## Opis choroby \*

## Definicja

Scleroderma is a rare autoimmune connective tissue disorder characterized by abnormal hardening of the skin and, sometimes, other organs. It is classified into two main forms: localized scleroderma and systemic sclerosis (SSc), the latter comprising three subsets; diffuse cutaneous SSc (dcSSc), limited cutaneous SSc (lcSSc) and limited SSc (lSSc) (see these terms).

Dane		
<b>Klasyfikacja</b> Grupa fenomenów		
<b>Kod ORPHA</b> 801	Kod OMIM -	Kod ICD10 -
Kod ICD11		
-		
<u>*Źródło</u>		
orphanet		