

Prisca 5.1.0.17
Date of report: 01/12/25

Patient data					
Name	Mrs. NEETA KAMBLE		Patient ID	0662511290167	
Birthday	03/04/87		Sample ID	B3947000	
Age at sample date	38.7		Sample Date	29/11/25	
Gestational age	13 + 0				
Correction factors					
Fetuses	1	IVF	yes	Previous trisomy 21 pregnancies	
Weight	66	diabetes	no		
Smoker	no	Origin	Asian		
Biochemical data			Ultrasound data		
Parameter	Value	Corr. MoM	Gestational age	12 + 6	
PAPP-A	2.44 mIU/mL	0.57	Method	CRL Robinson	
fb-hCG	36.97 ng/mL	1.02	Scan date	28/11/25	
Risks at sampling date			Crown rump length in mm		
Age risk	1:115		67		
Biochemical T21 risk	1:182		Nuchal translucency MoM		
Combined trisomy 21 risk	1:1052		0.71		
Trisomy 13/18 + NT	<1:10000		Nasal bone		
			present		
			Sonographer		
			NA		
			Qualifications in measuring NT		
			Sonographer		
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 1052 women with the same data, there is one woman with a trisomy 21 pregnancy and 1051 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 held 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