

Date of Report 08-11-2023
PRISCA 5.2.0.13

Patient Data			
Name	MRS MANI DIXIT	Patient ID	012311070136
Birthday	27-09-1986	Sample ID	11783184
Age at Sample date	37.1	Sample Date	07-11-2023
Gestational age	12+5		

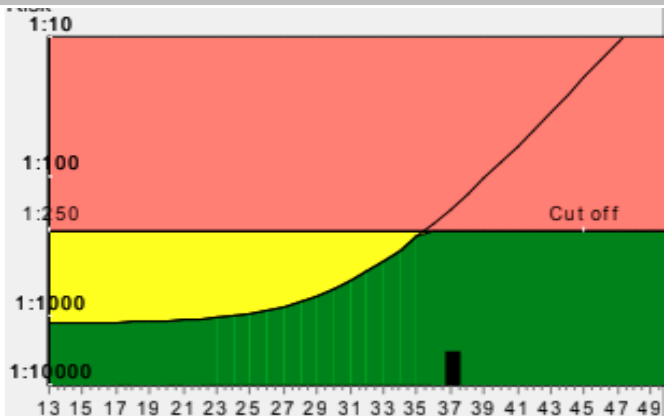
Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	73	Diabetes	NO	Pregnancies	unknown
Smoker	NO	Origin	Asian		

Biochemical Data			Ultrasound Data	
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Parameter	Value	Corr Mom		
PAPP-A	3.87 mIU/ml	0.84	Gestational age	12+4
fb-hCG	30.7 ng/ml	0.92	Method	CRL (<>Robinson)
			Scan date	06-11-2023

Risks at sampling date			Ultrasound Data	
Age Risk		1:168	Crown rump length in mm	62.1
Biochemical T21 risk		1:856	Nuchal translucency MoM	0.81
Combined trisomy 21 risk		1:4409	Nasal bone	PRESENT
Trisomy 13/18 + NT		<1:10000	Sonographer	DR INDRAJEET
			Qualifications in measuring NT	MD

Risk	Down's Syndrome Risk (Trisomy 21 Screening)
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The calculated risk for Trisomy 21 (with NT) is below the cut off, which represents a low risk.

After the result of the Trisomy 21 test (with NT) it is expected that among 4409 women with the same data, there is one woman with a trisomy 21 pregnancy and 4408 women with not affected pregnancies.

The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician. Please note that the risk calculations are statistical approaches and have no diagnostic value!

The patient combined risk presumes that NT measurement was done according to accepted guidelines (Prenat Diagn 18:511-523; 1998).

The calculated risk for Trisomy 13/18 (with NT) is <1:10000, which indicates a low risk

The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values

Risk Above Cut Off
 Risk above Age Risk
 Risk below Age risk