PRISCA 5.1.0.17

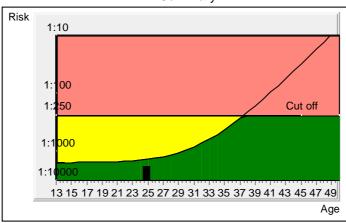
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6349/1, NICHOLSON ROAD, AMBALA CANTT

Results for:Sample noDate of report:MRS. MANPREET KAUR2203220602/AMB22-03-2022

Referring Doctors

Summary



Patient data	
Age at delivery	24.8
WOP	15 + 2
Weight	52 kg
Patient ID	
Ethnic origin	Asian

For MRS. MANPREET KAUR, born on 01-12-1997, a screening test was performed on the 21-03-2022. Prisca screens for Trisomy 21, Trisomy 18 and Neural tube defects (NTD).

MEASURED SERUM VALUES

	Value		Corr. MoMs
AFP	20.6	ng/ml	0.64
HCG	45617	miu/ml	1.09
uE3	0.35	ng/ml	0.85
Gestation a	ge	15+2	
Method		Scan	

The MoMs have been corrected according to: maternal weight

ethnic origin

Risks at term		
Biochemical risk for Tr.21	1:1783	
Age risk:	1:1390	
Neural tube defects risk	<1:10000	

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 1783 women with the same data, there is one woman with a trisomy 21 pregnancy and 1782 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 (0.64) is located in the low risk area for neural tube defects.





