

## Opis choroby \*

### Definicja

A rare X-linked syndromic intellectual disability characterized by intellectual disability of variable degree, behavioral anomalies (including autism, mood disorders, obsessive-compulsive behavior, and hetero- and auto-aggression), and epilepsy. Progressive neurological symptoms like movement disorders and spasticity, as well as subtle dysmorphic features have also been reported. Heterozygous females may be as severely affected as males.

### Dane

<b>Klasyfikacja</b> Choroba	<b>Synonimy</b> Raynaud-Claes syndrome Raynaud-Claes syndrome	
<b>Kod ORPHA</b> 485350	<b>Kod OMIM</b> 300114	<b>Kod ICD10</b> F78.1
<b>Kod ICD11</b> LD90		

---

### \*Źródło

orphanet