

PRISCA 5.1.0.17

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Results for:

Mrs. SARITHA

Sample no

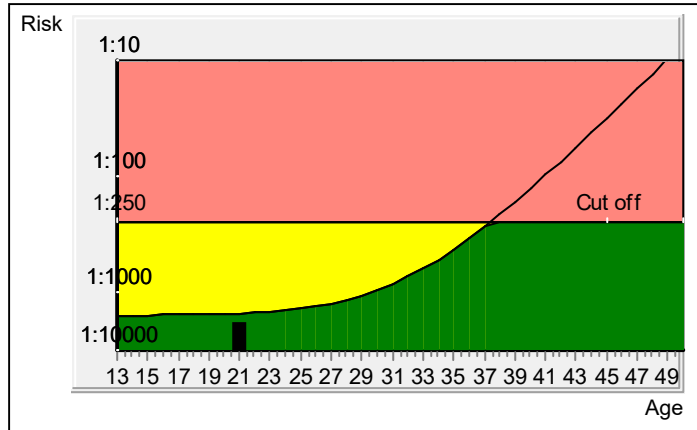
23043318

Date of report:

27/01/22

Referring Doctors

Summary



Patient data

Age at delivery 21.0
WOP 20 + 0
Weight 53 kg
Patient ID 0012201221261
Ethnic origin Asian

Risks at term

Biochemical risk for Tr.21 <1:10000
Age risk: 1:1519
Neural tube defects risk <1:10000

For Mrs. SARITHA, born on 16-06-2001, a screening test was performed on the 23-01-2022. Prisca screens for Trisomy 21, Trisomy 18 and Neural tube defects (NTD).

MEASURED SERUM VALUES

	Value	Corr. MoMs
AFP	49.16 ng/mL	0.74
HCG	7460.6 mIU/mL	0.40
uE3	1.56 ng/mL	0.97

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 more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy.

The HCG level is low.

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 (0.74) is located in the low risk area for neural tube defects.

Risk above
Cut off

Risk above
Age risk

Risk below
Age risk