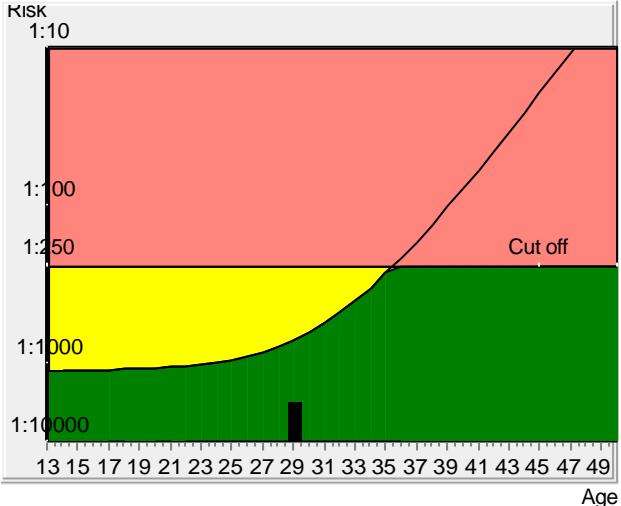


Patient data			
Name	Mrs. MADHU DEWANGAN	Patient ID	0622402130051
Birthday	25-01-1995	Sample ID	24134716
Age at sample date	29.1	Sample Date	13-02-2024
Gestational age	12 + 5		
Correction factors			
Fetuses	1	IVF	no
Weight	58	diabetes	no
Smoker	no	Origin	Asian
Biochemical data			
Parameter	Value	Corr. MoM	Ultrasound data
PAPP-A	12.14 mIU/mL	2.73	Gestational age 12 + 4
fb-hCG	35.66 ng/mL	0.88	Method CRL Robinson
Risks at sampling date			
Age risk	1:724		Scan date 12-02-2024
Biochemical T21 risk	<1:10000		Crown rump length in mm 64.1
Combined trisomy 21 risk	<1:10000		Nuchal translucency MoM 0.97
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 more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy.</p> <p>The PAPP-A level is high.</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