

Prisca 5.1.0.17
Date of report: 15/12/25

Patient data			
Name	Mrs. JHORAMANI HATI	Patient ID	0652512120040
Birthday	24/03/94	Sample ID	b4056370
Age at sample date	31.7	Sample Date	12/12/25
Gestational age	13 + 1		
Correction factors			
Fetuses	1	IVF	no
Weight	40	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	7.66 mIU/mL	0.96	Gestational age 12 + 4
fb-hCG	36.95 ng/mL	0.89	Method CRL Robinson
Risks at sampling date			Scan date 08/12/25
Age risk	1:518		Crown rump length in mm 62.9
Biochemical T21 risk	1:3840		Nuchal translucency MoM 1.05
Combined trisomy 21 risk	<1:10000		Nasal bone present
Trisomy 13/18 + NT	<1:10000		Sonographer NA
			Qualifications in measuring NT Sonographer
Risk			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 more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy. 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