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				Date of Report PRISCA	22/7/2023 5.2.0.13
Patient Data				1110011	
Name]	MRS. SWATI	Patient ID		012307210137
Birthday		4/6/2000	Sample ID		11778983
Age at term		23.7	Sample Date		21/7/2023
Gestational age		12+5			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	46 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom			12+3
PAPP-A	4.25 mIU/ml	0.54	Method		CRL (<>Robinson)
fb-hCG	23.4 ng/ml	0.6	Scan date		19/7/2023
Risks at sampling date			Crown rump length in mm 58.9		
Age Risk	1:1033		Nuchal translucency MoM 0.7		
Biochemical T21 risk		1:4284	Nasal bone		Present
Combined trisomy 21 ri	sk	<1:10000			
Trisomy 13/18 + NT		<1:10000			
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
1:10 1:100 1:250 Cut off 1:100			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 with NT test it is expected that among 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!		
1:10000 13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49 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 Risk above Age Risk		