

\*Free Home Sample Collection 9999 778 778

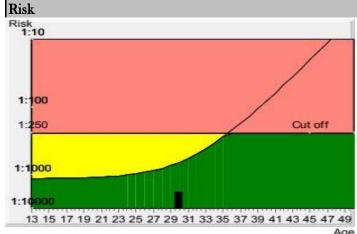


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Date of Report 19/11/19 PRISCA 50937

				3.0.2.37		
Patient Data						
Name	Mrs Anju Kumari		Patient ID		011911180096	
Birthday		22/12/1989	Sample ID		10303085	
Age at delivery		29.9	Sample Date		18/11/19	
Gestational age		13+2	2			
Correction factors						
Fetuses	1 Γ	VF	unknown	Previous trisomy 21	unknown	
Weight in kg	65 L	Diabetes	no	Pregnancies		
Smoker	no C	Origin	Asian			

Biochemical Data			Ultrasound Data		
Parameter	neter Value		Gestational age	13+0	
PAPP-A	2.31  mIU/ml	0.43	Method	CRL (<>Robinson)	
fb-hCG	18.4 ng/ml	0.44	Scan Date	16/11/19	
Risks at sampling date			Crown Rump Length (mm)	65.9	
Age Risk		1:671	Nuchal translucency MoM	0.71	
Biochemical Trisomy 21 Risk		1:2827	Nasal Bone	present	
Combined Trisomy 21 Risk		<1:10000	Sonographer	DR. ELLORA	
Trisomy 13/18 + NT		<1:10000	Qualification in measuring NT	C/R	



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

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