JITM Diagnostics

Patient data			
Name	MRS. NEHA		
Birthday	01/01/96	Sample ID	2501040238/NOD
Age at sample date	29.1	Sample Date	e 25/01/25
Gestational age	12 + 4		
Correction factors			
Fetuses 1	IVF	no	ı ,
Weight 58	diabetes	no	pregnancies
Smoker no	Origin	Asian	
Biochemical data		Ultrasound da	ata
Parameter Value	Corr. MoM	Gestational	age 12 + 4
PAPP-A 4.6 mIU/m	I 1.08	Method	CRL Robinson
fb-hCG 87.5 ng/ml	1.94	Scan date	25/01/25
Risks at sampling date		Crown rump length in mm 63.6	
Age risk	1:720	Nuchal translucency MoM 1.47	
Biochemical T21 risk	1:1090	[
Combined trisomy 21 risk	-		er DR. SUSHUM KUMAR VERMA
Trisomy 13/18 + NT	<1:10000	Qualification	ns in measuring NT M.D
KISK 1:10		Trisomy 21	ated risk for Trisomy 21 (with nuchal
1:100 1:1000 1:10000 13 15 17 19 21 23 25 27 29 31 33 38 Trisomy 13/18 + NT The calculated risk for trisomy 13 translucency) is < 1:10000, which risk.	Age	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 824 women with the same data, there is one woman with a trisomy 21 pregnancy and 823 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!	

Sign of Physician