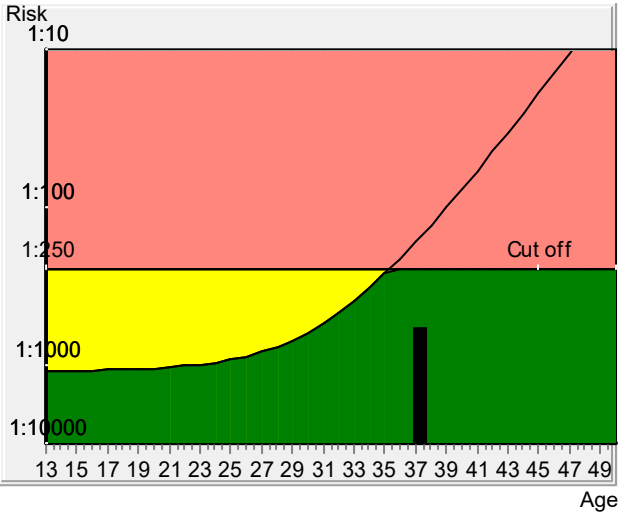


Prisca 5.1.0.17
Date of report: 07/10/25

NA

Patient data			
Name	Mrs. PRIYANKA SINGH	Patient ID	0482509270135
Birthday	11/06/88	Sample ID	B3675683
Age at sample date	37.3	Sample Date	27/09/25
Gestational age	11 + 5		
Correction factors			
Fetuses	1	IVF	no
Weight	66.6	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data			Ultrasound data
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	0.95 mIU/mL	0.39	11 + 5
fb-hCG	47.64 ng/mL	0.98	Method
			CRL Robinson
			Scan date
			27/09/25
			Crown rump length in mm
			51.5
			Nuchal translucency MoM
			0.72
			Nasal bone
			present
			Sonographer
			NA
			Qualifications in measuring NT
			NA
Risks at sampling date			Trisomy 21
Age risk		1:154	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 582 women with the same data, there is one woman with a trisomy 21 pregnancy and 581 women with not affected pregnancies. The PAPP-A level is low. 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!
Biochemical T21 risk		1:91	
Combined trisomy 21 risk		1:582	
Trisomy 13/18 + NT		1:8103	
			
Trisomy 13/18 + NT			
The calculated risk for Trisomy 13/18 (with nuchal translucency) is 1:8103, which represents a low risk.			

Sign of Physician

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