Opis choroby *

Definicja

A rare endocrine disease characterized by hyperinsulinemic hypoglycemia associated with the presence of autoantibodies to endogenous insulin without previous exposure to exogenous insulin. Patients usually present in adulthood with postprandial, fasting-, or exercise-induced hypoglycemia, often with pronounced neuroglycopenic symptoms. Laboratory investigations reveal markedly elevated serum insulin, as well as increased C-peptide and proinsulin. The condition may be associated with other autoimmune diseases, monoclonal gammopathy, and/or recent exposure to certain medications.

Dane

Klasyfikacja
Choroba

Synonimy
Hirata disease
Choroba Hirata

Kod ORPHA
411593

Kod OMIM
-

Kod ICD10
E16.1

Kod ICD11
-

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*Źródło

orphanet