

## **GENETIC TEST:** **X-linked creatine deficiency**

<b>FULL NAME:</b>	X-linked creatine deficiency
<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, DNA
<b>METHOD CATEGORY:</b>	Sequence analysis: entire coding region
<b>METHOD TECHNIQUE:</b>	Bi-directional Sanger Sequence analysis
<b>RIZIV CODE:</b>	565471-565482
<b>TURNAROUND TIME (MAXIMUM):</b>	1 - 2 months
<b>CREATED:</b>	02 Sep 2019 - 12:10
<b>CHANGED:</b>	25 Feb 2022 - 12:20
<b>URL:</b>	<a href="https://www.chuliege.be/jcms/c_7077895/fr/deficit-en-creatine-lie-a-l-x">https://www.chuliege.be/jcms/c_7077895/fr/deficit-en-creatine-lie-a-l-x</a>

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## RELATED CONTENT

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### Related Diseases

- X-linked creatine transporter deficiency

### Related Laboratories

- Centre de Génétique Humaine - CHU Sart-Tilman

### Related Analytes

- solute carrier family 6 member 8

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