

Risk Above Cut Off

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Risk below Age risk

 Date of Report
 04-11-2020

 PRISCA
 5.0.2.37

				TRISCA	5.0.2.37
Patient Data					
Name MRS VAISHALI			Patient ID		012011020101
Birthday		22-07-1999	Sample ID		10663403
Age at delivery		21.3	Sample Date		02/11/2020
Gestational age		12+0			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	74 Diabetes		unknown	Pregnancies	unknown
Smoker	Unknown Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	2	11+6
PAPP-A	$1.75 \mathrm{mIU/ml}$	0.61	Method		CRL(<>Robinson
fb-hCG	45.9 ng/ml	0.99	Scan date		02-11-2020
Risks at sampling date			Crown rump length in mm 51.1		
Age Risk 1:1049		Nuchal translucency MOM 0.87			
Biochemical T21 risk		1:2151	Nasal bone		Present
Combined Trisomy 21 Risk <1:10000		<1:10000	Sonographer		t.SANJEEV KUMAR SINGHAL
Trisomy 13/18 + NT		<1:10000	Qualification in	n measuring NT	MBBS,PGDUS,DMRD
Risk			Down's Syndro	ome Risk (Trisomy	21 Screening)
1:100 1:250 Cut off 1:1000 1:1000 1:10000 1:10000 1:10000 1:10000 Age			The calculated risk for Trisomy 21 (with NT) is below the cut off, which represents a low risk. After the result of the Trisomy 21 test (with NT) it is expected that among more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy and 9999 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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