

Prisca 5.1.0.17
Date of report: 31/01/25

NA

Patient data			
Name	Mrs. SUPRIYA KUMARI	Patient ID	0482501290183
Birthday	28/06/03	Sample ID	A2230252
Age at sample date	21.6	Sample Date	29/01/25
Gestational age	12 + 4		
Correction factors			
Fetuses	1	IVF	no
Weight	65	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	3.38 mIU/mL	0.92	
fb-hCG	56.12 ng/mL	1.39	
Risks at sampling date			
Age risk		1:1065	
Biochemical T21 risk		1:2580	
Combined trisomy 21 risk		1:261	
Trisomy 13/18 + NT		<1:10000	
			Gestational age 12 + 5
			Method CRL Robinson
			Scan date 30/01/25
			Crown rump length in mm 65
			Nuchal translucency MoM 0.96
			Nasal bone absent
			Sonographer NA
			Qualifications in measuring NT MD
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 261 women with the same data, there is one woman with a trisomy 21 pregnancy and 260 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

below cut off
 Below Cut Off, but above Age Risk
 above cut off