



Successful Treatment of Lipoid Proteinosis in a Young Adult with Isotretinoin

Hamad Alfahaad^{a++*}, Hamza Alshehri^{b#}, Ebtisam Aljerayan^{b†},
Albaraa Alwabel^{b†} and Abdullah Mushabeb Alfarwan^{b‡}

^a Department of Medicine, College of Medicine, Najran University, Saudi Arabia.

^b Ministry of Health, Saudi Arabia.

Authors' contributions

This work was carried out in collaboration among all authors. All authors read and approved the final manuscript.

Article Information

DOI: 10.9734/JPRI/2023/v35i107351

Open Peer Review History:

This journal follows the Advanced Open Peer Review policy. Identity of the Reviewers, Editor(s) and additional Reviewers, peer review comments, different versions of the manuscript, comments of the editors, etc are available here: <https://www.sdiarticle5.com/review-history/99557>

Case Report

Received: 25/02/2023

Accepted: 27/04/2023

Published: 01/05/2023

ABSTRACT

Lipoid proteinosis (LP) is an extremely rare autosomal recessive genodermatosis, characterized by persistent hoarseness and classical skin lesions. Amorphous hyaline depositions may affect any organ, but mainly the face and extremities. These depositions cause limitations in tongue movement, and hence, difficulty speaking. Biopsies of skin lesions usually show massive deposits of homogeneous, acidic, hyaline-like material at the epidermal junctions and superficial dermis that stains positive for periodic acid-Schiff stain (PAS-D). At present, there is no specific treatment for this extremely rare disease, but some studies have reported that treatment with acitretin and etretinate results in some improvement. This paper reports on the effectiveness of isotretinoin for the first time for treatment of this disease.

⁺⁺ Associate Professor and Consultant Dermatologist;

[#] Consultant Dermatologist & Dermatopathologist;

[†] Consultant Dermatologist;

[‡] Health Care Manager;

*Corresponding author: E-mail: hamadyam@gmail.com;

Keywords: Lipoid proteinosis; hoarseness; periodic acid schiff; isotretinoin.

1. INTRODUCTION

Lipoid proteinosis (LP), or Urbach-Wiethe disease, is a very rare autosomal recessive skin disease that initially presents with laryngeal deposition of amorphous vitreous material, leading to persistent hoarseness at birth or in early childhood, as expressed by a soft cry [1] that is accompanied by classic skin changes such as scarring, fragility, discomfort, papules and/or nodules infiltrating the eyelid [2].

The deposition of an amorphous vitreous substance on the tongue usually limits the tongue's movements and causes speech difficulties. In addition, these infiltrates can lead to the appearance of papules and scars resembling chickenpox [3]. It primarily affects the face and extremities. However, any organ in the body can be affected, such as the respiratory system, which causes airway obstruction, and the central nervous system, resulting in seizures [4].

There is currently no specific established therapeutic option to treat lipoid proteinosis, but this paper reports a positive response with treatment by isotretinoin.

2. CASE PRESENTATION

A 20-year-old man born to first degree consanguineous parents (48-year-old father and 43-year-old mother) presented with hoarseness that was noticed by his mother at birth and was clear when he was crying. This hoarseness has persisted until the current date. At the age of 17, recurrent skin lesions started to appear every 2-4 months on his face and upper extremities that left disfiguring lesions as scars when healed.

Clinical examinations revealed that the patient had hoarseness while talking. On the first examination, lesions (mainly on the face) were described as multiple papulopustular with atrophic scars similar to chickenpox (Fig. 1). Areas of friction (elbows, knees, hands, dorsae) developed verrucous hyperkeratotic plaques (Fig. 2). Typical beaded papules (monilia ornaments blepharosis) were also present on the eyelid margins (Fig. 3), and the patient had difficulty protruding his tongue (Fig. 4).



Fig. 1. Papulopustular lesions with atrophic scars



Fig. 2. Difficulty in tongue protrusion

The patient underwent ENT consultation, which reported vocal cord thickening and hyaline depositions at the larynx and posterior pharyngeal wall, oral cavity, and tongue. Laboratory investigations included a full blood count, measurement of serum glucose level, hepatic and renal function tests, and lipid profile, all of which were within normal levels. A skin biopsy from the upper back showed massive deposits of homogeneous, eosinophilic, hyaline-like material at the dermal-epidermal junctions and at the superficial dermis (H&E) that stains positive for periodic acid-Schiff stain (PAS-D) (Fig. 5).

Based on clinical and histopathological findings, the patient was diagnosed with a case of lipoid proteinosis. He was started on isotretinoin 20 mg/day in August 2021, with follow up examinations every two months. The drug dose was adjusted according to body weight at every follow-up visit. Baseline investigations were also conducted and repeated every two months. Six months following treatment initiation, the hoarseness partially improved, and the tongue could be partially protruded. The skin lesions

also decreased in number (Fig. 6). ENT follow up reported decreased vocal cord thickening and decreased hyaline deposits in the larynx and oral cavity. One year after follow-up, no new lesions had appeared, and a 70% improvement in voice quality was noted.



Fig. 3. Typical beaded papules occur on the eyelid margins "moniliform blepharitis"



Fig. 4. Verrucous hyperkeratotic plaques

3. DISCUSSION

Lipoid proteinosis is a rare autosomal recessive disease caused by a homozygous mutation of the Q276X in the ECM1 gene, resulting in deposition of hyaline-like material in multiple tissues [3]. Usually, hoarseness during crying in infancy is the initial symptom noticed, along with acne like cutaneous lesions, papulopustular lesions, beaded papules along the eyelids, and oral mucosal lesions appearing later in childhood, often after the age of 10 [5]. In agreement with this, the case study presented here shows a history of soft crying since birth, which continued as hoarseness when talking in early childhood. Skin lesions also appeared at age of 17 years as recurrent skin lesions every 2-4 months on his face and upper extremities that left disfiguring scars.

Cutaneous lesions of LP mainly affect sun exposed areas: the face, elbows and hands. They appear characteristically in the form of thickened facial skin with a waxy texture covered with yellow, infiltrated, flat plaques, and papules among disfiguring scars [6]. In the present case,

skin lesions appeared on the face and the upper extremities during puberty, which left post-healing scars in the form of disfiguring lesions. Infiltration of oral mucosa, the vocal cords and frenulum by an amorphous hyaline-like material also characterizes LP lesions, leading to hoarseness and restricted tongue protrusion. In addition, attacks of respiratory distress might occur, whereby the airway is affected [5].

Histopathological findings of LP include deposition of homogeneous, eosinophilic, hyaline-like material at the dermal-epidermal junctions and superficial dermis. In addition, the hyaline like-material is positive on periodic acid-Schiff stain (PAS-D) [7]. The skin biopsy from the upper back of the present case showed massive deposits of homogeneous, eosinophilic, hyaline-like material at the dermal-epidermal junctions and at the superficial dermis (H&E). The hyaline like-material was positive on periodic acid-Schiff stain (PAS-D).

Because of the rarity of lipoid proteinosis, there is a serious limitation in terms of trials of therapeutic options. Of those that do exist, some studies have reported good results with the treatment of acitretin and etretinate [8-10]. The present case was treated with isotretinoin and showed impressive improvement.

It is thought that retinoids modulate the metabolism of the connective tissue matrix of the dermal-epidermal junction and the dermis [11]. At present, no previous studies have been published to support the usefulness of isotretinoin on both cutaneous lesions and hoarseness [12]. Therefore, isotretinoin may be a superior treatment to acitretin and etretinate, the former of which has the qualities of a tolerance, ease of dose modification and minimal side effects over the long term compared to other variants of retinoid. In a similar case that used acitretin, there was no obvious cutaneous improvement; however, there was significant improvement in hoarseness [9]. In another two young girls in Egypt treated with acitretin, there was complete remission of cutaneous lesions and improvement of the hoarseness after one year [13]. Since there is limited experience with isotretinoin in treating LP, future studies are advised to evaluate and monitor its effects on LP patients despite our experience with this case providing impressive results.

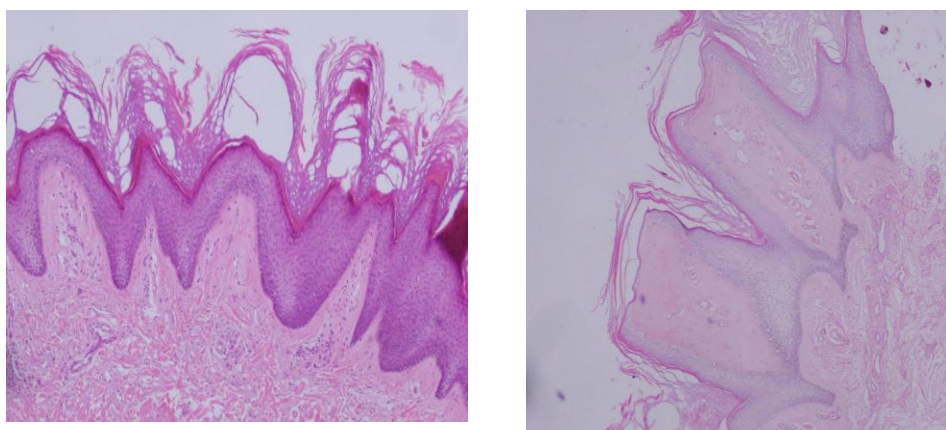


Fig. 5. Massive deposits of homogeneous, eosinophilic, hyaline-like material at the dermal-epidermal junctions and at the superficial dermis (H&E), that stains positive for periodic acid-Schiff stain (PAS-D)



Fig. 6. Post isotretinoin treatment showed improvement of skin lesions.

4. CONCLUSION

The efficacy of isotretinoin in the treatment of lipoid protenosis and the advantages of feasibility and shorter half-life make it superior to both acitretin and etrinitate, such that it can be safely discontinued immediately if serious complications or lack of response occur.

CONSENT

As per international standard or university standard, patient(s) written consent has been collected and preserved by the author(s).

ETHICAL APPROVAL

Ethical approval has been collected from IRB.

COMPETING INTERESTS

Authors have declared that no competing interests exist.

REFERENCES

1. Urbach E, Wiethe C. Lipoidosis cutis et mucosae. Virchows Archiv Für

- Pathologische Anatomie Und Physiologie Und Für Klinische Medizin. 1929;273(2): 285–319.
Available:<https://doi.org/10.1007/bf02158983>
2. Hamada T. Lipoid proteinosis. Clinical and Experimental Dermatology. 2002;27(8): 624–629.
Available:<https://doi.org/10.1046/j.1365-2230.2002.01143.x>
 3. Van Hougenhouck-Tulleken W, Chan I, Hamada T, Thornton H, Jenkins T, McLean WHI, McGrath JA, Ramsay M. Clinical and molecular characterization of lipoid proteinosis in Namaqualand, South Africa. British Journal of Dermatology. 2004;151(2):413–423.
Available:<https://doi.org/10.1111/j.1365-2133.2004.06076.x>
 4. McGrath JA. Lipoid proteinosis. Neurocutaneous Syndromes. 2015;317–322.
Available:<https://doi.org/10.1016/b978-0-444-62702-5.00023-8>
 5. L Ramsey M, Tschen JA, Wolf Jr JE. Lipoid proteinosis. International Journal of Dermatology. 1985;24(1):230–232.
Available:<https://doi.org/10.1111/j.1365-4362.1985.tb05443.x>
 6. Hu S, Kuo T.-tong, Hong HS. (). Lipoid proteinosis: Report of a possible localized form on both hands and wrists. International Journal of Dermatology. 2005;44(5):408–410.
Available:<https://doi.org/10.1111/j.1365-4632.2005.02031.x>
 7. Hamada T. Lipoid proteinosis maps to 1q21 and is caused by mutations in the extracellular matrix protein 1 gene (ECM1). Human Molecular Genetics. 2002;11(7): 833–840.
Available:<https://doi.org/10.1093/hmg/11.7.833>
 8. Toosi S, Ehsani AH. Treatment of lipoid proteinosis with acitretin: A case report. Journal of the European Academy of Dermatology and Venereology. 2009; 23(4):482–483.
Available:<https://doi.org/10.1111/j.1468-3083.2008.02928.x>
 9. Gündüz Ö, Şahiner N, Atasoy P, Şenyücel Ç. Acitretin treatment for lipoid proteinosis. Case Reports in Dermatological Medicine. 2012;2012:324506.
Available:<https://doi.org/10.1155/2012/324506>
 10. Gruber F, Manestar D, Stasic A, Grgurevic, Z. Treatment of lipoid proteinosis with etretinate. Acta dermato-venereologica. 1996;76(2):154–155.
Available:<https://doi.org/10.2340/0001555576154155>
 11. Daly TJ, Weston WL. Retinoid effects on fibroblast proliferation and collagen synthesis in vitro and on fibrotic disease in vivo. Journal of the American Academy of Dermatology. 1986;15(4):900–902.
Available:[https://doi.org/10.1016/s0190-9622\(86\)70248-x](https://doi.org/10.1016/s0190-9622(86)70248-x)
 12. Nickle SB, Peterson N, Peterson M. (). Updated Physician's Guide to the Off-label Uses of Oral Isotretinoin. The Journal of Clinical and Aesthetic Dermatology. 2014; 7(4):22–34.
 13. Bakry OA, Samaka RM, Houla NS, Basha MA. Two Egyptian cases of lipoid proteinosis successfully treated with acitretin. Journal of Dermatological Case Reports. 2014;8(1):29–34.
Available:<https://doi.org/10.3315/jdcr.2014.1168>

© 2023 Alfahaad et al.; This is an Open Access article distributed under the terms of the Creative Commons Attribution License (<http://creativecommons.org/licenses/by/4.0>), which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

Peer-review history:

The peer review history for this paper can be accessed here:
<https://www.sdiarticle5.com/review-history/99557>