

Prisca 5.1.0.17
Date of report: 17-07-2025

NA

Patient data			
Name		Mrs. BANITA BARIK, FETUS-A	
Patient ID		0652507140070	
Birthday		09-08-1997	
Sample ID		B2895048 FETUS-A	
Age at sample date		27.9	
Sample Date		14-07-2025	
Gestational age		12 + 3	
Correction factors			
Fetuses	2	IVF	no
Weight	54	diabetes	no
Smoker	no	Origin	Asian
Previous trisomy 21 pregnancies		unknown	
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	10.43 mIU/mL	1.30	12 + 3
fb-hCG	39.1 ng/mL	0.41	Method
Risks at sampling date			CRL Robinson
Age risk			14-07-2025
Biochemical T21 risk			61.8
Combined trisomy 21 risk			0.94
Trisomy 13/18 + NT			0.94
			Nasal bone
			present
			Sonographer
			NA
			Qualifications in measuring NT
			NA
Risk		Trisomy 21	
1:10		The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.	
1:100		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.	
1:250		The risk for this twin pregnancy has been calculated for a singleton pregnancy with corrected MoMs.	
1:1000		The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician.	
1:10000		Please note that risk calculations are statistical approaches and have no diagnostic value!	
13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49		The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)).	
Age		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: 17-07-2025

NA

Patient data				
Name	Mrs. BANITA BARIK, FETUS-B		Patient ID	0652507140070
Birthday	09-08-1997		Sample ID	B2895048 FETUS-B
Age at sample date	27.9		Sample Date	14-07-2025
Gestational age	13 + 0			
Correction factors				
Fetuses	2	IVF	no	Previous trisomy 21 pregnancies
Weight	54	diabetes	no	
Smoker	no	Origin	Asian	
Biochemical data			Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age	13 + 0
PAPP-A	10.43 mIU/mL	1.04	Method	CRL Robinson
fb-hCG	39.1 ng/mL	0.47	Scan date	14-07-2025
Risks at sampling date			Crown rump length in mm	
Age risk	1:813		70.5	
Biochemical T21 risk	<1:10000		Nuchal translucency MoM	
Combined trisomy 21 risk	<1:10000		1.08	
Trisomy 13/18 + NT	<1:10000		Nasal bone	
			present	
			Sonographer	
			NA	
			Qualifications in measuring NT	
			NA	
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
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