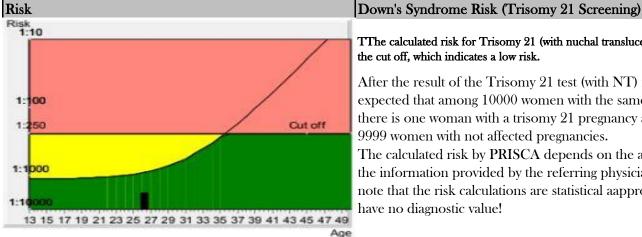
Date of Report 28/07/19 PRISCA 5.0.2.37

					TMSCA	3.0.2.37	
Patient Data							
Name		Mrs Reena		Patient ID		011907270131	
Birthday		15/05/1993		Sample ID		10620846	
Age at delivery		26.2		Sample Date		27/09/19	
Gestational age		12+6					
Correction factors							
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown	
Weight in kg	61.6	Diabetes		no	Pregnancies		
Smoker	no	Origin		Asian			

Biochemical Data			Ultrasound Data		
Parameter	ameter Value		Gestational age	12+5	
PAPP-A	$4.3~\mathrm{mIU/ml}$	0.88	Method	CRL (<>Robinson)	
fb-hCG	54.9 ng/ml	1.23	Scan Date	26/07/19	
Risks at sampling date			Crown Rump Length (mm)	64.2	
Age Risk		1:915	Nuchal translucency MoM	0.85	
Biochemical Trisomy 21	Risk	1:2635	Nasal Bone	present	
Combined Trisomy 21 F	Risk	<1:10000	Sonographer	DR.VIKAS GOYAL	
Trisomy 13/18 + NT		<1:10000	Qualification in measuring NT	DMRD	



TThe 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 10000 women with the same data, there is one woman with a trisomy 21 pregnancy and 9999 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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