

JITM Diagnostics

Patient data			
Name	MRS. KUMARI NIDHI	Patient ID	
Birthday	1/01/1993	Sample ID	2410032631/NOD
Age at sample date	31.8	Sample Date	17/10/2024
Gestational age	13 + 3		
Correction factors			
Fetuses	1	IVF	no
Weight	61.5	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	6.9 mIU/ml	1.24	13 + 3
fb-hCG	47.1 ng/ml	1.17	Method
			CRL Robinson
Risks at sampling date			Scan date
Age risk		1:517	17/10/2024
Biochemical T21 risk		1:3460	Crown rump length in mm
Combined trisomy 21 risk		<1:10000	75.1
Trisomy 13/18 + NT		<1:10000	Nuchal translucency MoM
			0.70
			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 more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy. 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