

The calculated risk for Trisomy 13/18 (with NT) is

<1:10000, which indicates a low risk

*Free Home Sample Collection 9999 778 778



on the risk assessment! Calculated risks have no diagnostic

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Date of Report 22/07/19
PRISCA 5.0.2.37

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Patient Data					
Name		Mrs Sapna	Patient ID		011907210091
Birthday		3/1/2001	Sample ID		10452175
Age at delivery		19	Sample Date		22/07/19
Gestational age		15+1			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	37 Diabetes		no	Pregnancies	unknown
Smoker	no Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	2	12+6
AFP	43.1 ng/ml	0.94	Method		CRL (<>Robinson)
uE3	$0.84~\mathrm{mIU/ml}$	1.57	Scan Date		
hCG	30334.5 ng/ml	0.57	Crown rump length (mm)		
Risks at sampling date			Nuchal translucency 1.4		
Age Risk	ge Risk 1:1550		Nuchal translucency MoM		0.84
Biochemical T21 Risk		<1:10000	Nasal Bone		Present
Combined Trisomy 21 Risk		<1:10000	Sonographer		DR.ALOK VARSHNEY
Trisomy 13/18 + NT		<1:10000	Qualifications in measuring NT		M.D,.D.N.B.
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
1:10			The calculated risk for Trisomy 21 (with NT) is below the cut off, which represents a low risk.		
1:100 1:250 Cut off 1:1000 1:1000 13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49 Age			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 the risk calculations are statistical aapproaches and have no diagnostic value!		
Trisomy 13/18 + NT			The laboratory cannot be hold responsible for their impact		

values