

PRISCA 5.1.0.17

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Results for:

Ms. SANTOSHI

Sample no

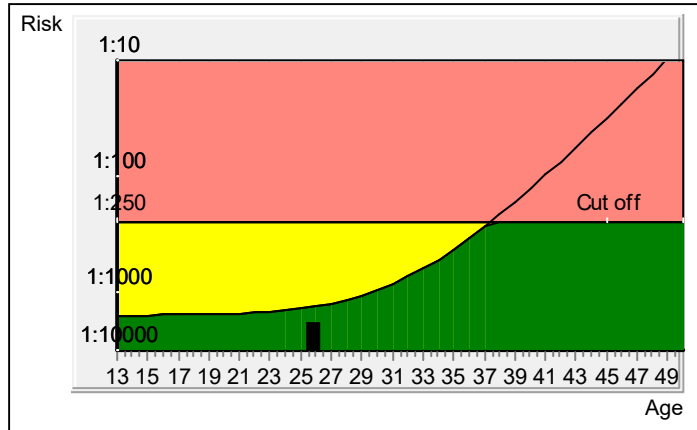
23540473

Date of report:

04-05-2022

Referring Doctors

Summary



Patient data

Age at delivery 25.8
WOP 20 + 0
Weight 49 kg
Patient ID 0012205040151
Ethnic origin Asian

Risks at term

Biochemical risk for Tr.21 1:3830
Age risk: 1:1332
Neural tube defects risk <1:10000

For Ms. SANTOSHI, born on 01-12-1996, a screening test was performed on the 04-05-2022. Prisca screens for Trisomy 21, Trisomy 18 and Neural tube defects (NTD).

MEASURED SERUM VALUES

	Value	Corr. MoMs
AFP	69.4 ng/mL	0.91
HCG	21414.22 mIU/mL	1.09
uE3	1.65 ng/mL	1.00

Gestation age 20+ 0
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 3830 women with the same data, there is one woman with a trisomy 21 pregnancy and 3829 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 (0.91) is located in the low risk area for neural tube defects.

Risk above
Cut off

Risk above
Age risk

Risk below
Age risk