

*Free Home Sample Collection 9999 778 778

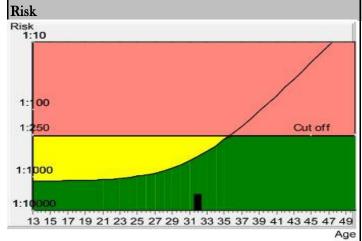


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Date of Report 10/12/2019
PRISCA 5 0 9 37

				PRISCA	5.0.2.37	
Patient Data						
Name		Mrs Akansha Sharma				061912080027
Birthday			10/1/1988	Sample ID		10560176
Age at delivery			31.9	Sample Date		08/12/19
Gestational age			13+4			
Correction factors						
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown
Weight in kg	73.2	Diabetes		no	Pregnancies	
Smoker	no	Origin		Asian		
Biochemical Data			Ultrasound Da	ata		

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Parameter	meter Value		Gestational age	12+5	
PAPP-A	2.76 mIU/ml		Method	CRL (<>Robinson)	
fb-hCG	p-hCG 17.4 ng/ml		Scan Date	2/12/2019	
Risks at sampling date			Crown Rump Length (mm)	63.8	
Age Risk		1:510	Nuchal translucency MoM	0.67	
Biochemical Trisomy 21 Risk		1:3938	Nasal Bone	present	
Combined Trisomy 21 Risk		<1:10000	Sonographer	DR SAVITA CHOPRA	
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 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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