

Patient data			
Name	Mrs. SHARDA DANGE	Patient ID	0662506020095
Birthday	02-11-1999	Sample ID	B2795301
Age at sample date	25.6	Sample Date	02-06-2025
Gestational age	12 + 2		
Correction factors			
Fetuses	1	IVF	no
Weight	49	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	1.93 mIU/mL	0.42	12 + 1
fb-hCG	42.5 ng/mL	0.90	Method
Risks at sampling date		CRL Robinson	
Age risk	1:927	Scan date	
Biochemical T21 risk	1:887	01-06-2025	
Combined trisomy 21 risk	1:5426	Crown rump length in mm	
Trisomy 13/18 + NT	<1:10000	57.3	
		Nuchal translucency MoM	
		0.80	
		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 5426 women with the same data, there is one woman with a trisomy 21 pregnancy and 5425 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.			

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