KOS DIAGNOSTIC LAB

6349/1, NICHOLSON ROAD, AMBALA CANTT

Prisca 5.0.2.37

Date of report: 27-04-2018

Patient data			
Name	MRS. INDU	Patient ID	
Birthday	30-09-1991	Sample ID	1804220640/AMB
Age at sample date	26.6	Sample Date	27-04-2018

Gestational age	13 + 3				
Correction factors					
Fetuses	1	IVF	no	Previous trisomy 21	no
Weight	70	diabetes	no	pregancies	
Smoker	no	Origin	Asian		

Biochemical data

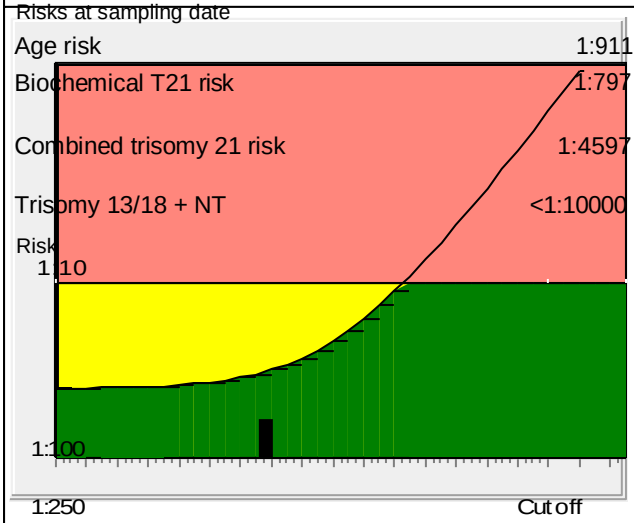
Parameter	Value	Corr. MoM
PAPP-A	3.1 mIU/ml	0.78
fb-hCG	52.1 ng/ml	1.81

Ultrasound data

Gestational age	13 + 0
Method	CRL Robinson
Scan date	24-04-2018
Crown rump length in mm	70.7
Nuchal translucency MoM	0.73
Nasal bone	present
Sonographer	.
Qualifications in measuring NT	MD

Risks at sampling date

Age risk	1:911
Biochemical T21 risk	1:797
Combined trisomy 21 risk	1:4597
Trisomy 13/18 + NT	<1:10000



Trisomy 21

The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.

After the result of the Trisomy 21 test (with NT) it is expected that among 4597 women with the same data, there is one woman with a trisomy 21 pregnancy and 4596 women with not affected pregnancies.

The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician. Please note that risk calculations are statistical approaches and have no diagnostic value!

The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)).

The laboratory can not be hold responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!

below cut off | Below Cut Off, but above Age Risk | above cut off

Trisomy 13/18 + NT

The calculated risk for trisomy 13/18 (with nuchal

translucency) is < 1:10000, which represents a low risk.

Sign of Physician