

Prisca 5.1.0.17
Date of report: 04-07-2025

N A

Patient data			
Name	Mrs. HEMA TWIN A	Patient ID	0772507030140
Birthday	03-07-1987	Sample ID	B2835764
Age at sample date	38.0	Sample Date	03-07-2025
Gestational age	12 + 3		
Correction factors			
Fetuses	2	IVF	no
Weight	63	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	5.98 mIU/mL	0.89	Gestational age 12 + 1
fb-hCG	43.85 ng/mL	0.48	Method CRL Robinson
Risks at sampling date			Scan date 01-07-2025
Age risk	1:133		Crown rump length in mm 58.6
Biochemical T21 risk	1:2996		Nuchal translucency MoM 0.91
Combined trisomy 21 risk	<1:10000		Nasal bone present
Trisomy 13/18 + NT	<1:10000		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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Patient data					
Name	Mrs. HEMA TWIN B		Patient ID	0772507030140	
Birthday	03-07-1987		Sample ID	B2835764	
Age at sample date	38.0		Sample Date	03-07-2025	
Gestational age	12 + 2				
Correction factors					
Fetuses	2	IVF	no	Previous trisomy 21 pregnancies	unknown
Weight	63	diabetes	no		
Smoker	no	Origin	Asian		
Biochemical data			Ultrasound data		
Parameter	Value	Corr. MoM	Gestational age	12 + 0	
PAPP-A	5.98 mIU/mL	0.95	Method	CRL Robinson	
fb-hCG	43.85 ng/mL	0.47	Scan date	01-07-2025	
Risks at sampling date			Crown rump length in mm	56.1	
Age risk	1:132		Nuchal translucency MoM	0.81	
Biochemical T21 risk	1:3629		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