

 Date of Report
 21/6/2023

 PRISCA
 5.1.0.17

Patient Data					
Name MRS. POONAM		Patient ID		012306190099	
Birthday		13/05/1996	Sample ID		11657089
Age at term		27.7	Sample Date		19/6/2023
Gestational age		13+0			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	2	12+3
PAPP-A	$4.28~\mathrm{mIU/ml}$	0.6	Method		CRL (<>Robinson)
fb-hCG	61.3 ng/ml	1.64	Scan date		15/06/2023
Risks at sampling date			Nuchal translucency (NT) 1.2		
Age Risk		1:868	Nuchal translucency MoM 0.7		0.78
Biochemical T21 risk		1:526	Nasal bone preser		present
Combined trisomy 21 risk		1:3199			
Trisomy 13/18		<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  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 21 test it is expected that among 3199 women with the same data, there is one woman with a trisomy 21 pregnancy and 3198 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		