

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 pseudohermafrodytyzm

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

*Źródło

orphanet