

## **Opis choroby \***

### Definicja

A rare autosomal recessive amino acid metabolism disorder characterized clinically by variable degrees of hyperammonemia, developing from about 3 years of age, and leading to progressive loss of developmental milestones and spasticity in the absence of treatment.

### Dane

Klasyfikacja	Synonimy	
Choroba	Arginase deficiency	
	Hyperargininemia	
	Niedobór arginazy	
	Hyperargininemia	
Kod ORPHA	Kod OMIM	Kod ICD10
90	207800	E72.2
Kod ICD11		
5C50.A2		

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### \*Źródło

orphanet