

Prisca 5.1.0.17
Date of report: 12-05-2025

N A

Patient data						
Name	Mrs. FARZANA PARWEEN		Patient ID	0472505110063		
Birthday	06-02-1993		Sample ID	B2788258		
Age at sample date	32.3		Sample Date	11-05-2025		
Gestational age	13 + 1					
Correction factors						
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	unknown	
Weight	65	diabetes	no			
Smoker	no	Origin	Asian			
Biochemical data			Ultrasound data			
Parameter	Value	Corr. MoM	Gestational age	13 + 1		
PAPP-A	2.61 mIU/mL	0.57	Method	CRL Robinson		
fb-hCG	36.52 ng/mL	1.05	Scan date	11-05-2025		
Risks at sampling date			Crown rump length in mm		71	
Age risk	1:474		Nuchal translucency MoM	0.79		
Biochemical T21 risk	1:713		Nasal bone	present		
Combined trisomy 21 risk	1:4132		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 4132 women with the same data, there is one woman with a trisomy 21 pregnancy and 4131 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