

 Date of Report
 24/8/2022

 PRISCA
 5.1.0.17

Patient Data					
Name MRS. AASTHA		Patient ID 012208220		012208220287	
Birthday	2/8/1992		Sample ID		11578513
Age at term 30.07		Sample Date 22/8/202		22/8/2022	
Gestational age		12+4			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	52.9 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	2	12+3
PAPP-A	5.3 mIU/ml	0.81	Method		CRL (<>Robinson)
fb-hCG	95.2 ng/ml	2.45	Scan date		22/8/2022
Risks at sampling date			Nuchal translucency 1.3		
Age Risk 1:642		1:642	Nuchal translucency MoM 0.86		
Biochemical T21 risk		1:285	Nasal bone		Present
Combined trisomy 21 risk 1:1596		1:1596			
		<1:10000			
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
1:100 1:250 1:1000 1:10000 1:110000 1:15 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 1596 women with the same data, there is one woman with a trisomy 21 pregnancy and 1595 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		