

JITM DIAGNOSTICS PVT LTD

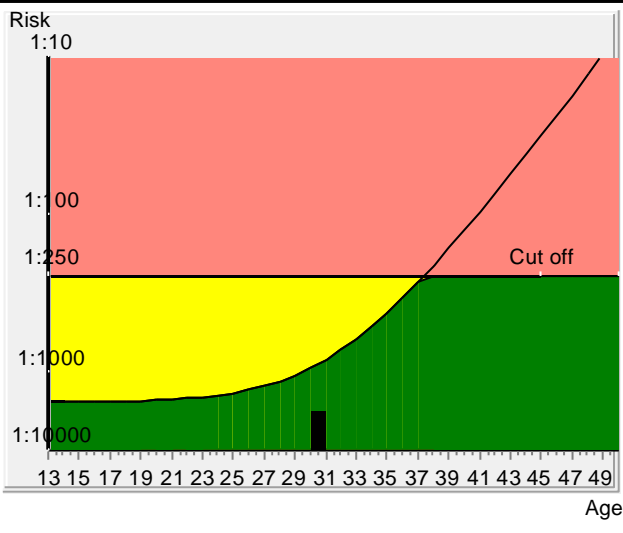
D-87 , Sector – 2 , NOIDA (UP) 201301

Prisca 5.1.0.17
Date of report: 08/01/23

Patient data			
Name	Mrs.POOJA RANI	Patient ID	4053733
Birthday	03/01/93	Sample ID	4053733
Age at delivery	30.5	Sample Date	08/01/23
Gestational age	12 + 3		

Correction factors			
Fetuses	1	IVF	unknown
Weight	55	diabetes	unknown
Smoker	unknown	Origin	Asian
		Previous trisomy 21 pregnancies	unknown

Biochemical data			Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age	12 + 1
PAPP-A	8.17 mIU/ml	2.21	Method	CRL Robinson
fb-hCG	11.9 ng/ml	0.32	Scan date	06/01/23
Risks at term			Crown rump length in mm	58
Age risk		1:905	Nuchal translucency MoM	0.66
Biochemical T21 risk		<1:10000	Nasal bone	unknown
Combined trisomy 21 risk		<1:10000	Sonographer	Dr.abhijit vipul Dr.abhijit vi
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 free beta HCG 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

