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				Date of Report PRISCA	1/7/2022 5.1.0.17
Patient Data					
Name	MRS. G	EETA KAUR	Patient ID		012206290209
Birthday		13/07/1996	Sample ID		11304219
Age at term		26.02	Sample Date		29/6/2022
Gestational age		13+4	ł		
Correction factors				•	
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	60 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational ag	e	13+2
PAPP-A	5.89 mIU/ml	0.75	Method		CRL (<>Robinson)
fb-hCG	68.2 ng/ml	2.49	Scan date		28/06/2022
Risks at sampling date			Nuchal translucency 1.2		
Age Risk		1:958	Nuchal translu	icency MoM	0.68
Biochemical T21 risk		1:339	Nasal bone		Present
Combined trisomy 21 risk		1:2064			
Trisomy 13/18 + NT		<1:10000			
Risk			Down's Syndr	ome Risk (Trisomy 21	Screening)
Risk 1:10 1:250 Cut off 1:1000 1:250 Cut off 1:1000 1:1000 1:1000 1:1000 1:1000 Trisomy 13/18+NT The calculated risk for Trisomy 13/18 (with NT) is <1:1000 (with NT) is			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 2064 women with the same data, there is one woman with a trisomy 21 pregnancy and 2063 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		
The calculated risk for T <1:10000 , which indicate		1 NT) is	on the risk ass	essment! Calculate	