

The spectrum of histomorphological features in psoriasis: a three years study

Arora D.¹, Mittal A.², Ahmad F.³, Dutta S.⁴, Awasthi S.⁵

¹Dr. Deepti Arora, Assistant Professor, ²Dr. Ankita Mittal, Assistant Professor, ³Dr. Faiyaz Ahmad, Associate Professor, ⁴Dr. Shyamoli Dutta, Professor, ⁵Dr. Seema Awasthi, Professor, all authors are affiliated with Department of Pathology, Teerthankar Mahaveer Medical College & Research Centre (TMMC & RC), Bagadpur, Moradabad, Uttar Pradesh, India.

Corresponding Author: Dr. Deepti Arora, Assistant Professor, Ankita Mittal, Assistant Professor, Department of Pathology, Teerthankar Mahaveer Medical College & Research Centre (TMMC & RC), Bagadpur, Moradabad, Uttar Pradesh, India., Email: deepti.a15@gmail.com

Abstract

Introduction: Psoriasis is a common relapsing chronic inflammatory dermatological disease associated with significant morbidity. Psoriasis is the prototype of a group of cutaneous disorders (psoriasiform dermatitides). Although histopathology is considered 'gold standard' for the diagnosis of psoriasis, at times even histopathological findings are confusing and inconclusive. **Aim of the study:** To study the histopathological findings at different stages of psoriasis and its subtypes. **Materials and Methods:** A total of 101 cases were studied for histopathological features. **Results:** Most of the cases were noted in the age group of 31 to 40 years (25.9%). There was a male predominance with male to female ratio of 2.7:1. Psoriasis vulgaris was the most common clinical type followed by Palmoplantar psoriasis and Erythrodermic psoriasis. Parakeratosis was the only consistent feature followed by dilated blood vessels/abnormal capillary pattern and elongation of rete ridges with thickening at lower ends. The other frequent features were acanthosis and agranulosis / hypogranulosis. **Conclusion:** Histopathology of psoriatic plaques exhibits heterogeneity and microscopic features will vary according to the stage of the disease. Though most of the predominant morphologic features of psoriasis are related to the epidermis, the presence of dilated blood vessels and/or abnormal capillary pattern is a constant finding in all stages of psoriasis and thus, can help in diagnosis of psoriatic lesion when in doubt.

Key words: Psoriasis, Parakeratosis, Acanthosis

Introduction

Psoriasis is a common relapsing chronic inflammatory dermatological disease associated with significant morbidity [1-3]. It is characterized by sharply demarcated, erythematous papules and plaques with abundant silvery white-scales [4]. Psoriasis is universal in occurrence affecting 1% to 3% of population worldwide [2,5]. There is paucity of data related to exact burden of psoriasis in India. However, in few separate hospital-based studies the incidence of psoriasis is reported as 0.44% - 2% and 2.3% [2,6].

Psoriasis is the prototype of a group of cutaneous disorders (psoriasiform dermatitides) that shows psoriasiform epidermal hyperplasia, defined as regular elongation of the rete ridges with preservation of the rete ridge-dermal papillae pattern [7,8]. Psoriasis has different clinical subtypes that may simulate various

other dermatological disorders. This presents a diagnostic challenge for the clinician and histopathological confirmation becomes mandatory [3]. The histologic diagnosis depends upon an aggregate of histologic criteria, some characteristic of psoriasis and others shared with other dermatoses [9]. The microscopic picture of psoriasis varies with the evolutionary stage of psoriatic lesion [10]. Although the histopathology is considered 'gold standard' for the diagnosis of psoriasis, at times even histopathological findings are confusing and inconclusive [3].

Aim of The Study- In this study, an attempt was made to establish profile of histopathological findings at different stages of psoriasis and its subtypes.

Materials and Methods

No ethical issues were involved in this study. This was a prospective; hospital-based, cross-sectional study and was carried out in the Department of Pathology of a

Manuscript received: 18th January 2019

Reviewed: 28th January 2019

Author Corrected: 6th February 2019

Accepted for Publication: 11th February 2019

Original Research Article

tertiary care medical institute in India for a period of three years. The study material constituted biopsy samples from patients of all age groups attending dermatology outpatient department (OPD) with the clinical diagnosis of psoriasis or with psoriasis as one of the differential diagnosis. Skin biopsies either as punch biopsy or incisional biopsy was done by the dermatologist as an out-patient procedure in the dermatology OPD.

The tissue specimen was immediately put in 10% buffered neutral formalin for fixation and was sent to the department of pathology. In the histopathology section, these specimens were fixed adequately and were submitted for routine histopathological processing. This was followed by cutting the sections at five-micron thickness and staining with routine hematoxylin and eosin stains. All the stained sections were observed

Results

Total of 101 biopsy samples from patients were evaluated for histological features of psoriasis. Most of the cases were noted in the age group of 31 to 40 years (25.9%) followed closely by age group of 41 to 50 years (23.8%). Only one case was noted in 0 to 10 years age group (0.9%). The youngest patient was 6 years old and oldest was 89 years old at the time of biopsy.

The study showed male predominance with male to female ratio of 2.7:1. (74 males and 27 females). Psoriasis vulgaris was the most common clinical type (89.3%) followed by Palmoplantar psoriasis (6.9%) and Erythrodermic psoriasis (2.9%) in the present study. Inverse psoriasis was the least common (0.9%) clinical type.

Among all histologic features observed, Parakeratosis was the only feature which was seen in all biopsy samples. Dilated blood vessels/abnormal capillary pattern (98.1%) and elongation of rete ridges with thickening at lower ends (98.1%) were second most common histologic feature after parakeratosis. The other frequent features were acanthosis (96.1%), pallor of upper layer of epidermis (88.1%), and agranulosis/hypogranulosis (83.1%).

Kogoj spongiform pustules were the least common histologic feature and were identified in only 9 (8.9%) out of 101 biopsy samples. The presence of spongiosis (10.8%) and exocytosis of neutrophils (20.7%) were also infrequent histopathological finding in biopsy samples of psoriatic patients. Histological features observed in the biopsy samples are summarized in table.1.

Table-1: Distribution of histologic features present in biopsy samples.

Histological features	No. of cases	Percentage (%)
Parakeratosis	101	100%
Hyperkeratosis	45	44.5%
Acanthosis (regular epidermal hyperplasia)	97	96.1%
Elongation of rete ridges with thickening at lower ends	99	98.1%
Suprapapillary thinning	42	41.5%
Agranulosis/ hypogranulosis	84	83.1%
Spongiosis	11	10.8%
Munro microabscess	30	29.7%
Kogoj spongiform pustule	9	8.9%
Exocytosis of neutrophils	21	20.7%

under light microscope and the histopathological findings were noted.

Inclusion criteria

1. Clinically suspected cases of psoriasis
2. Both genders and all age groups were included

Exclusion criteria

1. Inadequate biopsy samples (less than 4mm) and biopsies showing only epidermis or dermis on histologic examination were excluded.
 2. Skin biopsy done for cases other than psoriasis or suspected psoriasis
 3. Repeat skin biopsies within the study period
- No scoring system was used for this study. For statistical analysis, the percentages were done.

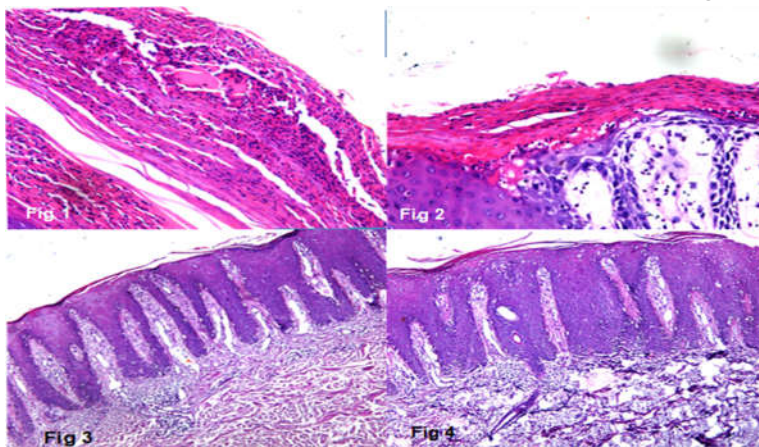


Fig 1: Munro micro abscesses (H & E Section, 10X)

Fig 2: Supra-papillary thinning with prominent blood vessels (H & E Section, 10X)

Fig 3: Shows elongation of rete ridges with thickening at lower ends (H & E Section, 10X)

Fig 4: Acanthotic epithelium (H & E Section, 10X)

Discussion

Psoriasis is a genetically determined erythematous-quamous dermatitis characterized by abnormal keratinocyte proliferation resulting in thickening of epidermis [8]. In its classic presentation, the disease comprises of well circumscribed reddish scaly papules and plaques typically on the elbow, knee and scalp, in addition to other cutaneous sites [11]. In this typical form, the diagnosis is straightforward based on clinical features alone and skin biopsy for histopathological examination is seldom required [8].

However, histologic analysis of skin biopsies can be helpful to confirm the diagnosis of psoriasis in its classic form and clinically atypical variants [11]. With this background, the present study was undertaken and biopsy samples of 101 psoriatic patients were systematically evaluated on histopathology for establishing the morphological features of psoriasis.

In our study, psoriasis vulgaris was the most common clinical type which accounted for 89.3% cases followed by palmoplantar psoriasis (6.9%) and erythrodermic psoriasis (2.9%). The inverse psoriasis was the least common (0.9%). Similar observations were found by other authors who reported maximum cases of psoriasis vulgaris ($\geq 85\%$) and almost similar number of cases of erythrodermic psoriasis (2-3%) and inverse psoriasis (1-2%) [12-15]. Thomas J et al reported higher incidence of palmoplantar psoriasis followed by palmoplantar psoriasis constituting 45% cases and 44% cases of psoriasis vulgaris [16]. Puri N et al found marginally higher cases of psoriasis vulgaris (32%) than guttate psoriasis (28%) followed by erythrodermic psoriasis (16%) and similar incidence of generalized pustular and palmoplantar psoriasis (12%) [17].

All the 101 biopsies were examined for the various morphological features described in literature for psoriasis in the present study. Parakeratosis was the consistent histopathological feature which was noted in all patients. Other authors in their respective studies have also described parakeratosis as the most consistent histopathological feature of psoriasis [8,11,15,18-20]. However, Mehta S et al and Puri N et al found parakeratosis in 52% and 56% cases respectively [3,17]. Hyperkeratosis was observed in 44.5% biopsies of psoriatic patients; the findings are comparable to the study by Puri N et al [17]. They observed hyperkeratosis in 64% cases. However, studies by Raghuvver C et al and Abdu NN et al reported hyperkeratosis in 89% and 88% cases respectively [14,21]. Bai et al, Chandanwale SS et al and Park JH et al found hyperkeratosis in all the biopsies of psoriasis patient [15,19,20].

In the present study, acanthosis (regular epidermal hyperplasia) was present in 96.1% of biopsies. Our results are in concordance with other studies in which acanthosis was observed in about 75% or more biopsies [3,14,17,19,22]. However, Park JH et al reported lesser prevalence of this feature (46.7%) as compared to other authors as well as the present study [20]. Elongation of rete ridges with thickening at the lower ends or 'club shaped' ridges was noticed in 98.1% biopsy samples. Our results are comparable with the studies conducted by Park JH et al and Kim BY et al [20,22]. Kim BY et al noticed elongation of rete ridges in 93% cases while Park JH et al reported thickening of lower ends in 100% biopsy samples. However few studies reported elongation of rete ridges in only 66-76% cases [14,17,21]. In a study conducted by Chandanwale SS et al they reported that slender long rete pegs favor the

Original Research Article

histopathological diagnosis of psoriasis which was not a consistent finding in our study [19]. In the present study, suprapapillary thinning was noted in 41.5% biopsy samples, findings similar to the studies conducted by Mehta S et al and Puri N et al [3,17]. This finding is in contrast to other studies which reported variable prevalence of this feature [15,20,22]. Ghasmeni et al showed thinning of suprapapillary plate to be significantly more common in psoriasis [23]. We have observed the feature of elongation of rete ridges with thickening at lower ends in almost all biopsies but the suprapapillary thinning was observed in only 41.5% cases. The suprapapillary epidermal thinning is a relative observation on comparison with markedly elongated rete ridges and therefore may not be observed in all cases of psoriasis.

Trozak explained the importance of club shaped rete ridges as characteristic of epidermis in psoriasis which helps in differentiating the disease from Psoriasiform dermatitis. He also included this feature in histological grading system for psoriasis developed by him [9]. Our findings also re-emphasize the club shaped rete ridges as characteristic of psoriasis even if not associated with suprapapillary thinning. In the present study, agranulosis/ hypogranulosis was noted in 83.1% biopsy samples. Similar results have been observed in previous studies which reported presence of agranulosis/ hypogranulosis in 68% - 100% cases. In contrast, a study conducted by Chandanwale SS et al found agranulosis in 32% cases [19]. The cause for absent granular layer or decreased/hypo granular layer is not known. However, it seems that this is probably related to epidermal hyperproliferation in earlier stage of disease, as later lesions of psoriasis may show intact granular layer [24,25].

In our study the spongiosis was present in 10.8% biopsies. Mehta S et al and Puri N et al found spongiosis in around 40% cases [3,17]. Bai S et al observed spongiosis in 91.7% [15]. Park JH et al noticed slight spongiosis in all the cases in their respective studies [20]. Thus, our results are in contrast to these studies. Munro microabscess was observed in 29.7% biopsies. Other studies observed Munro microabscess in approximately 50%-60% biopsies [3,14,15,19,21,22]. A study done by Park JH et al found this feature in almost all biopsies [20]. Kogoj spongiform pustules were seen in 8.9% biopsy samples. It was found that though Munro microabscess and Kogoj spongiform pustules are important diagnostic features of psoriasis, they are not always present on histopathology [1,9,18,26]. In the present study, dilated blood vessels/abnormal capillary pattern in dermal papillae were seen in 98.1% of biopsy samples. Our findings are in concordance with the

studies by Raghuveer C et al, Chandanwale et al and Abdu NN et al [14, 19,21] They reported dilated blood vessels in dermal papillae in 98% and 100% and 90% cases respectively. The studies by Mehta S et al and Puri N et al reported this feature in 80% and 54% cases respectively [3,17]. We have also observed that the dilated blood vessels/abnormal capillary pattern is seen in almost all cases of psoriasis indicating that in all stages of psoriasis, this is a consistent finding.

We observed exocytosis of neutrophils in 20.7% biopsy samples. Mehta S et al found this histologic feature in all biopsy samples, whereas, Puri N et al found exocytosis of neutrophils in 48% biopsies [3,17]. Our findings are in sharp contrast with the above studies. Griffin TD et al described exocytosis of neutrophils as a histopathological feature of acute psoriasis while chronic psoriatic plaques lack the presence of neutrophils [27]. The finding of neutrophil exocytosis appears to be a variable finding and may be limited to early or late stage. It may be seen as diagnostic clue or it may be of little help. We also did not find this finding in large number of biopsy specimens in our study.

Conclusions

Histopathology of psoriatic plaques though show some consistent features but will always exhibit heterogeneity and microscopic features will vary according to the stage of the disease. Most of the predominant morphologic features of psoriasis are related to the epidermis, with varying histopathological findings.

However, the presence of dilated blood vessels and/or abnormal capillary pattern is a constant finding in all stages of psoriasis. It is recommended to specifically look for these findings in cases of biopsies suspected of psoriasis as their presence can help in diagnosis of psoriatic lesions when in doubt.

Findings: Nil; **Conflict of Interest:** None initiated
Permission from IRB: Yes

References

1. Altman EM, Kamino H. Diagnosis: psoriasis or not? What are the clues? *Semin Cutan Med Surg.* 1999 Mar; 18 (1):25-35.
2. Dogra S, Yadav S. Psoriasis in India: prevalence and pattern. *Indian J Dermatol Venereol Leprol.* 2010 Nov-Dec; 76 (6):595-601. doi: 10.4103/0378-6323.72443.
3. Mehta S, Singal A, Singh N, et al. A study of clinic histopathological correlation in patients of psoriasis and psoriasiform dermatitis. *Indian J Dermatol Venereol Leprol.* 2009 Jan-Feb;75(1):100.

Original Research Article

4. Khalil FK, Keehn CA, Saeed S, et al. Verrucous psoriasis: a distinctive clinicopathologic variant of psoriasis. *Am J Dermatopathol.* 2005 Jun;27(3):204-7.
5. Linden KG, Weinstein GD. Psoriasis: current perspectives with an emphasis on treatment. *Am J Med.* 1999 Dec;107(6):595-605.
6. Kaur I, Handa S, Kumar B. Natural history of psoriasis: a study from the Indian subcontinent. *J Dermatol.* 1997 Apr;24(4):230-4.
7. Sehgal VN, Dogra S, Srivastava G, et al. Psoriasiform dermatoses. *Indian J Dermatol Venereol Leprol.* 2008 Mar-Apr;74(2):94-9.
8. Murphy M, Kerr P, Grant-Kels JM. The histopathologic spectrum of psoriasis. *Clin Dermatol.* 2007 Nov-Dec;25(6):524-8. DOI:10.1016/j.clindermatol.2007.08.005
9. Trozak DJ. Histologic grading system for psoriasis vulgaris. *Int J Dermatol.* 1994 May;33(5):380-1.
10. Micali G, Lacarrubba F, Musumeci ML, et al. Cutaneous vascular patterns in psoriasis. *Int J Dermatol.* 2010 Mar;49(3):249-56. doi: 10.1111/j.1365-4632.2009.04287.x.
11. Cribier BJ. Psoriasis under the microscope. *JEADV* 2006; 20(Suppl 2):3-9.
12. Asokan N, Prathap P, Ajithkumar K, et al. Pattern of psoriasis in a tertiary care teaching hospital in South India. *Indian J Dermatol.* 2011 Jan;56(1):118-9. doi: 10.4103/0019-5154.77575.
13. Golpour M, Hosseini SH, Khademloo M, et al. Depression and Anxiety Disorders among Patients with Psoriasis: A Hospital-Based Case-Control Study. *Dermatol Res Pract.* 2012;2012:381905. doi: 10.1155/2012/381905. Epub 2012 Jul 16.
14. Raghuvveer C, Shivanand DR, Rajashekar N. Clinico-histopathological Study of Psoriasis. *Int J Sci Stud* 2015;3(7):176-179.
15. Bai S, Sowmya S. Histopathologic diagnostic parameters of psoriasis; a clinicopathological study. *Int J Res Med Sci* 2016;4:1915-20.
16. Thomas J, Kumar NA, Manoharan D, Cynthia S, Prabu SKS, Ahmed NA. A study of comorbid conditions in psoriasis. *Journal of Pakistan Association of Dermatologists* 2009;19:200-2.
17. Puri N, Mahajan BB, Kaur S. Clinicohistopathological Correlation of Psoriasis in Acute Exacerbation. *2012;1:455. doi:10.4172/scientificreports.455*
18. De Rosa G, Mignogna C. The histopathology of psoriasis. *Reumatismo.* 2007;59 Suppl 1:46-8.
19. Chanadanwale SS, Panicker NK, Kulkarni SP, Shah KR, Kumar H, Sharma YK, et al. Morphometry analysis of psoriasis and psoriasiform dermatitis: A retrospective study of 50 cases. *Med J DY Patil Univ* 2015;8:43-7.
20. Park JH, Park YJ, Kim SK, et al. Histopathological Differential Diagnosis of Psoriasis and Seborrheic Dermatitis of the Scalp. *Ann Dermatol.* 2016 Aug;28(4): 427-32. doi: 10.5021/ad.2016.28.4.427. Epub 2016 Jul 26.
21. Abdu NN, Bugude G, Mallikarjun M, Deepadarshan K. Clinical and histopathological study of psoriasis. *Med Pulse International Journal of Medicine* 2017;4(2): 58-63.
22. Kim BY, Choi JW, Kim BR, et al. Histopathological findings are associated with the clinical types of psoriasis but not with the corresponding lesional psoriasis severity index. *Ann Dermatol.* 2015 Feb;27(1):26-31. doi: 10.5021/ad.2015.27.1.26. Epub 2015 Feb 3.
23. GhasemiBasir HR, Alirezaei P, Hamian Z, et al. Are quantitative histopathologic criteria capable of differentiating psoriasis from chronic dermatitis? *Clin CosmetInvestig Dermatol.* 2018 May 10;11:239-244. doi: 10.2147/CCID.S160697. eCollection 2018.
24. Barr RJ, Young EM Jr. Psoriasiform and related papulosquamous disorders. *J CutanPathol.* 1985 Oct; 12(5): 412-25.
26. Chowanec O, Jabłońska S, Beutner EH, et al. Earliest clinical and histological changes in psoriasis. *Dermatologica.* 1981;163(1):42-51.
26. Toussaint S, kamino H. Non-infectious erythematous papular and squamous diseases of the skin. In: Elder D, Elenitsas R, Jaworsky C, Johnson B, editors. *Lever's Histopathology of the skin.* 8th edition ed. Philadelphia: Lippincott-Raven; 1997. p. 156-63.
27. Griffin TD, Lattanand A, VanScott EJ. Clinical and histologic heterogeneity of psoriatic plaques. Therapeutic relevance. *ArchDermatol.* 1988Feb;124(2):216-20

How to cite this article?

Arora D, Mittal A, Ahmad F, Dutta S, Awasthi S. The spectrum of histomorphological features in psoriasis: a three years study. *Trop J Path Micro* 2019;5(2):58-62. doi:10.17511/jopm.2019.i02.02.