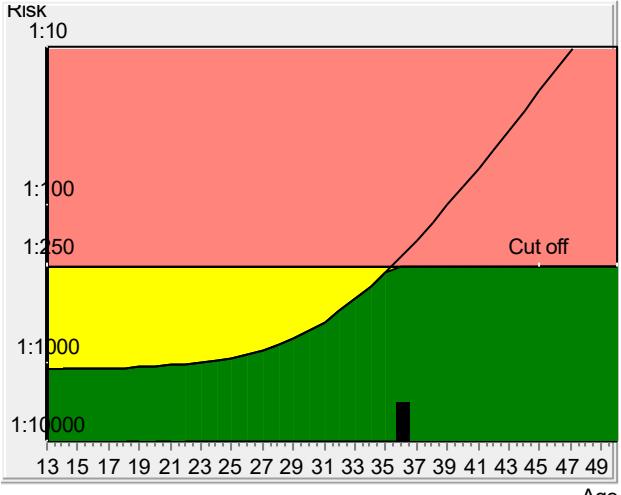


NA

Patient data			
Name	Mrs. S PARVEEN	Patient ID	0042508250009
Birthday	19/07/89	Sample ID	23168959
Age at sample date	36.1	Sample Date	25/08/25
Gestational age	12 + 0		
Correction factors			
Fetuses	1	IVF	no
Weight	54	diabetes	no
Smoker	no	Origin	Asian
Biochemical data			
Parameter	Value	Corr. MoM	Ultrasound data
PAPP-A	3.54 mIU/mL	0.90	Gestational age 11 + 6
fb-hCG	44.2 ng/ml	0.97	Method CRL Robinson
Risks at sampling date			
Age risk	1:208		Scan date 24/08/25
Biochemical T21 risk	1:1107		Crown rump length in mm 54.6
Combined trisomy 21 risk	1:5639		Nuchal translucency MoM 0.62
Trisomy 13/18 + NT	<1:10000		Nasal bone present
Sonographer 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 5639 women with the same data, there is one woman with a trisomy 21 pregnancy and 5638 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