

Dr.PADMA,M.B.B.S.,M.S

Patient data			
Name	Mrs. SRAVANI		Patient ID
Birth day	01/08/96	Sample ID	21000451
Age at sample date	24.6	Sample Date	20/02/21
Gestational age	13 + 2		
Correction factors			
Fetuses	1	IVF	no
Weight	65.4	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	6.98 mIU/mL	1.46	Method
fb-hCG	48.25 ng/mL	1.44	CRL Robinson
Risks at sampling date			Scan date
Age risk	1:1004		20/02/21
Biochemical T21 risk	1:5695		Crown rump length in mm
Combined trisomy 21 risk	<1:10000		73.2
Trisomy 13/18 + NT	<1:10000		Nuchal translucency MoM
			0.77
			Nasal bone
			present
			Sonographer
			DR NA
			Qualifications in measuring NT
			MD
Risk		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 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 hold 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