

Prisca 5.1.0.17
Date of report: 09/10/25

N A

Patient data					
Name	Mrs. PRIYANKA KUMARI		Patient ID	0522510070029	
Birthday	11/12/01		Sample ID	b3810366	
Age at sample date	23.8		Sample Date	06/10/25	
Gestational age	11 + 6				
Correction factors					
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	
Weight	61	diabetes	no		
Smoker	no	Origin	Asian		
Biochemical data			Ultrasound data		
Parameter	Value	Corr. MoM	Gestational age	11 + 6	
PAPP-A	1.12 mIU/mL	0.38	Method	CRL Robinson	
fb-hCG	47.85 ng/mL	0.99	Scan date	06/10/25	
Risks at sampling date			Crown rump length in mm		
Age risk	1:982		54		
Biochemical T21 risk	1:564		Nuchal translucency MoM		
Combined trisomy 21 risk	1:3609		0.56		
Trisomy 13/18 + NT	<1:10000		Nasal bone		
			present		
			Sonographer		
			N A		
			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 3609 women with the same data, there is one woman with a trisomy 21 pregnancy and 3608 women with not affected pregnancies. The PAPP-A level is low. 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
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