

 Date of Report
 6/1/2023

 PRISCA
 5.1.0.17

Patient Data					
Name		MRS. NIDHI	Patient ID		012301050157
Birthday		4/12/2001	Sample ID		11523772
Age at term		21.07	Sample Date		5/1/2023
Gestational age		13+3	3		
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	92 Diabete	S	NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom			13+2
PAPP-A	3.82 mIU/ml	0.86	Method		CRL (<>Robinson)
fb-hCG	16.9 ng/ml	0.67	Scan date		5/1/2023
Risks at sampling date			Nuchal translucency 1.2		
Age Risk 1:3		1:1104	Nuchal translucency MoM 0.6		
iochemical T21 risk <1:1		<1:10000	Nasal bone Prese		Present
Combined trisomy 21 risk		<1:10000			
Trisomy 13/18 + NT		<1:10000			
Risk			Down's Syndro	ome Risk (Trisomy 21	Screening)
1:100 1:250 1:1000 1:1000 1:100000 1:10000 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 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		
	Above Cut Off		Risk above Ag	e Risk	Risk below Age risk