

Prisca 5.1.0.17
Date of report: 03-08-2025

N A

Patient data			
Name	Mrs. SUNITA SHARMA	Patient ID	0622508020051
Birthday	01-07-1985	Sample ID	B2808849
Age at sample date	40.1	Sample Date	02-08-2025
Gestational age	12 + 0		
Correction factors			
Fetuses	1	IVF	no
Weight	59	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	3.12 mIU/mL	0.96	Gestational age 11 + 6
fb-hCG	45.27 ng/mL	0.96	Method CRL Robinson
Risks at sampling date			Scan date 01-08-2025
Age risk	1:75		Crown rump length in mm 53.4
Biochemical T21 risk	1:479		Nuchal translucency MoM 1.20
Combined trisomy 21 risk	1:1093		Nasal bone present
Trisomy 13/18 + NT	<1:10000		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 1093 women with the same data, there is one woman with a trisomy 21 pregnancy and 1092 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
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