

*Free Home Sample Collection 9999 778 778

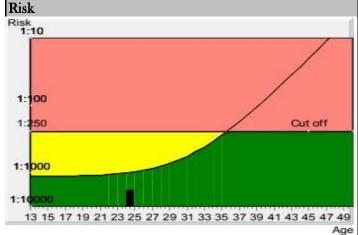


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Date of Report 8/5/2019

				5.0.2.37			
Patient Data							
Name		Mrs Neha Yadav		Patient ID		011905060239	
Birthday			20/12/1994	Sample ID		10412010	
Age at delivery			24.4	Sample Date		06/05/19	
Gestational age			13+0				
Correction factors							
Fetuses	1 I	VF		unknown	Previous trisomy 21	unknown	
Weight in kg	56 I	Diabetes		no	Pregnancies	unknown	
Smoker	no (Origin		Asian			
Biochemical Data			Ultrasound Data				

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Parameter	eter Value		Gestational age	12+6	
PAPP-A	3.61 mIU/ml	0.62	Method	CRL (<>Robinson)	
fb-hCG	48.68 ng/ml	1.08	Scan Date	6/5/2019	
Risks at sampling date			Crown Rump Length (mm)	65	
Age Risk		1:1002	Nuchal translucency MoM	0.54	
Biochemical Trisomy 21 Risk		1:1774	Nasal Bone	present	
Combined Trisomy 21 Risk		<1:10000	Sonographer	DR.RAHUL	
Trisomy 13/18 + NT		<1:10000	Qualification in measuring NT	DNB	



Down's Syndrome Risk (Trisomy 21 Screening)

TThe 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 approaches 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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