

## Opis choroby \*

### Definicja

A rare subtype of cerebrofacial arteriovenous metamerism characterized by unilateral arteriovenous malformations involving the cerebellum, pons, and mandible (lateral rhombencephalic group). The condition manifests in childhood. Common presenting signs and symptoms are progressive neurological deficit, hemorrhage, and cosmetic complaints like facial asymmetry.

### Dane

<b>Klasyfikacja</b>	<b>Synonimy</b>
Zespół wad wrodzonych	CAMS3 CAMS 3

<b>Kod ORPHA</b>	<b>Kod OMIM</b>	<b>Kod ICD10</b>
141199	-	Q28.2

**Kod ICD11**  
LA90.3Y

---

### \*Źródło

orphanet