DISEASE:
WAGR syndrome

NAME:	WAGR syndrome
DESCRIPTION:	A rare genetic disorder characterized by the association of total or partial aniridia, genitourinary anomalies (ranging from sexual ambiguity to ectopia testis), variable degrees of intellectual disability, and an increased risk of developing Wilms tumor. Glaucoma or cataract are also possible, and a minority of patients develop kidney failure. Other varaible findings may include obesity and duplicated halluces.
ORPHACODE:	893
SYNONYMS:	Del(11)(p13) Deletion 11p13 Monosomy 11p13 Wilms tumor-aniridia-genitourinary anomalies-intellectual disability syndrome
XREF(S):	Orphanet MeSH MeSH ICD-10 OMIM OMIM

ANALYTE(S):	BDNF WT1 PAX6
CREATED:	13 May 2019 - 01:02
CHANGED:	22 Jun 2023 - 16:14

Source URL: http://gentest.healthdata.be/disease/174

RELATED CONTENT

Related Genetic Tests

- Early-onset severe obesity
- Nephrotic syndrome, Focal Segmental Glomerulosclerosis (FSGS) , Alport syndrome and podocytopathy (gene panel)
- WAGR Syndrome

Related Laboratories

- Centre de Génétique Humaine - CHU Sart-Tilman
- Centre de Génétique-Institut de Pathologie et de Génétique (IPG)
- Centrum Menselijke Erfelijkheid - KUL

Related Analytes

- brain derived neurotrophic factor
- paired box 6
- WT1 transcription factor

Related Gene Panels

- Early-onset severe obesity (44 genes) - ULG
- Nephrotic syndrome, FSGS, Alport syndrome (76 genes) - IPG