

Prisca 5.1.0.17

Date of report: 08/04/25

N A

Patient data								
Name	Mrs. PRANJAL PAWAR	Patient ID	0662504070242					
Birthday	25/10/06	Sample ID	B2687778					
Age at sample date	18.5	Sample Date	07/04/25					
Gestational age	12 + 6							
Correction factors								
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	unknown			
Weight	49	diabetes	no					
Smoker	no	Origin	Asian					
Biochemical data								
Parameter	Value	Corr. MoM	Ultrasound data					
PAPP-A	3.09 mIU/mL	0.54	Gestational age 12 + 5					
fb-hCG	41.97 ng/mL	1.01	Method CRL Robinson					
Risks at sampling date								
Age risk	1:1117		Scan date 06/04/25					
Biochemical T21 risk	1:1564		Crown rump length in mm 65					
Combined trisomy 21 risk	1:9153		Nuchal translucency MoM 0.72					
Trisomy 13/18 + NT	<1:10000		Nasal bone unknown					
Sonographer N A								
Qualifications in measuring NT MD								
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 9153 women with the same data, there is one woman with a trisomy 21 pregnancy and 9152 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								
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