

Patient data			
Name	MRS. NIRMLA DEVI	Patient ID	1804220211/AMB
Birthday	01-01-1984	Sample ID	1804220211/AMB
Age at sample date	34.3	Sample Date	09-04-2018
Gestational age	12 + 5		
Correction factors			
Fetuses	1	IVF	no
Weight	62	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	1.75 mIU/ml	0.49	12 + 3
fb-hCG	51.5 ng/ml	1.52	Method
			CRL Robinson
			Scan date
			07-04-2018
			Crown rump length in mm
			61.1
			Nuchal translucency MoM
			1.20
			Nasal bone
			present
			Sonographer
			.
			Qualifications in measuring NT
			MD
Risks at sampling date		Trisomy 21	
Age risk	1:320	The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.	
Biochemical T21 risk	1:134	After the result of the Trisomy 21 test (with NT) it is expected that among 321 women with the same data, there is one woman with a trisomy 21 pregnancy and 320 women with not affected pregnancies.	
Combined trisomy 21 risk	1:321	The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician.	
Trisomy 13/18 + NT	<1:10000	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!	
Risk	1:10		
	1:100		
	1:250		
	1:1000		
	1:10000		
		Age	
		13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49	
		Cutoff	
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