

Prisca

5.2.0.13

Date of report: 16/01/2025

JITM Diagnostics

Patient data			
Name	MRS. SUMAN	Patient ID	
Birthday	1/01/2002	Sample ID	2501023280/NOD
Age at sample date	23.0	Sample Date	16/01/2025
Gestational age	12 + 0		
Correction factors			
Fetuses	1	IVF	no
Weight	59	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	2.28 mIU/ml	0.71	11 + 6
fb-hCG	30.1 ng/ml	0.63	Method
			CRL Robinson
			Scan date
			15/01/2025
Risks at sampling date		Crown rump length in mm	
Age risk	1:1010	55	
Biochemical T21 risk	1:7758	Nuchal translucency MoM	
Combined trisomy 21 risk	<1:10000	0.69	
Trisomy 13/18 + NT	<1:10000	Nasal bone	
		present	
		Sonographer	
		DR. PRERNA AGARWAL	
		Qualifications in measuring NT	
		M.D	
Risk		Trisomy 21	
		<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 more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy.</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>	
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