

SOWMYA REDDY

Patient data			
Name	Mrs. N. SANDHYA	Patient ID	0012102100246
Birthday	13/06/94	Sample ID	21007792
Age at sample date	26.7	Sample Date	10/02/21
Gestational age	11 + 1		
Correction factors			
Fetuses	1	IVF	no
Weight	68	diabetes	no
Smoker	no	Origin	Asian
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	3.67 mIU/mL	2.02	Gestational age
fb-hCG	63.41 ng/mL	1.18	Method CRL Robinson
Risks at sampling date		Scan date	
Age risk	1:833	10/02/21	
Biochemical T21 risk	<1:10000	Crown rump length in mm	
Combined trisomy 21 risk	<1:10000	45.3	
Trisomy 13/18 + NT	<1:10000	Nuchal translucency MoM	
Risk		0.80	
1:10		Nasal bone	
1:00		present	
1:250		Sonographer	
1:1000		NA	
1:10000		Qualifications in measuring NT	
13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49		NA	
Age			
Trisomy 13/18 + NT		Trisomy 21	
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 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 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!	

Sign of Physician

below cut off

Below Cut Off, but above Age Risk

above cut off