Prisca 5.2.0.13

Date of report: 6/11/2024

JITM Diagnostics

Patient data				
Name	MRS. SAKSHI			
Birthday	13/10/1997	Sample ID		2411009705/NOD
Age at sample date	27.1	Sample Date		6/11/2024
Gestational age	13 + 2			
Correction factors				
Fetuses 1	IVF	no	Previous trisomy 21	no
Weight 69.5	diabetes	no pregnancies		
Smoker no	Origin	Asian		
Biochemical data		Ultrasound data		
Parameter Value	Corr. MoM	Gestational age		13 + 1
PAPP-A 3.1 mIU/m	0.68	Method CRL Robinsor		
fb-hCG 24.1 ng/ml	0.61	Scan date 5/11/2024		
Risks at sampling date		Crown rump length in mm		72.6
Age risk	1:878	Nuchal translucency MoM		0.66
Biochemical T21 risk	1:6365	Nasal bone		present
Combined trisomy 21 risk	21 risk <1:10000		er	DR. NEERJA CHOPRA
Trisomy 13/18 + NT	<1:10000	Qualifications in measuring NT		M.D
Risk 1:10	1	Trisomy 21		
1:1000 1:10000 1:10000 13 15 17 19 21 23 25 27 29 31 33 33 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!			