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 Date of Report
 12/5/2022

 PRISCA
 5.1.0.17

Patient Data					
Name MRS. KANCHAN		Patient ID		012205110108	
Birthday		28/12/1993	Sample ID		11470655
Age at term		28.1	Sample Date		11/5/2022
Gestational age		11+5			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	54.4 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	e	11+4
PAPP-A	$1.94~\mathrm{mIU/ml}$	0.53	Method		CRL (<>Robinson)
fb-hCG	39.5 ng/ml	0.75	Scan date		10/5/2022
Risks at sampling date			Nuchal Translucency 0.8		
Age Risk		1:747	Nuchal Translucency MoM		0.62
Biochemical T21 risk		1:1917	Nasal Bone Presen		
Combined T21 risk		<1:10000			
Trisomy 13/18+NT		<1:10000			
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
1:100 1:250 Cut off 1:1000 1:1000 1:10000 13 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 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!  The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values		
Ris	k Above Cut Off		Risk above Ag	e Risk	Risk below Age risk