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					Date of Report PRISCA	19/8/2022 5.1.0.17
Patient Data						
Name		М	RS. MONIKA	. Patient ID		012208180037
Birthday			23/11/1998	Sample ID		11468850
Age at term			24.01	Sample Date		18/8/2022
Gestational age			13+1			
Correction factors						
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown
Weight in kg	52	Diabetes		NO	Pregnancies	unknown
Smoker	NO	Origin		Asian		
Biochemical Data				Ultrasound Data		
Parameter	Value		Corr Mom	Gestational ag	e	13+5
PAPP-A	4.66	mIU/ml	0.46	Method		CRL (<>Robinson)
fb-hCG	18.5	ng/ml	0.71	Scan date		17/8/2022
Risks at sampling date				Nuchal translucency 1.7		
Age Risk			1:1052	Nuchal translu	cency MoM	0.89
Biochemical T21 risk	emical T21 risk		1:1996	Nasal bone		Present
Combined trisomy 21 risk			<1:10000			
Trisomy 13/18 + NT			<1:10000			
Risk				Down's Syndrome Risk (Trisomy 21 Screening)		
Risk 1:10 1:250 1:250 1:1000 1:1000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:1000 1:1000 1:1000 1:250 1:100 1:250 1:100 1:250 1:100 1:250 1:100 1:250 1:100 1:250 1:100 1:250 1:100 1:250 1:100 1:250 1:100 1:250 1:100 1:250 1:100 1:250 1:100 1:250 1:1000 1:250 1:2				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				on the risk assessment! Calculated risks have no diagnostic values		
Risk A	bove Cı	ıt Off		Risk above Ag	e Risk	Risk below Age risk