

KRANTHI DIAGNOSTICS

Patient data			
Name	Mrs. VEERAMMA		Patient ID
Birth day	20-05-1996		Sample ID
Age at sample date	26.7		Sample Date
Gestational age	13 + 3		
Correction factors			
Fetuses	1	IVF	no
Weight	47	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	3.27 mIU/mL	0.44	13 + 0
fb-hCG	33.87 ng/mL	0.94	Method
			CRL Robinson
			Scan date
			10-02-2023
Risks at sampling date			Crown rump length in mm
Age risk			69
Biochemical T21 risk			Nuchal translucency MoM
1:874			0.81
Combined trisomy 21 risk			Nasal bone
1:5309			present
Trisomy 13/18 + NT			Sonographer
<1:10000			KRANTHI DIAGNOSTICS
			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 5309 women with the same data, there is one woman with a trisomy 21 pregnancy and 5308 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!	
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