Opis choroby *

Definicja

A rare, congenital, intestinal malformation morphological anomaly characterized by an egg-like, cystic, mucus-filled mass, composed of intestinal mucosal lining and smooth muscle tissue. Commonly it presents in childhood with symptoms of recurrent urinary tract infections, gastroenteritis, obstruction, perianal sepsis and rectal bleeding. Drainage of mucus or pus from the anus is also a typical presenting sign. The majority are found in the retro-rectal space where they communicate with, or are contiguous to, the rectum.

Dane

Klasyfikacja
Wada morfologiczna

Kod ORPHA
171220

Kod OMIM
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Kod ICD10
Q43.4

Kod ICD11
LB17.Y

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*Źródło

orphanet