

 Date of Report
 16/5/2023

 PRISCA
 5.2.0.13

Patient Data					
Name MRS. JYOTI		Patient ID		012305150148	
Birthday		13/08/1999	Sample ID		11668432
Age at term		24.2	Sample Date 15/5/20		15/5/2023
Gestational age		12+2			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	47 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom			12+2
PAPP-A	$5.2~\mathrm{mIU/ml}$	0.79	Method		CRL (<>Robinson)
fb-hCG	46.3 ng/ml	1.07	Scan date		15/5/2023
Risks at sampling date			Nuchal translucency 1.15		
Age Risk		1:999	Nuchal translucency MoM 0.7		0.76
Biochemical T21 risk		1:3191	Nasal bone Presen		
Combined trisomy 21 risk		<1:10000			
Trisomy 13/18 + NT		<1:10000			
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
1:100 1:1000 1:1000 1:100000 1:100000 1:100000 1:100000 1:100000 1:100000 1:100000 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 21with NT test 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		