

Prisca 5.1.0.17
Date of report: 09-06-2025

NA

Patient data			
Name	Mrs. PRITI KUMARI	Patient ID	0482506080160
Birthday	09-03-1988	Sample ID	B2710960
Age at sample date	37.2	Sample Date	08-06-2025
Gestational age	13 + 6		
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	12.96 mIU/mL	1.86	
fb-hCG	30.95 ng/mL	1.04	
Risks at sampling date			
Age risk		1:169	
Biochemical T21 risk		1:3140	
Combined trisomy 21 risk		1:9673	
Trisomy 13/18 + NT		<1:10000	
			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 9673 women with the same data, there is one woman with a trisomy 21 pregnancy and 9672 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