

*Free Home Sample Collection 9999 778 778

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				Date of Report PRISCA	4/6/2024 5.2.0.13
Patient Data					
Name					012406030196
Birthday		6/1/2000	Sample ID		11888312
Age at sample		24.4	Sample Date		3/6/2024
Gestational age		13+6			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	52.9 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	2	13+6
PAPP-A	6.80 mIU/ml	0.68	Method		CRL (<>Robinson)
fb-hCG	32.7 ng/ml	1.27	Scan date		3/6/2024
Risks at sampling date			Crown rump length in mm 78.1		
ge Risk 1:1028		Nuchal translucency MoM 0.57			
Biochemical T21 risk		1:1542	Nasal bone		Present
Combined trisomy 21 risk		1:8772	Sonographer		DR.PARVEEN
Trisomy 13/18 + NT		<1:10000	Qualifications	in measuring NT	C/R
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
1:10 1:250 1:250 Cut off 1:1000 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 test (with NT) it is expected that among 8772 women with the same data, there is one woman with a trisomy 21 pregnancy and 8771 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 patient combined risk presumes that NT measurement was done according to accepted guidelines (Prenat Diagn 18:511-523; 1998). The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values		
which indicates a low risk Risk Above Cut Off Risk				e Risk 📃 Ri	isk below Age risk