

JITM Diagnostics

Patient data			
Name	MRS. DEEPIKA	Patient ID	
Birthday	29/10/1989	Sample ID	2412050533/NOD
Age at sample date	35.2	Sample Date	31/12/2024
Gestational age	12 + 2		
Correction factors			
Fetuses	1	IVF	no
Weight	98	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	1.4 mlU/ml	0.72	12 + 1
fb-hCG	44.1 ng/ml	1.10	Method
			CRL Robinson
			Scan date
			30/12/2024
Risks at sampling date			Crown rump length in mm
Age risk		1:259	58
Biochemical T21 risk		1:618	Nuchal translucency MoM
Combined trisomy 21 risk		1:3397	0.66
Trisomy 13/18 + NT		<1:10000	Nasal bone
			present
			Sonographer
			DR. AMIT RAI
			Qualifications in measuring NT
			MD
			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 3397 women with the same data, there is one woman with a trisomy 21 pregnancy and 3396 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