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				Date of Report PRISCA	11/8/2023 5.2.0.13
Patient Data					
Name	MRS. MANPREET KAUR		Patient ID		012308090066
Birthday		1/1/1993	Sample ID		11059689
Age at term		31	Sample Date		9/8/2023
Gestational age		13+0			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom			12+0
PAPP-A	5.6 mIU/ml	0.79	Method		CRL (<>Robinson)
fb-hCG	66.4 ng/ml	1.77	Scan date		2/8/2023
Risks at sampling date			Crown rump length in mm 54.1		
Age Risk		1:608	Nuchal translucency MoM		0.7
Biochemical T21 risk		1:578	Nasal bone P		Present
Combined trisomy 21 risk		1:3312			
Trisomy 13/18 + NT		<1:10000			
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
Risk 1:10			The calculated risk for Trisomy 21 (with NT) is below the		
1:100 1:250 Cut off			<b>cut off, which represents a low risk.</b> After the result of the Trisomy 21 with NT test it is expected that among 3312 women with the same data, there is one woman with a trisomy 21 pregnancy and 3311 women with not affected pregnancies. The calculated risk by <b>PRISCA</b> depends on the accuracy of		
1:1000 1:10000 13 15 17 19 21 23 25 3 The calculated risk for <1:10000 , which indica	Trisomy 13/18 (with	1,	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		
Risl	k Above Cut Off		Risk above Ag	e Risk	Risk below Age risk