

 Date of Report
 18/6/2022

 PRISCA
 5.1.0.17

Patient Data					
Name MRS. SAVITA THAKUR		Patient ID		012206160232	
Birthday		14/01/1990	Sample ID		11621790
Age at term		32.1	Sample Date		16/6/2022
Gestational age		12+2			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	56.4 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	2	12+1
PAPP-A	$3.64~\mathrm{mIU/ml}$	0.68	Method		CRL(<>Robinson)
fb-hCG	46.8 ng/ml	1.15	Scan date		16/6/2022
Risks at sampling date			Nuchal Translucency 1		
Age Risk		1:447	Nuchal Translucency MoM 0.6		0.67
Biochemical T21 risk 1:847		1:847	Nasal Bone Presen		
Combined T21 risk 1:4744		1:4744			
Trisomy 13/18+NT		<1:10000			
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
1:100  1:250  Cut off  1:1000  1:10000  13 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 (with NT) it is expected that among 4744 women with the same data, there is one woman with a trisomy 21 pregnancy and 4743 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		
Ri	isk Above Cut Off		Risk above Age	e Risk	Risk below Age risk