

## **Opis choroby \***

### Definicja

Carnitine palmitoyltransferase II (CPT II) deficiency is an inherited metabolic disorder that affects mitochondrial oxidation of long chain fatty acids (LCFA). Three forms of CPT II deficiency have been described: a myopathic form, a severe infantile form and a neonatal form (see these terms).

### Dane

Klasifikacja	Synonimy
Choroba	CPT2
	CPT2
	CPTII
	Niedobór palmitylotransferazy karnityny 2
	CPTII
	Carnitine palmitoyltransferase deficiency type 2
<b>Kod ORPHA</b>	<b>Kod OMIM</b>
157	608836
<b>Kod ICD11</b>	<b>Kod ICD10</b>
5C52.00	E71.3

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\*[Źródło](#)

orphanet