

Patient data				
Name	Mrs. W O SGT S S KAPOOR ANJANA		Patient ID	0352203260058
Birthday	21-05-1989		Sample ID	nsk82072
Age at sample date	32.8		Sample Date	26-03-2022
Gestational age	13 + 2			
Correction factors				
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies
Weight	62	diabetes	no	
Smoker	no	Origin	Asian	
Biochemical data			Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age	13 + 0
PAPP-A	3.71 mIU/mL	0.73	Method	CRL Robinson
fb-hCG	39.21 ng/mL	1.15	Scan date	24-03-2022
Risks at sampling date			Crown rump length in mm	69.8
Age risk	1:430		Nuchal translucency MoM	0.74
Biochemical T21 risk	1:953		Nasal bone	present
Combined trisomy 21 risk	1:5248		Sonographer	Dr.PADMA,M.B.B.S.,M.S
Trisomy 13/18 + NT	<1:10000		Qualifications in measuring NT	Dr.PADMA,M.B.B.S.,M.S
Risk			Trisomy 21	
			The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk. After the result of the Trisomy 21 test (with NT) it is expected that among 5248 women with the same data, there is one woman with a trisomy 21 pregnancy and 5247 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 risk calculations are statistical approaches and have no diagnostic value! The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)). The laboratory can not be hold responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!	
Trisomy 13/18 + NT				
The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.				

Sign of Physician