KOS DIAGNOSTIC LAB

6349/1, NICHOLSON ROAD, AMBALA CANTT

Prisca 5.0.2.37

Date of report: 25-10-2018

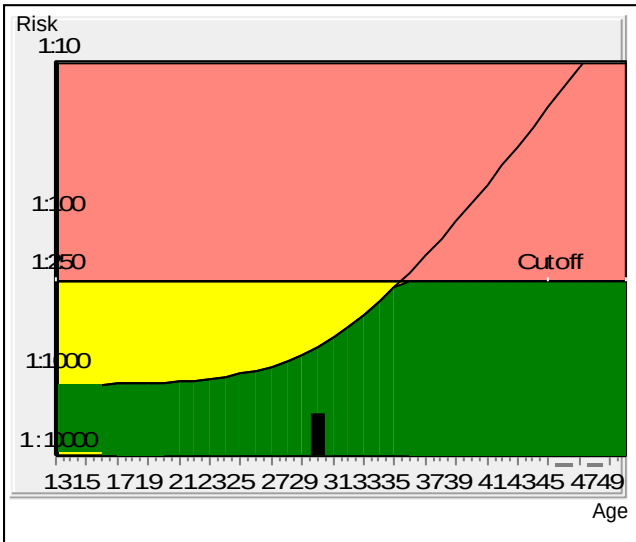
Patient data			
Name	MRS. HARMANJIT	Patient ID	1810220443/AMB
Birthday	12-11-1988	Sample ID	1810220443/AMB
Age at sample date	29.9	Sample Date	24-10-2018
Gestational age	12 + 4		

Correction factors					
Fetuses	1	IVF	no	Previous trisomy 21 pregancies	no
Weight	70.61	diabetes	no		
Smoker	no	Origin	Asian		

Parameter	Value	Corr. MoM
PAPP-A	2.1 mIU/ml	0.72
fb-hCG	74.4 ng/ml	2.23

Risks at sampling date	
Age risk	1:651
Biochemical T21 risk	1:285
Combined trisomy 21 risk	1:1721
Trisomy 13/18 + NT	<1:10000

Gestational age	12 + 2
Method	CRL Robinson
Scan date	22-10-2018
Crown rump length in mm	60.2
Nuchal translucency MoM	0.51
Nasal bone	present
Sonographer	DR. ARUN SHARMA
Qualifications in measuring NT	MD



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 1721 women with the same data, there is one woman with a trisomy 21 pregnancy and 1720 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!

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