## Sarita Vihar

Prisca 5.2.0.13

Date of report: 30/07/2024

Patient data				
Name	MRS. PINKI	Patient ID		
Birthday	30/01/1986		2407221285/AMB	
Age at sample date	38.5		e 27/07/2024	
Gestational age	12 + 0			
Correction factors				
Fetuses 1	IVF	yes	Previous trisomy 21 no	
Weight 64.1	diabetes	no	pregnancies	
Smoker no	Origin	Asian		
Biochemical data	Ultra		Ultrasound data	
Parameter Value	Corr. MoM	Gestational	age 12 + 0	
PAPP-A 7.45 mIU/m	l 2.56	Method	CRL Robinson	
fb-hCG 102 ng/ml	2.19	Scan date	27/07/2024	
Risks at sampling date		Crown rump length in mm 56.5		
Age risk	1:115	Nuchal translucency MoM 1.81		
Biochemical T21 risk	1:391	Nasal bone present		
Combined trisomy 21 risk	ned trisomy 21 risk 1:64		Sonographer	
Trisomy 13/18 + NT	<1:10000 Qualifications in measuring NT			
TAISIA			Trisomy 21	
1:1000 1:250			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 64 women with the same data, there is one woman with a trisomy 21 pregnancy and 63 women with not affected pregnancies.  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!	