

Prisca 5.1.0.17
Date of report: 31-07-2025

N A

Patient data			
Name Mrs. PRIYANKA MEHETRE TWIN A		Patient ID 0832507300077	
Birthday 10-12-1993		Sample ID b3411551	
Age at sample date 31.6		Sample Date 30-07-2025	
Gestational age 12 + 4			
Correction factors			
Fetuses 2	IVF no	Previous trisomy 21 pregnancies unknown	
Weight 72	diabetes no		
Smoker no	Origin Asian		
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age 12 + 2
PAPP-A	7.79 mIU/mL	1.29	Method CRL Robinson
fb-hCG	42.02 ng/mL	0.50	Scan date 28-07-2025
Risks at sampling date			Crown rump length in mm 59
Age risk 1:514			Nuchal translucency MoM 0.81
Biochemical T21 risk <1:10000			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 risk for this twin pregnancy has been calculated for a singleton pregnancy with corrected MoMs. 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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Date of report: 31-07-2025

N A

Patient data					
Name	Mrs. PRIYANKA MEHETRE TWIN 2.		Patient ID	0832507300077	
Birthday	10-12-1993		Sample ID	b3411551B	
Age at sample date	31.6		Sample Date	30-07-2025	
Gestational age	12 + 1				
Correction factors					
Fetuses	2	IVF	no	Previous trisomy 21 pregnancies	unknown
Weight	72	diabetes	no		
Smoker	no	Origin	Asian		
Biochemical data			Ultrasound data		
Parameter	Value	Corr. MoM	Gestational age	11 + 6	
PAPP-A	7.79 mIU/mL	1.54	Method	CRL Robinson	
fb-hCG	42.02 ng/mL	0.45	Scan date	28-07-2025	
Risks at sampling date			Crown rump length in mm	55	
Age risk	1:506		Nuchal translucency MoM	0.77	
Biochemical T21 risk	<1:10000		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 risk for this twin pregnancy has been calculated for a singleton pregnancy with corrected MoMs. 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