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Date of Report 23-01-19 DDISCA 50237

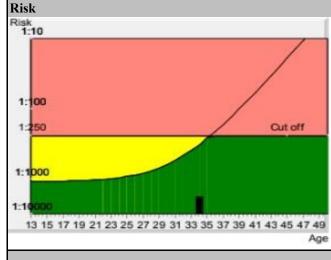
11 + 5

					PRISCA	5.0.2.37
Patient Data						
Name			Mrs Sudha	Patient ID		011901210358
Birthday			25-01-85	Sample ID		10364637
Age at delivery			34.0	Sample Date		23/01/2019
Gestational age			11+6			
Correction factors						
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown
Weight in kg	79.8	Diabetes		no	Pregnancies	
Smoker	no	Origin		Asian		
Biochemical Data				Ultrasound D	ata	

1 al allietei	v aruc	COII MOIII	Gestational age	11.5
PAPP-A	1.76 mIU/ml	0.84	Method	CRL (⇔Hadlock)
fb-hCG	35.63 ng/ml	0.80	Scan Date	21-01-19
Risks at sampling date			CRL	49.5
Age Risk		1:327	Nuchal translucency MoM	1.12
Biochemical T21 Risk		1:2260	Nasal Bone	Present
Combined Trisomy 21 Ris	k	1:6635	Sonographer	DR.RUBY RAHUL
Trisomy 13/18 + NT		<1:10000	Qualifications in measuring NT	CON. RADIOLOGIST

Gestational age

Corr Mom



Value

Trisomy 13/18 + NT

Parameter

The calculated risk for trisomy 13/18 (with nuchal translucency) is <1:10000, which represents a low risk.

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 6635 women with the same data, there is one woman with a trisomy 21 pregnancy and 6634 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 risk calculations are statistical approach and have no diagnostic values!

The patient combined risk presumes the NT measurement was done according to accepted guidelines The laboratory can not be held responsible for their impact on the risk assessment! Calculated value has no diagnostic value!