

Prisca 5.1.0.17
Date of report: 16-10-2024

N A

Patient data				
Name	Mrs. NIKITA NIRAJ VEER TWIN A		Patient ID	0372410140135
Birthday	04-07-1998		Sample ID	A1028335 TWIN A
Age at sample date	26.3		Sample Date	14-10-2024
Gestational age	13 + 1			
Correction factors				
Fetuses	2	IVF	no	Previous trisomy 21 pregnancies
Weight	62	diabetes	no	
Smoker	no	Origin	Asian	
Biochemical data			Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age	12 + 5
PAPP-A	12.1 mIU/mL	1.35	Method	CRL Robinson
fb-hCG	34.41 ng/mL	0.45	Scan date	11-10-2024
Risks at sampling date			Crown rump length in mm	
Age risk	1:919		65	
Biochemical T21 risk	<1:10000		Nuchal translucency MoM	
Combined trisomy 21 risk	<1:10000		0.91	
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 more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy. The risk for this twin pregnancy has been calculated for a singleton pregnancy with corrected MoMs. 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

Prisca 5.1.0.17
Date of report: 16-10-2024

N A

Patient data			
Name Mrs. NIKITA NIRAJ VEER TWIN B		Patient ID 0372410140135	
Birthday 04-07-1998		Sample ID A1028335 TWIN B	
Age at sample date 26.3		Sample Date 14-10-2024	
Gestational age 13 + 4			
Correction factors			
Fetuses 2	IVF no	Previous trisomy 21 pregnancies unknown	
Weight 62	diabetes no		
Smoker no	Origin Asian		
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age 13 + 1
PAPP-A	12.1 mIU/mL	1.16	Method CRL Robinson
fb-hCG	34.41 ng/mL	0.51	Scan date 11-10-2024
Risks at sampling date			Crown rump length in mm 71
Age risk 1:932			Nuchal translucency MoM 0.90
Biochemical T21 risk <1:10000			Nasal bone present
Combined trisomy 21 risk <1:10000			Sonographer N A
Trisomy 13/18 + NT <1:10000			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 more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy. The risk for this twin pregnancy has been calculated for a singleton pregnancy with corrected MoMs. 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