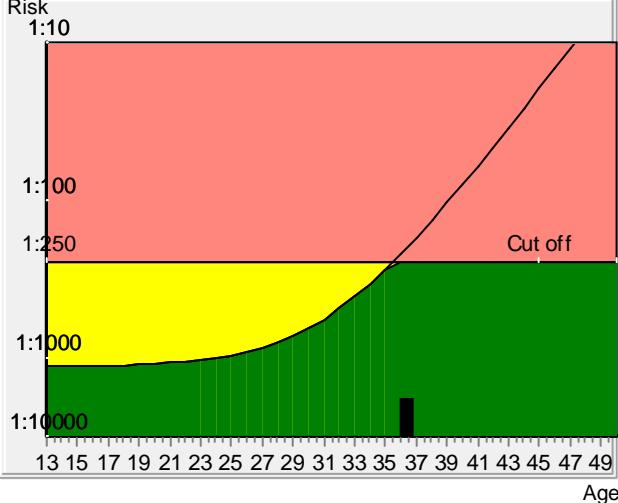


Prisca 5.2.0.13
Date of report: 31/10/21

Patient data					
Name	MUNNA DALAL DATTĀ	Patient ID			
Birthday	23/06/85	Sample ID	10032480		
Age at sample date	36.4	Sample Date	29/10/21		
Gestational age	12 + 6				
Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight	69	diabetes	no		
Smoker	no	Origin	Asian		
Biochemical data					
Parameter	Value	Corr. MoM			
PAPP-A	6.43 mIU/mL	1.44			
fb-hCG	22.08 ng/mL	0.70			
Risks at sampling date					
Age risk		1:202			
Biochemical T21 risk		1:5712			
Combined trisomy 21 risk		<1:10000			
Trisomy 13/18 + NT		<1:10000			
Ultrasound data					
Gestational age			11 + 1		
Method			CRL Robinson		
Scan date			17/10/21		
Crown rump length in mm			46.1		
Nuchal translucency MoM			0.87		
Nasal bone			unknown		
Sonographer			DR.B.SAHA.		
Qualifications in measuring NT			MD.		
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 more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy.					
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