

Prisca 5.1.0.17  
Date of report: 12-05-2025

N A

Patient data				
Name	Mrs. NAMRATA A1960740		Patient ID	0772505090101
Birthday	15-12-1997		Sample ID	A1960740
Age at sample date	27.4		Sample Date	08-05-2025
Gestational age	13 + 0			
Correction factors				
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies
Weight	58	diabetes	no	
Smoker	no	Origin	Asian	
Biochemical data			Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age	12 + 2
PAPP-A	1.37 mIU/mL	0.28	Method	CRL Robinson
fb-hCG	65.63 ng/mL	1.75	Scan date	03-05-2025
Risks at sampling date			Crown rump length in mm	
Age risk	1:849		60.2	
Biochemical T21 risk	>1:50		Nuchal translucency MoM	
Combined trisomy 21 risk	1:154		1.15	
Trisomy 13/18 + NT	1:5784		Nasal bone	
			present	
			Sonographer	
			N A	
			Qualifications in measuring NT	
			MD	
Trisomy 21				
<p><b>The calculated risk for Trisomy 21 (with nuchal translucency) is above the cut off, which indicates an increased risk.</b></p> <p>After the result of the Trisomy 21 test (with NT) it is expected that among 154 women with the same data, there is one woman with a trisomy 21 pregnancy and 153 women with not affected pregnancies.</p> <p>The PAPP-A level is low.</p> <p>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!</p> <p>The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)).</p> <p>The laboratory can not be hold responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!</p>				
Trisomy 13/18 + NT				
<p><b>The calculated risk for Trisomy 13/18 (with nuchal translucency) is 1:5784, which represents a low risk.</b></p>				

Sign of Physician