

Prisca 5.1.0.17
Date of report: 22-04-2025

N A

Patient data			
Name	Mrs. SHAGUFTA JABEEN	Patient ID	0482504210154
Birthday	08-03-2002	Sample ID	A1861622
Age at sample date	23.1	Sample Date	21-04-2025
Gestational age	11 + 3		
Correction factors			
Fetuses	1	IVF	no
Weight	72	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	0.55 mIU/mL	0.28	
fb-hCG	51.32 ng/mL	1.02	
Risks at sampling date			Gestational age
Age risk		1:986	10 + 6
Biochemical T21 risk		1:209	Method
Combined trisomy 21 risk		1:1390	CRL Robinson
Trisomy 13/18 + NT		<1:10000	Scan date
			17-04-2025
			Crown rump length in mm
			42
			Nuchal translucency MoM
			0.85
			Nasal bone
			present
			Sonographer
			N A
			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 1390 women with the same data, there is one woman with a trisomy 21 pregnancy and 1389 women with not affected pregnancies. The PAPP-A level is low. 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

below cut off	Below Cut Off, but above Age Risk	above cut off
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