

## **GENETIC TEST:** **Combined immunodeficiency (severe), X-linked**

<b>FULL NAME:</b>	Combined immunodeficiency (severe), X-linked
<b>TEST TYPE:</b>	Clinical
<b>TEST SPECIALTY:</b>	Molecular Genetics
<b>TEST PURPOSE:</b>	Post-natal Diagnosis
<b>SPECIMEN:</b>	Peripheral (whole) blood on EDTA
<b>METHOD CATEGORY:</b>	Sequence analysis: entire coding region
<b>METHOD TECHNIQUE:</b>	Next Generation Sequencing (NGS)
<b>RIZIV CODE:</b>	565456-565460
<b>TURNAROUND TIME (MAXIMUM):</b>	4-6 months
<b>CREATED:</b>	18 Jul 2019 - 12:05
<b>CHANGED:</b>	14 Oct 2021 - 10:13
<b>URL:</b>	<a href="https://laboboeken.nexuzhealth.com/pboek/internet/GHB%20CME/14676">https://laboboeken.nexuzhealth.com/pboek/internet/GHB%20CME/14676</a>

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## RELATED CONTENT

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### Related Diseases

- T-B+ severe combined immunodeficiency due to gamma chain deficiency

### Related Laboratories

- Centrum Menselijke Erfelijkheid - KUL

### Related Analytes

- interleukin 2 receptor subunit gamma

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