

Prisca 5.2.0.13

Date of report: 10/06/2024

Patient data			
Name	MRS. GUNJAN	Patient ID	
Birthday	16/09/1996	Sample ID	2406016733/NOD
Age at sample date	27.7	Sample Date	10/06/2024
Gestational age	13 + 3		
Correction factors			
Fetuses	1	IVF	no
Weight	63.7	diabetes	no
Smoker	no	Origin	Asian
Previous trisomy 21 pregnancies			no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	7.5 mIU/ml	1.40	13 + 1
fb-hCG	30.1 ng/ml	0.76	Method
			CRL Robinson
			Scan date
			8/06/2024
Risks at sampling date		Crown rump length in mm	
Age risk	1:839	71	
Biochemical T21 risk	<1:10000	Nuchal translucency MoM	
Combined trisomy 21 risk	<1:10000	0.73	
Trisomy 13/18 + NT	<1:10000	Nasal bone	
		present	
		Sonographer	
		DR. NEERJA CHOPRA	
		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 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

below cut off	Below Cut Off, but above Age Risk	above cut off
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