

Sporadic Pemphigus Foliaceus in a 75-year-old Filipino female: A Case Report

Joshua A. Arcaira, M.D.,¹ Lucia Lourdes O. Castro-Forés, M.D., FPD² Katrina M. Canlas-Estrella, M.D., FPDS³ Ma. Jasmin J. Jamora, M.D., FPDS⁴

¹Second Year Resident, Department of Dermatology, Makati Medical Center, Philippines

²Consultant, Department of Dermatology, Makati Medical Center, Philippines

³Immunodermatology Consultant, Department of Dermatology, Makati Medical Center, Philippines

⁴Dermatopathology Consultant, Department of Dermatology, Makati Medical Center, Philippines



Abstract

Pemphigus Foliaceus (PF) is an uncommon autoimmune blistering disease with male predominance in Japanese literature, and onset within the 4th – 6th decades of life. We report a case of an elderly Filipino woman who presented with a one-year history of crusted erosions on the face, trunk, and extremities. Most lesions began as erythematous patches and plaques that would progress into flaccid bullae and would easily erode. She was only maintained on irbesartan, atorvastatin, and multivitamins daily. There was no known familial history of blistering diseases. Skin punch biopsy revealed a vacuolar interface dermatitis and direct immunofluorescence showed granular intercellular staining of IgG and C3, anti-nuclear antibody staining of IgG, and granular vascular staining of fibrinogen, consistent with Pemphigus group. Anti-nuclear antibody level was negative and serum ELISA levels showed positivity to only Desmoglein 1, and negative for Desmoglein 3—thus, clinching the diagnosis of PF. She was started on oral prednisone therapy at 0.75mg/kg/day and azathioprine 50mg/day, along with ranitidine, loratadine, diphenhydramine, calcium + vitamin D3 supplements, and a topical compounding of clobetasol propionate cream, fusidic acid cream, and a bland emollient. Clinical improvement was observed at one month of treatment where most lesions became darkly erythematous to hyperpigmenting thin plaques and no recurrence of erosions. Prednisone dosage was then slowly tapered to 5mg/day of prednisone at 4 months of treatment. No untoward events developed during the course of therapy except for one episode of upper respiratory tract infection which resolved after one week of Cefixime. Only 100 cases of PF were reported in the Philippine Dermatological Society Health Information System from 2011 to 2018 with female to male ratio of 2:3. This report aims to elucidate the importance of prompt recognition and treatment of this condition to achieve good prognosis and low morbidity.



Biography:

Dr. Joshua A. Arcaira is the incoming deputy chief resident at Makati Medical Center, Department of Dermatology in the Philippines. He completed his bachelor's degree in Human Biology at De La Salle University - Manila with an honorable mention and graduated among the top 25th percentile of his batch in medical training at De La Salle Health Sciences Institute College of Medicine. Currently, he is interested in pursuing the subspecialties of Dermatologic Surgery and Dermatopathology.

[21st World Dermatology 2020](#); Tokyo, Japan - June 22 - 23, 2020.

Abstract Citation:

Joshua A. Arcaira, Sporadic Pemphigus Foliaceus in a 75-year-old Filipino female: A Case Report, World Dermatology 2020, 21st World Dermatology Congress; Tokyo, Japan – June 22 – 23, 2020

(<https://worlddermatology.conferenceseries.com/abstract/2020/sporadic-pemphigus-foliaceus-in-a-75-year-old-filipino-female-a-case-report>)