CASE SERIES

Prurigo pigmentosa following ketogenic diet and bariatric surgery: A growing association

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BACKGROUND

Prurigo pigmentosa (PP) is an uncommon inflammatory skin disorder. It is characterized by the development of erythematous urticated papules following a reticulate distribution. Although there is no clear cause for PP, multiple mechanical, inflammatory, and metabolic factors have been associated with the condition. One of these factors is the role of ketosis in inducing the eruption of PP.

Histopathologic examination varies during the different stages of PP and includes perivascular neutrophilic infiltrate, parakeratosis, spongiosis, and dyskeratotic cells in the epidermis, with later stages featuring melanophages in the superficial dermis. We report 4 new cases of PP from Saudi Arabia: 3 were temporally associated with initiating ketogenic diet, and 1 patient had undergone bariatric surgery 1 month before the onset of PP. We also review the literature regarding the association of ketosis, diet, and PP.

CASE REPORT

Patient 1

A 26-year-old woman, who was otherwise healthy, presented with itchy skin lesions that started 3 months after adhering to a ketogenic diet. Her weight at the initial visit was 82 kg. A skin examination found erythematous papules coalescing to form plaques arranged in a reticulated configuration involving the chest, with intermammary and submammary involvement, and the upper back (Fig 1). A skin biopsy found an acanthotic epidermis with parakeratosis and few dyskeratotic cells overlying a mild/focal vacuolar interface change at the dermoeidermal junction (Fig 2). The dermis showed a superficial and mid-dermal perivascular lymphocytic infiltrate with few eosinophils. The overall clinical and histologic findings were consistent with PP.

Patient 2

A 20-year-old healthy man developed a rash 2 months after starting the ketogenic diet. His weight was 78 kg, and upon examinations the patient had bilateral erythematous papules and plaques on the sides of the neck (Fig 3). Based on the results of skin biopsy and clinical presentation, the findings were in keeping with a diagnosis of PP.

Patient 3

A 22-year-old woman presented with a history of skin lesions appearing on the chest and neck. She started the ketogenic diet 4 months prior. Her weight was 84 kg at presentation, and a physical examination found grouped erythematous papules coalescing into plaques, mainly on the intermammary area with less involvement of the neck (Fig 4). The patient refused a skin biopsy for cosmetic concerns, and a diagnosis of PP was made based on clinical grounds owing to the typical presentation of the eruption.

Patient 4

A 24-year-old woman had itchy skin eruption 5 weeks after bariatric surgery. She weighed 79 kg, and clinical examination found scattered erythematous papules on the sides of the face, shoulders, and chest (Fig 5). The patient refused biopsy, but similar to the previous patient, a diagnosis of PP was based...
on the clinical presentation that was reinforced with the patient’s response to treatment.

All patients were treated with 100 mg/d of doxycycline for 4 weeks, and the ketogenic diet was discontinued for the 3 patients who were on it prior to the presentation. A clinical response was noted as early as 2 weeks from the initiation of treatment, which resolved with residual post inflammatory hyperpigmentation. There was no recurrence of PP at the 2-month follow-up. At 1-year follow up, patient 2 had a recurrence of PP that was in association with resuming the ketogenic diet (Fig 6). This patient also reported recent initiation of metformin, as he was found to be prediabetic.

DISCUSSION

The first report of PP was in 1971 by Nagashima et al.1 It disproportionally affects females in comparison to males, with a ratio of 2:1.2 The eruption of PP is characterized by erythematous pruritic coalescing of papules arranged in a reticulate configuration that is transient and later resolves with postinflammatory hyperpigmentation.3 Variations of the typical morphology of PP include pustular, bullous, and urticarial forms.2 Although most cases have been described initially in Japan, there is an increase in reporting of PP worldwide, which is hypothesized to be related to the increase in awareness of the condition.3 Histopathologic features of PP depend on the stage at which the lesions are biopsied. They typically show a perivascular neutrophilic infiltrate in the early stages, which later evolve to feature necrotic keratinocytes and spongiosis of the epidermis. They ultimately lead to the presence of melanophages and lymphocytic infiltrate of the superficial dermis.4

The exact cause of PP remains unknown. However, reports of the condition developing after certain exposures along with the nature of the disease’s behavior has led many investigators to hypothesize explanations to the appearance of such eruptions. Some examples include sweating,1,2 clothing friction, rubbing,3 contact dermatitis, and conditions like atopy and pregnancy.5 The metabolic associations with PP include diabetes mellitus,6 fasting,7 ketonemia,7 and dieting.8

A ketogenic diet consists of reducing the carbohydrate content in the diet (usually to less than 50 g/d) while increasing the fraction of fat and protein intake. There is resurgence in the interest to adopt this diet regimen because of the growing reports of its beneficial effect in areas like weight control, diabetes, and cardiovascular disease among others.9

Fig 1. PP: clinical presentation of patient 1. Reticulated erythematous papules and plaques on the chest.

Fig 2. PP: histopathologic examination of skin lesions of patient 1. Parakeratosis, acanthosis, few dyskeratotic cells, mild vacuolar change at the dermoepidermal junction, and superficial perivascular lymphocytic infiltrate.

Fig 3. PP: clinical presentation of patient 2. Erythematous papules and plaques on the neck.
Ketosis has frequently been linked to PP development in various reports.2,3,10 One of the earliest to note correlation was Teraki et al.7 They reported 10 cases, of which, 8 were found to have ketosis by laboratory findings precipitated by either dieting, loss of appetite, or insulin-dependent diabetes mellitus.7 Other conditions that have been reported to lead to ketosis included obesity,11 bariatric surgery,12 and hyperemesis gravidarum of pregnancy.5 It is important to consider PP, as its initial presentation may not be a straightforward diagnosis. For instance, a report of a patient that was initially thought to have an urticarial eruption later had PP diagnosed, caused by ketosis secondary to anorexia nervosa.13 Previous reports have also supported the association between anorexia nervosa and PP.8 There are frequent reports of ketosis secondary to diabetes mellitus.6 Similarly, and of interest to our case, diet is also linked to the development of PP caused by ketosis.2,7,8,14,15 In a recent report by Almaani et al,15 PP development in 2 of 3 Jordanian patients was linked to fasting during the month of Ramadan, whereas the other case was a result of strict dieting.15 Although these reports support an association between ketosis and the onset of PP, it is still not considered an established causative factor for the disease.2 It has been hypothesized that ketone bodies play a role in inducing perivascular inflammation, mainly through neutrophils, thus explaining the positive response to medications suppressing neutrophilic-mediated processes like dapsone and tetracyclines.12 Another mechanism that has been hypothesized is the direct alteration of intracellular processes owing to their entry into cells.12

Treatment of PP classically consists of oral antibiotics, including minocycline,2 dapsone,2,11 and doxycycline.14 Typically, the treatment is successful, but hyperpigmentation may persist beyond the acute eruption. One consideration when treating patients with history of dieting prior to PP development is to reinstitute a balanced diet as an adjunct to medical therapies, as it was found to help in the treatment of the disease.2

We report on 3 patients who were on a ketogenic diet and 1 patient who had bariatric surgery, all of whom had PP a few months after the precipitating factor was initiated/ performed. All patients were treated with doxycycline along with discontinuation of the ketogenic diet, which was associated with resolution of PP. It is prudent for the dermatologist to consider PP in the assessment and evaluation of
patients with risk factors for ketosis development, including those on diet or other metabolic disorders. Particularly, PP should be considered in patients on a ketogenic diet who have cutaneous eruptions, especially with the growing interest in adopting such dietary regimens.

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