Opis choroby *

Definicja

A rare spectrum of Mullerian duct anomalies characterized by congenital aplasia of the uterus and upper two-thirds of the vagina in otherwise phenotypically normal females. It can be classified as either Mayer-Rokitansky-Küster-Hauser (MRKH) syndrome type 1 (corresponding to isolated utero-vaginal aplasia) or MRKH syndrome type 2 (utero-vaginal aplasia associated with other malformations).

Dane

Klasyfikacja
Zespół wad wrodzonych

Synonimy
MRKH syndrome
Zespół MRKH
Zespół Rokitansky'ego
Rokitansky syndrome

Kod ORPHA
3109

Kod OMIM
601076

Kod ICD10
Q51.8

Kod ICD11
LB44.Y

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*Źródło

orphanet