

 Date of Report
 5/2/2023

 PRISCA
 5.1.0.17

Patient Data					
Name MRS. SHUBHANGI			Patient ID		012302040119
Birthday		9/1/1993	Sample ID		11481527
Age at term		31.5	Sample Date		4/2/2023
Gestational age		11+3			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 2	1 unknown
Weight in kg	38 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom			11+3
PAPP-A	$3.5~\mathrm{mIU/ml}$	0.6	Method		CRL (<>Robinson)
fb-hCG	51.7 ng/ml	0.89	Scan date		4/2/2023
Risks at sampling date			Nuchal translucency 0.8		
Age Risk		1:613	Nuchal translucency MoM		0.61
Biochemical T21 risk		1:1519	Nasal bone Presen		Present
Combined trisomy 21 risk		1:8478			
Trisomy 13/18 + NT		<1:10000			
Risk			Down's Syndro	ome Risk (Trisomy	21 Screening)
1:100 1:250 1:1000 1:1000 1:10000 1:110000 1:100000 1:100000 1:100000 1:100000 1:100000 1:100000 1:100000 1:100000000			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 8478 women with the same data, there is one woman with a trisomy 21 pregnancy and 8477 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		
Risk A	Above Cut Off		Risk above Age	e Risk	Risk below Age risk