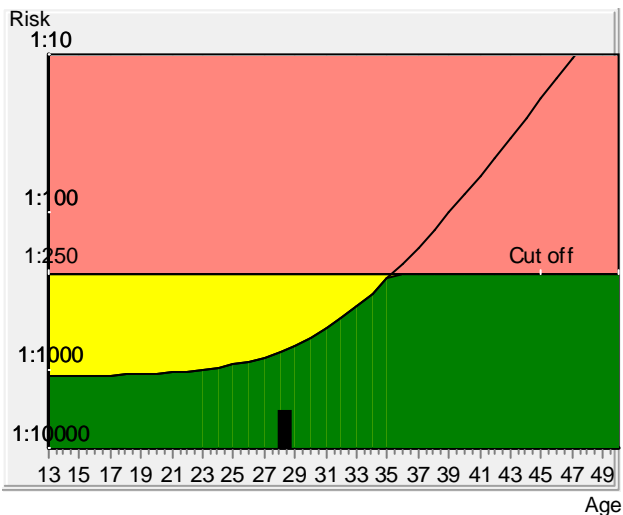


Prisca 5.2.0.13  
Date of report: 17/11/21

Patient data			
Name	RINA NASRIN		Patient ID
Birthday	13/08/93		Sample ID
Age at sample date	28.3		Sample Date
Gestational age	11 + 6		
Correction factors			
Fetuses	1	IVF	unknown
Weight	70	diabetes	no
Smoker	no	Origin	Asian
Previous trisomy 21		unknown	
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	2.7 mIU/mL	0.92	Method
fb-hCG	34.98 ng/mL	0.86	Scan date
Risks at sampling date			Crown rump length in mm
Age risk	1:758		Nuchal translucency MoM
Biochemical T21 risk	1:5397		Nasal bone
Combined trisomy 21 risk	1:2746		Sonographer
Trisomy 13/18 + NT	<1:10000		Qualifications in measuring NT
			DR.A.K.LAHIRI
			M.B.B.S
Trisomy 21			
<p><b>The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.</b></p> <p>After the result of the Trisomy 21 test (with NT) it is expected that among 2746 women with the same data, there is one woman with a trisomy 21 pregnancy and 2745 women with not affected pregnancies.</p> <p>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!</p> <p>The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)).</p> <p>The laboratory can not be hold responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!</p>			
Trisomy 13/18 + NT			
<p><b>The calculated risk for trisomy 13/18 (with nuchal translucency) is &lt; 1:10000, which represents a low risk.</b></p>			



Sign of Physician