Prisca 5.2.0.13

Date of report: 17/03/25

JITM Diagnostics

Patient data			
Name	MRS. PARUL P		
Birthday	16/04/94	Sample ID 2503032397/NOD	
Age at sample date	30.9	Sample Date 17/03/25	
Gestational age	10 + 4		
Correction factors			
Fetuses 1	IVF	no Previous trisomy 21 no	
Weight 74	diabetes	no pregnancies	
Smoker no	Origin	Asian	
		Ultrasound data	
Parameter Value	Corr. MoM	Gestational age 10 + 4	
PAPP-A 4.31 mIU/m	I 3.87	Method CRL Robinson	
fb-hCG 73.1 ng/ml	1.39		
Risks at sampling date		Crown rump length in mm 39.5	
Age risk	1:527	Nuchal translucency MoM 1.78	
Biochemical T21 risk	1:5584	Nasal bone present Sonographer DR. B. CHAUHAN MBBS MD	
Combined trisomy 21 risk			
Trisomy 13/18 + NT	<1:10000 Qualifications in measuring NT MD		
Risk 1:10		Trisomy 21	ated risk for Trisomy 21 (with nuchal
1:1000 1:10000 1:1100000 1:1100000 1:1100000 1:11000000 1:1100000000		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 1190 women with the same data, there is one woman with a trisomy 21 pregnancy and 1189 women with not affected pregnancies. 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!	