Opis choroby *

Definicja

A rare genetic, syndromic glomerular disorder characterized by the association of nephropathy presenting as persistent proteinuria or overt nephrotic syndrome, Wilms tumor and genitourinary structural defects. In addition, disorders of testicular development are common in subjects with 46,XY karyotype.

Dane

Klasyfikacja
Choroba

Synonimy
Drash syndrome
Guz Wilmsa i pseudohermfrodytyzm
Zespół Drasha
Wilms tumor-DSD syndrome
Wilms tumor-disorder of sex development syndrome

Kod ORPHA
220

Kod OMIM
194080

Kod ICD10
N04.1

Kod ICD11
LD2A.Y

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*Źródło

orphanet