

Prisca 5.1.0.17
Date of report: 11/12/25

Patient data			
Name Mrs. AROSHREE BINDHANI TWIN A		Patient ID 0652512090146	
Birthday 09/12/97		Sample ID B4056009 TWIN A	
Age at sample date 28.0		Sample Date 09/12/25	
Gestational age 12 + 0			
Correction factors			
Fetuses 2	IVF no	Previous trisomy 21 pregnancies unknown	
Weight 46	diabetes no		
Smoker no	Origin Asian		
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age 11 + 5
PAPP-A	5.37 mIU/mL	0.67	Method CRL Robinson
fb-hCG	48.52 ng/mL	0.43	Scan date 07/12/25
Risks at sampling date			Crown rump length in mm 53.1
Age risk 1:780			Nuchal translucency MoM 0.85
Biochemical T21 risk <1:10000			Nasal bone present
Combined trisomy 21 risk <1:10000			Sonographer NA
Trisomy 13/18 + NT <1:10000			Qualifications in measuring NT Sonographer
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

Prisca 5.1.0.17
Date of report: 11-12-2025

Patient data			
Name	Mrs. AROSHREE BINDHANI TWIN B	Patient ID	0652512090146
Birthday	09-12-1997	Sample ID	B4056009
Age at sample date	28.0	Sample Date	09-12-2025
Gestational age	11 + 6		
Correction factors			
Fetuses	2	IVF	no
Weight	46	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	5.37 mIU/mL	0.71	11 + 4
fb-hCG	48.52 ng/mL	0.42	Method
			CRL Robinson
			Scan date
			07-12-2025
Risks at sampling date			Crown rump length in mm
Age risk	1:776		51.1
Biochemical T21 risk	<1:10000		Nuchal translucency MoM
Combined trisomy 21 risk	<1:10000		0.80
Trisomy 13/18 + NT	<1:10000		Nasal bone
			present
			Sonographer
			NA
			Qualifications in measuring NT
			Sonographer
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