

## Centre for Human Metabolomics (CHM)

<b>Test:</b>	<b>Selective Lysosomal disorder screening - BLOOD SPOT SAMPLES (2x Guthrie cards)</b>
<b>NHRPL Tariff code:</b>	4268 x2
<b>Tariff (including VAT):</b>	R 2075.76 x 2 = R 4151.52
<b>Description:</b>	<b>Lysosomal storage disorders screen:</b> Fabry disease ( $\alpha$ -galactosidase), Krabbe disease/globoid cell leukodystrophy / galactosylceramide lipidosis (galactocerebrosidase), Gaucher disease ( $\beta$ -glucosidase), Niemann-Pick disease A / B (sphingomyelinase), MPS I / Hurler-, Hurler-Scheie-, Sheie syndrome ( $\alpha$ -L-iduronidase), Pompe / Glycogen storage defect type II ( $\alpha$ -glucosidase).
<b>Turnaround time:</b>	<ol style="list-style-type: none"> <li>1. 10 - 15 <b>work days</b> from receipt of sample at our laboratory (excluding public holidays and weekends).</li> <li>2. Abnormal results are not sufficient to conclusively establish a diagnosis of a particular disease.</li> <li>3. Genetic testing is required to confirm the diagnosis of a related disorder</li> </ol>
<b>Transit stability / Sample viability:</b>	<ol style="list-style-type: none"> <li>1. <b>Do not expose specimen to heat or direct sunlight.</b></li> <li>2. Do not stack wet specimens.</li> <li>3. Keep specimen dry.</li> <li>4. <b>AFTER the blood card samples are completely dry</b>, ensure that it is <b>transported within 36 hours after collection</b> to PLIEM laboratory.</li> <li>5. <b>If the samples are exposed to heat and not handled according to PLIEM laboratory's protocol, the samples will not be viable for the testing.</b></li> </ol>
<b>Comments:</b>	Specimens exposed to heat $>25^{\circ}\text{C}$ will not be viable for testing. Blood transfusion prior to collection may influence the analysis. Test is performed at accredited external laboratory in the USA.
<b>Sample required:</b>	<ol style="list-style-type: none"> <li>1. <b>2 x Blood collection cards (Guthrie cards / DBS)</b> by heel prick or fingerstick for a patient <math>&gt; 1</math> year of age.</li> <li>2. Allow blood to <b>dry on the filter paper at ambient temperature</b> in a <b>horizontal position for 3 hours. Required: Whatman Protein Saver 903 Paper.</b></li> </ol>
<b>Method:</b>	Enzyme assay on tandem mass spectrometry
<b>Reference ranges &amp; units:</b>	Gaucher: Beta-Glucosidase $> \text{or} = 1.75 \text{ nmol/mL/hr}$ Niemann-Pick A/B : Sphingomyelinase $> \text{or} = 2.5 \text{ nmol/mL/hr}$ Pompe disease: Alpha-Glucosidase $> \text{or} = 3 \text{ nmol/mL/hr}$ Fabry: Alpha-Galactosidase $> \text{or} = 2.75 \text{ nmol/mL/hr}$ MPS I: Alpha-L-Iduronidase $> \text{or} = 2.0 \text{ nmol/mL/hr}$ Galactocerebrosidase $> \text{or} = 0.4 \text{ nmol/mL/hr}$ <span style="float: right;">Krabbe:</span>
<b>Contact for results &amp; other enquiries:</b>	Sample reception and resulting
<b>Telephone number:</b>	018 299 2312 / 018 285 2652 (leave message)
<b>Fax number:</b>	018 299 2316
<b>E-mail address:</b>	<a href="mailto:pliem@nwu.ac.za">pliem@nwu.ac.za</a>
<b>Delivery address for samples:</b>	Center for Human Metabolomics (CHM), Sample reception (PLIEM/NBS/CRS) Building F3, Room Number G19, 11 Hoffmann street North West University, Potchefstroom, 2531

PLEASE NOTE: Collection, courier and administration costs are not included.  
Protocol for each individual test is available on our website: [www.pliem.co.za](http://www.pliem.co.za)

Valid: 1 January 2019 - 31 December 2019