

RAJINI MBBS.,DGO.,

Patient data			
Name	Mrs. FIRDOSE FETUS-B	Patient ID	0162105180003
Birthday	05-01-1997	Sample ID	21110652
Age at sample date	24.4	Sample Date	18-05-2021
Gestational age	12 + 4		
Correction factors			
Fetuses	2	IVF	no
Weight	79	diabetes	no
Smoker	no	Origin	Asian
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	6.87 mIU/mL	1.28	Gestational age
fb-hCG	54.25 ng/mL	0.66	Method
Risks at sampling date		CRL Robinson	
Age risk	1:988	Scan date	17-05-2021
Biochemical T21 risk	<1:10000	Crown rump length in mm	62
Combined trisomy 21 risk	<1:10000	Nuchal translucency MoM	0.75
Trisomy 13/18 + NT	<1:10000	Nasal bone	unknown
		Sonographer	DR NA
		Qualifications in measuring NT	MD
Risk		Trisomy 21	
1:10		The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.	
1:100		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.	
1:250		The risk for this twin pregnancy has been calculated for a singleton pregnancy with corrected MoMs.	
1:1000		The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician.	
1:10000		Please note that risk calculations are statistical approaches and have no diagnostic value!	
13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49		The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)).	
Age		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