KOS DIAGNOSTIC LAB

6349/1, NICHOLSON ROAD, AMBALA CANTT

Prisca 5.0.2.37

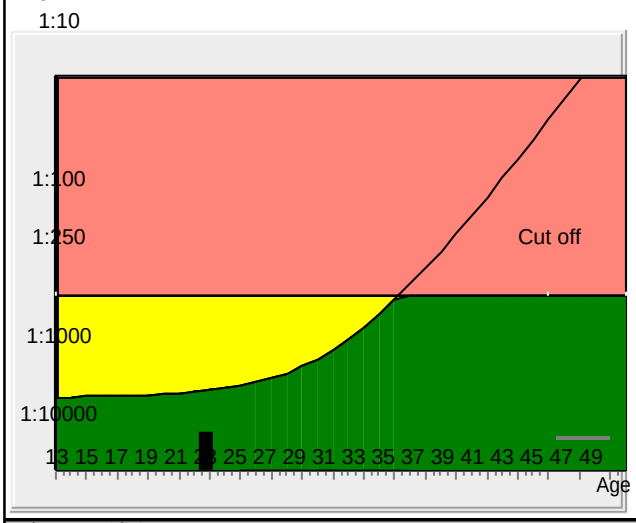
Date of report: 9-03-2019

Patient data			
Name	MRS. HARPREET	Patient ID	211903070004
Birthday	22/05/1996	Sample ID	211903070004
Age at sample date	22	Sample Date	07-03-2019
Gestational age	11 + 4		

Correction factors					
Fetuses	1	IVF	no	Previous trisomy 21 pregancies	no
Weight	48	diabetes	no		
Smoker	no	Origin	Asian		

Biochemical data			Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age	11 + 4
PAPP-A	1.9 mIU/ml	0.70	Method	
fb-hCG	30.1 ng/ml	0.69	Scan date	
			Crown rump length in mm	
Risks at sampling date			Nuchal translucency MoM	
Age risk		1:1000	Nasal bone	present
Biochemical T21 risk		1:6150	Sonographer	.
Combined trisomy 21 risk		<1:10000	Qualifications in measuring NT	MD
Trisomy 13/18 + NT		<1:10000		

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.

Sign of Physician

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