Prisca 5.1.0.17

Date of report: 26/12/22

Patient data					
Name Mrs	Mrs.ARJUMAND RAZVI		Patient ID		
Birthday	15/04/93	Sample ID		4053638	
Age at delivery	30.2	Sample Date	e	26/12/22	
Gestational age	13 + 3				
Correction factors					
Fetuses 1	IVF no Previous trisomy 21		unknown		
Weight 69.9	diabetes no pregnancies				
Smoker no	Origin Asian				
Biochemical data Ultrason			ata		
Parameter Value	Corr. MoM Gestational age 12 + 4		12 + 4		
PAPP-A 10 mIU/mI	2.51	Method CRL Robinson			
fb-hCG 38.8 ng/ml	1.35	Scan date 20/12/22			
Risks at term			length in mm	64	
Age risk	1:942	Nuchal translucency MoM		0.61	
Biochemical T21 risk	<1:10000			unknown	
Combined trisomy 21 risk	<1:10000	Sonographe	er	Dr.Rahul	
Trisomy 13/18 + NT	<1:10000	Qualifications in measuring NT FM		FMF	
TAIOK			Trisomy 21 The calculated risk for Trisomy 21 (with nuchal		
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.	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 PAPP-A level is high.  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