Opis choroby *

Definicja

A rare vaginal malformation characterized by the presence of a complete or incomplete septum dividing the vagina into two parallel cavities, resulting from failure of reabsorption of the midline uterine septum between the two fused Müllerian ducts during embryogenesis. Patients are often asymptomatic, but may present with menorrhagia, dysmenorrhea, dyspareunia, infertility, or spontaneous abortion. The condition may occur as an isolated malformation or in association with other Müllerian duct anomalies (such as septate uterus or uterus didelphys) or renal abnormalities.

Dane

Klasyfikacja
Podtyp kliniczny

Kod ORPHA
180157

Kod OMIM
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Kod ICD10
Q52.1

Kod ICD11
LB42.1

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*Źródło

orphanet