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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 <b>PRISCA</b> 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