Prisca 5.2.0.13

Date of report: 9/03/2025

JITM Diagnostics

Patient data MRS. PRIYANKA Patient ID				
Name	1/03/1994		Sample ID	
Birthday				
Age at sample date	31.0	Sample Date		9/03/2025
Gestational age	12 + 5			
Correction factors				
Fetuses 1	IVF	no	Previous trisomy 21	no
Weight 69	diabetes	no	pregnancies	
Smoker no	Origin	Asian		
Biochemical data		Ultrasound da	ıta	
Parameter Value	Corr. MoM	Gestational age 12 + 4		
PAPP-A 0.78 mIU/m	0.21	Method CRL Robinson		
fb-hCG 28.1 ng/ml	0.67	Scan date 8/03/2		8/03/2025
Risks at sampling date	. •		Crown rump length in mm	
Age risk	1:567	Nuchal translucency MoM		0.80
Biochemical T21 risk	1:119	Nasal bone		present
Combined trisomy 21 risk	ned trisomy 21 risk 1:843		Sonographer	
Trisomy 13/18 + NT	1:992 Qualifications in measuring NT			
1:100 1:250 1:1000 1:10000 13 15 17 19 21 23 25 27 29 31 33 39 Trisomy 13/18 + NT The calculated risk for Trisomy 1 translucency) is 1:992, which reports	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 843 women with the same data, there is one woman with a trisomy 21 pregnancy and 842 women with not affected pregnancies. The PAPP-A level is low. 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!			

Sign of Physician