## JITM Diagnostics

Patient data				
Name	MRS. APARNA			
Birthday	03/08/97	Sample ID 2505000082/NOD		
Age at sample date	27.7	Sample Date		01/05/25
Gestational age	12 + 1			
Correction factors				
Fetuses 1	IVF	no	Previous trisomy 21	no
Weight 70	diabetes	no	pregnancies	
Smoker no	Origin	Asian		
Biochemical data	U		Ultrasound data	
Parameter Value	Corr. MoM	Corr. MoM Gestational age		12 + 0
PAPP-A 8.7 mIU/m	I 3.10	Method CRL Robinson		
fb-hCG 41.1 ng/ml	0.92	Scan date 30/04/25		
sks at sampling date		Crown rump length in mm 56.		
Age risk	1:801	Nuchal trans	slucency MoM	0.60
Biochemical T21 risk	<1:10000	Nasal bone		present
Combined trisomy 21 risk	omy 21 risk <1:10000		r	DR. ANMOL NIGAM
Trisomy 13/18 + NT	<1:10000	Qualification	s in measuring NT	M.D
KISK 1:10		Trisomy 21	ited risk for Trisomy 2	
1:100 1:250 1:1000 1:10000 13 15 17 19 21 23 25 27 29 31 33 33 Trisomy 13/18 + NT The calculated risk for trisomy 13 translucency) is < 1:10000, which risk.	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 PAPP-A level is high.  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!			