Neonatal bullous impetigo = اﻠوﻠﻴﺪ ﻋﻨﺪ اﻠﻔﻘاﻌﻴﺔ اﻠﻘوﺒاء

Impetigo.
Two clinical patterns of impetigo are recognized: bullous and non-bullous. Bullous impetigo is caused by S. aureus. Currently, in industrialized nations, non-bullous impetigo is most commonly caused by S. aureus and less often by group A streptococcus. Group A streptococcus remains a common cause of non-bullous impetigo in developing nations.

**Infections and Toxin Syndromes Involving the Skin and Soft Tissues Caused by Staphylococcus aureus**

Sites of colonization (carrier state)

- Anterior nares
- Throat
- Axillae, perineum
- Hands
- Involved skin in individuals with atopic dermatitis
Sites of colonization in neonates (and sites of infection)

- Skin
- Umbilicus
- Circumcision site
- Conjunctivae

Superficial pyodermas

- Primary pyodermas

- Skin
Impetigo

Bullous impetigo

Erythema
Botryomycosis

- Hair follicles

Superficial folliculitis (follicular or Bockhart impetigo)
Neonatal bullous impetigo = اﻠوﻠﻴﺪ ﻋﻨﺪ اﻠﻔﻘاﻌﻴﺔ اﻠﻘوﺒاء

Folliculitis (sycosis barbae)

Furuncle (boil)
Carbuncle

- Intertriginous sites

Perianal dermatitis

- Digital infections
Paronychia

Blistering distal dactylitis

- After skin disruption
Neonatal bullous impetigo

Trauma (physical, thermal)

Foreign body (intravascular catheter, prosthetic device)

- Secondary pyoderma
- Impetiginization of dermatoses such as atopic dermatitis
- Pyoderma associated with systemic disease
- Job syndrome
- Chédiak-Higashi syndrome
- Chronic granulomatous disease

Invasive infections

- Lymphangitis, lymphadenitis
- Erysipelas
- Cellulitis
- Streptococcal gangrene
- Pyomyositis
- Bacteremia, septicemia

Metastatic skin infections associated with bacteremia (often S.

- Abscesses (superficial and deep)
- Septic vasculitis (pustular purpura)
Purpura fulminans

- Disseminated intravascular coagulation associated with staphylococcal bacteremia
- Meningococcemia-like syndrome

Staphylococcal toxin-associated syndromes

- Staphylococcal scarlet fever
- Staphylococcal scalded-skin syndromes
- Staphylococcal toxic-shock syndrome

NON-BULLOUS IMPETIGO.
The non-bullous type of impetigo accounts for more than 70 percent of cases of this form of pyoderma. It occurs in children of all ages as well as in adults. Intact skin is usually resistant to colonization or impetiginization, possibly due to absence of fibronectin receptors for teichoic acid moieties on S. aureus and group A streptococcus. Production of bacteriocins, produced by certain S. aureus strains (phage group 71) and highly bactericidal to group A streptococcus, may be responsible for the isolation of only S. aureus from some lesions initially caused by streptococci.

History.

In a typical sequence, S. aureus spreads from nose to normal skin (approximately 11 days later) and then develop into skin lesions (after another 11 days). Lesions commonly arise on the skin of the face (especially around the nares) or extremities after trauma. Nasal carriers of S. aureus can present with a very localized type of impetigo confined to the anterior nares and the adjacent lip area;
pruritus or soreness of the area is a common complaint. Conditions that disrupt the integrity of the epidermis, providing a portal of entry of impetiginization, include insect bites, epidermal dermatophytoses, herpes simplex, varicella, abrasions, lacerations, and thermal burns.

Cutaneous Lesions.

The initial lesion is a transient vesicle or pustule that quickly evolves into a honey-colored crusted plaque that can enlarge to greater than 2 cm in diameter. Surrounding erythema may be present. Constitutional symptoms are absent. Regional lymphadenopathy may be present in up to 90 percent of patients with prolonged, untreated infection. Untreated, the lesions may slowly enlarge and involve new sites over several weeks. In some individuals, lesions resolve spontaneously. In others, the lesions extend into the dermis, forming an ulcer (see Staphylococcal Ecthyma).