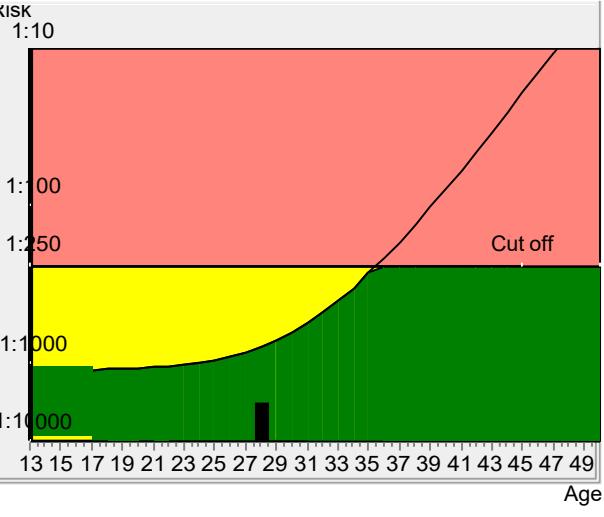


LAXMI RATNA

Patient data			
Name	Mrs. ASMA BEE	Patient ID	0012109280378
Birthday	23/09/93	Sample ID	21102978
Age at sample date	28.0	Sample Date	28/09/21
Gestational age	12 + 5		
Correction factors			
Fetuses	1	IVF	no
Weight	69	diabetes	no
Smoker	no	Origin	Asian
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	2.21 mIU/mL	0.61	Gestational age 12 + 0
fb-hCG	41.6 ng/mL	1.09	Method CRL Robinson
Risks at sampling date			
Age risk	1:800		Scan date 23/09/21
Biochemical T21 risk	1:1310		Crown rump length in mm 56.8
Combined trisomy 21 risk	1:3037		Nuchal translucency MoM 1.20
Trisomy 13/18 + NT	<1:10000		Nasal bone present
			Sonographer DR NA
			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.			
<p>After the result of the Trisomy 21 test (with NT) it is expected that among 3037 women with the same data, there is one woman with a trisomy 21 pregnancy and 3036 women with not affected pregnancies.</p> <p>The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician.</p> <p>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 held responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!</p>			
			
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