

## 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

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#### Kod ICD10

E16.1

#### Kod ICD11

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#### \*Źródło

orphanet