Pemphigus in the eastern region of Turkey

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Abstract

Introduction: Pemphigus refers to a group of rare autoimmune vesiculobullous diseases with high morbidity and mortality, mainly affecting the skin and the mucosae.

Aim: To evaluate the incidence of pemphigus in the Eastern region of Turkey by analysing the clinical characteristics of the patients including the drugs used in the treatment and their side-effect profiles and dosing schedules, duration of treatment, length of hospital stay, and the characteristics of the lesions.

Material and methods: The retrospective study included 130 pemphigus patients who were admitted to the in-and out-patient dermatology clinics at the University Medical School between January 2005 and October 2017. Age, gender, length of hospital stay, localization of the lesions, initial and maintenance dosages, treatment-related complications, comorbidities, family history, and smoking status were recorded for each patient.

Results: The 130 patients included 62 (47.69%) men and 68 (52.31%) women, of which 73 (56.15%) patients had a smoking history. The annual incidence rate of pemphigus was 1.036/100,000 population and pemphigus vulgaris (PV) was the most common clinical subtype of pemphigus in our patients (n = 122; 93.85%). The most common side effect of the treatment was oral candidiasis, followed by Cushingoid appearance.

Conclusions: The results indicated that the incidence of pemphigus is remarkably high in the Eastern region of Turkey and the associated factors including lifestyle changes should be taken into mind when considering these patients.

Key words: pemphigus, smoking, foods.

Introduction

The word ‘pemphigus’ takes its root from the Greek word ‘pemphig’, which literally means pustule [1]. Pemphigus refers to a group of rare autoimmune vesiculobullous diseases with high morbidity and mortality, mainly affecting the skin and the mucosae. Pemphigus vulgaris (PV) is the most common clinical subtype of pemphigus characterized by the presence of immunoglobulin G (IgG) antibodies that target the ectodomain of desmosomal cadherins, thus resulting in loss of intracellular adhesion between keratinocytes [2]. Pemphigus has been shown to have an incidence of 0.1–0.5/10,000 population with no gender preponderance. Moreover, pemphigus has a peak incidence between the fourth and sixth decades and is rarely seen in children and elderly patients [3]. The bullae in pemphigus easily rupture to form painful crusted erosions. Moreover, Nikolsky’s sign is a typical clinical sign of pemphigus [1].

Diagnosis of pemphigus is based on the clinical (flaccid bullae, vesicles, and erosions on the epithelium of mucous membranes and skin), typical histological (epidermal acantholysis), and fluorescent findings (circulating and bound skin antibodies directed against keratinocyte surface), followed by the Tzanck test which involves cytological examination of the material scraped from the floor of a bulla [2].

High-dose corticosteroids are often required in the treatment of pemphigus to achieve remission and to reduce the chronic recurrent course of the disease although death in all forms of pemphigus mostly results from the side effects of steroids. In addition, adjuvant immunosuppressive agents such as cyclophosphamide, azathioprine, and methotrexate can also be required during the course of the treatment [4].

Aim

The aim of this study was to evaluate the incidence of pemphigus in the Eastern region of Turkey by analysing the clinical characteristics of the patients including...
the drugs used in the treatment and their side-effect profiles and dosing schedules, duration of treatment, length of hospital stay, and the characteristics of the lesions.

**Material and methods**

The retrospective study included 130 pemphigus patients who were admitted to the in- and out-patient dermatology clinics at the Medical School between January 2005 and October 2017. This study was approved by the Clinical Ethics and Research Committee of the Yuzuncu Yil University, Faculty of Medicine (Date: 21.11.2017, Number: 12). Age, gender, length of hospital stay, localization of the lesions, initial and maintenance dosages, treatment-related complications, comorbidities, family history, smoking status, and the intake food suspected of triggering pemphigus were recorded for each patient. Diagnosis of pemphigus was established based on the clinical and histopathologic examinations followed by a direct immunofluorescence test. PV was diagnosed in the presence of flaccid bullae and vesicles on the skin, mucosal erosions, presence of Nikolsky’s sign, and typical localizations of the lesions. Additionally, histopathologic examination was performed to analyse the presence and degree of acantholysis and a direct immunofluorescence test was performed to analyse intercellular IgG and C3 deposits (chicken-wire appearance). On the other hand, pemphigus foliaceus (PF) was diagnosed in the presence of squamous, crusted erosions in a seborrheic distribution and subcorneal bullae, whereas pemphigus erythematosus was diagnosed in the presence of erythematous squamous plaques in the face and upper chest and serological detection of antinuclear antibodies, IgA pemphigus was diagnosed in the presence of vesicular pustules distributed in annular pattern and presence of IgA deposition in the direct immunofluorescence test, and paraneoplastic pemphigus was diagnosed based on Camisa and Helm’s criteria [5].

Control of disease activity was defined as the time when the established lesions began to heal, formation of new lesions ceased, non-epithelialized ulcerations and erosions were detected, and the Nikolsky’s sign turned negative. Remission was defined as the absence of new lesions while the patient was off therapy for a minimum period of 6 months.

**Statistical analysis**

Data were analysed using IBM SPSS Statistics for Windows 22.0 (Armonk, NY: IBM Corp.). The normality of distribution was tested using histogram plot and Kolmogorov-Smirnov test. Descriptive statistics were expressed as mean, standard deviation (SD), median, and minimum-maximum values. Pearson’s $\chi^2$ test and Fisher’s Exact Test were used for comparing 2 × 2 tables. Nonparametric data were evaluated using Kruskal-Wallis test and ordinal variables were compared using Spearman’s correlation coefficient. A $p$-value of $< 0.05$ was considered significant.

**Results**

The 130 patients included 62 (47.69%) men and 68 (52.31%) women. Of the 130 patients, 73 (56.15%) patients were smoking and 45 (34.62%) patients had a history of anxiety disorder or depression. When we analysed the patients according to food suspected of triggering pemphigus, we found that 90% (117/130) of patients had food intake including garlic, onion and leek (Table 1).

The 12-year incidence rate of pemphigus among our patients was 12.437/100,000 population, with an annual incidence rate of 1.036/100,000 population.

| Parameter                                      | $N$  | %   |
|-----------------------------------------------|------|-----|
| Gender                                        |      |     |
| Male                                          | 62   | 47.69|
| Female                                        | 68   | 52.31|
| Affected site                                 |      |     |
| Skin                                          | 8    | 6.15 |
| Mouth                                         | 23   | 17.69|
| Both                                          | 99   | 76.15|
| Smoking                                       |      |     |
| No                                            | 57   | 43.85|
| Yes                                           | 73   | 56.15|
| Intake of food suspected of triggering pemphigus (onion, garlic or leek) |       |     |
| No                                            | 13   | 10.00|
| Yes                                           | 117  | 90.00|
| Anxiety disorder or depression                |      |     |
| No                                            | 85   | 65.38|
| Yes                                           | 45   | 34.62|
Of the 130 patients, 122 (93.85%) patients were diagnosed with PV, including 56 (46%) men and 66 (54%) women with a mean age of 51.72 ± 12.71 years. In addition, 4 (3.08%) patients were diagnosed with PF who had a mean age of 54.75 ± 18.21 years, whereas the other pemphigus subtypes were detected in only 1 patient each (Table 2).

The positive DIF findings for pemphigus subtype were as follows:

- Positive DIF for pemphigus vulgaris; 95.9% (117/122): intercellular, intraepidermal IgG: 84.4% (103/122), and intercellular, intraepidermal C3 in 89.3% (109/122).
- Pemphigus foliaceus (4/4): intercellular, intraepidermal IgG in 75% (3/4), and intercellular, intraepidermal C3 in 73% (4/4). Interestingly, 100% (4/4) of the patients showed C3 deposits in the upper levels of the epidermis.
- Pemphigus erythematosus (1/1): intercellular, intraepidermal IgG in the patient (1/1) linear C3 deposits at the basement membrane zone in 100% (1/1).
- IgA pemphigus (1/1): intercellular, intraepidermal IgA deposits in 100% (1/1) and IgM deposits at the basement membrane zone in 100% (1/1).
- Paraneoplastic pemphigus (1/1): intercellular, intraepidermal IgG in the patient (1/1), intercellular, intraepidermal C3 and linear C3 deposits at the basement membrane zone in 100% (1/1).
- Pemphigus vegetans (1/1): intercellular, intraepidermal IgG: 100% (1/1) and intercellular, intraepidermal C3 in 100% (1/1).

Overall mean age was 52.20 ± 12.96 years, median hospital stay was 3 days, median diagnostic delay was 4 months, median initial steroid dosage was 80 mg, median duration of disease was 7 years, and the median length of time to disease control was 20 days (Table 3).

Mean duration of disease was 6.93 ± 3.28 years in PV patients as opposed to 6.50 ± 1.73 years in PF patients (Table 4).

Steroids alone were administered in all PF patients, whereas steroids + azathioprine were used in 56.2% and steroids alone were used in 23.97% of the PV patients (Table 5).

The most common side effect of the treatment was oral candidiasis, followed by Cushingoid appearance. Moreover, most of the patients were detected with more than one side effect (Table 6).

### Discussion

The results indicated that the incidence of pemphigus is considerably high in the Eastern region of Turkey. However, although the incidence rate in our study was found to be higher than that of Finland, Saudi Arabia, the Mediterranean Region in Turkey, Iran, and Greece, it was found to be lower than that of Israel. On the other hand, although the incidence rates in Iran and Greece were lower than the incidence rate in our study, they were found to be higher than those reported in the literature, which could be attributed to the geographical proximity of Turkey to Iran and Greece. The high incidence of pemphigus in our study could be explained by the high rate of consanguineous marriages in our region and the predisposition to pemphigus associated with various human leukocyte antigens (HLA) [6–11].

Pemphigus has been shown to have either an equal incidence in men and women or to have a female preponderance. However, Uzun et al. showed that pemphigus has a male preponderance in countries including

### Table 2. Distribution of pemphigus subtypes according to mean age and genders

| Pemphigus subtype | N  | %   | Mean age [years] | Gender |
|-------------------|----|-----|-----------------|--------|
| Vulgaris          | 122| 93.85| 51.72 ± 12.71   | 56 M/66 F |
| Foliaceus         | 4  | 3.08| 54.75 ± 18.21   | 4 M    |
| Erythematous      | 1  | 0.77| 77.00           | 1 F    |
| IgA               | 1  | 0.77| 51.00           | 1 F    |
| Paraneoplastic    | 1  | 0.77| 73.00           | 1 M    |
| Vegetans          | 1  | 0.77| 56.00           | 1 F    |
| Total             | 130| 100.00| 52.20 ± 1.96    | 62 M/68 F |

M – male, F – female.

Mean duration of disease was 6.93 ± 3.28 years in PV patients as opposed to 6.50 ± 1.73 years in PF patients (Table 4).

Steroids alone were administered in all PF patients, whereas steroids + azathioprine were used in 56.2% and steroids alone were used in 23.97% of the PV patients (Table 5).
Spain and Saudi Arabia [10–14]. On the other hand, auto-
immune diseases are seen at high rates in women. In our
study, men and women display a close rate, which means
autoimmunity is not the sole factor in the aetiology of
this disease. Pemphigus has been linked to different HLA
antigens in different populations. This association has
been mainly with class-II antigens [15]. Therefore, genetic
factors may play an important role in this disease.
Pemphigus usually has an onset between the fourth
and sixth decades [3]. However, the mean age of onset
has been reported to be lower in some countries such
as Iran (38 years) and Tunisia (36 years) [7, 14, 16]. In line
with the literature, the mean age of onset was 52.2 years
in our patients.
Pemphigus vulgaris (PV) is known to be the most
common subtype of pemphigus diseases. However, it
has been reported that pemphigus erythematous is
more common in Finland, PF is more common in Tuni-
sia, and fogo selvagem (endemic pemphigus foliaceus)
is more common in Brazil compared to other subtypes
[9, 16–18]. In line with the literature, PV was the most
common pemphigus subtype diagnosed in our pa-
tients. Moreover, the incidence of PV in our study was
remarkably higher than those reported in the literature
(122/130; 93.85%).
The bullae seen in PV can be localized in the mucosae
or skin, or both [2]. In our study, the bullae were mostly
localized in both the mucosae and skin (n = 99; 76.15%),
whereas the bullae localized in the oral mucosa alone
were present in 23 (17.69%) patients.

### Table 4. Distribution of pemphigus subtypes according to mean duration of disease

| Pemphigus subtype | Duration of disease [years] |
|-------------------|-----------------------------|
|                   | Mean | SD  | Median | Min. | Max. |
| Vulgaris          | 6.93 | 3.28| 7.00   | 1.00 | 12.00|
| Foliatecus        | 6.50 | 1.73| 6.50   | 5.00 | 8.00 |
| Erythematous      | 5.00 | –   | 5.00   | 5.00 | 5.00 |
| IgA               | 5.00 | –   | 5.00   | 5.00 | 5.00 |
| Paraneoplastic    | 5.00 | –   | 5.00   | 5.00 | 5.00 |
| Vegetans          | 7.00 | –   | 7.00   | 7.00 | 7.00 |

Kruskal-Wallis test; p = 0.949.

### Table 5. Distribution of pemphigus subtypes according to the drugs used in the treatment

| Pemphigus type   | Drugs                  |
|------------------|------------------------|
|                  | Steroid | Steroid + AZA | Steroid + MMP | Steroid + AZA + MMP | Steroid + AZA + IVIg | Steroid + AZA + MMP + IVIg | Steroid + AZA + MMP + CsA |
| Vulgaris         | 30      | 24.59         | 68            | 55.73            | 2                  | 1.63                       | 1                           | 0.82                       |
| Follatecus       | 4       | 100           |                |                  |                    |                            |                             |                            |
| Erythematous     | 1       | 100           |                |                  |                    |                            |                             |                            |
| IgA              | 1       | 100           |                |                  |                    |                            |                             |                            |
| Paraneoplastic   | 1       | 100           |                |                  |                    |                            |                             |                            |
| Vegetans         | 1       | 100           |                |                  |                    |                            |                             |                            |

AZA – azathioprine, MMP – mycophenolate mofetil, CsA – cyclosporine.

### Table 6. Side effects of the treatment detected in the patients

| Side effects        | N   | %   |
|---------------------|-----|-----|
| Oral candidiasis    | 101 | 77.70|
| Cushingoid appearance | 89  | 68.46|
| Weight gain         | 71  | 54.61|
| Striae              | 45  | 34.61|
| Bacterial infection | 41  | 31.53|
| Excessive hair growth | 33  | 25.38|
| Osteoporosis        | 29  | 22.30|
| Glaucoma/cataract   | 13  | 10.00|
| Avascular necrosis  | 11  | 8.46 |
Uzun et al. reported that painful and resistant oral ulcerations and erosions were one of the most important indicators of pemphigus in their patients and also noted that the disease was limited to oral mucosa in only 7.3% of the patients [11]. Similarly, these findings were consistent with the findings of our study, which implicates that PV should be considered in the differential diagnosis of the diseases that manifest only with erosions and ulcerations in the oral mucosa.

The literature indicates that the diagnosis of PV is often delayed due to various reasons. Bozdag and Bilgin reported a mean diagnostic delay of 4.8 months in their patients [19]. Similarly, we had a mean diagnostic delay of 4 months in our patients, which implies that direct immunofluorescence biopsy should be kept in mind in order to prevent a diagnostic error in the patients undergoing oral biopsy.

In our patients, the initial steroid dosage was 60–100 mg/day and the dosage was increased gradually in refractory patients. However, in line with the literature, the steroids were not ceased completely in any patient.

Adjuvant immunosuppressive agents are commonly used in the treatment of PV to reduce the dose of steroids and to achieve better control of the disease. These agents mostly include azathioprine, mycophenolate mophetil, cyclophosphamide, cyclosporine, and rituximab [2, 20]. In our study, azathioprine was the most common agent used in our patients, followed by mycophenolate mophetil, cyclosporine, and intravenous immunoglobulin (IVIg). However, no patient was treated with methotrexate, chlorambucil, cyclophosphamide, and rituximab.

The most common complications caused by pemphigus treatment include oral candidiasis, weight gain, Cushingoid appearance, bacterial infections, gastritis, osteoporosis, avascular necrosis, necrosis, cataract, and glaucoma [2, 19, 20]. In our study, azathioprine was the most common agent used in our patients, followed by mycophenolate mophetil, cyclosporine, and intravenous immunoglobulin (IVIg). However, no patient was treated with methotrexate, chlorambucil, cyclophosphamide, and rituximab.

The authors declare no conflict of interest.

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