin in epidermis. Dermascopy, a non invasive clinical technique aids in diagnosing early lesions of vitiligo and in assessing the stability of vitiligo.
Methods: An observational study of Dermascopy was conducted on 50 patients of vitiligo in Department of DVL, Basaveshwara Teaching and General Hospital, Kalaburagi.
Results: Out of 50 patients of vitiligo, majority belong to age group of 15-50 yrs with female preponderance. 120 lesions were analysed in 50 patients. 88 were clinically progressive and 32 were quiescent. Various patterns of vitiligo observed in our patients were vitiligo vulgaris 18, acral type 13, focal variant 9, mixed 8, segmental 2. On dermoscopy Marginal pigmentation was seen in 48%, perifollicular pigmentation in 32%, both patterns in 12% and none of the patterns were seen in 8%. Family history was seen in 9 patients. Dermoscopy features of disease activity in our study were Trichrome pattern, Petalloid pattern, Comet tail, Nebullous pattern, Polka dot. Leukotrichia was seen in 32 patients.

Conclusion: Dermoscopy aids in monitoring disease activity, treatment response and prognosis of disease. It helps in assessing the disease activity earlier than the clinical onset of disease instability.

Copyright © **Ambresh S Badad** *et al*, **2019**, this is an open-access article distributed under the terms of the Creative Commons Attribution License, which permits unrestricted use, distribution and reproduction in any medium, provided the original work is properly cited.

INTRODUCTION

Vitiligo is a common acquired dermatological disease occurring worldwide with an overall prevalence of 1%. However, its incidence ranges from 0.1 to > 8.8%¹⁻³ across the country and in different countries of the globe. Its characterized by white macules due to autoimmune destruction of melanocytes, with psychological impact owing to social burden. Normal skin has typical reticulate pigmentary pattern, which corresponds to pigmentation along rete ridges with pale areas corresponding to papillary dermis.⁴ Dermascopy facilitates in the diagnosis of altered reticulate pigmentation in vitiligo 3 and to assess evolution of stage of disease and the response to treatment.

Dermascopy, a noninvasive method aids in appreciating subtle features invisible to naked eye.

METHODS

A Prospective study was conducted in department of Dermatology, Basaveshwara Teaching and General Hospital, Kalaburagi from November 2018 to April 2019. All patients with vitiligo were included and patients with other hypopigmentory and depigmentary causes were excluded. Informed consent was taken from the patients, following which they were included in the study. A complete history regarding, duration of the disease, family history, history of Koebner's phenomenon, and history of associated other autoimmune disorders was taken. All patients were examined and were classified into stable and unstable vitiligo. Investigations were performed in patients when required and dermoscopic evaluation was done.

Different dermascopic parameters noted in our patients are:

Reticulate pigmentation Perifollicular pigmentation Perilesional pigmentation Absent pigment network Marginal pigmentation Trichrome Petalloid pattern Comet tail Nebullous pattern Polka dot Leucotrichia

RESULTS

Out of 50 patients of vitiligo, majority belong to age group of 15-50yrs with female preponderance. 120 lesions were

analysed in 50 patients. 88 were clinically progressive and 32 were quiescent. Various patterns of vitiligo observed in our patients were vitiligo vulgaris 18, acral type 13, focal variant 9, mixed 8, segmental 2.

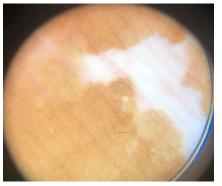


Figure 1 Amoeboid pattern

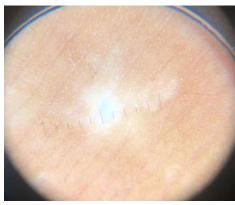


Figure 2 Nebulous pattern

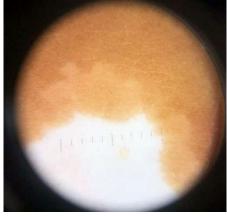


Figure 3 Trichrome pattern

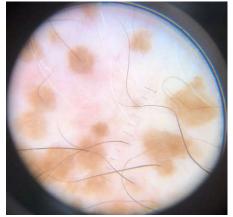


Figure 4 Perifollicular pigmentation



Figure 5 Telangiectasia

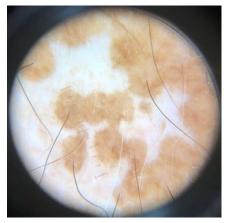


Figure 6 Leucotrichia

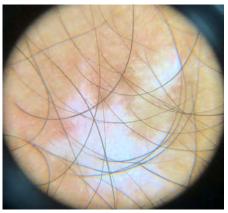


Figure 7 Starburst pattern

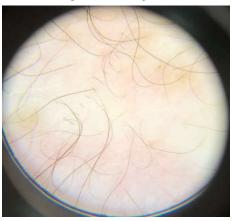


Figure 8 Absent pigment network



Figure 9 Reduced pigment network

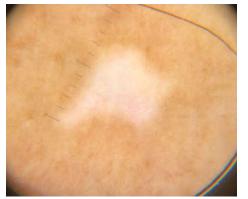


Figure 10 Comet tail pattern

On dermoscopy Marginal pigmentation was seen in 48%, perifollicular pigmentation in 32%, both patterns in 12% and none of the patterns were seen in 8%. Family history was seen in 9 patients. In our study vitiligo was seen in association with hypothyroidism in 6 patients and insulin dependent diabetes in 2 patients. Dermoscopy features of disease activity in our study were Trichrome pattern, Petalloid pattern, Comet tail, Nebullous pattern, Polka dot. Unstable patches were 74 and stable were 46. Leukotrichia was seen in 32 patients.

Additional signs seen in patients on treatment were erythema in 18 patients, telangiectasia in 20 patients, atrophy in 12 patients.

DISCUSSION

Vitiligo, acquired disorder with depigmented macules clinically and absence of functional melanocytes in epidermiss histologically. Evolving lesions of vitiligo which are difficult to be distinguished from other hypopigentory and depigmentary disorders can be diagnosed using dermoscopy and it also aids in diagnosing trichrome vitiligo, blue vitiligo.⁵⁻⁹.

Out of 50 patients, 29 were females and 21 were males. There is a preponderance of females in most series based on outpatient attendances, but the frequency in the population is probably the same in both sexes 10 .

A positive family history was seen in 9 patients.Different studies have reported association ranging from 6.25 % to 30%.^{11,12}.Vitiligo has polygenic or autosomal dominant inheritance pattern with incomplete penetrance and variable expression.¹³⁻¹⁵

Vitiligo vulgaris 26 was the most common type observed in our study followed by acral vitiligo 15, mucosal type in 3, focal in

5, segmental in 1, which is similar to reports of koranne *et al* ¹⁶ and sarin *et al* ¹⁷. Koebnerisation was seen in 11 patients in our study and nail involvement in 11 patients.

In our study vitiligo was seen in association with hypothyroidism in 6 patients, but it was reported as 12% by Gopal *et al* ¹⁸ and 1-7% in insulin dependent diabetes¹⁹, although it was seen in 2 patients in our study.

On dermoscopy Marginal pigmentation was seen in 48%, perifollicular pigmentation in 32%, both patterns in 12% and none of the patterns were seen in 8%. Study by Thatte and Khopkar showed 6.7% and 3.3% of patients with perifollicular and marginal pigmentation respectively. Family history was seen in 9 patients. Leukotrichia was seen in 32 pateinets. Its presence enhances diagnostic accuracy and signifies poorer prognosis mainly in segmental vitiligo.^{20,21} Dermoscopic findings associated with stability and repigmentation of vitiligo include marginal and perifollicular hyperpigmentation, reticular pigmentation and marginal reticular pigmentation. In our study, we noted reduced pigmentary network, absent pigmentary network in the evolving vitiligo lesions. Marginal hyperpigmentation was the most common pattern noted in our patients with stable vitiligo and trichrome pattern in unstable vitiligo patients.

CONCLUSION

Dermascopy helps in diagnosing evolving lesion of vitiligo, to assess disease activity, response to treatment and aids in prognosis of disease.

Declarations

Funding: No funding sources Conflict of interest: None declared Ethical approval: Approved

References

- 1. Shwartz RA, Janniger CK. Vitiligo. Cutis 1997; 60:239-44.
- 2. Hann SK, Park YK, Chun WH. Clinical feature of vitiligo. Clin Dermatol 1997;15:891-7
- 3. Alkhateeb A, Fain PR, Thody A, Bennett DC, Spritz RA. Epidemiology of vitiligo and associated autoimmune diseases in caucasian probands and their families. Pigment Cell Res 2003;16:208-14
- Haldar SS, Nischal KC, Khopkar US. Dermoscopy: Applications and patterns in Diseases of Brown skin. In: Khopkar U, editor. Dermoscopy and Trichoscopy in Diseases of Brown skin: Atlas and Short Text. New Delhi, India: Jaypee Brothers Medical publishers; 2012. p.16
- Bambroo M, Pande S, Khopkar US. Dermascopy in differentiation of idiopathic guttate hypomelanosis in: Khopkar U, editor. Dermoscopy and trichoscopy in diseases of brown skin:atlas and short text. New Delhi, India: Jaypee Brothers Medical Publishers; 2012.pp. 112-3
- 6. Gutte R, Khopkar US. Dermoscopy: differentiating evolving vitiligo from a hypopigmented patch of leprosy. In: Khopkar U, editor. Dermoscopy and trichoscopy in diseases of the brown skin: atlas and short

text. New Delhi, India: Jaypee Brothers Medical Publishers; 2012. pp. 112-3.

- Errichetti E, Stinco G. Dermoscopy in general dermatology: a practical overview. Dermatol Ther (Heidelb) 2016; 6:471-507.
- 8. Chandrashekar L. Dermatoscopy of blue vitiligo. Clin Exp Dermatol 2009; 34:e125-6.
- 9. Di Chiacchio NG, Ferreira FR, de Alvarenga ML, Baran R. Nail trichrome vitiligo: case report and literature review. Br j Dermatol 2013; 168:668-9.
- 10. Howitz J, Brodthagen H, Schwartz M et al. Prevalence of vitiligo. Arch Dermatol 1977; 113: 47–52.
- Hu Z, Liu JB, Ma SS, Yang S, Zhang XJ. Profile of childhood vitiligo in China: An analysis of 541 patients. Pediatr Dermatol 2006; 23:114-6.
- 12. Prcic S, Djuran V, Mikov A, Mikov I. Vitiligo in children. Pediatr Dermatol 2007; 24:666.
- Bleehen SS, Ebling FJ, Champion RH. Disorders of skin color. In: Champion RH, Burton JL, Ebling FJ, editors. Text book of Dermatology. London: Blackwell scientific publications: 1992, p.1561-622.
- 14. Moscher DB, Fitzpatrick TB, Hori Y, Ortonne JP. Disorders of pigmentation. In: Fitzpatrick TB, Isen AZ,Wolff K, Freedberg LM, Austen KF, editors. Dermatology in general medicine. New York: Mc Graw Hill;1993.p,903
- How to cite this article:

Ambresh S Badad *et al.*2019, A Study of Role of Dermoscopy In Disease Activity of Vitiligo Patients. *Int J Recent Sci Res.* 10(07), pp. 34000-34003. DOI: http://dx.doi.org/10.24327/ijrsr.2019.1007.3797

- Bolognia JL, Pawelek JM. Biology of hypopigmentation. J Am Acad Dermatol 1998; 19:217-55.
- Koranne RV. Sehgal VN, Sachdeva KG. Clinical profile of vitiligo in North India. *Indian J Dermatol Venereol Leprol* 1986;52:81-2
- 17. Sarin RC, Kumar AS. A clinical study of vitiligo. *Indian J Dermatol Venereol Leprol* 1977; 43:300-14.
- Gopal K, Rama Rao GR, Kumar YH, Appa Rao MV, Vasudev PS. Vitiligo: A part of a systemic autoimmune process. *Indian J Dermatol Venereol Leprol* 2007; 73:162 -5.
- Huggins RH, Janusz CA, Schwartz RA. Vitiligo: A sign of systemic disease. *Indian J Dermatol Venereol Leprol* 2006;72:68-71
- 20. Lee DY, Park JH, Lee JH, Yang JM, Lee ES.Is segmental vitiligo always associated with leukotrichia? Examination with digital portable microscope. *Int J Dermatol* 2009;48:1262
- Lee DY, Kim CR, Park JH, Lee JH. The incidence of leukotrichia in segmental vitiligo: Implication of poor response to medical treatment. *Int J Dermatol* 2011; 50:925-7.