

Patient data					
Name	Mrs. K SHRUTHI		Patient ID	0012207250449	
Birthday	19/06/93		Sample ID	23567370	
Age at sample date	29.1		Sample Date	25/07/22	
Gestational age	12 + 3				
Correction factors					
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	unknown
Weight	76	diabetes	no		
Smoker	no	Origin	Asian		
Biochemical data			Ultrasound data		
Parameter	Value	Corr. MoM	Gestational age	12 + 3	
PAPP-A	1.89 mIU/mL	0.66	Method	CRL Robinson	
fb-hCG	42.32 ng/mL	1.06	Scan date	25/07/22	
Risks at sampling date			Crown rump length in mm	62.4	
Age risk	1:714		Nuchal translucency MoM	0.66	
Biochemical T21 risk	1:1491		Nasal bone	unknown	
Combined trisomy 21 risk	1:8336		Sonographer	NA	
Trisomy 13/18 + NT	<1:10000		Qualifications in measuring NT	NAGA SAI TEJA	
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 8336 women with the same data, there is one woman with a trisomy 21 pregnancy and 8335 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