KOS DIAGNOSTIC LAB

6349/1, NICHOLSON ROAD, AMBALA CANTT

Prisca   5.0.2.37

Date of report: **24-04-2019**

Patient data			
Name	MRS. BEANT	Patient ID	231904230013
Birthday	28/09/1992	Sample ID	231904230013
Age at sample date	26	Sample Date	23-04-2019
Gestational age	13 + 5		

Correction factors					
Fetuses	1	IVF	no	Previous trisomy 21	no
Weight	42	diabetes	no	pregancies	
Smoker	no	Origin	Asian		

Biochemical data			Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age	13 + 5
PAPP-A	10.8 mIU/ml	1.96	Method	LMP
fb-hCG	35.5 ng/ml	1.04	Scan date	
Risks at sampling date			Crown rump length in mm	
Age risk		1:820	Nuchal translucency MoM	
Biochemical T21 risk		<1:10000	Nasal bone	present
Combined trisomy 21 risk		<1:10000	Sonographer	.
Trisomy 13/18 + NT		<1:10000	Qualifications in measuring NT	MD

Risk

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 more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy.

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!

Trisomy 13/18 + NT

The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.

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Sign of Physician

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