

Patient data			
Name	MRS. VANDANA	Patient ID	101901030020
Birthday	08-10-1996	Sample ID	101901030020
Age at sample date	27.0	Sample Date	03-01-2019
Gestational age	11 + 4		
Correction factors			
Fetuses	1	IVF	no
Weight	44	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	2.92 mIU/ml	1.24	11 + 4
fb-hCG	36.6 ng/ml	0.84	Method
			LMP
			Scan date
			Crown rump length in mm
			Nuchal translucency MoM
			Nasal bone
			present
			Sonographer
			.
			Qualifications in measuring NT
			MD
Risks at sampling date		Trisomy 21	
Age risk	1:824	The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.	
Biochemical T21 risk	<1:10000	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.	
Combined trisomy 21 risk	<1:10000	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

