

Prisca 5.1.0.17  
Date of report: 17/10/25

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Patient data			
Name	Mrs. P KAVITHA		Patient ID
Birth day	05/06/92		Sample ID
Age at sample date	33.4		Sample Date
Gestational age	13 + 4		
Correction factors			
Fetuses	1	IVF	no
Weight	52	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	5.3 mIU/mL	0.77	Method
fb-hCG	32.62 ng/mL	0.98	Scan date
Risks at sampling date			
Age risk	1:394		Crown rump length in mm
Biochemical T21 risk	1:1421		Nuchal translucency MoM
Combined trisomy 21 risk	1:7551		Nasal bone
Trisomy 13/18 + NT	<1:10000		Sonographer
			Qualifications in measuring NT
Risk			Trisomy 21
			<p><b>The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.</b></p> <p>After the result of the Trisomy 21 test (with NT) it is expected that among 7551 women with the same data, there is one woman with a trisomy 21 pregnancy and 7550 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><b>The calculated risk for trisomy 13/18 (with nuchal translucency) is &lt; 1:10000, which represents a low risk.</b></p>			

Sign of Physician