

 Date of Report
 10/4/2023

 PRISCA
 5.2.0.13

Patient Data					
ame MRS. POONAM KUMARI		Patient ID 012		012304090130	
Birthday	16/09/1998		Sample ID		11476625
Age at term 25		Sample Date 9/4/20		9/4/2023	
Gestational age		13+1			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	39 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom			12+6
PAPP-A	4.89 mIU/ml	0.44	Method		CRL (<>Robinson)
fb-hCG	62.2 ng/ml	1.69	Scan date		7/4/2023
Risks at sampling date			Nuchal translucency 1.2		
Age Risk		1:1000	Nuchal translu	cency MoM	0.72
Biochemical T21 risk		1:245	Nasal bone		Present
Combined trisomy 21 risk	ζ.	1:1630			
Trisomy 13/18 + NT		<1:10000			
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
1:100 1:250 Cut off 1:1000 1:10000 1:1			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 1630 women with the same data, there is one woman with a trisomy 21 pregnancy and 1629 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		