

Prisca 5.1.0.17
Date of report: 27-01-2026

Patient data			
Name	Mrs. POOJA MAURYA		Patient ID
Birthday	24-01-1994		Sample ID
Age at sample date	32.0		Sample Date
Gestational age	12 + 2		
Correction factors			
Fetuses	1	IVF	no
Weight	45	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	3.03 mIU/mL	0.61	12 + 2
fb-hCG	40.15 ng/mL	0.82	Method
			CRL Robinson
			Scan date
			25-01-2026
Risks at sampling date			Trisomy 21
Age risk	1:480		The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.
Biochemical T21 risk	1:1431		After the result of the Trisomy 21 test (with NT) it is expected that among 7784 women with the same data, there is one woman with a trisomy 21 pregnancy and 7783 women with not affected pregnancies.
Combined trisomy 21 risk	1:7784		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!
Trisomy 13/18 + NT	<1:10000		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

below cut off
 Below Cut Off, but above Age Risk
 above cut off