

## Opis choroby \*

### Definicja

An extremely rare congenital limb malformation syndrome, described in only 3 patients to date, characterized by the association of hypoplasia or aplasia of the hand and foot phalanges, hemivertebrae and various urogenital and/or intestinal abnormalities (i.e. dysgenesis of the urogenital tract and rectum). There have been no further descriptions in the literature since 1991.

### Dane

<b>Klasyfikacja</b>	<b>Synonimy</b>
Zespół wad wrodzonych	Johnson-Munson syndrome
	Zespół Johnsona i Munsona

<b>Kod ORPHA</b>	<b>Kod OMIM</b>	<b>Kod ICD10</b>
1112	207620	Q87.8

### Kod ICD11

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### \*Źródło

orphanet