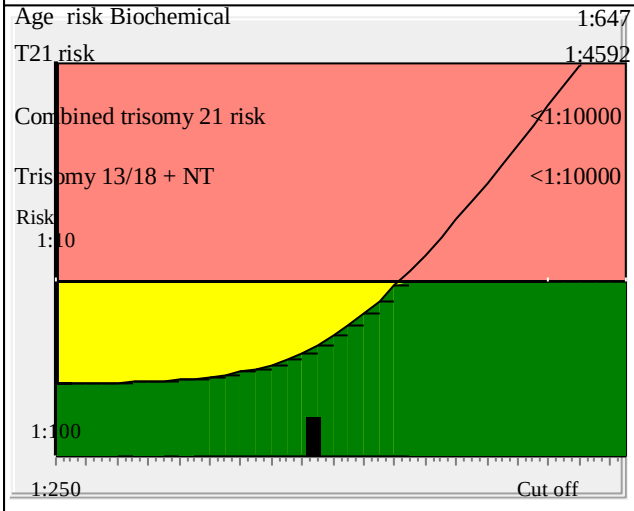
JITM DIAGNOSTICS

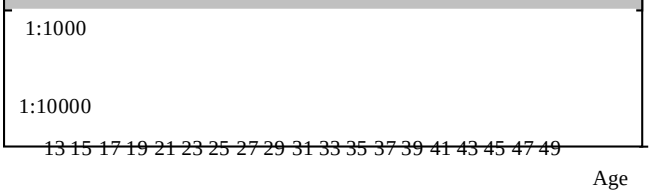
Date of report: 11/10/2023

Patient data			
Name	MRS. SAPNA	Patient ID	
Birthday	1/01/1994	Sample ID	2310018780/NOD
Age at sample date	29.8	Sample Date	10/10/2023
Gestational age	11 + 6		

Correction factors					
Fetuses	1	IVF	no	Previous trisomy 21	no
Weight	71	diabetes	no	pregnancies	
Smoker	no	Origin	Asian		

Biochemical data			Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age	11 + 3
PAPP-A	3.5 mIU/ml	1.46	Method	CRL Robinson
fb-hCG	60.1 ng/ml	1.31	Scan date	7/10/2023
Risks at sampling date			Crown rump length in mm	49.3
Age risk Biochemical		1:647	Nuchal translucency MoM	0.82
T21 risk		1:4592	Nasal bone	present
Combined trisomy 21 risk		<1:10000	Sonographer	DR. B.K. TYAGI
Trisomy 13/18 + NT		<1:10000	Qualifications in measuring NT	M.D





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 thesame data, there is one woman with a trisomy 21 pregnancy.
The calculated risk by PRISCA depends on the accuracyof the information provided by the referring physician.
Please note that risk calculations are statisticalapproaches and have no diagnostic value!
The patient combined risk presumes the NT measurementwas done according to accepted guidelines (Prenat Diagn18: 511-523 (1998)).
The laboratory can not be hold responsible for their impacton 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