

Prisca 5.1.0.17
Date of report: 16-02-2025

N A

Patient data					
Name	Mrs. A SRAVANI TWIN A		Patient ID	0012502150393	
Birthday	18-03-1988		Sample ID	A1749112	
Age at sample date	36.9		Sample Date	15-02-2025	
Gestational age	11 + 5				
Correction factors					
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	
Weight	71	diabetes	no		
Smoker	no	Origin	Asian		
Biochemical data			Ultrasound data		
Parameter	Value	Corr. MoM	Gestational age	11 + 5	
PAPP-A	2.26 mIU/mL	0.99	Method	CRL Robinson	
fb-hCG	39.05 ng/mL	0.82	Scan date	15-02-2025	
Risks at sampling date			Crown rump length in mm		
Age risk	1:169		53		
Biochemical T21 risk	1:1593		Nuchal translucency MoM		
Combined trisomy 21 risk	1:6309		0.99		
Trisomy 13/18 + NT	<1:10000		Nasal bone		
			present		
			Sonographer		
			N A		
			Qualifications in measuring NT		
			MD		
Risk			Trisomy 21		
			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 6309 women with the same data, there is one woman with a trisomy 21 pregnancy and 6308 women with not affected pregnancies. 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!		
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
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Prisca 5.1.0.17
Date of report: 16-02-2025

N A

Patient data				
Name	Mrs. A SRAVANI TWIN B		Patient ID	0012502150393
Birthday	18-03-1988		Sample ID	A1749112
Age at sample date	36.9		Sample Date	15-02-2025
Gestational age	12 + 0			
Correction factors				
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies
Weight	71	diabetes	no	
Smoker	no	Origin	Asian	
Biochemical data			Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age	12 + 0
PAPP-A	2.26 mIU/mL	0.87	Method	CRL Robinson
fb-hCG	39.05 ng/mL	0.87	Scan date	15-02-2025
Risks at sampling date			Crown rump length in mm	
Age risk	1:171		56	
Biochemical T21 risk	1:1062		Nuchal translucency MoM	
Combined trisomy 21 risk	1:4082		1.02	
Trisomy 13/18 + NT	<1:10000		Nasal bone	
			present	
			Sonographer	
			N A	
			Qualifications in measuring NT	
			MD	
Trisomy 21				
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 4082 women with the same data, there is one woman with a trisomy 21 pregnancy and 4081 women with not affected pregnancies. 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 held responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!				
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
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