

Prisca 5.1.0.17
Date of report: 14/02/25

N A

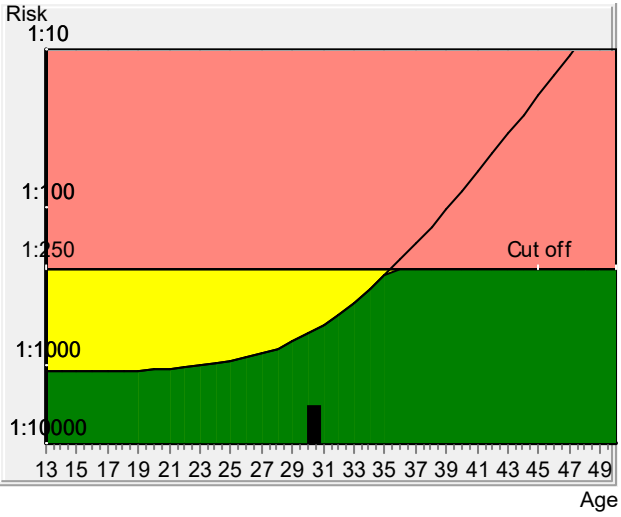
Patient data				
Name	Mrs. SIMRAN KAUR TWIN A		Patient ID	0372502130107
Birthday	29/08/94		Sample ID	A1641549
Age at sample date	30.5		Sample Date	13/02/25
Gestational age	12 + 3			
Correction factors				
Fetuses	2	IVF	no	Previous trisomy 21 pregnancies
Weight	60	diabetes	no	
Smoker	no	Origin	Asian	
Biochemical data			Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age	12 + 3
PAPP-A	5.9 mIU/mL	0.83	Method	CRL Robinson
fb-hCG	160.73 ng/mL	1.74	Scan date	13/02/25
Risks at sampling date			Crown rump length in mm	
Age risk	1:607		61	
Biochemical T21 risk	1:689		Nuchal translucency MoM	
Combined trisomy 21 risk	1:3882		0.70	
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 3882 women with the same data, there is one woman with a trisomy 21 pregnancy and 3881 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

below cut off
 Below Cut Off, but above Age Risk
 above cut off

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Patient data			
Name	Mrs. SIMRAN KAUR TWIN B	Patient ID	0372502130107
Birthday	29/08/94	Sample ID	A1641549
Age at sample date	30.5	Sample Date	13/02/25
Gestational age	12 + 2		
Correction factors			
Fetuses	2	IVF	no
Weight	60	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data			Ultrasound data
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	5.9 mIU/mL	0.88	12 + 2
fb-hCG	160.73 ng/mL	1.68	Method
			CRL Robinson
			Scan date
			13/02/25
			Crown rump length in mm
			59
			Nuchal translucency MoM
			0.78
			Nasal bone
			unknown
			Sonographer
			N A
			Qualifications in measuring NT
			MD
Risks at sampling date			Trisomy 21
Age risk		1:603	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 4686 women with the same data, there is one woman with a trisomy 21 pregnancy and 4685 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		1:848	
Combined trisomy 21 risk		1:4686	
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