

PRISCA 5.1.0.17

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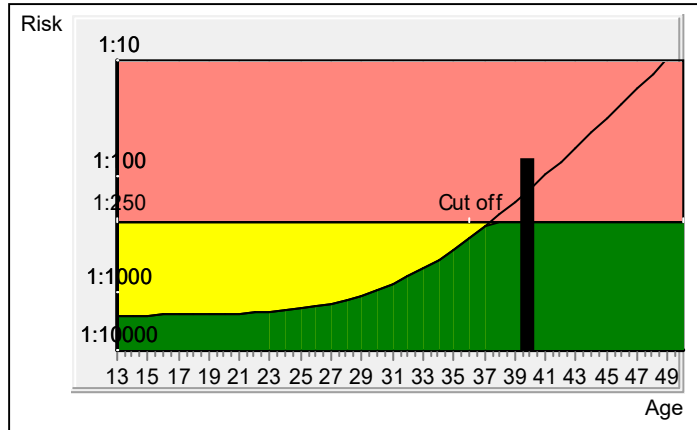
Results for:
Mrs. KHAMRUNNISA

Sample no
b4284687

Date of report:
20/12/25

Referring Doctors

Summary



Patient data	
Age at delivery	39.9
WOP	21 + 0
Weight	101 kg
Patient ID	0012512190404
Ethnic origin	Asian

Risks at term	
Biochemical risk for Tr.21	1:69
Age risk:	1:133
Neural tube defects risk	<1:10000

For Mrs. KHAMRUNNISA , born on 22-06-1986, a screening test was performed on the 19-12-2025. Prisca screens for Trisomy 21, Trisomy 18 and Neural tube defects (NTD).

MEASURED SERUM VALUES

	Value	Corr. MoMs
AFP	37.5 ng/mL	0.70
HCG	27302 mIU/mL	2.08
uE3	1.71 ng/mL	1.02

Gestation age 21+ 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 above the cut off which represents an increased risk.

After the result of the Trisomy 21 test it is expected that among 69 women with the same data, there is one woman with a trisomy 21 pregnancy and 68 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 (0.70) is located in the low risk area for neural tube defects.

Risk above
Cut off

Risk above
Age risk

Risk below
Age risk