Spectrum of autoimmune vesiculobullous diseases in Iran: a 13-year retrospective study

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Background: Autoimmune bullous diseases (ABDs) represent a group of rare, acquired disorders characterized by overlapping features with involvement of the skin and mucous membranes, resistance to treatment, and potential lethality that comprise pemphigus, bullous pemphigoid (BP), epidermolysis bullosa, dermatitis herpetiformis, and linear immunoglobulin A bullous dermatosis.

Aim: The main aim of this study was to identify the epidemiologic characteristics and clinical courses of these common diseases in Hamadan, Iran. Few surveys have been carried out to demonstrate the whole spectrum of ABDs in the literature. Notably, in Hamadan we are the first.

Materials and methods: This 13-year retrospective study was designed to evaluate all of documented data obtained from hospitalized patients with ABDs at Farshchian Hospital from October 1999 to October 2012. We collected information on epidemiologic data, clinical aspects, histologic findings, and therapy prescribed. Data were analyzed using SPSS.

Results: Of 168 patients, 78% had pemphigus. The age of patients at presentation ranged from 1 month to 115 years, with a mean of 47.5 ± 19.93 years. Mucosal or skin involvement of ABDs was statistically significant (P=0.001). The incidence of ABDs differed significantly based on anatomic location (P=0.003). We documented three deaths.

Conclusion: Compared to previous literature, our findings showed equal epidemiologic properties in Iran. Although pemphigus was the most common ABD followed by BP, it is expected that in line with the global trend, an increase in BP will be driven by population aging in Iran.

Keywords: autoimmune bullous disease, pemphigus, prevalence

Introduction
Autoimmune bullous diseases (ABDs) represent a group of rare, acquired disorders characterized by overlapping features, resistance to treatment, and potential lethality. The most important pathophysiological mechanisms in the skin of these patients have been identified as disruption of the host immune system, which leads to clinical blister formation. This group of diseases includes bullous pemphigoid (BP), dermatitis herpetiformis, linear immunoglobulin A (IgA) bullous dermatosis (LABD), and epidermolysis bullosa (EB).

Many published epidemiologic studies have focused on pemphigus, but few have encompassed the total spectrum of ABDs. Few surveys have been carried out to demonstrate the entire spectrum of ABDs in Iran. Notably, in Hamadan, Iran, we are the first. The main aim of this study was to identify the epidemiologic characteristics and clinical courses of these common diseases in Hamadan.
Materials and methods

The descriptive, retrospective, cross-sectional study was carried out in the Dermatology Department of Farshchian Hospital, Hamadan University of Medical Sciences, the regional secondary referral center for cutaneous diseases. This study conformed to the ethical guidelines of the 1975 Declaration of Helsinki. Hamadan University of Medical Sciences Institutional Review Board approved the research. Patient consent is not required for this retrospective study. All patients hospitalized at the department between October 1999 and October 2012 with a clinical diagnosis of ABDs were examined to characterize epidemiologic aspects of ABDs in Hamadan. Data were collected by means of record review, and the parameters of interest were surveyed in a protocol developed by the investigators. The criteria used for making the diagnosis were clinical features along with histopathologic findings; in addition, Direct immunofluorescence was properly used in many patients (but not all cases) when necessary. Documented information of bedridden patients included age, sex, bedridden duration, BD subtype, treatment modality, treatment duration, family history, mortality rate, and histopathologic results. A number of lesions approved on different sectors of the body and on the mucous membranes were detected. Details of documents were reviewed with regard to the clinical course of each patient at the onset of the disease, including age of patient in which the disease began, primary affected site, and season at onset. Thereby, the duration of the disease was calculated, ie, the interval between the onset of disease until the present. In this period, first complete remission and recurrence rate were noticed. Patients with incomplete or missing data in their records were excluded. The number of files excluded due to incompleteness was trivial, because we tried our best to avoid missing data. For this reason, two of us separately kept a careful check on each file for a second time to ensure that the findings were accurate. Therefore, we made sure we did not miss any important details.

Data were analyzed using SPSS version 20.0 (IBM Corporation, Armonk, NY, USA). Frequencies and percentages were calculated for qualitative variables. Means, mode, and standard deviation were calculated for quantitative variables. Student’s t-test and the χ2 test were used for statistical analysis. Fisher’s exact test was performed as necessary. Confidence rate higher than 95% was considered significant (P<0.05).

Results

In sum, 168 patients were diagnosed to have ABDs. The characteristics of the patients according to the type of disease are given in Table 1. There were 78 male patients (46.4%) and 90 female patients (53.6%), with a male:female ratio of 1:1.15. We found that women had a slightly higher rate of ABDs than men, which was not statistically significant (P=0.119). The distribution of patients in age-groups is shown in Figure 1. The youngest patient was a 1-month-old newborn with EB at birth, and the oldest patient was 115-years-old and had pemphigus. The most frequent ages for ABDs were 24 and 61 years, each of which represented seven patients (4.2%). The majority of the patients (20.8%) were in the fourth decade of their life.

Characteristics of the diseases are shown in Table 2. EB represented the most long-standing symptoms. The sites of involvement in the first episode were as follows: the two main sites were oral mucosa, with 69 cases, and front of the trunk, with 47 cases. Lower frequencies were seen on the head, posterior trunk, lower extremities, upper extremities, neck, and genitals with 29, 28, 19, 15, three, and one, respectively. The sum of these frequencies was greater than 168, which reflected disseminated lesions in some patients in the first episode. The incidence of ABDs differed significantly based on anatomic location (P=0.003). Phenotypes of mucocutaneous involvement in ABD patients revealed that 86 cases showed involvement of mucous membranes and skin surface concomitantly. In 59 patients, skin lesions were seen alone, and in 23 cases mucosal lesions were seen alone, which was statistically significant (P<0.001).

Table 1 Characteristics of the patients with different autoimmune bullous diseases (ABDs)

<table>
<thead>
<tr>
<th>ABD</th>
<th>Cases, n (%)</th>
<th>Male:female ratio</th>
<th>Mean age at onset (years ± SD)</th>
<th>Mean age at presentation (years ± SD)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Pemphigus</td>
<td>131 (78)</td>
<td>1:1.1</td>
<td>43.30±17.20</td>
<td>45.44±17.30</td>
</tr>
<tr>
<td>BP</td>
<td>21 (12.5)</td>
<td>1:1.1</td>
<td>71.19±14.06</td>
<td>72.23±13.77</td>
</tr>
<tr>
<td>DH</td>
<td>11 (6.5)</td>
<td>1:4.5</td>
<td>38.18±18.90</td>
<td>40.27±18.42</td>
</tr>
<tr>
<td>EB</td>
<td>5 (3)</td>
<td>1:4</td>
<td>5.2±10.54</td>
<td>13.42±10.44</td>
</tr>
<tr>
<td>LABD</td>
<td>0</td>
<td>–</td>
<td>–</td>
<td>–</td>
</tr>
<tr>
<td>Total</td>
<td>168 (100)</td>
<td>1:1.15</td>
<td>45.32±20.42</td>
<td>47.50±19.93</td>
</tr>
</tbody>
</table>

Abbreviations: SD, standard deviation; BP, bullous pemphigoid; DH, dermatitis herpetiformis; EB, epidermolysis bullosa; LABD, linear IgA bullous dermatosis.

Figure 1 Distribution of patients in age-groups.
The shortest interval between the presentation of lesions and diagnosis of disease was 1 month, whereas the longest was 121 months. That means this patient had been treated based on diagnoses other than ABDs for more than 10 years.

The first manifestations occurred most often in summer and autumn (60%), followed by spring and winter. The onset of disease and season did not correlate statically (P<0.273). Positive family history was seen only in three pemphigus patients. Other cases in our study had not any history of ABDs in their family, which was not statistically significant (P=0.979). A mismatch between clinical and histopathologic diagnosis was seen only in one pemphigus case.

With a mean of 9.8 months, EB had the most prolonged treatment duration. The most prevalent modality for treatment of ABDs was pulse therapy; 103 patients (61.3%) received corticosteroids as pulse therapy and 62 patients (36.9%) were treated with oral corticosteroids. Cyclophosphamide was added to steroids in 93 cases (55.4%) and azathioprine in nine cases (5.4%) as adjuvant therapy. The remaining 66 patients (39.2%) were treated with corticosteroids alone. Only three patients, including BP, dermatitis herpetiformis, and EB, were hospitalized without any treatment, and were just admitted for confirming the diagnoses.

All of the three cases who died from ABDs in our study had pemphigus: a 52-year-old woman and two men aged 45 and 75 years. The woman was treated with corticosteroid as pulse therapy in conjunction with cyclophosphamide. The men were treated with high-dose oral corticosteroids without adjuvant drugs. All of them died in our hospital from sepsisemia due to treatment complications.

**Table 2. Courses of the different autoimmune bullous diseases (ABDs)**

<table>
<thead>
<tr>
<th>ABD</th>
<th>Mean duration of disease (months ± SD)</th>
<th>Dx gap* (months ± SD)</th>
<th>Treatment gap (months ± SD)</th>
<th>Recurrence rate (mean ± SD)</th>
<th>Primary remission interval (months ± SD)</th>
<th>Skine lesions (mean ± SD)</th>
<th>Mean duration of the last treatment period (months ± SD)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Pemphigus</td>
<td>5.26±1.82</td>
<td>2.35±2.69</td>
<td>0.66±1.46</td>
<td>0.57±1.16</td>
<td>0.18±0.40</td>
<td>5.64±15.01</td>
<td>3.00±0.14</td>
</tr>
<tr>
<td>BP</td>
<td>2.11±3.19</td>
<td>3.63±4.15</td>
<td>4.00±2.45</td>
<td>0.00±1.0</td>
<td>2.40±1.36</td>
<td>3.00±0.14</td>
<td>2.00±0.0</td>
</tr>
<tr>
<td>DH</td>
<td>4.60±5.46</td>
<td>4.00±2.45</td>
<td>0.00±1.0</td>
<td>0.60±1.34</td>
<td>2.40±1.36</td>
<td>3.60±5.46</td>
<td>2.00±0.0</td>
</tr>
<tr>
<td>EB</td>
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<td>2.35±2.69</td>
<td>0.66±1.46</td>
<td>0.57±1.16</td>
<td>0.18±0.40</td>
<td>5.64±15.01</td>
<td>3.00±0.14</td>
</tr>
<tr>
<td>Total</td>
<td>8.21±1.49</td>
<td>4.06±1.69</td>
<td>0.31±1.48</td>
<td>0.68±1.76</td>
<td>2.69±1.53</td>
<td>3.00±0.14</td>
<td>2.06±0.48</td>
</tr>
</tbody>
</table>

Notes: Mean interval between lesion presentation and diagnosis of disease; Dx, diagnosis; sD, standard deviation; BP, bullous pemphigoid; Dh, dermatitis herpetiformis; E, epidermolysis bullosa.

Discussion

ABDs are uncommon, and the exact incidence and prevalence depend on the population studied. Similar to an earlier observation from our country, we found a female predilection in each group of ABDs. This was in common with other countries like Tunisia and Kuwait. The mean age of our patients at onset of the disease was 45.32±20.42 years, which was in line with a previous Iranian result (44.99±17.35 years). The mean age of the patients at presentation to our hospital (47.5 years) was comparable to Al-Sabah Hospital in Kuwait, with a similar ethnic background (42 years). It is noticeable that the differences in threshold for hospitalization of ABD patients in different countries may explain some of the variation in results. Indeed, ethnic, environmental, or genetic variation emphasized the different results in various countries.

What really allowed unambiguous differentiation of the ABDs was the characterization of their histology. Our...
results confirmed that our clinical diagnosis, almost always, was consistent with histopathologic findings. This evidence would provide a major clue as to the importance of clinical experience of specialists. We found that the average primary remission interval in all groups of ABDs was lower than 1 year. Therefore, repeated physical examinations during regular visits in the first year after treatment may improve the quality of management.

Pemphigus

We reported pemphigus as the most common ABD in Hamadan. It has been proposed by other investigators in some other countries, including northwestern Romania (58.6%), Tunisia (52.9%), Kuwait (47%), and Thailand (63.3%). Also, two Iranian studies reported the percentage as 86.1% and 83.6%. The mean age of pemphigus patients at the onset of disease has been reported as 38, 42, 45.8, and 53 years, and was 43.3 years in our study. The mean age of our pemphigus patients at presentation (45.44 years) was consistent with earlier research, at 44.6, 45.6, 52.5, and 62.43 years. Although it was not statistically significant, we found a female predominance in the pemphigus patients (male:female = 1:1.11). There have been similar results in other provinces of Iran and other countries. By a review of the literature, the male:female ratio in pemphigus was 1:1.9, 1:1.11, 1:1.33, 1:1.5, 1:1.53, 1:1.64, 1:1.75, 1:1.8, and 1:2.23.

In our pemphigus cases, the oral mucosa was the most common primary site of involvement in the beginning of disease (42.7%). This was consistent with Farshchian and Pilehvar (34.3%), Salamanpour et al (59.3%), Chams-Davatchi et al (62%), and Sadra-Ashkevari et al (63%). In the course of the disease, pemphigus most often affected (60%) both the mucous membranes and the skin in our study. This was similar to some other studies: 64%, 67.8%, 70%, and 16.7%. However, Abdolsamadi et al found that the oral mucosa in 44.9% of pemphigus patients was exclusively affected at presentation time, and oral and skin lesions were seen simultaneously in only 16.7% of cases. Farshchian and Pilehvar detected oral lesions with or without skin lesions in 83.8% of pemphigus patients. The mean duration of pemphigus (from the beginning of lesions until the last presentation) was 8.26 months in our findings. In contrast, Chams-Davatchi et al found 4.5 years for duration of the disease, and their mean follow-up duration was 3.8 years. Also, Tsankov et al showed that the mean duration of the pemphigus was 8.64 years for women and 6.42 years for men.

Pemphigus is a rare but chronic disease, and is associated with significant morbidity and mortality. In the present study, there were three deaths (2.29%) from a total of 131 pemphigus cases in the period of 13 years. This was in the range of other surveys, which demonstrated 1%–20%: 2.3% (36 deaths in 1,560 patients), 7.14% (eight deaths in 112 patients), 7.65% (18 deaths in 235 patients), 9.45% (seven deaths in 74 patients), 10.8% (four deaths in 37 patients), and 20% (eleven deaths in 55 patients). These eleven deaths were observed in Kerman, where Shamsadini et al showed that mortality rates were 7.3%, 13.2%, and 38.5% at 2, 6, and 10 years, respectively. Two other prospective studies, from Tehran (6.2%) and Taiwan (10.31%), have validated the high mortality rate due to pemphigus. Huang et al found that overall mortality among pemphigus patients was two times (2.36-fold) greater than that of the general population. Therefore, the survival rate in patients diagnosed with pemphigus was significantly lower than that expected in the general population. We found that sepsis was the commonest cause of deaths in pemphigus patients (100%), consistent with earlier studies.

The main information about the prognosis of pemphigus was from Halaji et al. They showed that old age, male sex, interval between beginning of lesions and diagnosis of more than 6 months, and more than ten skin lesions were accompanied by poor prognosis.

Bullous pemphigoid

Although BP was the second-most common ABD and the most common subepidermal BD in our series, predictable BP prevalence should be much higher than the number that was referred to us. It is important to remember the nonspecific BP clinical symptoms that may lead to underestimation. These points highlight the value of clinical diagnosis of BP in older patients. The mean age of BP patients was 82.6 years, and 74.5 years in France, the UK, Portugal, and the USA, respectively. These were a little higher than our result (72 years), though in Thailand it was 69.3 years. Likewise, another study in Iran reported 64.1 years. This would be interpreted as due to higher life expectancy in developed countries. This implies that as life expectancy increases, the incidence and the duration of disease may increase. In this way, a study has pointed out that life expectancy for Iran or Thailand is much lower than developed countries. Furthermore, we found that incidence of BP increased significantly with age. Several surveys confirmed this finding. Most authors agreed on general opinion considering increased BP incidence is attributable to a greater proportion of older persons in the general population. Broader analytic studies have been carried out, one in the USA and a population-based
A review of the literature revealed that BP was the most frequent ABD in the USA, Canada, and Europe: France, the UK, and Germany. Wong and Chua found that BP in Singapore was diagnosed three times more frequently than pemphigus. This is in contrast with other regions in which authors identified BP as the second-most prevalent ABD. For example, the ratio of BP to total ABDs was reported as 11.6% in Tehran, 22% in Kuwait, 23.6% in Tunisia, 34.5% in Romania, and 12.5% in our series. Therefore, BP is much more common in developed countries rather than developing countries like Iran. Bertam et al. diagnosed 27 BP cases (65%) among a total of 41 ABD patients in Germany. The mean age at disease onset was 74 years in this European city. North East Scotland appeared to have a relatively high incidence rate for Europe. The mortality rate due to BP in this region was considerable: 48% of patients with BP died within 2 years of diagnosis. Joly et al. found that the incidence of BP in France had increased threefold in the last 15 years. Relative to expected age- and sex-specific overall death rates in the general population in France, there was a more than sixfold excess of mortality among BP patients. That was assessed from a prospective cohort study. Among bullous skin diseases, pemphigoid was the leading cause of mortality in Canada. Although, BP mortality was much lower in the USA than Canada and Europe, it was higher than previous estimates.

Although ultrapotent topical corticosteroids are the first-line therapy for BP, all of our admitted patients, even BP cases, received systemic corticosteroids, and most of the protocols involved pulse therapy. The cause of this paradox is reasonable in a way that is clear. Mild-to-moderate BP patients usually do not need to be admitted. Therefore, our patients needed more aggressive treatment modalities. Jowkar et al. similarly showed that systemic drugs were administered to all BP patients; moreover, prednisolone monotherapy was the most frequent medication in a Shiraz hospital.

**Epidermolysis bullosa**

We detected EB as a very rare ABD (3%). Other authors have confirmed it as 0.5%, 0.6%, and 2.3%. EB cases were the youngest patients at the onset of the disease, but it was taking more time for diagnosis compared with other ABDs. This would be explained by delayed presentation to physicians or misdiagnosis with other infantile or childhood vesiculobullous diseases. Surprisingly, this time-consuming diagnosis was followed by immediate treatment. Noticeably, we found that EB had the longest disease course.

**Linear IgA bullous dermatosis**

LABD is rare in Western Europe, affecting mainly adults, but is more prevalent in Tunisia and Kuwait, with more children involved. Actually, it is the commonest ABD of childhood in Tunisia. Daneshpazhooh et al. found that in Iran, LABD was seen in only 0.4% of patients. Also, it is rare in Singapore. We observed no case of hospitalized LABD in our study.

**Limitations**

Data obtained by reviewing inpatient and outpatient records revealed that some cases may not reach our department, because they may be treated without hospitalization. Prospective studies or outpatient-plus-inpatient sampling may partially correct this bias.

**Conclusion**

Our findings compared to previous literature showed equal epidemiologic properties in Iran. Skin or mucosal involvement and anatomic location of lesions were two identifiable variables in the ABDs, with statistically significant value. Systemic corticosteroids, either with or without adjuvant drugs, were ordered for all of the patients, and most of them were cured. In our hospital, three patients died due to septicemia. In general, in most regions of our country, the whole spectrum of ABDs has not been described completely. Indeed, the epidemiologic characteristics must be distinguished in each state of Iran to demonstrate the therapeutic approach.

**Disclosure**

The authors report no conflicts of interest in this work.

**References**