

## Opis choroby \*

### Definicja

A rare syndrome characterized by gingival fibromatosis associated with progressive sensorineural hearing loss. It has been described in two families (with at least 16 affected members spanning five generations in one of the families, and five affected members spanning three generations in the other family). It is transmitted as an autosomal dominant trait.

### Dane

<b>Klasyfikacja</b>	<b>Synonimy</b>
Zespół wad wrodzonych	Gingival fibromatosis-progressive hearing loss syndrome Zespół Jonesa Jones syndrome

<b>Kod ORPHA</b> 2027	<b>Kod OMIM</b> 135550	<b>Kod ICD10</b> H90.3
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**Kod ICD11**  
LD2H.Y

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### \*Źródło

orphanet