

 Date of Report
 14/2/2023

 PRISCA
 5.1.0.17

Patient Data					
Name MRS. TAMANNA			Patient ID		012302130174
Birthday	thday 20/11/1999		Sample ID		11508845
Age at term 23.8		Sample Date		13/2/2023	
Gestational age		12+4			
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			12+4
PAPP-A	4.6  mIU/ml	0.93	Method		CRL (<>Robinson)
fb-hCG	35.8  ng/ml	1	Scan date		13/2/2023
Risks at sampling date			Nuchal translucency 2.1		
Age Risk		1:1025	Nuchal translucency MoM 1.31		
Biochemical T21 risk		1:5478	Nasal bone Presen		Present
Combined trisomy 21 risk		1:8352			
Trisomy 13/18 + NT		<1:10000			
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
1:100 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 8352 women with the same data, there is one woman with a trisomy 21 pregnancy and 8351 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		