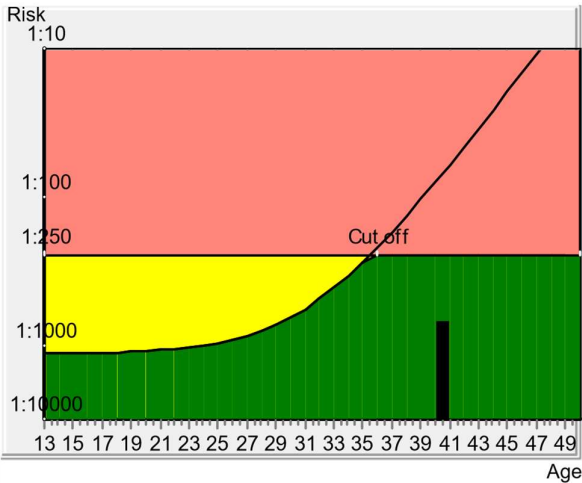


Patient data			
Name	MRS. SANTOSH KUMARI	Patient ID	
Birthday	14/12/1983	Sample ID	2405064900/NOD
Age at sample date	40.5	Sample Date	31/05/2024
Gestational age	12 + 6		
Correction factors			
Fetuses	1	IVF	no
Weight	59	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	2.38 mIU/ml	0.51	Gestational age 12 + 6
fb-hCG	37.5 ng/ml	0.86	Method CRL Robinson
Risks at sampling date			Scan date 31/05/2024
Age risk		1:71	Crown rump length in mm 66.9
Biochemical T21 risk		1:118	Nuchal translucency MoM 0.77
Combined trisomy 21 risk		1:688	Nasal bone present
Trisomy 13/18 + NT		<1:10000	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 688 women with the same data, there is one woman with a trisomy 21 pregnancy and 687 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

 below cut off	 Below Cut Off, but above Age Risk	 above cut off
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