

PRISCA 5.1.0.17

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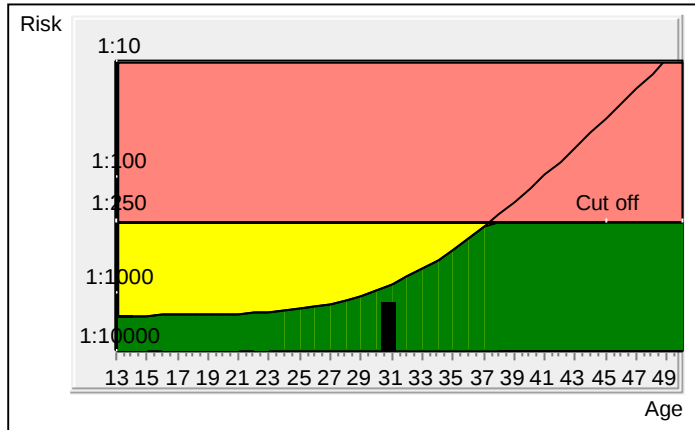
Results for:
Mrs. KOMAL PAILWAN

Sample no
A0162346

Date of report:
06-03-2024

Referring Doctors

Summary



Patient data		Risks at term	
Age at delivery	30.8	Biochemical risk for Tr.21	1:1240
WOP	19 + 6	Age risk:	1:876
Weight	62.2 kg	Neural tube defects risk	<1:10000
Patient ID	0662403040160		
Ethnic origin	Asian		

For Mrs. KOMAL PAILWAN, born on 06-10-1993, a screening test was performed on the 04-03-2024. Prisca screens for Trisomy 21, Trisomy 18 and Neural tube defects (NTD).		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 1240 women with the same data, there is one woman with a trisomy 21 pregnancy and 1239 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!	
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MEASURED SERUM VALUES		
	Value	Corr. MoMs
AFP	45.85 ng/mL	0.74
HCG	24144.5 mIU/mL	1.42
uE3	1.96 ng/mL	1.32
Gestation age	19+ 6	
Method	BPD Hadlock	
The MoMs have been corrected according to: maternal weight ethnic origin		

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 (0.74) is located in the low risk area for neural tube defects.	
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Risk above
Cut off

Risk above
Age risk

Risk below
Age risk