Physician:

Laboratory Report Online Version

Report Date: 23.10.2018

Patient Name: SAMPLE REPORT ONLY

Gender: Female Date of Birth: 01.01.1992

Nationality:

Your ID:

Remarks:

Test Request Code:

Sample ID: Patient IDNo:

Sampling Date / Time: 21.10.2018 / 00:00 Receipt Date / Time: 21.10.2018 / 00:00

Insurance:

Analysis Result Flag Units Reference Range

Special Tests (Serum)

First Trimester Screening

Date of Ultrasound	21/10/2018	
Number of Fetuses	1	
CRL	66.50	mm
NT	1.95	mm
Week of Gestation (PROV.)	13+2	Wk + D
Week of Gestation (CALC)	13+1	Wk + D
Body Weight	48.8	Kg
Ethnical Background	Caucasian	
Smoker	No	
Free Beta-HCG (ECL)"	35.78	IU/I
MoM (free B-hCG)	1.00	
PAPP-A (ECL)"	2.500	IU/I
MoM (PAPP-A)	0.52	
Trisomy 21 risk (maternal age)	1:778	
Trisomy 21 risk (all data)	1:3260	
Trisomy 13+18 risk (all data)	1:46552	

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

External Quality Control available on request.

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

Specialist Clinical Pathology (U/S) (DHA-P-0084548)

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

[^] 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).

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In consideration of the data concerning maternal age, anamnesis and week of pregnancy, the empirical risk for Down's syndrome for the actual pregnancy is 1:778.

The additional inclusion of the test results for free B-hCG and PAPP-A including NT reveals an adjusted risk for Down's syndrome of 1:3260.

The risk of Trisomy 21 not elevated

It is lower compared to the risk for a pregnancy in the same gestational state of a 35 year old woman (1:150) which is considered to be the cut- off for discussing further diagnostic possibilities such as NIPT.

The risk of Trisomy 13 + 18 not elevated

The statistic calculation was performed by the SSDwLab software (provided by Roche). The calculation considers the biochemical parameters and the reported clinical data (week of gestation, crown rump length, nuchal translucency, weight, age of mother). The risk calculation is based on the day of the blood withdrawal. Please note that risk factors are calculated reciprocal (for example the risk 1:300 is higher than 1:600).

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

Dr. Nehmat ElBanna **Specialist** Clinical Pathology (U/S) (DHA-P-0084548)

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