Opis choroby *

Definicja

A rare epidermal disease characterized by rough, dry skin with prominent, plate-like scaling. It is non-hereditary and usually arises during adulthood in the context of a variety of diseases or conditions, like various types of cancer, autoimmune diseases, endocrine disorders, nutritional deficiencies, but also as a side effect of certain medications. Severity depends on the underlying disease or condition.

Dane			
Klasyfikacja Choroba			
Kod ORPHA 454	Kod OMIM -	Kod ICD10 L85.0	
Kod ICD11 ED50.0			
<u>*Źródło</u>			
orphanet			