

Prisca 5.1.0.17
Date of report: 17/10/25

N A

Patient data					
Name		Mrs. P KAVITHA		Patient ID	0012510170126
Birthday		05/06/92		Sample ID	A1908040
Age at sample date		33.4		Sample Date	17/10/25
Gestational age		13 + 4			
Correction factors					
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	unknown
Weight	52	diabetes	no		
Smoker	no	Origin	Asian		
Biochemical data					
Parameter	Value	Corr. MoM	Ultrasound data		
PAPP-A	5.3 mIU/mL	0.77	Gestational age 13 + 3		
fb-hCG	32.62 ng/mL	0.98	Method CRL Robinson		
Risks at sampling date					
Age risk		1:394	Scan date 16/10/25		
Biochemical T21 risk		1:1421	Crown rump length in mm 76.6		
Combined trisomy 21 risk		1:7551	Nuchal translucency MoM 0.64		
Trisomy 13/18 + NT		<1:10000	Nasal bone present		
Risk 1:10			Sonographer N A		
1:100			Qualifications in measuring NT MD		
1:250			Trisomy 21		
1:1000			The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.		
1:10000			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.		
13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49		Age	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!		
Trisomy 13/18 + NT			The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)).		
The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.			The laboratory can not be held responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!		

Sign of Physician

 below cut off

 Below Cut Off, but above Age Risk

 above cut off