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				Date of Report PRISCA	16/10/2022 5.1.0.17
Patient Data					
Name		MRS. LAXMI	Patient ID		012210150073
Birthday		6/10/2001	Sample ID		11537565
Age at term		21.06	Sample Date		15/10/2022
Gestational age		13+3			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	53 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	e	13+3
PAPP-A	6.3 mIU/ml	0.72	Method		CRL (<>Robinson)
fb-hCG	18.5 ng/ml	0.62	Scan date		15/10/2022
Risks at sampling date			Nuchal translucency 1.6		
Age Risk 1:1105		1:1105	Nuchal translucency MoM 0.88		
Biochemical T21 risk	emical T21 risk 1:9255		Nasal bone Preser		
Combined trisomy 21 ris	k	<1:10000			
Trisomy 13/18 + NT		<1:10000			
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
Risk 1:10 1:250 Cut off 1:1000 1:1000 1:1000 1:10000 1:10000 1:10000			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 more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy and 9999 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 7 <1:10000 , which indicat	Trisomy 13/18 (with	Age	The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values Risk above Age Risk		