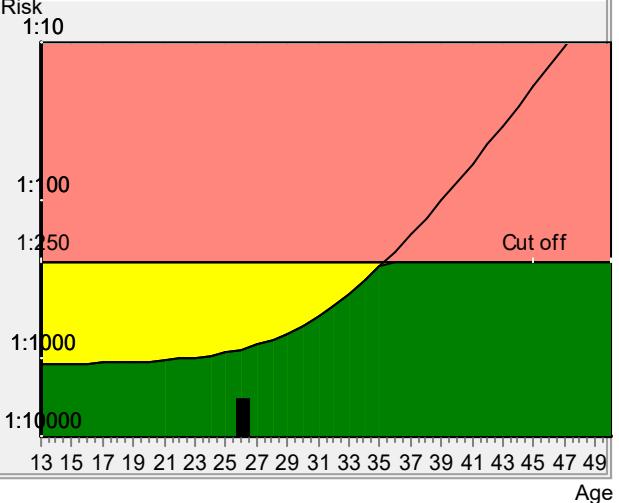


TAHERA FATHIMA

Patient data								
Name	Mrs. M BHARGAVI TWIN A	Patient ID	0012509260247					
Birthday	30-07-1999	Sample ID	B3533063					
Age at sample date	26.2	Sample Date	26-09-2025					
Gestational age	11 + 5							
Correction factors								
Fetuses	2	IVF	no	Previous trisomy 21 pregnancies	unknown			
Weight	61	diabetes	no					
Smoker	no	Origin	Asian					
Biochemical data								
Parameter	Value	Corr. MoM						
PAPP-A	4.89 mIU/mL	0.96						
fb-hCG	46.07 ng/mL	0.43						
Risks at sampling date								
Age risk	1:880		11 + 5					
Biochemical T21 risk	<1:10000		CRL Robinson					
Combined trisomy 21 risk	<1:10000		Scan date 26-09-2025					
Trisomy 13/18 + NT	<1:10000		Crown rump length in mm 53					
Ultrasound data								
Gestational age			Nuchal translucency MoM 0.85					
Method			Nasal bone present					
Scan date			Sonographer N A					
Crown rump length in mm			Qualifications in measuring NT MD					
Nuchal translucency MoM								
Nasal bone								
Sonographer								
Qualifications in measuring NT								
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 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

TAHERA FATHIMA

Patient data								
Name	Mrs. M BHARGAVI TWIN B	Patient ID	0012509260247					
Birthday	30-07-1999	Sample ID	B3533063					
Age at sample date	26.2	Sample Date	26-09-2025					
Gestational age	11 + 5							
Correction factors								
Fetuses	2	IVF	no	Previous trisomy 21 pregnancies	unknown			
Weight	61	diabetes	no					
Smoker	no	Origin	Asian					
Biochemical data								
Parameter	Value	Corr. MoM	Ultrasound data					
PAPP-A	4.89 mIU/mL	0.96	Gestational age					
fb-hCG	46.07 ng/mL	0.43	11 + 5					
Risks at sampling date								
Age risk	1:880		Method					
Biochemical T21 risk	<1:10000		CRL Robinson					
Combined trisomy 21 risk	<1:10000		Scan date					
Trisomy 13/18 + NT	<1:10000		26-09-2025					
Risk								
1:10			Crown rump length in mm					
1:100			52					
1:250			Nuchal translucency MoM					
1:1000			0.79					
1:10000			Nasal bone					
13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49			present					
			Sonographer					
			N A					
			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. <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 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