

Prisca 5.1.0.17
Date of report: 09-07-2025

NA

Patient data			
Name	Mrs. MAMTA YADAV	Patient ID	0622507080016
Birthday	01-10-1990	Sample ID	b2601248
Age at sample date	34.8	Sample Date	08-07-2025
Gestational age	10 + 6		
Correction factors			
Fetuses	1	IVF	no
Weight	62	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	1.59 mIU/mL	0.91	Gestational age 10 + 5
fb-hCG	54.13 ng/mL	0.93	Method CRL Robinson
			Scan date 07-07-2025
Risks at sampling date			Trisomy 21
Age risk	1:266		Crown rump length in mm 40.82
Biochemical T21 risk	1:1606		Nuchal translucency MoM 1.04
Combined trisomy 21 risk	1:5783		Nasal bone unknown
Trisomy 13/18 + NT	<1:10000		Sonographer NA
			Qualifications in measuring NT NA
			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 5783 women with the same data, there is one woman with a trisomy 21 pregnancy and 5782 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 held 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