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				Date of Report PRISCA	3/4/2023 5.2.0.13
Patient Data					
Name	MRS. JYOTI	W/O URMIT	Patient ID		012304010165
Birthday		25/05/2001	Sample ID		11640294
Age at term		22.3	Sample Date		1/4/2023
Gestational age		12+5			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	49 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom			12+4
PAPP-A	3.89 mIU/ml	0.53	Method		CRL (<>Robinson)
fb-hCG	31.4 ng/ml	0.82	Scan date		31/3/2023
Risks at sampling date			Nuchal translucency 1.1		
Age Risk	1:669		Nuchal translucency MoM 0.7		
Biochemical T21 risk		1:2184	Nasal bone	Present	
Combined trisomy 21 risk		<1:10000			
Trisomy 13/18 + NT		<1:10000			
Risk		Down's Syndrome Risk (Trisomy 21 Screening)			
Risk 1:10 1:250 Cut off 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 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 with NT test 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!		
The calculated risk for Trisomy 13/18 (with NT) is <1:10000 , which indicates a low risk			on the risk assessment! Calculated risks have no diagnostic values		
Risk A	bove Cut Off		Risk above Ag	e Risk	Risk below Age risk