

Prisca 5.1.0.17
Date of report: 08-08-2025

N A

Patient data					
Name	Mrs. VAISHNAVI JADHAV		Patient ID	0662508070046	
Birthday	06-05-2001		Sample ID	B3298110	
Age at sample date	24.3		Sample Date	07-08-2025	
Gestational age	13 + 2				
Correction factors					
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	unknown
Weight	52	diabetes	no		
Smoker	no	Origin	Asian		
Biochemical data			Ultrasound data		
Parameter	Value	Corr. MoM	Gestational age	13 + 1	
PAPP-A	4.56 mIU/mL	0.73	Method	CRL Robinson	
fb-hCG	51.1 ng/mL	1.41	Scan date	06-08-2025	
Risks at sampling date			Crown rump length in mm	71	
Age risk	1:1016		Nuchal translucency MoM	0.68	
Biochemical T21 risk	1:1396		Nasal bone	present	
Combined trisomy 21 risk	1:7917		Sonographer	N A	
Trisomy 13/18 + NT	<1:10000		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 7917 women with the same data, there is one woman with a trisomy 21 pregnancy and 7916 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