

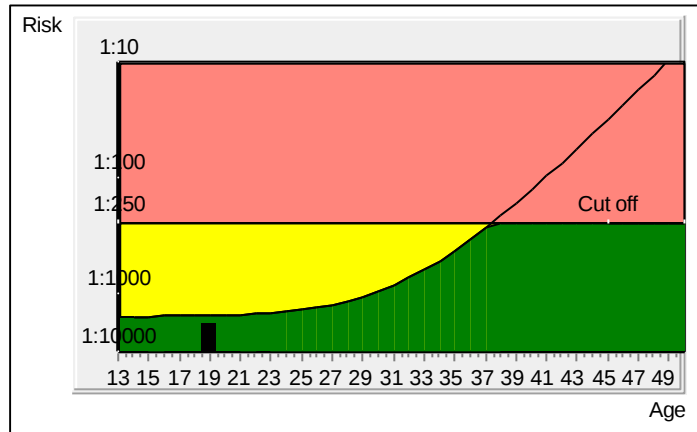
PRISCA 5.1.0.17

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Results for: Miss. RAJNANDANI KUMARI	Sample no A0814378	Date of report: 11-08-2024
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Referring Doctors

Summary



Patient data		Risks at term	
Age at delivery	18.9	Biochemical risk for Tr.21	1:5484
WOP	17 + 2	Age risk:	1:1552
Weight	42 kg	Neural tube defects risk	<1:10000
Patient ID	0482408090172		
Ethnic origin	Asian		

For Miss. RAJNANDANI KUMARI, born on 02-03-2006, a screening test was performed on the 09-08-2024. 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 5484 women with the same data, there is one woman with a trisomy 21 pregnancy and 5483 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	43.58 ng/mL	0.79
HCG	20937.07 mIU/mL	0.70
uE3	0.72 ng/mL	0.80
Gestation age	17+ 2	
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 (0.79) 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