

Biochemical Data

*Free Home Sample Collection 9999 778 778



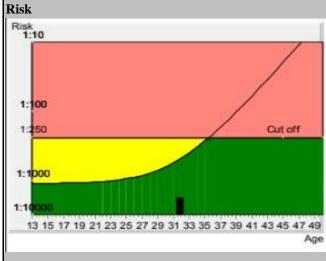
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Date of Report 21-12-18 PRISCA 5.0.2.37

					TRIBETT	5.0.2.51
Patient Data						
Name			Mrs Mona	Patient ID		011812180212
Birthday			12-03-87	Sample ID		10390484
Age at delivery			31.8	Sample Date		18/12/2018
Gestational age			12+4			
Correction factors						
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown
Weight in kg	59.8	Diabetes		no	Pregnancies	
Smoker	no	Origin		Asian		

Parameter	Value	Corr Mom	Gestational age	12+2
PAPP-A	4.15 mIU/ml	1.06	Method	CRL (<>Hadlock)
fb-hCG	28.56 ng/ml	0.64	Scan Date	17-12-18
Risks at sampling date			CRL	56.6
Age Risk		1:503	Nuchal translucency MoM	0.67
Biochemical T21 Risk		1:9269	Nasal Bone	Present
Combined Trisomy 21 Risk	ζ.	<1:10000	Sonographer	DR. A. JUNEJA
Trisomy 13/18 + NT		<1:10000	Qualifications in measuring NT	MD

Ultrasound Data



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

Trisomy 13/18 + NT

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 more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy.

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!