

**DISEASE:**  
**Sézary syndrome**

<b>NAME:</b>	Sézary syndrome
<b>DESCRIPTION:</b>	Sézary syndrome (SS) is an aggressive form of cutaneous T-cell lymphoma characterized by a triad of erythroderma, lymphadenopathy and circulating atypical lymphocytes (Sézary cells).
<b>ORPHACODE:</b>	3162
<b>SYNOMYS:</b>	Sézary lymphoma
<b>XREF(S):</b>	<a href="#">Orphanet</a> <a href="#">ICD-10</a> <a href="#">MeSH</a> <a href="#">MedDRA</a>
<b>ANALYTE(S):</b>	<a href="#">CTLA4</a> <a href="#">TNFRSF1B</a> <a href="#">CD28</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

- Primary immune deficiencies (gene panel)

### Related Laboratories

- Centrum Menselijke Erfelijkheid - KUL

### Related Analytes

- CD28 molecule
- cytotoxic T-lymphocyte associated protein 4
- TNF receptor superfamily member 1B

### Related Gene Panels

- Primary immune deficiencies (444 genes) - KUL

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