

Prisca 5.1.0.17
Date of report: 05/07/25

NA

Patient data			
Name	Mrs. B SWAROOPA	Patient ID	0012507050308
Birthday	15/06/95	Sample ID	B3106020
Age at sample date	30.1	Sample Date	05/07/25
Gestational age	13 + 4		
Correction factors			
Fetuses	1	IVF	no
Weight	42	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	4.67 mIU/mL	0.53	
fb-hCG	32.51 ng/mL	0.90	
Risks at sampling date			
Age risk		1:665	
Biochemical T21 risk		1:1153	
Combined trisomy 21 risk		1:6668	
Trisomy 13/18 + NT		<1:10000	
			Gestational age 13 + 2
			Method CRL Robinson
			Scan date 03/07/25
			Crown rump length in mm 74
			Nuchal translucency MoM 0.77
			Nasal bone present
			Sonographer NA
			Qualifications in measuring NT NA
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 6668 women with the same data, there is one woman with a trisomy 21 pregnancy and 6667 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