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				Date of Report PRISCA	2/2/2023 5.1.0.17
Patient Data					
Name		MRS. KAJAL	Patient ID		052302010030
Birthday		5/8/1997	Sample ID		11531918
Age at term		25.11	Sample Date		1/2/2023
Gestational age		13+1			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	44.6 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom			13+1
PAPP-A	4.63 mIU/ml	0.48	Method		CRL (<>Robinson)
fb-hCG	33.8 ng/ml	0.97	Scan date		1/2/2023
Risks at sampling date			Nuchal translucency 2.5		
Age Risk		1:959	Nuchal translucency MoM		1.45
Biochemical T21 risk		1:1097	Nasal bone		Present
Combined trisomy 21 risl	k	1:1005			
Trisomy 13/18 + NT		<1:10000			
Risk			Down's Syndro	ome Risk (Trisomy 21	Screening)
Risk 1:10 1:250 1:250 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:100 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 1005 women with the same data, there is one woman with a trisomy 21 pregnancy and 1004 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		
<1:10000 , which indicat		1 IN I) 15	Risk above Age Risk		