

## Opis choroby \*

### Definicja

Congenital Horner syndrome is a rare neurological disorder characterized by relative pupillary miosis and blepharoptosis, evident at birth, caused by interruption of the oculosympathetic innervation at any point along the neural pathway from the hypothalamus to the orbit. Often additional symptoms, such as enophthalmos, facial anhidrosis, iris heterochromia, conjunctival congestion, transient hypotonia and/or pupillary dilation lag, may be present. Association with birth trauma, neoplasms or vascular malformations has been reported.

### Dane

#### Klasyfikacja

Choroba

#### Synonimy

Congenital Claude-Bernard-Horner syndrome

Wrodzony zespół Claude'a, Bernarda i Hornera

#### Kod ORPHA

91413

#### Kod OMIM

143000

#### Kod ICD10

G90.2

#### Kod ICD11

8D8A.1

---

#### \*Źródło

orphanet