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# Stevens-Johnson syndrome

INSERM

## Source

*INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Stevens-Johnson syndrome. ORPHA:36426*

Stevens-Johnson syndrome is a limited form of toxic epidermal necrolysis (see this term) characterized by destruction and detachment of the skin epithelium and mucous membranes involving less than 10% of the body surface area.