

Prisca 5.1.0.17
Date of report: 25-07-2025

N A

Patient data				
Name	Mrs. A.ANUSHA. TWIN A.		Patient ID	0052507240014.
Birthday	10-05-2003		Sample ID	B2933652.
Age at sample date	22.2		Sample Date	24-07-2025
Gestational age	12 + 4			
Correction factors				
Fetuses	2	IVF	no	Previous trisomy 21 pregnancies
Weight	45	diabetes	no	
Smoker	no	Origin	Asian	
Biochemical data			Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age	12 + 4
PAPP-A	12.4 mIU/mL	1.18	Method	CRL Robinson
fb-hCG	42 ng/mL	0.42	Scan date	24-07-2025
Risks at sampling date			Crown rump length in mm	
Age risk	1:1052		63	
Biochemical T21 risk	<1:10000		Nuchal translucency MoM	
Combined trisomy 21 risk	<1:10000		0.86	
Trisomy 13/18 + NT	<1:10000		Nasal bone	
			present	
			Sonographer	
			N A	
			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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N A

Patient data			
Name	Mrs. A.ANUSHA. TWIN B..	Patient ID	0052507240014.
Birthday	10-05-2003	Sample ID	B2933652.
Age at sample date	22.2	Sample Date	24-07-2025
Gestational age	12 + 6		
Correction factors			
Fetuses	2	IVF	no
Weight	45	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	12.4 mIU/mL	1.06	
fb-hCG	42 ng/mL	0.45	
Risks at sampling date			
Age risk	1:1062	Gestational age	12 + 6
Biochemical T21 risk	<1:10000	Method	CRL Robinson
Combined trisomy 21 risk	<1:10000	Scan date	24-07-2025
Trisomy 13/18 + NT	<1:10000	Crown rump length in mm	68
		Nuchal translucency MoM	0.81
		Nasal bone	present
		Sonographer	N A
		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