

Prisca

5.2.0.13

Date of report: 10/09/2024

Patient data			
Name	MRS. MEENU	Patient ID	
Birthday	6/08/1993	Sample ID	2409018806/NOD
Age at sample date	31.1	Sample Date	9/09/2024
Gestational age	12 + 5		
Correction factors			
Fetuses	1	IVF	no
Weight	76.5	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	2.69 mIU/ml	0.83	Gestational age 12 + 4
fb-hCG	55.6 ng/ml	1.36	Method CRL Robinson
Risks at sampling date			Scan date 8/09/2024
Age risk		1:561	Crown rump length in mm 64.5
Biochemical T21 risk		1:1140	Nuchal translucency MoM 0.97
Combined trisomy 21 risk		1:4990	Nasal bone present
Trisomy 13/18 + NT		<1:10000	Sonographer DR. RASHMI BANSAL
			Qualifications in measuring NT M.D
		Trisomy 21 The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk. After the result of the Trisomy 21 test (with NT) it is expected that among 4990 women with the same data, there is one woman with a trisomy 21 pregnancy and 4989 women with not affected pregnancies. 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! 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