

Patient data			
Name	MRS. SWAPNIL KHANDELWAL		Patient ID
Birthday	30/04/1992		Sample ID
Age at sample date	32.7		Sample Date
Gestational age	12 + 4		
Correction factors			
Fetuses	1	IVF	no
Weight	60.7	diabetes	no
Smoker	no	Origin	Asian
Previous trisomy 21 pregnancies		no	
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	7.1 mIU/ml	1.76	
fb-hCG	90.3 ng/ml	2.03	
Risks at sampling date			
Age risk	1:429		
Biochemical T21 risk	1:1435		
Combined trisomy 21 risk	1:6844		
Trisomy 13/18 + NT	<1:10000		
			Gestational age 12 + 4 Method CRL Robinson Scan date 18/01/2025 Crown rump length in mm 63.2 Nuchal translucency MoM 0.80 Nasal bone present Sonographer DR. (MRS.) NEERJA CHOPRA Qualifications in measuring NT MD
			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 6844 women with the same data, there is one woman with a trisomy 21 pregnancy and 6843 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