# Pierwotna jamistość rdzenia
# Kod Orpha: 99856 Kod OMIM:

## Opis choroby *

**Definicja**
A rare central nervous system malformation characterized by a fluid-filled longitudinally oriented cavity (syrinx) within the spinal cord, which may or may not communicate with the central canal, does not have an ependymal lining, and is either idiopathic or seen as a familial malformation. Clinical manifestations in symptomatic patients include neuropathic pain, as well as sensory and motor disturbances. Typical presentations may be cape-like loss of pain and temperature sensation along the torso and arms, or disproportionately greater motor impairment in upper compared to lower extremities.

**Dane**

**Klasyfikacja**
Wada morfologiczna

**Synonimy**
Congenital syringomyelia
Wrodzona jamistość rdzenia

**Kod ORPHA**
99856

**Kod OMIM**
-

**Kod ICD10**
Q06.4

**Kod ICD11**
8D66.0

---

*Źródło

orphanet

## Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.

**Orphanet** - interntowa baza danych dotyczących rzadkich chorób i sierochych leków. ©INSERM 1999 - Dostępna na stronie www.orphanet.pl