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				Date of Report PRISCA	19/4/2023 5.2.0.13
Patient Data					
Name		MRS. KIRTI	Patient ID		012304170315
Birthday		15/4/1997	Sample ID		11635279
Age at term		26.6	Sample Date		17/4/2023
Gestational age		12+6			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	118 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom			12+5
PAPP-A	2.52 mIU/ml	0.98	Method		CRL (<>Robinson)
fb-hCG	27.3 ng/ml	0.95	Scan date		17/4/2023
Risks at sampling date			Nuchal translucency 1.3		
Age Risk		1:972	Nuchal translucency MoM		0.81
Biochemical T21 risk		1:6391	Nasal bone		Present
Combined trisomy 21 ris	sk	<1:10000			
Trisomy 13/18 + NT		<1:10000			
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
Risk 1:10 1:250 1:1000 1:1000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:100000 1:100000 1:10000 1:10000 1:10000 1:10000 1:1000	Trisomy 13/18 (with	Age	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 assessment! Calculated risks have no diagnostic values		
	Above Cut Off		Risk above Ag	e Risk	Risk below Age risk