

JITM Diagnostics

Patient data			
Name	MRS. VARSHA SHARMA	Patient ID	
Birthdate	29/09/1992	Sample ID	2410022374/NOD
Age at sample date	32.0	Sample Date	12/10/2024
Gestational age	13 + 1		
Correction factors			
Fetuses	1	IVF	no
Weight	74	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	1.51 mIU/ml	0.38	Gestational age 13 + 0
fb-hCG	62.8 ng/ml	1.60	Method CRL Robinson
Risks at sampling date			Scan date 11/10/2024
Age risk		1:492	Crown rump length in mm 69.9
Biochemical T21 risk		1:90	Nuchal translucency MoM 0.68
Combined trisomy 21 risk		1:616	Nasal bone present
Trisomy 13/18 + NT		<1:10000	Sonographer DR. ASHISH GARG
			Qualifications in measuring NT M.D
Risk			Trisomy 21
1:10			<p>The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.</p> <p>After the result of the Trisomy 21 test (with NT) it is expected that among 616 women with the same data, there is one woman with a trisomy 21 pregnancy and 615 women with not affected pregnancies.</p> <p>The PAPP-A level is low.</p> <p>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!</p> <p>The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)).</p> <p>The laboratory can not be hold responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!</p>
1:100			
1:250			
1:1000			
1:10000			
	Age		
Trisomy 13/18 + NT			
<p>The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.</p>			

Sign of Physician