

# JITM DIAGNOSTIC

Prisca

5.2.0.13

Date of report:

9/11/2024

Patient data			
Name	MRS. SHAGUN		Patient ID
Birthday	18/11/1996		Sample ID
Age at sample date	28.0		Sample Date
Gestational age	12 + 1		
Correction factors			
Fetuses	1	IVF	no
Weight	54.6	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	1.72 mIU/ml	0.46	Gestational age
fb-hCG	28.9 ng/ml	0.60	Method
			CRL Robinson
			Scan date
			7/11/2024
			Crown rump length in mm
			56.3
			Nuchal translucency MoM
			0.81
			Nasal bone
			present
			Sonographer
			DR. (MRS.) NEERJA CHOPRA
			Qualifications in measuring NT
			MD
Risks at sampling date			
Age risk		1:786	
Biochemical T21 risk		1:2134	
Combined trisomy 21 risk		<1:10000	
Trisomy 13/18 + NT		<1:10000	
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 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>		<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