

Prisca 5.1.0.17
 Date of report: 23/11/25

Patient data			
Name	Mrs. PRANJUL	Patient ID	0772511210142
Birthday	05/04/00	Sample ID	B4273497
Age at sample date	25.6	Sample Date	21/11/25
Gestational age	13 + 3		
Correction factors			
Fetuses	1	IVF	no
Weight	52	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	3.67 mIU/mL	0.56	
fb-hCG	35.41 ng/mL	1.02	
Risks at sampling date			
Age risk		1:962	
Biochemical T21 risk		1:1444	
Combined trisomy 21 risk		1:3847	
Trisomy 13/18 + NT		<1:10000	
			Gestational age 12 + 6
			Method CRL Robinson
			Scan date 17/11/25
			Crown rump length in mm 67.6
			Nuchal translucency MoM 1.17
			Nasal bone unknown
			Sonographer NA
			Qualifications in measuring NT Sonographer
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 3847 women with the same data, there is one woman with a trisomy 21 pregnancy and 3846 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