

Prisca 5.2.0.13
Date of report: 12/01/22

Patient data						
Name	SUSMITA ADHIKARI	Patient ID				
Birthday	23/10/98	Sample ID	00097747			
Age at sample date	23.2	Sample Date	10/01/22			
Gestational age	11 + 2					
Correction factors						
Fetuses	1	IVF	unknown	Previous trisomy 21		
Weight	58	diabetes	unknown	unknown		
Smoker	unknown	Origin	Asian			
Biochemical data						
Parameter	Value	Corr. MoM				
PAPP-A	0.82 mIU/mL	0.29				
fb-hCG	23.68 ng/mL	0.48				
Risks at sampling date						
Age risk	1:978	1:978				
Biochemical T21 risk	1:1011	1:1011				
Combined trisomy 21 risk	1:5974	1:5974				
Trisomy 13/18 + NT	1:3182	1:3182				
Ultrasound data						
Gestational age			11 + 1			
Method			CRL Robinson			
Scan date			09/01/22			
Crown rump length in mm			45.5			
Nuchal translucency MoM			0.88			
Nasal bone			unknown			
Sonographer			DR.KUNAL SEN.			
Qualifications in measuring NT			MBBS			
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 5974 women with the same data, there is one woman with a trisomy 21 pregnancy and 5973 women with not affected pregnancies.						
The PAPP-A level is low.						
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 held 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 1:3182, which represents a low risk.						

Sign of Physician

below cut off

Below Cut Off, but above Age Risk

above cut off