

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
			CRL Robinson
			Scan date
			3/09/2024
Risks at sampling date			Trisomy 21
Age risk		1:102	The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.
Biochemical T21 risk		1:94	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.
Combined trisomy 21 risk		1:425	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!
Trisomy 13/18 + NT		<1:10000	The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)).
			The laboratory can not be hold responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!
Trisomy 13/18 + NT			
The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.			

Sign of Physician

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