

Prisca 5.2.0.13
Date of report: 27/10/21

Patient data							
Name	SABANA KHATOON	Patient ID					
Birthday	04/03/86	Sample ID	10025587				
Age at sample date	35.6	Sample Date	23/10/21				
Gestational age	12 + 3						
Correction factors							
Fetuses	1	IVF	unknown	Previous trisomy 21			
Weight	66.9	diabetes	no	unknown			
Smoker	no	Origin	Asian				
Biochemical data		Ultrasound data					
Parameter	Value	Corr. MoM	Gestational age	12 + 2			
PAPP-A	2.37 mIU/mL	0.60	Method	CRL Robinson			
fb-hCG	19.38 ng/mL	0.54	Scan date	22/10/21			
Risks at sampling date							
Age risk	1:235						
Biochemical T21 risk	1:1610						
Combined trisomy 21 risk	1:8364						
Trisomy 13/18 + NT	<1:10000						
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 8364 women with the same data, there is one woman with a trisomy 21 pregnancy and 8363 women with not affected pregnancies.							
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 < 1:10000, which represents a low risk.							

Sign of Physician

below cut off

Below Cut Off, but above Age Risk

above cut off