

 Date of Report
 27/8/2022

 PRISCA
 5.1.0.17

Patient Data					
Name MRS. MAYA			Patient ID		012208260173
Birthday		1/1/1996	Sample ID		11545529
Age at term		27.01	Sample Date		26/8/2022
Gestational age 12+3					
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	55 Diabetes		NO	Pregnancies	unknown
Smoker N	O Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter Val	ıe	Corr Mom	Gestational age	2	12+3
PAPP-A 3.	87 mIU/ml	0.67	Method		CRL (<>Robinson)
fb-hCG 72	.5 ng/ml	1.83	Scan date		26/8/2022
Risks at sampling date			Nuchal translucency 1.2		
Age Risk		1:877	Nuchal translucency MoM 0.7		0.78
Biochemical T21 risk		1:516	Nasal bone Preser		Present
Combined trisomy 21 risk		1:3108			
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 3108 women with the same data, there is one woman with a trisomy 21 pregnancy and 3107 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		