

Prisca 5.2.0.13
Date of report: 27/10/21

Patient data			
Name	SABANA KHATOON	Patient ID	
Birthday	04/03/86	Sample ID	10025587
Age at sample date	35.6	Sample Date	23/10/21
Gestational age	12 + 3		
Correction factors			
Fetuses	1	IVF	unknown
Weight	66.9	diabetes	no
Smoker	no	Origin	Asian
Previous trisomy 21	unknown		
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	2.37 mIU/mL	0.60	
fb-hCG	19.38 ng/mL	0.54	
Risks at sampling date			Gestational age
Age risk	1:235		12 + 2
Biochemical T21 risk	1:1610		Method
Combined trisomy 21 risk	1:8364		CRL Robinson
Trisomy 13/18 + NT	<1:10000		Scan date
			22/10/21
			Crown rump length in mm
			60.3
			Nuchal translucency MoM
			0.70
			Nasal bone
			unknown
			Sonographer
			DR.RANJAN SUKLA GHOSH.
			Qualifications in measuring NT
			MBBS
Risk			Trisomy 21
			<p>The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.</p> <p>After the result of the Trisomy 21 test (with NT) it is expected that among 8364 women with the same data, there is one woman with a trisomy 21 pregnancy and 8363 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>The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.</p>			

Sign of Physician