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 Date of Report
 28/9/2022

 PRISCA
 5.1.0.17

Patient Data					
Name	MRS. PI	J SHPANJALI	Patient ID		012209270066
Birthday		31.12.1991	Sample ID		11621288
Age at term		31.01	Sample Date		27/9/2022
Gestational age		12+4			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	58.2 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	2	12+4
PAPP-A	5.34 mIU/ml	0.93	Method		CRL (<>Robinson)
fb-hCG	57.2 ng/ml	1.53	Scan date		27/9/2022
Risks at sampling date			Nuchal translucency 1.1		
Age Risk		1:587	Nuchal translucency MoM		0.7
Biochemical T21 risk		1:1155	Nasal bone Presen		
Combined trisomy 21 risk		1:2568			
Trisomy 13/18 + NT		<1:10000			
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
1:1000 1:250 Cut off 1:1000 1:10000 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 21with NT test it is expected that among 2568 women with the same data, there is one woman with a trisomy 21 pregnancy and 2567 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		