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				Date of Report PRISCA	8/2/2023 5.1.0.17
Patient Data					
Name	MRS. KIRAN	MEET KAUR	Patient ID		012302060324
Birthday		29/12/1991	Sample ID		11505320
Age at term		31.1	Sample Date		6/2/2023
Gestational age		11+5			
Correction factors				_	
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	71 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom			13+4
PAPP-A	4.28 mIU/ml	0.63	Method		CRL (<>Robinson)
fb-hCG	33.5 ng/ml	1.36	Scan date		7/2/2023
Risks at sampling date			Nuchal translucency 1.25		
ge Risk 1:579		Nuchal translucency MoM 0.68			
Biochemical T21 risk	1:627		Nasal bone Prese		Present
Combined trisomy 21 ris	k	1:3665			
Trisomy 13/18 + NT		<1:10000			
Risk			Down's Syndr	ome Risk (Trisomy 21	Screening)
Risk 1:10 1:10 1:250 Cut off 1:10000 1:1000 1:1			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 3665 women with the same data, there is one woman with a trisomy 21 pregnancy and 3664 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		