

 Date of Report
 15/4/2023

 PRISCA
 5.2.0.13

Patient Data					
Name MRS. PRAMILA SUBODH			Patient ID		012304140104
Birthday 23/05/1995		Sample ID		11476694	
Age at term		28.3	Sample Date		14/4/2023
Gestational age		13+6			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	50 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom			13+6
PAPP-A	$4.69~\mathrm{mIU/ml}$	0.44	Method		CRL (<>Robinson)
fb-hCG	15.6 ng/ml	0.59	Scan date		14/4/2023
Risks at sampling date			Nuchal translucency 1.2		
Age Risk		1:839	Nuchal translucency MoM 0.6		
Biochemical T21 risk		1:2080	Nasal bone Pres		Present
Combined trisomy 21 risk <1:10000		<1:10000			
Trisomy 13/18 + NT <1:10000		<1:10000			
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
1:100 1:250 1:1000 1:1000 1:10000 1:3 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49 Age 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 more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy and 9999 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		