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

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*Źródło

orphanet