

**DISEASE:**  
**Oculocerebrorenal syndrome of Lowe**

<b>NAME:</b>	Oculocerebrorenal syndrome of Lowe
<b>DESCRIPTION:</b>	A rare multisystem disorder characterized by congenital cataracts, glaucoma, intellectual disabilities, seizures, postnatal growth retardation and renal tubular dysfunction with chronic renal failure.
<b>ORPHACODE:</b>	534
<b>SYNOMYS:</b>	Lowe disease Lowe oculo-cerebro-renal dystrophy Lowe oculo-cerebro-renal syndrome Lowe oculocerebrorenal dystrophy Lowe syndrome OCRL Phosphatidylinositol 4,5-biphosphate 5-phosphatase deficiency
<b>XREF(S):</b>	<a href="#">Orphanet</a> <a href="#">OMIM</a> <a href="#">MeSH</a> <a href="#">MedDRA</a> <a href="#">ICD-10</a>
<b>ANALYTE(S):</b>	<a href="#">OCRL</a>
<b>CREATED:</b>	13 May 2019 - 01:02

**CHANGED:**

22 Jun 2023 - 16:14

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### Related Genetic Tests

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### Related Analytes

- [OCRL inositol polyphosphate-5-phosphatase](#)

### Related Gene Panels

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