

## **GENETIC TEST:** **Enzymatic dosage Pompe disease**

<b>FULL NAME:</b>	Enzymatic dosage Pompe disease
<b>DESCRIPTION:</b>	alpha-glucosidase
<b>TEST TYPE:</b>	Clinical
<b>TEST SPECIALTY:</b>	Biochemical Genetics
<b>TEST PURPOSE:</b>	Carrier diagnosis, Post-natal Diagnosis
<b>SPECIMEN:</b>	Skin fibroblasts
<b>METHOD CATEGORY:</b>	Unknown
<b>METHOD TECHNIQUE:</b>	Unknown
<b>RIZIV CODE:</b>	565574-565585
<b>TURNAROUND TIME (MAXIMUM):</b>	3 months
<b>CREATED:</b>	26 Aug 2019 - 16:37
<b>CHANGED:</b>	27 Jul 2022 - 11:27

URL:	<a href="https://laboguide.uzbrussel.be/laboguide#Analyses:Ziekte%20van%20Pompe&amp;&amp;&amp;&amp;789&amp;...">https://laboguide.uzbrussel.be/laboguide#Analyses:Ziekte%20van%20Pompe&amp;&amp;&amp;&amp;789&amp;...</a>
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## RELATED CONTENT

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### Related Diseases

- [Glycogen storage disease due to acid maltase deficiency](#)
- [Glycogen storage disease due to acid maltase deficiency, infantile onset](#)
- [Glycogen storage disease due to acid maltase deficiency, late-onset](#)

### Related Laboratories

- [Centrum Medische Genetica - UZ Brussel VUB](#)

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