

Zespół Stevensa i Johnsona

Kod Orpha: 36426 Kod OMIM: 608579

Opis choroby *

Definicja

A limited form of Stevens-Johnson syndrome/toxic epidermal necrolysis spectrum characterized by destruction and detachment of the skin epithelium, involving less than 10% of the body surface area, and mucous membranes. Onset usually occurs 4-28 days after administration of the causal medication and is most frequently associated with anticonvulsants, antibacterial sulfonamides, allopurinol, nevirapine, and oxicams (non-steroidal anti-inflammatory drugs), but many other medications have also been implicated. The disease is not induced by medication in 15% of cases. Histology is characterized by an epidermal necrolysis. Multiple disabling long-term sequelae (especially cutaneous, ocular and psychological) are frequent.

Dane

Klasifikacja

Podtyp kliniczny

Synonimy

Dermatostomatitis, Stevens Johnson type

Dermatostomatitis, Stevens Johnson type

Kod ORPHA

36426

Kod OMIM

608579

Kod ICD10

L51.1

Kod ICD11

EB13.0

[*Źródło](#)

orphanet

Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.

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