

Results for:
MRS. KAJAL KUMARI

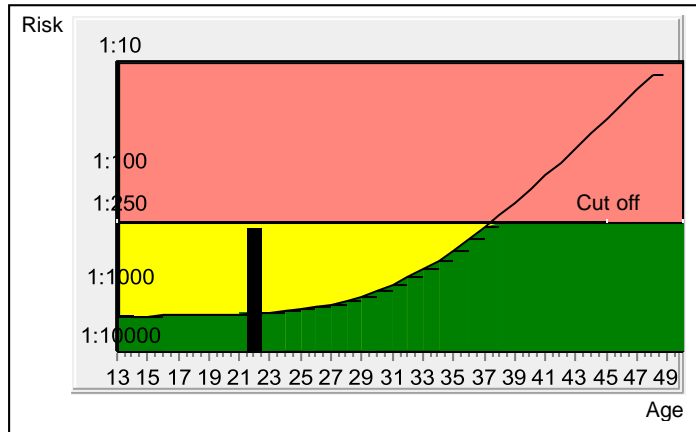
Sample no
2407025992/NOD

Date of report:
14/07/2024

Referring Doctors

JITM Diagnostics

Summary



Patient data	
Age at delivery	22.0
WOP	14 + 2
Weight	52 kg
Patient ID	
Ethnic origin	Asian

Risks at term	
Biochemical risk for Tr.21	1:280
Age risk:	1:1495
Neural tube defects risk	1:7235

For MRS. KAJAL KUMARI , born on 9/01/2003, a screening test was performed on the 13/07/2024. Prisca screens for Trisomy 21, Trisomy 18 and Neural tube defects (NTD).

MEASURED SERUM VALUES

	Value	Corr. MoMs
AFP	14.5 ng/ml	0.46
HCG	75879 mIU/ml	1.55
uE3	0.5 ng/ml	0.68

Gestation age 14+ 2
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 280 women with the same data, there is one woman with a trisomy 21 pregnancy and 279 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 (0.46) is located in the low risk area for neural tube defects.

Risk above Cut off

Risk above Age risk

Risk below Age risk