

 Date of Report
 18/12/2022

 PRISCA
 5.1.0.17

Patient Data					
Name	N	MRS. SUNITA	Patient ID		012212160223
Birthday		1/7/1993	Sample ID		11518667
Age at term		29.11	Sample Date		16/12/2022
Gestational age		13+2			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	52 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom			13+1
PAPP-A	$4.2~\mathrm{mIU/ml}$	0.5	Method		CRL (◇Robinson)
fb-hCG	15.6 ng/ml	0.49	Scan date		15/12/2022
Risks at sampling date			Nuchal translucency 2.4		
Age Risk 1		1:707	Nuchal translucency MoM 1.43		
Biochemical T21 risk 1:3485		1:3485	Nasal bone Present		
Combined trisomy 21 risk		1:3814			
Trisomy 13/18 + NT		1:3816			
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
1:100 1:250 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 1:3816, 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 3814 women with the same data, there is one woman with a trisomy 21 pregnancy and 3813 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		
Risk A	bove Cut Off		Risk above Age	e Risk	Risk below Age risk