

Prisca 5.1.0.17
Date of report: 17-05-2025

N A

Patient data			
Name		W/O VIKAS NEGI FETUS-A	
Patient ID		0692505160140	
Birthday		30-03-1991	
Sample ID		B2734627 FETUS-A	
Age at sample date		34.1	
Sample Date		16-05-2025	
Gestational age		13 + 3	
Correction factors			
Fetuses	2	IVF	no
Weight	61	diabetes	no
Smoker	no	Origin	Asian
Previous trisomy 21 pregnancies		unknown	
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	11.51 mIU/mL	1.13	13 + 3
fb-hCG	31.8 ng/mL	0.45	Method
			CRL Robinson
			Scan date
			16-05-2025
Risks at sampling date			
Age risk			1:338
Biochemical T21 risk			<1:10000
Combined trisomy 21 risk			<1:10000
Trisomy 13/18 + NT			<1:10000
			Crown rump length in mm
			76.5
			Nuchal translucency MoM
			0.80
			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 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

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