

Prisca 5.1.0.17
Date of report: 16-01-2025

KRANTHI DIAGNOSTICS

Patient data			
Name	Mrs. SUPRIYA		Patient ID
Birth day	06-01-1998		Sample ID
Age at sample date	27.0		Sample Date
Gestational age	13 + 6		
Correction factors			
Fetuses	1	IVF	no
Weight	63	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	5.22 mIU/mL	0.86	13 + 3
fb-hCG	22.29 ng/mL	0.78	Method
			CRL Robinson
			Scan date
			13-01-2025
Risks at sampling date			Crown rump length in mm
Age risk			77
Biochemical T21 risk			Nuchal translucency MoM
1:896			0.96
Combined trisomy 21 risk			Nasal bone
1:6920			present
Trisomy 13/18 + NT			Sonographer
<1:10000			N A
<1:10000			Qualifications in measuring NT
			MD
Risk			Trisomy 21
1:10			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!
1:100			
1:250			
1:1000			
1:10000			
Cut off			
Age			
13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49			
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