

N A

Patient data			
Name	Mrs. P JYOTHI	Patient ID	0012504290313
Birthday	24/07/98	Sample ID	B2528330
Age at sample date	26.8	Sample Date	29/04/25
Gestational age	13 + 6		
Correction factors			
Fetuses	1	IVF	no
Weight	54	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	3.39 mIU/mL	0.47	Gestational age 13 + 2
fb-hCG	28.96 ng/mL	0.96	Method CRL Robinson
Risks at sampling date			Scan date 25/04/25
Age risk		1:912	Crown rump length in mm 75
Biochemical T21 risk		1:965	Nuchal translucency MoM 0.76
Combined trisomy 21 risk		1:5835	Nasal bone present
Trisomy 13/18 + NT		<1:10000	Sonographer N A
			Qualifications in measuring NT MD
Risk			Trisomy 21
			<p>The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.</p> <p>After the result of the Trisomy 21 test (with NT) it is expected that among 5835 women with the same data, there is one woman with a trisomy 21 pregnancy and 5834 women with not affected pregnancies.</p> <p>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!</p> <p>The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)).</p> <p>The laboratory can not be hold responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!</p>
Trisomy 13/18 + NT			
<p>The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.</p>			

Sign of Physician