





Sarcoidosis

Sarcoidosis is a granulomatous disease, often systemic, of undetermined cause. A distinction is made
In subacute, transient sarcoidosis, erythema nodosum is associated with hilar adenopathy, fever, and,
In systemic sarcoidosis, cutaneous lesions are encountered in approximately one fourth of patients wh

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In the United States, this disorder is much more common and is more severe in African Americans . It is
The most common cutarlepous presions of sarcoid is is police of rownis redescontarpited prepartes and splacifates. With

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A rare form of sarcoidosis is its lichenoid variant, in which small, papular lesions occur . Very rare mani-
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Subcutaneous nodules of sarcoidosis are also rare. Originally described by Darier and Roussy, they r

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occur in association with other cutaneous lesion	or alone . Up to 80% to 90%	of patients with subcut
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Systemic sarcoidosis occasionally coexists with granuloma annulare . Cutaneous lesions of sarcoidosis
Histopathology.
The lesions of erythema nodosum occurring in subacute, transient sarcoidosis have the same histologic

Like lesions in other organs, the cutaneous lesions of chronic, persistent sarcoidosis are characterized by the lesions in other organs, the cutaneous lesions of chronic, persistent sarcoidosis are characterized by the lesions of chronic, persistent sarcoidosis are characterized by the lesions of chronic, persistent sarcoidosis are characterized by the lesions of chronic, persistent sarcoidosis are characterized by the lesions of chronic, persistent sarcoidosis are characterized by the lesions of chronic, persistent sarcoidosis are characterized by the lesions of chronic, persistent sarcoidosis are characterized by the lesions of chronic, persistent sarcoidosis are characterized by the lesions of chronic, persistent sarcoidosis are characterized by the lesions of chronic, persistent sarcoidosis are characterized by the lesions of chronic, persistent sarcoidosis are characterized by the lesions of chronic, persistent sarcoidosis are characterized by the lesions of chronic, persistent sarcoidosis are characterized by the lesions of chronic, persistent sarcoidosis are characterized by the lesions of chronic, persistent sarcoidosis are characterized by the lesions of chronic, persistent sarcoidosis are characterized by the lesions of chronic, persistent sarcoidosis are characterized by the lesions of chronic, persistent sarcoidosis are characterized by the lesions of chronic sarcoidos are characterized by the lesions	Sarcoidosis = 00000000
The papules, plaques, and lupus pernio-type lesions show variously sized aggregates of epithelioid cells	Like lesions in other organs, the cutaneous lesions of chronic, persistent sarcoidosis are characterized
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and, rarely, also giant cells . Typical sarcoidal granulomas are found in the ichthyosiform lesions , in ul
hypopigmented sarcoid may reveal granulomas, which may have a perineural component or fail to reve

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In typical cutaneous lesions of sarcoidosis, the well-demarcated islands of epithelioid cells contain few,

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Classically, sarcoid has been associated with only a sparse lymphocytic infiltrate, particularly at the	mar
oscinabilia staining is found in the center of some of the grapulomas. A roticulum stain of sarcoid	rovo
eosinophilic staining is found in the center of some of the granulomas . A reticulum stain of sarcoid	reve

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Systemic Lesions.	The lungs are the most commonly involved organ in the chronic, persistent type
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In about 25% of the nationts, equipy manifestations equipy most commonly obvenic ivide evolitic. Opleson
In about 25% of the patients, ocular manifestations occur, most commonly chronic iridocyclitis. Splenon

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Sarcoidosis, although usually a benign disease, is fatal in approximately 5% of patients . The most com-
Sarcoldosis, although distally a benigh disease, is fatal in approximately 37% of patients. The most con-

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The diagnosis of	of sarcoidosis in a p	atient with syste	emic disease is	based on clinic	al presentation,	biops

Pathogenesis . The cause of sarcoidosis/Miscantilenutworkurand the disabsence	Sarcoidosis = 000000000	
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Electron microscopic examination of epithelioid cells fails to show any evidence of b	acterial fragments, ι
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residual bodies of lysosomes. Asteroid bodies	consist of collagen showing the typical 64- to 70-nm period
Differential Diagnosis .	The histologic differentiation of sarcoidosis from lupus v
occasionally impossible. There is no absolute	histologic criterion by which the two diseases can be diffe

Foreign-body granulomas can also resemble sarcoidosis. Polariscopic examination in search o	f doubly :
Tuberculoid leprosy, which may show granulomas in association with only a sparse lymphocytic	c infiltrate