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				Date of Report PRISCA	13/2/2023 5.1.0.17
Patient Data					
Name	MRS. JYOTI 8370		Patient ID		012302100284
Birthday		11/2/1996	Sample ID		11476511
Age at term		27.6	Sample Date		11/2/2023
Gestational age		13+6			
Correction factors	T				
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	70 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom			13+5
PAPP-A	3.15 mIU/ml	0.44	Method		CRL (<>Robinson)
fb-hCG	57.6 ng/ml	2.44	Scan date		10/2/2023
Risks at sampling date			Nuchal translucency 2		
Age Risk	1:898		Nuchal translucency MoM 1.09		
Biochemical T21 risk	emical T21 risk 1:87		Nasal bone Present		
Combined trisomy 21 risk 1:143		1:143			
Trisomy 13/18 + NT		<1:10000			
Risk			Down's Syndro	ome Risk (Trisomy 21	Screening)
Risk 1:10 1:250 Cut off 1:1000 1:10000 1:10000 13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49 Age			The calculated risk for Trisomy 21 (with NT) is above the cut off, which represents a high risk. After the result of the Trisomy 21 with NT test it is expected that among 143 women with the same data, there is one woman with a trisomy 21 pregnancy and 142 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 Risk Above Cut Off			The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values Risk above Age Risk		