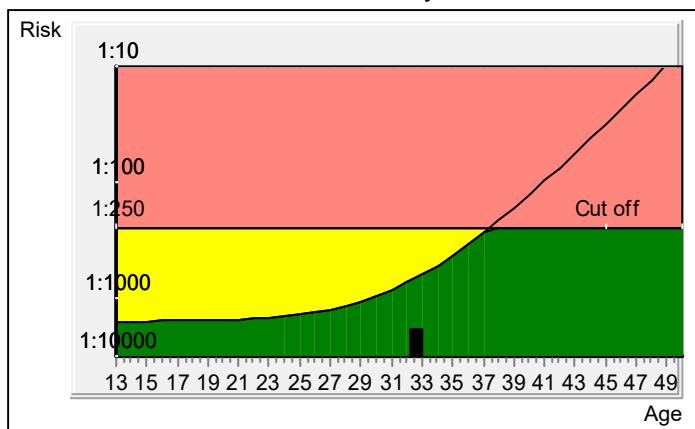


Results for:
Mrs. FARHANAZSample no
A1814119Date of report:
05/12/24

Referring Doctors

Summary



Patient data	
Age at delivery	32.6
WOP	20 + 1
Weight	67 kg
Patient ID	0012412040304
Ethnic origin	Asian

Risks at term	
Biochemical risk for Tr.21	1:4658
Age risk:	1:667
Neural tube defects risk	1:3445

For Mrs. FARHANAZ, born on 31-08-1992, a screening test was performed on the 04-12-2024. Prisca screens for Trisomy 21, Trisomy 18 and Neural tube defects (NTD).

MEASURED SERUM VALUES

	Value	Corr. MoMs
AFP	91.8 ng/mL	1.49
HCG	15196.07 mIU/mL	0.94
uE3	1.45 ng/mL	0.94
Gestation age	20+1	
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 4658 women with the same data, there is one woman with a trisomy 21 pregnancy and 4657 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 (1.49) is located in the low risk area for neural tube defects.



Risk above
Cut off



Risk above
Age risk



Risk below
Age risk