

*Free Home Sample Collection 9999 778 778

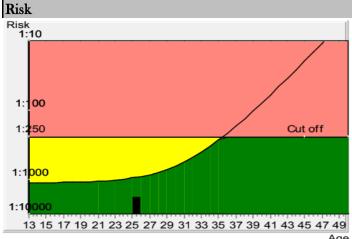


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Date of Report	09/02/2020
PRISCA	5.0.2.37

					IMSCA	3.0.2.07
Patient Data						
Name			Mrs Stuti	Patient ID		092002080005
Birthday			1/8/1994	Sample ID		10643871
Age at delivery			25.5	Sample Date		08/02/2020
Gestational age			12+4			
Correction factors						
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown
Weight in kg	85	Diabetes		no	Pregnancies	
Smoker	no	Origin		Asian		

Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	12+4	
PAPP-A	4.32 mIU/ml	1.45	Method	CRL (<>Robinson)	
fb-hCG	47.7 ng/ml	1.14	Scan Date	8/2/2020	
Risks at sampling date			Crown Rump Length (mm)	60	
Age Risk		1:940	Nuchal translucency MoM	0.77	
Biochemical Trisomy 2	21 Risk	1:9139	Nasal Bone	present	
Combined Trisomy 21	Risk	<1:10000	Sonographer	DR. KRITI RAJ	
Trisomy 13/18 + NT		<1:10000	Qualification in measuring NT	MBBS, MD	



Down's Syndrome Risk (Trisomy 21 Screening)

The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.

After the result of the Trisomy 21 test (with NT) it is expected that among more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy.

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!

Trisomy 13/18 + NT

The calculated risk for trisomy 13/18 (with nuchal translucency) is <1:10000, which represents a low risk.

The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values