

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
Weight	75	diabetes	no
Smoker	no	Origin	Asian
Biochemical data			
Parameter	Value	Corr. MoM	Ultrasound data
PAPP-A	1.89 mIU/mL	0.55	Gestational age 12 + 5
fb-hCG	39.07 ng/mL	1.08	Method CRL Robinson
Risks at sampling date			
Age risk	1:950		Scan date 10/09/25
Biochemical T21 risk	1:1185		Crown rump length in mm 65.7
Combined trisomy 21 risk	1:6977		Nuchal translucency MoM 0.66
Trisomy 13/18 + NT	<1:10000		Nasal bone present
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.			
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.			
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!			
The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)).			
The laboratory can not be held responsible for their impact on the risk assessment! Calculated risks have no diagnostic value!			
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