Date of report:	23/09/2024
Prisca	5.2.0.13

Patient data				
Name	MRS. SHALU	Patient ID		
Birthday	28/01/1988	Sample ID	2409047788/NOI	
Age at sample date	36.7	Sample Date	e 23/09/2024	
Gestational age	12 + 6			
Correction factors				
Fetuses 1	IVF	no Previous trisomy 21 no		
Weight 57	diabetes	no	pregnancies	
Smoker no	Origin	Asian		
Biochemical data		Ultrasound da	ata	
Parameter Value	Corr. MoM	Gestational age 12 + 5		
PAPP-A 3.7 mlU/m	nl 0.76	Method	CRL Robinso	
fb-hCG 20.1 ng/ml	0.46	Scan date	22/09/202	
Risks at sampling date		Crown rump length in mm 65		
Age risk	1:188		Nuchal translucency MoM 0.78	
Biochemical T21 risk			F	
Combined trisomy 21 risk	-		Sonographer	
Trisomy 13/18 + NT	<1:10000	Qualifications in measuring NT		
Risk 1:10		Trisomy 21	ated risk for Trisomy 21 (with nuchal	
1: 00 1: 250 Cut off 1: 1000 1: 100		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!		
The calculated risk for trisomy 13 translucency) is < 1:10000, which risk.				

Sign of Physician