

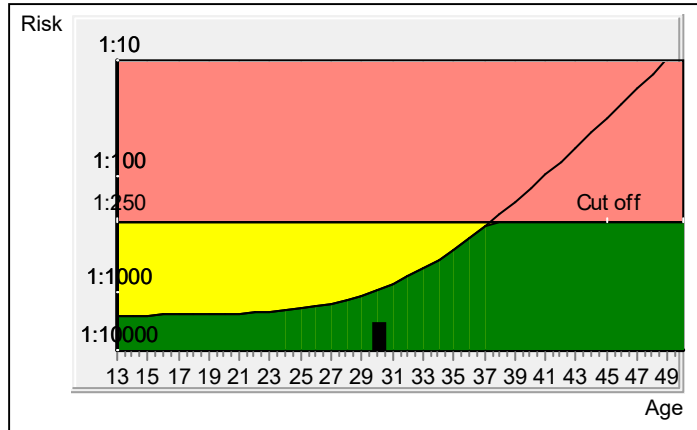
PRISCA 5.1.0.17

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Results for: Mrs. SANDHYA KUMARI	Sample no 24728092	Date of report: 21-09-2023
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Referring Doctors	NA
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Summary



Patient data	Risks at term
Age at delivery 30.2	Biochemical risk for Tr.21 1:2548
WOP 18 + 2	Age risk: 1:945
Weight 52.8 kg	Neural tube defects risk <1:10000
Patient ID 0472309200010	
Ethnic origin Asian	

For Mrs. SANDHYA KUMARI, born on 15-12-1993, a screening test was performed on the 20-09-2023. Prisca screens for Trisomy 21, Trisomy 18 and Neural tube defects (NTD).

MEASURED SERUM VALUES

	Value	Corr. MoMs
AFP	39.2 ng/mL	0.72
HCG	17797 mIU/mL	0.81
uE3	0.95 ng/mL	0.87

Gestation age 18+ 2
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 2548 women with the same data, there is one woman with a trisomy 21 pregnancy and 2547 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 (0.72) is located in the low risk area for neural tube defects.

Risk above
Cut off

Risk above
Age risk

Risk below
Age risk