JITM Diagnostics

Patient data						
Name	MRS. NITESH DAHIYA			Patient ID		
Birthday	02/08/95		Sample ID		2505000072/NOD	
Age at sample date	29.7			Sample Date)	30/04/25
Gestational age	11 + 4					
Correction factors						
Fetuses	1	IVF		no	Previous trisomy 21	no
Weight	64.5	diabetes		no	pregnancies	
Smoker	no	Origin		Asian		
Biochemical data				Ultrasound da	nta	
Parameter	Value		Corr. MoM	Gestational age		11 + 4
PAPP-A	2.6 mIU/m	I	1.11	Method		CRL Robinson
fb-hCG	95.1 ng/ml		1.95			30/04/25
Risks at sampling date				Crown rump length in mm 50		
Age risk			1:642	Nuchal translucency MoM		1.33
Biochemical T21 risk	1:1022		Nasal bone		present	
Combined trisomy 21 ri	abined trisomy 21 risk 1:1333			Sonographer DR. ANKIT KHANDELWAL MBBS DNB		
Trisomy 13/18 + NT			<1:10000		s in measuring NT	MD
1:100 1:250 1:1000 1:10000 13 15 17 19 21 23 25	1:100 1:250 Cut off			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 1333 women with the same data, there is one woman with a trisomy 21 pregnancy and 1332 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		/40 (with m	-	1		

The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low

risk.