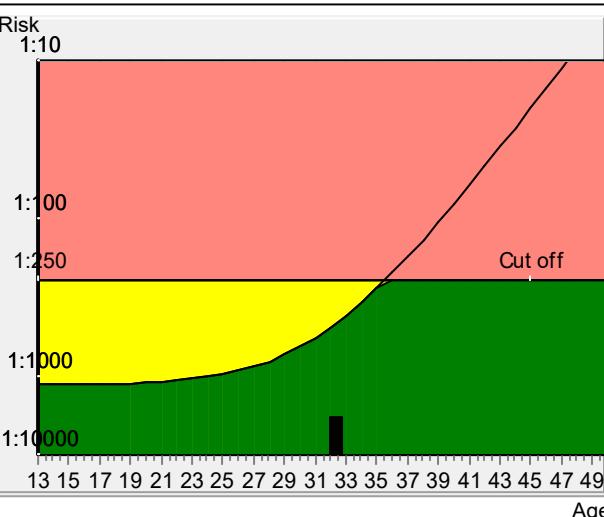


N A

Patient data			
Name	W/O ANSUL KUMAR	Patient ID	0692410230301
Birthday	13-07-1992	Sample ID	A1650085
Age at sample date	32.3	Sample Date	23-10-2024
Gestational age	13 + 1		
Correction factors			
Fetuses	1	IVF	no
Weight	77	diabetes	no
Smoker	no	Origin	Asian
Biochemical data			
Parameter	Value	Corr. MoM	Ultrasound data
PAPP-A	2.57 mIU/mL	0.69	Gestational age 13 + 1
fb-hCG	43.1 ng/mL	1.30	Method CRL Robinson
Risks at sampling date			
Age risk	1:473		Scan date 23-10-2024
Biochemical T21 risk	1:700		Crown rump length in mm 71
Combined trisomy 21 risk	1:3973		Nuchal translucency MoM 0.79
Trisomy 13/18 + NT	<1:10000		Nasal bone unknown
Sonographer N A			
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 3973 women with the same data, there is one woman with a trisomy 21 pregnancy and 3972 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 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