

\*Free Home Sample Collection 9999 778 778

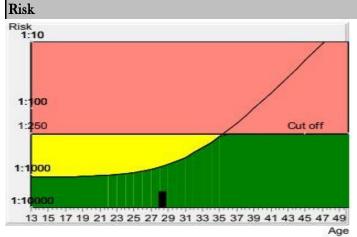


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Date of Report 7/10/2019 PRISCA 5.0.2.37

					TMSCA	5.0.2.57	
Patient Data							
Name		Mrs Pooja Yadav		Patient ID		011910050218	
Birthday			16/06/1991	Sample ID		10582667	
Age at delivery		28.3		Sample Date		05/10/19	
Gestational age			12+0				
Correction factors							
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown	
Weight in kg	45	Diabetes		no	Pregnancies		
Smoker	no	Origin		Asian			

Biochemical Data			Ultrasound Data		
Parameter	rameter Value		Gestational age	11+5	
PAPP-A	3.45  mIU/ml	0.67	Method	CRL (<>Robinson)	
fb-hCG	112.5 ng/ml	2.05	Scan Date	4/10/2019	
Risks at sampling date			Crown Rump Length (mm)	50.8	
Age Risk		1:759	Nuchal translucency MoM	0.88	
Biochemical Trisomy 21 Risk		1:349	Nasal Bone	present	
Combined Trisomy 21 Risk		1:1938	Sonographer	DR NITIN YADAV	
Trisomy 13/18 + NT		<1:10000	Qualification in measuring NT	MBBS, 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 1938 women with the same data, there is one woman with a trisomy 21 pregnancy and 1937 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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