

Prisca 5.1.0.17
Date of report: 19-11-2025

N A

Patient data							
Name	Mrs. SUNITA PANDHARE	Patient ID	0662511180205				
Birthday	08-06-1992	Sample ID	B3725921				
Age at sample date	33.4	Sample Date	18-11-2025				
Gestational age	13 + 0						
Correction factors							
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	unknown		
Weight	71	diabetes	no				
Smoker	no	Origin	Asian				
Biochemical data							
Parameter	Value	Corr. MoM					
PAPP-A	2.86 mIU/mL	0.73					
fb-hCG	36.97 ng/mL	1.05					
Risks at sampling date							
Age risk	1:381						
Biochemical T21 risk	1:1063						
Combined trisomy 21 risk	1:5766						
Trisomy 13/18 + NT	<1:10000						
Ultrasound data							
			Gestational age	13 + 0			
			Method	CRL Robinson			
			Scan date	18-11-2025			
			Crown rump length in mm	70			
			Nuchal translucency MoM	0.57			
			Nasal bone	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.							
After the result of the Trisomy 21 test (with NT) it is expected that among 5766 women with the same data, there is one woman with a trisomy 21 pregnancy and 5765 women with not affected pregnancies.							
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

below cut off

Below Cut Off, but above Age Risk

above cut off