

 Date of Report
 20/2/2023

 PRISCA
 5.1.0.17

Patient Data					
Name MRS. ANJANA			Patient ID		012302190089
Birthday 10/1/1991		Sample ID		11581048	
Age at term 32.7		Sample Date 19/2/202		19/2/2023	
Gestational age		12+4			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	68.7 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom			12+4
PAPP-A	$3.94~\mathrm{mIU/ml}$	0.84	Method		CRL (<>Robinson)
fb-hCG	27.3 ng/ml	0.77	Scan date		19/2/2023
Risks at sampling date			Nuchal translucency 1		
Age Risk		1:476	Nuchal translucency MoM 0.6		
Biochemical T21 risk		1:3532	Nasal bone Preser		Present
Combined trisomy 21 risk		<1:10000			
Trisomy 13/18 + NT		<1:10000			
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
1:100 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 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		