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				Date of Report PRISCA	30/1/2023 5.1.0.17
Patient Data					
Name		MRS. JYOTI	Patient ID		012301290036
Birthday		12/10/1993	Sample ID		011489007
Age at term		29.09	Sample Date		29/1/2023
Gestational age		12+4			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	55 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom			12+2
PAPP-A	3.45 mIU/ml	0.56	Method		CRL (<>Robinson)
fb-hCG	15.9 ng/ml	0.42	Scan date		26/1/2023
Risks at sampling date			Nuchal translucency 2.9		
Age Risk		1:702	Nuchal translucency MoM		1.93
Biochemical T21 risk		1:6678	Nasal bone Pre		Present
Combined trisomy 21 risk		1:914			
Trisomy 13/18 + NT		1:353			
Risk			Down's Syndry	ome Risk (Trisomy 21	Screening)
Risk 1:10 1:10 1:250 Cut off 1:10 1:1000			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 914 women with the same data, there is one woman with a trisomy 21 pregnancy and 913 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 laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values		
, which indicates a low r					Risk below Age risk