

Interesting Mimics

Pyoderma Gangrenosum



- An inflammatory, noninfectious ulcerative skin disease associated with several systemic illnesses including IBD, hematologic malignancies, and collagen vascular diseases. Invocation of lesions often caused by trauma including needle sticks. Systemic steroids are the mainstay of therapy.

Neutrophilic Dermatosi



- Sweet's syndrome is an inflammatory disorder manifesting as multiple painful erythematous plaques associated with fever. Associated with IBD, strep PNA, drugs, and hematologic malignancies. Lesions may occur in areas of skin trauma. Treat with steroids.

Cutaneous T Cell Lymphoma



- Mycosis fungoides is the most common type of CTCL, accounting for 50% of all primary cases. Look for erythematous patches and plaques with fine scale and papules and nodules. Diagnosis requires a punch biopsy or an excisional biopsy.

Mucormycosis



- Caused by the Phycomycetes species including Mucor, Rhizopus, and Absidia fungi. Can appear in immunocompromised patients including HIV patients, neutropenic patients and IV drug users. Look for a black, necrotic ulcer with a violaceous border.

Actinomycosis



- Caused by anaerobic bacteria from the actinomycetales group, usually Actinomyces israelii. Look for painful erythematous or violaceous nodules with overlying sinus tracts, fistulae, and grainy exudates. Tend to present at the law line or the thoracic or abdominal wall.

Inpatient Cellulitis

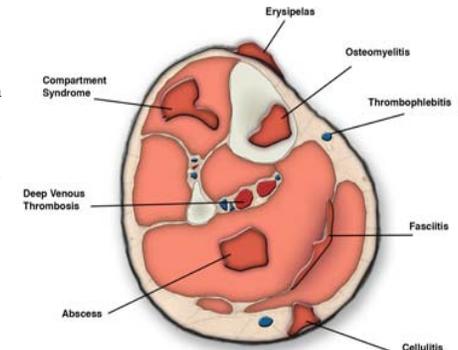
Clinical Evaluation

First, consider the differential with respect to the anatomic location and clinical context. An effective strategy is thinking about the contiguous anatomy

- **Limb**
 - **Infectious alternatives** include **necrotizing fasciitis**, bursitis, **osteomyelitis**, erythema migrans, compartment syndrome, **abscess**, ecthyma gangrenosum, **zoster**, toxic epidermal necrolysis
 - **Non-infectious alternatives** include compartment syndrome, **pyoderma gangrenosum**, nummular eczema, contact dermatitis, **deep venous thrombosis**, thrombophlebitis, **acute gout**, drug reaction, cutaneous T cell lymphoma, acute febrile neutrophilic dermatosis, lymphedema, erythema nodosum, stasis dermatitis, calciphylaxis, sarcoidosis, eosinophilic cellulitis, cutaneous GVHD
- **Trunk**
 - **Infectious alternatives** include **necrotizing fasciitis**, Fournier's gangrene, **focal abscess**, zoster, ecthyma gangrenosum
 - **Non-infectious alternatives** include **pyoderma gangrenosum**, nummular eczema, contact dermatitis, cutaneous T cell lymphoma, drug reaction, calciphylaxis, carcinoid erysipeloides, Paget disease of the breast, radiation recall
- **Face**
 - **Infectious alternatives** include necrotizing fasciitis, abscess, herpes simplex virus, zoster, cutaneous sinus tract, sinusitis, dental abscess, mucormycosis, actinomycosis
 - **Non-infectious alternatives** include eosinophilic cellulitis, contact dermatitis, angioedema, dermatomyositis, angiosarcoma, drug reaction, dacrocystitis (inflammation of lacrimal sac), cavernous sinus thrombosis

When considering a DDX...

- **CT** looking for gas is an excellent modality for **necrotizing fasciitis**; gas in absence of trauma or rim-enhancing collection is pathognomonic — *Raff, JAMA, 1016*
- **US** looking for **cutaneous abscesses** has a Sn 89-98% and Sp 64-88% — *Alsaawi, Eur J Em Med, 2015*
- **Osteomyelitis** can occur in the presence or absence of overlying skin ulceration although the presence of exposed bone has LR+ 9.2 — *Lipsky, Diab Metab Res Rev, 2012*
- For **oste**, the pooled Sn of **plain XR** is 54% and Sp 68%; LR+ is 2.3 and LR- is 0.63; the pooled Sn of **MRI** 77-100% and the Sp is 40-100%; LR+ 3.8 and LR- 0.14; **three-phase bone scan** has Sn 90% and Sp 46% — *Lipsky, Diab Metab Res Rev, 2012*
- **Blood cultures** are strongly recommended in very ill pts and those with neutropenia, malignancy, immunodeficiency—*HDSA, Clin Inf Dis, 2014*



- Compression US to r/o DVT is reasonable although the incidence of co-existent DVT in documented cellulitis is estimated at 3.1%; the rate of ipsilateral DVT is estimated at 0.5% - 0.75%; it is generally not recommended unless a high Wells, high clinical suspicion, or hypercoagulable state; can reconsider if no response to seemingly appropriate therapy — *Raff, JAMA, 2016; Gunderson, Thomb Res, 2013; Marz, BMC Inf dis, 2013*

