JITM SKILLS PVT.LTD
D-87, SECTOR-02,NOIDA 201301

Prisca 5.1.0.17

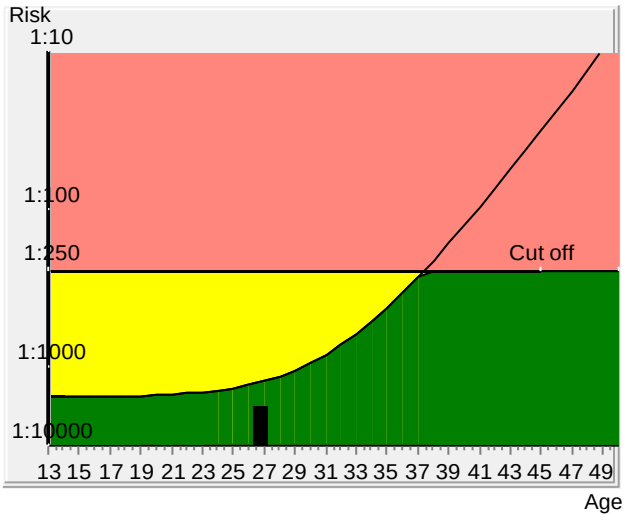
Date of report: 14-03-2023

Patient data			
Name	MRS. KAJAL DEVI	Patient ID	4248293
Birthday	01-01-1997	Sample ID	4248293
Age at delivery	26.7	Sample Date	13-03-2023
Gestational age	12 + 5		

Correction factors					
Fetuses	1	IVF	no	Previous trisomy 21	unknown
Weight	56	diabetes	no	pregnancies	

Biochemical data			Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age	12 + 2
PAPP-A	3 mIU/ml	0.74	Method	CRL Robinson
fb-hCG	33 ng/ml	0.94	Scan date	10-03-2023
			Crown rump length in mm	60.6

Risks at term		Ultrasound data	
Age risk	1:1270	Nuchal translucency MoM	0.95
Biochemical T21 risk	1:4601	Nasal bone	present
Combined trisomy 21 risk	<1:10000	Sonographer	NA
Trisomy 13/18 + NT	<1:10000	Qualifications in measuring NT	FMF



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

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