

JITM Diagnostics

Patient data			
Name	MRS. ARTI KUMARI	Patient ID	
Birthday	10/03/1989	Sample ID	2410011049/NOD
Age at sample date	35.6	Sample Date	6/10/2024
Gestational age	12 + 2		
Correction factors			
Fetuses	1	IVF	no
Weight	50	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	2.48 mIU/ml	0.56	12 + 1
fb-hCG	42.1 ng/ml	0.86	Method
			CRL Robinson
			Scan date
			5/10/2024
Risks at sampling date			Crown rump length in mm
Age risk		1:237	58.7
Biochemical T21 risk		1:517	Nuchal translucency MoM
Combined trisomy 21 risk		1:2766	0.88
Trisomy 13/18 + NT		<1:10000	Nasal bone
			present
			Sonographer
			DR. VIBHA 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 2766 women with the same data, there is one woman with a trisomy 21 pregnancy and 2765 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