Physician:

Dr. M. Jaksch Freiburg Medical Lab

Laboratory Report Online Version

Report Date: 17.11.2018

Patient Name: NBS abnormal Hb

Gender: Female
Date of Birth: 16.11.2018

Nationality: 17.11.2018

Your ID:

Test Request Code:

Sample ID:

Patient IDNo: 380524

Sampling Date / Time: 17.11.2018 / 00:00 Receipt Date / Time: 17.11.2018 / 12:19

1816

Remarks: Insurance:

Analysis Result Flag Units Reference Range

Exclusion of Metabolic Diseases (Filter card)

Week of gestation 39
Newborn screening* see text

No sign of the following disorders:

Amino Acid related disorders such as: Phenylketonuria (PKU),

Hyperphenylalaninaemia, Maple syrup disease etc.

Fatty acid oxidation disorders and other disorders recognized in the

acylcarnitine spectrum: e.g. MCAD, LCAD, VLCAD

Other screened disorders: e.g. Hypothyreosis, AGS, Biotinidase deficiency, Galactosaemia, G6PDH Deficiency, Cystic Fibrosis, Hemoglobinopathies

(HbS, HbC, HbD, HbE, HbH/HbBarts, HbO, HbLepore, HbG, beta-Thalassemia)

Methods: MS-MS, FIA, Photometry, Capillary Electrophoresis

Please note that the best time of collecting blood for NBS is 36-72 hours after birth.

Amino Acids (PKU etc.)*	normal	quantitative	normal
Acylcarnitines (Fatty acids)*	normal	quantitative	normal
TSH (Hypothyreodism)*	normal	quantitative	normal
17a-OH-Progesterone (AGS)*	normal	quantitative	normal
Galactose (Galactosemia)*	normal	quantitative	normal
Biotinidase (Biot.deficiency)*	normal	quantitative	normal
Glucose-6-PDH*	normal	quantitative	normal
Immunoreactive Trypsin*	normal	quantitative	normal
Hemoglobin Screening*	HbH/HbBarts	quantitative	normal

The Hb electrophoresis detected an abnormal hemoglobin in the HbH/Hb Barts-range (<1.0%).

Note:

Our reference values are adjusted to age and gender. Daily internal Quality Control within the required range (according to ISO 15189).

(according to ISO 15189). External Quality Control available on request.

^ non-accredited parameter

"This parameter is affected by Biotin intake of >5 mg
(RDI = 0.03mg)

*This investigation has been performed in a collaborating accredited laboratory (Germany).

Techn. Validation by Med. Technologist (Supervisor of the Department)

Dr. Nehmat ElBanna Specialist Clinical Pathology (U/S)

(DHA-P-0084548)

PD Dr. med. habil. M. Jaksch Associate Professor 6) Medical Director (DHA-LS-240710) Physician:

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380524

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Insurance:

Analysis

Result

Flag

Units

Reference Range

This finding of a hemoglobin-Barts, a tetramer of 4 gamma-chains, indirectly indicates an alpha-thalassemia (-a/aa). Clinical symptoms like microcytic hypochrome anemia can progress over years. Usually, there is no indication for therapy. For a further differentiation, we recommend an additional molecular genetic DNA analysis.

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PD Dr. med. habil. M. Jaksch **Associate Professor Medical Director** (DHA-LS-240710)

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