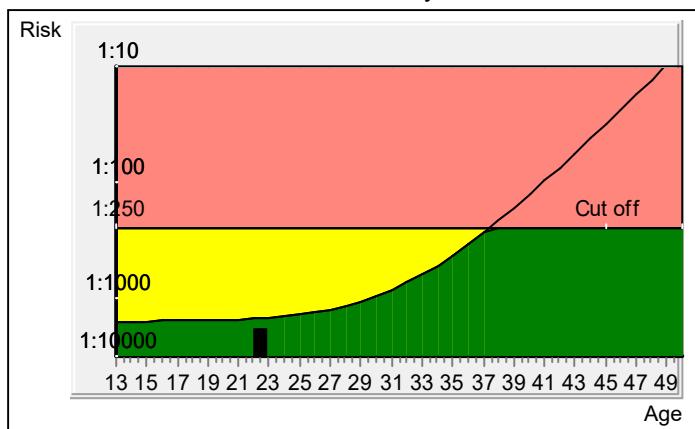


Results for:
Ms. KAVITA RUHELASample no
A1987953Date of report:
08-03-2025

Referring Doctors

Summary



Patient data

Age at delivery	22.4
WOP	14 + 2
Weight	67 kg
Patient ID	0382503070126
Ethnic origin	Asian

Risks at term

Biochemical risk for Tr.21	1:1853
Age risk:	1:1483
Neural tube defects risk	1:2520

For Ms. KAVITA RUHELA, born on 30-03-2003, a screening test was performed on the 07-03-2025. Prisca screens for Trisomy 21, Trisomy 18 and Neural tube defects (NTD).

MEASURED SERUM VALUES

	Value	Corr. MoMs
AFP	31.5 ng/mL	1.34
HCG	97038.6 mIU/mL	2.43
uE3	0.52 ng/mL	1.98
Gestation age	14+ 2	
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 1853 women with the same data, there is one woman with a trisomy 21 pregnancy and 1852 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 (1.34) is located in the low risk area for neural tube defects.



Risk above
Cut off



Risk above
Age risk



Risk below
Age risk