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Simultaneous occurrence of insulin-derived amyloidosis and acanthosis nigricans at the abdominal site of insulin injection

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1 Introduction

- 2 The most common complications of subcutaneous insulin injection are lipoatrophy,
- 3 lipohypertrophy, and localized allergic reactions. More rarely, skin abscesses, insulin edema,
- 4 insulin allergy, acanthosis nigricans (AN), and insulin-derived amyloidosis have been reported
- 5 at insulin injection sites(1). We report a patient with concomitant insulin-derived amyloidosis
- 6 and AN at the abdominal site of insulin injection.

7 Case report

- 8 A 68-year-old female patient with a history of hypertension, liver transplant for nonalcoholic
- 9 steatohepatitis cirrhosis and 30-year history of type 2 diabetes presented with growing
- 10 asymptomatic plaques on the abdomen. Her anti-rejection treatment consisted of
- mycophenolate mofetil and tacrolimus. Insulin glargine and insulin lispro were started 10 years
- before the visit. Abdominal plaques appeared at injection sites but treatment was continued 1.5
- years prior to presentation when treatment was changed to insulin glargine and insulin aspart
- due to worsening blood glucose control.
- Physical examination showed 2 large hyperkeratotic, hyperpigmented and indurated plaques
- symmetrically located on either side of the umbilicus. No masses were palpable. Along the
- superior edge, the plaques were confined by the liver transplantation scar (Figure 1a).
- 18 Examination of the back of the neck and armpits revealed AN.
- 19 Histologic examination of the abdomen showed hyperkeratosis and papillomatosis with mild
- 20 acanthosis, confirming AN (Figure 2a). Examination of the dermis revealed masses of
- 21 eosinophilic and amorphous material. The deposit stained orange-red with Congo red (Figure
- 22 2b) and showed green birefringence under polarized light (Figure 2c) that is diagnostic for
- amyloidosis. The amyloid deposit showed positive staining with anti-insulin and anti-serum
- amyloid A (SAA) antibodies.
- 25 Based on these findings, a diagnosis was made of insulin-derived amyloidosis associated with
- 26 AN at the site of insulin injection.
- 27 Surgical excision was considered, but not considered feasible in view of the absence of a
- palpable mass. Clobetasol propionate cream applied once daily resulted in no clinical change.
- 29 Topical adapatene 0,1% resulted in significant reduction in appearance of clinical plaques.
- 30 Induration however persisted (Figure 1b).

31 Discussion

- 32 Amyloid deposition at the injection site of porcine insulin was described for the first time in
- 1988 by Dische *et al.* (2). Since then, at least 86 cases of insulin-derived amyloidosis have been
- reported (3). Any type of insulin can cause insulin amyloidosis (3).
- 35 The usual clinical presentation is a rigid subcutaneous mass at the site of insulin injection.
- Nagase et al. reported 2 cases of insulin-derived amyloidosis without a palpable mass,
- 37 suggesting that its incidence may be underestimated(4).
- Patients often show poor glycemic control because of local degradation of insulin in amyloid
- or trapping of insulin in amyloid fibrils after injection(5). Patients may also present with severe
- 40 hypoglycemia due to erratic absorption from the affected site.

- Diagnosis is confirmed by histopathology with the presence of amyloid positively stained by
- an antibody against insulin(6).
- Lipohypertrophy is the most common differential diagnosis(6). This localized reaction site is
- less firm and solid than an amyloid tumor and regresses after cessation of insulin injections.
- 45 The treatment for insulin-derived amyloidosis is to avoid the injections at the sites of
- amyloidosis or surgery(6). Surgical excision can result in improved glycemic control in cases
- 47 of localized amyloid tumors.
- 48 AN at the site of insulin injection is another rare complication of insulin injection, with a dozen
- 49 cases reported(7). The patients are mostly men. The abdomen is the most commonly affected
- site. For 2 patients, plaques disappeared after rotating the injection site. The pathophysiology
- remains unknown, but may be the same as in benign AN. High concentrations of insulin in the
- 52 epidermis may bind to IGF-1 (insulin-like growth factor 1) receptors on keratinocytes,
- stimulating proliferation and inducing hyperkeratosis(8).
- To our knowledge, this is the third reported case of a simultaneous occurrence of insulin-
- derived amyloidosis and AN at the site of insulin injection. The first case reported a 59-year-
- old man whose dermatosis was located on the abdomen and was secondary to the use of human
- 57 insulin(9). The second case, published in 2014, reports on a 54-year-old man, and the
- 58 dermatosis also located on the abdomen. The insulin used was recombinant human insulin
- 59 without further details(10). The clinical presentation was similar to our patient.
- 60 Because our patient was seen many years after occurrence of AN and insulin-derived
- amyloidosis, it is difficult to know which pathology occurred first. We offer two hypotheses as
- 62 to why these two conditions might co-occur. Firstly, it is possible that the presence of
- 63 induration from amyloid led to more superficial insulin injection, with higher concentrations
- of insulin in the epidermis resulting in AN. Another possibility is that amyloid hindered
- absorption of insulin and poorer glycemic control, requiring increased doses of insulin and
- development of AN.
- 67 <u>Conclusion</u>
- 68 Although insulin-derived amyloidosis and AN are uncommon complications of insulin
- 69 injection, dermatologists must be aware of them. Prevention of these complication can be
- achieved by rotating insulin injection sites, self-monitoring, and use of fine needles.



Figure 1a: Hyperkeratotic plaques located on either side of the umbilicus



82 Figure 1b: Improvement after 4 months of topical retinoid treatment

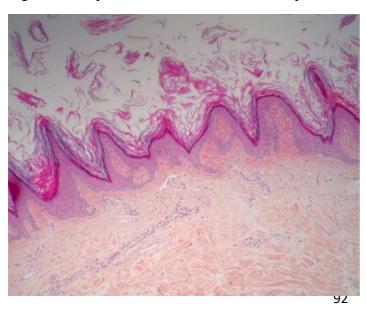


Figure 2a: Hyperkeratosis and papillomatosis (HES, x6,5)

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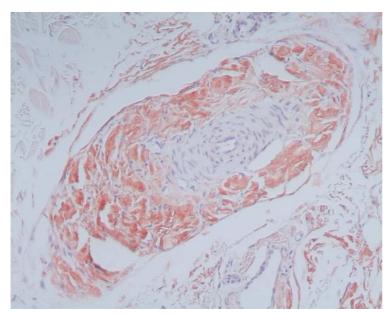


Figure 2b: Cutaneous amyloidosis : amyloid deposits Congo red positive (x15)

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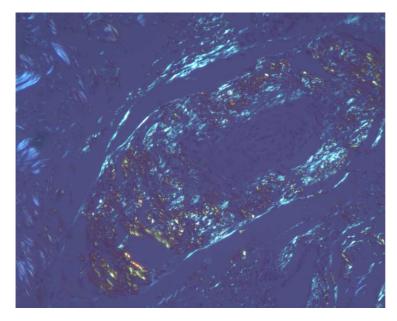


Figure 2c: Cutaneous amyloidosis : apple-green birefringence in polarized light (x15)

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