

Prisca 5.1.0.17
Date of report: 18-07-2025

N A

Patient data			
Name	Mrs. SABITA YADAV		Patient ID 0662507170246
Birthday	01-12-1998		Sample ID B2947122
Age at sample date	26.6		Sample Date 17-07-2025
Gestational age	13 + 3		
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	Gestational age 13 + 3
PAPP-A	2.71 mIU/mL	0.51	Method CRL Robinson
fb-hCG	34.04 ng/mL	1.04	Scan date 17-07-2025
Risks at sampling date			Crown rump length in mm 77
Age risk	1:908		Nuchal translucency MoM 1.01
Biochemical T21 risk	1:1010		Nasal bone present
Combined trisomy 21 risk	1:4344		Sonographer N A
Trisomy 13/18 + NT	<1:10000		Qualifications in measuring NT MD
Risk			Trisomy 21
The graph shows a risk curve that increases with age. At age 27, the risk is approximately 1:4344, which is below the cut-off (1:1000) but above the age risk (1:10000).			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 4344 women with the same data, there is one woman with a trisomy 21 pregnancy and 4343 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