

Prisca 5.1.0.17  
**Date of report:** 28-03-2025

N A

Patient data								
Name	A0833238	Patient ID	0312503270036					
Birthday	06-06-2001	Sample ID	Mrs. NIKITHA					
Age at sample date	23.8	Sample Date	27-03-2025					
Gestational age	13 + 5							
Correction factors								
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	unknown			
Weight	55	diabetes	no					
Smoker	no	Origin	Asian					
Biochemical data								
Parameter	Value	Corr. MoM	Ultrasound data					
PAPP-A	4.98 mIU/mL	0.73	Gestational age 13 + 4					
fb-hCG	32.52 ng/mL	1.04	Method CRL Robinson					
Risks at sampling date								
Age risk	1:1046		Scan date 26-03-2025					
Biochemical T21 risk	1:2960		Crown rump length in mm 78					
Combined trisomy 21 risk	<1:10000		Nuchal translucency MoM 0.84					
Trisomy 13/18 + NT	<1:10000		Nasal bone present					
Risk								
1:10			Sonographer N A					
1:100			Qualifications in measuring NT MD					
1:250			Trisomy 21					
1:1000			The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.					
1:10000			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.					
13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49			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