

Prisca 5.1.0.17
Date of report: 27-10-2025

NA

Patient data			
Name	Mrs. SNEHA KUJUR	Patient ID	0442510250017
Birthday	29-10-2000	Sample ID	B3779813
Age at sample date	25.0	Sample Date	25-10-2025
Gestational age	12 + 0		
Correction factors			
Fetuses	1	IVF	no
Weight	53	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	5.16 mIU/mL	1.41	
fb-hCG	46.83 ng/mL	0.95	
Risks at sampling date		Gestational age	
Age risk		11 + 6	
Biochemical T21 risk		Method	
<1:10000		CRL Robinson	
Combined trisomy 21 risk		Scan date	
<1:10000		24-10-2025	
Trisomy 13/18 + NT		Crown rump length in mm	
<1:10000		54	
		Nuchal translucency MoM	
		0.77	
		Nasal bone	
		present	
		Sonographer	
		NA	
		Qualifications in measuring NT	
		NA	
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 more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy. 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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