

 Date of Report
 4/7/2022

 PRISCA
 5.1.0.17

				TRISCIT	
Patient Data					
Name MRS. KAJAL SHARMA			Patient ID		012207010061
Birthday		11/9/1999	Sample ID		11446062
Age at term		23.02	Sample Date		1/7/2022
Gestational age 13+1					
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	66 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	2	13+1
PAPP-A	6.2 mIU/ml	1.02	Method		CRL (<>Robinson)
fb-hCG	86.1 ng/ml	2.83	Scan date		1/7/2022
Risks at sampling date			Nuchal translucency 1.16		
Age Risk 1:1057		1:1057	Nuchal translucency MoM 0.76		
Biochemical T21 risk		1:531	Nasal bone Prese		
Combined trisomy 21 risk	ζ	- 1:3033			
Trisomy 13/18 + NT		- <1:10000			
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
Risk 1:10 1:250 Cut off 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 21with NT test it is expected that among 3033 women with the same data, there is one woman with a trisomy 21 pregnancy and 3032 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		