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				Date of Report PRISCA	7/1/2023 5.1.0.17
Patient Data					
Name		MRS. NISHA	Patient ID		012301060118
Birthday		8/4/1990	Sample ID		11523773
Age at term		33.01	Sample Date		6/1/2023
Gestational age		12+4			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	58 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom			12+4
PAPP-A	2.79 mIU/ml	0.48	Method		CRL (<>Robinson)
fb-hCG	15.8 ng/ml	0.42	Scan date		6/1/2023
Risks at sampling date			Nuchal translucency 1.3		
Age Risk		1:427	Nuchal translu	cency MoM	0.82
Biochemical T21 risk		1:2653	Nasal bone		Present
Combined trisomy 21 risk		<1:10000			
Trisomy 13/18 + NT		<1:10000			
Risk			Down's Syndro	ome Risk (Trisomy 21	l Screening)
Risk 1:10 1:100 1:250 Cut off			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.		
1:1000 1:10000 13 15 17 19 21 23 25 23 Trisomy 13/18+NT	41 43 45 47 49 Age	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			
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		
Risk Above Cut Off Risk above Age Risk Risk below Age risk					