

*Free Home Sample Collection 9999 778 778



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 Date of Report
 27-10-2023

 PRISCA
 5.2.0.13

Patient Data						
Name	ame MRS JYOTI SHARMA			Patient ID		012310260010
Birthday			22-09-1994	Sample ID		11787544
Age at Sample date 29.1				Sample Date		26-10-2023
Gestational age		11+6				
Correction factors						
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown
Weight in kg	58	Diabetes		NO	Pregnancies	unknown
Smoker	NO	Origin		Asian		
Biochemical Data				Ultrasound Data		
Parameter	Value		Corr Mom	Gestational ago	2	11+5
PAPP-A	3.58	mIU/ml	0.83	Method		CRL (<>Robinson)
fb-hCG	48.5	ng/ml	1.08	Scan date		25-10-2023
Risks at sampling date				Crown rump length in mm 50.		
Age Risk			1:699	Nuchal translucency MoM		0.66
Biochemical T21 risk			1:2389	Nasal bone		PRESENT
Combined trisomy 21 risk			<1:10000	Sonographer		DR VIKAS GOYAL
Trisomy 13/18 + NT			<1:10000	Qualifications	in measuring NT	MBBS
Risk				Down's Syndrome Risk (Trisomy 21 Screening)		
1:100 1:250 Cut off 1:1000 1:1000 1:1000 1:1000 The calculated risk for Trisomy 13/18 (with NT) is <1:10000,				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 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 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 assessment!	Calculated risks have no	diagnostic values
Ris	k Above Cı	ıt Off		Risk above Ag	e Risk	Risk below Age risk