

Prisca 5.1.0.17  
Date of report: 12/12/24

NA

Patient data				
Name	Mrs. POOJA BHARGAVA		Patient ID	0442412110008
Birthday	10/11/86		Sample ID	a1851959
Age at sample date	38.1		Sample Date	10/12/24
Gestational age	12 + 1			
Correction factors				
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies
Weight	77	diabetes	no	
Smoker	no	Origin	Asian	
Biochemical data			Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age	12 + 1
PAPP-A	1.48 mIU/mL	0.59	Method	CRL Robinson
fb-hCG	43.58 ng/mL	1.03	Scan date	10/12/24
Risks at sampling date			Crown rump length in mm	
Age risk	1:128		58.6	
Biochemical T21 risk	1:220		Nuchal translucency MoM	
Combined trisomy 21 risk	1:700		1.11	
Trisomy 13/18 + NT	<1:10000		Nasal bone	
			present	
			Sonographer	
			NA	
			Qualifications in measuring NT	
			MD	
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 700 women with the same data, there is one woman with a trisomy 21 pregnancy and 699 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