

Prisca

5.1.0.17

Date of report:

20/11/24

NA

Patient data			
Name	Mrs. KALPANA MISHRA TWIN A		Patient ID
Birthday	16/11/91	Sample ID	A0323189 TWIN A
Age at sample date	33.0	Sample Date	16/11/24
Gestational age	12 + 6		
Correction factors			
Fetuses	2	IVF	no
Weight	71	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	5.34 mIU/mL	0.69	Method
fb-hCG	82.87 ng/ml	1.10	CRL Robinson
Risks at sampling date			Scan date
Age risk	1:412		12/11/24
Biochemical T21 risk	1:879		Crown rump length in mm
Combined trisomy 21 risk	1:2905		60
Trisomy 13/18 + NT	<1:10000		Nuchal translucency MoM
			1.09
			Nasal bone
			present
			Sonographer
			NA
			Qualifications in measuring NT
			MD
		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 2905 women with the same data, there is one woman with a trisomy 21 pregnancy and 2904 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

Prisca

5.1.0.17

Date of report:

20/11/24

NA

Patient data				
Name	Mrs. KALPANA MISHRA TWIN B		Patient ID	0782411160013
Birthday	16/11/91		Sample ID	A0323189 TWIN B
Age at sample date	33.0		Sample Date	16/11/24
Gestational age	13 + 3			
Correction factors				
Fetuses	2	IVF	no	Previous trisomy 21 pregnancies
Weight	71	diabetes	no	
Smoker	no	Origin	Asian	
Biochemical data			Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age	12 + 6
PAPP-A	5.34 mIU/mL	0.53	Method	CRL Robinson
fb-hCG	82.87 ng/ml	1.25	Scan date	12/11/24
Risks at sampling date			Crown rump length in mm	68
Age risk	1:420		Nuchal translucency MoM	1.05
Biochemical T21 risk	1:354		Nasal bone	present
Combined trisomy 21 risk	1:1381		Sonographer	NA
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 1381 women with the same data, there is one woman with a trisomy 21 pregnancy and 1380 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