

# Iminoglicynuria

**Kod Orpha: 42062 Kod OMIM: 242600**

## Opis choroby \*

### Definicja

A rare inborn error of metabolism characterized by elevated levels of imino acids (proline, hydroxyproline) and glycine in urine due to defective reabsorption in the kidney. The condition is considered benign and not associated with any specific clinical phenotype. Mode of inheritance is autosomal recessive.

### Dane

### Klasyfikacja

Choroba

### Kod ORPHA

42062

### Kod OMIM

242600

### Kod ICD10

E72.0

### Kod ICD11

5C60.Y

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### [\\*Źródło](#)

orphanet

## Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.