

Prisca 5.2.0.13
Date of report: 8/09/2024

Patient data			
Name	MRS. MEENU SHARMA	Patient ID	
Birthday	25/07/1985	Sample ID	2409220225/AMB
Age at sample date	39.1	Sample Date	7/09/2024
Gestational age	13 + 1		
Correction factors			
Fetuses	1	IVF	no
Weight	64	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	2.61 mIU/ml	0.55	Gestational age
fb-hCG	50.6 ng/ml	1.24	Method
			Scan date
Risks at sampling date			Crown rump length in mm
Age risk		1:102	Nuchal translucency MoM
Biochemical T21 risk		1:94	Nasal bone
Combined trisomy 21 risk		1:425	Sonographer
Trisomy 13/18 + NT		<1:10000	Qualifications in measuring NT
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 425 women with the same data, there is one woman with a trisomy 21 pregnancy and 424 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

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