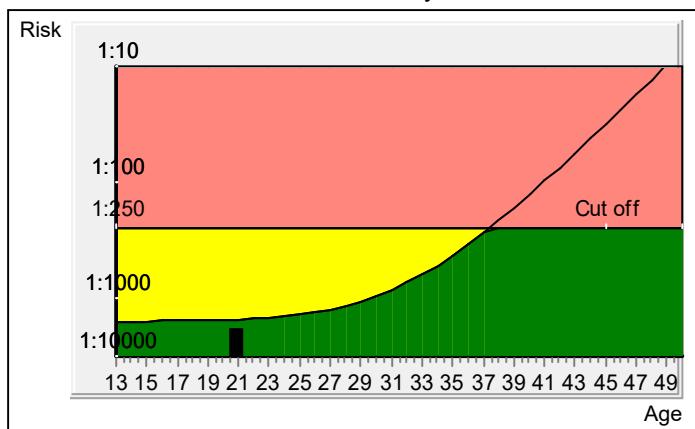


Results for:  
Mrs. PAYAL SHRESHTHSample no  
A1060125Date of report:  
18-07-2024

Referring Doctors

## Summary



## Patient data

Age at delivery	20.8
WOP	14 + 4
Weight	68 kg
Patient ID	0672407150161
Ethnic origin	Asian

## Risks at term

Biochemical risk for Tr.21	<1:10000
Age risk:	1:1522
Neural tube defects risk	1:78

For Mrs. PAYAL SHRESHTH, born on 13-03-2004, a screening test was performed on the 15-07-2024. Prisca screens for Trisomy 21, Trisomy 18 and Neural tube defects (NTD).

## MEASURED SERUM VALUES

	Value	Corr. MoMs
AFP	59.11 ng/mL	2.42
HCG	35598.3 mIU/mL	0.95
uE3	0.53 ng/mL	1.78
Gestation age	14+ 4	
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 more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy.

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



Risk above  
Cut off



Risk above  
Age risk



Risk below  
Age risk