Prisca 5.2.0.13

Date of report: 16/01/2025

JITM Diagnostics

Patient data				
Name	MRS. SUMAN			
Birthday	1/01/2002			2501023280/NOD
Age at sample date	23.0)	16/01/2025
Gestational age	12 + 0			
Correction factors				
Fetuses 1	IVF	no	Previous trisomy 21	no
Weight 59	diabetes	no	pregnancies	
Smoker no	Origin	Asian		
Biochemical data		Ultrasound da	ata	
Parameter Value	Corr. MoM	Gestational age 11 + 6		
PAPP-A 2.28 mIU/n	nl 0.71	Method CRL Robinson		
fb-hCG 30.1 ng/ml	0.63	Scan date		15/01/2025
Risks at sampling date		Crown rump	length in mm	55
Age risk	sk 1:1010		slucency MoM	0.69
Biochemical T21 risk	1:7758	Nasal bone		present
Combined trisomy 21 risk <1:10000		Sonographe		DR. PRERNA AGARWAL
Trisomy 13/18 + NT	<1:10000		s in measuring NT	M.D
Risk 1:10 1:1000 1:10000 1:10000 1:10000 1:10000 Trisomy 13/18 + NT The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low 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!		