

K.NAGESWARI RAO

Patient data			
Name	Mrs. JYOTHI		Patient ID
Birth day	28-04-1989		Sample ID
Age at sample date	32.1		Sample Date
Gestational age	11 + 4		
Correction factors			
Fetuses	1	IVF	no
Weight	57.7	diabetes	no
Smoker	no	Origin	Asian
			Previous trisomy 21 pregnancies
			unknown
Biochemical data			Ultrasound data
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	1.98 mIU/mL	0.73	11 + 4
fb-hCG	51.23 ng/mL	0.98	Method
			CRL Robinson
			Scan date
			31-05-2021
			Crown rump length in mm
			51
			Nuchal translucency MoM
			1.09
			Nasal bone
			unknown
			Sonographer
			DR NA
			Qualifications in measuring NT
			MD
Risks at sampling date			Trisomy 21
Age risk	1:460		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 4653 women with the same data, there is one woman with a trisomy 21 pregnancy and 4652 women with not affected pregnancies. 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!
Biochemical T21 risk	1:1439		
Combined trisomy 21 risk	1:4653		
Trisomy 13/18 + NT	<1:10000		
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