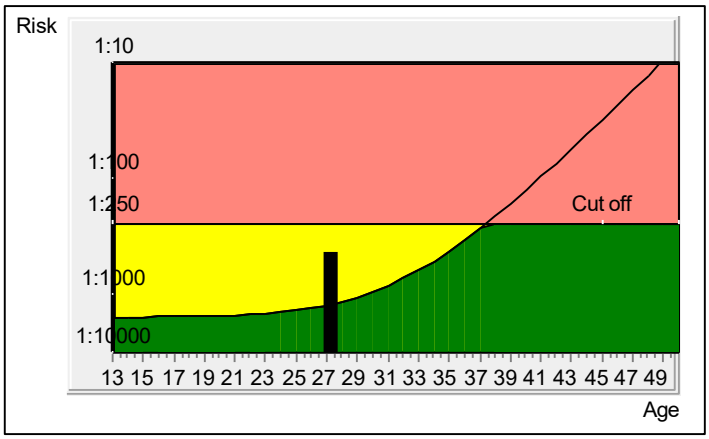
Results for: MRS. RAVNEET KAUR	Sample no 2410045862/NOD	Date of report: 24/10/2024
Referring Doctors	JITM Diagnostics	

Summary



Patient data	
Age at delivery	27.3
WOP	14 + 1
Weight	43 kg
Patient ID	
Ethnic origin	Asian

Risks at term	
Biochemical risk for Tr.21	1:442
Age risk:	1:1226
Neural tube defects risk	1:6370

For MRS. RAVNEET KAUR , born on 8/01/1998, a screening test was performed on the 24/10/2024. Prisca screens for Trisomy 21, Trisomy 18 and Neural tube defects (NTD).

MEASURED SERUM VALUES

	Value		Corr. MoMs
AFP	28.1	ng/ml	0.78
HCG	165123	mIU/ml	2.90
uE3	1.13	ng/ml	1.51
Gestation age		14+ 1	
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 442 women with the same data, there is one woman with a trisomy 21 pregnancy and 441 women with not affected pregnancies.
The HCG level is high.
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 (0.78) is located in the low risk area for neural tube defects.

Risk above Cut off

Risk above Age risk

Risk below Age risk