

Prisca 5.1.0.17
Date of report: 12-09-2024

N A

Patient data					
Name	Mrs. HUMAIRA SUMAIN		Patient ID	0372409050028	
Birthday	08-01-1998		Sample ID	A0428373	
Age at sample date	26.7		Sample Date	05-09-2024	
Gestational age	13 + 2				
Correction factors					
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	
Weight	67	diabetes	no		
Smoker	no	Origin	Asian		
Biochemical data			Ultrasound data		
Parameter	Value	Corr. MoM	Gestational age	13 + 1	
PAPP-A	6.31 mIU/mL	1.36	Method	CRL Robinson	
fb-hCG	58.36 ng/mL	1.75	Scan date	04-09-2024	
Risks at sampling date			Crown rump length in mm		
Age risk	1:902		71		
Biochemical T21 risk	1:2756		Nuchal translucency MoM		
Combined trisomy 21 risk	1:8720		1.07		
Trisomy 13/18 + NT	<1:10000		Nasal bone		
			unknown		
			Sonographer		
			N A		
			Qualifications in measuring NT		
			MD		
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 8720 women with the same data, there is one woman with a trisomy 21 pregnancy and 8719 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