## Opis choroby *

Definicja

Olivopontocerebellar atrophy-deafness syndrome is characterised by infancy-onset olivopontocerebellar atrophy, sensorineural deafness and speech impairment. It has been described in less than 15 children. Most cases were sporadic, but autosomal recessive inheritance was suggested in three cases.

Dane

| **Klasyfikacja** | Synonimy |
| --- | --- |
| Zespół wad wrodzonych | Olivopontocerebellar atrophy-hearing loss syndrome |

| **Kod ORPHA** | **Kod OMIM** | **Kod ICD10** |
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| 2732 | - | Q04.8 |

**Kod ICD11**
LD2H.Y

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*Źródło

orphanet