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				Date of Report PRISCA	5/5/2022 5.1.0.17
Patient Data					
Name	Ν	ARS. LALITA	Patient ID		012205030055
Birthday		15/02/1995	Sample ID		11447233
Age at term		27.8	Sample Date		3/5/2022
Gestational age		13+1			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	60.1 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	2	13+0
PAPP-A	3.62 mIU/ml	0.65	Method		CRL (<>Robinson)
fb-hCG	48.6 ng/ml	1.12	Scan date		2/5/2022
Risks at sampling date			Nuchal Translucency 1.13		
Age Risk	ge Risk 1:865		Nuchal Translucency MoM 0.6		0.67
Biochemical T21 risk 1:15		1:1533	Nasal Bone Prese		Present
Combined T21 risk		1:8679			
Trisomy 13/18+NT		<1:10000			
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
Risk 1:10			cut off, which	d risk for Trisomy 21 (v represents a low risk.	
1:100 1:250 Cut off 1:1000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10			After the result of the Trisomy 21 test it is expected that among 8679 women with the same data, there is one woman with a trisomy 21 pregnancy 8678 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!		
Trisomy 13/18+NT The calculated risk for Trisomy 13/18 (with NT) is <1:10000 , which indicates a low risk			The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values		
Ri	sk Above Cut Off		Risk above Ag	e Risk	Risk below Age risk