

Prisca 5.1.0.17
Date of report: 13/03/25

N A

Patient data				
Name	Mrs. BINITA HAZARIKA		Patient ID	0692503100190
Birthday	01/09/87		Sample ID	B2519729
Age at sample date	37.5		Sample Date	10/03/25
Gestational age	12 + 6			
Correction factors				
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies
Weight	55	diabetes	no	
Smoker	no	Origin	Asian	
Biochemical data			Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age	12 + 6
PAPP-A	3.67 mIU/mL	0.73	Method	CRL Robinson
fb-hCG	29.76 ng/mL	0.75	Scan date	10/03/25
Risks at sampling date			Crown rump length in mm	
Age risk	1:152		67	
Biochemical T21 risk	1:881		Nuchal translucency MoM	
Combined trisomy 21 risk	1:4557		0.71	
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. After the result of the Trisomy 21 test (with NT) it is expected that among 4557 women with the same data, there is one woman with a trisomy 21 pregnancy and 4556 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!				
Risk 1:10 1:100 1:250 1:1000 1:10000 Age 13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49 Cut off				
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