PRISCA 5.0.2.37

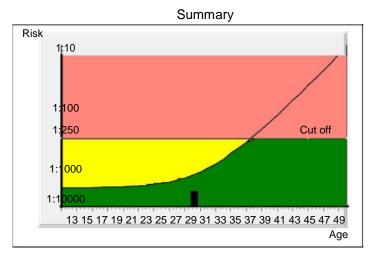
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6349/1, NICHOLSON ROAD, AMBALA CANTT

Results for:	Sample no	Date of report:
MRS. HARJOT KAUR	1610220062/AMB	05-10-2016

Referring Doctors

KOS DIAGNOSTIC LAB



Patient data	
Age at delivery	30.2
WOP	16+ 3
Weight	60 kg
Patient ID	

Risks at term	
Biochemical risk for Tr.21	1:3006
Age risk:	1:940
Neural tube defects risk	1:5118

For MRS. HARJOT KAUR, born on 26-12-1986, a screening test was performed on the 04-10-2016. Prisca screens for Trisomy 21, Trisomy 18 and Neural tube defects (NTD).

MEASURED SERUM VALUES

	Value	Corr. MoMs
AFP	60.5 ng/ml	1.45
HCG	47647 mIU/ml	1.58
uE3	1 ng/ml	1.35
Gestation age	16+ 3	
Method	Scan	

The MoMs have been corrected according to: maternal weight ethnic origin

TRISOMY 21 SCREENING

The calculated risk for Trisomy 21 is below the cut off which represents a low risk.

After the result of the Trisomy 21 test it is expected that among 3006 women with the same data, there is one woman with a trisomy 21 pregnancy and 3005 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!

TRISOMY 18 SCREENING

The calculated risk for trisomy 18 is < 1:10000, which indicates a low risk.

NEURAL TUBE DEFECTS (NTD) SCREENING

The corrected MoM AFP (1.45) is located in the low risk area for neural tube defects.





