DISEASE:
Autosomal dominant Alport syndrome

NAME:	Autosomal dominant Alport syndrome
ORPHACODE:	88918
XREF(S):	Orphanet MeSH OMIM ICD-10
ANALYTE(S):	COL4A3 COL4A4
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RELATED CONTENT

Related Genetic Tests

- Nephrotic syndrome, Focal Segmental Glomerulosclerosis (FSGS) , Alport syndrome and podocytopathy (gene panel)

Related Laboratories

- Centre de Génétique-Institut de Pathologie et de Génétique (IPG)

Related Analytes

- collagen type IV alpha 3 chain
- collagen type IV alpha 4 chain

Related Gene Panels

- Alport (X-linked and recessive) (3 genes) - UZA
- Nephrotic syndrome, FSGS, Alport syndrome (76 genes) - IPG