

Prisca 5.1.0.17
Date of report: 01-04-2025

N A

Patient data			
Name	Mrs. NIKITA GAUR		Patient ID 0372503310062
Birthday	09-10-1994		Sample ID B2401729
Age at sample date	30.5		Sample Date 31-03-2025
Gestational age	13 + 2		
Correction factors			
Fetuses	1	IVF	no
Weight	60	diabetes	no
Smoker	no	Origin	Asian
			Previous trisomy 21 pregnancies unknown
Biochemical data			Ultrasound data
Parameter	Value	Corr. MoM	Gestational age 11 + 1
PAPP-A	2.43 mIU/mL	0.46	Method CRL Robinson
fb-hCG	33.1 ng/mL	0.96	Scan date 16-03-2025
Risks at sampling date			Crown rump length in mm 46
Age risk	1:624		Nuchal translucency MoM 1.11
Biochemical T21 risk	1:635		Nasal bone present
Combined trisomy 21 risk	1:2110		Sonographer N A
Trisomy 13/18 + NT	<1:10000		Qualifications in measuring NT MD
Risk			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 2110 women with the same data, there is one woman with a trisomy 21 pregnancy and 2109 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 hold 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