

## Opis choroby \*

### Definicja

A rare stiff person syndrome spectrum disorder characterized by limb and truncal rigidity, stimulus-sensitive spasms, myoclonus, hyperekplexia, autonomic disturbance, and brainstem involvement or other neurological defects. The condition is progressive and potentially life-threatening, especially due to respiratory failure. It may be associated with the presence of glycine receptor or glutamic acid decarboxylase antibodies, as well as thymomas or lymphomas.

### Dane

<b>Klasyfikacja</b>	Synonimy
Podtyp kliniczny	PERM PERM

<b>Kod ORPHA</b>	<b>Kod OMIM</b>	<b>Kod ICD10</b>
438266	184850	G04.8

**Kod ICD11**  
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### \*Źródło

orphanet