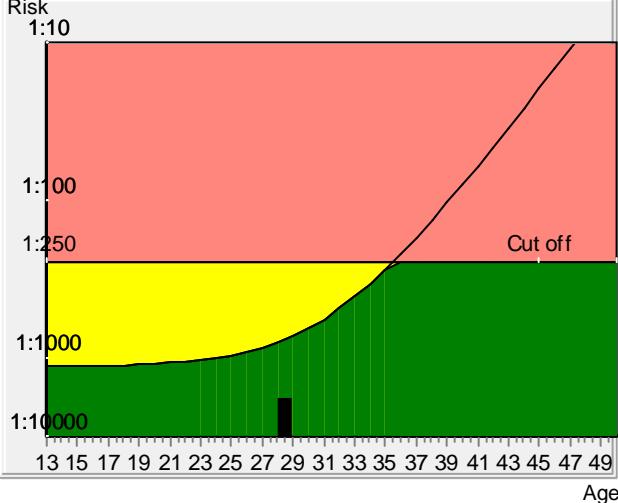


Prisca 5.2.0.13
Date of report: 01/12/21

Patient data					
Name	HANUFA BIBI	Patient ID			
Birthday	05/07/93	Sample ID	10061028		
Age at sample date	28.4	Sample Date	29/11/21		
Gestational age	12 + 6				
Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight	50	diabetes	no		
Smoker	no	Origin	Asian		
Biochemical data					
Parameter	Value	Corr. MoM			
PAPP-A	7.63 mIU/mL	1.17			
fb-hCG	22.53 ng/mL	0.64			
Risks at sampling date					
Age risk		1:776			
Biochemical T21 risk		<1:10000			
Combined trisomy 21 risk		<1:10000			
Trisomy 13/18 + NT		<1:10000			
Ultrasound data					
			Gestational age	12 + 6	
			Method	CRL Robinson	
			Scan date	29/11/21	
			Crown rump length in mm	67	
			Nuchal translucency MoM	1.18	
			Nasal bone	unknown	
			Sonographer	DR.PIYALI.DAS	
			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 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