



BACILLARY

ANGIOMATOSIS

BARTONELLA INFECTION IN THE IMMUNOSUPPRESSED

Epidemiology

BA is most commonly seen in patients with acquired immunodeficiency syndrome (AIDS) and a CD4 count less than 50 cells/mm³, with an incidence of 1.2 cases per 1000 at-risk patients. Patients with other forms of immunosuppression, including patients with leukemia and recipients of organ transplants, have been reported. Uncommonly, human immunodeficiency virus-negative and nonimmunosuppressed persons develop BA. In the cases of BA in immunocompetent persons, however, although the lesions were proliferative vascular papules, they were limited in number, the affected persons had limited or no systemic involvement and a benign course. There is no predisposition in terms of race, sex, or age.

BACILLARY ANGIOMATOSIS AT

AGLANCE

- Etiology: *Bartonella henselae* and *B. quintana*
- Acquired from infected cats
- Most commonly in patients with acquired immunodeficiency syndrome but also in other forms of immunosuppression
- Pyogenic granuloma-like and subcutaneous nodules
- Hyperpigmented plaques in African Americans
- May be associated with hepatic and systemic lesions
- Treatment: erythromycin or doxycycline

Etiology and Pathogenesis

Both *B. henselae* (the CSD bacillus) and *B. quintana* (the agent of trench fever) have been identified as causative agents of BA. At one end of the clinical spectrum, classic CSD is seen in young, immunocompetent hosts as a limited infection. At the other end, BA is seen in patients who are severely immunocompromised as a systemic disease. Thus, it is the immunocompetence of the host and the bacterial load that dictates the clinical manifestations of the disease. Moreover,

immunocompromised
patients with BA develop an
angioproliferative
response in response to intracellular hypoxia. The presence of intracellular bacilli induces
hypoxia-inducible factor-1 that in turn induces vascular endothelial cell growth
factor,
leading to vascular proliferation.

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BA caused by *B.*
henselae
is acquired from infected cats and is a manifestation of CSD in the
immunocompromised
host.

Peliosis
hepatitis is exclusively associated with *B.*
henselae
infection.

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In
contrast,
patients with BA caused by *B.*
quintana
develop subcutaneous masses and
lytic
bone lesions.

Clinical Manifestations

CUTANEOUS LESIONS

The incubation period for BA is unknown. In AIDS patients, the clinical constellation includes
fever, cutaneous or subcutaneous vascular lesions, lymphadenopathy, and/or abdominal

symptoms. The most common

cutaneous

morphologies of BA are (1)

pyogenic

granuloma

-like lesions, (2) subcutaneous nodules, and (3)

hyperpigmented

indurated

plaques. The same patient may have several morphologies. Lesions resembling

pyogenic

granuloma

can range in size from 1 mm to many centimeters and are dusky-red in color with a

collarette

of scale and peripheral satellite lesions . The lesions are firm, bleed easily, and are often tender.

They occur on skin and mucosa. Subcutaneous nodules can range from distinct nodules to

diffuse swellings

with or without induration and are also often tender. Hyperpigmented plaques are most

commonly seen in African Americans with BA and are oval in shape; they are several

centimeters in diameter with indistinct borders. Large,

fungating

masses rarely occur. Patients with BA may have few to thousands of lesions with the number of

lesions gradually increasing over time. Additional

immunosuppression

with chemotherapeutic agents may be followed by a shower of

miliary

skin lesions.

Box 182-2 Differential Diagnosis of Bacillary Angiomatosis

Most Likely

- Pyogenic granuloma
- Kaposi sarcoma
- Angioma

Consider

- Chronic herpes simplex
- Hypertrophic scars
- Nocardiosis

Always Rule Out

- Amelanotic melanoma
- Squamous cell carcinoma
- Basal cell carcinoma
- Dermatofibroma protuberans
- Merkel cell carcinoma

RELATED PHYSICAL FINDINGS

In addition to cutaneous lesions, other organ systems may be affected. Hepatic and splenic vascular lesions can occur concomitantly with or independently of cutaneous

lesions and can be a cause of significant blood loss and anemia.

Bartonella

infection, especially that caused by B.

quintana

, can affect bone and soft tissues. Lesions of the central nervous system have been reported and can result in

neurologic

and psychiatric disorders.

Bacteremia

, chronic fevers, and pulmonary and gastrointestinal lesions have also been reported.

Radiologic

studies usually identify areas of systemic involvement. Ocular vascular proliferative lesions can produce loss of vision.

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There are several reports of patients with other cutaneous diseases concomitant with BA, as well as the simultaneous existence of BA and another infection within the same lesion. Several patients have been reported with both BA and Kaposi sarcoma. Cytomegalovirus, Epstein-Barr virus, Cryptococcus neoformans, and Mycobacterium avium-intracellulare have been found within lesions of BA.

Differential Diagnosis

Laboratory Findings

Patients with AIDS and BA are anemic and may have elevated liver function tests (characteristically, lactic acid dehydrogenase and alkaline phosphatase are more elevated than hepatocellular enzymes). Blood cultures are positive for Bartonella sp. in approximately one-half of BA patients. B. henselae and B. quintana can be cultured from skin lesions. The organisms grow slowly and may not be detected without prolonged culture (more than 1 month). PCR of affected tissue is virtually always positive if lesions are histologically

characteristic. The vast majority of cases are diagnosed histologically , with identification of the causative bacteria by Warthin -Starry staining.

Histopathology

Lesions of BA have the general features of a lobular capillary hemangioma (pyogenic granuloma) , but in contrast to a pyogenic granuloma , the endothelial cells are often larger and polygonal; they may have marked atypia . There is a prominent inflammatory

infiltrate, with significant numbers of neutrophils as well as leukocytoclastic debris . Polymorphonuclear leukocytes (PMNs) are scattered throughout the lesion, as opposed to classic pyogenic granuloma lesions in which the PMNs are at or near the surface, even if the pyogenic granuloma is eroded and impetiginized

. There is usually a finely granular pink to purple material in areas of PMN infiltration adjacent to blood vessels. This represents large clumps of bacteria, best visualized with a modified Warthin

-Starry stain. Standard tissue Gram stain and the Warthin

-Starry stain used for syphilis do not stain the organisms. If the diagnosis cannot be confirmed with special stains, electron microscopy may be used. The lack of spindle cells, atypically shaped vascular channels, and hyaline globules distinguish BA from Kaposi sarcoma. Lesions of BA in tissues other than liver show similar histologic features.

Clinical Course

In the immunocompromised host, the natural history of untreated BA is gradually progressive disease, with increasing numbers of skin lesions and involvement of many visceral organs.

Untreated, severely immunocompromised patients might die of their infection.

Treatment

Erythromycin, 500 mg four times a day, or doxycycline, 100 mg twice a day for 3 months, is the treatment of choice for BA. Other antibiotics thought to be effective are

minocycline

, tetracycline,

chloramphenicol

,

azithromycin

, and

roxithromycin

. For

peliosis

hepatitis

, 4 months of treatment is recommended. Some patients require life-long suppressive therapy.

Relapses have been reported, especially with shorter treatment courses. A

Jarisch-Herxheimer

reaction not uncommonly occurs after initiation of therapy. Most patients respond rapidly to antibiotic therapy. The rapid disappearance of the vascular lesions may relate to the effects of the

macrolides

and

tetracyclines

on protein synthesis, stopping the production of vascular growth factors. If patients are not treated for a sufficient period, they are likely to relapse, despite the fact that their skin lesions vanish after a few weeks of treatment. The additional treatment is required to sterilize visceral or

hematologic

reservoirs of bacteria.