

Case Report

Open Access

Localised cutaneous Amyloidosis Revealed by Immunohistochemistry: an Interesting Diagnostic Tool

Derqaoui Sabine^{1*}, Essaoudi Mohamed Amine² and Elktaibi Abderrahim²

¹Department of Pathology, Mohamed V Military Instruction Hospital, Rabat, Morocco

²Faculty of Medicine and Pharmacy of Rabat, Mohamed V University, 10100, Rabat, Morocco

*Corresponding author

Derqaoui Sabine, Department of Pathology, Mohamed V Military Instruction Hospital, Rabat, Morocco. E-Mail: s.derqaoui@gmail.com

Received: April 03, 2022; **Accepted:** April 11, 2022; **Published:** April 25, 2022

Keywords: Cutaneous Amyloidosis, Histopathology, Immunohistochemistry

Case Presentation

A 85-year-old woman had a lightly pruritic firm plaque on the forearm for 28 years. She had no major medical trouble and she didn't receive any medical treatment before visiting dermatology's department. Biopsy specimen from the lesion revealed an amorphous a cellular eosinophilic deposits (black star) in the papillary dermis (Figure 1 A and B). The overlying epidermis was of normal thickness without hyperkeratosis (Figure 1 A and B). The deposits were negative for Congo red staining (Figure 1C), without typical green birefringence on polarizing microscopy (Figure 1D). On immunohistochemistry, they stained diffusely positive with pan cytokeratin AE1/AE3 (Figure 2).

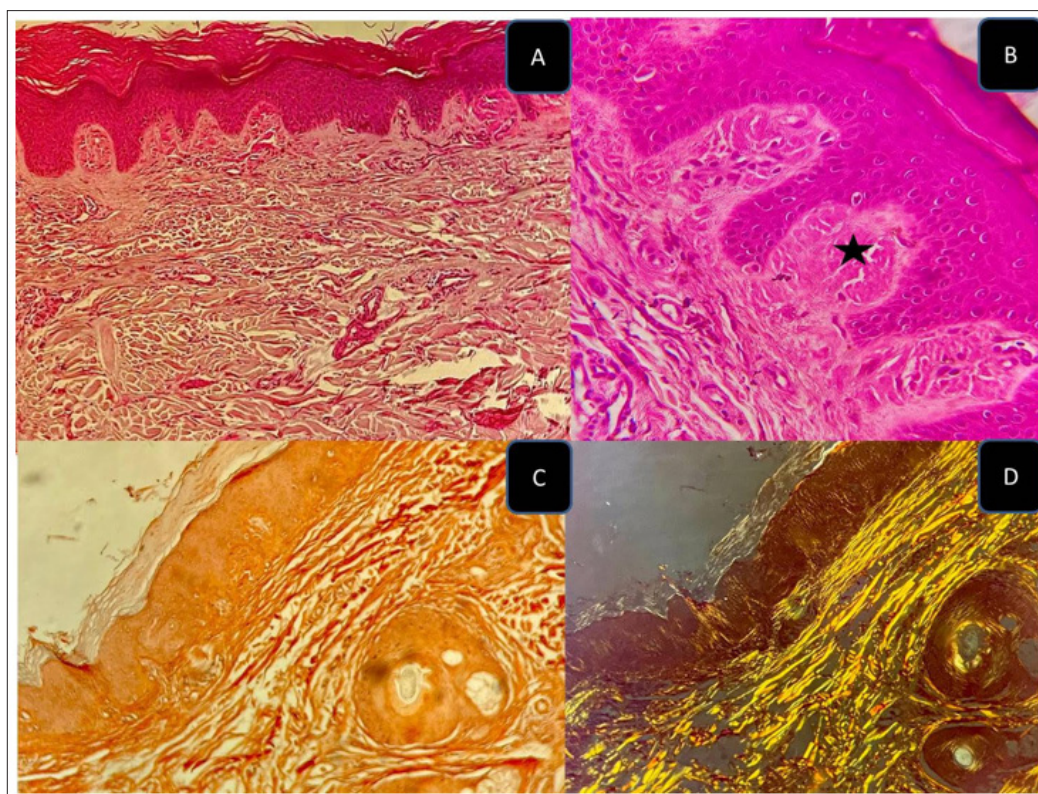


Figure 1: Amyloid Dermal Deposits : HE Stainig at Low Power (A) and High Power (B). These Deposits are Negative for Congo Red (C), Without Typical Green Birefringence on Polarizing Microscopy (D)

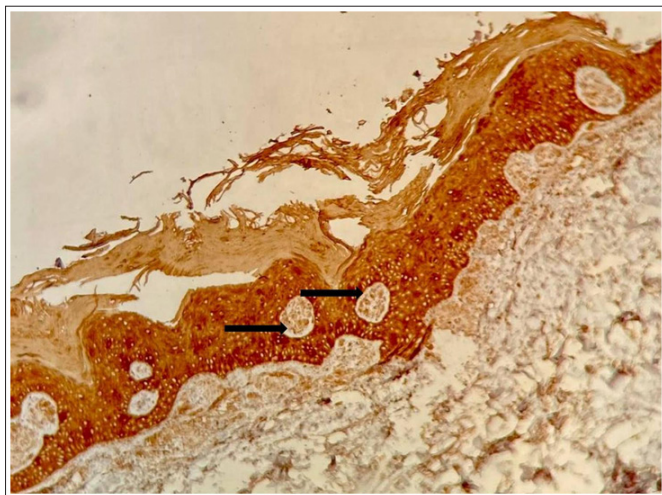


Figure 2: Immunohistochemistry : Positive Staining for pan cytokeratin AE1/AE3

Discussion

Localised cutaneous amyloidosis (LCA) is defined by the deposition of amyloid in the skin with the absence of systemic involvement. LCA can be divided into: primary and secondary LCA. The latter, is observed in several inflammatory and neoplastic skin disorders such as seborrheic keratosis, Bowen's disease and basal cell carcinoma [1,2]. Primary cutaneous amyloid is a chronic pruritic condition [1]. On histology, PCA shows amorphous eosinophilic deposits in the papillary dermis. Definite diagnosis requires special stains and sometimes immunohistochemistry. Amyloid is usually positive with Congo red staining; and shows green birefringence in polarizing microscopy [3]. However, Congo red staining might be negative; as in the present case. Immunohistochemistry for cytokeratins (high molecular weight cytokeratins and ck5/6) represents a useful diagnostic tool in LCA, because Congo red staining may not detect small deposits. In the study of et al. The authors recommend performing IHC for high molecular weight cytokeratins to exclude the diagnosis, when Congo red is negative [4]. These findings confirm the origin of amyloid: epidermal keratins due to epidermal long term damage [3].

Declarations

The authors declare no competing interests

Funding Resource: Not Applicable

Informed consent was obtained

References

1. Sari Aslani F, Kargar H, Safaei A, Jowkar F, Hosseini M, et al. (2020) Comparison of Immunostaining with Hematoxylin-Eosin and Special Stains in the Diagnosis of Cutaneous Macular Amyloidosis. *Cureus* 12: e7606.
2. Chang YT, Liu HN, Wang WJ, Lee DD, Tsai SF (2004) A study of cytokeratin profiles in localized cutaneous amyloids. *Archives of Dermatological Research* 296: 83-88.
3. Kentaro Izumi, Ken Arita, Keita Horie, Daichi Hoshina, Hiroshi Shimizu (2014) Localized Cutaneous Amyloidosis Associated with Poikiloderma with Mucin Degeneration. *Acta Derm Venereol* 94: 225-226.
4. Ahmed Abdullah Alhumidi, Amany Abdulgader Fathaddin (2015) The Utility Of Congo Red Stain And Cytokeratin Immunostain In The Detection Of Primary Cutaneous Amyloidosis. *The Internet Journal of Pathology* 17: 1-7.