

Risk Above Cut Off



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Risk below Age risk

 Date of Report
 24-06-2021

 PRISCA
 5.0.2.37

				FMSCA	3.0.2.37
Patient Data					
Name		MRS CHARU	Patient ID		012106220176
Birthday		26-05-1991	Sample ID		10823675
Age at delivery		30.1	Sample Date		22/06/2021
Gestational age		12+2			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	89 Diabete	S	unknown	Pregnancies	unknown
Smoker	Unknown Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	2	12+0
PAPP-A	5.35 mIU/ml	2.12	Method		CRL(<>Robinson
fb-hCG	98.5 ng/ml	2.3	Scan date		21-06-2021
Risks at sampling date			Crown rump length in mm 53.7		
Age Risk		1:634	Nuchal translucency MOM		0.98
Biochemical T21 risk		1:1890	Nasal bone		Present
Combined Trisomy 21	Risk	1:7074	Sonographer		DR.PRINCY SETHI
Trisomy 13/18 + NT		<1:10000	Qualification in	n measuring NT	MBBS
Risk 1:10			Down's Syndrome Risk (Trisomy 21 Screening)		
			The calculated risk for Trisomy 21 (with NT) is below the cut off, which represents a low risk.		
1:: 00 1::250			After the result of the Trisomy 21 test (with NT) it is expected that among more than 7074 women with the same data, there is one woman with a trisomy 21 pregnancy and 7073 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 the risk calculations are statistical aapproaches and have no diagnostic value!		
Trisomy 13/18 + NT The calculated risk for Trisomy 13/18 (with NT) is <1:10000, which indicates a low risk			The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values		
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