

P.PRATHIBHA

Patient data					
Name	Mrs. G.SWATHI	Patient ID	0082203160017		
Birthday	19-05-1997	Sample ID	23019196		
Age at sample date	24.8	Sample Date	16-03-2022		
Gestational age	12 + 5				
Correction factors					
Fetuses	1	IVF	no		
Weight	56	diabetes	no		
Smoker	no	Origin	Asian		
Biochemical data		Ultrasound data			
Parameter	Value	Corr. MoM			
PAPP-A	2.69 mIU/mL	0.58	Gestational age		
fb-hCG	46.57 ng/mL	1.14	Method		
Risks at sampling date		CRL Robinson			
Age risk	1:975	Scan date			
Biochemical T21 risk	1:1262	16-03-2022			
Combined trisomy 21 risk	1:6075	Crown rump length in mm			
Trisomy 13/18 + NT	<1:10000	66			
Risk		Nuchal translucency MoM			
1:10		0.95			
1:100		Nasal bone			
1:250		unknown			
1:1000		Sonographer			
1:10000		DR NA			
13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49		Qualifications in measuring NT			
Age		MD			
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 6075 women with the same data, there is one woman with a trisomy 21 pregnancy and 6074 women with not affected pregnancies.					
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 held responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!					
Trisomy 13/18 + NT					
The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.					

Sign of Physician

 below cut off

 Below Cut Off, but above Age Risk

 above cut off