

Prisca 5.1.0.17
Date of report: 08-03-2025

N A

Patient data				
Name	Mrs. BHAVYA SRI TWIN A		Patient ID	0842503070027
Birthday	12-09-1998		Sample ID	A2144795
Age at sample date	26.5		Sample Date	07-03-2025
Gestational age	11 + 5			
Correction factors				
Fetuses	2	IVF	no	Previous trisomy 21 pregnancies
Weight	75	diabetes	no	
Smoker	no	Origin	Asian	
Biochemical data			Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age	11 + 5
PAPP-A	9.97 mIU/mL	2.52	Method	CRL Robinson
fb-hCG	48.08 ng/mL	0.48	Scan date	07-03-2025
Risks at sampling date			Crown rump length in mm	
Age risk	1:863		52	
Biochemical T21 risk	<1:10000		Nuchal translucency MoM	
Combined trisomy 21 risk	<1:10000		0.50	
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 PAPP-A level is high. 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
---	--	---

Prisca 5.1.0.17
Date of report: 08-03-2025

N A

Patient data			
Name	Mrs. BHAVYA SRI TWIN B		Patient ID
Birthday	12-09-1998		Sample ID
Age at sample date	26.5		Sample Date
Gestational age	11 + 6		
Correction factors			
Fetuses	2	IVF	no
Weight	75	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	9.97 mIU/mL	2.36	11 + 6
fb-hCG	48.08 ng/mL	0.49	Method
			CRL Robinson
			Scan date
			07-03-2025
Risks at sampling date			Crown rump length in mm
Age risk			54
Biochemical T21 risk			Nuchal translucency MoM
<1:10000			0.42
Combined trisomy 21 risk			Nasal bone
<1:10000			present
Trisomy 13/18 + NT			Sonographer
<1:10000			N A
			Qualifications in measuring NT
			MD
Risk			Trisomy 21
1:10			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!
1:100			
1:250			
1:1000			
1:10000			
Cut off			
Age			
13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49			
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