

*Free Home Sample Collection 9999 778 778

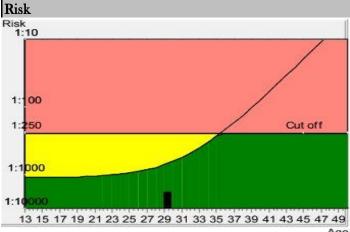


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Date of Report 1/1/2020 PRISCA 5.0.2.37

					TMSCA	3.0.2.07	
Patient Data							
Name		Mrs Kamla		Patient ID		051912310029	
Birthday		16/08/1990		Sample ID		10501472	
Age at delivery		29.4		Sample Date		31/12/19	
Gestational age			12+2				
Correction factors							
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown	
Weight in kg	74	Diabetes		no	Pregnancies		
Smoker	no	Origin		Asian			

Biochemical Data			Ultrasound Data		
Parameter	rameter Value		Gestational age	12+2	
PAPP-A	$1.69~\mathrm{mIU/ml}$		Method	CRL (<>Robinson)	
fb-hCG	b-hCG 19.85 ng/ml		Scan Date	31/12/19	
Risks at sampling date	e		Crown Rump Length (mm)	55.5	
Age Risk		1:689	Nuchal translucency MoM	0.82	
Biochemical Trisomy 21 Risk		1:5040	Nasal Bone	present	
Combined Trisomy 21 Risk		<1:10000	Sonographer	DR. KAVITA	
Trisomy 13/18 + NT		<1:10000	Qualification in measuring NT	MD	



After the result of the Trisomy 21 test (with NT) it is expected that among more than 10000 women with the

below the cut off, which indicates a low risk.

same data, there is one woman with a trisomy 21 pregnancy. The calculated risk by PRISCA depends on the accuracy of

Down's Syndrome Risk (Trisomy 21 Screening)

the information provided by the referring physician. Please note that the risk calculations are statistical aapproaches and have no diagnostic value!

The calculated risk for Trisomy 21 (with nuchal translucency) is

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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