

## Opis choroby \*

### Definicja

A rare genetic multiple congenital anomalies/dysmorphic syndrome characterized by global developmental delay and intellectual disability, infantile hypotonia, microcephaly, movement disorder, and impaired balance. More variable manifestations are hearing loss, cortical visual impairment, abnormalities of fingers and/or toes, congenital cardiac anomalies, kyphoscoliosis, dysmorphic facial features, abnormal sleep pattern, and seizures, among others.

### Dane

<b>Klasyfikacja</b>	<b>Synonimy</b>
Zespół wad wrodzonych	You-Hoover-Fong syndrome
	Zespół You'a i Hoover-Fong

<b>Kod ORPHA</b>	<b>Kod OMIM</b>	<b>Kod ICD10</b>
488642	616954	Q87.8

**Kod ICD11**  
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### \*Źródło

orphanet