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 Date of Report
 24/5/2022

 PRISCA
 5.1.0.17

Patient Data					
Jame MRS. HEMLATA		Patient ID		052205210038	
Birthday		15/09/1990	Sample ID		11464465
Age at term		32.01	Sample Date		21/5/2022
Gestational age		12+6			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	62.8 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	2	12+6
PAPP-A	$2.92~\mathrm{mIU/ml}$	0.61	Method		CRL (⇔Robinson)
fb-hCG	18.5 ng/ml	0.42	Scan date		21/05/2022
Risks at sampling date			Nuchal Translucency 1.2		
Age Risk		1:516	Nuchal Translucency MoM 0.73		
Biochemical T21 risk		1:6001	Nasal Bone Present		
Combined T21 risk		<1:10000			
Trisomy 13/18+NT		<1:10000			
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
Risk 1:10 1:250 1:1000 1:10000 1:1			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 and 9999 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		