

Prisca 5.1.0.17
Date of report: 22/11/25

Patient data								
Name		Mrs. SHWETA JAGTAP		Patient ID	0662511200055			
Birthday		18/11/95		Sample ID	B3728432			
Age at sample date		30.0		Sample Date	20/11/25			
Gestational age		12 + 5						
Correction factors								
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	unknown			
Weight	55	diabetes	no					
Smoker	no	Origin	Asian					
Biochemical data								
Parameter	Value	Corr. MoM	Ultrasound data					
PAPP-A	1.63 mIU/mL	0.34	Gestational age 12 + 2					
fb-hCG	40.01 ng/mL	0.97	Method CRL Robinson					
Risks at sampling date								
Age risk		1:650	Scan date 17/11/25					
Biochemical T21 risk		1:284	Crown rump length in mm 60.2					
Combined trisomy 21 risk		1:1868	Nuchal translucency MoM 0.70					
Trisomy 13/18 + NT		<1:10000	Nasal bone present					
			Sonographer NA					
			Qualifications in measuring NT Sonographer					
Trisomy 21								
The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk. <p>After the result of the Trisomy 21 test (with NT) it is expected that among 1868 women with the same data, there is one woman with a trisomy 21 pregnancy and 1867 women with not affected pregnancies.</p> <p>The PAPP-A level is low.</p> <p>The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician.</p> <p>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 held responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!</p>								
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