

Prisca 5.1.0.17
Date of report: 13/10/25

N A

Patient data				
Name	Mrs. ANJALI RATHOD TWIN A		Patient ID	0672510120117
Birthday	18/03/04		Sample ID	B3745002 TWIN A
Age at sample date	21.6		Sample Date	12/10/25
Gestational age	13 + 0			
Correction factors				
Fetuses	2	IVF	no	Previous trisomy 21 pregnancies
Weight	48	diabetes	no	
Smoker	no	Origin	Asian	
Biochemical data			Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age	12 + 6
PAPP-A	12.9 mIU/mL	1.12	Method	CRL Robinson
fb-hCG	36.85 ng/mL	0.42	Scan date	11/10/25
Risks at sampling date			Crown rump length in mm	
Age risk	1:1081		67.9	
Biochemical T21 risk	<1:10000		Nuchal translucency MoM	
Combined trisomy 21 risk	<1:10000		1.46	
Trisomy 13/18 + NT	<1:10000		Nasal bone	
			unknown	
			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. ANJALI RATHOD TWIN B		Patient ID 0672510120117	
Birthday 18/03/04		Sample ID B3745002 TWIN B	
Age at sample date 21.6		Sample Date 12/10/25	
Gestational age 13 + 0			
Correction factors			
Fetuses 2	IVF no	Previous trisomy 21 pregnancies unknown	
Weight 48	diabetes no		
Smoker no	Origin Asian		
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age 12 + 6
PAPP-A	12.9 mIU/mL	1.12	Method CRL Robinson
fb-hCG	36.85 ng/mL	0.42	Scan date 11/10/25
Risks at sampling date			Crown rump length in mm 68.3
Age risk 1:1081			Nuchal translucency MoM 0.87
Biochemical T21 risk <1:10000			Nasal bone unknown
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