

**DISEASE:**  
**Autoimmune lymphoproliferative syndrome**

<b>NAME:</b>	Autoimmune lymphoproliferative syndrome
<b>DESCRIPTION:</b>	A rare, inherited disorder characterized by non-malignant lymphoproliferation, multilineage cytopenias, and a lifelong increased risk of Hodgkin's and non-Hodgkin's lymphoma.
<b>ORPHACODE:</b>	3261
<b>SYNONYMS:</b>	ALPS Canale-Smith syndrome
<b>XREF(S):</b>	<a href="#">Orphanet</a> <a href="#">OMIM</a> <a href="#">MeSH</a> <a href="#">OMIM</a> <a href="#">OMIM</a> <a href="#">OMIM</a> <a href="#">MedDRA</a> <a href="#">ICD-10</a>

<b>ANALYTE(S):</b>	<u>CASP10</u> <u>FAS</u> <u>FAS</u> <u>FASLG</u> <u>PRKCD</u> <u>RASGRP1</u>
<b>CREATED:</b>	13 May 2019 - 01:02
<b>CHANGED:</b>	22 Jun 2023 - 16:14

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Source URL: <http://gentest.healthdata.be/disease/1392>

## RELATED CONTENT

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### Related Genetic Tests

- [Autoimmune lymphoproliferative syndrome](#)
- [Autoimmune lymphoproliferative syndrome type 1A](#)
- [Primary immune deficiencies \(gene panel\)](#)

### Related Laboratories

- [Centrum Menselijke Erfelijkheid - KUL](#)

### Related Analytes

- [caspase 10](#)
- [Fas cell surface death receptor](#)
- [Fas ligand](#)
- [protein kinase C delta](#)
- [RAS guanyl releasing protein 1](#)

### Related Gene Panels

- [Immunogenetics \(21 genes\)](#)
- [Primary immune deficiencies \(444 genes\) - KUL](#)