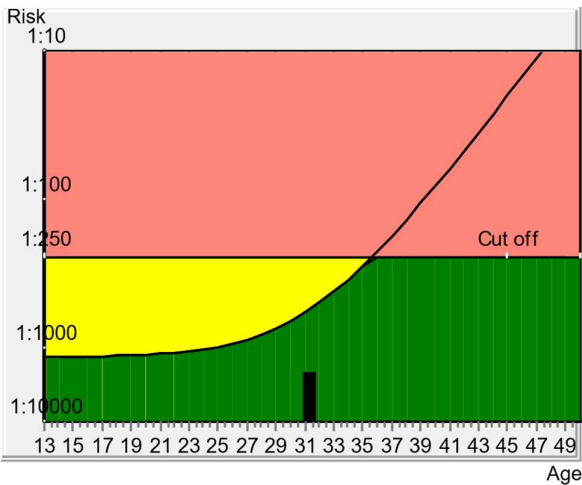


Patient data			
Name	MRS. PALLAVI	Patient ID	
Birthday	17/02/1993	Sample ID	2406019711/NOD
Age at sample date	31.3	Sample Date	11/06/2024
Gestational age	13 + 5		
Correction factors			
Fetuses	1	IVF	no
Weight	65	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	5.18 mIU/ml	0.89	Gestational age 12 + 5
fb-hCG	102 ng/ml	2.67	Method CRL Robinson
Risks at sampling date			Scan date 4/06/2024
Age risk		1:562	Crown rump length in mm 65.7
Biochemical T21 risk		1:249	Nuchal translucency MoM 0.72
Combined trisomy 21 risk		1:1461	Nasal bone present
Trisomy 13/18 + NT		<1:10000	Sonographer DR. JASWINDER SINGH
			Qualifications in measuring NT MD
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 1461 women with the same data, there is one woman with a trisomy 21 pregnancy and 1460 women with not affected pregnancies.</p> <p>The free beta HCG level is high.</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
--	--	--