

Prisca 5.1.0.17
Date of report: 26/09/25

N A

Patient data			
Name	Mrs. KRAPA KEVAT	Patient ID	0442509250009
Birthday	30/09/06	Sample ID	B3476368
Age at sample date	19.0	Sample Date	23/09/25
Gestational age	12 + 0		
Correction factors			
Fetuses	1	IVF	no
Weight	41	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	2.39 mIU/mL	0.49	
fb-hCG	43.52 ng/mL	0.80	
Risks at sampling date			
Age risk	1:1080	Gestational age	12 + 0
Biochemical T21 risk	1:1898	Method	CRL Robinson
Combined trisomy 21 risk	1:6550	Scan date	23/09/25
Trisomy 13/18 + NT	<1:10000	Crown rump length in mm	55.4
		Nuchal translucency MoM	1.09
		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 6550 women with the same data, there is one woman with a trisomy 21 pregnancy and 6549 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 hold 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
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