

Prisca 5.1.0.17
Date of report: 24/12/24

NA

Patient data			
Name		W/O DERHASAT NARZARY FETUS-B	
Patient ID		0692412230046	
Birthday		30/06/95	
Sample ID		A1760113 FETUS-B	
Age at sample date		29.5	
Sample Date		23/12/24	
Gestational age		13 + 2	
Correction factors			
Fetuses	2	IVF	no
Weight	56	diabetes	no
Smoker	no	Origin	Asian
Previous trisomy 21 pregnancies		unknown	
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	4.55 mIU/mL	0.43	
fb-hCG	19.8 ng/mL	0.26	
Gestational age		13 + 2	
Method		CRL Robinson	
Scan date		23/12/24	
Crown rump length in mm		73.9	
Nuchal translucency MoM		0.79	
Nasal bone		present	
Sonographer		NA	
Qualifications in measuring NT		MD	
Risks at sampling date		Trisomy 21	
Age risk	1:705		
Biochemical T21 risk	1:4815		
Combined trisomy 21 risk	<1:10000		
Trisomy 13/18 + NT	1:3496		
		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 free beta HCG level is low. 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 1:3496, which represents a low risk.	

Sign of Physician