

## Opis choroby \*

### Definicja

A rare autosomal dominant association characterized clinically by juvenile cataract associated with bilateral microcornea, and renal glucosuria without other renal tubular defects.

### Dane

<b>Klasyfikacja</b> Choroba	Synonimy Juvenile cataract-microcornea-renal glycosuria syndrome
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<b>Kod ORPHA</b> 247794	<b>Kod OMIM</b> 612018	<b>Kod ICD10</b> E88.8
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**Kod ICD11**  
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### \*Źródło

orphanet