

## Opis choroby \*

### Definicja

A rare arthrogyposis syndrome characterized by the association of multiple congenital joint contractures (of the large joints, fingers and toes) and hyperkeratosis (i.e. thick, scaling and fissured skin), with death occurring in early infancy. There have been no further reports in the literature since 1993.

### Dane

|                       |   |
|-----------------------|---|
| <b>Klasyfikacja</b>   | <b>Synonimy</b>   |
| Zespół wad wrodzonych | Johnston-Aarons-Schelley syndrome<br>Zespół Johnstona, Aaronsa i Schelley |

|                  |                 |                  |
|------------------|-----------------|------------------|
| <b>Kod ORPHA</b> | <b>Kod OMIM</b> | <b>Kod ICD10</b> |
| 1485             | 208158          | Q68.8            |

### Kod ICD11

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### \*Źródło

orphanet