

PRISCA 5.1.0.17

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Results for:
Ms. SANTOSHI PARMANIK

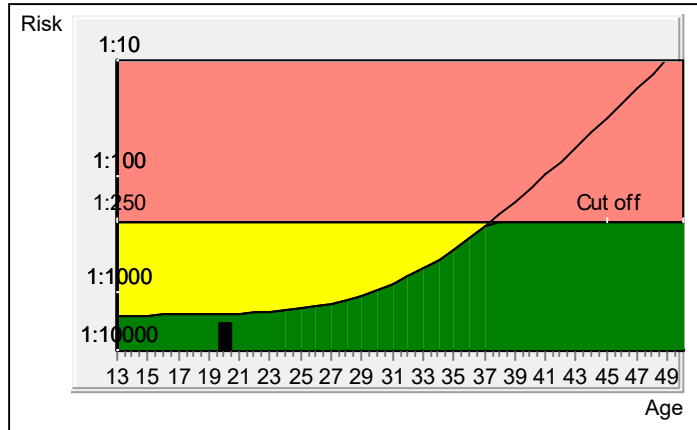
Sample no
25059777

Date of report:
14/12/23

Referring Doctors

NA

Summary



Patient data	
Age at delivery	20.1
WOP	20 + 0
Weight	56 kg
Patient ID	0502312130003
Ethnic origin	Asian

Risks at term	
Biochemical risk for Tr.21	1:6082
Age risk:	1:1535
Neural tube defects risk	<1:10000

For Ms. SANTOSHI PARMANIK, born on 02-04-2004, a screening test was performed on the 12-12-2023. Prisca screens for Trisomy 21, Trisomy 18 and Neural tube defects (NTD).

MEASURED SERUM VALUES

	Value	Corr. MoMs
AFP	63.72 ng/mL	0.93
HCG	15247.04 mIU/mL	0.84
uE3	1.41 ng/mL	0.89

Gestation age 20+ 0
Method BPD Hadlock

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 6082 women with the same data, there is one woman with a trisomy 21 pregnancy and 6081 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 (0.93) is located in the low risk area for neural tube defects.

Risk above
Cut off

Risk above
Age risk

Risk below
Age risk