

Prisca 5.1.0.17
Date of report: 21/10/24

N A

Patient data			
Name	Mrs. PARVEEN TWIN A	Patient ID	0042410200005
Birthday	04/02/98	Sample ID	23125539 TWIN A
Age at sample date	26.7	Sample Date	20/10/24
Gestational age	12 + 3		
Correction factors			
Fetuses	2	IVF	no
Weight	50	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	4.1 mIU/mL	0.47	12 + 3
fb-hCG	63.25 ng/mL	0.64	Method
			CRL Robinson
			Scan date
			20/10/24
			Crown rump length in mm
			62
			Nuchal translucency MoM
			1.00
			Nasal bone
			unknown
			Sonographer
			N A
			Qualifications in measuring NT
			MD
Risks at sampling date		Trisomy 21	
Age risk	1:874	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 9693 women with the same data, there is one woman with a trisomy 21 pregnancy and 9692 women with not affected pregnancies. 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!	
Biochemical T21 risk	1:2201		
Combined trisomy 21 risk	1:9693		
Trisomy 13/18 + NT	<1:10000		
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. PARVEEN TWIN B		Patient ID
Birth day	04/02/98		Sample ID
Age at sample date	26.7		Sample Date
Gestational age	12 + 4		
Correction factors			
Fetuses	2	IVF	no
Weight	50	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	4.1 mIU/mL	0.44	Method
fb-hCG	63.25 ng/mL	0.66	Scan date
Risks at sampling date			Crown rump length in mm
Age risk	1:878		Nuchal translucency MoM
Biochemical T21 risk	1:1760		Nasal bone
Combined trisomy 21 risk	1:9244		Sonographer
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 9244 women with the same data, there is one woman with a trisomy 21 pregnancy and 9243 women with not affected pregnancies. 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