

Zespół znamienia Beckera

Kod Orpha: 64755 Kod OMIM: 604919

Opis choroby *

Definicja

A rare, syndromic, benign, epidermal nevus syndrome characterized by the association of a Becker nevus (i.e. circumscribed, unilateral, irregularly shaped, hyperpigmented macules, with or without hypertrichosis and/or acneiform lesions, occurring predominantly on the anterior upper trunk or scapular region) with ipsilateral breast hypoplasia or other, typically hypoplastic, skeletal, cutaneous, and/or muscular defects, such as pectoralis major hypoplasia, supernumerary nipples, vertebral defects, scoliosis, limb asymmetry, odontomaxillary hypoplasia and lipoatrophy.

Dane

Klasyfikacja

Choroba

Synonimy

Pigmentary hairy epidermal nevus
Owłosione znamię naskórkowe

Kod ORPHA

64755

Kod OMIM

604919

Kod ICD10

D22.5

Kod ICD11

LC02

[*Źródło](#)

orphanet

Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.