KOS DIAGNOSTIC LAB

6349/1, NICHOLSON ROAD, AMBALA CANTT

Prisca 5.0.2.37

Date of report: 22-12-2018

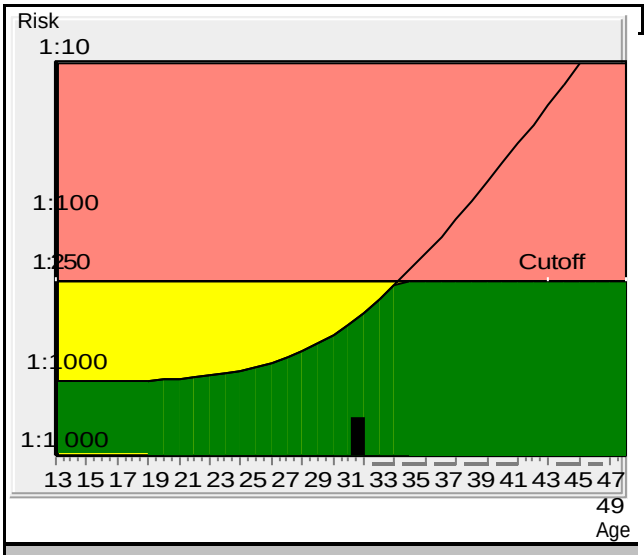
Patient data			
Name	Ms.SIMRANJEET	Patient ID	
Birthday	05-04-1986	Sample ID	131812210005
Age at sample date	32.6	Sample Date	21-12-2018
Gestational age	11 + 5		

Correction factors					
Fetuses	1	IVF	no	revious trisomy 21 pregancies	no
Weight	68	diabetes	no		
Smoker	no	Origin	Asian	ta	

Parameter	Value	Corr. MoM
PAPP-A	2.9 mIU/ml	1.61
fb-hCG	26.1 ng/ml	0.64

Risks at sampling date	
Age risk	1:419
Biochemical T21 risk	<1:10000
Combined trisomy 21 risk	<1:10000
Trisomy 13/18 + NT	<1:10000

Gestational age	11 + 5
Method	CRL Robinson
Scan date	20-12-2018
Crown rump length in mm	50
Nuchal translucency MoM	1.19
Nasal bone	present
Sonographer	.
Qualifications in measuring NT	MD



Trisomy 13/18 + NT

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

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!

Sign of Physician

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