Opis choroby *

Definicja

A rare neural tube defect characterized by cystic dilatation of the central canal of the spinal cord, herniating posteriorly through a dorsal spinal defect. The malformation can occur anywhere along the spinal cord but appears to be more frequent in the posterior cervical and the lumbosacral region. It may be an isolated anomaly or be associated with other defects, including anorectal and genitourinary anomalies, or sacral agenesis.

Dane

Klasyfikacja
Grupa fenomenów

Kod ORPHA
268813

Kod OMIM
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Kod ICD10
Q05.9

Kod ICD11
LA02.02

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*Źródło

orphanet