

Prisca 5.1.0.17
Date of report: 06/09/24

N A

Patient data			
Name		Mrs. RASHMI PATORIYA	
Patient ID		0672409050077	
Birthday		01/06/93	
Sample ID		A1182682	
Age at sample date		31.3	
Sample Date		05/09/24	
Gestational age		13 + 4	
Correction factors			
Fetuses	1	IVF	no
Weight	65	diabetes	no
Smoker	no	Origin	Asian
Previous trisomy 21 pregnancies		unknown	
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	16.48 mIU/mL	3.10	
fb-hCG	46.37 ng/mL	1.50	
Gestational age		13 + 3	
Method		CRL Robinson	
Scan date		04/09/24	
Crown rump length in mm		77	
Nuchal translucency MoM		1.06	
Nasal bone		present	
Sonographer		N A	
Qualifications in measuring NT		MD	
Risks at sampling date		Trisomy 21	
Age risk	1:564	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 PAPP-A level is high. 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!	
Biochemical T21 risk	1:4973		
Combined trisomy 21 risk	<1:10000		
Trisomy 13/18 + NT	<1:10000		
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