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				Date of Report PRISCA	15/1/2023 5.1.0.17
Patient Data					
Name		MRS. SANA	Patient ID		012301140038
Birthday		13/04/1994	Sample ID		11523783
Age at term		29.02	Sample Date		14/1/2023
Gestational age		13+0			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	78 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom			13+0
PAPP-A	5.6 mIU/ml	1.19	Method		CRL (<>Robinson)
fb-hCG	18.9 ng/ml	0.62	Scan date		14/1/2023
Risks at sampling date			Nuchal translucency 1		
Age Risk 1:754		Nuchal translucency MoM 0.59			
Biochemical T21 risk	Γ21 risk <1:10000		Nasal bone Presen		
Combined trisomy 21 ri	sk	<1:10000			
Trisomy 13/18 + NT		<1:10000			
Risk			Down's Syndr	ome Risk (Trisomy 21	Screening)
Risk 1:10 1:100 1:250 Cut off 1:1000 1:1000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:1000 1:10000 1:10000 1:1000 1:10000 1:10000 1:10000 1:10000 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! The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values		