

Prisca 5.1.0.17  
Date of report: 01/10/24

N A

Patient data				
Name	Mrs. PRIYANKA CHOUDHARI		Patient ID	0712409300006
Birthday	14/08/97		Sample ID	A0976711
Age at sample date	27.1		Sample Date	30/09/24
Gestational age	12 + 6			
Correction factors				
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies
Weight	45	diabetes	no	
Smoker	no	Origin	Asian	
Biochemical data			Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age	12 + 5
PAPP-A	14.2 mIU/mL	2.26	Method	CRL Robinson
fb-hCG	77.73 ng/mL	1.82	Scan date	29/09/24
Risks at sampling date			Crown rump length in mm	
Age risk	1:861		66	
Biochemical T21 risk	1:4709		Nuchal translucency MoM	
Combined trisomy 21 risk	<1:10000		1.07	
Trisomy 13/18 + NT	<1:10000		Nasal bone	
			present	
			Sonographer	
			N A	
			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 more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy.</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