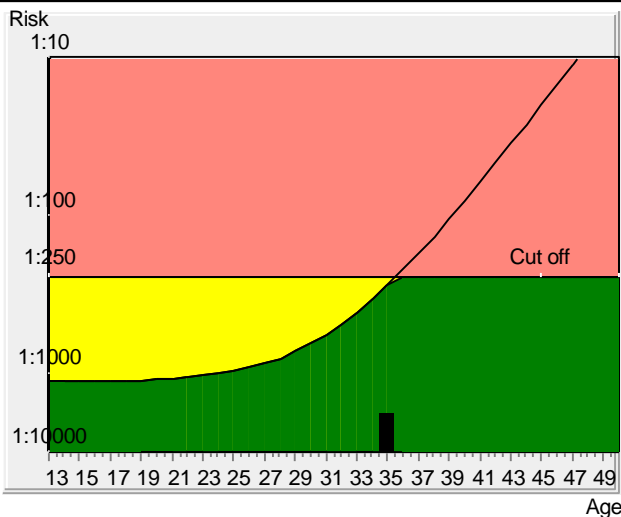


Patient data			
Name	MRS. SUNITA		Patient ID
Birthday	5/03/1990		Sample ID
Age at sample date	34.9		Sample Date
Gestational age	13 + 1		
Correction factors			
Fetuses	1	IVF	no
Weight	55.75	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	3.56 mIU/ml	0.63	Gestational age
fb-hCG	25.1 ng/ml	0.59	Method
Risks at sampling date			CRL Robinson
Age risk	1:285		Scan date
Biochemical T21 risk	1:1907		22/01/2025
Combined trisomy 21 risk	1:9053		Crown rump length in mm
Trisomy 13/18 + NT	<1:10000		68.6
			Nuchal translucency MoM
			0.92
			Nasal bone
			present
			Sonographer
			..
			Qualifications in measuring NT
			..
Trisomy 21			
<p>The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.</p> <p>After the result of the Trisomy 21 test (with NT) it is expected that among 9053 women with the same data, there is one woman with a trisomy 21 pregnancy and 9052 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>The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.</p>			



Sign of Physician