## Opis choroby *

Definicja

A rare dysostosis with predominant vertebral involvement characterized by paraspinal ligament ossification (most pronounced in the lower thoracic region), osteophytosis, marginal sacroiliac joint sclerosis, and punctate hyperkeratosis on the soles and palms. Patients may be asymptomatic or present mild to moderate back pain. There have been no further descriptions in the literature since 1969.

Dane

**Klasyfikacja**
Zespół wad wrodzonych

**Kod ORPHA**
2206

**Kod OMIM**
106400

**Kod ICD10**
M48.1

**Kod ICD11**
LD24.H

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*Źródło

orphanet