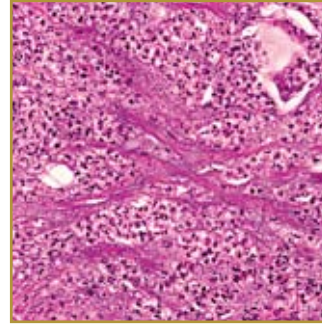


PYODERMA GANGRENOSUM



EPIDEMIOLOGY: Occurs in 1 out of 100,000 people each year

ETIOLOGY: Unknown; Trauma to skin induces new lesions

PATHOGENESIS: Neutrophil dysfunction (i.e., defects in chemotaxis or hyperreactivity) has been suggested

CLINICAL: Classic - Deep ulceration with a violaceous border that overhangs the ulcer bed. Atypical - Has a vesiculopustular component. Usually only at the border, erosive or superficially ulcerated

HISTOLOGY: Massive neutrophilic infiltration, hemorrhage, and necrosis of the overlying epidermis

PYODERMA GANGRENOSUM (PG) is an uncommon cause of ulcerations in the skin which may affect other organs. PG is classified into two types Classic or Atypical. Classic pyoderma gangrenosum consists of deep ulcerations with a purple border. The lesions often occur on the legs, but may occur anywhere on the body. Atypical pyoderma gangrenosum has more of a fluid filled component around the border. The lesions most often occur on the dorsa of the hands, the face or the extensor part of the forearms. There is no specific cause for PG, however skin trauma can result in new lesions on the skin pathology. Treatment options include wound care with antibacterial ointments, corticosteroids to relieve inflammation or immunosuppressants.

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