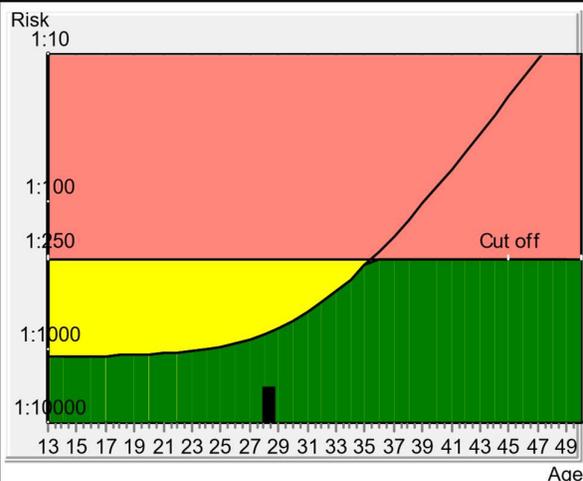


Patient data			
Name	MRS. PREETI (B)	Patient ID	2312036509/NOD (B)
Birthday	12/09/95	Sample ID	2312036509/NOD (B)
Age at sample date	28.3	Sample Date	22/12/23
Gestational age	12 + 5		
Correction factors			
Fetuses	2	IVF	no
Weight	54	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	8.92 mIU/ml	0.98	12 + 4
fb-hCG	40.9 ng/ml	0.42	Method
			CRL Robinson
			Scan date
			21/12/23
Risks at sampling date			Crown rump length in mm
Age risk		1:781	63
Biochemical T21 risk		<1:10000	Nuchal translucency MoM
Combined trisomy 21 risk		<1:10000	0.68
Trisomy 13/18 + NT		<1:10000	Nasal bone
			present
			Sonographer
			DR. PRIYANKA GHURE
			Qualifications in measuring NT
			M.D
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 more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy.</p> <p>The risk for this twin pregnancy has been calculated for a singleton pregnancy with corrected MoMs.</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 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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