

 Date of Report
 23/2/2023

 PRISCA
 5.1.0.17

Patient Data					
Name MRS. AYESHA			Patient ID		012302220096
Birthday 10/2/1992		Sample ID		11514481	
Age at term 31.6		Sample Date 22/2/202		22/2/2023	
Gestational age		13+0			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	67 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom			12+6
PAPP-A	$3.26~\mathrm{mIU/ml}$	0.57	Method		CRL (<>Robinson)
fb-hCG	27.5 ng/ml	0.87	Scan date		21/2/2023
Risks at sampling date			Nuchal translucency 1.7		
Age Risk		1:572	Nuchal translucency MoM 1.0		
Biochemical T21 risk		1:1305	Nasal bone Preser		Present
Combined trisomy 21 risk		1:4966			
Trisomy 13/18 + NT		<1:10000			
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
1:100 1:1000 1:1000 1:100000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:1000			The calculated risk for Trisomy 21 (with NT) is below the cut off, which represents a low risk. After the result of the Trisomy 21 with NT test it is expected that among 4966 women with the same data, there is one woman with a trisomy 21 pregnancy and 4965 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		