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				Date of Report PRISCA	27/2/2023 5.2.0.13
Patient Data					
Name	MRS	5. ANUPAMA	Patient ID		012302250369
Birthday		16/01/1991	Sample ID		11574541
Age at term		32.7	Sample Date		25/2/2023
Gestational age		12+6			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	49 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
<b>Biochemical Data</b>			Ultrasound Data		
Parameter	Value	Corr Mom			12+3
PAPP-A	4.72 mIU/ml	0.61	Method		CRL (<>Robinson)
fb-hCG	64.9 ng/ml	1.77	Scan date		23/2/2023
Risks at sampling date			Nuchal translucency 0.97		
Age Risk 1:481		1:481	Nuchal translucency MoM 0.		0.64
Biochemical T21 risk		1:244	Nasal bone		Present
Combined trisomy 21 ris	sk	1:1498			
Trisomy 13/18 + NT		<1:10000			
Risk			Down's Syndr	ome Risk (Trisomy 21	Screening)
Risk 1:10 1:10 1:250 Cut off 1:10000 1:1000 1:1000 1:1000 1:10000 1:10000 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:10			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 1498 women with the same data, there is one woman with a trisomy 21 pregnancy and 1497 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		