Prisca 5.2.0.13

Date of report: 13/12/2024

	DIMPLE	D-4:4 ID	
Birthday 19/		Patient ID	
	/03/1997	Sample ID	2412021882/NOD
Age at sample date	27.7	Sample Date	e 12/12/2024
Gestational age	13 + 1		
Correction factors			
Fetuses 1 IVF		no	Previous trisomy 21 no
Weight 58.6 diabetes		no	pregnancies
Smoker no Origin		Asian	
Biochemical data		Ultrasound da	ata
Parameter Value C	Corr. MoM	Gestational	age 13 + 1
PAPP-A 5.3 mIU/ml	1.00	Method	CRL Robinson
fb-hCG 16.8 ng/ml	0.40	Scan date 12/12/2024	
Risks at sampling date	Cro		o length in mm 72.2
Age risk	1:831	Nuchal translucency MoM 0.72	
Biochemical T21 risk	<1:10000	Nasal bone present	
Combined trisomy 21 risk <			er DR. (MRS.) NEERJA CHOPRA
Trisomy 13/18 + NT			ns in measuring NT MD
1:1000 1:250 Cut off 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:100000 1:100000 1:100000 1:100000 1:100000000		Trisomy 21 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 free beta HCG level is low. 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!	