Prisca 5.2.0.13

Date of report: 17/11/2024

Patient data					
Name	MRS. SHIVANI	Patient ID			
Birthday	5/04/2001	Sample ID 2411030971/NOD			
Age at sample date	23.6	Sample Date	e	16/11/2024	
Gestational age	12 + 0				
Correction factors					
Fetuses 1	IVF	no	Previous trisomy 21	no	
Weight 36	diabetes	no	pregnancies		
Smoker no	Origin	Asian			
Biochemical data	cal data		Ultrasound data		
Parameter Value	Corr. MoM	Gestational	age	11 + 5	
PAPP-A 8.6 mIU/m	1.53	Method		CRL Robinson	
fb-hCG 124.6 ng/ml	2.15	Scan date		14/11/2024	
Risks at sampling date	•		length in mm	53	
Age risk	1:993	Nuchal translucency MoM		0.78	
Biochemical T21 risk	1:2246			present	
Combined trisomy 21 risk				DR. AMIT KUMAR	
Trisomy 13/18 + NT	<1:10000 C		is in measuring NT	M.D	
1:10 1:250 1:1000 1:10000 1:10000 13 15 17 19 21 23 25 27 29 31 33 35 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