

Prisca 5.1.0.17
Date of report: 23-07-2025

N A

Patient data			
Name	Mrs. YOGITA SAHU		Patient ID
Birthday	12-04-1987	Sample ID	
Age at sample date	38.3	Sample Date	
Gestational age	13 + 2		
Correction factors			
Fetuses	1	IVF	no
Weight	57	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	3.28 mIU/mL	0.58	
fb-hCG	30.1 ng/mL	0.86	
Risks at sampling date			
Age risk		1:128	
Biochemical T21 risk		1:313	
Combined trisomy 21 risk		1:1754	
Trisomy 13/18 + NT		<1:10000	
			Gestational age 13 + 1 Method CRL Robinson Scan date 21-07-2025 Crown rump length in mm 71 Nuchal translucency MoM 0.79 Nasal bone present Sonographer N A Qualifications in measuring NT MD
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 1754 women with the same data, there is one woman with a trisomy 21 pregnancy and 1753 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