

Prisca 5.1.0.17
Date of report: 30/09/25

N A

Patient data			
Name	Mrs. JYOTI KUMARI	Patient ID	0482509290183
Birthday	15/11/91	Sample ID	B3737145
Age at sample date	33.9	Sample Date	29/09/25
Gestational age	13 + 4		
Correction factors			
Fetuses	1	IVF	no
Weight	63	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	2.88 mIU/mL	0.52	
fb-hCG	32.52 ng/mL	1.04	
Risks at sampling date			
Age risk		1:357	
Biochemical T21 risk		1:430	
Combined trisomy 21 risk		1:2067	
Trisomy 13/18 + NT		<1:10000	
			Gestational age 13 + 3
			Method CRL Robinson
			Scan date 28/09/25
			Crown rump length in mm 76.7
			Nuchal translucency MoM 0.96
			Nasal bone present
			Sonographer N A
			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 2067 women with the same data, there is one woman with a trisomy 21 pregnancy and 2066 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

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