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 Synonimy

Choroba Drash syndrome

Guz Wilmsa i pseudohermfrodytyzm

Zespół Drasha

Wilms tumor-DSD syndrome

Wilms tumor-disorder of sex development

syndrome

Kod ORPHA Kod OMIM Kod ICD10

220 194080 N04.1

Kod ICD11 LD2A.Y

*Źródło

orphanet