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				Date of Report PRISCA	3/8/2022 5.1.0.17
Patient Data					
Name	MRS	5. PRIYANKA	Patient ID		012208010126
Birthday		21/03/1994	Sample ID		11220920
Age at term		28.1	Sample Date		1/8/2022
Gestational age		13+1			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	46 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational ag	2	12+0
PAPP-A	4.32 mIU/ml	0.47	Method		CRL (<>Robinson)
fb-hCG	31.4 ng/ml	0.91	Scan date		25/7/2022
Risks at sampling date			Nuchal translucency 1.2		
Age Risk		1:786	Nuchal translu	cency MoM	0.85
Biochemical T21 risk	1:937		Nasal bone	ne Present	
Combined trisomy 21 risk	_	1:5428			
Trisomy 13/18 + NT		<1:10000			
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
Risk 1:10 1:250 1:250 Cut off 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:100 1:100 1:100 1:100 1:250 Cut off 1:100 1:250 Cut off 1:100 1:250 Cut off 1:100 1:250 Cut off 1:100 1:250 Cut off 1:100			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 5428 women with the same data, there is one woman with a trisomy 21 pregnancy and 5427 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		
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	Above Cut Off		Risk above Ag	e Risk 📃 F	Risk below Age risk