

PRISCA 5.1.0.17

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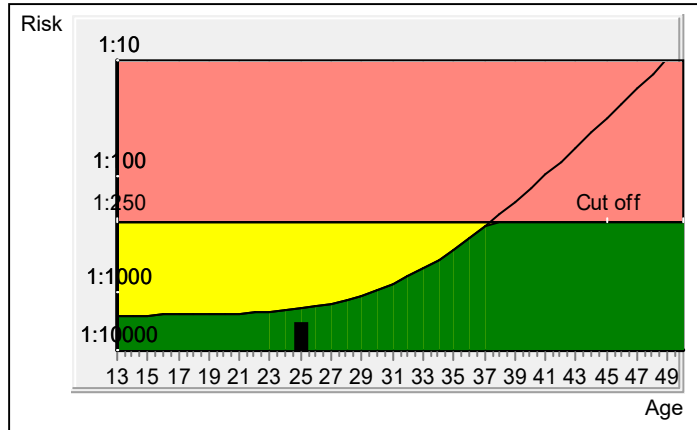
Results for:
Mrs. SANTHOSHA

Sample no
B4145809

Date of report:
11-01-2026

Referring Doctors

Summary



Patient data	
Age at delivery	25.0
WOP	16 + 0
Weight	63.6 kg
Patient ID	0012601100351
Ethnic origin	Asian

Risks at term	
Biochemical risk for Tr.21	1:2651
Age risk:	1:1377
Neural tube defects risk	1:1792

For Mrs. SANTHOSHA , born on 15-06-2001, a screening test was performed on the 10-01-2026. Prisca screens for Trisomy 21, Trisomy 18 and Neural tube defects (NTD).

MEASURED SERUM VALUES

	Value	Corr. MoMs
AFP	54.4 ng/mL	1.68
HCG	69693 mIU/mL	2.38
uE3	0.58 ng/mL	1.11

Gestation age 16+ 0
Method CRL Robinson

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 2651 women with the same data, there is one woman with a trisomy 21 pregnancy and 2650 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 (1.68) is located in the low risk area for neural tube defects.

Risk above
Cut off

Risk above
Age risk

Risk below
Age risk