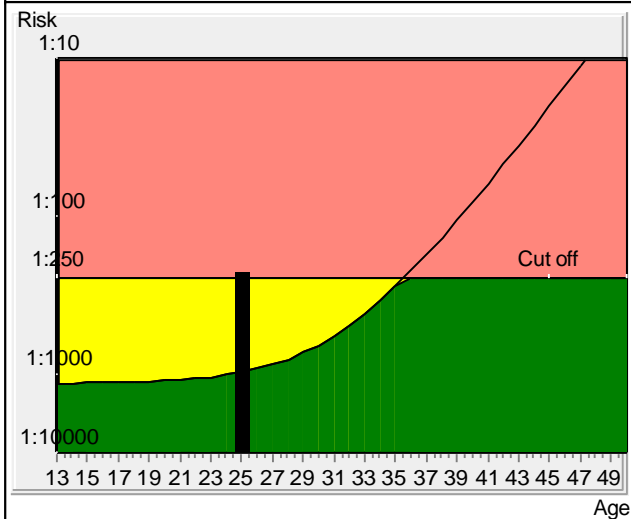


Patient data			
Name	MRS. KANCHAN KASTUR	Patient ID	
Birthday	5/06/1999	Sample ID	2406036585/NOD
Age at sample date	25.0	Sample Date	20/06/2024
Gestational age	13 + 2		

Correction factors			
Fetuses	1	IVF	no
Weight	38.4	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no

Biochemical data			Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age	11 + 5
PAPP-A	2.5 mIU/ml	0.28	Method	CRL Robinson
fb-hCG	112 ng/ml	2.30	Scan date	9/06/2024
Risks at sampling date			Crown rump length in mm	52.5
Age risk		1:984	Nuchal translucency MoM	0.57
Biochemical T21 risk		>1:50	Nasal bone	present
Combined trisomy 21 risk		1:229	Sonographer	DR. VIBHA BANSAL
Trisomy 13/18 + NT		<1:10000	Qualifications in measuring NT	M.D



Trisomy 21

The calculated risk for Trisomy 21 (with nuchal translucency) is above the cut off, which indicates an increased risk.

After the result of the Trisomy 21 test (with NT) it is expected that among 229 women with the same data, there is one woman with a trisomy 21 pregnancy and 228 women with not affected pregnancies.

The PAPP-A level is low.

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

 below cut off	 Below Cut Off, but above Age Risk	 above cut off
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