

Patient data								
Name	Ms. SEEMA KUMARI	Patient ID	0472601150075					
Birthday	14-10-2002	Sample ID	B2490130					
Age at sample date	23.3	Sample Date	15-01-2026					
Gestational age	13 + 0							
Correction factors								
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	unknown			
Weight	41	diabetes	no					
Smoker	no	Origin	Asian					
Biochemical data								
Parameter	Value	Corr. MoM	Ultrasound data					
PAPP-A	1.71 mIU/mL	0.23	Gestational age 12 + 6					
fb-hCG	38 ng/mL	0.89	Method CRL Robinson					
Risks at sampling date								
Age risk	1:1040		Scan date 14-01-2026					
Biochemical T21 risk	1:162		Crown rump length in mm 68.4					
Combined trisomy 21 risk	1:736		Nuchal translucency MoM 1.04					
Trisomy 13/18 + NT	1:2418		Nasal bone unknown					
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 736 women with the same data, there is one woman with a trisomy 21 pregnancy and 735 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 1:2418, which represents a low risk.								

Sign of Physician