

Prisca 5.1.0.17
Date of report: 22-04-2025

N A

Patient data						
Name	Ms. NAJRINA BEGUM		Patient ID	0692504210109		
Birthday	20-08-1996		Sample ID	B2765422		
Age at sample date	28.7		Sample Date	21-04-2025		
Gestational age	13 + 2					
Correction factors						
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	unknown	
Weight	64	diabetes	no			
Smoker	no	Origin	Asian			
Biochemical data			Ultrasound data			
Parameter	Value	Corr. MoM	Gestational age	13 + 1		
PAPP-A	3.4 mIU/mL	0.69	Method	CRL Robinson		
fb-hCG	33.1 ng/mL	0.98	Scan date	20-04-2025		
Risks at sampling date			Crown rump length in mm		71.6	
Age risk	1:768		Nuchal translucency MoM	0.78		
Biochemical T21 risk	1:2164		Nasal bone	present		
Combined trisomy 21 risk	<1:10000		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 more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy. 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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