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				Date of Report PRISCA	19/6/2023 5.2.0.13
Patient Data					01210110
Name	MI	RS.BHAWNA	Patient ID		012306180103
Birthday		15/6/1986	Sample ID		11668058
Age at term		30.6	Sample Date		18/6/2023
Gestational age		13+0			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	75 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom			12+5
PAPP-A	4.29 mIU/ml	0.87	Method		CRL (<>Robinson)
fb-hCG	22.8 ng/ml	0.75	Scan date		16/6/2023
Risks at sampling date			Crown rump length in mm 63.7		
Age Risk 1:174		Nuchal translucency MoM 0.8			
Biochemical T21 risk	emical T21 risk 1:1499		Nasal bone Preser		
Combined trisomy 21 r	risk	1:7436			
Trisomy 13/18 + NT		<1:10000			
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
1:10 1:100 1:250 Cut off			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 7436 women with the same data, there is one woman with a trisomy 21 pregnancy and 7435 women with not affected pregnancies.		
1:1000 1:10000 13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49 The calculated risk for Trisomy 13/18 (with NT) is <1:10000, which indicates a low risk			The calculated risk by <b>PRISCA</b> 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		
	k Above Cut Off		Risk above Ag	e Risk	Risk below Age risk