

Prisca 5.1.0.17  
Date of report: 31-01-2026

Patient data				
Name	Mrs. CHANDANI DEVI		Patient ID	0522601300044
Birthday	22-02-1998		Sample ID	B4381833
Age at sample date	27.9		Sample Date	29-01-2026
Gestational age	11 + 5			
Correction factors				
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies
Weight	56	diabetes	no	
Smoker	no	Origin	Asian	
Biochemical data			Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age	10 + 5
PAPP-A	1.35 mIU/mL	0.45	Method	CRL Robinson
fb-hCG	46.21 ng/mL	0.90	Scan date	22-01-2026
Risks at sampling date			Crown rump length in mm	
Age risk	1:776		41	
Biochemical T21 risk	1:838		Nuchal translucency MoM	
Combined trisomy 21 risk	1:4825		0.86	
Trisomy 13/18 + NT	<1:10000		Nasal bone	
			present	
			Sonographer	
			NA	
			Qualifications in measuring NT	
			Sonographer	
Trisomy 21				
<p><b>The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.</b></p> <p>After the result of the Trisomy 21 test (with NT) it is expected that among 4825 women with the same data, there is one woman with a trisomy 21 pregnancy and 4824 women with not affected pregnancies.</p> <p>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!</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 hold responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!</p>				
Trisomy 13/18 + NT				
<p><b>The calculated risk for trisomy 13/18 (with nuchal translucency) is &lt; 1:10000, which represents a low risk.</b></p>				

Sign of Physician

below cut off
  Below Cut Off, but above Age Risk
  above cut off