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				Date of Report PRISCA	31/8/2022 5.1.0.17
Patient Data				1100011	
Name	M	RS. KASHISH	Patient ID		012208300296
Birthday		16.5.1998	Sample ID		11585849
Age at term		24.09	Sample Date		30/8/2022
Gestational age		12+0			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	77 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational ag	e	12+0
PAPP-A	4.56 mIU/ml	1.4	Method		CRL (<>Robinson)
fb-hCG	71.6 ng/ml	1.81	Scan date		30/8/2022
Risks at sampling date			Nuchal translucency 1		
Age Risk	e Risk 1:971		Nuchal translucency MoM 0.71		
Biochemical T21 risk		1:2911	Nasal bone		Present
Combined trisomy 21 ris	k	<1:10000			
Trisomy 13/18 + NT		<1:10000			
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
Risk 1:10 1:100 1:250 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 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		
	Above Cut Off		Risk above Ag	e Risk	Risk below Age risk