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					Date of Report PRISCA	12/6/2023 5.1.0.17
Patient Data						
Vame MRS. PALLAVI				Patient ID		012306110221
Birthday			5/1/2002	Sample ID		11644947
Age at term			21.1	Sample Date		11/6/2023
Gestational age			13+1			
Correction factors						
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown
Weight in kg	53	Diabetes		NO	Pregnancies	unknown
Smoker	NO	Origin		Asian		
Biochemical Data				Ultrasound Data		
Parameter	Value		Corr Mom	Gestational age	2	13+0
PAPP-A	5.4	mIU/ml	0.68	Method		CRL (<>Robinson)
fb-hCG	31.9 ng/ml		0.97	Scan date		10/6/2023
Risks at sampling date				Nuchal translucency (NT) 2.3		
Age Risk			1:1088	Nuchal translucency MoM		1.34
Biochemical T21 risk		1:3017	Nasal bone		present	
Combined trisomy 21 risk			1:4287			
Trisomy 13/18			<1:10000			
Risk				Down's Syndrome Risk (Trisomy 21 Screening)		
Risk 1:10 1:200 1:450 Cut off 1:1000 1:1000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:1000 1:1000 Age Trisomy 13/18+NT The calculated risk for Trisomy 13/18 (with NT) is <1:10000 + 1:1000 + 1:100000 + 1:10000000 + 1:10000000000				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 it is expected that among 4287 women with the same data, there is one woman with a trisomy 21 pregnancy and 4286 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		
	bove Ci			Risk above Ag	e Risk	Risk below Age risk