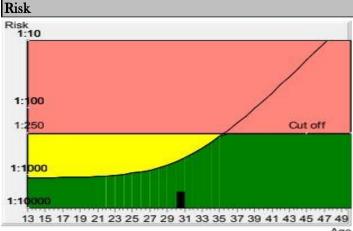
Date of Report
 24/06/19

 PRISCA
 5.0,2.37

					TRISCIT	0.0.2.07	
Patient Data							
Name	Mrs Kamaljit Kaur		Patient ID		011906220343		
Birthday		11/12/1998		Sample ID		10609295	
Age at delivery		30.5		Sample Date		22/06/19	
Gestational age			12+4				
Correction factors							
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown	
Weight in kg	57	Diabetes		no	Pregnancies		
Smoker	no	Origin		Asian			

Biochemical Data			Ultrasound Data		
Parameter	nrameter Value		Gestational age	12+3	
PAPP-A	PAPP-A 2.65 mIU/ml		Method	CRL (<>Robinson)	
fb-hCG	66.99 ng/ml	1.42	Scan Date	22/06/19	
Risks at sampling dat	ze		Crown Rump Length (mm)	59.5	
Age Risk		1:604	Nuchal translucency MoM	0.9	
Biochemical Trisomy	21 Risk	1:400	Nasal Bone	present	
Combined Trisomy 2	21 Risk	1:2169	Sonographer	DR.RUBY RAHUL	
Trisomy 13/18 + NT		<1:10000	Qualification in measuring NT	CON. RADIOLOGIST	
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		



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 2169 women with the same data, there is one woman with a trisomy 21 pregnancy and 2168 women with not affected pregnancies.

The calculated risk by PRISCA depends on the accuracy of

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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