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				Date of Report PRISCA	12/3/2023 5.2.0.13
Patient Data					
Name	Ν	MRS. SAKSHI	Patient ID		012303110213
Birthday		24/11/1994	Sample ID		11487575
Age at term		28.9	Sample Date		11/3/2023
Gestational age		12+0			
Correction factors				<b>1</b>	
Fetuses	1 IVF		unknown	Previous trisomy	21 unknown
Weight in kg	65 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom			12+0
PAPP-A	4.5 mIU/ml	1.12	Method		CRL (<>Robinson)
fb-hCG	19.8 ng/ml	0.48	Scan date		11/3/2023
Risks at sampling date			Nuchal translucency 1.26		
Age Risk 1:760		1:760	Nuchal translucency MoM 0.9		
Biochemical T21 risk <1:10000		<1:10000	Nasal bone Presen		
Combined trisomy 21 risk <1:10000					
Trisomy 13/18 + NT		<1:10000			
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
Risk 1:10 1:250 Cut off 1:250 Cut off 1:1000 1:250 Cut off 1:1000 1:250 Cut off 1:1000 1:250 Cut off 1:1000 1:250 Cut off Trisomy 13/18+NT			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! The laboratory cannot be hold responsible for their impact on the risk calculated risk by preserve the referring physician.		
The calculated risk for Trisomy 13/18 (with NT) is <1:10000 , which indicates a low risk			on the risk assessment! Calculated risks have no diagnostic values		
Ri	sk Above Cut Off		Risk above Ag	e Risk	Risk below Age risk