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				Date of Report PRISCA	3/3/2023 5.2.0.13
Patient Data					
Name	MRS. PO	OOJA SINGH	Patient ID		012303010433
Birthday		29/09/1998	Sample ID		11514789
Age at term		24.1	Sample Date		1/3/2023
Gestational age		12+0			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	48 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom			11+6
PAPP-A	2.45 mIU/ml	0.43	Method		CRL (<>Robinson)
fb-hCG	18.9 ng/ml	0.41	Scan date		1/3/2023
Risks at sampling date			Nuchal translucency 0.8		
Age Risk	1:966		Nuchal translucency MoM 0.5		
Biochemical T21 risk	1:4568		Nasal bone	ne Present	
Combined trisomy 21 risk		<1:10000			
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:			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 Trisomy 13/18 (with NT) is <1:10000 , which indicates a low risk			The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values		
Risk A	Above Cut Off		Risk above Ag	e Risk	Risk below Age risk