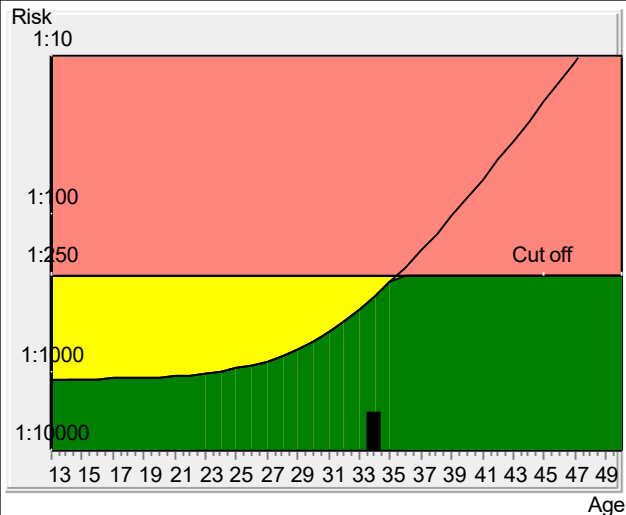


JITM Diagnostics

Patient data			
Name	MRS. MANISHA	Patient ID	
Birth day	5/01/1991	Sample ID	2412037930/NOD
Age at sample date	34.0	Sample Date	23/12/2024
Gestational age	12 + 4		
Correction factors			
Fetuses	1	IVF	no
Weight	70	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	3.76 mIU/ml	1.11	Gestational age 10 + 5
fb-hCG	34.1 ng/ml	0.80	Method CRL Robinson
Risks at sampling date			Scan date 10/12/2024
Age risk		1:338	Crown rump length in mm 41
Biochemical T21 risk		1:4255	Nuchal translucency MoM 0.95
Combined trisomy 21 risk		<1:10000	Nasal bone present
Trisomy 13/18 + NT		<1:10000	Sonographer DR. ANKUR KUMAR SINHA DMRD DNB
			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.
 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.
 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!
 The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)).
 The laboratory can not be hold responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!

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