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Date of Report 14-11-18 PRISCA 5.0.2.37

				PRISCA	5.0.2.37	
Patient Data						
Name		Mrs Heena	Patient ID		051811130001	
Birthday		05-12-93	Sample ID		10384031	
Age at delivery		24.9	Sample Date		13/11/18	
Gestational age		12+1				
Correction factors						
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown	
Weight in kg	40.6 Diabetes		unknown	Pregnancies	unknown	
Smoker	Unknown Origin		Asian			
Biochemical Data			Ultrasound Data			
Parameter	Value	Corr Mom	Gestational ag	e	12+0	
PAPP-A	4.25 mIU/ml	0.82	Method		CRL (<>Robinson)	
fb-hCG	62.41 ng/ml	1.15	Scan Date		12-11-18	
Risks at sampling date			Crown rump lo	ength (mm)	52.6	
Age Risk		1:951	Nuchal translu	icency MoM	0.7	
Biochemical T21 Risk		1:2747	Nasal Bone		Present	
Combined Trisomy 21 I	Risk	<1:10000	Sonographer		DR.VENKAT REDDY M	
Trisomy 13/18 + NT		<1:10000	Qualifications	in measuring NT	MD	
Risk			Down's Syndrome Risk (Trisomy 21 Screening)			
Risk 1:10			The calculated risk for Trisomy 21 (with NT) is below the cut off, which represents a low risk.			
1:100 1:250 1:1000 1:10000 13 15 17 19 21 23 25 3	27 29 31 33 35 37 39	Cut off 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 calculated risk for Trisomy 13/18 (with NT) is			•	The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic		

<1:10000, which indicates a low risk

values

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