Patient data						
Name	MRS. MOHINI GUPTA			Patient ID		
Birthday	4/06/1996			Sample ID		2406034072/NOD
Age at sample date	28.0			Sample Date		18/06/2024
Gestational age	12 + 5					
Correction factors						
Fetuses	1 IVF		no	Previous trisomy 21	no	
Weight	52	diabetes		no	pregnancies	
Smoker	no	Origin		Asian		
Biochemical data				Ultrasound data		
Parameter	Value		Corr. MoM	Gestational age		12 + 5
PAPP-A	1.7 mIU/ml		0.33	Method		CRL Robinson
fb-hCG	137 ng/ml		2.97	Scan date		18/06/2024
Risks at sampling date			Crown rump length in mm		65.1	
Age risk	1:798			Nuchal translucency MoM		0.59
Biochemical T21 risk	>1:50			Nasal bone		present
Combined trisomy 21 risk	bined trisomy 21 risk 1:164			Sonographer .		
Trisomy 13/18 + NT <1:10000			Qualifications in measuring NT			
Risk				Trisomy 21		
1:10				The calculated risk for Trisomy 21 (with nuchal		

1:1000 1:250 Cut off 1:10000 1:110000 1:100000 1:100

The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.

The calculated risk for Trisomy 21 (with nuchal translucency) is above the cut off, which indicates an increased risk.

After the result of the Trisomy 21 test (with NT) it is expected that among 164 women with the same data, there is one woman with a trisomy 21 pregnancy and 163 women with not affected pregnancies.

The free beta HCG level is high.

The PAPP-A 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!

Sign of Physician