

PRISCA 5.1.0.17

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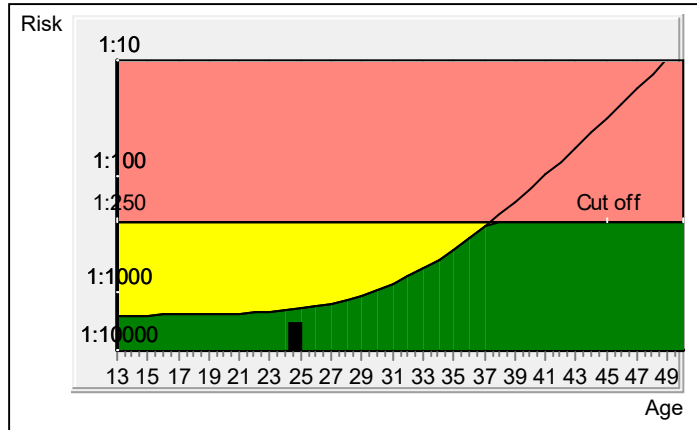
Results for:
Mrs. SANGITA KUMARI

Sample no
L2308838

Date of report:
20-08-2022

Referring Doctors

Summary



Patient data		Risks at term	
Age at delivery	24.7	Biochemical risk for Tr.21	<1:10000
WOP	18 + 1	Age risk:	1:1394
Weight	48 kg	Neural tube defects risk	1:1497
Patient ID	0482208160121		
Ethnic origin	Asian		

For Mrs. SANGITA KUMARI, born on 05-05-1998, a screening test was performed on the 16-08-2022. 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 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!	
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MEASURED SERUM VALUES		
	Value	Corr. MoMs
AFP	104.38 ng/mL	1.83
HCG	21544.22 mIU/mL	0.91
uE3	1.25 ng/mL	1.16
Gestation age	18+ 1	
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 (1.83) 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