

Prisca 5.1.0.17
Date of report: 15/09/25

NA

Patient data				
Name	Mrs. AMRUTA WADGHULE		Patient ID	0662509110141
Birthday	14/03/00		Sample ID	B3403095
Age at sample date	25.5		Sample Date	11/09/25
Gestational age	12 + 6			
Correction factors				
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies
Weight	75	diabetes	no	
Smoker	no	Origin	Asian	
Biochemical data			Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age	12 + 5
PAPP-A	1.89 mIU/mL	0.55	Method	CRL Robinson
fb-hCG	39.07 ng/mL	1.08	Scan date	10/09/25
Risks at sampling date			Crown rump length in mm	
Age risk	1:950		65.7	
Biochemical T21 risk	1:1185		Nuchal translucency MoM	
Combined trisomy 21 risk	1:6977		0.66	
Trisomy 13/18 + NT	<1:10000		Nasal bone	
			present	
			Sonographer	
			N A	
			Qualifications in measuring NT	
			MD	
Risk			Trisomy 21	
			<p>The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.</p> <p>After the result of the Trisomy 21 test (with NT) it is expected that among 6977 women with the same data, there is one woman with a trisomy 21 pregnancy and 6976 women with not affected pregnancies.</p> <p>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!</p> <p>The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)).</p> <p>The laboratory can not be hold responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!</p>	
Trisomy 13/18 + NT				
<p>The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.</p>				

Sign of Physician

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