Prisca 5.2.0.13

Date of report: 27/12/2024

Patient data				
Name MRS. TAF	e MRS. TARA JAIPAL 10306578			
Birthday	1/01/1996		Sample ID 2412044917/NOD	
Age at sample date 29.0		Sample Date 26/12/2024		
Gestational age 13 + 3				
Correction factors				
Fetuses 1	IVF	no	Previous trisomy 21	l no
Weight 42.7	diabetes	no	pregnancies	
Smoker no	Origin	Asian		
Biochemical data		Ultrasound da	ata	
Parameter Value	Corr. MoM	Gestational age 13		13 + 0
PAPP-A 11.2 mIU/m	1.32	Method CRL Robin		CRL Robinson
fb-hCG 37.9 ng/ml	0.83	Scan date		23/12/2024
Risks at sampling date		Crown rump length in mm		70.5
Age risk	1:747	Nuchal translucency MoM		0.91
Biochemical T21 risk	<1:10000	Nasal bone		present
Combined trisomy 21 risk <1:10000		Sonographer DR. VINEET NAVANI MD		
Trisomy 13/18 + NT <1:10000		Qualifications in measuring NT MD Trisomy 21		
1:100 1:250 1:1000 1:10000 1:3 15 17 19 21 23 25 27 29 31 33 3 Trisomy 13/18 + NT The calculated risk for trisomy 13 translucency) is < 1:10000, which risk.	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!			

Sign of Physician