Review Article
The Significance of Scalp Involvement in Pemphigus: A Literature Review

Marta Sar-Pomian, Lidia Rudnicka, and Małgorzata Olszewska

Department of Dermatology, Medical University of Warsaw, Warsaw, Poland

Correspondence should be addressed to Małgorzata Olszewska; małgorzata.olszewska@wum.edu.pl

Received 14 November 2017; Accepted 18 February 2018; Published 25 March 2018

Academic Editor: Emiliano Antiga

Copyright © 2018 Marta Sar-Pomian et al. This is an open access article distributed under the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

1. Introduction

Pemphigus is a group of autoimmune bullous diseases, in which anti-desmoglein autoantibodies induce the loss of keratinocyte adhesion, leading to blister formation. Two main types of pemphigus may be distinguished: pemphigus vulgaris (PV) and pemphigus foliaceus (PF). Desmoglein 3 is considered to be the main autoantigen in pemphigus vulgaris. Additionally, anti-desmoglein 1 autoantibodies may be detected in more than half of patients with pemphigus vulgaris [1]. Desmoglein 1 is the main autoantigen of pemphigus foliaceus. Mucosal lesions with or without skin lesions are observed in pemphigus vulgaris, whereas in pemphigus foliaceus only cutaneous involvement is observed. The diagnosis of pemphigus is based on the clinical picture, direct and indirect immunofluorescence, and histopathology as well as anti-desmoglein 1 and 3 enzyme-linked immunoassay.

The frequency of scalp involvement in the course of pemphigus is estimated at 16–60% [2–4]. According to literature data, the scalp is the first location in 9–15% of patients with pemphigus [3, 4]. Several cases of alopecia in the course of pemphigus have been described [5]. The significance of the distribution of desmogleins in hair follicles for scalp involvement [6] and a potential use of direct immunofluorescence of plucked hairs are discussed in the literature [7]. The significance of scalp involvement for the course of pemphigus remains controversial.

The aim of this systematic review was to collect and critically analyze data concerning the types of alopecia occurring in pemphigus, the usability of direct immunofluorescence of plucked hairs as well as trichoscopy in diagnosing pemphigus, the histopathological picture of scalp involvement in pemphigus, and the responsiveness to treatment in patients with scalp involvement in the course of pemphigus.

2. Material and Methods

A systematic review of the literature was conducted in three electronic medical databases: PubMed, Embase, and Web of Science (till February 10, 2018). The search with the conjunction of search terms “pemphigus” AND “scalp OR hair”, “pemphigus” AND “alopecia”, “desmoglein” AND “hair” returned a total number of 1595 articles. Only studies conducted in humans and English-language articles were...
included. Non-full-text publications, duplicates, and non-relevant articles were excluded. Publications on endemic pemphigus foliaceus and paraneoplastic pemphigus were excluded. Eventually, 77 manuscripts found in the databases were included (Figure 1). We reviewed published data related to desmoglein distribution in hair follicles, as well as the clinical manifestations, histopathological, immunopathological, and trichoscopic features of scalp involvement in pemphigus. The response of scalp lesions to treatment was analyzed.

3. Results and Discussion

3.1. Desmoglein Expression in Hair Follicles. Desmogleins are the main autoantigens of pemphigus. Desmogleins (Dsgs) are calcium-dependent glycoproteins belonging to the desmosomal cadherin superfamily. Four types of desmogleins have been described. Desmoglein 1 (Dsg1) is expressed in the inner root sheath and the innermost layers of the outer root sheath, as well as epidermal suprabasal cells [6]. Hypotrichosis with microscopically normal hairs was observed in a mutation of Dsg1 gene causing SAM syndrome (severe dermatitis, multiple allergies, and metabolic wasting) [8]. Desmoglein 2 (Dsg2), an antigen of a less prominent role in pemphigus, is highly expressed in the least differentiated cells of the cutaneous epithelium, including the hair follicle bulge of the fetus and adult, bulb matrix cells, and the basal layer of the outer root sheath (ORS) [6]. The expression of desmoglein 3 in the hair follicle depends on the keratinization type. In the infundibulum of the hair follicle keratinization resembles the one in the epidermis (epidermal keratinization), whereas in the isthmus the granular layer is absent (trichilemmal keratinization) [6]. In the areas of trichilemmal keratinization of the hair follicle desmoglein 3 is expressed throughout all layers of the ORS, while in the areas of epidermal keratinization its expression is limited mainly to the basal layer. Moreover, there is a strong expression of Dsg3 detected in the medulla [6]. Desmoglein 4 (Dsg4) is mainly expressed in the hair shaft cortex, the lower hair cuticle, and the upper inner root sheath [9, 10]. Some authors suggest that desmoglein 4 may play a role as an additional autoantigen in pemphigus vulgaris [9]. Mutations in Dsg4 gene are responsible for localized autosomal recessive hypotrichosis [11], monilethrix [12], and in some cases an overlap of both [13]. Schaffer et al. reported three cases of localized autosomal recessive hypotrichosis and monilethrix in combination with scalp erosions presenting mutations in the gene for desmoglein 4 [14].
3.2. Clinical Manifestations of Scalp Involvement in Pemphigus.
In the course of pemphigus vulgaris and pemphigus foliaceus scalp lesions manifest as erosions and crusted or scaly plaques (Figures 2 and 3). Cases of pemphigus vegetans, a variant of pemphigus vulgaris, with lesions localized on the scalp were also described [15–18]. In five cases scalp lesions constituted an isolated manifestation of pemphigus vegetans [15–18]. Unilesional pemphigus vulgaris on the scalp was also described in two cases. In both patients the development of the lesions followed a surgical intervention, in one of the patients a cochlear implantation [19] and a hair transplant in the other [20]. It was hypothesized that trauma may facilitate the overpresentation of epidermal antigens by disturbing the dermoeidermal junction [21]. Scalp erosions limited to the scalp were also reported in the course of pemphigus foliaceus [22, 23]. According to our literature search, no cases of pemphigus erythematous limited to the scalp were reported to date.

Hair Loss in Pemphigus. Scalp lesions in pemphigus may lead to alopecia. Nonscarring alopecia was reported to be an unusual manifestation of scalp pemphigus [24]. Veraitch et al. described 5 patients with pemphigus vulgaris and progressive hair loss with alopecic patches [5]. In this study, apart from characteristic histopathological features of pemphigus, bacterial culture revealed the presence of Staphylococcus epidermidis, Staphylococcus aureus, and Klebsiella pneumoniae. The authors suggested that the combination of anti-desmoglein autoantibody-mediated acantholysis with a concomitant infection may cause the weakening of the hair follicle anchorage and result in alopecia [5]. Hadayer et al. reported a case of a patient with pemphigus vulgaris, in whom alopecia was also accompanied by Staphylococcus aureus infection [24]. Our literature search revealed only one case report of nonscarring alopecia occurring in pemphigus foliaceus [25]. In this publication, the widespread erosions and alopecia were the first manifestations of the disease in a 14-year-old girl. The authors suggested that the occurrence of alopecia in this case could have been associated with an extremely high anti-Dsg1 antibody concentration, equaling 2017 U/ml (cut off value < 14 U/ml). Recently, Yoshida et al. described a case of transition from pemphigus foliaceus to pemphigus vulgaris [26]. The authors observed a correlation between scalp lesions and anti-Dsg3 antibody concentration in enzyme-linked immunoassay [26].

To date, scarring alopecia has been described in 7 patients with pemphigus vulgaris [27–32]. Five of them developed tufted hair folliculitis in the course of prolonged scalp involvement [28–32]. In 5 cases bacterial culture was positive, showing the presence of Staphylococcus aureus [30–33] or Staphylococcus epidermidis [28]. It is unclear whether bacteria may act as an additional factor leading to alopecia. Gaitanis et al. [27] and Lapiere et al. [33] described cases of pemphigus vulgaris in which scarring alopecia was not accompanied by tufting. In both cases the development of cicatricial alopecia was associated with long-lasting and extensive scalp lesions.

Delmonte et al. first observed that in 3 patients with pemphigus vulgaris anagen hairs with intact root sheaths could be easily removed from both lesional and perilesional scalp by a gentle pull test [34]. The authors of this case report called that phenomenon “normal anagen effluvium.” According to Delmonte et al., the loss of anagen hairs with preserved root sheaths is a manifestation of the subclinical involvement of the hair follicle and may be regarded as a hair equivalent to Nikolsky’s sign of the skin [34]. According to Daneshpazhooh et al. anagen hair loss is a common feature of active pemphigus vulgaris. In this article, intact anagen hairs covered by root sheaths were obtained by gentle pull test in 59 of 96 (61.5%) patients with pemphigus vulgaris [35]. Lesional anagen hair loss was observed in 51 of 57 (89.5%) patients, whereas nonlesional anagen hair loss was present in 27 of 57 (47.4%) patients. Mean anagen hair count was significantly higher in severe cases (6.83 +/- 7.89 hairs) as compared to patients with pemphigus vulgaris with a moderate course of the disease (1.06 +/- 1.94 hairs). The authors of this study reported that anagen hair loss was an independent predictor of pemphigus vulgaris severity [35]. In a recent study of Fard et al. performed in 52 patients with newly diagnosed pemphigus vulgaris, a significant correlation between anagen hair loss and the concentration of anti-desmoglein 1 antibody concentration as well as scalp and skin Pemphigus Disease Area Index scores was observed [36].

3.3. Measurement Methods for Scalp Involvement in Pemphigus.
Scalp lesions in pemphigus may differ in size, number, and activity. Objective measurement tools are needed to evaluate scalp involvement in patients with pemphigus. As regards the most commonly used validated measurement...
Table 1: Pemphigus Disease Area Index (PDAI) for the scalp.

| Scalp | Postinflammatory hyperpigmentation or erythema from resolving lesion |
|-------|---------------------------------------------------------------|
| 0—absent | 0—absent |
| 1—one quadrant | 1—present |
| 2—two quadrants | |
| 3—three quadrants | |
| 4—affecting the whole skull | |
| 10—at least one lesion > 6 cm | |

methods evaluating pemphigus disease activity, scalp involvement is separately assessed only with Pemphigus Disease Area Index (PDAI) [37]. The following factors are taken into consideration in scalp PDAI: the distribution of the lesions in scalp quadrants and the size of the lesions as well as their quality, such as the presence of blisters, erosions, or new erythema for active lesions and postinflammatory hyperpigmentation or erythema after resolving lesions (Table 1). In Autoimmune Bullous Skin Disorder Intensity Score (ABSIS) the extent of skin lesions is assessed with the “rule of nine,” where the head and neck area represents 9% of the body surface area. The percentage of body surface area is multiplied by the weighting factor, depending on the activity of the lesions (factor 1.5 for bullae and erosions, factor 1.0 for dry erosions, and factor 0.5 for reepithelialized erosions) [38].

In Pemphigus Vulgaris Activity Score (PVAS) only the presence or absence of scalp involvement is evaluated, whereas the quality and number of lesions as well as the presence of Nikolsky’s sign are not evaluated separately for the scalp [39].

According to Rahbar et al. PDAI may be recommended for use in multicenter studies because it has the highest interrater reliability as well as the highest convergent validity as compared to ABSIS and PVAS [37].

3.4. Histopathological Features of Scalp Involvement in Pemphigus. Intraepidermal blister formation due to acantholysis is a characteristic histopathological finding of pemphigus. Acantholysis is located over the basal layer in pemphigus vulgaris and forms the so-called “tombstone” pattern on the floor of the blister [19]. In pemphigus foliaceus, acantholysis is located under the corneal layer [25]. Acantholysis may also be observed in the structures of hair follicles (Figure 4) [24, 27, 40, 41]. Veraitch et al. observed acantholysis between keratinocytes of the outer root sheath extending from the infundibulum to the suprabulbar level in anagen hair follicles of pemphigus vulgaris lesions [5]. A case report of pemphigus foliaceus by Mlynek et al. described acantholysis which was observed in the infundibular outer root sheath, sparing its infrainfundibular part [25]. In our recent study performed on 32 patients with pemphigus, we observed that follicular acantholysis is a common feature of both types of pemphigus occurring in 12/17 (70.6%) patients with pemphigus vulgaris and 12/15 (80.0%) patients with pemphigus foliaceus [42]. We also confirmed that pemphigus vulgaris and pemphigus foliaceus differ as regards the depth of acantholysis. Acantholysis extended down the entire length of the outer root sheath of the hair follicle in pemphigus vulgaris, whereas in pemphigus foliaceus acantholysis was restricted to the infundibulum [42]. Acantholysis was also observed in vellus hair follicles in 12/17 (70.6%) patients with pemphigus vulgaris and 9/15 (60.0%) patients with pemphigus foliaceus [42]. Furthermore, acantholysis was present in sebaceous gland ducts in 3/17 (17.6%) patients with pemphigus vulgaris and 1/15 (6.7%) patients with pemphigus foliaceus as well as sweat gland ducts in 2/17 (11.8%) and 2/15 (13.3%) patients, respectively [42]. We also observed the miniaturization of sebaceous glands, which was seen in 14/17 (82.4%) patients with pemphigus vulgaris and 12/15 (80%) patients with pemphigus foliaceus. Whether this feature is specific for pemphigus or is associated with other comorbidities, for example, androgenetic alopecia, requires further evaluation [42].

Perifollicular infiltrates were observed in long-standing scalp lesions of patients with pemphigus vulgaris [28]. Superficial inflammatory infiltrates consisting of lymphocytes and neutrophils were observed by Veraitch et al. who reported 5 patients with pemphigus vulgaris and progressive hair loss [5]. Interestingly, the infiltrates did not reach the lowermost bulbar region of the hair follicle. Moreover, hair follicle density and hair cycle were unaffected [5].

As regards the course of pemphigus, perivascular inflammatory infiltrates in scalp lesions commonly consisted of lymphocytes [28, 30]; in rare cases they were accompanied by eosinophils [15, 33]. In our study neutrophils were the most common inflammatory cells observed in the epidermis, occurring in about 40% of patients with pemphigus vulgaris and
60% of patients with pemphigus foliaceus [42]. Mononuclear inflammatory infiltrates were predominant in the dermis, occurring in 82% of patients with pemphigus vulgaris and 93% of patients with pemphigus foliaceus [42]. The presence of keratinization disorders was another common histopathological finding observed in scalp pemphigus [42]. Parakeratosis was observed in pemphigus foliaceus [22]. In our study the hyperkeratinization of follicular ostia was observed in 53% of PV patients and 60% of PF patients [42].

3.5. Direct Immunofluorescence (DIF) of Plucked Hair. The direct immunofluorescence of plucked hairs may be helpful in the diagnosis of pemphigus (Figure 5). The hairs for direct immunofluorescence are obtained by plucking with forceps [7]. Daneshpazhooh et al. reported that the sensitivity of the direct immunofluorescence of plucked hair equals 91% (100 of 110 patients with PV) [43]. Positive intercellular staining of the outer root sheath of anagen hairs was also reported by Rao et al in 85% (17/20) of patients with active pemphigus vulgaris [44]. In this study the direct immunofluorescence of plucked hairs was positive in all patients who had scalp lesions and in 77.7% of patients with no scalp lesions in the course of pemphigus. A negative result of direct immunofluorescence was observed in 3 patients, with two of them being elderly men (mean age: 71 years) and one with mucosal lesions only. In a study conducted by Alexandru et al. [45] the sensitivity of the direct immunofluorescence of anagen plucked hair equaled 100% (12/12 patients with pemphigus vulgaris). In this study 4 out 12 patients had no scalp involvement and one patient was in clinical remission [45]. Daneshpazhooh et al. suggested that the direct immunofluorescence of plucked hairs is a suitable alternative to the traditional DIF of the skin or mucous membranes [43]. The value of the direct immunofluorescence of telogen hairs remains controversial. In the study by Alexandru et al. [45] the direct immunofluorescence of plucked telogen hairs from 3 patients with pemphigus vulgaris was negative, whereas Tanasilovic et al. [46] reported a positive immunofluorescence of outer root sheath in both anagen and telogen hairs of all patients with active pemphigus vulgaris (27/27) and pemphigus foliaceus (8/8), irrespective of scalp involvement. The authors reported that the direct immunofluorescence of plucked telogen hairs was negative in all patients with pemphigus vulgaris or pemphigus foliaceus in remission [46]. According to European guidelines on pemphigus management, plucked hairs may be utilized for DIF for the diagnosis of pemphigus in specialized laboratories [47].

Rao et al. reported on the applicability of the direct immunofluorescence of plucked hairs in monitoring pemphigus activity [48]. The authors observed a positive direct immunofluorescence of plucked hair in a patient with a clinical remission and a negative result of direct immunofluorescence of the skin, suggesting that plucked hair is even a more reliable substrate for direct immunofluorescence than the skin [48]. However, the study was performed on a total of 5 patients with pemphigus vulgaris, no anti-desmoglein enzyme-linked immunoassay was performed, and indirect immunofluorescence was performed on two fixed serum dilutions, which constitute the limitations of the study [48]. A negative direct immunofluorescence of plucked hairs was reported to be a marker of remission [7]. In a recent study of Rai and Harikumar the usability of hair DIF in the assessment of immunological remission was evaluated in a total of 30 patients with pemphigus vulgaris in clinical remission [49]. The clinical remission was defined as no new/nonhealing lesions for at least 6 months in patients being off therapy or being treated with maximum dose of 10 mg of prednisolone daily with or without immunosuppressive therapy such as azathioprine in the dose of 50 mg daily or cyclophosphamide in the dose of 50 mg daily [49]. Positive hair DIF was observed in 14 patients, whereas positive DIF of the skin was present in 10 patients [49]. In 6 patients, hair DIF was positive, while DIF of the skin was negative, which may indicate that hair DIF is a better tool in evaluating immunological remission in patients with pemphigus vulgaris [49]. Daneshpazhooh et al. suggest repeating the DIF of plucked hairs in patients in clinical remission until the negativization and afterwards, performing a conventional DIF [7].

3.6. Trichoscopy of Pemphigus Lesions. Three full-text articles about trichoscopy (dermoscopy of the scalp and hairs) were identified.

Firstly, Pirmez published a case report which described the presence of acantholytic casts as a trichoscopic feature of pemphigus vulgaris. The author regarded this feature as Nikolsky’s sign of the scalp indicating the activity of the disease [50].

In 2014, we described the trichoscopic features of scalp involvement in 9 patients with pemphigus vulgaris and 10 patients with pemphigus foliaceus [51]. The most common trichoscopic features of pemphigus were extravasations occurring in 95% (18/19) of patients with pemphigus and yellow haemorrhagic crusts present in 58% (11/19) of all patients. White polygonal structures were characteristic of pemphigus foliaceus and were visualized in 60% (6/10) of patients with pemphigus foliaceus, whereas vascular abnormalities were more common in pemphigus vulgaris and correlated with the severity of pemphigus measured with scalp PDAI.

In 2017, our group described trichoscopic features of 26 patients with pemphigus vulgaris and 17 patients with...
pemphigus foliaceus [52]. We concluded that the presence of extravasations (20/26, 76.9%), yellow haemorrhagic crusts (19/26, 73.1%), dotted vessels with whitish halo (6/26, 23.1%), circular vessels (6/26, 23.1%), or polymorphic vessels (6/26, 23.1%) can suggest a diagnosis of pemphigus vulgaris [52]. The presence of dotted vessels, dotted vessels with whitish halo, clustered dotted vessels, dotted vessels forming “string of pearls,” and lace-like vessels correlated positively with PDAI scalp activity score [52]. The combination of extravasations (12/17, 70.6%), yellow haemorrhagic crusts (12/17, 70.6%), yellow diffuse scaling (9/17, 52.9%), and tubular scaling (7/17, 41.2%) can indicate a diagnosis of pemphigus foliaceus [52].

3.7. Response to the Treatment. Cases of treatment-resistant scalp lesions in the course of pemphigus were reported [19, 24, 28–32, 53]. Kamphausen et al. [53] reported a case of a 48-year-old man with a severe involvement of the scalp with erosions, crusting, vegetative granulations, and alopecia. The patient was treated with protein A immunoadsorption, intravenous rituximab, intravenous dexamethasone pulses, and oral azathioprine. Complete healing of lesions was observed after 12 weeks [53]. Mlynèk et al. reported a case of pemphigus foliaceus not responding to oral prednisolone at the dose of 1 mg/kg daily, with the intolerance of azathioprine, manifesting as a significant elevation of liver enzymes. The patient was treated with oral prednisolone at a dose of 1 mg/kg daily combined with intravenous immunoglobulins at a dose of 2 g/kg over 5 days in two cycles given at a 4-week interval, followed by eight immunopherases. After a 5-month follow-up the healing of scalp lesions with complete hair regrowth was observed. However, new lesions were still, although sporadically, observed [25].

Contrary to the cases above, Tyros et al. reported a successful treatment of pemphigus foliaceus localized on the scalp and the nose in a patient treated with pimecrolimus cream once daily for 4 weeks as monotherapy [22]. Good response to another topical calcineurin inhibitor, tacrolimus, as monotherapy was observed by Termeer et al. [23].

In our recent paper, we described the efficacy of perilesional and intralesional triamcinolone acetonide injections in two patients with residual scalp involvement in the course of pemphigus vulgaris [54]. Complete clearance of scalp lesions was achieved after, respectively, one and two perilesional and intralesional triamcinolone acetonide (10 mg/ml) injections, showing its possible usability in the treatment of recalcitrant scalp lesions in pemphigus [54].

In conclusion, to date there have not been sufficient convincing data to prove that scalp involvement in pemphigus is associated with poor response to treatment.

Detailed literature data on scalp involvement in pemphigus are shown in Table S1.

4. Conclusions

The scalp is a common location for pemphigus because of a high load of pemphigus antigens distributed in hair follicles. A high concentration of pemphigus antigens in hair follicles makes them a perfect substrate for direct immunofluorescence. Another useful diagnostic method is trichoscopy, especially in cases of pemphigus lesions limited to the scalp. The evaluation of response to treatment of patients with a severe scalp involvement requires large prospective observational studies.

Conflicts of Interest

The authors declare that there are no conflicts of interest regarding the publication of this paper.

Supplementary Materials

Table S1: literature data on scalp involvement in pemphigus. (Supplementary Materials)

References

[1] X. Ding, V. Aoki, J. M. Mascaro Jr., A. Lopez-Swiderski, L. A. Diaz, and J. A. Fairley, “Mucosal and mucocutaneous (generalized) pemphigus vulgaris show distinct autoantibody profiles,” Journal of Investigative Dermatology, vol. 109, pp. 592–596, 1997.
[2] S. R. Arya, A. G. Valand, and K. Krishna, “A clinico-pathological study of 70 cases of pemphigus,” Indian Journal of Dermatology, Venereology and Leprology, vol. 65, pp. 168–171, 1999.
[3] N. Esmaili, C. Chams-Davatchi, M. Valikhanlil et al., “Pemphigus vulgaris in Iran: A clinical study of 140 cases,” International Journal of Dermatology, vol. 46, no. 11, pp. 1166–1170, 2007.
[4] R. Salamanpour, H. Shahkar, M. R. Namazi, and M. R. Rahman-Shenas, “Epidemiology of pemphigus in south-western Iran: A 10-year retrospective study (1991–2000),” International Journal of Dermatology, vol. 45, no. 2, pp. 103–105, 2006.
[5] O. Veraitch, M. Ohyama, J. Yamagami, and M. Amagai, “Alopecia as a rare but distinct manifestation of pemphigus vulgaris,” Journal of the European Academy of Dermatology and Venereology, vol. 27, no. 1, pp. 86–91, 2013.
[6] H. Wu, J. R. Stanley, and G. Cotsarelis, “Desmoglein isotype expression in the hair follicle and its cysts correlates with type of keratinization and degree of differentiation,” Journal of Investigative Dermatology, vol. 120, no. 6, pp. 1052–1057, 2003.
[7] M. Daneshpazhooh, Z. S. Naraghi, A. Ramezani, A. Ghanadan, N. Esmaili, and C. Chams-Davatchi, “Direct immunofluorescence of plucked hair for evaluation of immunologic remission in pemphigus vulgaris,” Journal of the American Academy of Dermatology, vol. 65, no. 6, pp. e173–e177, 2011.
[8] L. Samuelov, O. Sarig, R. M. Haron et al., “Desmoglein 1 deficiency results in severe dermatitis, multiple allergies and metabolic wasting,” Nature Genetics, vol. 45, no. 10, pp. 1244–1248, 2013.
[9] A. Klujuic, H. Bazzi, J. P. Sundberg, A. Martinez-Mir, R. O'Shaughnessy, M. G. Mahoney et al., “Desmoglein 4 in hair follicle differentiation and epidermal adhesion: evidence from inherited hypotrichosis and acquired pemphigus vulgaris,” Cell, vol. 113, pp. 249–260, 2003.
[10] H. Bazzi, A. Getz, M. G. Mahoney et al., “Desmoglein 4 is expressed in highly differentiated keratinocytes and trichocytes in human epidermis and hair follicle,” Differentiation, vol. 74, no. 2-3, pp. 129–140, 2006.
[11] A. Ullah, S. I. Raza, R. H. Ali et al., “A novel deletion mutation in the DSG4 gene underlies autosomal recessive hypotrichosis.
with variable phenotype in two unrelated consanguineous families,” Clinical and Experimental Dermatology, vol. 40, no. 1, pp. 78–84, 2015.

[12] M. Kato, A. Shimizu, Y. Yokoyama et al., “An autosomal recessive mutation of DSG4 causes monilethrix through the ER stress response,” Journal of Investigative Dermatology, vol. 135, no. 5, pp. 1253–1260, 2015.

[13] J.-M. Wang, Y.-J. Xiao, and Y.-H. Liang, “Novel D323G mutation of DSG4 gene in a girl with localized autosomal recessive hypotrichosis clinically overlapped with monilethrix,” International Journal of Dermatology, vol. 54, no. 10, pp. 1163–1168, 2015.

[14] J. V. Schaffer, H. Bazzi, A. Vitebsky et al., “Mutations in the desmoglein 4 gene underlie localized autosomal recessive hypotrichosis with monilethrix hairs and congenital scalp erosions,” Journal of Investigative Dermatology, vol. 126, no. 6, pp. 1286–1291, 2006.

[15] S. Lehrhoff, K. Miller, M. Fischer, H. Kamino, and S. Meehan, “Localized pemphigus with vegetative features,” Dermatology Online Journal, vol. 18, no. 12, p. 11, 2012.

[16] M. Mori, G. Mariotti, V. Grandi, S. Gunnella, and V. Maio, “Pemphigus vegetans of the scalp,” Journal of the European Academy of Dermatology and Venereology, vol. 30, no. 2, pp. 368–370, 2016.

[17] I. Danopoulou, P. Stavropoulos, A. Stratigos, E. Chatziolou, A. Chiou, S. Georgala et al., “Pemphigus vegetans confined to the scalp,” International Journal of Dermatology, vol. 45, pp. 1008–1009, 2006.

[18] G. Ferrara, C. Massone, I. Zalaudek, and G. Argenziano, “Unilateral pemphigus vulgaris of the scalp,” Dermatology Online Journal, vol. 15, no. 10, p. 9, 2009.

[19] G. Oretti, D. Giordano, F. Di Lella, P. Gradoni, E. Zendri, and T. Ferri, “Unilateral pemphigus vulgaris of the scalp after cochlear implantation,” American Journal of Otologyngology, vol. 32, no. 1, pp. 80–81, 2011.

[20] K. Balighi, M. Daneshpazhooh, A. Azizpour, Y. Yoshida, K. Ishii, and A. Ishiko, “Alopecia developed in a persistent pemphigus vulgaris showing features of tufted hair folliculitis,” Annals of Dermatology, vol. 23, no. 4, pp. 523–525, 2011.

[21] M. Daneshpazhooh, M. Fatehnejad, Z. Rahbar, M. Mirshams-Shahshahani et al., “Pemphigus disease activity measurements: Pemphigus disease area index, autoimmune bullous skin disorder intensity score, and pemphigus vulgaris activity score,” JAMA Dermatology, vol. 150, no. 3, pp. 266–272, 2014.

[22] C. C. Termeer, K. Technau, M. Augustin, and J. C. Simon, “Topical tacrolimus (protopic) for the treatment of localized pemphigus foliaceous with topical pimecrolimus,” Case Reports in Dermatological Medicine, vol. 2013, Article ID 489618, 3 pages, 2014.

[23] C. C. Termeer, K. Technau, M. Augustin, and J. C. Simon, “Topical tacrolimus (protopic) for the treatment of localized pemphigus foliaceous,” Journal of the European Academy of Dermatology and Venereology, vol. 18, pp. 636–637, 2004.

[24] N. Hadayer, Y. Ramot, A. Maly, and A. Zlotogorski, “Pemphigus vulgaris with loss of hair on the scalp,” International Journal of Trichology, vol. 3, no. 3, pp. 157–158, 2013.

[25] A. Mlynek, M. Bür, A. Bauer, and M. Meurer, “Juvenile pemphigus foliaceus associated with severe nonscarring alopecia,” British Journal of Dermatology, vol. 161, no. 2, pp. 472–474, 2009.

[26] K. Yoshida, K. Ishii, and A. Ishiko, “Alopecia developed in a transitional case from pemphigus foliaceus to pemphigus vulgaris,” The Journal of Dermatology, 2017.

[27] G. Gaitanis, K. Patmanidis, K. Skandalis, I. Alexis, A. Zioga, and I. D. Bassukas, “Scarring alopecia in pemphigus vulgaris: A rare or underdiagnosed presentation?” European Journal of Dermatology, vol. 23, no. 2, pp. 253–255, 2013.

[28] D. K. Ko, I. S. Chae, K. H. Chung, J. S. Park, and H. Chung, “Persistent pemphigus vulgaris showing features of tufted hair folliculitis,” Annals of Dermatology, vol. 23, no. 4, pp. 865–866, 1995.

[29] S. C. Rackett, M. J. Rothe, D. M. Hoss, C. M. Grin-Jorgensen, and J. M. Grant-Kels, “Treatment-resistant pemphigus vegetans of the scalp,” International Journal of Dermatology, vol. 34, pp. 78–84, 1995.

[30] V. Petronic-Rosic, A. Krunic, M. Mijuskovic, and S. Vesic, “Tufted hair folliculitis: A pattern of scarring alopecia?” Journal of the American Academy of Dermatology, vol. 41, no. 1, pp. 112–114, 1999.

[31] U. Jappe, K. Schröder, D. Zillikens, and D. Petzoldt, “Tufted hair folliculitis associated with pemphigus vulgaris,” Journal of the European Academy of Dermatology and Venereology, vol. 17, no. 2, pp. 223–226, 2003.

[32] S. Saijo and H. Tagami, “Tufted hair folliculitis developing in a recalcitrant lesion of pemphigus vulgaris,” Journal of the American Academy of Dermatology, vol. 38, no. 5, pp. 857–859, 1998.

[33] K. Lapiere, S. Caers, and J. Lambert, “A case of long-lasting localized pemphigus vulgaris of the scalp,” Dermatology, vol. 209, pp. 162–163, 2004.

[34] S. Delmonte, M. T. Semino, A. Parodi, and A. Rebora, “Normal anagen effluvium: a sign of pemphigus vulgaris,” British Journal of Dermatology, vol. 142, pp. 1244–1245, 2000.

[35] M. Daneshpazhooh, H. R. Mahmoudi, S. Rezakhanlou et al., “Loss of normal anagen hair in pemphigus vulgaris,” Journal of Investigative Dermatology, vol. 40, no. 5, pp. 485–488, 2015.

[36] G. D. Fard, H. Khosravi, A. Ghayoumi et al., “Anagen hair loss, anti-desmoglein 1, and pemphigus disease area index: a significant relationship?” Journal of the German Society of Dermatology, vol. 15, no. 9, pp. 946–948, 2017.

[37] Z. Rahbar, M. Daneshpazhooh, M. Mirshams-Shahshahani et al., “Pemphigus disease activity measurements: Pemphigus disease area index, autoimmune bullous skin disorder intensity score, and pemphigus vulgaris activity score,” JAMA Dermatology, vol. 150, no. 3, pp. 266–272, 2014.

[38] M. Pfütze, A. Niedermeier, M. Hertl, and R. Eming, “Introducing a novel Autoimmune Bullous Skin Disorder Intensity Score (ABISIS) in pemphigus,” European Journal of Dermatology, vol. 17, pp. 4–11, 2007.

[39] C. Chams-Davatchi, Z. Rahbar, M. Daneshpazhooh et al., “Pemphigus vulgaris activity score and assessment of convergent validity,” Acta Medica Iranica, vol. 51, no. 4, pp. 224–230, 2013.

[40] M. Mahalingam, “Folliculc acantholyis: A subtle clue to the early diagnosis of pemphigus vulgaris,” American Journal of Dermatopathology, vol. 27, no. 3, pp. 237–239, 2005.

[41] C. Ohata, K. Akamatsu, N. Imai, and Y. Sumimura, “Localized pemphigus foliaceus exclusively involving the follicular infundibulum: A novel peau d’orange appearance,” European Journal of Dermatology, vol. 21, no. 3, pp. 392–395, 2011.

[42] M. Sar-Pomian, J. Czuwara, L. Rudnicka, and M. Olszewska, “Miniaturization of sebaceous glands: A novel histopathological finding in pemphigus vulgaris and pemphigus foliaceus of the scalp,” Journal of Cutaneous Pathology, vol. 44, no. 10, pp. 835–842, 2017.
in pemphigus vulgaris,” *Journal of the European Academy of Dermatology and Venereology*, vol. 23, no. 2, pp. 129–131, 2009.

[44] R. Rao, K. Dasari, S. Shenoi, and C. Balachandran, “Demonstration of pemphigus-specific immunofluorescence pattern by direct immunofluorescence of plucked hair,” *International Journal of Dermatology*, vol. 48, no. 11, pp. 1187–1189, 2009.

[45] A. Alexandru, S. Zurac, C. M. Salavastru et al., “Direct immunofluorescence on hair follicles - Present and future perspectives,” *American Journal of Dermatopathology*, vol. 35, no. 4, pp. 472–476, 2013.

[46] S. Tanasilovic, L. Medenica, and S. Popadic, “Direct immunofluorescence of the outer root sheath in anagen and telogen hair in pemphigus vulgaris and pemphigus foliaceus,” *Australian Journal of Dermatology*, vol. 55, no. 4, pp. e74–e76, 2014.

[47] M. Hertl, H. Jedlickova, S. Karpati et al., "Pemphigus. S2 Guideline for diagnosis and treatment - Guided by the European Dermatology Forum (EDF) in cooperation with the European Academy of Dermatology and Venereology (EADV)," *Journal of the European Academy of Dermatology and Venereology*, vol. 29, no. 3, pp. 405–414, 2015.

[48] R. Rao, K. Dasari, S. Shenoi, C. Balachandran, and P. Dinesh, “Monitoring the disease activity in pemphigus by direct immunofluorescence of plucked hair: A pilot study,” *Indian Journal of Dermatology*, vol. 58, no. 2, p. 164, 2013.

[49] R. Rai and M. Harikumar, “Comparison of direct immunofluorescence of plucked hair and skin for evaluation of immunological remission in pemphigus,” *Indian Dermatology Online Journal (IDOJ)*, vol. 8, no. 5, p. 319, 2017.

[50] R. Pirmez, “Acantholytic hair casts: A dermoscopic sign of pemphigus vulgaris of the scalp,” *International Journal of Trichology*, vol. 4, no. 3, pp. 172-173, 2012.

[51] M. Sar-Pomian, M. Kurzeja, L. Rudnicka, and M. Olszewska, “The value of trichoscopy in the differential diagnosis of scalp lesions in pemphigus vulgaris and pemphigus foliaceus,” *Anais Brasileiros de Dermatologia*, vol. 89, no. 6, pp. 1007–1012, 2014.

[52] M. Sar-Pomian, L. Rudnicka, and M. Olszewska, “Trichoscopy – a useful tool in the preliminary differential diagnosis of autoimmune bullous diseases,” *International Journal of Dermatology*, vol. 56, no. 10, pp. 996–1002, 2017.

[53] I. Kamphausen, F. Schulze, E. Schmidt, D. Zillikens, and M. Kunz, “Treatment of severe pemphigus vulgaris of the scalp with adjuvant rituximab and immunoadsorption,” *European Journal of Dermatology*, vol. 22, no. 6, pp. 786–787, 2012.

[54] M. Sar-Pomian, J. Czuwara, T. Grygorowicz et al., “Efficacy of perilesional and intralosional triamcinolone acetonide injections in pemphigus vulgaris lesions of the scalp: An effective therapeutic option,” *Clinical and Experimental Dermatology*, vol. 43, pp. 168–170, 2018.