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Patient Data Name				PRISCA	5.1.0.17
Name					
	MRS	5. PRIYANKA	Patient ID		102301170028
Birthday		5/12/1992	Sample ID		11556097
Age at term		30.07	Sample Date		17/1/2023
Gestational age		13+3			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	57 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom			13+1
PAPP-A	4.84 mIU/ml	0.61	Method		CRL (<>Robinson)
fb-hCG	32.7 ng/ml	1.12	Scan date		15/1/2023
Risks at sampling date			Nuchal translucency 1.7		
Age Risk	e Risk 1:657		Nuchal translucency MoM 0.93		
Biochemical T21 risk		1:982	Nasal bone		Present
Combined trisomy 21 risk	X	1:4535			
Trisomy 13/18 + NT		<1:10000			
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
Risk 1:10 1:250 1:250 1:1000 1:1000 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 4535 women with the same data, there is one woman with a trisomy 21 pregnancy and 4534 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		