

Prisca 5.1.0.17
Date of report: 08-02-2025

NA

Patient data			
Name	Mrs. SNEHLATA KUMARI	Patient ID	0522502060032
Birth day	21-08-1997	Sample ID	b2570390
Age at sample date	27.5	Sample Date	05-02-2025
Gestational age	11 + 2		
Correction factors			
Fetuses	1	IVF	no
Weight	56	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	1.94 mIU/mL	0.79	
fb-hCG	102.95 ng/mL	1.85	
Risks at sampling date			Gestational age
Age risk	1:792		11 + 2
Biochemical T21 risk	1:684		Method
Combined trisomy 21 risk	1:3941		CRL Robinson
Trisomy 13/18 + NT	<1:10000		Scan date
			05-02-2025
			Crown rump length in mm
			47.9
			Nuchal translucency MoM
			0.54
			Nasal bone
			present
			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 3941 women with the same data, there is one woman with a trisomy 21 pregnancy and 3940 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