

Prisca 5.1.0.17  
Date of report: 18-03-2025

N A

Patient data				
Name	Mrs. PRIYANKA KHAIRE TWIN B		Patient ID	0372503170027
Birthday	06-09-1997		Sample ID	A1552771
Age at sample date	27.5		Sample Date	17-03-2025
Gestational age	12 + 4			
Correction factors				
Fetuses	2	IVF	no	Previous trisomy 21 pregnancies
Weight	50	diabetes	no	
Smoker	no	Origin	Asian	
Biochemical data			Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age	12 + 2
PAPP-A	3.67 mIU/mL	0.40	Method	CRL Robinson
fb-hCG	116.43 ng/mL	1.22	Scan date	15-03-2025
Risks at sampling date			Crown rump length in mm	
Age risk	1:828		59	
Biochemical T21 risk	1:325		Nuchal translucency MoM	
Combined trisomy 21 risk	1:1614		0.97	
Trisomy 13/18 + NT	<1:10000		Nasal bone	
			present	
			Sonographer	
			N A	
			Qualifications in measuring NT	
			MD	
Risk			Trisomy 21	
			<p><b>The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.</b></p> <p>After the result of the Trisomy 21 test (with NT) it is expected that among 1614 women with the same data, there is one woman with a trisomy 21 pregnancy and 1613 women with not affected pregnancies.</p> <p>The PAPP-A level is low.</p> <p>The risk for this twin pregnancy has been calculated for a singleton pregnancy with corrected MoMs.</p> <p>The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician.</p> <p>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 &lt; 1:10000, which represents a low risk.</b></p>				

Sign of Physician