JITM Diagnostics

Patient data				
Name	MRS. PRIYANKA			
Birthday	15/07/91	Sample ID		2505034338/NOD
Age at sample date	33.8	Sample Date		18/05/25
Gestational age	12 + 5			
Correction factors				
Fetuses 1	IVF	no	Previous trisomy 21	no
Weight 71.2	diabetes	no	pregnancies	
Smoker no	Origin	Asian		
Biochemical data		Ultrasound da	ata	
Parameter Value	Corr. MoM	Gestational age		12 + 4
PAPP-A 1.85 mIU/m	0.52	Method		CRL Robinson
fb-hCG 55.3 ng/ml	1.33	Scan date		17/05/25
Risks at sampling date		Crown rump length in mm		63.9
Age risk	1:349	Nuchal translucency MoM		0.73
Biochemical T21 risk	1:243			present
Combined trisomy 21 risk	1:1488			DR. NEERJA CHOPRA
Trisomy 13/18 + NT	<1:10000	Qualifications in measuring NT Trisomy 21		M.D
1:100 1:250 1:1000 1:10000 13 15 17 19 21 23 25 27 29 31 33 3 Trisomy 13/18 + NT The calculated risk for trisomy 13 translucency) is < 1:10000, which risk.	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 1488 women with the same data, there is one woman with a trisomy 21 pregnancy and 1487 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!			