

Prisca 5.1.0.17
Date of report: 28/01/26

Patient data			
Name	Mrs. PAYAL YADAV	Patient ID	0852601230107
Birthday	20/04/98	Sample ID	b4072723
Age at sample date	27.8	Sample Date	23/01/26
Gestational age	12 + 2		
Correction factors			
Fetuses	1	IVF	no
Weight	72	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	2.77 mIU/mL	0.96	12 + 2
fb-hCG	40.2 ng/mL	0.96	Method
			CRL Robinson
			Scan date
			23/01/26
Risks at sampling date			Trisomy 21
Age risk	1:804		Crown rump length in mm
Biochemical T21 risk	1:5010		60.2
Combined trisomy 21 risk	<1:10000		Nuchal translucency MoM
Trisomy 13/18 + NT	<1:10000		0.77
			Nasal bone
			present
			Sonographer
			NA
			Qualifications in measuring NT
			Sonographer
			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 more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy. 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

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