

 Date of Report
 21/4/2023

 PRISCA
 5.2.0.13

Patient Data					
Name	MRS. PARUL	W/O MOHIT	Patient ID		012304190268
Birthday		25/06/1997	Sample ID		11645059
Age at term		26.2	Sample Date		19/4/2023
Gestational age		13+2			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	51 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom			13+0
PAPP-A	4.94 mIU/ml	0.57	Method		CRL (<>Robinson)
fb-hCG	67.2 ng/ml	2.12	Scan date		18/04/2023
Risks at sampling date			Nuchal translucency 1.3		
Age Risk		1:948	Nuchal translucency MoM		0.99
Biochemical T21 risk		1:265	Nasal bone Presen		
Combined trisomy 21 risk 1:3		- 1:1698			
Trisomy 13/18 + NT		< 1:10000			
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
1:100 1:250 1:1000 1:10000 1:1			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 with NT test it is expected that among 1698 women with the same data, there is one woman with a trisomy 21 pregnancy and 1697 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		values Risk above Ag	e Risk R	isk below Age risk