PRISCA 5.1.0.17

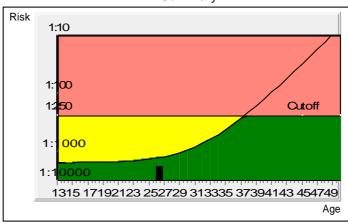
Licenced for: KOS DIAGNOSTIC LAB

6349/1, NICHOLSON ROAD, AMBALA CANTT

Results for:Sample noDate of report:MRS. MANJIT KAUR1910220963/AMB18-10-2019

Referring Doctors

Summary



Patient	data
Age at delivery	26.3
WOP	15 + 4
Weight	52 kg
Patient ID	
Ethnic origin	Asian

For MRS. MANJIT KAUR, born on 30-11-1993, a screening test was performed on the 17-10-2019. Prisca screens for Trisomy 21, Trisomy 18 and Neural tube defects (NTD).

MEASURED SERUM VALUES

	Value		Corr. MoMs
AFP	34.7	ng/ml	0.85
HCG	22183	mIU/mI	0.59
uE3	0.53	ng/ml	0.95
Gestation ag Method	е	15+ 4 Scan	

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

Risks at term		
Biochemical risk for Tr.21	1:7713	
Age risk:	1:1296	
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 7713 women with the same data, there is one woman with a trisomy 21 pregnancy and 7712 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 (0.85) is located in the low risk area for neural tube defects.





