

**DISEASE:**  
**Isolated congenital sclerocornea**

<b>NAME:</b>	Isolated congenital sclerocornea
<b>DESCRIPTION:</b>	A rare corneal disorder characterized by non-inflammatory, non-progressive, bilateral ingrowth of vascularized, opaque scleral tissue into the peripheral cornea, obliterating the corneoscleral limbus and scleral sulcus. The condition is not associated with other ocular abnormalities.
<b>ORPHACODE:</b>	91490
<b>XREF(S):</b>	<a href="#">Orphanet</a> <a href="#">ICD-10</a> <a href="#">OMIM</a>
<b>ANALYTE(S):</b>	<a href="#">GJA8</a>
<b>CREATED:</b>	13 May 2019 - 01:02
<b>CHANGED:</b>	22 Jun 2023 - 16:14

---

Source URL: <http://gentest.healthdata.be/disease/3204>

## RELATED CONTENT

---

### Related Genetic Tests

- Cataract (gene panel)

### Related Laboratories

- Centrum Medische Genetica - UZ Gent

### Related Analytes

- gap junction protein alpha 8

### Related Gene Panels

- Cataract - UGent
- test

---

Source URL: <http://gentest.healthdata.be/disease/3204>