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				Date of Report PRISCA	15/1/2023 5.1.0.17
Patient Data					
Name	MRS. POOJ	A CHAUHAN	Patient ID		012301130199
Birthday		5/7/1985	Sample ID		11515891
Age at term		37.11	Sample Date		13/1/2023
Gestational age		12+3			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	64 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom			12+2
PAPP-A	4.1 mIU/ml	0.84	Method		CRL (<>Robinson)
fb-hCG	27.8 ng/ml	0.74	Scan date		13/1/2023
Risks at sampling date			Nuchal translucency 1.1		
Age Risk	e Risk 1:150		Nuchal translucency MoM 0.73		
Biochemical T21 risk	1:1237		Nasal bone Presen		
Combined trisomy 21 ris	k	1:2572			
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 1:10000 1:10000 1:10000 1:10000 1:10000 Trisomy 13/18+NT The calculated risk for Trisomy 13/18 (with NT) is <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 2572 women with the same data, there is one woman with a trisomy 21 pregnancy and 2571 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 7 <1:10000 , which indicat		n NT) is	on the risk ass	essment! Calculated risl	