

## **Opis choroby \***

Definicja

A rare pervasive developmental disorder characterized by the presence of a unilateral angioma on the face and autistic developmental problems including language delay and atypical social interactions. The disease may initially resemble Sturge-Weber syndrome.

Dane

### **Klasyfikacja**

Zespół wad wrodzonych

**Kod ORPHA**  
137911

**Kod OMIM**  
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**Kod ICD10**  
F84.8

**Kod ICD11**

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\*[Źródło](#)

[orphanet](#)