

Prisca 5.1.0.17  
Date of report: 02/12/25

Patient data			
Name Mrs. P.KRISHNA VENI TWIN A		Patient ID 0042512020002	
Birthday 02/07/07		Sample ID B3981594 TWIN A	
Age at sample date 18.4		Sample Date 01/12/25	
Gestational age 11 + 4			
Correction factors			
Fetuses 2	IVF no	Previous trisomy 21 pregnancies unknown	
Weight 54	diabetes no		
Smoker no	Origin Asian		
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age 11 + 4
PAPP-A	4.3 mIU/mL	0.78	Method CRL Robinson
fb-hCG	47.85 ng/mL	0.41	Scan date 01/12/25
Risks at sampling date			Crown rump length in mm 51.2
Age risk 1:1068			Nuchal translucency MoM 0.58
Biochemical T21 risk <1:10000			Nasal bone present
Combined trisomy 21 risk <1:10000			Sonographer NA
Trisomy 13/18 + NT <1:10000			Qualifications in measuring NT Sonographer
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 more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy.</p> <p>The risk for this twin pregnancy has been calculated for a singleton pregnancy with corrected MoMs.</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

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Date of report: 02/12/25

Patient data			
Name Mrs. P.KRISHNA VENI TWIN B		Patient ID 0042512020002	
Birthday 02/07/07		Sample ID B3981594 TWIN B	
Age at sample date 18.4		Sample Date 01/12/25	
Gestational age 11 + 3			
Correction factors			
Fetuses 2	IVF no	Previous trisomy 21 pregnancies unknown	
Weight 54	diabetes no		
Smoker no	Origin Asian		
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age 11 + 3
PAPP-A	4.3 mIU/mL	0.84	Method CRL Robinson
fb-hCG	47.85 ng/mL	0.40	Scan date 01/12/25
Risks at sampling date			Crown rump length in mm 48.5
Age risk 1:1062			Nuchal translucency MoM 0.68
Biochemical T21 risk <1:10000			Nasal bone present
Combined trisomy 21 risk <1:10000			Sonographer NA
Trisomy 13/18 + NT <1:10000			Qualifications in measuring NT Sonographer
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 more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy.</p> <p>The risk for this twin pregnancy has been calculated for a singleton pregnancy with corrected MoMs.</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

below cut off
  Below Cut Off, but above Age Risk
  above cut off