

PRISCA 5.1.0.17

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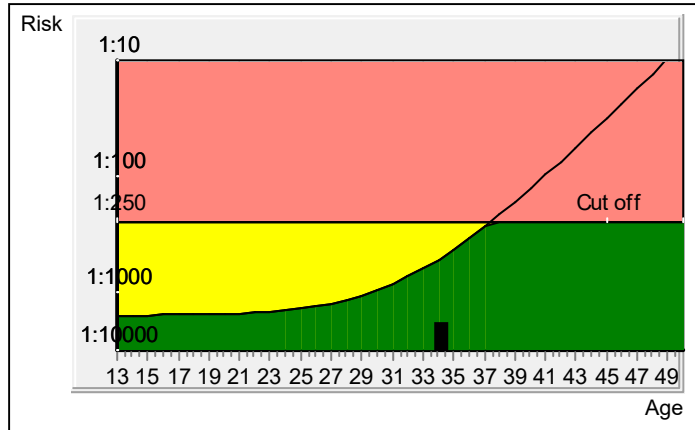
Results for:
Mrs. HOMAİN WEİNGKEN

Sample no
24933877

Date of report:
09-11-2023

Referring Doctors

Summary



Patient data	
Age at delivery	34.1
WOP	18 + 2
Weight	58 kg
Patient ID	0692311060059
Ethnic origin	Asian

Risks at term	
Biochemical risk for Tr.21	1:5266
Age risk:	1:509
Neural tube defects risk	1:527

For Mrs. HOMAİN WEİNGKEN, born on 14-02-1990, a screening test was performed on the 05-11-2023. Prisca screens for Trisomy 21, Trisomy 18 and Neural tube defects (NTD).

MEASURED SERUM VALUES

	Value	Corr. MoMs
AFP	105.6 ng/mL	2.09
HCG	19888.89 mIU/mL	0.96
uE3	1.12 ng/mL	1.06

Gestation age 18+ 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 5266 women with the same data, there is one woman with a trisomy 21 pregnancy and 5265 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 (2.09) is located in the low risk area for neural tube defects.

Risk above
Cut off

Risk above
Age risk

Risk below
Age risk