

Prisca 5.1.0.17
Date of report: 22-02-2025

N A

Patient data			
Name	Mrs. SHANTI MAURYA		Patient ID
Birthday	01-05-1992		Sample ID
Age at sample date	32.8		Sample Date
Gestational age	12 + 3		
Correction factors			
Fetuses	1	IVF	no
Weight	57	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	2.08 mIU/mL	0.51	12 + 2
fb-hCG	41.12 ng/mL	0.94	Method
			CRL Robinson
			Scan date
			20-02-2025
Risks at sampling date			Trisomy 21
Age risk			1:420
Biochemical T21 risk			1:600
Combined trisomy 21 risk			1:2192
Trisomy 13/18 + NT			<1:10000
			Crown rump length in mm
			60
			Nuchal translucency MoM
			1.07
			Nasal bone
			present
			Sonographer
			N A
			Qualifications in measuring NT
			MD
Risk Cut off Age 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 2192 women with the same data, there is one woman with a trisomy 21 pregnancy and 2191 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 21 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