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 Date of Report
 26/5/2022

 PRISCA
 5.1.0.17

Patient Data					
Name MRS. SHAILA YADAV		Patient ID		012205250083	
Birthday		1/1/1981	Sample ID		11446103
Age at term 41.1		Sample Date		25/5/2022	
Gestational age		12+2			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	65 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	2	12+1
PAPP-A	$4.38~\mathrm{mIU/ml}$	1.17	Method		CRL (<>Robinson)
fb-hCG	86.7 ng/ml	1.86	Scan date		24/05/2022
Risks at sampling date			Nuchal Translucency 1.34		
Age Risk	1:58		Nuchal Translucency MoM 0.9		0.91
Biochemical T21 risk		1:106	Nasal Bone Preser		Present
Combined T21 risk 1		1:495			
Trisomy 13/18+NT		<1:10000			
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
1:100 1:250 1:1000 1:10000 1:10000 1:3 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49 Age  Trisomy 13/18+NT The calculated risk for Trisomy 13/18 (with NT) is <1:10000 , which indicates a low risk			The calculated risk for Trisomy 21 (with NT) is below the cut off, which represents a low risk.  After the result of the Trisomy 21 test (with NT) it is expected that among 495 women with the same data, there is one woman with a trisomy 21 pregnancy and 494 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 the risk calculations are statistical aapproaches and have no diagnostic value!  The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values		