

Prisca 5.1.0.17
Date of report: 12/12/25

Patient data			
Name	Mrs. JYOTIMA BARIK TWIN A	Patient ID	0622512100089
Birthday	02/06/95	Sample ID	A1448879 TWIN A
Age at sample date	30.5	Sample Date	10/12/25
Gestational age	13 + 1		
Correction factors			
Fetuses	2	IVF	no
Weight	45	diabetes	no
Smoker	no	Origin	Asian
Biochemical data			
Parameter	Value	Corr. MoM	Ultrasound data
PAPP-A	7.83 mIU/mL	0.60	Gestational age 13 + 0
fb-hCG	36.95 ng/mL	0.43	Method CRL Robinson
Risks at sampling date			
Age risk	1:617		Scan date 09/12/25
Biochemical T21 risk	1:6607		Crown rump length in mm 70.7
Combined trisomy 21 risk	<1:10000		Nuchal translucency MoM 0.90
Trisomy 13/18 + NT	<1:10000		Nasal bone present
			Sonographer NA
			Qualifications in measuring NT Sonographer
Risk			
1:10			Trisomy 21
1:100			The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.
1:250			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.
1:1000			The risk for this twin pregnancy has been calculated for a singleton pregnancy with corrected MoMs.
1:10000			The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician.
13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49	Age		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 held 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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Date of report: 12/12/25

Patient data			
Name	Mrs. JYOTIMA BARIK	Patient ID	0622512100089
Birthday	02/06/95	Sample ID	A1448879 TWIN B
Age at sample date	30.5	Sample Date	10/12/25
Gestational age	12 + 4		
Correction factors			
Fetuses	2	IVF	no
Weight	45	diabetes	no
Smoker	no	Origin	Asian
Biochemical data			
Parameter	Value	Corr. MoM	Ultrasound data
PAPP-A	7.83 mIU/mL	0.75	Gestational age 12 + 3
fb-hCG	36.95 ng/mL	0.37	Method CRL Robinson
Risks at sampling date			
Age risk	1:605		Scan date 09/12/25
Biochemical T21 risk	<1:10000		Crown rump length in mm 62.3
Combined trisomy 21 risk	<1:10000		Nuchal translucency MoM 1.12
Trisomy 13/18 + NT	<1:10000		Nasal bone present
			Sonographer NA
			Qualifications in measuring NT Sonographer
Trisomy 21			
The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk. <p>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.</p> <p>The free beta HCG level is low.</p> <p>The risk for this twin pregnancy has been calculated for a singleton pregnancy with corrected MoMs.</p> <p>The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician.</p> <p>Please note that risk calculations are statistical approaches and have no diagnostic value!</p> <p>The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)).</p> <p>The laboratory can not be held responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!</p>			
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