

\*Free Home Sample Collection 9999 778 778

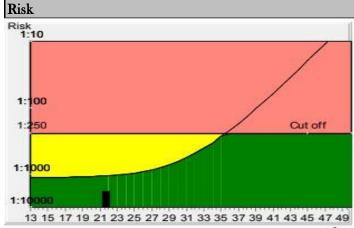


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Date of Report 7/10/2019 PRISCA 50937

					FMSCA	3.0.2.37	
Patient Data							
Name		Mrs Ranjana		Patient ID		011910030241	
Birthday		7/2/1998		Sample ID		10572689	
Age at delivery		21.7		Sample Date		05/10/19	
Gestational age		12+5					
Correction factors							
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown	
Weight in kg		Diabetes		no	Pregnancies		
Smoker	no	Origin		Asian			

Biochemical Data			Ultrasound Data		
Parameter	rameter Value		Gestational age	12+2	
PAPP-A	4.88  mIU/ml	0.92	Method	CRL (<>Robinson)	
fb-hCG	o-hCG 131.2 ng/ml		Scan Date	3/10/2019	
Risks at sampling date			Crown Rump Length (mm)	57.2	
Age Risk		1:1069	Nuchal translucency MoM	0.47	
Biochemical Trisomy 21 Risk		1:597	Nasal Bone	present	
Combined Trisomy 21 Risk		1:3445	Sonographer	DR. POONAM YADAV	
Trisomy 13/18 + NT		<1:10000	Qualification in measuring NT	C/R	



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

Down's Syndrome Risk (Trisomy 21 Screening)

After the result of the Trisomy 21 test (with NT) it is expected that among 3445 women with the same data, there is one woman with a trisomy 21 pregnancy and

3444 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!

## 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

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