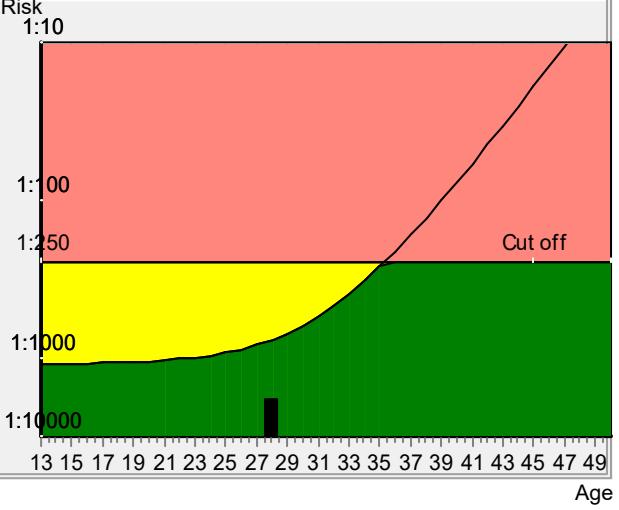


Patient data								
Name	Mrs. CHANDANI DEVI	Patient ID	0522601300044					
Birthday	22-02-1998	Sample ID	B4381833					
Age at sample date	27.9	Sample Date	29-01-2026					
Gestational age	11 + 5							
Correction factors								
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	unknown			
Weight	56	diabetes	no					
Smoker	no	Origin	Asian					
Biochemical data								
Parameter	Value	Corr. MoM	Ultrasound data					
PAPP-A	1.35 mIU/mL	0.45	Gestational age 10 + 5					
fb-hCG	46.21 ng/mL	0.90	Method CRL Robinson					
Risks at sampling date								
Age risk	1:776		Scan date 22-01-2026					
Biochemical T21 risk	1:838		Crown rump length in mm 41					
Combined trisomy 21 risk	1:4825		Nuchal translucency MoM 0.86					
Trisomy 13/18 + NT	<1:10000		Nasal bone present					
Sonographer NA								
Qualifications in measuring NT Sonographer								
Trisomy 21								
<b>The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.</b> <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 held responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!</p>								
								
Trisomy 13/18 + NT								
<b>The calculated risk for trisomy 13/18 (with nuchal translucency) is &lt; 1:10000, which represents a low risk.</b>								

Sign of Physician

 below cut off

 Below Cut Off, but above Age Risk

 above cut off