JITM Diagnostics

Patient data			
Name	MRS. AMRITA	Patient ID	
Birthday	23/04/1992	Sample ID	2412035313/NOD
Age at sample date	32.7	Sample Date	20/12/2024
Gestational age	13 + 2		

Correction factors					
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	no
Weight	72.5	diabetes	no		
Smoker	no	Origin	Asian		

Biochemical data			Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age	13 + 2
PAPP-A	6.8 mIU/ml	1.57	Method	CRL Robinson
fb-hCG	40.1 ng/ml	1.03	Scan date	20/12/2024

Risks at sampling date			
Age risk	1:445	Crown rump length in mm	73
Biochemical T21 risk	1:6244	Nuchal translucency MoM	0.88
Combined trisomy 21 risk	<1:10000	Nasal bone	present
Trisomy 13/18 + NT	<1:10000	Sonographer	DR. RAJINDER NANDA
		Qualifications in measuring NT	M.D

Risk

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

	below cut off		Below Cut Off, but above Age Risk		above cut off