

A Case of Herpes Gestationis

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ÖZET

Herpes gestationis (HG), gebelikle görülen nadir bir cilt hastalığıdır. 60.000 gebelikte 1 sıklıkta görülmektedir. 24 yaşında 32 haftalık bir gebede HG tanısı konulmuş; klinik, laboratuvar ve doku bulguları yönünden incelenerek sonuçları tartışılmıştır.

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SUMMARY

Herpes gestationis (HG) is an uncommon blistering disease of pregnancy and the Puerperium. Its incidence has been estimated at about 1 in 60.000 pregnancies.

24-year old pregnant woman with herpes gestationis at 32 weeks gestation was seen in our clinic. Following case presentation, HG was discussed in all aspects.

Key Words: Herpes gestationis, pregnancy.

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Herpes gestationis (HG) is an uncommon blistering disease of pregnancy and the puerperium (1). Classically, it presents as an intensely pruritic erythematous vesiculobullous eruption that is often widespread and recurs in subsequent pregnancies. Its incidence has been estimated at about 1 in 60.000 pregnancies. Since most medical centers have only a limited experience with HG, we would like to present a case of HG seen in our clinic.

Report of a Case

A 24-year-old multigravida (G₇ P1051) presented to our clinic with complain of severe pruritis and cutaneous eruption at 32 weeks gestation (Fig 1 and 2). Skin changes first developed at 28 weeks gestation. Topical applications of corticosteroids and oral antihistaminic medication did not control her pruritis and skin lesions.

Her past medical history was unremarkable except in review of her past obstetric history revealed that she had had five first-trimester spontaneous abortions after a full-term spontaneous vaginal delivery.

On her physical examination, skin changes involved chiefly the abdomen, back and extremities and formed confluent edematous and erythematous pla-

ques with scattered bullae and vesicles. There were no systemic symptoms.

Hb was 10.2g/dl, WBC 6000/ml and platelets 24×10^3 /ml. FBS was 70 mg/dl. creatinin was 0,5 mg/dl. Liver function tests and thyroid function tests were normal. IgA 343 mg/dl, IgM 402 mg/dl, C₃ 144 mg/dl, C₄ 45.6 mg/dl and ANA was negative.

Skin biopsy revealed subepidermal bullae with partial destruction of the deep layers of the epidermis. There was an inflammatory infiltrate of lymphocytes, histiocytes and eosinophils below the epidermis and in the perivascular regions. Direct immunofluorescence of skin demonstrated IgA deposition along the BMZ (Fig 3).

Systemic corticosteroid treatment started at full dose (40 mg/day) and tapered gradually and maintained at a dose of 15 mg every other day. The response was prompt with symptomatic improvement within 48 hours followed rapidly by clinical sign of improvement.

The mother delivered a healthy child at term. The child did not develop any cutaneous eruption at all.



3 weeks after delivery cutaneous lesions disappeared leaving no traces, and did not recur during the next five months of observation. The steroid therapy was discontinued after delivery.

Comment

Herpes gestationis is a blistering disease of pregnancy and postpartum period. The time of onset is variable, although it is much more common to begin in the second or third trimester of pregnancy, with acute flares immediately after delivery or with the first menstrual period. It persists for several weeks to a few months. There is a tendency, for it to begin earlier in the course of subsequent pregnancies. There is history of at least one spontaneous abortion in one third of the cases.

Lesions may present as papules, urticarial plaques, vesicle, bullae, crusts, and excoriations. Preferred sites of involvement are the abdomen, particularly the periumbilical area, and the extremities.

Though the clinical picture, histopathological findings and the course of the disease may suggest the diagnosis of HG, it is sometimes difficult to differentiate this entity from dermatitis herpetiformis, erythema multiforme, bullous drug eruptions, bullous pemphigoid, and other dermatoses of pregnancy (3). The application of immunological techniques as additional diagnostic criteria has proved to be very useful in precisely delineating HG and separating it from other bullous diseases. Immunofluorescent (IF) investigations of perilesional and of normal appearing skin reveal the presence of the third complement (C₃), with or without IgG, in a linear band-like distribution along the basement membrane zone (BMZ). A circulating factor in the serum of these patients, termed the herpes gestationis factor (HGF), has also been shown (6). Other immunoreactants occasionally deposited along the BMZ included IgA, IgE, C₁, C₂, C₃, properdin and fibrinogen (7). It has been noted that there is a high frequency of HLA-B8 and DR3 and an increased incidence of autoimmune thyrotoxicosis in patients with HG (8).



There appears to be a considerable incidence of fetal morbidity and mortality associated with HG. In one study, of 39 cases of HG, there were eight, premature deliveries, three were eight premature deliveries, three stillbirths and one spontaneous abortion at 4 1/2 months gestation (7).

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