Prisca 5.2.0.13

Date of report: 08/11/24

Patient data				
Name	MRS. HEENA	Patient ID		
Birthday	11/11/87	Sample ID	2411013656/NOD	
Age at sample date	37.0	Sample Date	e 07/11/24	
Gestational age	13 + 2			
Correction factors				
Fetuses 1	IVF	no	Previous trisomy 21 no	
Weight 62	diabetes	no	pregnancies	
Smoker no	Origin	Asian		
Biochemical data	The state of the s		Ultrasound data	
Parameter Value	Value Corr. MoM Gestational		age 13 + 1	
PAPP-A 11.4 mIU/m	1 2.18	Method	CRL Robinson	
fb-hCG 62.8 ng/ml	1.54	Scan date	06/11/24	
Risks at sampling date			Crown rump length in mm 71	
Age risk	1:176	Nuchal translucency MoM 0.79		
Biochemical T21 risk	1:1447	Nasal bone	•	
·		Sonographer		
Trisomy 13/18 + NT			Qualifications in measuring NT	
Risk 1:10		Trisomy 21	ated risk for Trisomy 21 (with nuchal	
1:1000 1:250 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.	Age /18 (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 6466 women with the same data, there is one woman with a trisomy 21 pregnancy and 6465 women with not affected pregnancies. 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