

Prisca 5.1.0.17
Date of report: 31/12/25

Patient data				
Name	Mrs. PRATIKSHA AUTADE TWIN A		Patient ID	0672512300169
Birthday	22/02/03		Sample ID	D0052144 TWIN A
Age at sample date	22.9		Sample Date	30/12/25
Gestational age	12 + 2			
Correction factors				
Fetuses	2	IVF	no	Previous trisomy 21 pregnancies
Weight	59	diabetes	no	
Smoker	no	Origin	Asian	
Biochemical data			Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age	12 + 2
PAPP-A	10.48 mIU/mL	1.54	Method	CRL Robinson
fb-hCG	40.01 ng/mL	0.42	Scan date	30/12/25
Risks at sampling date			Crown rump length in mm	
Age risk	1:1026		59.7	
Biochemical T21 risk	<1:10000		Nuchal translucency MoM	
Combined trisomy 21 risk	<1:10000		1.09	
Trisomy 13/18 + NT	<1:10000		Nasal bone	
			present	
			Sonographer	
			NA	
			Qualifications in measuring NT	
			Sonographer	
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: 31/12/25

Patient data			
Name	Mrs. PRATIKSHA AUTADE TWIN E	Patient ID	0672512300169
Birthday	22/02/03	Sample ID	D0052144 TWIN B
Age at sample date	22.9	Sample Date	30/12/25
Gestational age	12 + 2		
Correction factors			
Fetuses	2	IVF	no
Weight	59	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	10.48 mIU/mL	1.54	
fb-hCG	40.01 ng/mL	0.42	
Risks at sampling date			
Age risk	1:1026	Gestational age	12 + 2
Biochemical T21 risk	<1:10000	Method	CRL Robinson
Combined trisomy 21 risk	<1:10000	Scan date	30/12/25
Trisomy 13/18 + NT	<1:10000	Crown rump length in mm	59.9
		Nuchal translucency MoM	0.96
		Nasal bone	present
		Sonographer	NA
		Qualifications in measuring NT	Sonographer
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