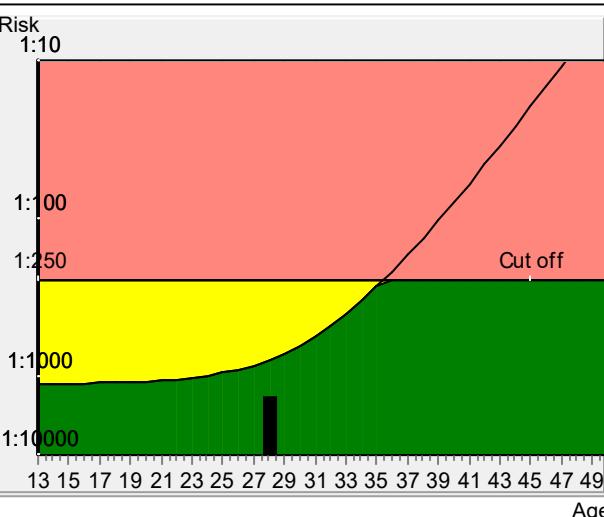


Prisca 5.1.0.17  
**Date of report:** 10-11-2025

NA

Patient data								
Name	Mrs. MANJU GUPTA	Patient ID	0872511060031					
Birthday	07-10-1997	Sample ID	A1961912					
Age at sample date	28.1	Sample Date	06-11-2025					
Gestational age	12 + 4							
Correction factors								
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	unknown			
Weight	47	diabetes	no					
Smoker	no	Origin	Asian					
Biochemical data								
Parameter	Value	Corr. MoM	Ultrasound data					
PAPP-A	1.97 mIU/mL	0.37	Gestational age 12 + 3					
fb-hCG	40.58 ng/mL	0.90	Method CRL Robinson					
Risks at sampling date								
Age risk	1:791		Scan date 05-11-2025					
Biochemical T21 risk	1:496		Crown rump length in mm 62.63					
Combined trisomy 21 risk	1:1361		Nuchal translucency MoM 1.18					
Trisomy 13/18 + NT	1:9592		Nasal bone present					
Sonographer NA								
Qualifications in measuring NT NA								
Trisomy 21								
<b>The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.</b> <p>After the result of the Trisomy 21 test (with NT) it is expected that among 1361 women with the same data, there is one woman with a trisomy 21 pregnancy and 1360 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								
<b>The calculated risk for Trisomy 13/18 (with nuchal translucency) is 1:9592, which represents a low risk.</b>								

Sign of Physician