

Prisca 5.1.0.17
Date of report: 13-01-2025

NA

Patient data				
Name	Mrs. SNEHAL NILESH JAGTAP FETUS-B		Patient ID	0662501120080
Birthday	12-07-1999		Sample ID	A0918479 FETUS-B
Age at sample date	25.5		Sample Date	12-01-2025
Gestational age	12 + 6			
Correction factors				
Fetuses	2	IVF	no	Previous trisomy 21 pregnancies
Weight	68	diabetes	no	
Smoker	no	Origin	Asian	
Biochemical data			Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age	12 + 5
PAPP-A	3.57 mIU/mL	0.49	Method	CRL Robinson
fb-hCG	59.36 ng/mL	0.74	Scan date	11-01-2025
Risks at sampling date			Crown rump length in mm	
Age risk	1:950		65	
Biochemical T21 risk	1:2061		Nuchal translucency MoM	
Combined trisomy 21 risk	1:4998		1.20	
Trisomy 13/18 + NT	<1:10000		Nasal bone	
			unknown	
			Sonographer	
			NA	
			Qualifications in measuring NT	
			MD	
Risk			Trisomy 21	
			<p>The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.</p> <p>After the result of the Trisomy 21 test (with NT) it is expected that among 4998 women with the same data, there is one woman with a trisomy 21 pregnancy and 4997 women with not affected pregnancies.</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>The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.</p>				

Sign of Physician