

**DISEASE:**  
**Thyroid hypoplasia**

<b>NAME:</b>	Thyroid hypoplasia
<b>DESCRIPTION:</b>	Thyroid hypoplasia is a form of thyroid dysgenesis (see this term) characterized by incomplete development of the thyroid gland that results in primary congenital hypothyroidism (see this term), a permanent thyroid deficiency that is present from birth.
<b>ORPHACODE:</b>	95720
<b>XREF(S):</b>	<a href="#">Orphanet</a> <a href="#">OMIM</a> <a href="#">OMIM</a> <a href="#">MedDRA</a> <a href="#">ICD-10</a>
<b>ANALYTE(S):</b>	<a href="#">SLC26A4</a> <a href="#">TSHR</a> <a href="#">PAX8</a>
<b>CREATED:</b>	13 May 2019 - 01:02
<b>CHANGED:</b>	22 Jun 2023 - 16:14

## RELATED CONTENT

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### Related Genetic Tests

- Thyroid disgenesis (38 genes)

### Related Laboratories

- Centrum Medische Genetica - UZ Brussel VUB

### Related Analytes

- paired box 8
- solute carrier family 26 member 4
- thyroid stimulating hormone receptor

### Related Gene Panels

- Thyroid disgenesis (38 genes) - VUB

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Source URL: <http://gentest.healthdata.be/disease/3463>