

Prisca 5.1.0.17
Date of report: 25-07-2025

N A

Patient data					
Name	Mrs. KRISHNA SAMANTA. TWIN A.		Patient ID	0622507220096.	
Birthday	15-10-2002		Sample ID	A1820144.	
Age at sample date	22.8		Sample Date	22-07-2025	
Gestational age	13 + 5				
Correction factors					
Fetuses	2	IVF	no	Previous trisomy 21 pregnancies	unknown
Weight	58	diabetes	no		
Smoker	no	Origin	Asian		
Biochemical data			Ultrasound data		
Parameter	Value	Corr. MoM	Gestational age	13 + 4	
PAPP-A	8.51 mIU/mL	0.72	Method	CRL Robinson	
fb-hCG	32.72 ng/mL	0.49	Scan date	21-07-2025	
Risks at sampling date			Crown rump length in mm	78	
Age risk	1:1077		Nuchal translucency MoM	0.53	
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

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Date of report: 25-07-2025

N A

Patient data			
Name Mrs. KRISHNA SAMANTA. TWIN B..		Patient ID 0622507220096.	
Birthday 15-10-2002		Sample ID A1820144.	
Age at sample date 22.8		Sample Date 22-07-2025	
Gestational age 13 + 3			
Correction factors			
Fetuses 2	IVF no	Previous trisomy 21 pregnancies unknown	
Weight 58	diabetes no		
Smoker no	Origin Asian		
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age 13 + 2
PAPP-A	8.51 mIU/mL	0.79	Method CRL Robinson
fb-hCG	32.72 ng/mL	0.45	Scan date 21-07-2025
Risks at sampling date			Crown rump length in mm 75
Age risk 1:1068			Nuchal translucency MoM 0.49
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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