

Prisca 5.2.0.13  
Date of report: 12/02/22

Patient data			
Name	FIROJA KHATUN	Patient ID	
Birthday	20/10/93	Sample ID	10128638
Age at sample date	28.3	Sample Date	12/02/22
Gestational age	12 + 4		
Correction factors			
Fetuses	1	IVF	unknown
Weight	75	diabetes	unknown
Smoker	unknown	Origin	Asian
		Previous trisomy 21	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	2.98 mIU/mL	0.82	
fb-hCG	27.74 ng/mL	0.84	
Risks at sampling date			Gestational age
Age risk	1:775		12 + 2
Biochemical T21 risk	1:4669		Method
Combined trisomy 21 risk	1:5951		CRL Robinson
Trisomy 13/18 + NT	<1:10000		Scan date
			10/02/22
			Crown rump length in mm
			58.9
			Nuchal translucency MoM
			1.37
			Nasal bone
			unknown
			Sonographer
			DR.SUKANTA GHOSAL.
			Qualifications in measuring NT
			M.D.
Risk			Trisomy 21
			<p><b>The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.</b></p> <p>After the result of the Trisomy 21 test (with NT) it is expected that among 5951 women with the same data, there is one woman with a trisomy 21 pregnancy and 5950 women with not affected pregnancies.</p> <p>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!</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 hold responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!</p>
Trisomy 13/18 + NT			
<p><b>The calculated risk for trisomy 13/18 (with nuchal translucency) is &lt; 1:10000, which represents a low risk.</b></p>			

Sign of Physician