

\*Free Home Sample Collection 9999 778 778



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 Date of Report
 18-12-2023

 PRISCA
 5.2.0.13

					TRIBUTI	0.2.0.10
Patient Data						
Name			MRS POOJA	Patient ID		012312170079
Birthday			19-06-1996	Sample ID		11785622
Age at Sample date			27.5	Sample Date		17-12-2023
Gestational age 13+1						
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 age	2	13+0
PAPP-A	5.1	mIU/ml	1.01	Method		CRL (<>Robinson)
fb-hCG	39.7	ng/ml	1.36	Scan date		16-12-2023
Risks at sampling date				Crown rump length in mm 68.3		
Age Risk			1:847	Nuchal translucency MoM		0.93
Biochemical T21 risk			1:2637	Nasal bone PRESE		PRESENT
Combined trisomy 21 risk			<1:10000	Sonographer DR DEEP		DR DEEPIKA
Trisomy 13/18 + NT			<1:10000	Qualifications in measuring NT		MD
Risk				Down's Syndrome Risk (Trisomy 21 Screening)		
1:100 1:250 Cut off 1:1000 1:10000 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 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		
Ris	k Above Cı	ıt Off		Risk above Age	e Risk	Risk below Age risk
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