

\*Free Home Sample Collection 9999 778 778

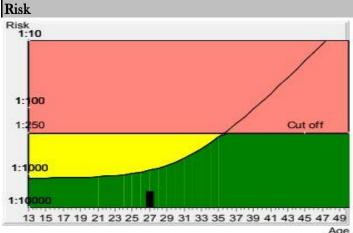


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Date of Report 10/9/2019 PRISCA 5 0 9 37

			PRISCA		5.0.2.37		
Patient Data							
Name		Mrs Suman		Patient ID		011909080249	
Birthday		21/09/1992				10586091	
Age at delivery		27		Sample Date		08/09/19	
Gestational age		13+3					
Correction factors							
Fetuses	1 I	IVF		unknown	Previous trisomy 21	unknown	
Weight in kg	45 I	Diabetes		no	Pregnancies		
Smoker	no (	Origin		Asian			

Biochemical Data			Ultrasound Data		
Parameter Value		Corr Mom	Gestational age	13+0	
PAPP-A	8.61  mIU/ml	1.01	Method	CRL (<>Robinson)	
fb-hCG	50.05 ng/ml	1.07	Scan Date	6/9/2019	
Risks at sampling date			Crown Rump Length (mm)	73.6	
Age Risk		1:888	Nuchal translucency MoM	0.99	
Biochemical Trisomy 21 Risk		1:4772	Nasal Bone	present	
Combined Trisomy 21 Risk		<1:10000	Sonographer	DR. RUPANG	
Trisomy 13/18 + NT		<1:10000	Qualification in measuring NT	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 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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