

# Mnogi Niedobór karboksylazy

Kod Orpha: 148 Kod OMIM:

## Opis choroby \*

### Definicja

A group of inborn errors of biotin metabolism characterized by reduced activities of biotin-dependent enzymes resulting in a wide spectrum of symptoms, including feeding difficulty, breathing difficulties, lethargy, seizures, skin rash, alopecia, and developmental delay. This group includes biotinidase deficiency and biotin holocarboxylase synthetase deficiency.

### Dane

#### Klasyfikacja

Grupa fenomenów

#### Synonimy

MCD

MCD

#### Kod ORPHA

148

#### Kod OMIM

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#### Kod ICD10

E53.8

#### Kod ICD11

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#### [\\*Źródło](#)

orphanet

## Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.