

PRISCA 5.1.0.17

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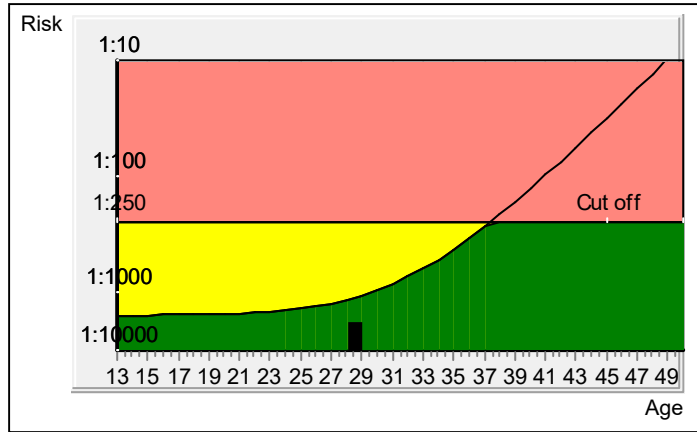
Results for:
Mrs. SANGITA PAWAR

Sample no
A0544096

Date of report:
18/06/24

Referring Doctors

Summary



Patient data	
Age at delivery	28.5
WOP	14 + 2
Weight	43 kg
Patient ID	0662406140240
Ethnic origin	Asian

Risks at term	
Biochemical risk for Tr.21	1:5309
Age risk:	1:1116
Neural tube defects risk	1:7015

For Mrs. SANGITA PAWAR, born on 01-06-1996, a screening test was performed on the 14-06-2024. Prisca screens for Trisomy 21, Trisomy 18 and Neural tube defects (NTD).

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 5309 women with the same data, there is one woman with a trisomy 21 pregnancy and 5308 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!

MEASURED SERUM VALUES

	Value	Corr. MoMs
AFP	28 ng/mL	0.85
HCG	37663.45 mIU/mL	0.71
uE3	0.65 ng/mL	2.15
Gestation age	14+ 2	
Method	Scan	

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

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.

Risk above Cut off

Risk above Age risk

Risk below Age risk